1	Putting RFMix and ADMIXTURE to the test in a complex admixed population				
2					
3	Caitlin Uren <sup>1\$</sup> , Eileen G. Hoal <sup>1</sup> , Marlo Möller <sup>1</sup>				
4					
5	<sup>1</sup> DST-NRF Centre of Excellence for Biomedical Tuberculosis Research, South				
6	African Medical Research Council Centre for Tuberculosis Research, Division of				
7	Molecular Biology and Human Genetics, Faculty of Medicine and Health Science				
8	Stellenbosch University, Cape Town, South Africa				
9					
10	\$ Correspondence should be addressed to:				
11	Dr. Caitlin Uren, Room 4036, 4 <sup>th</sup> Floor Education Building, Francie van Zijl Drive				
12	DST-NRF Centre of Excellence for Biomedical Tuberculosis Research, South Africar				
13	Medical Research Council Centre for Tuberculosis Research, Division of Molecula				
14	Biology and Human Genetics, Faculty of Medicine and Health Sciences,				
15	Stellenbosch University, Cape Town, 8000, South Africa				
16	Phone: 021 938 9692				
17	E-mail: caitlinu@sun.ac.za				
18					
19	Running title: Accuracy of global & local ancestry				
20					
21	ORCID ID's:				
22	CU - 0000-0003-2358-0135				
23	EGH - 0000-0002-6444-5688				
24	MM - 0000-0002-0805-6741				
25					

## 26 Abstract

27

Global and local ancestry inference in admixed human populations can be performed 28 29 using computational tools implementing distinct algorithms, such as RFMix and ADMIXTURE. The accuracy of these tools has been tested largely on populations 30 with relatively straightforward admixture histories but little is known about how well 31 32 they perform in more complex admixture scenarios. Using simulations, we show that RFMix outperforms ADMIXTURE in determining global ancestry proportions in a 33 34 complex 5-way admixed population. In addition, RFMix correctly assigns local ancestry with an accuracy of 89%. The increase in reported local ancestry inference 35 accuracy in this population (as compared to previous studies) can largely be 36 37 attributed to the recent availability of large-scale genotyping data for more 38 representative reference populations. The ability of RFMix to determine global and local ancestry to a high degree of accuracy, allows for more reliable population 39 40 structure analysis, scans for natural selection, admixture mapping and case-control association studies. This study highlights the utility of the extension of computational 41 tools to become more relevant to genetically structured populations, as seen with 42 RFMix. This is particularly noteworthy as modern-day societies are becoming 43 increasingly genetically complex and some genetic tools are therefore less 44 45 appropriate. We therefore suggest that RFMix be used for both global and local ancestry estimation in complex admixture scenarios. 46

47

48

Keywords: South Africa; local ancestry inference; population genetics; RFMix;
ADMIXTURE

## 51 Introduction

52

Admixture, the exchange of genetic material between distinct populations, is a 53 54 hallmark of modern society - it can occur between closely or distantly related populations (both genetically and geographically) (1000 Genomes Project 55 Consortium et al. 2012). This exchange of genetic material leads to population 56 structure; the pattern, timing and extent has been investigated in detail in a number 57 of populations (1000 Genomes Project Consortium et al. 2012; Gurdasani et al. 58 59 2015; Uren et al. 2016). Such studies on southern African populations are particularly noteworthy as this area is postulated to be the geographical origin of 60 modern humans and therefore investigating population structure in modern southern 61 62 African populations may reveal more about the area's rich history (Henn et al. 2011). 63

Correctly and efficiently determining ancestral proportions in an admixed population 64 65 is possible by using computational and statistical algorithms that adapt to a variety of demographic scenarios (Alexander et al. 2009; Maples et al. 2013; Brown and 66 Pasaniuc 2014). Furthermore, the ability to determine the ancestral origin of a 67 particular chromosomal region in an admixed individual has enabled the mapping of 68 69 the origins of genetic risk factors in complex disease i.e. admixture mapping 70 (Freedman et al. 2006; Cheng et al. 2009; Daya et al. 2014). The majority of the computational and statistical tools used for global and local ancestry were however 71 tested on and tailored to 2- to 3-way admixed populations. The extension to a 72 73 complex 5-way admixed population and the evaluation of the resulting accuracy as we present here, has rarely been done. (Daya et al. 2014; Uren et al. 2016). 74

75

76 A South African population with unique genetic ancestry and 5-way admixuture (the 77 South African Coloured (SAC) population as termed in the South African census) received ancestral contributions from Bantu-speaking African (~30%), KhoeSan 78 79 (~30%), European (~20%), East Asian (~10%) and South East Asian populations (~10%) (de Wit et al. 2010; Chimusa et al. 2013; Uren et al. 2016). The admixture 80 began approximately 15 generations ago and followed a continuous migration model 81 82 (Uren et al. 2016). This number, mode and timing of admixture events is unique and creates a highly complex population. Although we are able to describe the 83 84 demographic model for most populations, there are some gaps in knowledge. This may include not knowing which populations or specific geographical locations are the 85 best proxies for the true ancestral populations. Therefore, any studies investigating 86 87 an association between genetics and disease risk needs to be able to correctly 88 account for population or even individual admixture proportions within the limits of the availability of current genetic data. 89

