

Caitlin J Curry, James N Derr

cjcurry@cvm.tamu.edu; jderr@cvm.tamu.edu Interdisciplinary Program of Genetics, Department of Veterinary Pathobiology College of Veterinary Medicine and Biomedical Sciences Texas A&M University, College Station, TX, USA



A Century of Conservation Genetics

Abjection Lion (Panthera leo)



UNT HEALTH SCIENCE CENTER

> Universiteit Leiden



Genetic diversity is examined using fourteen microsatellites with primers designed to be closer to the target region and specific to the lion (miniSTRs). Preliminary statistical analyses include determining the number of alleles per locus and calculation of expected heterozygosity (*GenAlEx*), structural analysis (*Structure*) and determining the most likely number of clusters following the ΔK method (*Structure Harvester*).

We determined the genetic architecture of both historical (>100 years ago; n=155) and contemporary (2000 to present; n=557) African lion populations across the traditional range states in Africa (*Figure 1*). Both datasets were analyzed using the same methods allowing for a more direct comparison over time than has previously been employed. The historical lion dataset is DNA isolated from high quality and welldocumented while museum specimens the contemporary lion dataset is from modern material and data from several recently published studies. Combining the datasets of historical as well as contemporary populations provides quantitative measures on the extent of change in the genetic diversity of lions over the past 100 years and provides a basis for assessing the genetic health of the African lion.





Figure 1

All samples are categorized into 6 conventionally recognized regions: • North (Historical Only) • West • Central • East • Southeast • Southern

Additional sample categories:
India (Asiatic Lion – Panthera leo persica)
Unknown (Historical Only – Date Provide, No Location)

Leo391 Leo506 Overall Leo077 Leo085 Leo098 Leo126 Leo247 Leo224 Leo281 Leo006 Leo008 Leo031 Leo045 Leo230 %AS 97.5% 97.8% 95.6% 98.1% 100.0% 98.5% 92.5% 100.0% 100.0% 99.8% 99.0% 86.6% 99.5% 98.3% 99.3% 508 408 470 464 455 405 410 441 499 469 501 504 415 393 o 476 Na 20 8 9 12 19 12 11 9 16 10 13 10 9 10 약 0.75 0.78 0.71 0.84 0.40 0.39 0.75 0.65 0.72 0.77 0.82 0.82 0.72 0.74 0.81 He %AS 83.3% 97.4% 76.8% 88.4% 85.2% 98.1% 76.8% 60.6% 38.1% _T 87.1% 95.5% 91.6% 99.4% 77.4% 94.2% 129 152 142 132 154 119 59 St 135 151 119 137 148 120 146 94 16 r 16 Na 24 15 12 21 10 16 11 24 13 11 18 22 10 0.81 0.59 0.83 0.86 0.83 0.84 0.82 0.82 0.90 0.81 0.73 0.80 0.84 0.85 0.84

%AS = Amplification Success; Na = No. of Different Alleles; He = Expected Heterozygosity

The historical and modern populations both have

Table 1 Genetic Diversity Overall and By Locus







high levels of overall genetic diversity (*Table 1*). They also exhibit similar trends in population structure (*Figure 2*). Heterozygosity by region (*Figure 3*) supports observations from recent studies of a precipitous decline in population size occurring in West and Central Africa.

Further analysis will include the addition of mtDNA haplotypes. A phylogeographic analysis will be performed along with estimation of effective population size (Ne) and calculation of genetic differentiation between regions (Fst).

Intro