
Genetic Management of Wild and Translocated Black Rhinoceros in South Africa's KwaZulu-Natal Region

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To my parents, Louis and Linda Anderson

Biology is a science of three dimensions. The first is the study of each species across all levels of biological organization, molecule to cell to organism to population to ecosystem. The second dimension is the diversity of all species in the biosphere. The third dimension is the history of each species in turn, comprising both its genetic evolution and the environmental change that drove the evolution. Biology, by growing in all three dimensions, is progressing toward unification and will continue to do so.

-Edward O. Wilson



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ABSTRACT

The African black rhinoceros (*Diceros bicornis*) is critically endangered. Like other megafauna, the species is managed in parks and is often translocated to expand their range into reserves where they have been extirpated. Management of genetic variation has been identified as an important consideration in long-term management plans for many wild and captive endangered species including black rhino. In this thesis I examined the contemporary levels of genetic variation within the black rhinoceros (*D. b. minor*) in KwaZulu-Natal (KZN), South Africa, and specifically the relict source population at Hluhluwe-iMfolozi Game Reserve (HiP), and how this information can be incorporated into management decisions to improve the long-term viability and persistence of the population. Previous studies have examined levels of genetic variation and differentiation among the three black rhino subspecies (*D. b. minor*, *D. b. michaeli* and *D. b. bicornis*) in an attempt to resolve their taxonomy and to establish baseline genetic assessments for managing populations. However, there has been a lack of genetic information based on the variable mitochondrial DNA (mtDNA) control region of the KZN metapopulation and a direct comparison of microsatellite variability between the *D. b. minor* populations of KZN and Zimbabwe.

The specific objectives of this study were to: (1) determine the DNA sequence of the mtDNA control region of three subspecies and estimate the level of variation within the HiP source and KZN metapopulation and compare the results with *D. b. minor* outside KZN and the other two subspecies; (2) use ten microsatellite DNA markers to estimate the levels of heterozygosity and

allelic diversity in the HiP source and KZN metapopulation and compare results to previously published microsatellite data (specifically native Zimbabwe *D. b. minor*; and (3) use VORTEX Population Viability Analysis (PVA) and HiP vital rates to model the effects of increasing population size and supplementation, and investigate what management scenarios would be most effective for minimizing the loss of genetic variation caused by genetic drift with HiP.

MtDNA showed evidence of a bottleneck in the KZN *D. b. minor* metapopulation. The KZN metapopulation were fixed for a single haplotype (n=65), compared to six haplotypes (n=11) in native Zimbabwe *D. b. minor*. *D. b. michaeli* (n = 21) samples had 13 haplotypes, while the *D. b. bicornis* (n = 4) samples had one. Additionally, a haplotype network showed a discernable pattern of separation amongst the three subspecies with the KZN population positioned with the *D. b. minor* populations of Zimbabwe. While it was expected that the KZN *D. b. minor* would cluster together with the Zimbabwe *D. b. minor* because they are the same subspecies, the haplotype network provides further supporting evidence of a bottleneck in the KZN metapopulation.

The microsatellite DNA results from the KZN metapopulation also indicated a likely bottleneck pattern and possible inbreeding. The KZN metapopulation was out of Hardy-Weinberg Equilibrium, monomorphic at one locus, showed excess of homozygosity at five out of ten microsatellite DNA loci, and had 28% less genetic variation at microsatellite DNA loci and lower allele frequency than the native Zimbabwe *D. b. minor*. Modified M-ratio results indicated that all three of the subspecies had been through a bottleneck. There is no pre-decline genetic information available for the KZN metapopulation, so it

is unclear if the KZN region has historically always had low genetic variation or if the low levels were caused by a recent population decline.

No translocations are made into HiP. Modelling results of a simulated *D. b. minor* source population indicated that if no translocations into the reserve are made, expected heterozygosity (H_E) would decrease ~25% over ~100 black rhino generations (BRGs). Increasing the size of the modelled population slowed the rate of loss with the mean H_E decreasing by ~10% over ~100 BRGs. Models of supplementations made with a pair of black rhino (one female and one male) from the KZN metapopulation made every ten gestational years, maintained the mean H_E ($H_E = 0.45$) of the population over ~100 BRGs, but increased ~30% when supplemented with individuals from Zimbabwe. PVA results indicated that increasing gene flow through supplementation is effective and does not require a large number of individuals or need to be frequent.

If KZN has always had low levels of genetic variability, then based on the model findings, serial translocations made with KZN metapopulation rhinos into the HiP source population would be recommended to slow the rate of loss caused by drift and to maintain current levels of genetic variation. If, however, the low levels of variation were caused by a recent decline in population size, then according to the model, supplementations with native Zimbabwe *D. b. minor* would not only decrease the rate of loss of genetic variation, but would increase levels of genetic variation.

This research highlights the importance of shifting focus from increasing the number of individuals within a population to that of quality (e.g. levels of genetic variation) as black rhino move into recovery. Techniques like serial translocation and supplementation can help maintain current levels of genetic

diversity and prevent further loss of variation in the KZN source and metapopulation, which will enable managers to improve long-term African black rhinoceros conservation efforts. These techniques can also be integrated into active management schemes for other large conservation-reliant species. More specifically, those in small, remnant populations with limited reserve or range sizes in order to increase long-term survival and population persistence.

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CHAPTER ONE

Introduction



Photo by Rosalynn Anderson-Lederer

Introduction

The evolutionary process is dependent on genetic variation in order for populations to adapt to changes occurring in the environment (e.g. diseases, predators, climate change) (Lacy 1987b). Large breeding populations are able to maintain high levels of variation and evolutionary potential (Lacy 1987b; Bijlsma et al. 2000), but small populations are prone to decreases in the level of genetic variation and inbreeding depression. This increases the risk of extinction and reduces their evolutionary potential (Lacy 1987b; Frankham et al. 1999; Garner et al. 2005; Frankham 2005). It is unclear which, or how many, traits will be affected by inbreeding depression, or how long before adverse genetic effects manifest in small populations (Hogg et al. 2006). However, when genetic diversity is lost in small populations, the rate at which a population recovers is primarily contingent on mutation rate (Lynch 1996), which can take a considerable number of generations (Allendorf and Leary 1986a).

Many African wild animal populations have gone through gradual or sudden population declines for reasons that include overexploitation, habitat loss and disease (e.g. rinderpest) (Western and Vigne 1985; Simonsen et al. 1998; Harley et al. 2005). Historic population sizes and declines, however, are seldom well documented. Europeans recorded wildlife sightings while exploring and hunting the continent, but documentation was general with basic descriptions, usually kept in personal journals (Tingley and Beissinger 2009). These records are valuable for estimating historic ranges, but their use is limited when estimating historic population sizes and declines (Shoo et al. 2006; Rookmaaker 2007; Tingley and Beissinger 2009). It was often only in extreme

cases of species decline that detailed population information was recorded and is available (Shaffer et al. 1998).

Genetic Variation in Post-Decline Populations

Many threatened or endangered species that have experienced population declines are now conservation-reliant (see: Miller et al. 1988; Walters 1991; Tyus and Saunders 2000; Jamieson et al. 2006; Johnson et al. 2010). The level of genetic variation remaining after a severe population decline is determined by past and current population sizes, and pre-decline levels of diversity (Frankham et al. 2002). An ongoing loss of diversity is expected if population recovery is slow (Nei et al. 1975).

To determine loss of genetic variation as a consequence of decline, an assessment of pre-decline gene flow and variability is required, but seldom possible (Briskie and Mackintosh 2004). Museum samples can be used to apprise missing data (greater prairie chicken (*Tympanuchus cupido*) Bouzat et al. 1998; whooping crane (*Grus americana*) Glen et al. 1999), but when post-decline levels of genetic variation are low and no museum specimens are available it may be difficult to determine whether the low variability was due to the reduction in population size or a general response to demographic and environmental differences (Bouzat et al. 1998).

If pre-decline levels of genetic variation are unknown, post-decline population levels should still be sampled and monitored. The collected data will establish baseline levels of variation, assisting in future reintroductions and parentage analyses in founder populations (Schwartz et al. 2006).

1.2 Study Species

The Rhinocerotidae (meaning “nose horns”) are the second largest living land animals after elephants (Kingdon 1997). They are in the order Perissodactyla (odd-toed ungulates), which includes the Tapiridae (Tapirs) and Equidae (Horses) (Silberman and Fulton 1979; Lacombat 2005). First appearing approximately 56 to 34 million years ago (MYA), Rhinocerotidae included some 26 different genera in Eurasia spreading to North America and later, in the Miocene Epoch (approximately 23 to 5 MYA), from Asia into Africa (Lacombat 2005). The earliest genera of Rhinocerotidae went extinct during early Oligocene (37 MYA), and Rhinocerotidae declines at the end of the Miocene are attributed to climate change (Lacombat 2005). Today there are five extant rhinoceros species: three Asian; Sumatran (*Dicerorhinus sumatrensis*; critically endangered A2abd; C1+2a(i) ver 3.1 (van Strien et al. 2008a)), Javan (*Rhinoceros sondaicus*; critically endangered C2a(i); D ver 3.1 (van Strien et al. 2008b)) and Indian (*Rhinoceros unicornis*; vulnerable B1ab(iii) (Talukdar et al. 2008)) and two African; black (*Diceros bicornis*) and white (*Ceratotherium simum*; near threatened ver 3.1 (Emslie 2012)) (Owen-Smith 1988).

The Black Rhinoceros (Diceros bicornis)

Unlike their Asian counterparts, the African black rhino lack incisors and canine teeth (Kingdon 1997; Emslie and Brooks 1999). Their brachyodont teeth (low crown) enable them to browse on coarse plant material like leaves, twigs, branches and long grass (Kingdon 1997; Lacombat 2005). They have two horns which have a dermal origin and are comprised of laminated keratinaceous filaments (compressed hair and fingernail matter) that grow throughout the

animal's life (Silberman and Fulton 1979; Lacombat 2005). Black rhino have a muscular finger-like prehensile lip used for browsing (Skinner and Smithers 1990). They range in height from 1.4 – 1.8 m (55 – 71 in.) at the shoulder and vary in weight from 1000 - 1800 kg (2200 – 3970 lbs); their body length ranges from 2.9 – 3.75 m (114 – 148 in.) (Kingdon 1997; Emslie and Brooks 1999).

Black rhino are not actually black, but generally grey; they may appear to vary in colour as a result of mud or dust bathing (Kingdon 1997). Wallowing may help reduce body temperature and protects rhino from ectoparasites (e.g. ticks and biting flies). Distinguishing the sex of a rhinoceros in the field can be difficult because males have undescended testes and, therefore, lack a scrotum (Kingdon 1997). The genitalia of both sexes face backwards and they are capable of projecting urine up to three to four meters (Schenkel and Schenkel-Hulliger 1969). Black rhino tend to be asocial and while female home ranges overlap, males tend to live in mutually exclusive home ranges (Owen-Smith 1988; Conway and Goodman 1989). Both sexes can live up to 30 – 35 years in the wild, but that is extended to 45+ years in captivity (Owen-Smith 1988). One black rhino generation is approximately 14 years (Brooks and Adcock 1997).

Gestation is approximately 15.33 months (Schenkel and Schenkel-Hulliger 1969; Owen-Smith 1988; Bertschinger 1994) with infants weighing between 27 – 45 kg (60 – 100 lbs) at birth. Females give birth to a single offspring that is able to stand and walk shortly after birth. They start to suckle within 3 - 4 hours of birth and are able start eating solid food (grass and non-woody plants) within 10 days (Schenkel and Schenkel-Hulliger 1969; Owen-Smith 1988; Bertschinger 1994). The mean intercalving time was shown to be between 30 and 44 months within HiP (Bertschinger 1994).

Black Rhino Taxonomy

Before wildlife managers can design effective conservation plans they must first resolve and understand the target species' taxonomy. Identification inaccuracies can lead to inadequate protection for some species resulting in extinction and/or the possibility of unwanted hybridization (e.g. black wildebeest (*Connochaetes gnou*) Grobler et al. 2011) of sister taxon when translocating animals for supplementation or reintroductions (Allendorf and Luikart 2007).

Taxonomic resolution includes two processes: (1) listing and priority setting involving legislation; and (2) recovery planning and *in situ* and *ex situ* conservation actions (Mace 2004). Once a species has been listed, conservation managers can turn their attention to figuring out why the species is in decline and implementing strategies for arresting and mitigating the effects of the decline (Mace 2004).

Zukowsky (1965) described 17 separate black rhino subspecies based on photos, literature, museum skull specimens and zoo animals. Groves' (1967) study of a smaller number of skulls narrowed the number of subspecies down to seven, based on size and morphology. However, subsequent but unpublished data collected on southern Africa black rhino skulls claim to refute Groves' findings (du Toit 1987). Groves and Grubb (2011) have increased the subspecies number up to eight to include *Diceros bicornis bicornis* (Linnaeus, 1758; from the Cape north to Kuruman, South Africa to southern Namibia); *D. b. chobiensis*, (Zukowsky 1965; the Okavango region of Botswana) *D. b. minor* (Drummond, 1976; KZN, South African north to north-west Tanzania and the south-west borders of Kenya); *D. b. occidentalis* (Zukowsky, 1922; northern Namibia and southern Angola); *D. b. michaeli* (Zukowsky 1965; north-west

Tanzania into eastern Kenya); *D. b. brucii* (Lesson, 1842; Somalia, western Somalia and northern Sudan); *D. b. ladoensis* (Groves 1967; Kenya Rift Valley north-west into southern Sudan); and *D. b. longipes* (Zukowsky 1949; south-western Chad, northern Cameroon and north-east Nigeria).

Black Rhino “Ecotypes”

In the context of a poorly resolved black rhino taxonomy, a meeting of rhino managers in 1986 proposed that for better metapopulation management, formal recognition of vernacular “ecotypes” be accepted and applied (Rookmaaker 1995, 2005, 2011). In 1990, taxonomic accuracy was set aside for pragmatic reasons and ecotype designations were officially recognized (Rookmaaker 2005; du Toit 2006a). Even though more than one subspecies occupied each ecotype region before the decline in black rhino numbers, in 1990 there was only one subspecies remaining in each of the regions. The three remaining recognized ecotypes and the corresponding subspecific names of subspecies remaining within the ecotypes (Eastern (*D. b. michaeli*): Ethiopia, Kenya, Rwanda, Somalia, Sudan, Tanzania, Uganda; South-western (*D. b. bicornis*): Angola, Botswana, Namibia, South Africa; and South-central (*D. b. minor*): Angola, Botswana, Congo Malawi, Mozambique, South Africa, Swaziland, Tanzania, Zambia, Zimbabwe) are now used interchangeably.

The critically endangered A2abcd ver 3.1 (Emslie 2011) *D. b. michaeli* (Eastern) is mostly found in Kenya, although small numbers have been accounted for in Rwanda and Tanzania. There is an extralimital or ‘insurance’ population in South Africa that was founded with individuals from Kenya. This subspecies is, however, probably nationally extinct in Ethiopia (Amin et al.

2006; Emslie and Brooks 1999). The vulnerable D1 ver 3.1 (Emslie 2011) *D. b. bicornis* (South-western) are found in Namibia and parts of South Africa, and are presumed nationally extinct in Angola and Botswana (Emslie and Brooks 1999). The critically endangered *D. b. minor* A2abcd ver 3.1 (Emslie 2011) (South-central) are mainly found in South Africa with smaller numbers in Zimbabwe (native), Tanzania (native), Swaziland (reintroduced), Malawi (reintroduced), Zambia (reintroduced), and Botswana (reintroduced). They are believed to be extinct in Angola and Mozambique (Amin et al. 2006). The last sighting of five remaining members of a fourth subspecies *D. b. longipes* (Western 'ecotype') was in 2001 in Northern Cameroon. This ecotype was declared 'probably extinct' in 2006 and subsequently declared 'extinct' in 2011 (Emslie 2011). While each of the ecotypes occupy different regions in Africa, there are no impervious geographical boundaries between them (Emslie and Brooks 1999). With the three remaining subspecies already red-listed, managers are addressing the second taxonomic process of how best to manage each subspecies.

Species Decline

Black rhino once numbered in the hundreds of thousands but they have suffered an extraordinary decline in the last century, disappearing more quickly than any other large mammal (Hitchins 1975; Western and Vigne 1985; Ashley et al. 1990; Swart et al. 1994; Harley et al. 2005). Human hunting is the primary reason for their decline (Western and Vigne 1985; Emslie and Brooks 1999; Harley et al. 2005). Black rhino were (and continue to be) killed for trophies,

meat, hides (for shields and good luck charms) and for their horns (traditional medicines and handles for daggers).

Prior to 2004, it was assumed that historic black rhino distribution extended from the Cape of South Africa, north to Somalia and Ethiopia, west to Senegal and Guinea (Figure 1.1). Rookmaaker (2004) revised the extent of

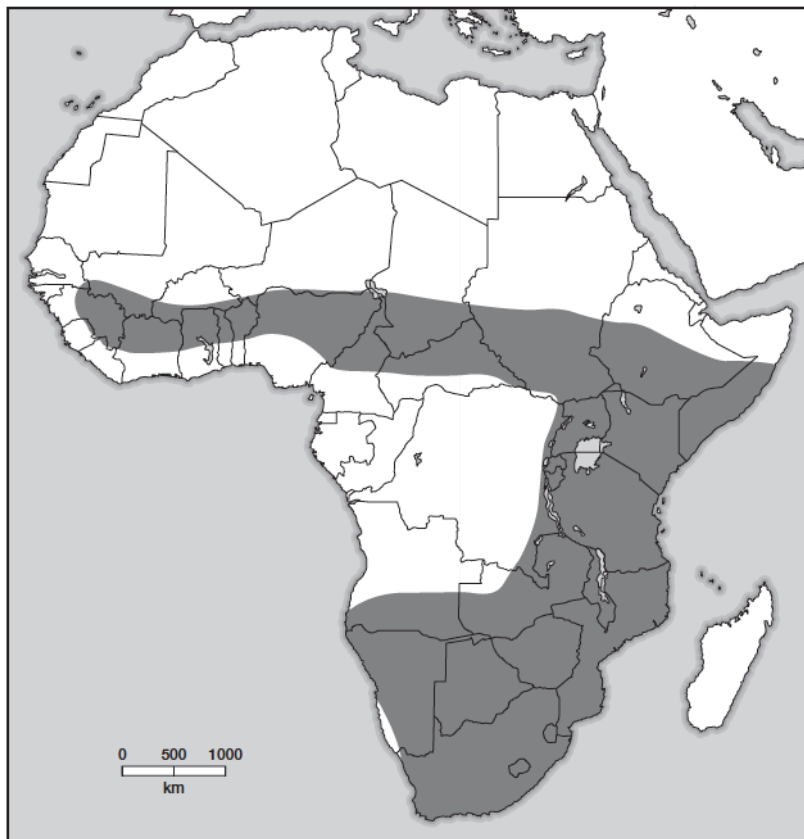


Figure 1.1: Previous 'pre-1900' distribution map of African black rhino. (Map from Emslie and Brooks 1999)

historic distribution in West Africa (only to as far as Nigeria, possibly the southwest of Niger) based on a study of bibliographical and iconographic literature (Figure 1.2). Revising the extent of the black rhino historic range to the west may help some aspects of taxonomic resolution.

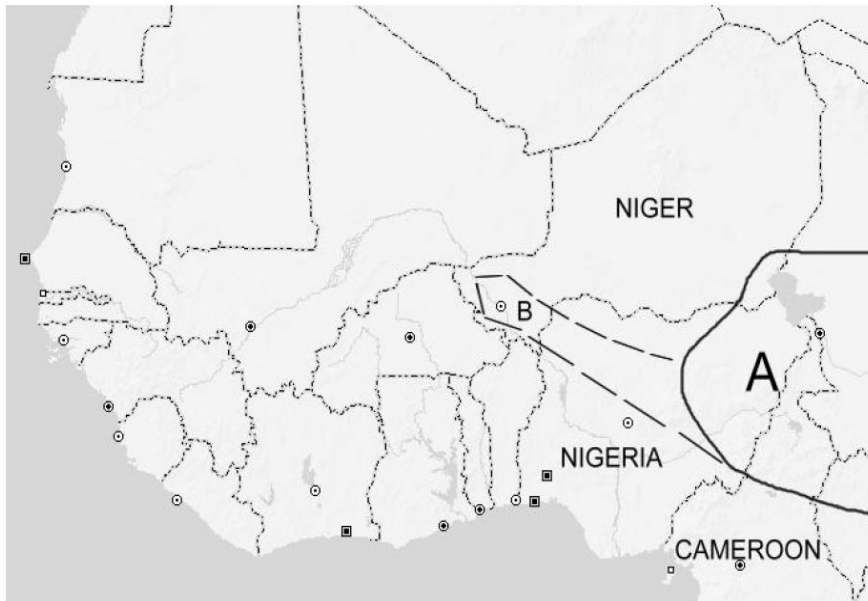


Figure 1.2: Revised 'pre-1900' distribution map of black rhino in West Africa. Area A is the likely extent of the black rhino to the west. Area B is the maximum possible extent of the range. (Map from Rookmaaker, 2004)



Figure 1.3: Black rhino game reserves in KZN. Ndumo Game Reserve, Tembe Elephant Reserve, Pongola Game Reserve, Ithala Game Reserve, Mkuze Game Reserve, Thanda Game Reserve, Phinda Resource Reserve, Zululand Rhino Reserve, Zululand Rhino Reserve, Ubizane Wildlife Reserve, eMakhosini Heritage Park, HiP, Eastern Shores, Weenan Game Reserve

From 1970 to 1995, black rhino numbers were reduced from approximately 65,000 to an estimated 2,400 (Emslie et al. 2009). However, by the end of 2010, *in situ* conservation methods increased the total number of black rhino to 4,880 (Emslie 2011). The number of *D. b. minor* in South Africa increased from a mere 110 reported in 1930 to 1,684 by the end of 2010 (Emslie 2011). While the number of *D. b. minor* in South Africa appears to be increasing, a recent spike in poaching has decreased the number of *D. b. minor* in Zimbabwe to approximately 431 (Emslie 2011).

D. b. minor were extirpated from Kruger National Park (KNP), South Africa in the 1930's, but have been reintroduced through translocations from South Africa and Zimbabwe. Ferreira et al. (2011) estimated that there were, approximately 627 *D. b. minor* in KNP, which would make it the largest *D. b. minor* population in Africa. Their results were based on block surveys consisting of 155 individual black rhino visual encounters (Ferreira et al. 2011). Information gained by estimating population sizes is valuable and is often required to justify the implementation of management schemes (Tacha et al. 1982), but as management budgets allow, more precise methods of determining population size should be utilized.

Hluhluwe-iMfolozi Game Park (HiP), in the KwaZulu-Natal Region of South Africa used estimates for management purposes from 1998 to 2008 only to discover that their estimates were inflated by nearly 50% (Clinning et al. 2009). To improve accuracy of *D. b. minor* numbers, HiP currently employs a Priority Species Monitor (specialized ranger), who spends up to three months in each of its five sections visually locating as many black rhino as possible. Recent monitoring put the number of *D. b. minor* within HiP at roughly 220 (Clinning et

al. 2009) or 13% of the total *D. b. minor* in South Africa and approximately half of the *D. b. minor* in KwaZulu-Natal (Emslie 2011). HiP is the focal site for this project (Figure 1.3).

1.3 Hluhluwe-iMfolozi Game Park (HiP), Source Population

HiP is home to the largest relict population of the critically endangered *D. b. minor*. The park covers ~96,000 hectares (ha) and consists of two reserves, Hluhluwe and iMfolozi, which were previously managed separately but are now managed as a single game park with a connecting corridor between the two reserves. HiP is further broken down into five administration sections: Makhamisa (Wilderness-southernmost), Mbhuzane (South-west), Masinda (south-central), Nqumeni (north-central) and Manzibomvu (north). Field rangers regularly patrol each section and a game-proof fence surrounds the entire park. Located in the KwaZulu-Natal (KZN) region, it was one of the first game parks established in South Africa. Before becoming a game park in 1897, HiP was used as a hunting ground for King Shaka kaSenzangakhona, founder of the Zulu Nation (Brooks 2000). The park is also home to many other extant African wildlife species, including all of the “Big Five” (buffalo, elephant, lion, leopard and rhino) and has been a major source population for *D. b. minor* since the early 1960’s (Table 1.1).

Along with its historical significance, HiP is instrumental in the conservation of important southern African species. Southern white rhino (*Ceratotherium simum simum*), once found in large numbers across southern Africa, had been hunted to near extinction (Emslie and Brooks 1999). By the end of the 19th century ~200 individuals remained in South Africa; most were found

in the iMfolozi section of HiP (Rookmaaker 2000). Protection was set up for them and through *ex situ* breeding programmes their numbers grew. As of December 2010, there were ~20,170 white rhino in the wild (Emslie 2011).

Table 1.1: Translocation history of D. b. minor from HiP from 1962 - 2008. (Adapted from Hitchens 1984, Hall-Martin & Knight 1994 and Brooks & Adcock 1997)

Game Park	Year(s)	Number
Eastern Shores	1984	7
Elandsdrift	1994	5
Great Fish River	1989-1991	10
Ithala Game Reserve	1977-1984	17
Kruger National Park	1971-1982	55
Lapalala Game Reserve	1990-1993	11
Marakele National Park	1993-2002	2
Maremani Nature Reserve	2003	5
Mauricedale Game Ranch	2007	1
Malilangwe Wildlife Reserve	1998	27
Ndumo Game Reserve	1962-2008	18
Phinda Game Reserve	2004-2008	9
Phongola Game Reserve	2006	5
Pilanesberg Game Reserve	1981-1983	18
Rhinolands	1995	6
Sable Game Reserve	1991	4
Shamwari Game Reserve	1993	4
Somkhanda Game Reserve	2007	8
Sutton Game Reserve	2006	3
Tembe National Elephant Park	1994	13
Weenen Game Reserve	1983-2007	3
Western Shores	2003	4
Zulu Nyala Game Reserve	2004	1
Zululand Game Reserve	2005	18

1.4 Genetics

Conservation Genetics

Anthropogenic activities are the primary reason for species decline and extinction (e.g. habitat destruction and overexploitation) (Caughley 1994; Allendorf and Luikart 2007). The current levels of extirpations and the

widespread decline in the number of taxon (e.g. bear (*Ursus arctos*), bighorn sheep (*Ovis canadensis*)), are prompting conservation managers to incorporate reintroductions into species recovery plans (e.g. giant tortoises (*Geochelone nigra hoodensis*) and cactus (*Opuntia megasperma* var. *megasperma*) Gibbs et al. 2008; tuatara (*Sphenodon guntheri*) Nelson et al. 2002; Przewalski horse (*Equus ferus przewalskii*) Van Dierendonck and Wallis de Vries 1996). If the remaining number of individuals in a population of an endangered species is extremely small, genetic considerations may be overlooked for pragmatic reasons (e.g. necessity to increase the number of individuals). However, if populations are in recovery, genetic diversity should not be neglected. If a population is small enough to be conservation-reliant, it is likely to be prone to loss of genetic diversity and inbreeding, which may affect evolutionary potential (Frankham et al. 1999; Frankham 2005), meaning that a loss in the level of genetic variation might limit a population's suitability or adaptability to its environment (Soulé 1980; Allendorf and Leary 1986b). Management of black rhino has mainly focused on protecting existing populations, creating new populations through the means of reintroductions, translocations and supplementations, and through captive breeding programmes (Emslie et al. 2007).

Black rhino genetic studies have examined levels of genetic variation and differentiation among and between the subspecies (Ashley et al. 1990; Swart et al. 1994; Harley et al. 2005), but the definition of "acceptable levels" of genetic variation differ between studies. As researchers, we need to provide quantitative evidence to wildlife managers in a way that helps them in the field to secure the evolutionary potential of the black rhinoceros (e.g. identifying individuals that are ideal translocation candidates). Despite work in the field of

black rhino genetics over the past two decades, field managers only have a general idea of how their subspecific populations differ genetically from other populations.

1.4.1 Black Rhinoceros Genetics

As the black rhino species recovery progresses, it is now necessary for metapopulation management to shift in emphasis from size and growth to population quality indicators, such as levels of genetic variation. Despite an increase in the total number of black rhino, sample sizes for genetic studies tend to have been small due to the difficulty in collecting genetic material. Small sample sizes typically underestimate diversity measures like allelic richness (also called allelic diversity) and heterozygosity. Nei (1978) demonstrated that when estimating average heterozygosity (measure of genetic variation within a population), a small number of individuals could be used if a large number of loci (more than 50) were used in the study and the average heterozygosity is low. Nei (1978) also established that a small number of individuals could be sampled for determining genetic distance if the genetic distance was large and the average heterozygosity of the two species being compared was low.

The following are studies that have included black rhino. Most of the studies vary in sample size, examine fewer than 50 loci, and the genetic distance between the subspecies is not large. The combination of these limitations has occasionally lead to contradictory results. While each of these studies has contributed to the understanding of rhino genetics, there are still information gaps.

Allozymes

Allozymes are soluble protein-coding enzymes, usually taken from blood, kidney or liver, mixed with a buffer and separated by their charge or molecular weight on an electrophoresis gel. Allozyme markers are used to evaluate genetic variation, population structure and gene flow (Lowe et al. 2004). They are inexpensive, easy to detect and generally selectively neutral (Lowe et al. 2004); however they are limited in that they do not directly measure the amount of DNA variation within a population (Conner and Hartl 2004).

Merenlender et al. (1989) observed a significant lack of genetic variability across 25-30 loci in the four rhino taxa (*C. s. simum*, *C. s. cottoni*, *D. bicornis* and *R. unicornis*) they examined and more specifically, low amounts of genetic variation within *D. b. michaeli* black rhino samples (0.013 observed heterozygosity) from Kenya and east Africa (Table 1.2), concluding that the low levels of genetic variation was likely caused by recent historic demographic bottlenecks. While the authors suggested that the results were probably not important for short-term conservation goals, they recognized that their sample sizes were small (<10 samples for each taxa), which may have biased results.