90

The first step in a study design aimed at finding a link between ancestry and disease 91 (such as genome-wide association studies and admixture mapping) is to understand 92 the ancestral composition of the study population. Ancestral origins and contributions 93 94 to the 5-way admixed South African population have been estimated but there have 95 been very few studies that have investigated the accuracy of the results generated by the computational algorithm used (de Wit et al. 2010; Chimusa et al. 2013; 96 Petersen et al. 2013; Daya et al. 2013; Uren et al. 2016). Here we have set out to 97 98 test the accuracy of global and local ancestry inference in one of the most complex admixed populations world-wide, using newly available dense genotyping data. A 99 100 simulated 5-way admixed population is generated and global and local ancestry

101 estimates are compared to the true values to determine the accuracy of the 102 computational algorithm.

103

104 Methods

105

# 106 Data merging and filtering

107

108 KhoeSan genotype data from Martin and colleagues (Martin *et al.* 2017) was merged 109 with the PAGEII dataset (Wojcik *et al.* 2018). In order to increase the number of 110 European and South East Asian reference samples in the dataset, the data was 111 merged with Gujarati Indian and European genetic data from the 1000 Genomes 112 Project (1000 Genomes Project Consortium (2010) 2010).

113

Preliminary data filtering included a filter for minor allele frequency (0.003), missingness per genotype (max 0.05) and missingness per individual (max 0.01). A total of ~776k SNPs passed these filters and formed the initial merged dataset. Further data filtering is described in the appropriate sections below. Data was phased using SHAPEIT2 utilizing a recombination map averaged across European and African populations (The International HapMap Consortium 2007; O'Connell *et al.* 2014). A summary of the populations in the final dataset can be seen in Table 1.

121

### 122 Simulations

123

124 The computational workflow is summarized in Figure 1. A subset of reference 125 individuals from the final merged dataset described in Table 1 was used to generate

126 a simulated dataset using admix-simu (Williams 2016). A demographic model 127 consisting of the ancestry proportions described above and a continuous migration model starting at 15 generations ago (Uren et al. (2016)), was used to generate a 128 129 simulated 5-way admixed population (Uren et al. 2016). This simulation results in a 130 heterogenous population, reminiscent of a real-world SAC population. The simulation does not take post-admixture selection into account since it is highly unlikely that 350 131 132 years would result in distinct selection signals, rather, the inherent selection signals in the source populations will be transferred in a random manner to the simulated 133 134 admixed population. Genotype as well as local ancestry calls were generated for this simulated dataset from real reference haplotypes, thus capturing the complexity of 135 this heterogenous 5-way admixed South African population. 136

137

138

# 139 Software choices

140

Although there are a number of software programs that are able to estimate global ancestry, ADMIXTURE is the most utilized. Reasons for this include the ability to include related individuals in one run and to generate accurate admixture proportions using relatively low-density SNP-array data (Alexander *et al.* 2009). The other widely used global ancestry algorithm, STRUCTURE has been shown to overestimate admixture proportions in complex populations (Cheng *et al.* 2017).

147

148 RFMix was chosen as the local ancestry inference algorithm of choice as it allows for 149 parameter optimization given the number of ancestral populations, has the inherent 150 ability to calculate local and global ancestry simultaneously, allows for array-based

input data as well as whole genome sequencing data, and has a proven track record
with admixed populations (Maples *et al.* 2013; Padhukasahasram 2014).
Furthermore, during a preliminary study by Daya and colleagues, RFMix was shown
to be the most accurate tool for local ancestry estimation in this 5-way admixed
South African population (Daya *et al.* 2014).

156

# 157 GAI accuracy

158

Reference individuals not included in the dataset used for the simulation, were allocated to the dataset used for global and local ancestry inference. Global ancestry proportions were determined by ADMIXTURE (Alexander *et al.* 2009) and RFMix (Maples *et al.* 2013).

163

The ADMIXTURE analysis was performed in an unsupervised manner after filtering the dataset for linkage disequilibrium as per recommendations in the ADMIXTURE manual (50kb window size, step size of 10kb and R<sup>2</sup> threshold of 0.1). Relatedness in the reference dataset was assessed using king (Manichaikul *et al.* 2010) and all second degree relatives were removed prior to admixture analysis.

169

RFMix was run using default parameters, a time since admixture of 15 generations
(in line with the simulation) as well as 3 expectation-maximization (EM) iterations
(further EM iterations were not shown to increase accuracy (Maples *et al.* 2013)).
The correlation of the two methods by means of the Root Mean Squared Error was
performed in R.

175

# 176 LAI accuracy

177

Local ancestry calls were generated by RFMix using the same parameters asdescribed in the previous section.