Swart et al. (1994) narrowed their focus to four southern African black rhino populations (*D. b. minor*, Zambezi Valley, Zimbabwe (n=90); *D. b. bicornis*, Etosha, Namibia (n=6); *D. b. minor*, HiP (n=25) and *D. b. minor*, Mkuze (n=34), South Africa) (Table 1.2). All populations were in Hardy-Weinberg Equilibrium (HWE), six out of 30 loci were polymorphic and one locus was sex linked (*Gp-5*). When the 24 monomorphic loci were excluded from calculations, the proportion of heterozygote individuals for the four populations was between 0.036 – 0.059 with expected heterozygosity between 0.003 – 0.02. The Zimbabwe samples had

Table 1.2: Genetic studies that include black rhino; Type of study, Allozyme using plasma and red blood cells (RBC), Restriction Fragment Length Polymorphism (RFLP) using mitochondrial DNA (mtDNA) and microsatellites (msats) for black rhino (BR) and white rhino (WR); Number of restriction sites and number of loci used in the study, Sample Size, Species/Subspecies, Sample Country of Origin, Expected heterozygosity (H_e), Observed heterozygosity (H_o), Number of mtDNA haplotypes

Author	Type of Study	Number of sites / loci	Sample Size	Species/ Subspecies	Sample Origin	$\sim H_e$	$\sim H_o$	No. of mtDNA Haplotypes
Merenlender et al. 1989	Allozyme (plasma)	25-31	9	<i>D. b. michaeli</i>	Kenya/East Africa	0.013	-	-
			23	<i>C. s. simum</i>	South Africa	0.013	-	-
			7	<i>C. s. cottoni</i>	Sudan	0.019	-	-
			3	<i>R. unicornis</i>	India	0.000	-	-
Swart et al. 1994	Allozyme (plasma & RBC)	30 (6 polymorphic)	6	<i>D. b. bicornis</i>	Namibia	0.013	0.053	-
			105	<i>D. b. minor</i>	Zimbabwe	0.020	0.059	-
			34	<i>D. b. minor</i>	South Africa (HiP)	0.003	0.038	-
			25	<i>D. b. minor</i>	South Africa (Mkuze)	0.012	0.036	-
Swart & Ferguson 1997	Allozyme (plasma & RBC)	30 (6 polymorphic)	21	<i>D. b. bicornis</i>	Namibia	-	0.053	-
			135	<i>D. b. minor</i>	Zimbabwe	-	0.062	-
			25	<i>D. b. minor</i>	South Africa (HiP)	-	0.038	-
			34	<i>D. b. minor</i>	South Africa (Mkuze)	-	0.037	-
Ashley et al. 1990	RFLP (mtDNA)	18 (4 polymorphic)	11	<i>D. b. michaeli</i>	Kenya	-	-	1 (type B ₁)
			11	<i>D. b. minor</i>	Zimbabwe	-	-	2 (type A & B ₂)
			1	<i>D. b. minor</i>	South Africa	-	-	1 (type A)
O'Ryan et al. 1993	RFLP (mtDNA)	18 (monomorphic for <i>D. b. minor</i>)	4	<i>C. s. simum</i>	South Africa	-	-	2
			22	<i>D. b. minor</i>	South Africa (HiP & Mkuzi)	-	-	1
			2	<i>D. b. minor</i>	Zimbabwe	-	-	1
O'Ryan et al. 1994	RFLP (mtDNA)	16 (3 polymorphic)	5	<i>D. b. bicornis</i>	Namibia	-	-	1 (type B ₁)
			1	<i>D. b. michaeli</i>	South Africa	-	-	1 (type B ₂)
			26	<i>D. b. minor</i>	South Africa	-	-	1 (type A)
			1	<i>D. b. chobiensis</i>	Namibia	-	-	1 (type A)
Brown & Houlden 2000	Sequencing (mtDNA)	2 (forward and reverse) mt15996L (forward) mt1652H (reverse)	2	<i>D. b. michaeli</i>	Captive Born	-	-	2
			9	<i>D. b. minor</i>	Zimbabwe & Captive Born	-	-	5
Brown & Houlden 1999	Msat	11	7	<i>D. b. bicornis</i>	Unknown	0.686	-	-
		DB1, DB4, DB5, DB14,	2	<i>D. b. michaeli</i>	Captive Born	0.682	-	-
		DB23, DB30, DB42, DB44, DB49, DB52, DB66	5	<i>D. b. minor</i>	Zimbabwe & Captive Born	0.594	-	-
Cunningham et al. 1999	Msat	5 BR3, BR4, BR6, BR17, BR20	72	<i>D. bicornis</i>	Unknown	0.580	-	-
Garnier et al. 2001	Msat	10 BR4, BR6, BR17 (Cunningham et al. 1999)	35	<i>D. b. minor</i>	Zimbabwe	0.616	0.726	-
		DB1, DB5, DB23, DB44, DB49, DB52, DB66 (Brown & Houlden 1999)						
Harley et al. 2005	Msat	9	19	<i>D. b. michaeli</i>	Kenya	0.675	0.731	-
		BR4, BR6, BR17 (Cunningham et al. 1999)	47	<i>D. b. minor</i>	South Africa & Zimbabwe	0.459	0.436	-
		1	1	<i>D. b. chobiensis</i>	southern Angola	-	-	-
		DB1, DB14, DB44, DB49, DB52, DB66	1	<i>D. b. longipes</i>	Cameroon	-	-	-
		(Brown & Houlden 1999)	53	<i>D. b. bicornis</i>	Namibia	0.505	0.523	-
Nielsen et al. 2008	Msat	5 (WR) 12 (BR)	22	<i>C. s. simum</i>	South Africa (HiP)	0.420	0.436	-
		WR: AY138542, AY138543, AY138544, AY138545, AY138541 BR: AF12972, AF129726, AF129727, AF129729, AF129730, AF129734, AF129732, AY606078, AY606079, AY606080, AY606082, AY606083 (Nielsen et al. 2008)	6	<i>D. b. minor</i>	South Africa (HiP)	0.372	0.322	-
Karsten et al. 2011	Msat	10	77	<i>D. b. minor</i>	South Africa & Zimbabwe	0.44	0.38	-
		BR4, BR6, BR17 (Cunningham et al. 1999)	4	<i>D. b. bicornis</i>	Namibia	0.43	0.46	-
		DB1, DB14, DB49, DB66 (Brown & Houlden 1999)	4	<i>D. b. michaeli</i>	South Africa & Tanzania	0.54	0.54	-
		AY606078, AY606080, AY606083 (Nielsen et al. 2008)						
Muya et al. 2011	Msat	BR4, BR6, BR17 (Cunningham et al. 1999) DB1, DB5, DB30, DB44, DB52, DB66 (Brown & Houlden 1999)	145	<i>D. b. michaeli</i>	Kenya	0.69	0.70	-
Van Coeverden de Groot et al. 2011	Msat	DB1, DB44, DB52, DB66 (Brown & Houlden 1999) BR4, BR6, BR17 (Cunningham et al. 1999) BIRh2B, BIRh37D (Van Coeverden de Groot et al. 2011)	144	<i>D. b. bicornis</i>	Namibia	0.51	0.52	-

the highest gene diversity, while Mkuze had the lowest. The authors suggested that the two KZN populations (HiP and Mkuze) were not genetically depauperate and many southern African black rhino populations had similar genetic variation to populations of “some outbreeding mammal species”, although the authors do not site examples of said outbred species. Swart et al. (1994) also propose that the level of genetic variation in the southern African populations was representative of the heterozygosity present before drastic population declines and that this would be advantageous to captive breeding programmes.

Swart and Ferguson (1997) studied two subspecies of black rhino (one *D. b. bicornis* and three *D. b. minor* populations) (Table 1.2). They concluded that the four populations were conspecific isolated remnants of a large ancestral population; none of the populations belonged to discrete subspecies but were instead part of a west-to-east ‘genetic continuum’ where by the Etosha (Namibia) and KZN (South Africa) populations are the extremes, but mere subsets of the Zambezi (Zimbabwe) population. They concluded that short-term genetic management for the species was unnecessary due to large genetic variation and no evidence of inbreeding or excess in homozygosity. However, they concluded that the levels of variation in the Zimbabwe *D. b. minor* population indicate that it is the only population of black rhino to retain pre-bottleneck levels of genetic variation. They recommended immediate genetic management in order to maintain the level of variability in the Zambezi, Zimbabwe population.

Mitochondrial DNA (mtDNA)

MtDNA consists of a haploid circular molecule found in the cellular mitochondria of most eukaryotes. It is typically maternally inherited in mammals and lacks recombination due to the nature of its replication process. MtDNA is more sensitive to changes in population demography because it has a quarter the effective population size (N_e) compared with nuclear loci. MtDNA has a relatively high mutation rate and shows higher levels of polymorphism compared to many nuclear genes making it useful when looking for patterns of genetic differentiation (Moritz et al. 1987). Studies of mtDNA can be used effectively in long-term and short-term management of populations, more specifically to (1) measure genetic variation in recently declining populations (2) define Evolutionarily Significant Units (ESUs) and (3) to ascertain evolutionary or phylogenetic conservation value of populations (Moritz 1994). Studies on black rhino mtDNA include the use of restriction maps and direct DNA sequencing of the control region and 12S rRNA which have indicated differences between the black rhino subspecies and suggest there may be population differentiation (Harley et al. 2005).

Restriction Enzymes Analysis of mtDNA

Restriction fragment length polymorphisms (RFLP) are used to evaluate genetic variation, population structure and gene flow; however, restriction enzymes identify differences in the sequence of DNA, not the expressed proteins (Lowe et al. 2004). RFLP results are repeatable and a considerable amount of variation can be identified if the right combination of restriction enzymes is developed.

Unfortunately, this method can be expensive, time consuming and combining results from different labs can be difficult (Lowe et al. 2004).

Ashley et al. (1990) examined mtDNA RFLPs of black rhinos from three different geographic populations: Zimbabwe (*D. b. minor*, n=11), South Africa (*D. b. minor*, n=1) and Kenya (*D. b. michaeli*, n=11) (Table 1.2). They found a small amount of intraspecific variation, with only three mtDNA haplotypes; one unique haploptype in Kenya (*D. b. michaeli*), one unique haploptype in Zimbabwe (*D. b. minor*) and one shared with Zimbabwe and South Africa (*D. b. minor*). The DNA sequences of all three haplotypes were similar which led the authors to conclude that the subspecies had recently shared a common ancestor.

O'Ryan and Harley (1993) used mtDNA restriction maps using 18 restriction endonucleases and estimated that the time divergence from a common ancestor for black rhino and white rhino was 3.4×10^6 years ago. Black rhino samples were from HiP (n=16), Mkuze (n=6) and Zimbabwe (n=2) (Table 1.2). White rhino samples (n=4) were from HiP. There were no polymorphic sites detected in the restriction map of the black rhino and only one polymorphic site in the white rhino. The results contrasted with those of Ashley et al. (1990) who also used a restriction-fragment size approach, but were consistent with Merenlender et al. (1989) allozyme findings of small intraspecific variation.

O'Ryan et al. (1994), again using a mtDNA restriction map, analyzed differentiation among black rhino subspecies and populations. Their subspecies samples included *D. b. minor* (n=26), *D. b. bicornis* (n=5), *D. b. michaeli* (n=1) and *D. b. chobiensis* (n=1) (Table 1.2). In their findings, they recommended discarding *D. b. chobiensis* as a subspecies and placing it in the south-central

ecotype with *D. b. minor*. Their results showed little variation with only two site differences between 33 individuals from three different geographic regions and monomorphic mtDNA restriction maps within the same geographic region.

MtDNA Sequencing

DNA sequencing typically involves the amplification of a segment of DNA (e.g. mitochondrial DNA control region) via polymerase chain reaction (PCR) and it is a direct measure of genetic variation.

Tougard et al. (2001) sequenced the mtDNA 12S rRNA in order to establish where Sumatran rhino fit within the other four extant Asian and African species. They narrowed the divergence time of the species to approximately 26 million years ago with the Sumatran rhino forming a sister group of the genus *Rhinoceros*. Later, Fernando et al. (2006) used mtDNA 12S rRNA and control region sequences to examine the genetic divergence and level of variation within and between two extant Javan rhino populations based on samples from all five extant rhino species. They established that the two populations each formed a discrete ESU and recommended independent management of each population.

Brown and Houlden (2000) sequenced the non-coding mtDNA control region of captive *D. b. michaeli* (n=2) and wild captured Zimbabwe *D. b. minor* (n=9) to examine evolutionary relationships. Five haplotypes were found in the nine *D. b. minor* samples with a haplotype diversity of 0.86 (Table 1.2). Both *D. b. minor* and *D. b. michaeli* were shown to be reciprocally monophyletic, meaning that all members of a lineage share a more recent common ancestor, with a divergence time between the two subspecies ranging from 0.92 to 1.3 MYA.

DNA Microsatellites

DNA microsatellites are a type of co-dominant DNA marker used extensively in contemporary population and evolutionary genetic studies. (Slatkin 1995b). They are tandem repeats usually between one and five base pairs long (Jarne and Lagoda 1996) that are typically non-coding and are not influenced by the selection process (Slatkin 1995). Microsatellites are found throughout an organism's genome and are surrounded by unique DNA sequences, which enable primers to be designed and used to amplify each locus separately (Jarne and Lagoda 1996). The mutation rate of microsatellites is important because the rate of change and model of mutation help determine population structure (Jarne and Lagoda 1996). Microsatellite mutations are approximately 10^{-5} to 10^{-2} higher than seen in allozymes. Mutations occur through slippage through increases and decreases in the number of unit repeats (Jarne and Lagoda 1996).

Brown and Houlden (1999) and Cunningham et al. (1999) were the first to specifically isolate microsatellite sequences from black rhinos followed by Nielsen et al. (2008) and Van Coeverden de Groot et al. (2011) (Table 1.2). Brown and Houlden (1999) designed 11 microsatellite marker primers to assess genetic diversity within an Australian *ex situ* breeding and conservation programme of *D. b. michaeli* (originally from Kenya (n=2) and *D. b. minor* (originally from Zimbabwe (n=5)). The mean expected heterozygosity (H_E) over 11 loci for *D. b. minor* was 0.594 ± 0.068 and 0.682 ± 0.085 for *D. b. michaeli*.

Garnier et al. (2001) conducted a genetic analysis using ten microsatellites and used DNA from faecal samples to increase the understanding of the mating system, reproductive skew and effective population size of a *D. b. minor* population (n=35) in Save Valley, Zimbabwe

(Table 1.2). This was the first study to provide genetic evidence of both polygyny and a male reproductive skew in black rhino. The Save Valley population was founded with individuals from Zambezi, Zimbabwe, which Swart and Ferguson (1997) suggested had the highest level of genetic variation of all black rhino populations and probably reflect that of a pre-bottleneck population. The published genotypes provide a genetic record that will enable conservation managers of other Zambezi founder populations to compare the levels of genetic variations within their own populations. These data will also be helpful in the future to the Save Valley population as a decrease in the level of variability will be easily identifiable.

Harley et al. (2005) used nine of the black rhino microsatellite markers to establish baseline information regarding levels of genetic diversity and population differentiation in black rhino subspecies (*D. b. bicornis* (n=53); *D. b. minor* (n=47); *D. b. michaeli* (n=19) and the now extinct *D. b. longipes* (n=1) and *D. b. chobiensis* (n=1)) (Table 1.2). They found that *D. b. michaeli* had the highest level of genetic diversity with an expected heterozygosity (H_E) of 0.675 followed by *D. b. bicornis* ($H_E = 0.505$) and *D. b. minor* ($H_E = 0.459$). The authors point out that the *D. b. minor* results may indicate a level of population substructure due to samples coming from a number of games reserves in South Africa and Zimbabwe. The possible population substructure of *D. b. minor* was not important to their study however, and was therefore, not investigated.

Nielsen et al. (2008) designed 21 microsatellites for both black and white rhino, seven of which were polymorphic and were used to distinguish the two species from each other (Table 1.2). The authors do not specify with which subspecies of black rhino (n=6) they were working, but an assumption can be

made that the subspecies is *D. b. minor* as the authors stated that the samples were from HiP.

When comparing results among genetic studies with differing sample sizes, expected heterozygosity is preferred, as it takes into account sample size variation. Nielsen et al. (2008) reported their microsatellite observed heterozygosity ($H_0 = 0.322$), instead of H_E (0.372) for two out of three of the comparisons with other black rhino studies and also compared their results against different black rhino subspecies. Since the aim of their study was to differentiate black rhino samples from white rhino samples, identifying differences between black rhino subspecies was not considered important. That being said, they concluded that the wild black rhino population in South Africa retains a moderate degree of allele diversity. They reached this conclusion by stating that their black rhino $H_0 = 0.322$ (n=6) was lower than Brown and Houlden's (1999) $H_0 = 0.660$, however the H_0 the authors reported for Brown and Houlden (1999) was the mean H_0 of two black rhino subspecies (*D. b. minor*, $H_0 = 0.594$ (n = 5) and *D. b. michaeli*, $H_0 = 0.682$ (n=2)). Brown and Houlden (1999) used 11 microsatellites, six of which were used by Nielsen et al. (2011). When comparing their microsatellite results to Garnier et al.'s (2001) *D. b. minor* (n = 35), the authors used Garnier et al.'s (2001) $H_0 = 0.726$ instead of $H_E = 0.62$. For their last result comparison, Nielsen et al. (2008) compared their $H_E = 0.372$ with that of Harley et al. (2005), except they only reported the results for Harley et al.'s (2005) *D. b. michaeli*, $H_E = 0.68$ (n= 19), which had the highest H_E of each of the three subspecies in Harley et al's (2005) study (*D. b. minor*, $H_E = 0.46$ (n=46); *D. b. bicornis*, $H_E = 0.51$ (n=53)) (Table 1.2). Finally, due to the nature of their study, the H_E and H_0 results Neilsen et al. (2008) reported for the

black rhino in their study included the use of both black rhino and white rhino loci instead of separating out the data between the two species. When specific white rhino marker results are excluded, the black rhino H_E and H_O increase to 0.439 and 0.411 respectively, much higher than the reported mean H_E and H_O of 0.372 and 0.322, respectively.

Muya et al. (2011) focused on 12 of 16 extant *D. b. michaeli* populations in Kenya using both mtDNA control region sequencing and nine microsatellite loci (Table 1.2). They confirmed previous studies (Harley et al. 2005) of moderate to high levels of genetic diversity in their *D. b. michaeli* metapopulation, reporting a mean mtDNA haplotype diversity (h) of 0.73 ± 0.14 with mean microsatellite H_E and H_O of 0.70 ± 0.087 and 0.69 ± 0.034 respectively.

Van Coeverden de Groot et al. (2011) used nine polymorphic microsatellite loci to examine genetic diversity and structure of *D. b. bicornis* (n=144) individuals of Etosha National Park, Namibia (Table 1.2); a population that experienced a significant population increase due to increased protection. The results were to be utilized as a baseline with which conservation managers can measure changes in the level of genetic variation in the future. Mean expected heterozygosity for the samples was 0.51, similar to levels published for *D. b. bicornis* by Harley et al. (2005).

Karsten et al. (2011) used 10 microsatellites to evaluate levels of genetic diversity, differentiation and inbreeding among *D. b. minor* (n = 74) in seven game reserves in KZN, South Africa and a single population of *D. b. minor* (n = 3) in Zimbabwe that was founded (and is managed separately from native Zimbabwe *D. b. minor*) with black rhino from KZN. They also compared the

results to *D. b. bicornis* ($n = 4$) and *D. b. michaeli* ($n = 4$) (Table 1.2). They found low levels of differentiation among KZN metapopulation and their microsatellite variation over 10 loci in *D. b. minor* ($H_E = 0.44$) was lower than for the other two subspecies ($H_E = 0.54$ for *D. b. michaeli* and $H_E = 0.43$ for *D. b. bicornis*). The authors stated that the KZN *D. b. minor* values still fell within the range of other large mammals across Africa (African buffalo (*Syncerus caffer*) $H_E = 0.76$ and $H_O = 0.73$ (van Hooft et al. 2000) and $H_E = 0.58$ and $H_O = 0.52$ (Simonsen et al. 1998), African elephant (*Loxodonta africana africana*) $H_E = 0.96$ and $H_O = 0.37$ (Whitehouse and Harley 2001), Black wildebeest (*Connochaetes gnou*) $H_E = 0.35$ (Grobler et al. 2005), Blue wildebeest (*C. taurinus*) $H_E = 0.65$, (Grobler et al. 2005), Cape Mountain zebra (*Equus zebra zebra*) $H_E = 0.38$ and $H_O = 0.24$ (Moodley and Harley 2005), Hartmann's mountain zebra (*E. z. hartmannae*) $H_E = 0.54$ and $H_O = 0.48$ (Moodley and Harley 2005)). While it is true that the $H_E = 0.44$ for *D. b. minor* falls within the 0.38 – 0.96 of the other large mammals they listed, caution should be exercised when comparing levels of genetic variation across species since experiments that do not include the entire genome (e.g. microsatellite) represent a small portion of total DNA (Selander and Johnson 1973). The authors concluded that the use of translocations within the KZN metapopulation has helped *D. b. minor* retain acceptable amounts of genetic diversity and further concluded there was no need to change how black rhino in KZN were managed. They, however, missed an opportunity to compare regional levels of microsatellite variation for *D. b. minor* of KNZ to native Zimbabwe *D. b. minor* by not comparing their findings to the genotype data provided in Garnier et al.'s (2001) paper.

The levels of genetic variation within and between populations of *D. b. minor* from different regions (e.g. South Africa and Zimbabwe) should be examined separately with comparable and repeatable methods (e.g. similar microsatellite DNA markers) to ensure accuracy. With successful reintroductions of South African black rhino continuing into areas where they have been extirpated, expansion will eventually mean that regional populations may meet and become part of the same management scheme. However, genetic considerations (e.g. differentiation) need to be addressed to ensure that long-term viability of regional populations is not compromised by outbreeding.

1.5 Management

Translocation and Reintroductions

A translocation is the well-planned movement of animals from one part of their range to another (IUCN 1987). It is a powerful tool used by conservation managers to reintroduce animals to areas where they have been extirpated or have undergone dramatic declines in their distribution ranges (e.g. big horn sheep (*Ovis canadensis*), Singer et al. 2000; black bears (*Ursus americanus*) Smith and Clark 1994); to genetically augment existing populations (white-spotted charr (*Salvelinus leucomaenis*) Yamamoto et al. 2006) and establish extralimital populations to reduce the possibility of species loss from catastrophe (e.g. Rarotonga Monarch (*Pomarea dimidiata*) (Griffith et al. 1989; Robertson et al. 2006)).

Reduced levels of heterozygosity and allelic diversity in founder populations is well documented (Nei et al. 1975; Leberg 1992; Keller and Waller 2002), but for pragmatic reasons reintroduced populations of threatened and

endangered species are typically founded with a small number of individuals (Griffith et al. 1989; Kerley et al. 2003). Unfortunately, those individuals are usually sourced from populations that have themselves been through a bottleneck. Since reintroduced populations often only retain a small portion of the genetic variation from the source at functional and neutral loci, if the source population has experienced a genetic bottleneck, the rate of loss of genetic variation in the founder population may be accelerated (Bijlsma et al. 2000; Keller and Waller 2002; Leberg 1993). Establishing founder populations with individuals from different populations of the same subspecies (different genetic stock) or with high levels of genetic variation may be more likely to be successful (e.g. increased fitness)(Leberg 1993).

Reintroductions of megaherbivores like elephants and rhino require different treatment than smaller animals for various reasons. Some of these include: (1) the nature of plant abundance and disturbance due to their size, (2) they do not persist outside of conservation areas, (3) they are charismatic species and attract attention (e.g. tourism, conservation), and (4) management techniques are well-developed (Kerley et al. 2003). The primary reason for establishing founder populations of megaherbivores is because parks and reserves can no longer support (e.g. nutritionally) an increase in the number of individuals in established populations.

The first successful black rhino translocation from HiP was to Ndumo Game Reserve in 1962 (Table 1.1), coordinated under the Natal Parks Game and Fish Preservation Board (Hitchins 1984). Successful translocations from HiP continue today under current management of Ezemvelo KwaZulu-Natal Wildlife. Like HiP, Mkuze Game Reserve (MGR) is home to the only other relict

population of *D. b. minor* (n= 45; D. Kelly, pers. comm.) in South Africa. However, MGR is smaller than HiP (38,000 ha compared to ~96,000 ha) and translocations out of MGR occur less frequently than from HiP.

Receiving sites for black rhino translocations include other KZN reserves capable of supporting black rhino (e.g. Ndumo Game Park, Tembe Elephant Reserve, Ithala Game Reserve, Zululand Rhino Reserve, Pongola Game Reserve, Phinda Resource Reserve, Weenan Game Reserve and Eastern Shores Game Reserve; Figure 1.3). As per recommended guidelines, when black rhino populations within smaller, established KZN reserves approach 75% of their estimated carrying capacities, individuals are removed and used for reintroductions elsewhere. In addition to parks within KZN, receiving translocation sites also include parks outside KZN like Kruger National Park, Pilanesberg Game Reserve and across borders (e.g. Zimbabwe, Mozambique and Swaziland; Hitchins 1984, Emslie et al. 2009). As translocation success rates improve, it becomes increasingly necessary to study the genetic diversity of various black rhino populations in order to plan long-term management of the species (Emslie and Brooks 1999).

The Need for a Paradigm Shift in Rhino Conservation Practice

As black rhino species recovery continues, the focus of conservation managers on population growth (number of black rhino) will need to be replaced by that of population quality (e.g. genetic variation and N_e). Marked decreases in levels of genetic variation in small and recovering populations could decrease fitness or limit the long-term capacity of a population to respond to changes in the environment (Westemeier et al. 1998). The shift in focus is important because if

genetic factors are disregarded, it could lead to inappropriate recovery strategies (Frankham 2005). If rhino poaching can be reduced and current population trends continue, in 20 years black rhino are likely to be regarded as a conservation success, much as the southern white rhino conservation is today.

While it is possible to have high levels of genetic variation represented by a few individuals forming a founder population, variation may be lost if the population remains small (Lande and Barrowclough 1987). Currently, black rhino from HiP are used to reintroduce the species back into its historic range. However, no individuals immigrate (naturally or assisted through translocations) back into HiP, meaning that the HiP population is unable to benefit from the expansion in the metapopulation, which would slow or stop a decline in levels of genetic variation. HiP has been through a population reduction and may have lost a significant amount of allelic diversity, in which case HiP could benefit from translocations either from the KZN metapopulation or possible native Zimbabwe populations back into the reserve to replenish diversity. Comparing levels of genetic variation within HiP and the KZN metapopulation against native Zimbabwe populations may help with developing management schemes to prevent loss of genetic variability within the *D. b. minor* subspecies.

Rhino Management Groups

In my thesis, I will refer to several different organizations that make recommendations regarding black rhino management. Here I describe the international agencies and how they are associated with each other. There is

considerable crossover between the groups as individuals are often affiliated with more than one organization that drafts black rhino management guidelines.

The International Union for Conservation of Nature (IUCN)

The International Union for Conservation of Nature is a multinational organization dedicated to the conservation of nature and natural resources. The IUCN is divided into specialized groups like the Species Survival Commission (SSC), a science-based network that provides feedback to the IUCN on biodiversity, species concerns, and dispenses recommendations to specialized conservation projects. The African Rhino Specialist Group (AfRSG) is a working group within the SSC network. The group meets every two years to update rhino statistics, set priorities for populations of rhino and generate 'Action Plans'. AfRSG members are also usually involved in regional projects (Emslie et al. 2007; Emslie and Brooks 1999; Emslie et al. 2009).

Regional Rhino Conservation Groups Affiliated with the IUCN

Southern African Development Community (SADC) is an organization of southern Africa nations including Angola, Botswana, Malawi, Mozambique, Namibia, South Africa, Swaziland, Tanzania, Zambia and Zimbabwe (Emslie et al. 2009) established to promote improved standards of living in member states (SADC 2010). Food, Agriculture and Natural Resources (FANR 2010), a department within the SADC oversees the SADC Wildlife Programme of Action (WPA). The SADC WPA is responsible for managing several projects, one being the SADC Regional Programme for Rhino Conservation that focuses on conservation efforts of both black and white rhinos in southern Africa. The SADC Rhino Management Group (RMG) concentrates on black rhino efforts and

implementing conservation plans in South Africa, Namibia, Swaziland and Zimbabwe. Members of the IUCN-AfRSG are an integral part of the SADC RMG (Emslie and Brooks 1999; Emslie et al. 2007; Emslie et al. 2009).

World Wildlife Fund

The World Wildlife Fund (WWF) South African Black Rhino Range Expansion Project (BRREP) began in 2003 and is currently directed by Dr. Jacques Flamand and coordinated with Ezemvelo KwaZulu-Natal Wildlife. The project is ongoing and focuses on successful reintroduction and translocation of black rhino from HiP and MGR source populations to areas where they have been extirpated.

1.6 Recommended Guidelines for Black Rhino Management

The Status Survey and Conservation Action Plan for the African Rhino (SSCAP) (Emslie and Brooks 1999) and the IUCN's Guidelines for the *in situ* Re-introduction and Translocation of African and Asian Rhinoceros (Emslie et al. 2009) are examples of management plans detailing necessary concerns for conservation managers including genetic monitoring. The SSCAP incorporates guidance for protection, ascertaining sex and age structure of populations, estimating population sizes, recording mortalities and estimating carrying capacities, each of which is necessary as part of a comprehensive plan. It also stresses the importance of genetic diversity maintenance. Recommendations also include founding new populations in areas that formerly supported the subspecies with cohorts consisting of at least 20 individuals. In addition, guidelines also suggest that when translocating rhino to established populations, the newly translocated individuals have as little "genetic similarity"

as possible with the receiving population. They also advise that if possible, one rhino per generation should be introduced and accurate stud books be maintained.

The IUCN re-introduction guidelines (Emslie et al. 2009) give detailed criteria for African and Asian rhino translocations for both *in situ* and *ex situ* conservation schemes. It spells out important details needed before (i.e., harvesting in existing populations, selecting rhino for translocation, nominating new locations), during (i.e., logistics, veterinary care, holding) and after (i.e., monitoring, protection) translocations take place. Regarding genetic management, the IUCN guidelines are similar to the SSCAP in that translocated individuals should be unrelated if they are being introduced to an existing population, founder populations need to be of the same subspecies that were in that particular historical range, the number of individuals in a new population should be at least 20, and newly founded populations should be carefully monitored.

Local agencies like Ezemvelo KZN Wildlife (EKZNW) and South Africa National Parks (SANParks) together with private and community conservancies work hard to manage black rhino populations. The local private and government management groups create management plans and follow recommended guidelines to the very best of their ability considering the financial and manpower constraints many game reserves face.

1.7 Thesis Structure

The specific objectives of this study were to:

- 1) Determine the sequence of the mitochondria DNA (mtDNA) control region of three black rhino subspecies, estimate the level of variation within the HiP source and KZN metapopulation and compare the results to native Zimbabwe *D. b. minor* and the other black rhino subspecies.
- 2) Use ten microsatellite DNA markers to estimate the levels of heterozygosity and allelic diversity in HiP and KZN metapopulation and compare the results to previously published microsatellite data of native Zimbabwe *D. b. minor* as well as the other black rhino subspecies to determine whether or not HiP still has an appropriate level of genetic variation to use to establish founder populations of *D. b. minor*.
- 3) Perform a Population Viability Analysis (PVA) based on genetic data from HiP and vital rates to model the effects of increasing population size and supplementation to determine which management scenarios would be most effective for minimizing the loss of genetic variation.

When attempting to ascertain historic gene flow, evaluate a species demographic limits and gain understanding into a species population structure, mtDNA is an excellent metric with which to start (Rubinoff and Holland 2005). MtDNA is haploid, generally maternally inherited in mammals, lacks recombination and has a high mutation rate. Chapter Two examines sequences of the highly variable mtDNA control region of the *D. b. minor* source (HiP) and metapopulation of KZN and compares them against previously published native

Zimbabwe *D. b. minor* samples, as well as, samples of *D. b. bicornis* from Namibia and *D. b. michaeli* from Kenya. Identifying the level of mtDNA variation within HiP is the first step in determining whether or not HiP remains an appropriate source for *D. b. minor* for South African founder populations. This chapter is published (Appendix A) as:

Anderson-Lederer, R.M., Linklater, W.L., and P.A. Ritchie (2012) Limited mitochondrial DNA variation within South Africa's black rhino (*Diceros bicornis minor*) population and implications for management. African Journal of Ecology 50(4): 404-413

I have changed the format to conform to a suggested thesis format.

Chapter Three continues the investigation of genetic structure by examining ten microsatellite DNA loci to estimate the levels of heterozygosity and allelic diversity in HiP and the KZN metapopulation comparing them against a small number of samples of *D. b. bicornis* from Namibia and *D. b. michaeli* from Kenya as well as previously published *D. b. minor* microsatellite data including native Zimbabwe populations. This chapter is in preparation for submission to Journal of Zoology as:

Anderson-Lederer, R.M., Linklater, W.L., and P.A. Ritchie (In prep) Low levels of microsatellite DNA variation and possible management considerations for black rhino (*Diceros bicornis minor*) in KwaZulu-Natal, South Africa

MtDNA and microsatellite DNA results help describe past and current genetic variation as well as assisting conservationists with understanding mechanisms that are responsible for variation in allele frequencies (Conner and Hartl 2004), but they are only part of a larger management picture. Chapter Four builds on the previous chapters by incorporating microsatellite DNA and HiP vital rate information into a VORTEX population viability analysis (PVA). A

PVA was used to model the effects of increasing population size versus supplementation to determine which management scenarios would be most effective for maintaining or minimizing the loss of genetic variation in a source population.

Chapter Five is a synopsis of work completed. My findings contribute to the knowledge base already accumulated for the KZN *D. b. minor* metapopulation and will assist wildlife managers improve conservation plans as *D. b. minor* enter recovery. Results from the data chapters are reviewed and recommendations for management of black rhino are made. Shortcomings of this project are also pointed out and suggestions for future work are addressed.

CHAPTER TWO

Limited mitochondrial DNA variation within South Africa's black rhino (*D. b. minor*) population and implications for management

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Photo by Rosalynn Anderson-Lederer

2.1 Abstract

The taxonomy of African black rhinoceros (*Diceros bicornis*) remains unresolved. Maintaining levels of genetic diversity, and species rescue by reintroduction and restocking requires its resolution. I compared the sequences of the mitochondrial DNA (mtDNA) control region for a total of 101 *D. bicornis* from three subspecies: *D. b. minor*, *D. b. michaeli* and *D. b. bicornis*. A single unique haplotype was found within the 65 *D. b. minor* samples from KwaZulu-Natal (KZN) Province, South Africa, 55 of which came from Hluhluwe-iMfolozi Game Park (HiP) and Mkuze Game Reserve (MGR) source populations. However, six different haplotypes were represented in 11 *D. b. minor* samples from Zimbabwe. Similarly, published autosomal microsatellite data indicate low levels of diversity within the KZN *D. b. minor* populations. The low levels of mtDNA diversity within the KZN metapopulation point to the possible need for genetic supplementation. However, there is a need to determine whether the low levels of genetic variation within KZN *D. b. minor* is a result of the recent bottleneck or if KZN historically always had low diversity.

2.2 Introduction

Species conservation depends on identifying genetically distinct groups or management units and implementing strategies to retain genetic variation. Genetically distinct populations can contain unique genetic variation and/or they can be locally adapted to their habitat. Mixing them with other populations may break up genetically complex traits and, in some cases, lead to outbreeding depression (Templeton 1986; O'Ryan et al. 1994). Alternatively, genetic differences between populations can also result from strong genetic drift caused

by population fragmentation and declining population sizes (Frankham et al. 2002; Allendorf and Luikart 2007). When the genetic structure and historic pattern of gene flow of a species has been described, reintroduction methods can be used to secure locally adapted populations or restocking used for genetic supplementation.

Variation in mitochondrial DNA (mtDNA) is a particularly useful metric for determining population structure and history (Moritz 1994). The control region of mtDNA is highly variable and it can often be used to resolve phylogenetic relationships between closely related taxa or for describing the genetic structure within species (Moritz et al. 1987). MtDNA is maternally inherited and so does not recombine (Hayashi et al. 1985), which means it reflects a quarter the effective population size (N_e) compared with nuclear loci and hence it is more sensitive to changes in population demography.

The black rhinoceros (*Diceros bicornis*: Perissodactyla) once ranged across the African continent and numbered in the hundreds of thousands (Western and Vigne 1985). By 1969, their numbers had declined to ~65,000 (Muya and Ouge 2000) and, during the last century, the species disappeared faster than any other large mammal (Hitchins 1975; Western and Vigne 1985). The major causes for their decline have been anthropogenic, primarily illegal hunting (Western and Vigne 1985; Emslie and Brooks 1999; Amin et al. 2006). Nevertheless, conservation efforts have seen *in situ* black rhino numbers increase from a low of 2,475 individuals in 1993 to approximately 4,880 in 2010 (Emslie 2011).

Three extant black rhino subspecies are recognised across Africa, including approximately 742 *D. b. michaeli* (Eastern black rhino), 1,922 *D. b.*

bicornis (South-western black rhinoceros) and 2,216 *D. b. minor* (South-central black rhinoceros) (Emslie 2011). Appraisal of the black rhino subspecies was initially based on skull measurements (Zukowsky 1965; Groves 1967; du Toit 1987), however, uncertainty regarding taxonomy remained (du Toit 1987). Although there are apparently no impervious geographic boundaries or reproductive barriers between the subspecies, they occupy different areas with distinct habitats and climates (Harley et al. 2005; Emslie and Brooks 1999). With no historical records of migration and the extent of gene flow between the subspecies unknown, some authors have speculated that each subspecies may have genetic or behavioural adaptations to their local environments (Emslie and Brooks 1999; Harley et al. 2005). Their suggestion regarding genetic differences was confirmed through recent mtDNA and autosomal DNA analyses (Merenlender et al. 1989; Ashley et al. 1990; O'Ryan and Harley 1993; O'Ryan et al. 1994; Swart and Ferguson 1997; Brown and Houlden 1999, 2000; Nielsen et al. 2008; Karsten et al. 2011; Muya et al. 2011). Thus, current black rhino management policy is for each subspecies to be managed separately in order to maintain possible local adaptive traits and minimize the risk of outbreeding depression (Templeton 1986; O'Ryan et al. 1994; Brown and Houlden 2000; Harley et al. 2005).

The largest remnant population of the critically endangered (IUCN 2008) *D. b. minor* subspecies is in Hluhluwe-iMfolozi Game Park (HiP) (n=~220 Clinning *et al.* 2009) in KwaZulu-Natal (KZN) Province, South Africa (Figure 2.1). KZN black rhino have been separated from other populations to the north (e.g. Zimbabwe) since at least the latter half of the 19th century (Swart et al. 1994). HiP and the smaller remnant in Mkuze Game Reserve (MGR) (n=~45 D.

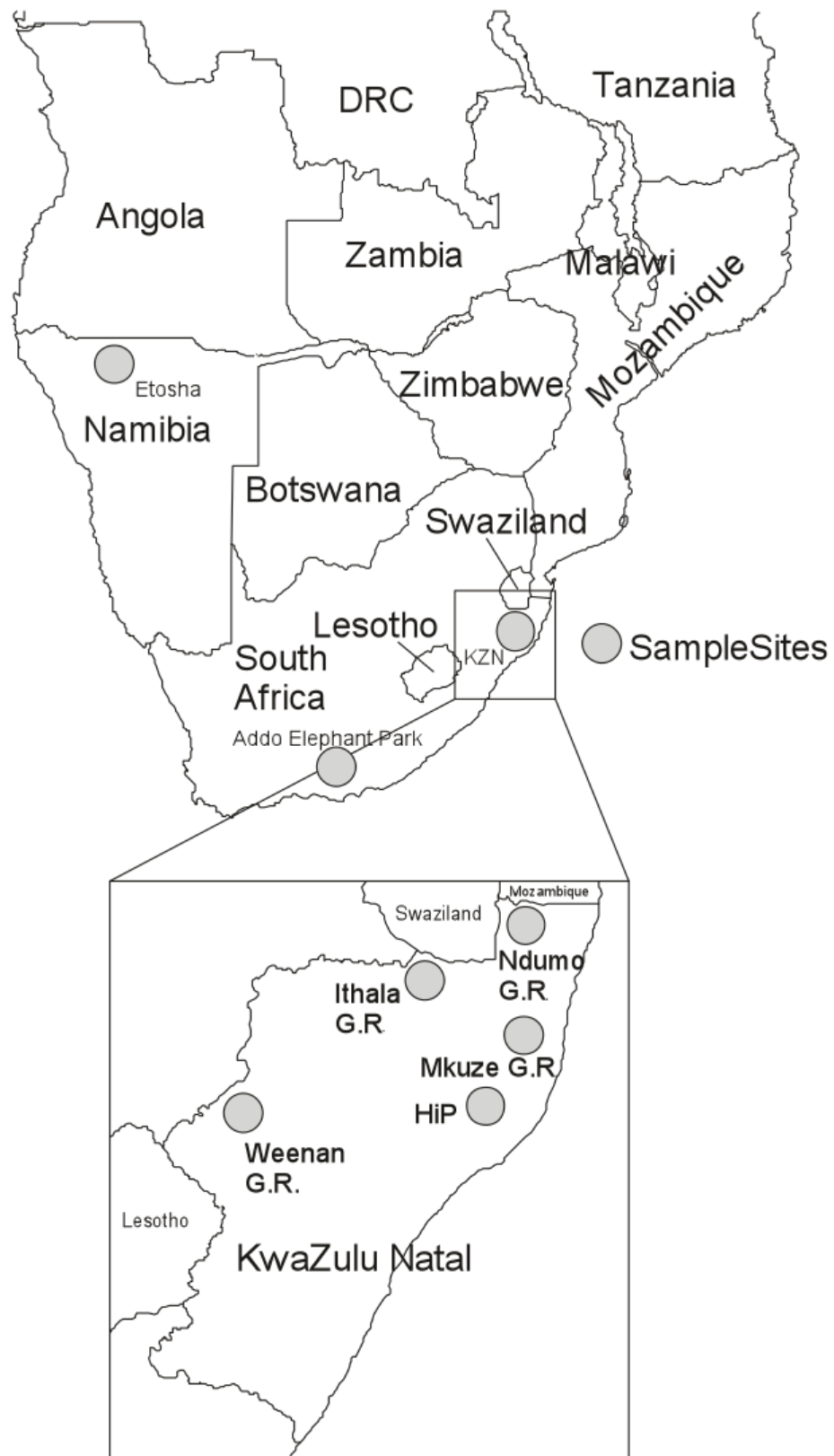


Figure 2.1: Map of Southern Africa showing black rhinoceros sample sites. Inset showing KwaZulu-Natal Game Reserves (Ndumo Game Reserve, Ithala Game Reserve, Mkuze Game Reserve, Hluhluwe-iMfolozi Game Park (HiP) and Weenan Game Reserve)

Kelly pers. comm.) have been sources for metapopulation expansion and genetic management by reintroduction and re-stocking. Translocations from HiP to other KZN reserves first began in 1962, expanded to other South African provinces and later to other African nations (e.g., Zimbabwe, Zambia, Hitchins 1984; Emslie et al. 2009). The potential now exists for KZN *D. b. minor* to be mixed with *D. b. minor* in or from other smaller African populations, especially those in Zimbabwe, if they are not too genetically divergent. Although the KZN population will likely be strategic to the subspecies recovery throughout the African continent (Emslie and Brooks 1999), no study has yet compared the mtDNA sequences of the KZN *D. b. minor* metapopulation with populations outside South Africa.

The aim of this study was to use mtDNA control region sequences (406 bp) to determine the level of variation within the *D. b. minor* source population at HiP (n=50) and compare it with the KZN metapopulation (n=15) and *D. b. minor* populations outside South Africa (n=11) and the other black rhino subspecies (*D. b. michaeli* n=21, *D. b. bicornis* n= 4). I considered the implications of the findings for the long-term management of *D. b. minor* and made recommendations for possible future research.

2.3 Materials and Methods

Sampling

Samples of blood and pinna ear tissue were collected from individuals of *D. b. minor* in the KZN province in South Africa (n=65), *D. b. michaeli* in Addo Elephant National Park, South Africa (n=1) and *D. b. bicornis* in Namibia's Northern Region (n=4) (Figure 2.1). The samples were acquired

opportunistically during routine translocation and ear notching (for identification) events from 2002 to 2009. Blood samples were stored in cryovials containing 1mL of DMSO/EDTA/Tris/salt solution (Seutin et al. 1991).

DNA Sequencing and Analysis

DNA extraction.

Seventy microlitres of the preserved blood solution or a 3mm x 3mm piece of pinna ear tissue was digested in an SDS/proteinase-K solution. After dissolution, a standard phenol-chloroform DNA extraction and ethanol precipitation was conducted following the procedure of Sambrook et al. (1989).

PCR and DNA sequencing

A fragment of the mitochondrial DNA control region (406bp) was amplified using the primers mt15996L (5'-TCCACCATCAGCACCCAA-AGC-3') (Campbell et al. 1995; Brown and Houlden 2000) and mt16502H (5'- TTTG-ATGGCCCTGAAGTAAGAACCA- 3') (Brown and Houlden 2000; Moro et al. 1998). PCR amplifications using 1-2 µL of DNA template were carried out in 25µL volumes with 67 mM Tris pH 8.8, 16mM (NH₄)₂SO₄, 1.5 mM MgCl₂, 0.4 µg/ml BSA, 0.4 µL of each of the forward and reverse primer, 200 µM of each dNTP, and 0.5 to 1 units of BIOTAQ DNA polymerase (Bioline). Thermal cycling was carried out using an Eppendorf Mastercycler for; 94°C 2 min, (94°C 3 min, 50-54°C 30 sec, 72°C 2 min), repeated for 30-40 cycles, followed by a final step of 72°C 3 min.

PCR products were electrophoresed in agarose gel and a molecular weight standard was used to determine the size of amplified products. Products of the correct size were purified using column purification (Roche) or ExoSAP-

IT (GE Healthcare Lifesciences) and their DNA sequence determined using an ABI 3730 Genetic Analyzer (Massey University Genome Service).

For comparison, 11 *D. b. minor* sequences stored in GenBank (Accession numbers AF187825 - AF187827 & AF187829 - AF187831, (Brown and Houlden 2000); AY742832 & AY742833 (Fernando et al. 2006)) originally sampled from Zimbabwe and zoos in Australia and the United States were added to the data set, in addition to 20 *D. b. michaeli* samples (Accession numbers AF187834 & AF187835, (Brown and Houlden 2000); AY742830 & AY742831, (Fernando et al. 2006); FJ227484 - FJ227498, (Muya et al. 2011)) originally sampled from Kenya and zoos in Australia and the United States.

Data Analysis.

The 101 mitochondrial DNA sequences were edited by eye and then aligned using Clustal W (Larkin et al. 2007). Homogeneity of base compositions was tested using PAUP 4.0b (Swofford 2002). DnaSP v 5.10.1 (Rozas et al. 2003) was used to calculate haplotype diversity (h), nucleotide diversity (π) and standard deviation (SD) within the subspecies. The level of sequence divergence within and between populations was estimated using a pairwise distance analysis in MEGA 5.1 (Tamura et al. 2011), and standard errors were calculated using a bootstrap procedure. A statistical parsimony haplotype network was calculated with NETWORK 4.610 (Bandelt et al. 1999).

2.4 Results

The sequence of the mtDNA control region was determined for a total of 70 individual black rhinos as follows: *D. b. minor* samples: 50 from HiP, eight from Ithala, five from MGR, one from Ndumo Game Reserve, one from the

Johannesburg Zoo (Accession number JN593089) and 11 sequences from Genbank (Accession numbers AF187826 - AF187831, AY742832 - AY742833 & AF187832 - AF187833); *D. b. michaeli* samples: one from Addo Elephant Park (Accession number JN5930090) and 20 from Genbank (Accession number FJ227483 - FJ227498, AY742830 - AY742831 & AF187834 - AF187835) and four samples for *D. b. bicornis* from Namibia's northern region (Accession numbers JN593091-JN593094) (Table 2.1).

The 101 aligned sequences were 363bp long with 31 polymorphic sites; there was an average pairwise difference of $4\% \pm 1\%$ between *D. b. michaeli* and *D. b. minor* $4.5\% \pm 1.1\%$ between *D. b. michaeli* and *D. b. bicornis* and $2.3\% \pm 0.8\%$ between *D. b. minor* and *D. b. bicornis*. No insertions or deletions were observed.

Considering each subspecies separately, the greatest level of diversity was recorded in *D. b. michaeli* (n=21), which contained 13 haplotypes and showed comparatively high nucleotide diversity ($\pi = 0.011 \pm 0.00106$) and haplotype diversity ($h = 0.958 \pm 0.026$) (Table 2.2). The lowest level of diversity within subspecies was seen in the Namibian *D. b. bicornis* samples (n=4) where only one unique haplotype was found, however this was based on a small sample size and might not represent the total amount of genetic variation within the population. The pooled KZN samples and Genbank sequences of all *D. b. minor* individuals (n=79) contained seven haplotypes and haplotype diversity (h) was 0.267 ± 0.067 and a nucleotide diversity (π) of 0.002 ± 0.00063 . The eight *D. b. minor* Zimbabwe sequences from Brown and Houlden (2000) and two from Fernando et al. (2006) had shared haplotypes (Table 2.3), however there were no shared haplotypes with the KZN samples.

Table 2.1: Rhinoceros subspecies and sources analyzed for mtDNA variation

Subspecies	Sample Size	Sample/Sequence Source	Reference
<i>D.b.minor</i>	50	Hluhluwe-iMfolozi Game Park, KNZ Region, Accession number JN593089	This study
	8	Itala Game Park, KNZ Region	This study
	5	Mkuze Game Park, KNZ Region	This study
	1	Ndumo Game Park, KNZ Region	This study
	1	Johannesburg Zoo	This study
	6	Chete National Park, Zimbabwe, Accession number AF187825 - AF187827 & AF187829 - AF187831	Brown & Houlden 2000
	2	Zambezi Valley, Zimbabwe, Accession numbers AY742832-AY742833	Fernando et al. 2006
	1	Captive born, San Diego Zoo Accession number AF187832	Brown & Houlden 2000
	1	Captive born, Milwaukee Zoo, Accession number AF187833	Brown & Houlden 2000
	1	Captive born, Western Plains Zoo Accession number AF187828	Brown & Houlden 2000
	16	Kenya, Accession numbers FJ227483-FJ227498	Muya 2008
	2	Solio Game Reserve, Kenya, Accession numbers AY742830-AY742831	Fernando et al. 2006
<i>D.b.michaeli</i>	1	Cincinnati Zoo, Accession number AF187834	Brown & Houlden 2000
	1	Taronga Zoo, Accession number AF187835	Brown & Houlden 2000
	1	Addo Elephant Park, South Africa Accession number JN593090	This study
	4	Etoshia, Namibia Accession numbers JN593091-JN593094	This study
	4	HiP & London Zoo Accession numbers AF187836-AF187839	Brown & Houlden 2000

Table 2.2: Mitochondrial DNA D-loop sequence variability within subspecies. Sample size (n), Number of haplotypes (H), haplotype diversity (h), Nucleotide diversity (π), Standard deviation (SD)

	Genetic Variability				π	(SD)
	n	H	h	(SD)		
Subspecies:						
<i>D.b.minor</i> (aggregate)	76	7	0.267	0.067	0.0023	0.001
KZN meta- population	65	1	-	-	-	-
Zimbabwe samples	11	6	0.855	0.085	0.0074	0.00124
<i>D.b.michaeli</i>	21	13	0.952	0.024	0.0112	0.00106
<i>D.b.bicornis</i>	4	1	-	-	-	-

Table 2.3: Summary statistics for the mitochondrial DNA control region sequence variability in each subspecies and haplotype identifiers used in Figure 2.2. Sample size (n), Haplotypes, labeled A - U (H), haplotype diversity (h), Nucleotide diversity (π), Standard deviation (SD), Number of segregating sites (S)

Subspecies & Population:		Genetic Variability					
		H	h	(SD)	π	(SD)	S
<i>D. b. minor</i>	Pooled <i>D. b. minor</i> samples	A - G	0.267	0.067	0.002	0.00063	6
	Hluhluwe-iMfolozi Game Park (KZN)	A	-	-	-	-	-
	Accession number JN593089						
	Itala Game Park (KZN)	A	-	-	-	-	-
	Mkuze Game Park (KZN)	A	-	-	-	-	-
	Ndumo Game Park (KZN)	A	-	-	-	-	-
	Johannesburg Zoo	A	-	-	-	-	-
	Chete National Park, Zimbabwe,	B, C & D	0.733	0.155	0.004	0.00056	4
	Accession number						
	AF187825 - AF187827 & AF1878329 - AF187831						
	Zambezi Valley, Zimbabwe, Accession numbers	E & G	1	0.5	0.008	0.00413	3
	AY742832 - AY742833						
	Captive born, San Diego Zoo	E	-	-	-	-	-
	Accession number AF187832						
	Captive born, Western Plains Zoo	B	-	-	-	-	-
	Accession number AF187828						
	Captive born, Milwaukee Zoo, Accession number AF187833	F	-	-	-	-	-
<i>D. b. michaeli</i>	Pooled <i>D. b. michaeli</i> samples	H - T	0.958	0.026	0.011	0.00106	15
	Kenya, Accession numbers	I - S	0.952	0.031	0.011	0.00119	15
	FJ227483-FJ227498						
	Solio Game Reserve, Kenya	J - T	1	0.5	0.011	0.00413	3
	Accession numbers						
	AY742830-AY742831						
	Cincinnati Zoo,	H	-	-	-	-	-
	Accession number AF187834						
	Taronga Zoo,	I	-	-	-	-	-
	Accession number AF187835						
	Addo Elephant Park, South Africa	T	-	-	-	-	-
	Accession number JN593090						
<i>D.b.bicornis</i>	Pooled <i>D. b. bicornis</i> samples	U	0	-	-	-	-
	Etosha, Namibia	U	-	-	-	-	-
	Accession numbers						
	JN593091-JN593094						

The haplotype network (Figure 2.2) shows a clear pattern of the separation among the three currently recognised subspecies with the KZN population falling out with the *D. b. minor* populations of Zimbabwe. Our finding of no more than three base pair substitutions between adjacent haplotypes within the *D. b. michaeli* subspecies is consistent with Muya et al. (2011). There is significant separation between the *D. b. minor* and *D. b. bicornis* with eight base pair substitutions as well as between *D. b. minor* and *D. b. michaeli* with nine base pair substitutions.

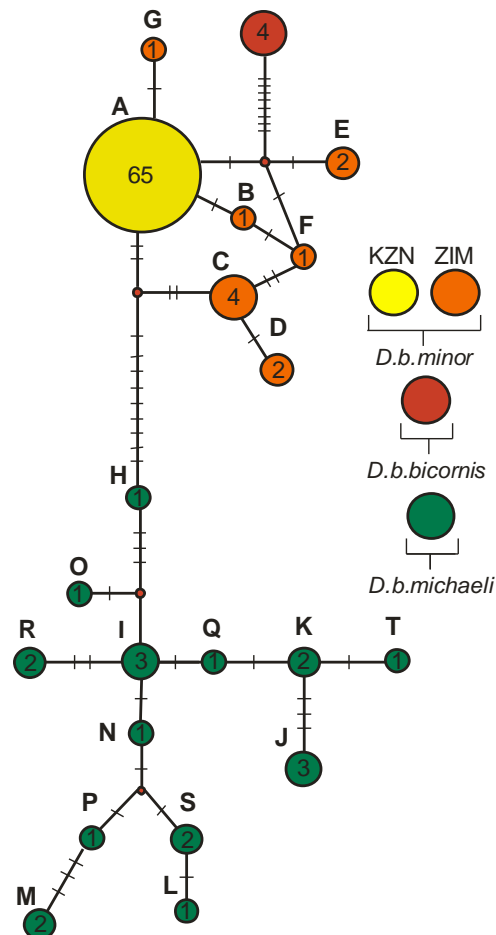


Figure 2.2: Statistical parsimony haplotype network calculated with Network Software for *D. bicornis*. KZN refers to the pooled *D. b. minor* samples within KZN (Ndumo Game Reserve, Ithala Game Reserve, Mkuze Game Reserve, Hluhluwe-iMfolozi Game Park (HiP))

2.5 Discussion

I showed that the KZN population of *D. b. minor* is fixed for a single mtDNA haplotype, like most Sumatran rhino (*Dicerorhinus sumatrensis*) populations (Morales et al. 1997). However, unlike the Sumatran rhino populations that have occupied separate land masses for more than 10,000 years (Morales et al. 1997), it has been widely assumed that the KZN *D. b. minor* population has been separate from other *D. b. minor* populations only recently (i.e., caused by anthropogenic settlement and habitat modification during the 19th century, Swart et al. 1994). The single mtDNA haplotype in KZN *D. b. minor* raises the question of whether the KZN remnant population lost genetic variation recently due to the population bottleneck, or has it been a genetically separate lineage for longer than previously thought?

MtDNA has a smaller effective population size (N_e) compared to nuclear loci and is one of the first genetic markers to show the genetic signature of a demographic decline. The likelihood of two or more mtDNA haplotypes persisting within an isolated population is reduced to $p < 0.1$ over $4N_{ef}$ generations and the population is expected to become monophyletic after $4N_{ef}$ generations (Avice et al. 1984; Mucci et al. 1999). If this holds true for the KZN *D. b. minor* then recent population decline and fragmentation would have increased the rate of drift and might be responsible for lack of haplotype diversity within the KZN black rhinoceros. Examples of monomorphic haplotypes occurring from severe bottlenecks are well documented in several species. For instance, the Whooping Crane (*Grus americana*) once found throughout North America had six haplotypes in 10 pre-bottleneck museum samples, but only one haplotype persisted in the remnant post-bottleneck

population of 14 (Glenn et al. 1999). Such rapid declines in genetic variation have also occurred amongst southern Africa's other large mammals. For example, three small remaining remnant populations of Cape mountain zebra (*Equus zebra zebra*) each contain a single, unique haplotype but larger Namibian populations of closely related Hartmann's mountain zebra (*E. z. hartmannae*) have as many as 11 different haplotypes (Moodley and Harley 2005; Watson and Chadwick 2007). Another case in point is the loss of genetic diversity at mitochondrial and Y-chromosome loci observed in small, managed populations of Cape buffalo in Kenya and Uganda which was attributed to restricted gene flow into protected areas (Van Hooft et al. 2002).

Low genetic variation is not always a consequence of recent anthropogenic fragmentation. An alternative hypothesis is that low levels of mtDNA and autosomal variation are a result of long-term demographic separation, historically small population sizes and local adaptation. For example, despite having lower mtDNA and autosomal DNA variation, there was no evidence of a genetic bottleneck in the Yellowstone National Park, U.S.A. grizzly bear (*Ursus arctos*) population compared to surrounding grizzly bear populations (Miller and Waits 2003). Although Yellowstone's large population is embedded within the species' range, Miller and Waits (2003) attribute the lower genetic variation to restricted gene flow into the area from the north. The common impala (*Aepyceros melampus melampus*) of KZN also exhibited population differentiation from populations in the Limpopo Province just 490 km north. Schwab et al. (2012) attributed the genetic divergence to a narrow zone of unsuitable habitat below the eastern escarpment of the Drakensberg Mountains that impeded dispersal between the two provinces.

Genetic replenishment by restocking and outbreeding is recommended in cases where anthropogenically induced fragmentation has caused a loss in genetic diversity and an increase in genetic divergence. For example, “genetic rescue” has been recommended for the Cape zebra (Moodley and Harley 2005; Watson and Chadwick 2007). However, where differences amongst genetically depauperate populations might be of natural origin, population management may need to take into account local adaptation and the possibility of outbreeding depression. Resolving the question regarding KZN *D. b. minor* mtDNA and autosomal DNA genetic structure being a recent or old event is important for guiding management plans (Rookmaaker 2005).

Microsatellite DNA markers were previously used to assess the levels of genetic variation amongst *D. b. minor* populations. Harley et al. (2005) found appreciable amounts of variation within the *D. b. minor* subspecies using nine microsatellite loci (Table 2.4). They recommended that as long as heterozygosity and allele numbers stayed at ‘current’ levels, no management policy change was necessary.

Based on a survey of 10 microsatellite DNA loci (Table 2.4), Karsten et al. (2011) found low levels of genetic variation within the KZN *D. b. minor*, but concluded that it was not cause for concern. They reached their conclusion based on (1) the similarity of allelic diversity and heterozygosity between the KZN *D. b. minor* population and the other subspecies and (2) a higher level of diversity within the black rhinoceros metapopulation compared to those found in other large African mammals. In their study, H_E estimates for the *D. b. bicornis* and *D. b. michaeli* subspecies (each based on only four samples) were substantially lower than those reported by Harley et al. (2005) (Table 2.4).

Thus, estimates for *D. b. bicornis* and *D. b. michaeli* in Karsten et al. (2011) are probably underestimates. Moreover, comparisons with other large African mammals should be made cautiously. Lions in the Serengeti Plains and Ngorongoro Crater have an H_E of 0.54 and 0.46, on par with black rhinos in Harley et al. (2005), yet unlike the Serengeti Plains lions the Ngorongoro Crater lions have a marked decrease in their reproductive rate attributed to inbreeding depression levels of genetic diversity and differentiation within and among the KZN metapopulation reported by this mtDNA study and published autosomal microsatellite data.

The likelihood of outbreeding depression in supplemented populations of the same species is low if they have the same karyotype, have been isolated for less than 500 years, and occupy similar environments (Frankham et al. 2011). Houck et al. (1995) identified variation in chromosome morphology (number of submetacentric elements) between *D. b. minor* and *D. b. michaeli* zoo samples and recommended further studies to investigate possible differences in geographically separated populations of each subspecies in the wild. Furthermore, twenty-seven KZN *D. b. minor* were translocated to Malilangwe, Zimbabwe in 1997 where they were managed separately and not outbred with any Zimbabwe populations. The translocated population thrived with a growth rate of 8.3% per annum (R. du Toit pers. comm.) cf. 3.4% over a 10-year period (1999 – 2008) in HiP (Clinning et al. 2009). The success of the translocated KZN *D. b. minor* in Zimbabwe alleviated concerns about the adaptability of KZN rhino to Zimbabwe. The only remaining concern is whether or not the populations have been genetically isolated for longer than previously considered.

Table 2.4: Microsatellite results from Harley *et al.* 2005 and Karsten *et al.* 2011. Expected heterozygosity (H_E), Observed heterozygosity (H_O)

Study	Microsatellites Used	Subspecies	Geographic region	Sample Size	H_E	H_O
Harley <i>et al.</i> 2005	BR4, BR6, BR17	<i>D. b. minor</i>	South Africa and Zimbabwe	46	0.46	0.436
	(Cunningham <i>et al.</i> 1999)	<i>D. b. bicornis</i>	Namibia	53	0.51	0.523
	DB1, DB14, DB44, DB49, DB52, DB66 (Brown & Houlden 1999)	<i>D. b. michaeli</i>	South Africa	19	0.68	0.731
Karsten <i>et al.</i> 2011	BR4, BR6, BR17	<i>D. b. minor</i>	South Africa	77	0.44	0.38
	(Cunningham <i>et al.</i> 1999)	<i>D. b. bicornis</i>	Namibia	4	0.43	0.46
	DB1, DB14, DB49, DB66 (Brown & Houlden 1999)	<i>D. b. michaeli</i>	South Africa and Tanzania	4	0.54	0.54
	AY606078, AY606080, AY606083 (Nielsen <i>et al.</i> 2008)					

I recommend five research tasks to assist in resolving the genetic structure of southern Africa's black rhino as a guide to future management. (1) Determine historic levels of genetic variation using museum or collection samples. (2) Investigate whether there is evidence of inbreeding depression within the HiP and KZN metapopulation. (3) Conduct a karyotype analysis on *D. b. minor* in KZN and Zimbabwe to determine whether chromosomal differences exist. (4) Increase the mtDNA sample size of the Zimbabwe *D. b. minor* population. Considering the high level of variation in the small sample size of the Zimbabwe sequences, a larger sample size of *D. b. minor* from that region might show that the KZN haplotype (A) is also there. (5) Lastly, genetic supplementation experiments should be implemented cautiously and systematically. A mixed population should be founded with at least 20 animals as suggested by du Toit (2006a), perhaps using the Malilangwe, Zimbabwe translocation event as a template or more recent guidelines (Linklater et al. 2012; Linklater et al. 2011). The translocated KZN *D. b. minor* rhinos in Malilangwe have not yet been outbred with the Zimbabwe rhinos (R. du Toit pers. comm.) but might be with the F1 and F2 offspring carefully monitored for signs of reduction in reproductive fitness (outbreeding depression). If the research tasks we have recommended are completed and there is evidence of historic gene flow between KZN and Zimbabwe *D. b. minor* and no signs of outbreeding depression in the experimentally mixed population, then KZN *D. b. minor* is a candidate for genetic supplementation using progeny from Zimbabwe populations.

2.6 Acknowledgements

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CHAPTER THREE

Low levels of microsatellite DNA variation and possible management considerations for black rhino (*Diceros bicornis minor*) in KwaZulu-Natal, South Africa



Photo by Rosalynn Anderson-Lederer

3.1 Abstract

When preparing management plans based on genetic information, it is helpful to validate results with discrete tests to confirm outcomes. Mitochondrial DNA (mtDNA) combined with microsatellite DNA markers can assist managers in making conservation decisions based on understanding the genetic structure of a species. Previous studies indicated that the expected heterozygosity in the black rhino (*Diceros bicornis minor*) metapopulation of KwaZulu-Natal (KZN) including the source population of Hluhluwe-iMfolozi (HiP), South Africa is within the range of other large animals across Africa and no changes in management policy were necessary. However, recent mtDNA findings of one unique haplotype (n=65) in the KZN metapopulation compared to six haplotypes (n=11) in native Zimbabwe *D. b. minor* suggest otherwise. I used 10 microsatellites and found that the KZN metapopulation was out of Hardy-Weinberg Equilibrium and showed excess homozygosity at five loci. I confirmed mtDNA findings that the South African metapopulation has lower genetic variation than the native Zimbabwe *D. b. minor* population indicating that current conservation plans might need to be modified to prevent further genetic decay. A loss of genetic diversity might be arrested by either (1) increasing population numbers to accommodate needed growth by expanding habitat and reserve sizes, or (2) carrying out a serial translocation scheme between the metapopulation of smaller populations including the source population Hluhluwe-iMfolozi and native *D. b. minor* from Zimbabwe to generate an artificially larger single population. Implementing these recommended changes could help reduce further genetic loss and maintain the levels of genetic variability

in the HiP source and KZN metapopulation.

3.2 Introduction

Understanding the genetic structure of wild populations provides conservation managers with valuable insight into the design of management plans for reintroduction and supplementation. Levels of genetic variability are known to vary among populations, making it a perfect tool for determining the underlying structure of a natural population. While not always feasible, the level of genetic variability should be quantified using a range of DNA markers, and decisions should be based on corroborated results (Moritz 1994; Waits et al. 1998; Manceau et al. 1999). For example, recommended guidelines for the management of the Scandinavian brown bear (*Ursus arctos*) (Taberlet and Bouvet 1994; Waits et al. 2000) and western North American caribou (*Rangifer tarandus*) (Weckworth et al. 2012) are the consensus of mtDNA and microsatellite data.

Microsatellite and mtDNA are complimentary metrics for examining the genetic structure and the level of diversity of a populations (Toews and Brelsford 2012). MtDNA is maternally inherited and has one quarter of the effective size (N_e) of a nuclear diploid locus, which makes it more sensitive to changes in population size. Microsatellite DNA markers, on the other hand, enable a better coverage of the genome and give a more precise estimate of the level of genetic variation in a population. Typically highly polymorphic, microsatellites are used extensively in genetic studies (Bruford and Wayne 1993; Jarne and Lagoda 1996; Forstmeier et al. 2012).

African black rhino (*Diceros bicornis*) once ranged across the African continent in large numbers (Ashley et al. 1990; Lacomat 2005), but are now endangered (Emslie 2011). The genetic structure of the species has been extensively studied (ALLOYZMES: Merenlender et al. (1989); Ashley et al. (1990); Swart et al. (1994), MTDNA SEQUENCING: Brown and Houlden (2000); Muya et al. (2011); Anderson-Lederer et al. (2012) MICROSATELLITES: Brown and Houlden (1999); Cunningham et al. (1999); Garnier et al. (2001); Harley et al. (2005); Karsten et al. (2011); Muya et al. (2011)), yet confusion remains regarding subspecific nomenclature and the grouping of subspecies into 'ecotypes' (Zukowsky 1965; Groves 1967; du Toit 1986, 1987; Rookmaaker 1995, 2005).

Hluhluwe-iMfolozi Game Park (HiP) (Fig. 3.1) in the KwaZulu-Natal (KZN) province of South Africa has the largest remnant population of critically endangered (IUCN 2008) *D. b. minor* (n~220; Clinning et al. 2009). Successful translocations of *D. b. minor* from HiP to other KZN reserves began in 1962, later expanding to other South African provinces and African nations (e.g., Zimbabwe, Zambia, Swaziland) (Hitchins 1984; Emslie et al. 2009). While translocations have resulted in expansion and growth of the KZN *D. b. minor* metapopulation, the HiP source remains small, with no translocations back into the population, making it vulnerable to loss of genetic variation. Harley et al. (2005) and Karsten et al. (2011) demonstrated that *D. b. minor* subspecies had lower microsatellite variability than the other two subspecies (*D. b. michaeli* and *D. b. bicornis*). More specifically, Anderson-Lederer et al. (2012) established that the KZN metapopulation of *D. b. minor* are fixed for a single unique haplotype whereas six haplotypes (n=11) were

identified in native Zimbabwe *D. b. minor*. The mtDNA results could be used to imply that genetic variability may be decreasing within HiP.

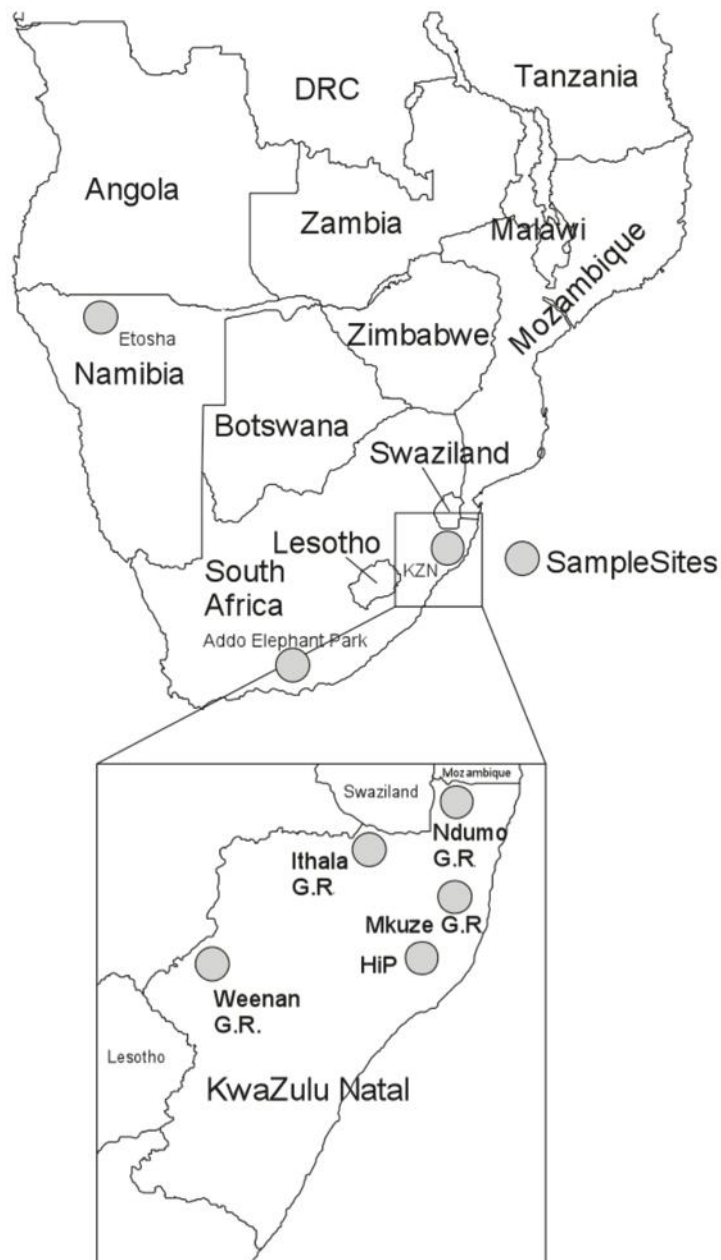


Figure 3.1: Map of southern Africa with sample sites indicated by circles. Inset of KwaZulu-Natal with sample sites Hluhluwe-iMfolozi Game Park (HiP), Mkuze Game Reserve, Ndumo Game Reserve, Ithala Game Reserve and Weenan Game Reserve.

Table 3.1: Characteristics of 10 microsatellite markers used to type 127 black rhinoceros and the number of alleles found in 118 KZN *D. b. minor*

Locus ID	Repeat Size	Primer pair sequence 5'-3'	Author	Tag	Size (bp)	Annealing Temp (C°)	No. of alleles
BR4	(CA)19	(F) CCC CTA AAT TCT AGG AAC AC (R) CCA AAG ACC ACC AGT AAT TC	(Cunningham <i>et al.</i> 1999)	M-13 VIC	124-146	49	5
BR6	(CA)15	(F) TCA TTT CTT TGT TCC CCA TAG CAC (R) AGC AAT ATC CAC GAT ATG TGA AGG	"	M-13 VIC	126-158	51	4
BR17	(AT)6(GT)18	(F) ACT AGC CCT CCT TTC ATC AG (R) GCA TAT TGT AAG TGC CCC AG	"	Fluoro VIC	123-135	60	3
DB1	(CA)14	(F) AGA TAA TAA TAG GAC CCT GCT CCC (R) GAG GGT TTA TTG TGA ATG AGG C	(Brown & Houlden 1999)	Fluoro FAM	121-127	60	2
DB5	(CA)13	(F) GAC CCC CAT GTT CAC TGC (R) AGG TCC ATC CAT TTT GTC CC	"	Fluoro FAM	185-204	60	4
DB30	(CA)21	(F) GCG ACT ATG ACA TAC AAC TAT CTA C (R) GGT CA AGG ATT ATT CTG ACT AGC	"	Fluoro VIC	201-205	64	4
DB44	(CA)4G(CA)16	(F) GGT GGA ATG TCA AGT AGC GG (R) CTT GTT GCC CCA TCC CTG	"	M-13 VIC	170-184	64	1
DB49	(CA)14	(F) GTC AGG CAT TGG CAG CAA G (R) CAG GGT AAG TGG GGG TGC	"	Fluoro FAM	152-162	64	4
DB52	(CA)21	(F) CAT GTG AAA TGG ACC GTC AGG (R) ATT TCT GGG AAG GGG CAG G	"	Fluoro FAM	210-220	64	3
DB66	(CA)7TA(CA)16	(F) CCA GGT GAA GGG TCT TAT TAT TAG C (R) GGA TTG GCA TGG ATG TTA CC	"	Fluoro FAM	187-205	58	3

In light of recent mtDNA findings, re-examination of the levels of variation using microsatellite markers in the KZN *D. b. minor* metapopulation and more specifically the HiP source population is required. The aim of this study was to use ten microsatellite DNA markers (Table 3.1) to investigate levels of heterozygosity and allelic diversity in HiP *D. b. minor*. I then compared these results to the KZN metapopulation and previously published microsatellite and sequenced mtDNA control region data. I used the results to make recommendations for translocation, reintroduction and supplementation for KZN's *D. b. minor* source and metapopulation.

3.3 Methods

Sampling

Samples of blood and/or ear tissue were collected from a total of 127 individuals of *D. b. minor* in the KZN province in South Africa, *D. b. michaeli* in Addo Elephant

National Park and *D. b. bicornis* in Namibia's Northern Region (Table 3.2). The samples were acquired opportunistically during routine translocation and ear notching (for identification) events from 2002 to 2009. Blood and tissue samples were stored in cryovials containing 1mL of DMSO/EDTA/Tris/salt solution (Seutin et al. 1991) or RNAlater[®] Solution (Life Technologies).

Table 3.2: Subspecies of *Diceros bicornis* and corresponding populations and sample size (N). All *D. b. minor* samples are from the KwaZulu-Natal metapopulation

Subspecies	Population	N
<i>D. b. minor</i>	Eastern Shores	2
	Hluhluwe-iMfolozi Park	97
	Ithala Game Reserve	10
	Johannesburg Zoo	1
	Mkuze Game Reserve	6
	Ndumo Game Reserve	1
	Tembe Elephant Park	1
	KZN Populations Combined	118
<i>D. b. michaeli</i>	Addo Elephant Park, South Africa	3
<i>D. b. bicornis</i>	Waterberg National Park, Namibia	6

Genetic Analyses

I extracted genomic DNA using DNeasy kits (Qiagen Inc.) following the manufacturer's recommended protocol. Ten polymorphic microsatellite loci (Table 3.1) reported by Cunningham et al. (1999) and Brown and Houlden (1999) were chosen based on their reliability and were amplified by polymerase chain reaction

(PCR). One primer from each pair was labeled using the M-13 tag methodology (Schuelke 2000) or directly labeled with a fluorescent dye (6-FAM or VIC, Invitrogen). The resultant PCR products were analysed on a 3730 automated sequencer using the GS-500 LIZ size standard and the GENESCAN software (Applied Biosystems). Samples that did not amplify for all loci were removed from the data set. Alleles were visualized and analysed using GENEMAPPER software ver. 3.7 (Applied Biosystems), then results were then confirmed with GENEMARKER software (Softgenetics). MICROCHECKER ver. 2.2.0.3 (Van Oosterhout, 2004) assessed possible reasons for deviation of HWE, which include null alleles (one or more alleles that fail to amplify during PCR), large allele dropout (small alleles amplify better than large alleles) and scoring errors due to stutter (slight changes that occur in the allele sizes during PCR).

Hardy-Weinberg Equilibrium (HWE) (non-random association of alleles within diploid individuals), linkage disequilibrium (non-random association of alleles at different loci) and heterozygote excess and deficiency were estimated using GENEPOP ver. 1.2 (Raymond and Rousset 1995; Rousset 2008). The Markov Chain parameters for the locus-by-locus pair-wise tests for gametic disequilibrium utilized 1000 dememorizations, 100 batches, and 5000 iterations per batch. Statistical significance (P-value) was corrected for multiple testing using the False Discovery Rate (FDR) procedure (Benjamini and Hochberg 1995). ARLEQUIN ver. 3.5 (Excoffier et al. 2005) was used to assess the level of population differentiation between the three subspecies based on Wright's (1965) pairwise F_{ST} , which is derived from the variances of allele frequency and Slatkin's (1995) R_{ST} , which

calculates the fraction of total variance of allele size that exists between populations (Balloux and Lugon-Moulin 2002). The level of significance was assessed at 1000 permutations. I also examined D_{est} (Jost 2008), which is an estimate for actual differentiation, with SMOGD ver. 1.2.5 (Crawford 2010) using 1000 bootstrap replicates. Allelic richness (A_r , a measure of the number of alleles corrected for different sample sizes) was calculated for each subspecies and each loci with HP-Rare ver. 1.0 (Kalinowski 2005). Inbreeding coefficient F_{IS} was analysed using FSTAT (Goudet 1995).

Genetic Structure

STRUCTURE ver. 2.3.3 (Pritchard et al. 2000), a Bayesian model-based clustering software implementing the Markov Chain Monte Carlo (MCMC) method was used to identify distinct genetic patterns in the subspecies populations and designate individuals to one or more genetic clusters (K). One potential criticism of STRUCTURE is that the output of this programme can be difficult to interpret when levels of population structure are low. Therefore data was analysed using the LOCPRIOR setting within STRUCTURE, which uses the sampling locations of individuals to assist the clustering process, thereby generating more accurate estimates of K (Hubisz et al. 2009). Ten STRUCTURE runs for each value of K were carried out (one to five for the *D. bicornis* subspecies and one to eight for the KZN *D. b. minor* metapopulation) for 1000000 iterations and a burn-in time of 100000 iterations for both data sets. Since the three subspecies sampled may have mixed ancestry, admixture ancestry model was chosen (Pritchard et al. 2007). Allele frequencies were correlated among populations and assumed different values of F_{ST} for the different

subpopulations. STRUCTURE HARVESTER ver. A.1 (Earl and vonHoldt 2012) which applies the Evanno method (Evanno et al. 2005) was used to visualize STRUCTURE output.

Testing for a Genetic Bottleneck

I examined signatures of a reduction in population size using BOTTLENECK ver. 1.2.2 (Cornuet and Luikart 1996; Luikart et al. 1998; Piry et al. 1999). This analysis is designed to detect a recent bottleneck occurring within the past $2N_e - 4N_e$ generations, assuming the populations were in mutation-drift equilibrium. Piry et al. (1999) recommends the Wilcoxon two-tailed sign-rank test within BOTTLENECK, which accounts for both heterozygosity excess and deficiency for data with less than 20 polymorphic loci and where effective population sizes may have remained constant for long periods of time. The three mutation models within the Wilcoxon test are the infinite allele model (IAM), the stepwise mutation model (SMM) and the two-phase model (TPM); microsatellite mutation was set at 95% single-step mutation rate and 5% multiple step mutation along with the variance among multiple steps of 12 (Piry et al. 1999). BOTTLENECK also tested for a mode-shift of the allele frequency distribution, because when a population has recently been through a bottleneck, rare alleles are typically lost causing a distortion in allele frequencies at selectively neutral loci (Luikart et al. 1998). Since Harley et al. (2005) found a significant departure from HWE and a slight overall homozygous excess in their mixed Zimbabwe/South Africa samples of *D. b. minor* (n=46), I also checked for a bottleneck using the Garza-Williamson index or M-ratio (Garza and Williamson 2001) implemented in ARLEQUIN ver. 3.5 (Excoffier et al. 2005). The M-ratio ($M =$

k/r where k = number of alleles and r = overall range in fragment sizes) can identify a bottleneck even when data is out of equilibrium and a modified M-ratio is used if any of the loci are monomorphic.

3.4 Results

All ten microsatellite loci were amplified successfully. There was evidence of null alleles at loci DB5 and DB1 in the KZN *D. b. minor* samples (n=118). The two loci were removed for KZN metapopulation comparisons, but neither loci was excluded from the data set for comparison between the three subspecies, since null alleles were not present in either *D. b. micheali* or *D. b. bicornis*. Results for *D. b. bicornis* samples (n=6) were monomorphic at two loci and there was no evidence for scoring error due to stuttering, large allele dropout or null alleles. The *D. b. micheali* samples (n=3) had too few alleles at each locus to perform the same tests. No significant linkage disequilibrium was observed for any pairs of loci after FDR correction.

The KZN *D. b. minor* population (n=118) was not in HWE. Five of the ten loci showed deviation from HWE (DB30, DB1, BR17, DB5 and DB52) (Table 3.3) and locus DB44 was monomorphic. Three loci showed deviation from HWE after an FDR correction (DB30, DB5 and DB52). The *D. b. micheali* samples (n=3) were in HWE, but were monomorphic at two loci (DB1 & DB44) (Table 3.3). The *D. b. bicornis* samples (n=6) were in HWE but monomorphic at one locus (BR4) (Table 3.3).

Table 3.3: H_e and H_o per locus, per subspecies with associated P-values for *D. b. bicornis* (n = 6), *D. b. michaeli* (n = 3), KZN *D. b. minor* (n = 118) and native Zimbabwe *D. b. minor* (n = 35). Zimbabwe data are from Garnier et al. (2001). Bold data highlight regional variations between *D. b. minor* subspecies. (M: monomorphic, no value)

Locus	<i>D. b. bicornis</i>			<i>D. b. michaeli</i>			KZN <i>D. b. minor</i>			Zimbabwe <i>D. b. minor</i>	
	H_e	H_o	P-value	H_e	H_o	P-value	H_e	H_o	P-value	H_e	H_o
DB30	0.62	1.00	1.00	0.33	0.33	-	0.63	0.60	0.002	not tested	
DB1	0.59	0.67	0.76	M	-	-	0.33	0.25	0.009	0.46	0.69
BR17	0.41	0.50	1.00	0.60	0.33	0.20	0.20	0.17	0.029	0.61	0.69
DB66	0.74	0.83	0.19	0.93	1.00	1.00	0.64	0.66	0.598	0.72	0.91
DB5	0.67	1.00	1.00	0.73	0.33	0.21	0.64	0.52	0.001	0.58	0.69
BR6	0.32	0.33	1.00	0.87	0.67	0.33	0.55	0.56	0.768	0.80	0.81
BR44	0.68	0.33	0.05	M	-	-	M	-	-	0.47	0.47
BR4	M	-	-	0.87	1.00	1.00	0.66	0.58	0.109	0.60	0.69
DB49	0.64	0.20	0.50	0.73	1.00	1.00	0.55	0.58	0.288	0.70	0.81
DB52	0.53	0.50	0.76	0.80	0.67	0.46	0.49	0.50	0.003	0.77	0.91
mean	0.52			0.59			0.47			0.62	

The F_{ST} results indicated differentiation between the three subspecies over the 10 microsatellite loci examined with values ranging from 0.091 - 0.20 (Table 3.4). F_{ST} results for the three largest sample sets within the KZN metapopulation (HiP, Mkuze & Ithala) indicated very little differentiation with the KZN samples with values between 0.001 – 0.03 (Table 3.4). R_{ST} results for the three subspecies were lower than those reported for F_{ST} and ranged from 0.02 – 0.10 (Table 3.4) suggesting lower genetic differentiation between the subspecies. Allelic differentiation as expressed by D_{est} was minimal between the three subspecies ranging between 0.05 and 0.10 (Jost 2009)(Table 3.4).

Table 3.4: A) Pairwise F_{ST} values B) Pairwise R_{ST} values C) Pairwise D_{est} values for *D. bicornis*

A) F_{ST}		
	<i>D. b. michaeli</i>	<i>D. b. minor</i>
<i>D. b. minor</i>	0.09094 (p = 0.01802 ± 0.0121)	
<i>D. b. bicornis</i>	0.20116 (p = 0.00301 ± 0.0091)	0.19101 (p = 0.000 ± 0.000)
B) R_{ST}		
	Mkuze	HiP
HiP	0.002044 (p = 0.11712 ± 0.0237)	
Ithala	0.001111 (p = 0.30631 ± 0.0388)	0.02846 (p = 0.02703 ± 0.0194)
C) D_{est}		
	<i>D. b. michaeli</i>	<i>D. b. minor</i>
<i>D. b. minor</i>	0.04675	
<i>D. b. bicornis</i>	0.06652	0.09656

H_E averaged over all 10 loci for the three subspecies were between 0.47 ± 0.22 and 0.59 ± 0.35 , while H_0 were between 0.49 ± 0.17 and 0.67 ± 0.31 (Table 3.5). H_E and H_0 for KZN source populations were as follows: HiP = 0.45 ± 0.24 and 0.48 ± 0.23 ; Mkuze; 0.46 ± 0.22 and 0.49 ± 0.25 respectively. When loci DB1 and DB5 were removed from the KZN source populations for analysis (HiP and Mkuze) H_E was only slightly changed: HiP = 0.45 ± 0.25 ; Mkuzi = 0.43 ± 0.23 . H_E and H_0 for the *D. b. minor* Zimbabwe data published by Garnier et al. (2001) was 0.62 ± 0.13 and 0.72 ± 0.13 respectively (Table 3.5). A_r for the three subspecies was between 2.18 and 2.90 (Table 3.5) while F_{IS} was between -0.032 ± 0.46 and 0.054 ± 0.12 (Table 3.5).

Table 3.5: Microsatellite results from this study, Harley *et al.* 2005, Karsten *et al.* 2011, Garnier *et al.* 2001, Muya *et al.* 2011 and Van Coeverden de Groot *et al.* 2011. N, number of individuals analysed; H_E , expected heterozygosity (for polymorphic loci); H_O , observed heterozygosity (for polymorphic loci); SD, standard deviation; A_r , allelic richness (averaged over loci); F_{IS} , inbreeding coefficient.

Study	Microsatellites Used	Country	Subspecies	N	H_E	SD	H_O	SD	A_r	F_{IS}	SD
This study	BR4, BR6, BR17	South Africa	<i>D. b. minor</i>	118	0.47	±0.22	0.49	±0.17	2.18	0.054	±0.12
	(Cunningham <i>et al.</i> 1999)	Namibia	<i>D. b. bicornis</i>	6	0.52	±0.22	0.60	±0.30	2.34	-0.032	±0.46
	DB1, DB5, DB30, DB44, DB49,	South Africa	<i>D. b. michaeli</i>	3	0.59	±0.35	0.67	±0.31	2.90	0.111	±0.39
	DB52, DB66 (Brown & Houlden 1999)										
Harley <i>et al.</i> 2005	BR4, BR6, BR17	South Africa & Zimbabwe	<i>D. b. minor</i>	47	0.46	-	0.44	-		-	-
	(Cunningham <i>et al.</i> 1999)	Namibia	<i>D. b. bicornis</i>	53	0.51	-	0.52	-		-	-
	DB1, DB14, DB44, DB49,	South Africa	<i>D. b. michaeli</i>	19	0.68	-	0.73	-		-	-
	DB52, DB66 (Brown & Houlden 1999)										
Karsten <i>et al.</i> 2011	BR4, BR6, BR17	South Africa & Zimbabwe	<i>D. b. minor</i>	77	0.44	-	0.38	-		0.14	-
	(Cunningham <i>et al.</i> 1999)	Namibia	<i>D. b. bicornis</i>	4	0.43	-	0.46	-		0.09	-
	DB1, DB14, DB49, DB66	South Africa & Tanzania	<i>D. b. michaeli</i>	4	0.54	-	0.54	-		0.16	-
	(Brown & Houlden 1999) AY606078, AY606080, AY606083 (Nielsen <i>et al.</i> 2008)										
Garnier <i>et al.</i> 2001	BR4, BR6, BR17 (Cunningham <i>et al.</i> 1999) DB1, DB5, DB23, DB44, DB49, DB52, DB66 (Brown & Houlden 1999)	Zimbabwe	<i>D. b. minor</i>	35	0.616	-	0.726	-		-	-
Muya <i>et al.</i> 2011	BR4, BR6, BR17 (Cunningham <i>et al.</i> 1999) DB1, DB5, DB30, DB44, DB52, DB66 (Brown & Houlden 1999)	Kenya	<i>D. b. michaeli</i>	145	0.69	±0.03	0.70	±0.09		0.046	±0.09
Van Coeverden de Groot <i>et al.</i> 2011	DB1, DB44, DB52, DB66 (Brown & Houlden 1999) BR4, BR6, BR17 (Cunningham <i>et al.</i> 1999) BIRh2B, BIRh37D (Van Coeverden de Groot <i>et al.</i> 2011)	Namibia	<i>D. b. bicornis</i>	144	0.51	-	0.52	-		-	-

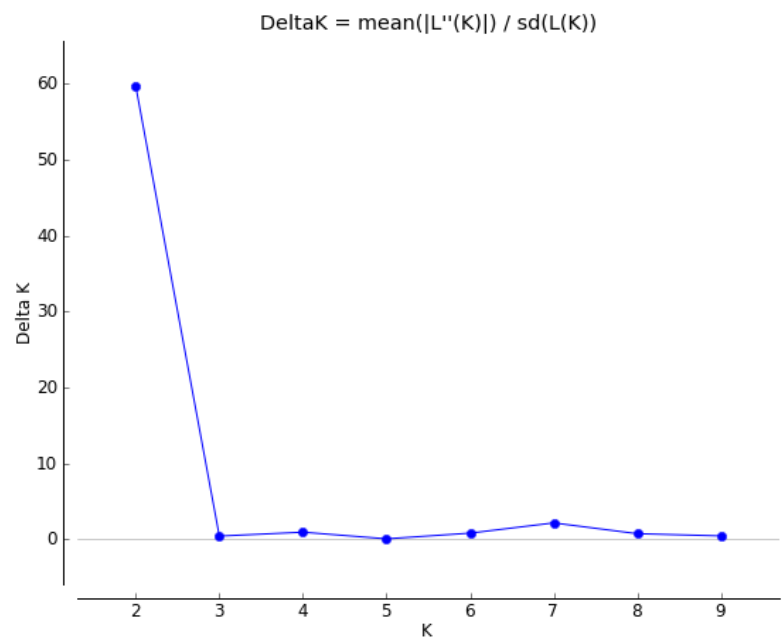
Genetic Structure

The STRUCTURE analysis indicated, the maximum mean log likelihood value of -2094.54 for the three *D. bicornis* subspecies was most likely $K = 2$ (-2195.10 for $K = 1$ and -2106.98 for $K = 3$). Utilizing the Evanno method (Evanno et al. 2005), the highest value of ΔK also indicated that the number of clusters was likely $K = 2$ (Fig. 3.2A). The maximum mean log likelihood value of -1776.88 indicated that the number of clusters for three largest populations in KZN (HiP, Mkuze and Ithala) was $K = 1$ (-1808.80 for $K = 2$), utilizing the Evanno method (Evanno et al. 2005), the highest value of ΔK inferred that the number of clusters was $K = 2$ (Fig. 3.2B). However, ΔK is based on the second order rate of change with respect to the likelihood associated with K and is not a suitable method for detecting if the true K of a population is $K = 1$ (Evanno et al. 2005).

Bottleneck

Wilcoxon two-tailed sign-rank test for bottleneck gave conflicting results. *D. b. minor* showed a classic L-shaped allele frequency distribution, and both the SMM and TPM were both in mutation–drift equilibrium (no bottleneck detected), however the IAM was out of equilibrium (indicating bottleneck). The *D. b. michaeli* and *D. b. bicornis* populations could not be assessed, as there were too few samples for the Wilcoxon and mode-shift tests. Modified M-ratio (Excoffier et al. 2005) results indicated that all three populations have been through significant bottlenecks: *D. b. minor* 0.26494, *D. b. michaeli* 0.22761 and *D. b. bicornis* 0.25238 (results <0.7 indicate bottleneck; Excoffier et al. 2005). Not all of the loci in this study were polymorphic, so the results of the modified M-ratio were reported.

A



B

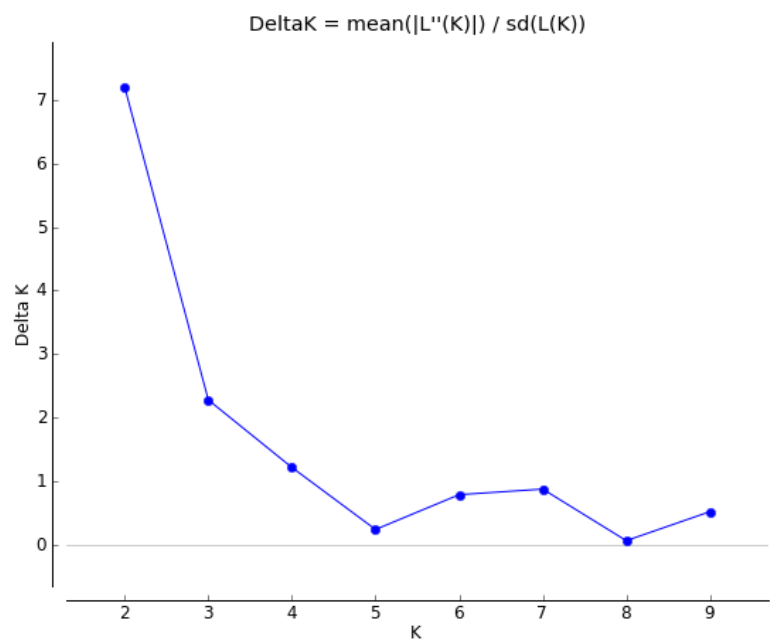


Figure 3.2: STRUCTURE population genetic structure output for *D. bicornis*. Determination of the number of clusters using ΔK for values of K from 1 to 10. A) Three *D. bicornis* subspecies, K = 2. B) KZN sample set, K = 2.

3.5 Discussion

The KZN *D. b. minor* population had an excess of homozygotes at five out of ten microsatellite loci and deviated from HWE expectations. This was consistent (excess homozygotes) with those reported by Harley et al. (2005) whose *D. b. minor* samples were a combination of both South Africa and Zimbabwe individuals. Examination of eight out of 10 of the same microsatellite loci (Table 3.3) as Harley et al. (2005) and ~81% of their samples collected from KZN are probable contributing factors to the similar outcomes.

The R_{ST} values for this data set should be viewed with skepticism since R_{ST} does not perform well with a small sample size (≤ 10) or a small number of loci (≤ 20), as is the case with this study (Gaggiotti et al. 1999). In addition a histogram of allele sizes for the largest population (HiP; not shown) revealed multiple peaks, indicating deviation from the assumption of stepwise mutation, further indicating that R_{ST} is probably not a suitable measure for this data set. Although based on the IAM, F_{ST} outperforms R_{ST} in cases such as this where sample sizes are small (Gaggiotti et al. 1999). F_{ST} and D_{est} results suggest differentiation at the subspecies level, but very low differentiation at the KZN metapopulation level.

Identifying the cause of homozygote excess and departure from HWE may be difficult. The most commonly reported reasons a population can have excess homozygotes at microsatellite loci include scoring errors, the presence of null (non-amplifying) alleles (Pemberton et al. 1995), sampling more than one population (i.e., the Wahlund effect) and inbreeding (Castric et al. 2002). The possibility of scoring errors was eliminated by MICROCHECKER (Van Oosterhout, 2004). Samples that did not amplify for all loci were removed from the analysis.

All remaining individuals in the data set amplified for all loci, indicating that there were no homozygotes for a null allele. This implies that the null alleles were only present at a low frequency and should not have significantly contributed to deviation from HWE.

Departure from HWE may also be present in a population with a low number of effective breeders (reproductive skew) (Luikart and Cornuet 1999). Garnier et al. (2001) found a high reproductive skew (~53%) in Zimbabwe *D. b. minor* that might be related to spatial distribution and linked to variations in fertility levels in each sex. A reproductive skew would affect the level of genetic variation in a small population more than it would in a larger population, especially if only a small number of males (as was the case in Zimbabwe black rhino) are contributing to reproductive output (Garnier et al. 2001).

STRUCTURE results suggest no significant population subdivision between the three subspecies and less within the KZN metapopulation. However, Rodriguez-Ramilo and Wang (2012) advised using caution interpreting output from STRUCTURE. Closely related individuals should be removed from datasets before conducting analyses, otherwise Hardy-Weinberg and linkage disequilibrium may skew results of genetic structure of the population. Unfortunately, there was no way of controlling the samples for this parameter since no pedigrees or studbooks are kept on the wild populations. I do however, have more confidence in the KZN STRUCTURE results than for the subspecies since the subspecies sample size for *D. b. michaeli* was only $n = 3$.

The indeterminate *D. b. minor* BOTTLENECK results might be attributed to testing less than 20 loci. Increasing the number of loci may provide discernable results, but since BOTTLENECK was designed to detect a recent bottleneck within

the past $2N_e$ - $4N_e$ generations, the IAM results are likely indicative of a bottleneck occurring more than $4N_e$ generations ago.

Loss of Genetic Variability within HiP

Reductions in microsatellite variation and bottleneck signature were not unexpected given that black rhino across Africa suffered rapid geographic and population size declines over the last century. However, the genetic variation of the KZN *D. b. minor* was significantly lower than that of the native Zimbabwe *D. b. minor* (KZN: H_E 0.47 ± 0.22 ; Zimbabwe: H_E 0.62 ± 0.13) (Table 3.3 & 3.5), which could be a consequence of low population numbers persisting for many generations (Harley et al. 2005). Although the use of translocations of *D. b. minor* between game reserves in KZN has aided in the retention of current levels of genetic variation (Karsten et al. 2011), there are no translocations of *D. b. minor* into the HiP source population. Lower microsatellite variability coupled with a fixed mtDNA haplotype (Anderson-Lederer et al. 2012), especially in HiP may signal a need for management to intervene to prevent further genetic decay within this valuable source population. If the level of genetic variability continues to decline without intervention (e.g. genetic rescue), it could lead to a reduction in adaptability (evolutionary potential) and increase the risk of inbreeding depression in HiP (Lacy 1987b; Burger and Lynch 1995). This has occurred in other species such as the black-footed rock-wallaby island populations (BFRW) (*Petrogale lateralis*) (Eldridge et al. 1999) and harbour seals (*Phoca vitulina*) (Coltman et al. 1998).

Swart and Ferguson (1997) speculated that native Zimbabwe black rhino populations were the only *D. b. minor* to retain pre-bottleneck levels of genetic

variation. It is unclear why the small native Zimbabwe populations have retained genetic variation through a severe bottleneck while KZN black rhino have not. To prevent a further loss of genetic variation and increasing risk of inbreeding depression within KZN but specifically HiP, the following steps could be taken: (1) rapidly increasing population numbers by increasing reserve sizes or (2) serial translocations amongst the KZN metapopulation reserves, including back into the HiP source population, perhaps including replenishment using native Zimbabwe *D. b. minor*.

Population Increases through Land Acquisitions and Serial Translocations

The recommendation by Emslie (2001) for rapid growth as a buffer against black rhino poaching would also apply to precluding the effects of low genetic variability. Avoidance of a loss of allelic variation through rapid expansion has been documented in other animals. Despite only 13 European rabbits (*Oryctolagus cuniculus*) being imported to Australia in 1859, there was no significant genetic difference between the contemporary introduced rabbit population and the population of European rabbits in France (Zenger et al. 2003). Researchers suspect that the initial population did not experience a decrease in levels of genetic variation because at no time were there enough generations at small sizes to lose significant diversity.

While HiP could increase in size by connecting adjacent reserves in northern KZN, the area of land in question may not be sufficiently large enough for increasing population numbers to levels necessary to arrest the loss of genetic variation. Mkuze Game Reserve (MGR) is connected via a corridor to Greater St. Lucia Wetland Park that extends along the east coast from Kosi Bay

south to Mapelane encompassing Eastern Shores (one of this study's sample sites, Figure 3.4). Connecting MGR (including Greater St. Lucia Wetland Park from Sodwana National Park south to Mapelane) with close neighbours Phinda, Thanda, and founder populations established by the Black Rhino Range Expansion Project (Pongola, Mnyawane and Zululand Game Reserves) would create an area that is very roughly 2,806 km². Adding HiP (~40km away from Zululand Game Reserves) into the conglomeration of reserves by establishing a corridor through existing subtropical fruit and sugarcane fields would increase the area to ~3,745 km². Using the average black rhino/km² of 0.22 for northern KwaZulu-Natal reserves (Adcock, K. pers. comm.) the approximate carrying capacity (CC) for an area that size would be about 824, almost doubling the 430 CC for HiP. Procuring enough land for black rhino management to increase HiP to a size that would allow for rapid growth is unrealistic (Goodman 2001). A compromise to increasing land area of HiP may be found in serial translocations between HiP and other reserves to replicate immigration and emigration for each reserve. Translocations from HiP have been used as a successful black rhino management tool for KZN black rhinos since 1962 (Hitchins 1984; Hall-Martin and Knight 1994; Emslie et al. 2009). Rhinos moved back to HiP would allow the source population to benefit genetically from the growth in other reserves effectively reinstating a single large genetic population. There are still costs associated with translocation and not all of them are financial, but may also include short-term social disruption, reduction in breeding performance and death during capture and post



Figure 3.3: Black rhino game reserves in KZN. Ndumo Game Reserve, Tembe Elephant Reserve, Pongola Game Reserve, Ithala Game Reserve, Mkuze Game Reserve, Thanda Game Reserve, Phinda Resource Reserve, Zululand Rhino Reserve, Ubizane Wildlife Reserve, eMakhosini Heritage Park, HiP, Eastern Shores, Weenan Game Reserve

translocation (Hitchins 1984; Hall-Martin and Knight 1994; Adcock et al. 1998; Linklater et al. 2011). However, benefits to KZN *D. b. minor* through use of this type of adaptive management may outweigh associated costs of all types (Van Houtan et al. 2009).

Genetic Replenishment

Managers may also consider genetic replenishment by introducing native *D. b. minor* from Zimbabwe to KZN. The native *D. b. minor* population in Zimbabwe

has higher mtDNA variation (six haplotypes; $n=11$) than the KZN metapopulation (one mtDNA haplotype; $n=65$) (Anderson-Lederer et al. 2012) and has a higher microsatellite H_E and H_O than KZN (Table 3.4). Although there may be concerns with how well the native Zimbabwe rhinos would thrive in KZN, the reverse scenario was successfully tested. In 1997, twenty-seven native KZN *D. b. minor* were translocated to Malilangwe, Zimbabwe where they were managed separately and not outbred with native Zimbabwe populations. The translocated KZN rhinos thrived with a growth rate of 8.3% per annum (du Toit 2001). The success of the translocated KZN *D. b. minor* to Zimbabwe may translate to native Zimbabwe *D. b. minor* being successfully translocated to KZN. However, before any native Zimbabwe *D. b. minor* are introduced to the KZN source populations, genetic supplementation experiments with a mixed population of KZN and native Zimbabwe *D. b. minor* should be cautiously and systematically established. F1 and F2 offspring of the mixed population could then be carefully monitored for signs of reduction in reproductive fitness (outbreeding depression), even though the likelihood of outbreeding depression in supplemented populations of the same species is low if they have the same karyotype, have been isolated for less than 500 years, and occupy similar environments (Frankham et al. 2011). An example of this type of mixed population is in Kruger National Park where 15 native Zimbabwe and 82 native KZN *D. b. minor* were introduced to the southern section of the nearly 2 million ha park from 1971 to 1988 (Hall-Martin and Castley 2001). The population would be ideal to study if stud books were kept and genetic samples collected during ear notching for identification events.

Possible Inbreeding Depression

Experiments with mixed KZN and native Zimbabwe *D. b. minor* populations may also offer insight and solutions to possible inbreeding depression being expressed within the HiP population. Inbreeding depression and the way it impacts wild populations varies across taxa, populations and environments (Keller and Waller 2002). Garner et al. (2005) found that in populations that experienced one or more demographic threats (e.g. population declines, bottlenecks, reduction of population range) the level of genetic variation (>20% reduction in heterozygosity) was affected. Indeed, as the impact of genetic drift increases in small fragmented populations, genetic loss will reduce the range of possible adaptive responses in stressful environments (Bijlsma and Loeschcke 2012). Unfortunately, most stress resistance alleles have lower frequencies in populations and as genetic erosion takes place, those “rare” alleles have a higher probability of being lost (Bijlsma and Loeschcke 2012). While no outward signs of inbreeding depression have been identified, the average growth rate of the HiP population was only 3.4% per annum (1999 – 2008) (Clinning et al. 2009). This is quite low when compared to the 6.75% per annum in KZP (Ferreira et al. 2011) and 8.3% per annum (du Toit 2001) for the 27 translocated KZN *D. b. minor* to Zimbabwe mentioned earlier. The low growth rate and homozygote excess could be a result of genetic erosion or simply that Zimbabwe has a greater annual rain fall than South Africa which may increase the natural resources available to the black rhinos creating an environment more favourable to higher birth rates (Berkeley and Linklater 2010).

3.6 Conclusion

It is important to detect losses in the level of genetic variation in KZN, especially HiP where no translocations into the population take place. As poaching continues to threaten rhino populations across Asia and Africa, problems associated with managing small isolated populations grow with it. KZN black rhino managers have an opportunity to take necessary steps to curtail or stop inbreeding depression before it becomes detrimental to the metapopulation. With the KZN metapopulation exhibiting a single mtDNA haplotype and lower expected and observed heterozygosities than the native Zimbabwe *D. b. minor* (Table 3.3 & 3.5) suggesting inbreeding, a more detailed management plan including data from this study may be required to prevent further loss of genetic variation, especially within HiP. The natural migration process that black rhino were afforded before the 1600's may no longer be possible, but increasing reserve size, translocations of native *D. b. minor* from Zimbabwe to KZN or serial translocations from amongst the KZN metapopulation back into HiP may aid in maintaining current levels or increasing the overall genetic diversity of the source population that will translate to more diversity in the KZN metapopulation.

3.7 Acknowledgements

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CHAPTER FOUR

Population Viability Analysis of Black Rhino (*Diceros bicornis minor*) in Hluhluwe-iMfolozi Game Park, KwaZulu-Natal, South Africa



Photo by Rosalynn Anderson-Lederer

4.1 Abstract

Once the loss of genetic diversity is identified in an endangered population, it is important for conservation managers to develop a plan to arrest further loss.

Population viability analyses (PVA) are stochastic computer simulations used to predict the probability of future population persistence or extinction.

Researchers use PVAs to visualize quantitative data to create informed management plans for endangered species. Hluhluwe-iMfolozi Game Park (HiP) in KwaZulu-Natal (KZN) is home to the largest remnant population of black rhino (*Diceros bicornis minor*) in South Africa and is a primary source for metapopulation expansion in South Africa. Unfortunately, HiP has significantly lower levels of mitochondrial and microsatellite DNA genetic variability than native *D. b. minor* populations in Zimbabwe. In this study, Vortex PVA was used to model population increases and supplementations into HiP with individuals from the KZN metapopulation and Zimbabwe. If current management remains unchanged, the PVA predicted a loss in the mean expected heterozygosity of ~25% over ~100 black rhino generations (BRGs). Doubling the size of the modelled population decreased the rate of loss of the mean H_E by ~10% over ~100 BRGs. When supplementations of one female and one male black rhino from the KZN metapopulation were made every ten gestational years, the mean H_E of the population was maintained (~ 0.45) over ~100 BRGs, but increased ~30% when supplemented with one female and one male from Zimbabwe. PVA results indicate that artificial game park expansion through supplementation is effective and does not require a large number of individuals or frequent translocation. Based on these results, HiP managers should consider

incorporating a supplementation regime into current management plans to prevent further loss of genetic diversity within this valuable source population.

4.2 Introduction

The task of restoring populations of vulnerable and endangered species while maintaining genetic diversity is challenging because remnant populations are inevitably small. When a species which had an historically large and widespread population is fragmented into smaller isolated populations, genetic drift can quickly eliminate genetic variability (Lacy 1987b). In response to the problem of small populations the idea of a minimum viable population (MVP) size was introduced. Unfortunately, population sizes required to prevent variability loss tend to be significantly larger than the targets set by conservation managers and organizations (Traill et al. 2010). Nevertheless, increasing population sizes to match the MVP is not always possible due to habitat size limits. Adding conspecific individuals (supplementations) to the population (IUCN 1987), however, can be carried out via serial translocations. Supplementations via serial translocations among small populations would create an artificial metapopulation that would make the effective population size (N_e) large enough to match the MVP size. The N_e could in turn prevent the loss of genetic diversity in small, fragmented populations (Waite et al. 2005).

Similar to MVPs, population viability analyses (PVA) use a stochastic computer simulation to forecast the likelihood of future population persistence or extinction using species specific life-history data (vital rates: e.g. age, reproductive rates, mortality, breeding system) (Boyce 1992). Conservation managers typically use the outcomes of PVAs to visualize quantitative data (e.g.

demographic, ecological, and genetic) to establish policy priorities and develop realistic targets (e.g. fiscal, technical, personnel use) (Possingham et al. 1993; Lindenmayer et al. 1993).

The species life-history data used to create models can be difficult to obtain from small populations of some species, which means PVA models have an inherent uncertainty in their results (Shaffer 1990; Lindenmayer et al. 1993). That being said, retrospective PVAs performed on birds (e.g. black-capped chickadee (*Parus atricapillus*), mammals (e.g. Cape hunting dog (*Cynomys ludovicianus*), fish (brook trout (*Salvelinus fontinalis*) and reptiles (sage-brush lizard (*Sceloporus graciosus*)) were proven to be a reliable and effective tool for managing endangered species (Brook et al. 2000). Several PVA models for wild and captive black rhino populations have also been reported.

Analyses of founder members for captive populations, demographic stability, and loss of genetic variability in reserves in Kenya and Tanzania as well as the viability of captive black rhinos have been investigated using VORTEX PVA (Lacy 1987a; Foose 1987; Moehlman et al. 1996). In addition, conservation strategies including carrying capacity, population structure and density-dependence (Swart et al. 1990; Adcock 2001; Crooms et al. 2002; Dunn et al. 2007) have been investigated using a range of MVP and PVA modelling techniques. With a black rhino generation time of c. 14 years (Brooks and Adcock 1997) it is too early to corroborate most of the simulated results with real populations that have been modelled. Nonetheless, population estimates of black rhino in the Ngorongoro Crater, Tanzania in 2006 (Mills et al. 2006) were comparable with predicted PVA estimates from simulations made in 1996 (Moehlman et al. 1996) confirming the predicted outcome of the black rhino

population. In addition, the retrospective study by Crooms et al. (2002) examining structure and density-dependence models demonstrated that out of five deterministic models tested, Fowler's translocation model (Fowler 1981) best fitted the Hluhluwe-iMfolozi (HiP) and Mkuze Game Reserve (MGR) population and translocation censuses (HiP 1990 - 1998; MGR 1989 - 1998), again showing that computer models based on black rhino data can produce realistic population outcomes.

The HiP *D. b. minor* population is important in the black rhino recovery programme because it is the largest endemic and remnant population of *D. b. minor* in South Africa. Recent findings of significantly lower microsatellite DNA variation in KwaZulu-Natal (KZN) *D. b. minor* than native Zimbabwe *D. b. minor* (KZN: $H_E = 0.47$, $H_O = 0.49$; Zimbabwe: $H_E = 0.65$, $H_O = 0.72$; Chapter 3) and low mtDNA variation (KZN: one haplotype, $n=65$; Zimbabwe: six haplotypes, $n=11$ Anderson-Lederer et al. 2012) strongly suggest a loss of genetic diversity within HiP and the KZN metapopulation requiring alterations to current management strategies to preserve genetic diversity.

There are several possible management responses that could be taken to reduce the loss of genetic variation. One such strategy is to expand the size of the game reserve. This has been proposed for HiP by connecting it via corridors with other neighbouring game parks including Zululand Game Reserve, Mkuze Game Reserve, Phinda Resource Reserve and Greater St. Lucia Wetland Park. This strategy however, has limits including the cost of purchasing land, increased management expenses (e.g. additional monitoring, fencing) and the logistics of creating corridors. In addition, the land must fulfill strict black rhino nutritional requirements for optimal breeding at pre-set carrying capacities

(CC) enforced by state agencies (Hall-Martin and Castley 2001; Emslie et al. 2009). It is unclear, though how much area would be required to slow the rate of loss in the level of genetic variation for the population.

Another strategy is serial translocation that exchanges individuals between small populations allowing each population to benefit from immigration. Limitations for this strategy involve costs associated with translocations. Expenses include, but are not limited to the capture and holding of animals (e.g. trucks, helicopters, fuel, transportation crates, darting medications), as well as legal and biological considerations (e.g. age and sex of animals being moved) (Emslie et al. 2009). If the translocation takes place across international borders, the process can be even more complicated by government involvement (Emslie et al. 2009). It is uncertain however, how many individuals would need to be moved between populations to slow the rate of loss in the level of genetic variation for the population.

PVA modelling can be used to determine which management scenario might be most effective. When choosing a PVA programme for modelling possible management scenarios, Lindenmayer et al. (1995) suggests selection criteria be based on (1) the primary objectives of the study and (2) the strengths, limitations and assumptions of the programme and how these correspond to the traits, life-history parameters, quality and quantity of available data for the species being modelled. The criteria selection is important because not all PVAs calculate outcomes in the same way and what may be appropriate for one type of population may not be appropriate for others (e.g. closed population versus metapopulation, long-lived species versus short-lived). VORTEX has been rigorously examined in peer-reviewed studies and

population forecasts have been shown to be accurate when sufficient and accurate species specific life-history data are available (Brook et al. 1997; Brook et al. 1999; Brook et al. 2000; Coulson et al. 2001).

The objective of this study was to use a PVA to examine a set of management strategies and determine the most effective management plan for preventing the loss of genetic variation within the HiP population. Included in the analyses are scenarios that increase the population size and supplementations made with individuals representing the KZN metapopulation and individuals representing the native Zimbabwe *D. b. minor* population. The results of the models are used to make management recommendations for the HiP population.

4.3 Methods

Available life-history data of the HiP population of *D. b. minor* were incorporated into the PVA model. When specific data for HiP black rhino were unavailable, information from other populations of black rhino were used from published literature and unpublished reports. Life-history data and model parameters were set as follows (Table 4.1):

4.3.1 Species Description

Number of iterations

The model used a random number generator so that none of the repeated simulations would be the same (Miller and Lacy 2005); therefore, 100 iterations is usually adequate to uncover tendencies (Lacy 1993); however, between 500

Table 4.1: Vortex scenario parameters

Vortex Parameters	Value	Source
Species Description		
Iterations	1000	This study
No. of years	1000 'gestational' (1342 actual)	This study
Duration of each "year"	490 days (adjusted to accommodate length of black rhino pregnancy)	This study
Extinction def:	one sex remains	This study
No. of Populations	1	This study
Inbreeding depression	Yes & No	This study
Reproductive System		
Polygynous	Polygynous	Garnier et al. 2001
Age of First offspring for females	6⁺ (gestational years)	Owen-Smith 1988
Age of First offspring for males	7⁺ (gestational years)	Lent & Fike 2003
Maximum Age of Reproduction	27⁺ (gestational years)	Shenkle & Shenkle 1969
Maximum no. of broods per year	1	Owen-Smith 1988
Maximum no. of progeny per brood	1	Owen-Smith 1988
Sex ratio at birth in % males	53	Owen-Smith 1988
Density Dependent Reproduction	= (32-((32-28)*((N/K)^8)))*(N/0+N)[*]	Berkley & Linklater 2010
Reproductive Rates		
% Adult females breeding	32%	This study
environmental variation in % breeding	3	Clinning 2009
Mortality Rates Female in %		
from age 0 - 1	15%⁺ (adjusted from 11%)*	Clinning 2009
from age 1 - 4	4%⁺ (adjusted from 3%)*	Adcock and Emslie 2003
from age 4-6	8%⁺ (adjusted from 5.7%)*	Adcock and Emslie 2003
Mortality from age 6 and above	5%⁺ (adjusted from 3.5%)*	Owen-Smith 1988
Mortality rate Male %		
Mortality from age 0 - 1	15%⁺ (adjusted from 11%)*	Owen-Smith 1988
from age 1 - 4	4%⁺ (adjusted from 3%)*	Adcock and Emslie 2003
from age 4-6	8%⁺ (adjusted from 5.7%)*	Adcock and Emslie 2003
Mortality from age 6 and above	10%⁺ (adjusted from 7.3%)*	Owen-Smith 1988
Mate Monopolization		
% males breeding in pool	100	
% males successfully siring offspring	37.1 (Calculated by Vortex)	
Mean # of mates/successful sire	1.2 (Calculated by Vortex)	
Initial population size		
Stable Age Distribution	300	Fanayo et al. 2005
Carrying Capacity		
Carrying capacity (K)	430 & 860[‡]	Emslie 2009
SD in K due to environmental variation	43, 86	Emslie 2009
Harvest		
Population Harvested?	Yes	Emslie 2009
First year of harvest?	1	Emslie 2009
Last year of harvest?	1000	Emslie 2009
Interval between harvests	1	Emslie 2009
Optional Criterion for harvest	=(W+X)≥(K/2)	
Supplementation		
first year of supplementation	1F, 1M; 2F, 2M	This study
last year of supplementation	1	This study
interval between supplementations	Varied	This study
Genetic Mangement		
Number of microsatellite loci examined	1, 2, 5, & 10 years (gestational)	This study
Number of microsatellite loci examined	10 from KZN pop. & 10 from Zim. pop	This study, Garnier et al. 2001

⁺ Variation in age and mortality greatly affect PVA outcomes, younger ages and older mortality rates increase reproductive output over an animal's life; ^{*}Based on density dependent variables, see text for details; ^{*} Percentages are adjusted to reflect a longer year of 490 days; [‡]Carrying capacities based on HiP CC, see text for details

and 1000 iterations are encouraged to provide more rigorous results (Miller and Lacy 2005). One thousand iterations were chosen for the models run in this study.

Duration of a year

The length of a year was adjusted from 365 days (default) to 490 days in order to satisfy ‘maximum number of broods per year’ in the “Reproductive System” section of the parameter settings (Lacy, R. pers. comm.). Entries made for ‘maximum number of broods per year’ must be a whole number. The gestation period for a black rhino is 460 days (15.33 months) (Linklater 2007). In order to enter an integer (i.e. 1) instead of a fraction (i.e. 0.8), a ‘year’ was adjusted to reflect 490 days to accommodate one brood per year, plus an additional 30 days, the minimum time required to become pregnant again. This was done to avoid over-estimating the number of births in the simulations. The adjusted year (460 + 30 days) is referred to as the ‘gestational year’.

Number of years

Simulations were run for 1000 gestational years to see how genetic variation changed over a lengthy time period. One thousand gestational years translates to 1342 calendar years. According to the Conservation Plan for the Black Rhinoceros in South Africa (Brooks and Adcock 1997) one black rhino generation (BRG) is c. 14 years. One thousand gestational years enabled visualization of approximately 96 BRGs.

Inbreeding depression

Previous black rhino microsatellite DNA studies concluded that the level of genetic variation in *D. b. minor* was not low enough to be of concern (Harley et

al. 2005; Karsten et al. 2011). However, since KZN *D. b. minor* has only one mtDNA haplotype (Anderson-Lederer et al. 2012) and homozygote excess was observed in the 10 microsatellite DNA loci examined (Chapter Three), scenarios with and without inbreeding were run to compare how inbreeding depression influences the outcome for the simulated population. In scenarios with inbreeding depression, a default value of lethal equivalents (3.14) was selected; the default is based on Ralls et al. (1988) survey of 40 mammal populations (Miller and Lacy 2005). When modelling inbreeding depression, the model reduced the survival of offspring only in the first year, which caused the results of inbreeding depression to be conservative (Miller and Lacy 2005).

Catastrophes

No catastrophes (e.g. drought and disease) were modelled in these scenarios. Environmental variation is reflected in other parameters, and this project is focused on establishing baseline genetic results for increasing population sizes and supplementation, not addressing how catastrophes affect the population.

4.3.2 Reproductive System

Age of first offspring for females

Female black rhino first give birth between ages 6.5 and 8.5 (Owen-Smith 1988). Using the average of 7.5, age was adjusted to 6 years based on a gestational year ($7.5 \text{ years} \times 365.25 \text{ days} / 490 \text{ days} = 5.59 \text{ years}$, ~ 6). This number appears high because black rhino females become sexually mature as early as 3.5 - 4 years old (Schenkel and Schenkel-Hulliger 1969), but the model population is assumed to be near carrying capacity (CC). Large mammal density-dependence is expected to be weak except near CC, where it is reflected

in low reproductive rates especially in younger females which exhibit delayed first reproduction (Fowler 1981). An increase in the age of first offspring will mean a reduced reproductive output over the life of the female, but the mean age was chosen to reflect conservative outcomes for the simulated population.

Age of first offspring for males

Males successfully reproduce at approximately 9 years old (Owen-Smith 1988; Bertschinger 1994; Lent and Fike 2003). Age was adjusted from 9 years to 7 years based on a gestational year ($9 \text{ years} \times 365.25 \text{ days} / 490 \text{ days} = 6.70 \text{ years}$, ~ 7).

Maximum age of reproduction, number of progeny per year

Black rhino females have one offspring per pregnancy. The maximum age of reproduction is approximately 37 years old (Schenkel and Schenkel-Hulliger 1969; Owen-Smith 1988). Age was adjusted to 28 based on a gestational year ($37 \text{ years} \times 365.25 \text{ days} / 490 \text{ days} = 27.58$, ~ 28).

Sex ratio at birth – in % male

The sex ratio of black rhino across combined age groups averages to approximately 1:1 (Hillman-Smith and Groves 1994); however, the proportion of males detected soon after birth is slightly higher (53%) (Emslie et al. 2009; Berkeley and Linklater 2010).

Density-dependent reproduction

Large-bodied species show a life-history strategy that includes slow growth rates with fitness components (e.g. infant mortality, reproductive rates) that are affected as populations near CC (Eberhardt 1977; Fowler 1981; Gaillard et al.

2000). Density-dependence first affects the youngest members of a population with infant and juvenile mortality highest when populations approach CC (Gaillard et al. 2000). Fecundity of young females is next to be affected, followed by adult females and then adult (male and female) survival (Gaillard et al. 2000). The influence of density-dependence has been observed in several large ungulate species, including wildebeest (Mduma et al. 1999), caribou (Messier et al. 1988; Tews et al. 2007), wild reindeer (Skogland 1985), roe deer (Kjellander et al. 2004) and northern fur seals (Fowler 1990).

The function used for modelling was *Density-Dependent Reproduction* = $(32 - ((32 - 28) * (N/K)^8)) * (N / (0 + N))$, where N = population size, K = carrying capacity and P = population identifier, based on the following:

- % Breeding at Low Density, P (0): 32 (Clinning et al. 2009)
- % Breeding at Carrying Capacity (Maximum Age), P (K): 28 (Miller and Lacy 2005)
- Allee Parameter A: 0 Vortex manual: (Miller and Lacy 2005)
- Steepness Parameter B: 8 Vortex manual: (Miller and Lacy 2005)

4.3.3 Reproductive Rates

% Adult Females Breeding

The percent of adult females breeding is set automatically based on the density-dependence variables entered in the model.

4.3.4 Mortality Rates

Mortality rates were based on Owen-Smith's (1988) observations from HiP. Any age groups that were not documented by Owen-Smith were supplemented with mortality rates for black rhino from South Africa and Namibia from Adcock and

Emslie (2003). All mortality rates were adjusted to accommodate the gestational year (Table 4.1).

4.3.5 Initial Population Size

Between 1930 – 2009, black rhino numbers at HiP ranged between ~130 to ~400 individuals (Emslie 2011; Fanayo et al. 2005), although Clinning et al. (2009) determined that past census numbers were over-estimated, in some years by as much as 48%. Since it is impossible to know the exact number of individuals in a wild population, 300 was entered as the initial population size and the default ‘stable age distribution’ was chosen.

4.3.6 Carrying Capacity

Management cannot increase a reserve’s CC unless the size of the reserve is increased through land acquisition. However, an understanding of how a population’s size is affected by its CC is important in order to decrease effects of density-dependence when drafting black rhino management plans (Adcock 2001). Owen-Smith (2001) defines CC as the number of individuals a population can sustain (relying on resources in the area) that remains constant due to births cancelling out deaths. Brooks and Adcock (1997) estimate that the CC for HiP black rhino is 430 individuals. Their estimates are based on frost ratings, vegetation and approximate annual rainfall each year. Carrying capacity for the simulated population was therefore set at 430 for baseline scenarios. If the simulated population were to increase in size (e.g. HiP merging with neighbouring game reserves), it may be able to double its CC (~860); therefore, an 860 CC was also used to visualize how increasing the size of the simulated population would affect the population from a genetic perspective.

4.3.7 Harvest (*Capturing individuals to relocate to other reserves*)

Harvest criteria

Harvests occurred every gestational year (490 days).

Optional criteria for harvest

Parameters were set to only harvest if the total population size was 50% or more of the CC (Emslie et al. 2009). The function used for this was *Optional Criteria* $= (W+X) \geq (K/2)$, where X is females in the population, W is males in the population and K is the carrying capacity (Lacy, R. pers. comm.). If during any year of a harvest the total population was less than 50% of the carrying capacity, no harvest took place (Miller and Lacy 2005).

Number of female and male of each age to be harvested

As suggested by translocation studies, the model harvested only adult females and males (Linklater et al. 2011; Linklater et al. 2012) because young translocated black rhino experience higher mortality rates than their adult counterparts and young females have lower fecundity rates after translocations. The SADC recommends harvest rates of 5% - 8% for black rhino (Emslie 2001). The percentage of black rhino removed from the modelled population was set at 4% of the total population per gestational year (which translates to 5.4% for a calendar year). This percentage was chosen because it was conservative and also because it is the harvesting goal of HiP (Clinning et al. 2009). If harvesting took place during a particular year (population size was more than 50% of the carrying capacity), the 4% was split between females and males (e.g. 2% of the total population was removed from the adult females (≥ 6 years old) and 2% of the total population was removed from the adult males (≥ 7 years old). The

functions used for this were *Females Harvested* = $X * \%$ and *Males Harvested* = $W * \%$, where X was females, W was males, and % was half of the percent harvested that year.

4.3.8 Supplementation

Frequency of supplementations

Supplementations occurred every year (E1Y), two years (E2Y), 5 years (E5Y) and ten years (E10Y).

Number of females and males supplemented

Supplementations were made with one female and one male (1F, 1M) and two females and two males (2F, 2M) at each of the time periods simulated.

4.3.9 Genetic Management

Data from 10 microsatellite DNA loci for the KZN metapopulation (Chapter 3) with an H_E of 0.47 were imported to the PVA to reflect the current level of genetic diversity for the modelled population (Table 4.2).

Supplementations were made with two different types of supplemental individuals, those representing the KZN metapopulation (H_E of 0.47; Table 4.2) and those representing the native Zimbabwe *D. b. minor* population (H_E of 0.62; Table 4.3).

Table 4.2: Allele frequencies for the 10 microsatellite DNA loci examined in Chapter 3 for HiP *D. b.*

Number of neutral loci imported to																
Vortex:	10															
Number of alleles for each loci:	4	2	3	3	4	4	1	5	4	3						
Frequency distributions for loci 1:	0.4	0.154	0.43	0.02												
loci 2:	0.79	0.208														
loci 3:	0.07	0.034	0.89													
loci 4:	0.39	0.419	0.19													
loci 5:	0.45	0.390	0.08	0.08												
loci 6:	0	0.603	0.28	0.11												
loci 7:	1															
loci 8:	0.43	0.346	0.19	0.03	0.01											
loci 9:	0.01	0.042	0.41	0.53												
loci 10:	0.01	0.389	0.6													

Table 4.3: Allele frequencies for the 10 microsatellite DNA loci examined by Garnier *et al.* (2001) for native Zimbabwe *D. b. minor*

Number of neutral loci imported to										
Vortex:		10								
Number of alleles for each loci:		4	7	4	2	4	2	3	4	5
Frequency distributions for loci 1:		0.242	0.0561	0.015	0.182					
loci 2:		0.273	0.106	0.212	0.273	0.03	0.061	0.045		
loci 3:		0.394	0.015	0.485	0.106					
loci 4:		0.348	0.652							
loci 5:		0.197	0.030	0.606	0.167					
loci 6:		0.682	0.318							
loci 7:		0.076	0.682	0.242						
loci 8:		0.424	0.061	0.258	0.258					
loci 9:		0.258	0.318	0.106	0.227	0.091				
loci 10:		0.167	0.121	0.409	0.176	0.030				

4.4 Results

4.4.1 Levels of Genetic Variation with No Supplementation

When current levels of genetic variation observed at 10 microsatellite DNA loci in the KZN metapopulation (H_E of 0.47) were assigned to the modelled population (CC 430) with no inbreeding depression and no supplementation, the population decreased in size from 300 where it then stabilised to approximately 259 individuals (Figure 4.1). In the model population with no supplementation and a CC of 430, mean expected heterozygosity (H_E) averaged over 10 microsatellite loci decreased by 3%, 11% and 23% over 19, 48 and 96

BRGs respectively (200, 500 and 1000 gestational years) (Table 4.4). When the model population's CC was doubled to 860, to simulate the outcome of increasing the size of the reserve (e.g. merging with other reserves), genetic variation decreased over time, but at a slower rate than the 430 CC (Table 4.5). The mean H_E averaged over 10 microsatellite loci was maintained for approximately 24 BRGs (250 gestational years), but then began to decline. There was a decrease in the mean H_E of 4% and 11% over 48 and 96 BRGs respectively (500 and 1000 gestational years respectively) (Table 4.5).

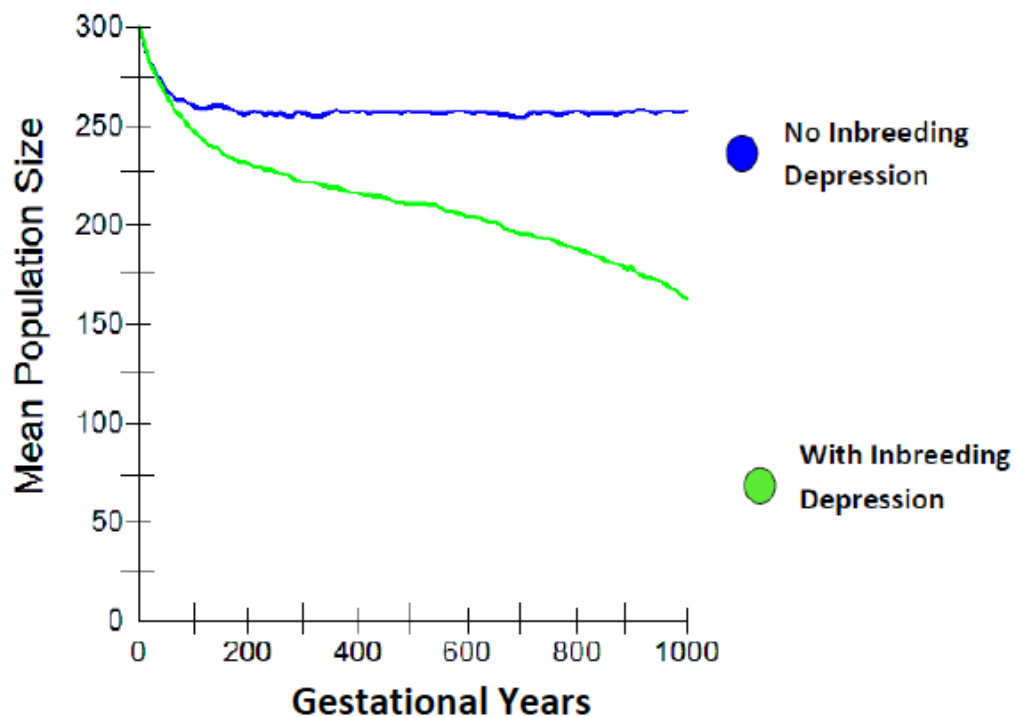


Figure 4.1: The mean population sizes for a modelled population with a CC of 430, with and without inbreeding depression, and a 4% harvesting rate per gestational year (translates to 5.4% in a calendar year).

Table 4.4: Expected and observed heterozygosities (H_E and H_O) with standard error for HiP *D. b. minor*; with and without inbreeding depression at 430 carrying capacity for years 50, 100, 150, 200, 250, 500, 700 & 1000 (5, 10, 14, 19, 24, 48, 67, and 96 black rhino generations respectively); with mean population sizes and standard error. The 4% harvest is for a gestational year, which translates to 5.4% in a calendar year.

430 carrying capacity, 4% Harvest						
Year	~ Pop Size	No Inbreeding Depression, No Genetic Supplementation		~ Pop Size	Inbreeding Depression, No Genetic Supplementation	
		H_E	H_O		H_E	H_O
50	267 ± 1.25	0.459 ± 0.000	0.462 ± 0.000	263 ± 1.21	0.459 ± 0.000	0.462 ± 0.000
100	260 ± 1.19	0.451 ± 0.001	0.453 ± 0.001	247 ± 1.09	0.453 ± 0.001	0.455 ± 0.001
150	257 ± 1.16	0.446 ± 0.001	0.448 ± 0.001	238 ± 1.01	0.447 ± 0.001	0.450 ± 0.001
200	258 ± 1.21	0.438 ± 0.001	0.441 ± 0.001	230 ± 0.91	0.440 ± 0.001	0.444 ± 0.001
250	256 ± 1.20	0.432 ± 0.001	0.435 ± 0.001	226 ± 0.90	0.432 ± 0.001	0.436 ± 0.001
500	259 ± 1.23	0.402 ± 0.001	0.404 ± 0.001	211 ± 0.77	0.397 ± 0.001	0.400 ± 0.001
700	256 ± 1.18	0.379 ± 0.001	0.381 ± 0.001	199 ± 0.82	0.370 ± 0.001	0.373 ± 0.001
1000	257 ± 1.18	0.347 ± 0.002	0.349 ± 0.002	165 ± 1.39	0.339 ± 0.002	0.333 ± 0.002

Table 4.5: Expected and observed heterozygosities (H_E and H_O) with standard error for HiP *D. b. minor*; with and without inbreeding depression at 860 carrying capacity for years 50, 100, 150, 200, 250, 500, 700 & 1000 (5, 10, 14, 19, 24, 48, 67, and 96 black rhino generations respectively); with mean population sizes and standard error. The 4% harvest is for a gestational year, which translates to 5.4% in a calendar year.

860 carrying capacity, 4% Harvest						
Year	~ Pop Size	No Inbreeding Depression, No Genetic Supplementation		~ Pop Size	Inbreeding Depression, No Genetic Supplementation	
		H_E	H_O		H_E	H_O
50	473 ± 1.46	0.461 ± 0.000	0.462 ± 0.000	470 ± 1.48	0.461 ± 0.000	0.463 ± 0.000
100	503 ± 1.85	0.458 ± 0.000	0.459 ± 0.000	491 ± 1.85	0.458 ± 0.000	0.459 ± 0.000
150	514 ± 2.09	0.454 ± 0.000	0.456 ± 0.000	488 ± 1.85	0.454 ± 0.000	0.455 ± 0.000
200	517 ± 2.14	0.451 ± 0.001	0.452 ± 0.001	478 ± 1.65	0.450 ± 0.001	0.452 ± 0.001
250	519 ± 2.09	0.448 ± 0.001	0.449 ± 0.001	470 ± 1.59	0.447 ± 0.001	0.449 ± 0.001
500	516 ± 2.04	0.432 ± 0.001	0.433 ± 0.001	452 ± 1.33	0.430 ± 0.001	0.432 ± 0.001
700	516 ± 2.04	0.419 ± 0.001	0.421 ± 0.001	442 ± 1.25	0.416 ± 0.001	0.418 ± 0.001
1000	517 ± 2.11	0.402 ± 0.001	0.403 ± 0.001	425 ± 1.28	0.395 ± 0.001	0.397 ± 0.001

4.4.2 Level of Variation with Supplementations

Supplementations of one pair (1F, 1M) of KZN *D. b. minor* made to the simulated population (CC 430) approximately every BRG (10 gestational years, 13.4 calendar years) showed a negligible increase in mean population size (Figure 4.2). Supplementations made with one pair (1F, 1M) of KZN rhino every one and

two gestational years maintained the mean H_E (~ 0.45) through 96 BRGs (1000 gestational years) (Table 4.6; Figures 4.3 A-B). When supplementations of the pair were reduced to every five and ten gestations years, 96% and 92% of H_E averaged over 10 microsatellite loci was maintained respectively (Table 4.6; Figures 4.3 C-D). Supplementations made with one pair (1F, 1M) of Zimbabwe rhino every one, two, five and 10 gestational years increased the mean H_E by 34%, 27%, 31%, and 29% every 96 BRGs (1000 gestational years) (Table 4.6; Figures 4.3 A-D).

The difference in variation between the non-supplemented model population with a 430 CC and the model supplemented with one pair of KZN *D. b. minor* every one, two, five and 10 years is 32%, 30%, 25% and 19% respectively after 96 BRGs (1000 gestational years) (Tables 4.4 & 4.6). When the number of individuals supplemented to the model population from the KZN metapopulation was increased to two pairs (2F, 2M), every one, two, five and ten gestational years, there was only a slight improvement in the increase in genetic variation over the single pair scenarios (Table 4.6, Figure 4.4A). However, similar to the single pair, supplementations made with individuals from the native Zimbabwe population, increasing the pair number to two greatly improved the level of diversity within the model population (Table 4.6, Figure 4.4B).

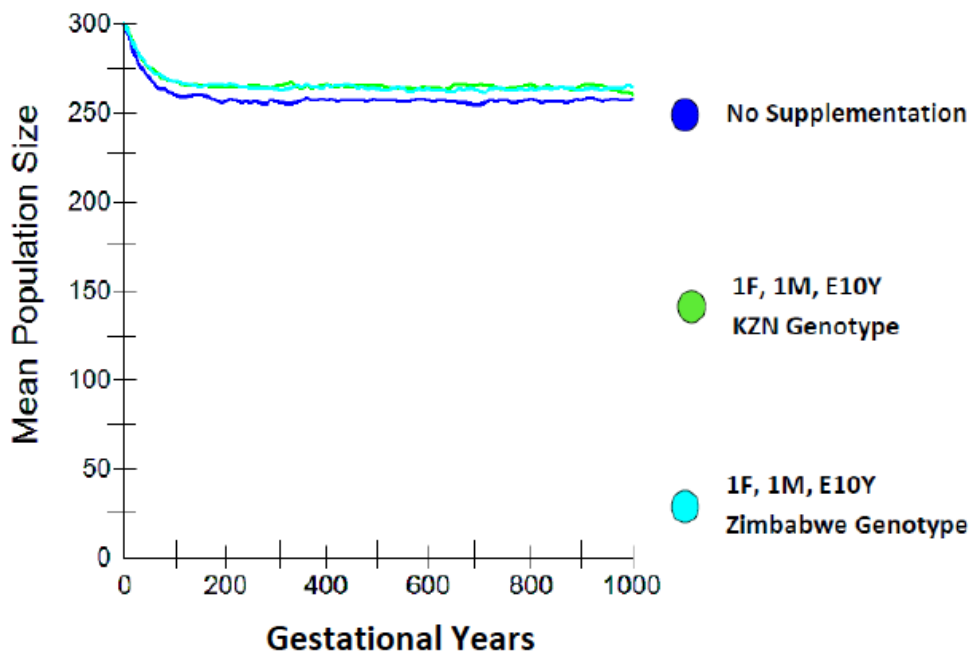


Figure 4.2: The mean population sizes for a modelled population with a CC of 430, without inbreeding depression, with a 4% harvesting rate (translates to 5.4% in a calendar year) with and without supplementations. Supplementations were made with one female and one male (1F, 1M) every ten years (E10Y; 13.4 years for a calendar year). Supplemented individuals were either assigned the KZN genotype or the native *D. b. minor* Zimbabwe genotype respectively.

Table 4.6: Mean population sizes and expected heterozygosity (H_E) with standard error for a population of *D. b. minor*; with various supplementations; without inbreeding depression at 430 carrying capacity for years 50, 100, 150, 200, 250, 500, 700 & 1000 (5, 10, 14, 19, 24, 48, 67, and 96 black rhino generations respectively). The 4% harvest is for a gestational year, which translates to 5.4% in a calendar year. Supplementations (Supp.) were made with one male and one female (1F, 1M), two females and two males (2F, 2M) at intervals of every one year (E1Y), two years (E2Y), five years (E5Y) and 10 years (E10Y) with individuals with the KwaZulu-Natal genotype (KZN GT) and Zimbabwe genotype (Zim. GT).

Supp. w/ 1F, 1M, E1Y - KZN GT				Supp. w/ 1F, 1M, E1Y - Zim. GT				Supp. w/ 2F, 2M, E1Y - KZN GT				Supp. w/ 2F, 2M, E1Y - Zim. GT			
Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E	
50	328 ± 1.10	0.461 ± 0.000		50	326 ± 1.12	0.605 ± 0.000		50	356 ± 0.83	0.462 ± 0.000		50	358 ± 0.85	0.644 ± 0.000	
100	329 ± 1.04	0.460 ± 0.000		100	328 ± 1.06	0.643 ± 0.000		100	358 ± 0.81	0.462 ± 0.000		100	358 ± 0.83	0.641 ± 0.000	
150	329 ± 1.05	0.459 ± 0.000		150	330 ± 1.07	0.644 ± 0.000		150	357 ± 0.83	0.461 ± 0.000		150	357 ± 0.82	0.619 ± 0.000	
200	331 ± 1.05	0.459 ± 0.000		200	329 ± 1.08	0.633 ± 0.000		200	356 ± 0.85	0.461 ± 0.000		200	357 ± 0.85	0.602 ± 0.000	
250	332 ± 1.06	0.459 ± 0.000		250	329 ± 1.07	0.620 ± 0.000		250	355 ± 0.86	0.461 ± 0.000		250	359 ± 0.82	0.593 ± 0.000	
500	330 ± 1.10	0.458 ± 0.000		500	330 ± 1.05	0.586 ± 0.000		500	356 ± 0.81	0.462 ± 0.000		500	357 ± 0.80	0.582 ± 0.000	
700	329 ± 1.09	0.458 ± 0.000		700	329 ± 1.05	0.580 ± 0.000		700	356 ± 0.84	0.461 ± 0.000		700	358 ± 0.82	0.582 ± 0.000	
1000	330 ± 1.08	0.458 ± 0.000		1000	330 ± 1.09	0.579 ± 0.000		1000	358 ± 0.85	0.462 ± 0.000		1000	356 ± 0.84	0.583 ± 0.000	
Supp. w/ 1F, 1M, E2Y - KZN GT				Supp. w/ 1F, 1M, E2Y - Zim. GT				Supp. w/ 2F, 2M, E2Y - KZN GT				Supp. w/ 2F, 2M, E2Y - Zim. GT			
Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E	
50	300 ± 1.31	0.460 ± 0.000		50	302 ± 1.28	0.555 ± 0.001		50	326 ± 1.12	0.461 ± 0.000		50	327 ± 1.09	0.605 ± 0.000	
100	299 ± 1.27	0.457 ± 0.000		100	300 ± 1.26	0.604 ± 0.001		100	327 ± 1.05	0.459 ± 0.000		100	328 ± 1.07	0.642 ± 0.000	
150	297 ± 1.25	0.454 ± 0.000		150	299 ± 1.27	0.628 ± 0.000		150	327 ± 1.06	0.458 ± 0.000		150	328 ± 1.07	0.644 ± 0.000	
200	300 ± 1.32	0.454 ± 0.000		200	298 ± 1.28	0.637 ± 0.000		200	332 ± 1.08	0.458 ± 0.000		200	330 ± 1.09	0.632 ± 0.000	
250	299 ± 1.22	0.453 ± 0.000		250	298 ± 1.25	0.637 ± 0.000		250	330 ± 1.04	0.459 ± 0.000		250	328 ± 1.05	0.620 ± 0.000	
500	298 ± 1.21	0.452 ± 0.001		500	299 ± 1.28	0.605 ± 0.000		500	328 ± 1.08	0.458 ± 0.000		500	329 ± 1.06	0.586 ± 0.000	
700	298 ± 1.28	0.452 ± 0.001		700	298 ± 1.28	0.585 ± 0.001		700	329 ± 1.05	0.459 ± 0.000		700	329 ± 1.13	0.579 ± 0.000	
1000	299 ± 1.28	0.452 ± 0.001		1000	297 ± 1.28	0.575 ± 0.001		1000	327 ± 1.09	0.458 ± 0.000		1000	328 ± 1.04	0.579 ± 0.000	
Supp. w/ 1F, 1M, E5Y - KZN GT				Supp. w/ 1F, 1M, E5Y - Zim. GT				Supp. w/ 2F, 2M, E5Y - KZN GT				Supp. w/ 2F, 2M, E5Y - Zim. GT			
Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E	
50	281 ± 1.30	0.460 ± 0.000		50	281 ± 1.28	0.506 ± 0.001		50	294 ± 1.31	0.460 ± 0.000		50	295 ± 1.29	0.541 ± 0.001	
100	276 ± 1.27	0.455 ± 0.000		100	272 ± 1.31	0.540 ± 0.001		100	291 ± 1.31	0.457 ± 0.000		100	291 ± 1.30	0.588 ± 0.001	
150	275 ± 1.31	0.451 ± 0.001		150	273 ± 1.25	0.565 ± 0.001		150	291 ± 1.29	0.454 ± 0.000		150	291 ± 1.27	0.615 ± 0.001	
200	274 ± 1.30	0.449 ± 0.001		200	273 ± 1.27	0.584 ± 0.001		200	291 ± 1.28	0.452 ± 0.001		200	291 ± 1.32	0.628 ± 0.001	
250	275 ± 1.27	0.446 ± 0.001		250	275 ± 1.29	0.598 ± 0.001		250	287 ± 1.27	0.451 ± 0.001		250	289 ± 1.30	0.633 ± 0.001	
500	273 ± 1.31	0.438 ± 0.001		500	273 ± 1.29	0.617 ± 0.001		500	289 ± 1.31	0.449 ± 0.001		500	290 ± 1.31	0.612 ± 0.001	
700	274 ± 1.29	0.435 ± 0.001		700	271 ± 1.28	0.608 ± 0.001		700	291 ± 1.27	0.448 ± 0.001		700	289 ± 1.33	0.591 ± 0.001	
1000	272 ± 1.27	0.434 ± 0.001		1000	275 ± 1.27	0.588 ± 0.001		1000	291 ± 1.29	0.448 ± 0.001		1000	291 ± 1.26	0.575 ± 0.001	
Supp. w/ 1F, 1M, E10Y - KZN GT				Supp. w/ 1F, 1M, E10Y - Zim. GT				Supp. w/ 2F, 2M, E10Y - KZN GT				Supp. w/ 2F, 2M, E10Y - Zim. GT			
Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E		Year	Mean Pop. Size	Mean H_E	
50	275 ± 1.33	0.458 ± 0.000		50	274 ± 1.26	0.483 ± 0.000		50	283 ± 1.30	0.459 ± 0.000		50	281 ± 1.28	0.506 ± 0.001	
100	268 ± 1.23	0.453 ± 0.000		100	268 ± 1.24	0.500 ± 0.000		100	274 ± 1.25	0.454 ± 0.000		100	274 ± 1.24	0.538 ± 0.001	
150	265 ± 1.24	0.448 ± 0.001		150	265 ± 1.23	0.517 ± 0.001		150	271 ± 1.23	0.450 ± 0.001		150	274 ± 1.33	0.563 ± 0.001	
200	266 ± 1.24	0.443 ± 0.001		200	265 ± 1.28	0.530 ± 0.001		200	273 ± 1.30	0.447 ± 0.001		200	272 ± 1.25	0.582 ± 0.001	
250	265 ± 1.27	0.439 ± 0.001		250	264 ± 1.21	0.542 ± 0.001		250	270 ± 1.22	0.444 ± 0.001		250	275 ± 1.29	0.595 ± 0.001	
500	266 ± 1.21	0.425 ± 0.001		500	263 ± 1.22	0.577 ± 0.001		500	271 ± 1.22	0.437 ± 0.001		500	272 ± 1.27	0.614 ± 0.001	
700	266 ± 1.26	0.417 ± 0.417		700	262 ± 1.25	0.583 ± 0.001		700	273 ± 1.31	0.435 ± 0.001		700	273 ± 1.29	0.606 ± 0.001	
1000	260 ± 1.22	0.412 ± 0.001		1000	265 ± 1.25	0.580 ± 0.001		1000	275 ± 1.27	0.434 ± 0.001		1000	274 ± 1.27	0.587 ± 0.001	

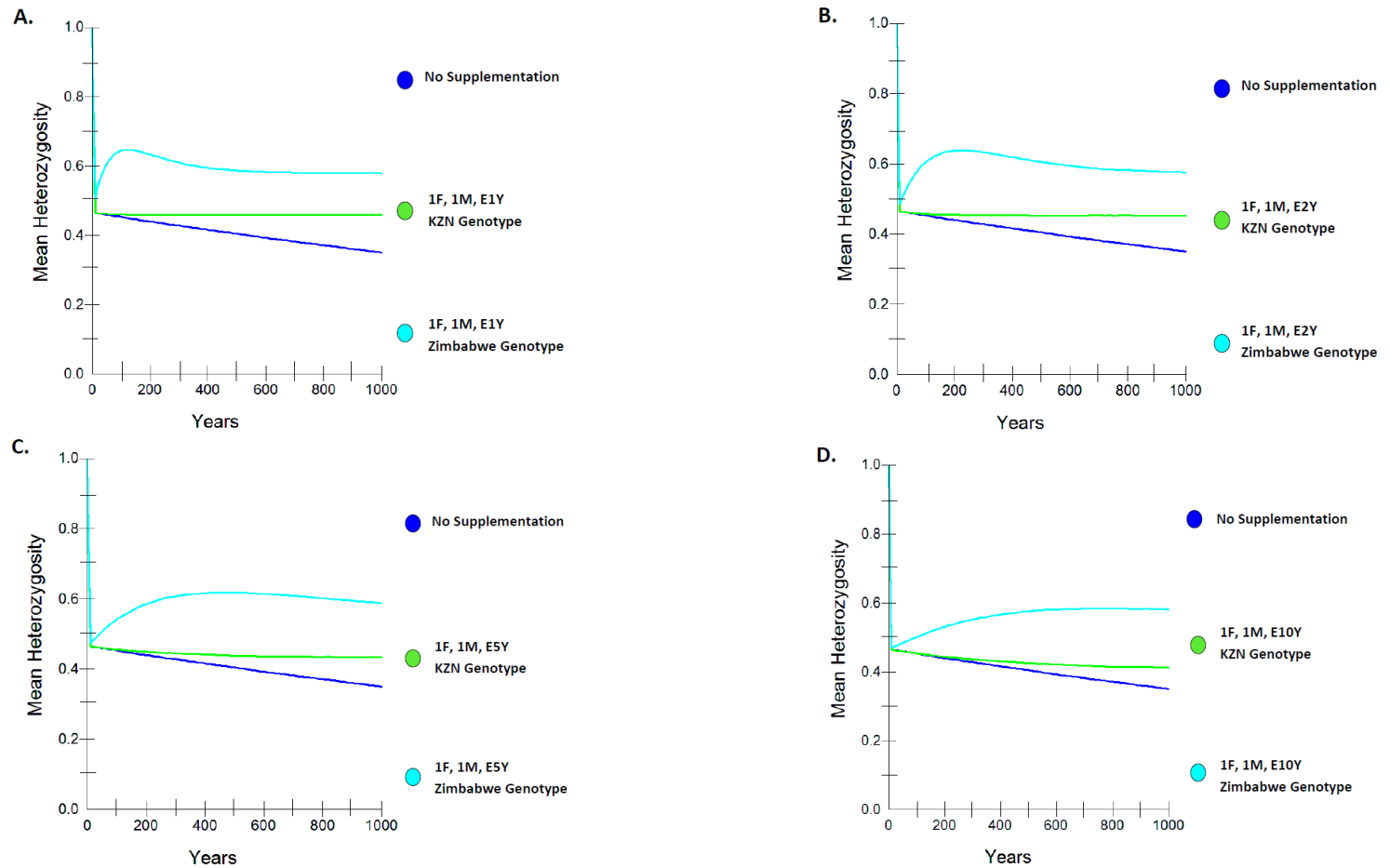
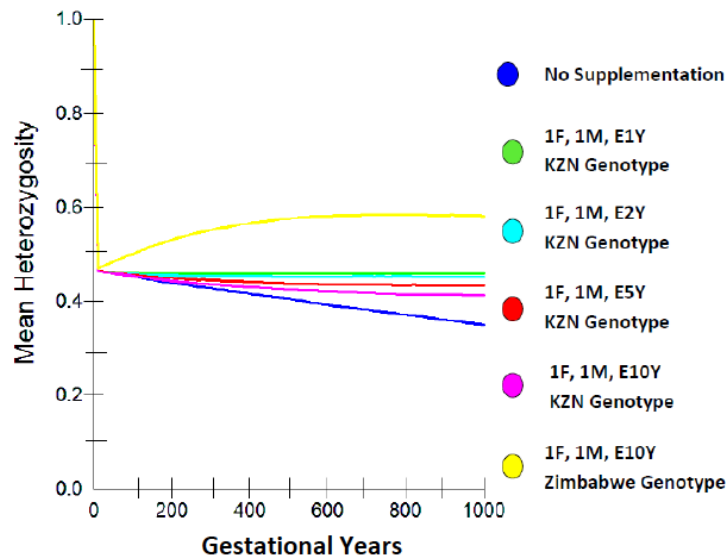


Figure 4.3: Mean expected heterozygosity for a modelled population with a carrying capacity of 430, without inbreeding depression, with a 4% harvesting rate per gestational year (translates to 5.4% in a calendar year) with and without supplementations. Supplementations were made with one female and one male (1F, 1M). Supplemented individuals were either from the KZN metapopulation or the native *D. b. minor* Zimbabwe population. A) Supplementations made every gestational year (1EY). B) Supplementations made every two gestational years (E2Y). C) Supplementations made every five gestational years (E5Y). D) Supplementations made every 10 gestational years (E10Y)

A.



B.

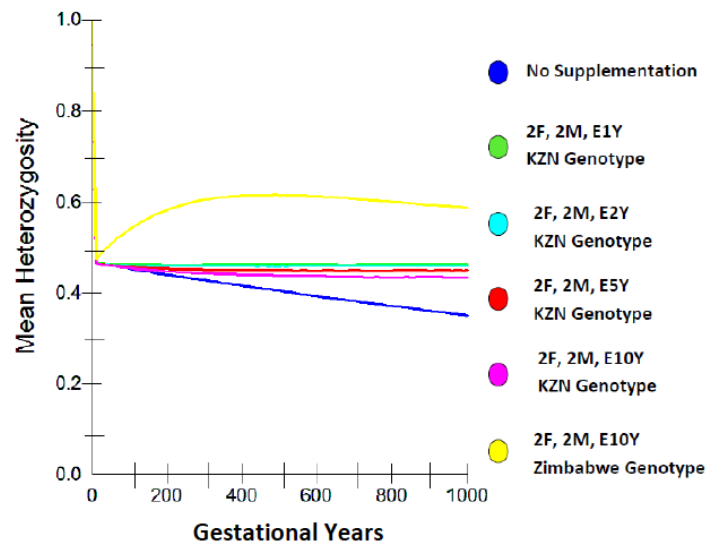


Figure 4.4 Mean expected heterozygosity for a modelled population with a CC of 430, without inbreeding depression, with a 4% harvesting rate per gestational year (translates to 5.4% in a calendar year) with and without supplementations. Supplemented individuals were either from the KZN metapopulation or the native *D. b. minor* Zimbabwe metapopulation. A. Supplementations made with one female and one male (1F, 1M) every one (E1Y), two (E2Y), five (E5Y) and ten (E10Y) gestational years. B. Supplementations were made with two females and two males (2F, 2M) every one (E1Y), two (E2Y), five (E5Y) and ten (E10Y) gestational years.

4.5 Discussion

The PVA analysis showed that the population size could be increased to the level needed to prevent the loss of genetic variation by expanding the size of the reserve or through supplementations. Doubling the model population size helped maintain the mean H_E , even after approximately 96 BRGs (H_E : 0.470 to 0.402 averaged over 10 microsatellite loci; 1000 gestational years /1342 calendar years), which was better than the significant loss seen when the population remained at 430 CC over the same time period (H_E = 0.470 to 0.347 averaged over 10 microsatellite loci; 1000 gestational years /1342 calendar years). Nevertheless, models also indicate that when supplementations were made with individuals from the KZN metapopulation the mean H_E improved more effectively than increasing the population size, even when supplementations only took place every ten gestational years (13.4 calendar years). However, the greatest effect on slowing the loss and increasing the mean H_E was when supplementations were made with native *D. b. minor* Zimbabwe individuals.

Currently, no translocations are made back into HiP. The only way to increase the size of the population is to expand the reserve beyond its current boundaries, which can be expensive and not always feasible. The SADC RMG supports sourcing same-subspecies individuals for founding populations from different original genetic sources including more than one source population and/or country (Emslie et al. 2009), which has been done successfully in Zambia with 25 *D. b. minor* sourced from several populations in South Africa (e.g. Kruger National Park, Eastern Cape, HiP, Markarele)(Chomba and Matandiko 2011). Guidelines do not, offer recommendations specifically for

supplementing source populations. I recommend that this gap in guidelines for managing source populations be addressed, especially when populations are unable to expand and have no natural or assisted immigration. Increasing the number of black rhino in HiP could slow the loss of genetic variation. No decrease in genetic variation has been found in small populations that experienced rapid population expansion in habitats with virtually unlimited CCs (e.g. 13 European rabbits (*Oryctolagus cuniculus*) imported to Australia in 1859; Zenger et al. 2003). Increasing the number of black rhino in the population can be done either by (1) physically increasing the size of the reserve allowing the CC to increase, thereby increasing the effective size (N_e) (2) using other reserves to mimic expansion by performing serial translocations between HiP and the metapopulation or other sources of *D. b. minor*. In this way, *D. b. minor* are both emigrating from and immigrating to HiP and the population can take advantage of an artificial increase, thereby limiting further deterioration of genetic variation.

Increasing black rhino numbers by increasing the physical size of HiP will probably help to decrease the rate of loss of genetic diversity, but it will not increase variation as effectively as genetic supplementation. Furthermore, the limits associated with increasing the size of the reserve (e.g. land acquisition, facilitating corridors) make this option less appealing to conservation managers. Supplementations on the other hand are relatively inexpensive by comparison and translocations in KZN occur on an annual basis, so this method could be easily incorporated into existing management plans.

Emslie (1994) states that there is no proof that a long-term decrease in heterozygosity of black rhinos will automatically diminish future performance.

While it is true that no obvious signs of inbreeding depression within HiP have been reported, the single mtDNA haplotype and 30% lower microsatellite DNA variation than native Zimbabwe *D. b. minor* and lower growth rates compared to other *D. b. minor* populations suggests that a precautionary approach to managing HiP may be necessary. Preserving current levels of genetic variation within HiP is important and by avoiding further genetic decay HiP can continue to improve as a source for restocking other populations demographically and genetically. An example of this was observed in an isolated population of Scandinavian grey wolves (*Canis lupus*: Vila et al. 2003). A single male immigrant led to an increase in the mean H_E from 0.49 to 0.62 of the small population (n=16) over a five-year period (Vila et al. 2003). The HiP population is much larger than the wolf example, but a similar change can be seen in the model population using supplemented individuals from the native *D. b. minor* Zimbabwe population.

It is clear from the PVA results that supplementation was more effective at arresting the rate of loss of genetic variation than maintaining the CC at current levels or increasing the size of the reserve. However, deciding whether to use black rhino from the KZN metapopulation or native Zimbabwe *D. b. minor* for supplementations is not straightforward. Unfortunately, it is unknown if the low levels of mtDNA and microsatellite DNA variation in HiP were anthropogenically induced or a result from a long-term demographic separation that caused historically small population sizes and local adaptation (Chapters Two and Three). If the population has always had low levels of variation, then supplementations with individuals from the KZN metapopulation would be recommended to avoid possible outbreeding depression that could compromise

the population's evolutionary potential. However, if it can be proven that the low variation is a consequence of overexploitation by humans and the population was continuous through Zimbabwe, then supplementations with native Zimbabwe *D. b. minor* would be recommended to restore genetic variation possibly to near pre-decline levels. The only way to answer this question is by examining samples of KZN black rhino from before the decline. PVAs model the best-case scenario but wild populations do not necessarily respond in the same way as a simulated population.

4.6 Conclusion

Detecting losses in the level of genetic diversity in an endangered source population like HiP calls attention to the importance of developing conservation strategies that prevent such losses in other endangered species. While, managing genetic risks by facilitating gene flow with minimal intervention is preferable, at this time expanding park borders is not a viable option for HiP. The HiP source population only has one mtDNA haplotype (n=65) (Anderson-Lederer et al. 2012) an H_E of 0.47, has a homozygote excess at five out of 10 microsatellite DNA loci and is out of HWE (Chapter Three). In addition, it is subjected to poaching pressure. While modelled results are only as reliable as the species life-history data used to create scenarios, management has the opportunity to shift from a focus from not only increasing numbers of black rhino, but ensuring the future of the species by incorporating results from PVAs into management schemes. This could improve the genetic health of HiP black rhino in a practical and evidence-based way. PVA results suggest that pseudo-metapopulation expansion through supplementations are effective and are not

required to be large or frequent (one female and one male every BRG). The supplementations can be accomplished inexpensively via the metapopulation using serial translocations. Most endangered large mammals no longer have the freedom to migrate, but with assistance via translocations and genetic supplementation, counteracting low levels of genetic variation that many small, fragmented populations exhibit is possible.

CHAPTER FIVE

Managing genetic diversity in the *D. b. minor* metapopulation of KZN: Thesis summary and applications



Photo by Rosalynn Anderson-Lederer

5.1 Introduction

Many threatened species are now conservation-reliant after suffering severe declines in population size (see: Miller et al. 1988; Walters 1991; Tyus and Saunders 2000; Jamieson et al. 2006; Johnson et al. 2010). The continued survival of these species requires *in situ* and *ex situ* intervention by conservation teams. Unfortunately, for pragmatic reasons population genetic considerations are often not a priority when trying to increase the inevitably low number of individuals remaining in small and fragmented populations. However, when populations are in the process of recovery, wildlife managers need to shift their focus from protecting and increasing numbers to addressing population quality indicators such as the loss of genetic variation.

A loss of genetic variation in small and recovering populations could result in decreased fitness (inbreeding depression) or limit the long-term capacity of a population to respond to changes in the environment (Westemeier et al. 1998). Without shifting focus to the genetic variation in the small populations, the genetic risks may be overlooked and populations might continue to be vulnerable to extinction even though their size is increasing (Frankham 2005).

This thesis research contributes to a broader understanding of the genetic structure of the metapopulation of *D. b. minor* in KwaZulu-Natal (KZN), South Africa. It also demonstrates how management can incorporate a simple supplemental regime in a source population, like Hluhluwe-iMfolozi Game Park (HiP), to maintain and/or increase levels of genetic variation within the endangered, but recovering metapopulation.

5.2 Summary of findings

5.2.1 Chapter Two:

Anderson-Lederer, R.M., Linklater, W.L., and P.A. Ritchie (2012) Limited mitochondrial DNA variation within South Africa's black rhino (*Diceros bicornis minor*) population and implications for management. African Journal of Ecology 50(4): 404-413. (Appendix A)

Mitochondrial DNA (mtDNA) control region sequences (406 bp) were examined to determine the level of variation within the KZN *D. b. minor* source population at HiP (n=50) and KZN metapopulation (n=15) compared to *D. b. minor* populations outside South Africa (n=11; native Zimbabwe populations) and the two other black rhino subspecies (*D. b. michaeli* n=21, *D. b. bicornis* n= 4). The KZN source (HiP) and metapopulation had a single haplotype. However, six different haplotypes were represented in the 11 native *D. b. minor* individuals from Zimbabwe. The *D. b. michaeli* samples (n=21) had 13 haplotypes and the *D. b. bicornis* samples (n=4) had one haplotype. The single mtDNA haplotype in the KZN source and metapopulation coupled with previously published low levels of *D. b. minor* mixed population (KZN and native Zimbabwe) microsatellite DNA suggest a recent population decline and fragmentation. Small population numbers combined with fragmentation could have increased the rate of drift and may be responsible for the lack of haplotype diversity within the KZN HiP source and metapopulation. Further investigation of the native Zimbabwe *D. b. minor* mtDNA should be considered because the KZN haplotype may be present, but not yet detected. If the KZN haplotype is found in the Zimbabwe populations, management may also want to explore the possibility of

outbreeding the native Zimbabwe *D. b. minor* populations with the translocated KZN *D. b. minor* population in Malilangwe, Zimbabwe.

5.2.2 Chapter Three:

Low levels of microsatellite DNA variation and possible management considerations for black rhino (*Diceros bicornis minor*) in KwaZulu-Natal, South Africa (In prep)

Ten microsatellite DNA loci were examined in the three *D. bicornis* subspecies:

D. b. minor from the KZN source and metapopulation (n=118), *D. b. michaeli* (n=3) and *D. b. bicornis* (n=6). Results were compared with previously published findings for microsatellite DNA loci from a native Zimbabwe *D. b. minor* populations. The KZN source and metapopulation was out of Hardy-Weinberg Equilibrium and showed excess homozygosity at five loci. Direct comparison of 10 microsatellite DNA loci results confirmed that the South African metapopulation has lower genetic variation than the native Zimbabwe *D. b. minor* populations (H_E : 0.47 and 0.65 respectively, and lower number of alleles per locus), further indicating that current conservation plans may need to be modified to prevent additional genetic decay within KZN.

Additional loss of genetic variation and the possible risk of inbreeding depression could be prevented by rapidly increasing population numbers by increasing reserve sizes. Since procuring enough land to increase populations to sizes that would enable rapid growth is expensive and not always feasible, managers may consider serial translocations between HiP and other KZN reserves to replicate immigration and emigration. Moving rhino back into HiP would enable the source to take advantage of growth in other reserves and

would essentially reinstate a single large genetic population. However, since native Zimbabwe *D. b. minor* have higher levels of genetic variation (mitochondrial and microsatellite DNA loci) than the KZN *D. b. minor*, managers could also consider genetic replenishment using native Zimbabwe individuals as supplements.

5.2.3 Chapter Four:

Population Viability Analysis of Black Rhino (*Diceros bicornis minor*) in Hluhluwe-iMfolozi Game Park, KwaZulu-Natal, South Africa

The KZN *D. b. minor* source and metapopulation have low mitochondrial DNA (Chapter Two) and microsatellite DNA (Chapter Three) variation. It was proposed in Chapters Two and Three that increasing the size of the HiP source to accommodate rapid growth may slow the rate of genetic drift and help to maintain current levels of genetic variation. It was also suggested that supplementation with individuals from the KZN metapopulation or native *D. b. minor* Zimbabwe populations may be equally effective at preserving current levels of genetic variability. To test whether one or both of the recommendations are viable options, a population viability analysis (PVA) was conducted.

The PVA modelled population increases and supplementations into HiP with individuals from the KZN metapopulation and Zimbabwe. With no change in management strategies, the PVA predicted a progressive loss in the mean expected heterozygosity of 23% over 96 black rhino generations (BRGs). Opportunities to increase the HiP population size by connecting the reserve

through corridors to neighbouring reserves was modelled by increasing the carrying capacity (CC) of the population. Doubling the CC helped to decrease the rate of loss of the mean H_E by 11% over 96 BRGs. When supplementations of one female and one male black rhino from the KZN metapopulation were made every ten gestational years, the mean H_E of the population was maintained ($H_E \sim 0.45$) over 96 BRGs, but increased 29% when supplemented with native *D. b. minor* from Zimbabwe.

The PVA results show that serial translocation between populations is a powerful tool that can be used to decrease the rate of loss of genetic variability. In addition, it does not require a large number of individuals or to be frequent and corresponds with current management practices.

5.3 Conservation Implications

Data obtained in this study will allow field managers to forecast the likely changes in the levels of genetic variation within the relict HiP source population. Genetic variation in a population arises through mutation or gene flow and is typically lost either passively through genetic drift or actively through natural selection (Amos and Harwood 1998). The rate of loss of genetic variation in a population depends on its effective population size (N_e) and the amount of time (number of generations) that the population has been isolated (Frankham 1997; Palstra and Ruzzante 2008). The loss of genetic variation caused by genetic drift increases in a population that has gone through a bottleneck and strong drift will continue to erode genetic variation if a population's size is unable to increase (Nei et al. 1975; Allendorf and Leary 1986b). Bottlenecks occur across many taxa, but origin, severity and population recovery times vary (i.e. they can

develop rapidly over a short period of time in small populations or take several generations in large populations) (England et al. 2003).

Heterozygosity and allelic diversity (also called allelic richness) in populations are correlated when there is a mutation-drift balance, but after a short-term bottleneck correlation is disrupted because heterozygosity is only slightly affected (Allendorf 1986). Rare alleles on the other hand are typically lost when population sizes decrease, which means allelic diversity is expected to decrease faster than heterozygosity (Allendorf 1986; England et al. 2003).

The low levels of heterozygosity and allelic diversity, and the similar F_{ST} and R_{ST} results in HiP compared to native Zimbabwe *D. b. minor* are indicative of a recent (although not necessarily within $2N$ to $4N$ generations) decrease in HiP's population size. Recent bottlenecks have also been linked to reduced or low levels of genetic variation in other rhino taxa (Merenlender et al. 1989; Dinerstein and Mccracken 1990; Harley et al. 2005; Fernando et al. 2006), which is not unexpected given the high level of poaching that has occurred during the last century.

Regardless of whether the decreased level of genetic variation in the relict *D. b. minor* HiP population is due to the 20th century bottleneck or historic demographic separation, it is necessary to take corrective action to prevent any further loss. The HiP population is small, fenced and has had no immigration since the latter half of the 19th century (Swart et al. 1994), yet it is used as a source for *D. b. minor* subspecies expansion. HiP is also subjected to an annual harvest that targets removal of ~5% of the total population each year. The removals are meant to keep the population below its CC, but this practice ultimately stagnates the size of the population, which could be having

detrimental effects on the level of genetic variation.

This study has demonstrated that despite *D. b. minor* numbers increasing, HiP is genetically a relatively small population ($n \sim 220$ Clinning et al. 2009) and genetic drift has most likely caused genetic variability to be lost faster than would be expected in what might appear to be a larger metapopulation. Managers of any small, endangered source population should evaluate conservation plans once a species is classified as in recovery, to evaluate the levels of genetic variation and adjust priorities from quantity (number of individuals) to quality (evolutionary potential).

Using serial translocations and supplementations to maintain the levels of genetic variation in the relict HiP source population has important implications. Managers of other small, remnant populations of conservation reliant species may be able to avoid the expense of acquiring land to ‘expand’ their population, by using serial translocations and supplementation to increase gene flow. Supplementation proved to be an effective tool for a remnant population of greater prairie chickens (*Tympanuchus cupido pinnatus*) that experienced 35 years of population declines and a reduction in genetic diversity (Westemeier et al. 1998). Despite three decades of intensive management that focused on and successfully increased the number of individuals in the population, the mean population fitness decreased to alarming levels and it was only turned around when the remnant population was supplemented with individuals from a larger population with higher levels of genetic variation (Westemeier et al. 1998).

With regards to genetic supplementation of new and established populations, management of African rhinos (black and white), the SADC RMG

recommend that cohorts used for supplementations consist of individuals with “as little genetic similarity with the receiving population as possible” (Emslie and Brooks 1999). They also recommend that when establishing new populations, animals should not be from the same genetic source (Emslie and Brooks 1999) sensu North Luangwa National Park, Zambia (Chomba and Matandiko 2011) and Kruger National Park, South Africa (Hall-Martin and Knight 1994). These recommendations are important because if individuals with similar genotypes are used to establish or augment populations it could have a negative impact on fitness and long-term evolutionary potential of the receiving population (Eldridge et al. 1999; Moritz 2002).

Rhino guidelines encourage the mixing of populations of the same subspecies (du Toit 2006b; Emslie and Brooks 1999), however, there are instances of black rhino management taking measures to prevent the same subspecies from different regions from intermixing (i.e. the 27 KZN *D. b. minor* translocated to Malilangwe, Zimbabwe in 1997 that are not outbred with native Zimbabwe black rhino). There is a risk of over splitting the subspecies if *D. b. minor* populations (KZN and native Zimbabwe) are treated as separate subspecies. Managing *D. b. minor* as one large metapopulation (e.g. similar to the translocations into Krueger National Park from South Africa and Zimbabwe) across borders may aid with the overall management goals of retaining genetic diversity, but may not be feasible due to manpower (e.g. needed for anti-poaching) and budget constraints (e.g. small GDP economies).

5.4 Future research directions

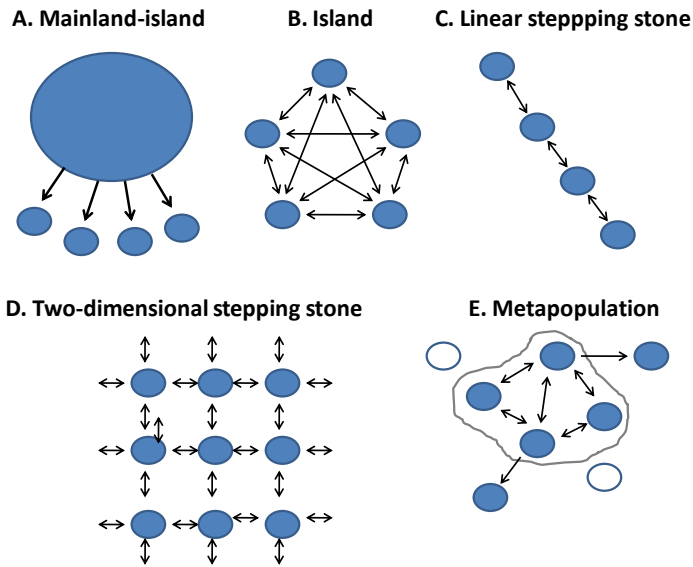
5.4.1 *Historic Samples*

Determining whether the current levels of genetic diversity in an endangered species population are significantly different from historic levels is difficult because it requires 'before and after' samples of the population (Leberg 2002). If both historic and current samples could be obtained, it would confirm whether or not a decline in the level of genetic variation has occurred (Roy et al. 1994). Without temporal genetic data, assessments of the amount of genetic diversity lost in populations can only be inferred by comparing them to examples of closely related species, or populations of the same species in different geographic locations (Leberg 2002).

Sampling historic levels of genetic variation may also provide some insight into the past population structure and the amount of fragmentation that has occurred. Frankham et al. (2002) described five fragmented population structures (Figure 5.1) that have various impacts on levels of variation within populations ranging from negligible to severe, depending on the structure of and migration patterns between the fragments. Identifying the historic pattern of fragmentation could help managers plan metapopulations and allow duplication of past gene flow patterns through the creation of corridors or through translocations.

Information derived from historic samples could also reveal that very little loss in the level of genetic variation has occurred and that the populations were always small and isolated. These findings would assist when implementing management practices that might lead to outbreeding depression

Figure 5.1: The five structures of fragmented populations. (From Frankham et al. 2002)



(Leberg 2002). In the case of rhino, historic samples might be obtained through zoo collections from the progeny of individuals originally captured in populations that have since been extirpated. They might also be collected from museum samples if documentation regarding the sample's origin is verified.

5.4.2 Harvest for Population Growth

As large wild populations of threatened and endangered species begin to recover, it will become necessary for managers to address limited reserve sizes and the need for harvesting populations when they reach maximum carrying capacities.

Translocations are used for increasing the viability of a species.

Specifically, the technique is used to (1) move individuals between wild populations to bolster genetic heterogeneity of small populations, (2) move wild individuals to captivity (also called capture or collection) and (3) to move

individuals or cohorts from captivity to the wild (also called reintroduction or release) (Griffith et al. 1989; Tenhumberg et al. 2004).

Apart from megaherbivores, translocations are not typically used to remove (or ‘harvest’) individuals from endangered populations that have reached their carrying capacity in order to stimulate continued population growth (see Appendix B for PVA results pertaining to HiP harvest rates) (Emslie et al. 2007; Emslie et al. 2009; Emslie 2001). Because the use of harvesting to facilitate breeding may become more popular as large species conservation programmes move into a recovery phase, shifting priorities to that of preserving levels of genetic variation will be important and warrants a more thorough examination (Lubow 1996).

5.4.3 Translocation Cohort Sizes and Composition

As translocation techniques are improved, reintroductions are becoming more important in the management of threatened species. There is little species specific information available, however, about the recommended number and composition (e.g. number of males and females, ages, parent-offspring) of individuals necessary in a cohort to minimize losses in levels of genetic diversity (Tracy et al. 2011).

Most researchers do not define what an ‘acceptable’ or ‘adequate’ number equates to in numeric terms, which makes planning the composition of cohorts for successful translocations and maintenance of levels of genetic diversity difficult for conservation managers. While Griffith et al. (1989) only refer to ‘large’ versus ‘small’ cohort sizes, they found that translocations had higher success rates among individuals and cohorts that were native game

species (86%), herbivores (77%), and wild-caught individuals (75%). Knapp and Dryer (1998) recommend using individuals that are a 'genetic match' or that already have local adaptations to the environment to improve the success rates. They found that individuals used for reintroductions that had adaptations to local conditions had higher fitness and lower mortality than individuals that were translocated to areas they were not adapted to (see Appendix C for PVA results pertaining to HiP cohort sizes).

As field managers are increasingly using translocations and reintroductions as tools for managing wild populations, improving the success of reintroductions while maintaining levels of genetic variation is an area that requires further investigation.

5.4.4 Functionally important genetic variation and Conservation Genomics

To successfully conserve an endangered species, it is important to: (1) identify different species within taxa, (2) identify evolutionarily significant units (ESUs) within species, and (3) identify management units (MUs) within ESUs (Moritz 1994; Hedrick et al. 2001). Examining the relative levels of genetic variation at neutral markers like mitochondrial or microsatellite DNA loci, can reveal historic population characteristics that support ESU and MU designation (Bos et al. 2008), however maintaining adaptive variation is important to the conservation objective that supports ESU and MU designations (Hedrick et al. 2001).

While demographic factors (e.g. genetic drift and inbreeding) play a crucial role in the degree of variation in neutral genes, some functional (coding) genes are under selection (Sommer 2003). Major histocompatibility complex

(MHC) genes are functional genes that are primarily responsible for recognizing foreign proteins, presenting them to immune cells and initiating an immune response (Pirotney and Oliver 2006). MHC genes have been connected with individual fitness, population viability and evolutionary potential in changing environments, which makes them ideal for studying adaptive genetic diversity (Strand 2011). However, due to the complexity of working with MHC genes (organization of MHC loci found in model species differs from non-model species), many large-scale studies of wild populations have been hindered (Strand 2011).

MHC primers for this study (Dbminor-MHC Alpha1-2 Forward: CCTCCTCCTGCTCTCGG and Dbminor-MHC Alpha 1-2 Reverse: CCACAGCCGCCCACTTCTGG) were developed using Geneious v4.8 (Drumond et al. 2010). Remaining DNA originally extracted for Chapters Two and Three was used with the new primer pair. The forward primer was tagged with an M-13 tag (Schuelke 2000). Polymerase Chain Reaction (PCR) amplifications using 1-2 μ L of DNA template were carried out in 25 μ L volumes with 67 mM Tris pH 8.8, 16mM $(\text{NH}_4)_2\text{SO}_4$, 1.5 mM MgCl_2 , 0.4 $\mu\text{g}/\text{ml}$ BSA, 0.4 μL of each of the forward and reverse primer, 200 μM of each dNTP, and 0.5 to 1 units of BIOTAQ DNA polymerase (Bioline). Thermal cycling was carried out using an Eppendorf Mastercycler for; 94°C 2 min, (94°C 3 min, 50-54°C 30 sec, 72°C 2 min), repeated for 30-40 cycles, followed by a final step of 72°C 3 min. PCR products were electrophoresed in agarose gel and a molecular weight standard was used to determine the size of amplified products. The resultant PCR products were analysed on a 3730 automated sequencer using the GS-500 LIZ size standard and the GENESCAN software (Applied Biosystems) I experienced significant

difficulties amplifying MHC genes to a level of quality that would allow clear sequencing for this project.

Even though neutral and functional genes provide complimentary information about the recent evolutionary history of populations, only a small number of studies have actually used a combination of the markers to examine the relative levels of genetic variation in populations (Bos et al. 2008).

Conservation genomics is a new field that utilises recent advances in DNA sequence techniques to discover genomic regions that are adaptively important for populations in particular habitats (Allendorf et al. 2010).

Most conservation genetic studies use neutral markers that are unable to answer questions about the impact of a small effective population size (N_e) on functionally important genetic variation or whether microsatellite DNA variation provides a good representation of the genome wide levels of variation (Ouborg et al. 2010). Genomics will enable the field of conservation genetics to understand both the mechanistic (e.g. genetic and cellular operations and how they affect organismal development, ecology and evolution) and inventorial (functional, e.g. improve ability to take genealogical stock of biological resources at all levels in the phylogenetic hierarchy) aspects of at risk species (Avis 2010). The diagram in Figure 5.3 shows how conservation genomics can address a range of conservation genetic issues that previous techniques have been unable to address.

Conservation genetic analyses are typically based on neutral loci that only provide details of a very small part of an organism's genome, however with advances in conservation genomics (e.g. SNPs), researchers will be able to analyze genome-wide data that will translate into practical information that can

significantly improve the way endangered species are managed. However, this new technical approach is expensive and requires detailed computational analyses, which will limit its uptake in the short term.

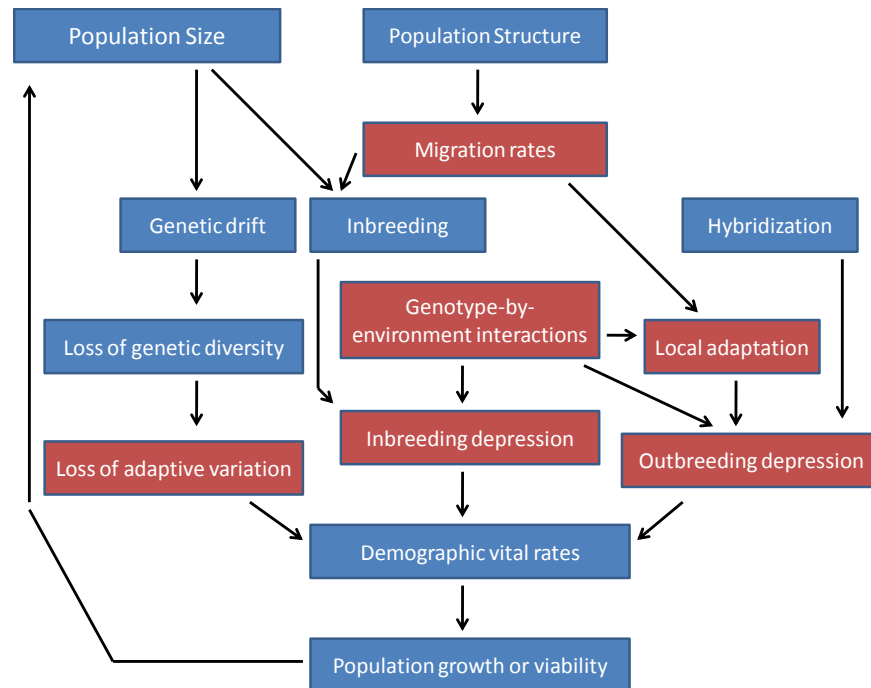


Figure 5.3: A diagram of interacting factors in the conservation of natural population. Traditional conservation genetics (neutral markers), provides direct estimates of some interacting factors (blue). Conservation genomics can address a wider range of factors (red). It also promises more precise estimates of neutral processes (blue) and understanding of the specific genetic basis of all of the factors. (From Allendorf et al. 2010)

5.6 Summary

In this study, I have added to the understanding of the genetic structure of the KZN remnant source (HiP) and metapopulation, highlighted genetic differences between KZN and native Zimbabwe *D. b. minor* as well as made recommendations for maintaining the current levels of genetic heterozygosity within HiP. This is timely as the black rhino species is recovering. If poaching

pressure can be abated, black rhino numbers may be able to match those of the southern white rhino that have been the most successful of the rhino taxa to recover from a severe bottleneck. I anticipate that this research will help to improve management of black rhino populations and contribute to conservation biology on a broader scale across species.

As rapid changes in the field continue, conservation genetics will bring to light historic and current stochastic and demographic changes occurring in wild populations. Techniques are advancing turning genetics and genomics into incredibly powerful tools for assisting in the management of populations across all levels (individual, population, and species). By wildlife managers taking research findings and incorporating them into management plans, the field of conservation genetics and genomics can help slowly reverse the process of destruction that human induced changes have caused.

Appendix A

African Journal of Ecology

Limited mitochondrial DNA variation within South Africa's black rhino (*Diceros bicornis minor*) population and implications for management

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Abstract

The taxonomy of African black rhinoceros (*Diceros bicornis*) remains unresolved. Maintaining levels of genetic diversity and species rescue by reintroduction and restocking requires its resolution. We compared the sequences of the mitochondrial DNA (mtDNA) control region for a total of 101 *D. bicornis* from three subspecies: *D. b. minor*, *D. b. michaeli* and *D. b. bicornis*. A single unique haplotype was found within the 65 *D. b. minor* samples from KwaZulu-Natal (KZN) Province, South Africa, 55 of which came from Hluhluwe-iMfolozi Game Park (HiP) and Mkuzi Game Reserve (MGR) source populations. However, six different haplotypes were represented in eleven *D. b. minor* samples from Zimbabwe. Similarly, published autosomal microsatellite data indicate low levels of diversity within the KZN *D. b. minor* populations. The low levels of mtDNA diversity within the KZN metapopulation point to the possible need for genetic supplementation. However, there is a need to determine whether the low levels of genetic variation within KZN *D. b. minor* are a result of the recent bottleneck or whether KZN historically always had low diversity.

Key words: conservation genetics, control region, *Diceros bicornis*, D-loop, genetic rescue

Résumé

La taxonomie du rhinocéros noir d'Afrique (*Diceros bicornis*) n'est pas encore résolue. Pour préserver le taux de diversité génétique et pouvoir sauvegarder l'espèce par des réintroductions et des repeuplements, il faut résoudre cette

question. Nous avons comparé les séquences de la région de contrôle de l'ADN mitochondrial (ADNmt) d'un total de 101 *D. bicornis* appartenant aux trois sous-espèces *D. b. minor*, *D. b. michaeli* et *D. b. bicornis*. Nous avons trouvé un seul et unique haplotype pour les 65 échantillons venant de *D. b. minor* de la province du KwaZulu-Natal (KZN) en Afrique du Sud, dont 55 venaient de populations sources du Parc de Hluhluwe-iMfolozi (HiP) et de la Réserve de Faune de Mkuzi (MGR). Par contre, il y avait six haplotypes différents dans 11 échantillons de *D. b. minor* venant du Zimbabwe. Les données publiées sur les microsatellites autosomaux indiquent, elles aussi, un faible taux de diversité au sein des populations de *D. b. minor* du KZN. Le faible taux de diversité de l'ADNmt dans la métapopulation du KZN indique un éventuel besoin de supplémentation génétique. Cependant, il faut d'abord déterminer si le faible taux de variation génétique chez les *D. b. minor* du KZN est un résultat des réductions récentes ou si le KZN a toujours eu une faible diversité.

Introduction

Species conservation depends on identifying genetically distinct groups or management units and implementing strategies to retain genetic variation. Genetically distinct populations can contain unique genetic variation, and/or they can be locally adapted to their habitat. Mixing them with other populations may break up genetically complex traits and, in some cases, lead to outbreeding depression (Templeton, 1986; O'Ryan, Flamand & Harley, 1994). Alternatively, genetic differences between populations can also result from strong genetic drift caused by population fragmentation and declining

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population sizes (Frankham, Ballou & Briscoe, 2002; Allendorf & Luikart, 2007). When the genetic structure and historic pattern of gene flow of a species has been described, reintroduction methods can be used to secure locally adapted populations or restocking used for genetic supplementation.

Variation in mitochondrial DNA (mtDNA) is a particularly useful metric for determining population structure and history (Moritz, 1994). The control region of mtDNA is highly variable, and it can often be used to resolve phylogenetic relationships between closely related taxa or for describing the genetic structure within species (Moritz, Dowling & Brown, 1987; Kidd & Friesen, 1998). MtDNA is maternally inherited and so does not recombine (Hayashi, Tagashira & Yoshida, 1985), which means it reflects a quarter the effective population size (N_e) compared with nuclear loci and hence it is more sensitive to changes in population demography.

The black rhinoceros (*Diceros bicornis*; Perisodactyla) once ranged across the African continent and numbered in the hundreds of thousands (Western & Vigne, 1985). By 1969, their numbers had declined to ~65,000 (Muya & Ouge, 2000), and during the last century, the species disappeared faster than any other large mammal (Hitchins, 1975; Western & Vigne, 1985). The major causes for their decline have been anthropogenic, primarily illegal hunting (Western & Vigne, 1985; Emslie & Brooks, 1999; Amin *et al.*, 2006). Nevertheless, conservation efforts have seen *in situ* black rhino numbers increase from a low of 2475 individuals in 1993 to approximately 4880 in 2010 (Emslie, 2011).

Three extant black rhino subspecies are recognized across Africa, including approximately 742 *D. b. michaeli* (Eastern black rhino), 1922 *D. b. bicornis* (South-western black rhinoceros) and 2216 *D. b. minor* (South-central black rhinoceros) (Emslie, 2011). Appraisal of the black rhino subspecies was initially based on skull measurements (Zukowsky, 1964; Groves, 1964; du Toit, 1987); however, uncertainty regarding taxonomy remained (du Toit, 1987). Although there are apparently no impervious geographic boundaries or reproductive barriers between the subspecies, they occupy different areas with distinct habitats and climates (Harley *et al.*, 2005; Emslie & Brooks, 1999). With no historical records of migration and the extent of gene flow between the subspecies unknown, some authors have speculated that each subspecies may have genetic or behavioural adaptations to

their local environments (Emslie & Brooks, 1999; Harley *et al.*, 2005). Their suggestion regarding genetic differences was confirmed through recent mtDNA and autosomal DNA analyses (Merenlender *et al.*, 1989; Ashley, Melnick & Western, 1990; O'Ryan & Harley, 1993; O'Ryan, Flamand & Harley, 1994; Swart & Ferguson, 1997; Brown & Houlden, 1999, 2000; Nielsen *et al.*, 2008; Karsten *et al.*, 2011; Muya *et al.*, 2011). Thus, current black rhino management policy is for each subspecies to be managed separately in order to maintain possible local adaptive traits and minimize the risk of outbreeding depression (Templeton, 1986; O'Ryan, Flamand & Harley, 1994; Brown & Houlden, 2000; Harley *et al.*, 2005).

The largest remnant population of the critically endangered (IUCN, 2008) *D. b. minor* subspecies is in Hluhluwe-Imfolozi Game Park (HiP) ($n = \sim 220$ Clinning *et al.*, 2009) in KwaZulu-Natal (KZN) Province, South Africa (Fig. 1). KZN black rhinos have been separated from other populations to the north (e.g. Zimbabwe) since at least the latter half of the 19th century (Swart *et al.*, 1994). HiP and the smaller remnant in Mkuze Game Reserve (MGR) ($n = \sim 45$ D. Kelly, personal communication) have been sources for metapopulation expansion and genetic management by reintroduction and restocking. Translocations from HiP to other KZN reserves first began in 1962, expanded to other South African provinces and later to other African nations (e.g. Zimbabwe, Zambia) (Hitchins, 1984; Emslie, Amin & Kock, 2009). The potential now exists for KZN *D. b. minor* to be mixed with *D. b. minor* in or from other smaller African populations, especially those in Zimbabwe, if they are not too genetically divergent. Although the KZN population will likely be strategic to the subspecies recovery throughout the African continent (Emslie & Brooks, 1999), no study has yet compared the mtDNA sequences of the KZN *D. b. minor* metapopulation with populations outside South Africa.

The aim of our study was to use mtDNA control region sequences (406 bp) to determine the level of variation within the *D. b. minor* source population at HiP ($n = 50$) and compare it with the KZN metapopulation ($n = 15$) and *D. b. minor* populations outside South Africa ($n = 11$) and the other black rhino subspecies (*D. b. michaeli* $n = 21$, *D. b. bicornis* $n = 4$). We considered the implications of our findings for the long-term management of *D. b. minor* and make recommendations for possible future research.



Fig 1 Map of southern Africa showing black rhinoceros sample sites. Inset showing KwaZulu-Natal study Game Reserves (Nduro G.R., Ithala G.R., Mkuze G.R., Hluhluwe-Imfolozi Game Park (HiP) and Weenan G.R.).

Materials and methods

Sampling

Samples of blood and pinna ear tissue were collected from individuals of *D. b. minor* in the KZN province in South Africa ($n = 65$), *D. b. michaeli* in Addo Elephant National Park, South Africa ($n = 1$), and *D. b. bicornis* in Namibia's northern region ($n = 4$) (Fig. 1). The samples were acquired opportunistically during routine translocation and ear notching (for identification) events from 2002 to 2009. Blood samples were stored in cryovials

containing 1 ml of DMSO/EDTA/Tris/salt solution (Seutin, White & Boag, 1991).

DNA sequencing and analysis

DNA extraction. Seventy micro-litres of the preserved blood solution or a 3×3 mm piece of pinna ear tissue was digested in an SDS/proteinase-K solution. After dissolution, a standard phenol-chloroform DNA extraction and ethanol precipitation was conducted following the procedure of Sambrook, Fritsch & Maniatis (1989).

PCR and DNA sequencing. A fragment of the mitochondrial DNA control region (406 bp) was amplified using the primers mt15996L (5'-TCCACCATCAGCACCCAA AGC-3') (Campbell *et al.*, 1995; Brown & Houlden, 2000) and mt16502H (5'-TTTG-ATGGCCCTGAAGTAAGAACA-3') (Moro *et al.*, 1998; Brown & Houlden, 2000). PCR amplifications using 1–2 μ l of DNA template were carried out in 25 μ l volumes with 67 mM Tris pH 8.8, 16 mM $(\text{NH}_4)_2\text{SO}_4$, 1.5 mM MgCl_2 , 0.4 $\mu\text{g ml}^{-1}$ BSA, 0.4 μ l of each of the forward and reverse primer, 200 μM of each dNTP and 0.5–1 units of BIOTAQ DNA polymerase (Bioline USA Inc., Taunton, MA, USA). Thermal cycling was carried out using an Eppendorf Mastercycler at 94°C for 2 min, (94°C for 3 min, 50–54°C for 30 s, 72°C for 2 min), repeated for 30–40 cycles, followed by a final step of 72°C for 3 min.

PCR products were electrophoresed in agarose gel, and a molecular weight standard was used to determine the size of amplified products. Products of the correct size were purified using column purification (Roche Corporate Communications, Basel, Switzerland) or ExoSAP-IT (GE Healthcare, Waukesha, WI, USA), and their DNA sequence was determined using an ABI 3730 Genetic Analyzer (Massey Genome Service, Palmerston North, New Zealand).

For comparison, eleven *D. b. minor* sequences stored in GenBank (accession numbers AF187825 - AF187827 & AF187829 - AF187831, Brown & Houlden, 2000; AY742832 & AY742833, Fernando *et al.*, 2006) originally sampled from Zimbabwe and zoos in Australia and the United States were added to the data set, in addition to 20 *D. b. michaeli* samples (accession numbers AF187834 & AF187835, Brown & Houlden, 2000; AY742830 & AY742831, Fernando *et al.*, 2006; FJ227484 - FJ227498, Muya *et al.*, 2011) originally

sampled from Kenya and zoos in Australia and the United States.

Data analysis. The 101 mitochondrial DNA sequences were edited by eye and then aligned using Clustal W (Larkin *et al.*, 2007). Homogeneity of base compositions was tested using PAUP 4.10b (Swofford, 2002). DnaSP v 5.10.1 (Rozas *et al.*, 2003) was used to calculate haplotype diversity (h), nucleotide diversity (p) and standard deviation (SD) within the subspecies. The level of sequence divergence within and between populations was estimated using a pairwise distance analysis in MEGA 5.1 (Tamura *et al.*, 2011), and standard errors were calculated using a bootstrap procedure. A statistical parsimony haplotype network was calculated using NETWORK 4.610 (Bandelt, Forster & Rohl, 1999).

Results

The sequence of the mtDNA control region was determined for a total of 70 individual black rhinos as follows: *D. b. minor* samples: 50 from HiP, eight from Itala, five from MGR, one from Ndumo Game Reserve, one from

the Johannesburg Zoo (accession number JN593089) and eleven sequences from GenBank (accession numbers AF187826 - AF187831, AY742832 - AY742833 & AF187832 - AF187833); *D. b. michaeli* samples: one from Addo Elephant Park (Accession number JN5930090) and 20 from GenBank (accession numbers FJ227483 - FJ227498, AY742830 - AY742831 & AF187834 - AF187835) and four samples for *D. b. bicornis* from Namibia's northern region (accession numbers JN593091-JN593094) (Table 1).

The 101 aligned sequences were 363 bp long with 31 polymorphic sites, and there was an average pairwise difference of 4% ($\pm 1\%$) between *D. b. michaeli* and *D. b. minor*, 4.5% ($\pm 1.1\%$) between *D. b. michaeli* and *D. b. bicornis* and 2.3% ($\pm 0.8\%$) between *D. b. minor* and *D. b. bicornis*. No insertions or deletions were observed.

Considering each subspecies separately, the greatest level of diversity was recorded in *D. b. michaeli* ($n = 21$), which contained thirteen haplotypes and showed comparatively high nucleotide diversity ($p = 0.011 \pm 0.00106$) and haplotype diversity ($h = 0.958 \pm 0.026$) (Table 2). The lowest level of diversity within a subspecies was seen in the Namibian *D. b. bicornis* samples

Table 1 Rhinoceros subspecies and sources analysed for mtDNA variation

Subspecies	Sample size	Sample/sequence source	References
<i>Diceros bicornis minor</i>	50	Hluhluwe-Imfolozi Game Park, KNZ Region, Accession number JN593089	This study
	8	Itala Game Park, KNZ Region	This study
	5	Mkuze Game Park, KNZ Region	This study
	1	Ndumo Game Park, KNZ Region	This study
	1	Johannesburg Zoo	This study
	6	Chete National Park, Zimbabwe, Accession numbers AF187825-AF187827 & AF187829-AF187831	Brown & Houlden, 2000;
	2	Zambezi Valley, Zimbabwe, Accession numbers AY742832-AY742833	Fernando <i>et al.</i> , 2006;
	1	Captive born, San Diego Zoo Accession number AF187832	Brown & Houlden, 2000;
	1	Captive born, Milwaukee Zoo, Accession number AF187833	Brown & Houlden, 2000;
	1	Captive born, Western Plains Zoo Accession number AF187828	Brown & Houlden, 2000;
<i>D. b. michaeli</i>	16	Kenya, Accession numbers FJ227483-FJ227498	Muya 2011
	2	Solio Game Reserve, Kenya, Accession numbers AY742830-AY742831	Fernando <i>et al.</i> , 2006;
	1	Cincinnati Zoo, Accession number AF187834	Brown & Houlden, 2000;
	1	Taronga Zoo, Accession number AF187835	Brown & Houlden, 2000;
	1	Addo Elephant Park, South Africa Accession number JN593090	This study
<i>D. b. bicornis</i>	4	Etosha, Namibia, Accession numbers JN593091-JN593094	This study
<i>Ceratotherium simum</i>	4	HiP & London Zoo, Accession numbers AF187836-AF187839	Brown & Houlden, 2000

	n	Genetic variability				
		H	h	(SD)	π	(SD)
Subspecies						
<i>Diceros bicornis minor</i> (aggregate)	76	7	0.267	0.067	0.0023	0.001
KZN metapopulation	65	1	—	—	—	—
Zimbabwe samples	11	6	0.855	0.085	0.0074	0.001
<i>D. b. michaeli</i>	21	13	0.952	0.024	0.0112	0.001
<i>D. b. bicornis</i>	4	1	—	—	—	—

Sample size (n), Number of haplotypes (H), haplotype diversity (h), Nucleotide diversity (π), Standard deviation (SD).

(n = 4) where only one unique haplotype was found; however, this was based on a small sample size and might not represent the total amount of genetic variation within the population. The pooled KZN samples and GenBank sequences of all *D. b. minor* individuals (n = 79) contained seven haplotypes, and haplotype diversity (h) was 0.267 ± 0.067 and a nucleotide diversity (π) of 0.002 ± 0.00063 . The eight *D. b. minor* Zimbabwe sequences from Brown & Houlden (2000) and two from Fernando *et al.* (2006) had shared haplotypes (Table 3); however, there were no shared haplotypes with the KZN samples.

The haplotype network (Fig. 2) shows a clear pattern of the separation amongst the three currently recognized subspecies with the KZN population falling out with the *D. b. minor* populations of Zimbabwe. Our finding of no more than three base pair substitutions between adjacent haplotypes within the *D. b. michaeli* subspecies is consistent with the finding by Muya *et al.* (2011). There is a significant separation between *D. b. minor* and *D. b. bicornis* with eight base pair substitutions as well as between *D. b. minor* and *D. b. michaeli* with nine base pair substitutions.

Discussion

We showed that the KZN population of *D. b. minor* is fixed for a single mtDNA haplotype, like most Sumatran rhino (*Dicerorhinus sumatrensis*) populations (Morales *et al.*, 1997). However, unlike the Sumatran rhino populations that have occupied separate land masses for more than 10,000 years (Morales *et al.*, 1997), it has been widely assumed that the KZN *D. b. minor* population has been separate from other *D. b. minor* populations only recently (i.e. caused by anthropogenic settlement and habitat modification during the 19th century, Swart *et al.*, 1994). The single mtDNA haplotype in KZN

D. b. minor raises the question of whether the KZN remnant population lost genetic variation recently because of the population bottleneck or has been a genetically separate lineage for longer than previously thought.

MtDNA has a smaller effective population size (N_e) compared to nuclear loci and is one of the first genetic markers to show the genetic signature of a demographic decline. The likelihood of two or more mtDNA haplotypes persisting within an isolated population is reduced to $P < 0.1$ over $4N_e$ generations, and the population is expected to become monophyletic after $4N_e$ generations (Avice, Neigel & Arnold, 1984; Mucci *et al.*, 1999). If this holds true for the KZN *D. b. minor*, then recent population decline and fragmentation would have increased the rate of drift and might be responsible for the lack of haplotype diversity within the KZN black rhinoceros. Examples of monomorphic haplotypes occurring from severe bottlenecks are well documented in several species. For instance, the Whooping Crane (*Grus americana*) once found throughout North America had six haplotypes in ten prebottleneck museum samples, but only one haplotype persisted in the remnant postbottleneck population of 14 (Glenn, Stephan & Braun, 1999). Such rapid declines in genetic variation have also occurred amongst southern Africa's other large mammals. For example, three small remaining remnant populations of Cape mountain zebra (*Equus zebra zebra*) each contain a single, unique haplotype but larger Namibian populations of closely related Hartmann's mountain zebra (*E. z. hartmannae*) have as many as eleven different haplotypes (Moodley & Harley, 2006; Watson & Chadwick, 2007). Another case in point is the loss of genetic diversity at mitochondrial and Y-chromosome loci observed in small, managed populations of Cape buffalo in Kenya and Uganda, which was attributed to restricted gene flow into protected areas (Van Hooft, Groen & Prins, 2002).

Table 2 MtDNA D-loop sequence variability within subspecies

Table 3 Summary statistics for the mtDNA control region sequence variability in each subspecies and haplotype identifiers used in Fig. 2

Subspecies & population	n	Genetic variability					
		H	h	(SD)	π	(SD)	S
<i>Diceros bicornis minor</i>	76	A-G	0.267	0.067	0.002	0.00063	6
Pooled <i>D. b. minor</i> samples							
Hluhluwe-iMfolozi Game Park (KZN) Accession number JN593089	50	A	—	—	—	—	—
Itala Game Park (KZN)	8	A	—	—	—	—	—
Mkuze Game Park (KZN)	5	A	—	—	—	—	—
Ndumo Game Park (KZN)	1	A	—	—	—	—	—
Johannesburg Zoo	1	A	—	—	—	—	—
Chete National Park, Zimbabwe, Accession numbers AF187825-AF187827 & AF1878329-AF187831	6	B, C & D	0.733	0.155	0.004	0.00056	4
Zambezi Valley, Zimbabwe, Accession numbers AY742832-AY742833	2	E & G	1	0.5	0.008	0.00413	3
Captive born, San Diego Zoo, Accession number AF187832	1	E	—	—	—	—	—
Captive born, Western Plains Zoo, Accession number AF187828	1	B	—	—	—	—	—
Captive born, Milwaukee Zoo, Accession number AF187833	1	F	—	—	—	—	—
<i>D. b. michaeli</i>	21	H-T	0.958	0.026 0.011	—	0.00106	15
Pooled <i>D. b. michaeli</i> samples							
Kenya, Accession numbers FJ227483-FJ227498	16	1-S	0.952	0.031	0.011	0.00119	15
Solio Game Reserve, Kenya, Accession numbers, AY742830-AY742831	2	J-T	1	0.5	0.011	0.00413	3
Cincinnati Zoo, Accession number AF187834	1	H	—	—	—	—	—
Taronga Zoo, Accession number AF187835	1	I	—	—	—	—	—
Addo Elephant Park, South Africa, Accession number JN593090	1	T	—	—	—	—	—
<i>D. b. bicornis</i>	4	U	0	—	—	—	—
Pooled <i>D. b. bicornis</i> samples							
Etosha, Namibia, Accession numbers JN593091-JN593094	4	U	—	—	—	—	—

Sample size (n), Haplotypes, labelled A–U (H), haplotype diversity (h), Nucleotide diversity (π), Standard deviation (SD), Number of segregating sites (S).

Low genetic variation is not always a consequence of recent anthropogenic fragmentation. An alternative hypothesis is that low levels of mtDNA and autosomal variation are a result of long-term demographic separation, historically small population sizes and local adaptation. For example, despite having lower mtDNA and autosomal DNA variation, there was no evidence of a genetic bottleneck in the Yellowstone National Park, U.S. A. grizzly bear (*Ursus arctos*) population compared to surrounding grizzly bear populations (Miller & Waits, 2003). Although Yellowstone's large population is embedded within the species' range, Miller & Waits (2003) attribute the lower genetic variation to restricted gene flow into the area from the north. The common impala (*Aepyceros melampus melampus*) of KZN also exhibited population

differentiation from populations in the Limpopo Province just 490 km north. Schwab *et al.* (2012) attributed the genetic divergence to a narrow zone of unsuitable habitat below the eastern escarpment of the Drakensberg Mountains that impeded dispersal between the two provinces.

Genetic replenishment by restocking and outbreeding is recommended in cases where anthropogenically induced fragmentation has caused a loss in genetic diversity and an increase in genetic divergence. For example, 'genetic rescue' has been recommended for the Cape zebra (Moodley & Harley, 2006; Watson & Chadwick, 2007). However, where differences amongst genetically depauperate populations might be of natural origin, population management may need to take into account local adaptation and the possibility of outbreeding depression.

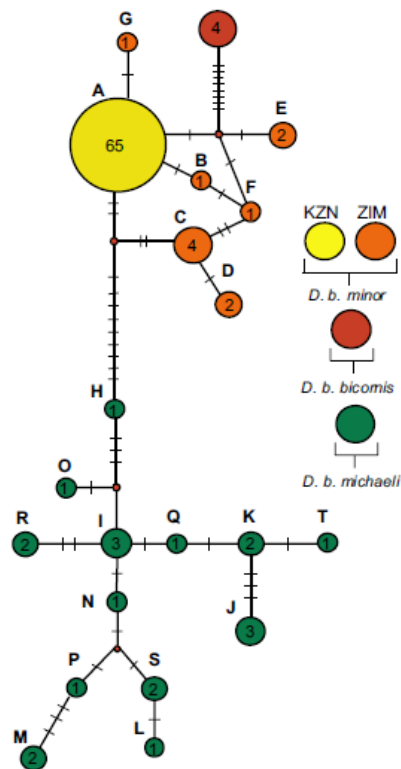


Fig 2 Statistical parsimony haplotype network calculated with Network Software for *Diceros bicornis*. KZN refers to the pooled *D. b. minor* samples within KwaZulu-Natal (KZN) (Ndumo Game Reserve, Ithala Game Reserve, Mkuze Game Reserve, Hluhluwe-Imfolozi Game Park (HIP) and Weenan Game Reserve as well as the sample from the Johannesburg Zoo). ZIM refers to the *D. b. minor* samples from Zimbabwe. Circles with numbers refer to the number of sample with corresponding haplotype from Table 3. The small dots denote the mutational step, and the cross bars are the number of base pair differences.

Resolving the question regarding KZN *D. b. minor* mtDNA and autosomal DNA genetic structure being a recent or old event is important for guiding management plans (Rookmaaker, 2005).

Microsatellite DNA markers were previously used to assess the levels of genetic variation amongst *D. b. minor* populations. Harley *et al.* (2005) found appreciable amounts of variation within the *D. b. minor* subspecies using nine microsatellite loci (Table 4). They recommended that as long as heterozygosity and allele num-

bers stayed at 'current' levels, no management policy change was necessary.

Based on a survey of ten microsatellites (Table 4), Karsten *et al.* (2011) found low levels of genetic variation within the KZN *D. b. minor*, but concluded that it was not cause for concern. They reached their conclusion based on (i) the similarity of allelic diversity and heterozygosity between the KZN *D. b. minor* population and the other subspecies; and (ii) a higher level of diversity within the black rhinoceros metapopulation compared to those found in other large African mammals. In their study, H_e estimates for the *D. b. bicornis* and *D. b. michaeli* subspecies (each based on only four samples) were substantially lower than those reported by Harley *et al.* (2005) (Table 4). Thus, estimates for *D. b. bicornis* and *D. b. michaeli* in the study by Karsten *et al.* (2011) are probably underestimates. Moreover, comparisons with other large African mammals should be made cautiously. Lions in the Serengeti Plains and Ngorongoro Crater have an H_e of 0.54 and 0.46, on par with black rhinos in the study by Harley *et al.* (2005), yet unlike the Serengeti Plains lions the Ngorongoro Crater lions have a marked decrease in their reproductive rate attributed to inbreeding depression caused by an anthropogenic bottleneck (Brown & Houlden, 1999; Driscoll *et al.* 2002).

HIP *D. b. minor* may be exhibiting signs of inbreeding depression owing to low reproductive rates that are not meeting the 0.25 fecundity rate desired by black rhino managers (Clinning *et al.*, 2009). If historic gene flow is verified and inbreeding is detected, there may be a need for genetic supplementation. Supplementation would also be supported based on genetic distances between the subspecies, low levels of genetic diversity and differentiation within and amongst the KZN metapopulation reported by this mtDNA study and published autosomal microsatellite data.

The likelihood of outbreeding depression in supplemented populations of the same species is low if they have the same karyotype, have been isolated for less than 500 years and occupy similar environments (Frankham *et al.*, 2011). Houck *et al.* (1995) identified the variation in chromosome morphology (number of submetacentric elements) between *D. b. minor* and *D. b. michaeli* zoo samples and recommended further studies to investigate possible differences in geographically separated populations of each subspecies in the wild. Furthermore, twenty-seven KZN *D. b. minor* were translocated to Malilangwe, Zimbabwe in 1997 where

Table 4 Microsatellite results from Harley *et al.*, 2005 and Karsten *et al.*, 2011. Expected heterozygosity (H_e), Observed heterozygosity (H_o)

Study	Microsatellites used	Subspecies	Geographic region	Sample size	H_e	H_o
Harley <i>et al.</i> , 2005	BR4, BR6, BR17	<i>Diceros bicornis minor</i>	South Africa and	46	0.46	0.436
	(Cunningham <i>et al.</i> 1999)	<i>D. b. bicornis</i>	Zimbabwe	53	0.51	0.523
	DB1, DB14, DB44, DB49, DB52, DB66 (Brown & Houlden, 1999)	<i>D. b. michaeli</i>	Namibia	19	0.68	0.731
Karsten <i>et al.</i> , 2011;	BR4, BR6, BR17	<i>D. b. minor</i>	South Africa	77	0.44	0.38
	(Cunningham <i>et al.</i> 1999)	<i>D. b. bicornis</i>	Namibia	4	0.43	0.46
	DB1, DB14, DB49, DB66 (Brown & Houlden, 1999)	<i>D. b. michaeli</i>	South Africa and	4	0.54	0.54
	AY606078, AY606080, AY606083 (Nielsen <i>et al.</i> , 2008)		Tanzania			

they were managed separately and not outbred with any Zimbabwe populations. The translocated population thrived with a growth rate of 8.3% per annum (R. du Toit, personal communication) cf. 3.4% over a 10-year period (1999–2008) in HiP (Clinning *et al.*, 2009). The success of the translocated KZN *D. b. minor* in Zimbabwe alleviated concerns about the adaptability of KZN rhino to Zimbabwe. The only remaining concern is whether the populations have been genetically isolated for longer than previously considered.

We recommend five research tasks to assist in resolving the genetic structure of southern Africa's black rhino as a guide to future management: (i) determine historic levels of genetic variation using museum or collection samples; (ii) investigate whether there is evidence of inbreeding depression within the HiP and KZN metapopulation; (iii) conduct a karyotype analysis on *D. b. minor* in KZN and Zimbabwe to determine whether chromosomal differences exist; (iv) increase the mtDNA sample size of the Zimbabwe *D. b. minor* population. Considering the high level of variation in the small sample size of the Zimbabwe sequences, a larger sample size of *D. b. minor* from that region might show that the KZN haplotype (A) is also there; and (v) lastly, genetic supplementation experiments should be implemented cautiously and systematically. A mixed population should be founded with at least 20 animals as suggested by du Toit (2006), perhaps using the Malilangwe, Zimbabwe translocation event as a template or more recent guidelines (Linklater *et al.*, 2011, 2012). The translocated KZN *D. b. minor* rhinos in Malilangwe have not yet been outbred with the Zimbabwe rhinos (R. du Toit, personal communication) but might be with the F1 and F2 offspring carefully monitored for

signs of reduction in reproductive fitness (outbreeding depression). If the research tasks we have recommended are completed and there is evidence of historic gene flow between KZN and Zimbabwe *D. b. minor* and no signs of outbreeding depression in the experimentally mixed population, then KZN *D. b. minor* is a candidate for genetic supplementation using progeny from Zimbabwe populations.

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Appendix B

In order for harvests to be effective and to ensure that over-harvest does not occur, an accurate census is necessary. HiP black rhino managers recently re-evaluated the number of *D. b. minor* in the reserve. They discovered that black rhino population estimates reported for 1998 to 2008 were over estimated (Clinning et al. 2009). Inaccurate counts during those years led to harvesting of up to 10% of the total population, which may be responsible for the low growth rate of the population (Clinning et al. 2009). Excessive harvesting can affect endangered populations that tend to already be small and more than likely have a smaller effective population size (N_e) (Allendorf et al. 2008). Harvesting at levels above 10% will decrease population sizes and counteract efforts to stimulate growth (Owen-Smith 1987). HiP management has done its best to adhere to the 5% - 8% harvesting rate suggested by the SADC RMG. However, PVA modelling results suggest that managers should consider altering harvesting rates to 3% per gestational year (4.0% for actual year) so the population can reach its growth rate potential (e.g. higher stable population numbers and increased genetic heterozygosity; Table A.1). While the SADC RMG establishes recommended harvesting criteria for black rhino management, it must be appreciated that those guidelines are not necessarily suitable for every population. If lower harvesting rates for HiP was implemented (<5%), management might see an increase a slight increase in reproductive rates.

Consideration should be taken when interpreting SADC RMG guidelines and calculating harvest regimes that not all populations are the same (e.g. different levels of heterozygosity, reproductive skew, etc.). If managers have

ample and accurate life-history data on the population in question, a computer simulation should be carried out.

Table A.1: Expected heterozygosity (H_E) with standard error for a modelled *D. b. minor* population; with and without inbreeding at 430 carrying capacity for years 50, 100, 150, 200, 250, 500, 700 & 1000; with mean population sizes and standard error. The 2% harvest is for a gestational year, which translates to 2.6% in a calendar year; 3% (4.0% in a calendar year); 4% (5.4% in a calendar year) and 5% (6.7% in a calendar year); 6% (8% in a calendar year); 7% (9.3% in a calendar year); 8% (10.7% in a calendar year); 9% (12% in a calendar year); 10% (13.4% in a calendar year).

430 carrying capacity, 2% Harvest					430 carrying capacity, 3% Harvest					430 carrying capacity, 4% Harvest				
Year	Mean Pop. Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	~ Pop Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	Mean Pop. Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding
50	334 ± 1.11	0.460 ± 0.000	331 ± 1.11	0.460 ± 0.00	50	309 ± 1.28	0.460 ± 0.000	308 ± 1.25	0.459 ± 0.000	50	267 ± 1.25	0.459 ± 0.000	263 ± 1.21	0.459 ± 0.000
100	336 ± 1.10	0.455 ± 0.000	329 ± 1.14	0.456 ± 0.00	100	308 ± 1.27	0.454 ± 0.000	297 ± 1.36	0.453 ± 0.000	100	260 ± 1.19	0.451 ± 0.001	247 ± 1.09	0.453 ± 0.001
150	338 ± 1.11	0.450 ± 0.001	322 ± 1.18	0.450 ± 0.001	150	311 ± 1.30	0.449 ± 0.001	286 ± 1.38	0.448 ± 0.001	150	257 ± 1.16	0.446 ± 0.001	238 ± 1.01	0.447 ± 0.001
200	339 ± 1.06	0.446 ± 0.001	317 ± 1.21	0.446 ± 0.001	200	309 ± 1.29	0.444 ± 0.001	276 ± 1.39	0.442 ± 0.001	200	258 ± 1.21	0.438 ± 0.001	230 ± 0.91	0.440 ± 0.001
250	339 ± 1.04	0.441 ± 0.001	315 ± 1.25	0.440 ± 0.001	250	308 ± 1.30	0.438 ± 0.001	267 ± 1.39	0.435 ± 0.001	250	256 ± 1.20	0.432 ± 0.001	226 ± 0.90	0.432 ± 0.001
500	337 ± 1.07	0.418 ± 0.001	289 ± 1.30	0.415 ± 0.001	500	310 ± 1.27	0.413 ± 0.001	238 ± 1.21	0.406 ± 0.001	500	259 ± 1.23	0.402 ± 0.001	211 ± 0.77	0.397 ± 0.001
700	339 ± 1.05	0.401 ± 0.001	273 ± 1.32	0.396 ± 0.001	700	308 ± 1.29	0.394 ± 0.001	212 ± 1.07	0.381 ± 0.001	700	256 ± 1.18	0.379 ± 0.001	199 ± 0.82	0.370 ± 0.001
1000	337 ± 1.10	0.377 ± 0.001	236 ± 1.45	0.363 ± 0.001	###	308 ± 1.30	0.366 ± 0.001	180 ± 1.34	0.341 ± 0.002	1000	257 ± 1.18	0.347 ± 0.002	165 ± 1.39	0.339 ± 0.002

430 carrying capacity, 5% Harvest					430 carrying capacity, 6% Harvest					430 carrying capacity, 7% Harvest				
Year	~ Pop Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	~ Pop Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	Mean Pop. Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding
50	236 ± 0.83	0.458 ± 0.000	236 ± 0.87	0.458 ± 0.000	50	220 ± 0.60	0.457 ± 0.000	218 ± 0.57	0.457 ± 0.000	50	213 ± 0.49	0.458 ± 0.000	210 ± 0.47	0.458 ± 0.000
100	232 ± 0.79	0.452 ± 0.001	223 ± 0.71	0.451 ± 0.001	100	217 ± 0.53	0.450 ± 0.001	211 ± 0.5	0.450 ± 0.001	100	211 ± 0.49	0.450 ± 0.001	206 ± 0.48	0.449 ± 0.001
150	232 ± 0.76	0.444 ± 0.001	218 ± 0.64	0.444 ± 0.001	150	217 ± 0.56	0.443 ± 0.001	208 ± 0.54	0.442 ± 0.001	150	212 ± 0.48	0.443 ± 0.001	203 ± 0.50	0.442 ± 0.001
200	231 ± 0.77	0.437 ± 0.001	214 ± 0.63	0.437 ± 0.001	200	218 ± 0.54	0.436 ± 0.001	206 ± 0.56	0.434 ± 0.001	200	210 ± 0.50	0.435 ± 0.001	200 ± 0.55	0.433 ± 0.001
250	231 ± 0.75	0.428 ± 0.001	211 ± 0.67	0.429 ± 0.001	250	217 ± 0.55	0.428 ± 0.001	204 ± 0.55	0.427 ± 0.001	250	210 ± 0.46	0.428 ± 0.001	197 ± 0.59	0.425 ± 0.001
500	231 ± 0.78	0.394 ± 0.001	200 ± 0.72	0.393 ± 0.001	500	217 ± 0.54	0.394 ± 0.001	193 ± 0.69	0.390 ± 0.001	500	211 ± 0.49	0.390 ± 0.001	187 ± 0.74	0.388 ± 0.001
700	230 ± 0.75	0.370 ± 0.001	185 ± 0.95	0.363 ± 0.001	700	219 ± 0.56	0.369 ± 0.001	179 ± 0.94	0.362 ± 0.002	700	211 ± 0.48	0.364 ± 0.001	176 ± 0.91	0.360 ± 0.001
1000	230 ± 0.75	0.337 ± 0.002	151 ± 1.42	0.318 ± 0.001	###	217 ± 0.55	0.334 ± 0.002	140 ± 1.54	0.317 ± 0.002	1000	211 ± 0.48	0.327 ± 0.002	134 ± 1.57	0.314 ± 0.002

430 carrying capacity, 8% Harvest					430 carrying capacity, 9% Harvest					430 carrying capacity, 10% Harvest				
Year	Mean Pop. Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	~ Pop Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding	Year	Mean Pop. Size	H_E : No Inbreeding	Mean Pop. Size	H_E : Inbreeding
50	207 ± 0.42	0.457 ± 0.000	205 ± 0.42	0.457 ± 0.000	50	203 ± 0.42	0.456 ± 0.000	200 ± 0.42	0.456 ± 0.000	50	199 ± 0.39	0.457 ± 0.000	198 ± 0.42	0.456 ± 0.000
100	206 ± 0.46	0.449 ± 0.001	201 ± 0.46	0.449 ± 0.001	100	203 ± 0.41	0.447 ± 0.001	198 ± 0.45	0.448 ± 0.001	100	199 ± 0.42	0.449 ± 0.001	194 ± 0.44	0.448 ± 0.001
150	207 ± 0.44	0.441 ± 0.001	198 ± 0.49	0.440 ± 0.001	150	203 ± 0.40	0.439 ± 0.001	195 ± 0.47	0.440 ± 0.001	150	199 ± 0.41	0.440 ± 0.001	192 ± 0.49	0.440 ± 0.001
200	206 ± 0.44	0.432 ± 0.001	196 ± 0.54	0.432 ± 0.001	200	201 ± 0.42	0.431 ± 0.001	192 ± 0.50	0.432 ± 0.001	200	199 ± 0.40	0.433 ± 0.001	190 ± 0.51	0.432 ± 0.001
250	206 ± 0.43	0.425 ± 0.001	194 ± 0.57	0.425 ± 0.001	250	202 ± 0.41	0.422 ± 0.001	191 ± 0.59	0.424 ± 0.001	250	199 ± 0.39	0.424 ± 0.001	190 ± 0.51	0.423 ± 0.001
500	206 ± 0.43	0.389 ± 0.001	185 ± 0.65	0.387 ± 0.001	500	202 ± 0.44	0.385 ± 0.001	181 ± 0.74	0.385 ± 0.001	500	199 ± 0.39	0.387 ± 0.001	180 ± 0.71	0.383 ± 0.001
700	205 ± 0.44	0.362 ± 0.001	172 ± 0.91	0.358 ± 0.001	700	202 ± 0.42	0.357 ± 0.002	168 ± 0.98	0.355 ± 0.001	700	199 ± 0.41	0.360 ± 0.001	165 ± 0.99	0.353 ± 0.001
1000	206 ± 0.42	0.326 ± 0.002	130 ± 1.60	0.308 ± 0.002	###	202 ± 0.43	0.320 ± 0.002	125 ± 1.57	0.307 ± 0.002	1000	199 ± 0.41	0.322 ± 0.002	118 ± 1.63	0.302 ± 0.002

Appendix C

It has already been established that cohorts consisting of individuals from HiP that are destined for reintroduction or established population supplementation within KZN have been successful as South African numbers of *D. b. minor* have steadily increased since translocations and reintroductions began in 1962. Whether or not the success can be attributed to local adaptations however could be debated as the cohort of 27 KZN *D. b. minor* translocated to Mililangwe, Zimbabwe in 1997 had a higher reproductive rate than local KZN *D. b. minor* and no reported increases in mortality.

When running PVA computer simulations for Chapter Four, I also ran models to test the SADC RMG recommendation that cohorts used for reintroductions consist of at least 20 individuals. The results were interesting in that the cohorts with as few as 16 individuals was large enough to maintain a population from a 'numbers' standpoint (Table A2). Populations founded with 16 individuals from HiP had an 83% probability of success. The success rate increased to 88% when 20 individuals were modelled.

Table A2: Probability of success or extinction for founder populations with sizes from 5- 30 individuals. The HiP results take into account the mean expected heterozygosity for each population using microsatellite DNA loc from Chapter Three with a carrying capacity of the founding reserve 50.

HiP Founders					
Number of Individuals in Starting Population	Pops. that Went Extinct	Pops. That Survived	Probability of Extinction	Probability of Success	Mean to First Extinction (in years)
5	383	117	0.766 ± 0.0189	0.2340 ± 0.0189	37 ± 3.21
6	331	169	0.6620 ± 0.0212	0.3380 ± 0.0212	54 ± 4.70
7	297	203	0.5940 ± 0.0220	0.4060 ± 0.0220	63 ± 5.22
8	258	242	0.516 ± 0.0223	0.4840 ± 0.0223	57 ± 4.63
9	210	290	0.4200 ± 0.0221	0.5800 ± 0.0221	88 ± 7.44
10	213	287	0.4260 ± 0.0221	0.5740 ± 0.0221	79 ± 6.31
11	169	331	0.3380 ± 0.0212	0.6620 ± 0.0212	90 ± 7.65
12	148	352	0.2960 ± 0.0204	0.7040 ± 0.0204	130 ± 10.49
13	153	347	0.3060 ± 0.0206	0.6940 ± 0.0206	119 ± 9.73
14	129	371	0.2580 ± 0.0196	0.7420 ± 0.0196	131 ± 11.75
15	108	392	0.2160 ± 0.0184	0.7840 ± 0.0184	174 ± 14.07
16	86	414	0.1720 ± 0.0169	0.8280 ± 0.0169	171 ± 16.13
17	85	415	0.1700 ± 0.0168	0.8300 ± 0.0168	174 ± 16.30
18	88	412	0.1760 ± 0.0170	0.8240 ± 0.0170	194 ± 15.63
19	78	422	0.1560 ± 0.0162	0.8440 ± 0.0162	218 ± 17.88
20	62	438	0.1240 ± 0.0147	0.8760 ± 0.0147	173 ± 15.89
21	79	421	0.1580 ± 0.0163	0.8420 ± 0.0163	209 ± 16.31
22	61	439	0.1220 ± 0.0146	0.8780 ± 0.0146	212 ± 16.19
23	62	438	0.1240 ± 0.0147	0.8760 ± 0.0147	235 ± 19.39
24	64	436	0.1280 ± 0.0149	0.8720 ± 0.0149	246 ± 17.20
25	62	438	0.1240 ± 0.0147	0.8760 ± 0.0147	235 ± 16.80
26	55	445	0.1100 ± 0.0140	0.8900 ± 0.0140	278 ± 19.40
27	57	443	0.1140 ± 0.0142	0.8860 ± 0.0142	252 ± 17.61
28	45	455	0.0900 ± 0.0128	0.9100 ± 0.0128	245 ± 19.17
29	63	437	0.1260 ± 0.0148	0.8740 ± 0.0148	253 ± 17.32
30	53	447	0.1060 ± 0.0138	0.8940 ± 0.0138	294 ± 18.86

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