180

The ability to correctly assign local ancestry was calculated in two ways. The first determined the global accuracy i.e. how often the computational tool assigned the correct ancestry (as per the simulations) and the second looked at this accuracy per ancestral population (Atkinson 2018). These accuracy estimators were then averaged over all individuals in the simulated 5-way admixed dataset.

186

187

### 188 Data Availability

189

190 No new genetic data was generated for this study however all reference data 191 supporting the findings of this study are available via the original publication.

192

193

### 194 Results and discussion

195

The aim of this study was to determine the accuracy of global and local ancestry inference. In order to do this, a highly complex 5-way admixed population was simulated. The local and global ancestry estimates were then compared to the true simulated data.

200

#### 201

## 202 Global Ancestry Inference Accuracy

The genetic diversity inherent in an admixed South African population was simulated using 5 reference populations (see Methods). The average ancestry proportions across these individuals were in line with what is seen in the real-world (Table 2) (Uren *et al.* 2016). The simulations provided the basis with which the global ancestry proportions as calculated by ADMIXTURE (Alexander *et al.* 2009) and RFMix (Maples *et al.* 2013) could be compared.

209

Unsupervised admixture analysis of the simulated dataset by ADMIXTURE and 210 211 RFMix confirmed that the simulated 5-way admixed population is highly 212 heterogenous. Average ancestral proportions for both computational tools are given in Table 2. The comparisons across the 5 ancestries for each simulated individual 213 214 are also depicted in Figure 2. Root Mean Squared Errors (RMSE) (RFMix vs 215 Simulation and ADMIXTURE vs simulation) were calculated for each ancestry. As per the RMSE's, RFMix outperforms ADMIXTURE in correctly estimating admixture 216 proportions in the 5-way admixed population, with the exception of East Asian 217 ancestry where the accuracy is equal. ADMIXTURE over-estimates the Bantu-218 219 speaking African contribution and under-estimates the KhoeSan ancestral 220 proportions. ADMIXTURE also overestimates European ancestry and underestimates South East Asian ancestry. This is most likely due to inherent 221 European ancestry present in South East Asian populations and similarly, Bantu-222 speaking ancestry in the KhoeSan reference population. It is likely that if more 223 homogenous reference populations were chosen, this trend would be negated but, 224

as previously mentioned, most modern day populations are admixed and thereforecomputational tools should be able to account for this within the algorithms.

227

In addition, we hypothesize that the discrepancy in admixture proportions between RFMix and ADMIXTURE can also be attributed to the increase in prior information given to RFMix in order to determine admixture proportions i.e. phase and recombination rate.

232

233

# 234 Local Ancestry Inference Accuracy

235 Beyond global ancestry proportions, the simulation of a 5-way admixed population 236 resulted in known local ancestry calls, to which calls by a computational tool can be 237 compared. The ancestral origin of each parental chromosomal region was determined using RFMix. RFMix has been shown to outperform other computational 238 239 tools in the estimation of local ancestry in complex admixture scenarios (Daya et al. 2014). The local ancestry calls by RFMix were compared to the "true" simulated 240 241 ancestral origin of each region (Figure 3). The overall local ancestry inference accuracy across all individuals and ancestries is ~89%; 88% accurate in calling 242 243 Bantu-speaking African ancestry, 87% calling KhoeSan ancestry, 95% calling 244 European ancestry, 86% calling East Asian ancestry and 85% calling South East Asian ancestry. The statistical significance of RFMix's ability to call a specific 245 ancestry over another was assessed. RFMix is able to call European ancestry more 246 247 precisely than any of the African or Asian ancestries and it was able to call KhoeSan ancestry more accurately than East Asian ancestry (Figure 3). This is consistent with 248

what was previously found and confirms that these algorithms are not tailored toAfrican populations (Maples *et al.* 2013).

251

252 The local ancestry inference accuracy estimates presented here are substantially higher than previously obtained for this 5-way admixed South African population 253 (Daya et al. 2014). This increased accuracy can be attributed largely to the recent 254 availability of large-scale genotyping array data from the KhoeSan population which 255 256 is used as a reference for this admixed population. This, in addition to the overall 257 higher SNP density, increased the accuracy from ~70% (as previously reported (Daya et al. 2014)) to ~89%. As new datasets become available and the overlap 258 259 between datasets increases, we envisage this accuracy increasing even further.

260

261

# 262 Conclusion

263

In conclusion, the findings presented here detail the accuracy of global and local 264 ancestry inference of one of the most complex populations worldwide, which puts 265 ADMIXTURE and RFMix to the ultimate test. Due to the accuracy and versatility of 266 267 RFMix in determining global and local ancestry in a single program, it should be the 268 algorithm of choice to characterize more complex admixture scenarios. The inclusion of accurate admixture proportions as a covariate in association studies is vital, and it 269 270 is our opinion that researchers studying complex admixed populations should use 271 RFMix for this purpose.

272

Furthermore, we demonstrate that computational tools *are* able to decipher the complex African genetic history with a high degree of accuracy, but there is still some room for improvement regarding the tailoring of computational tools to handle diverse, admixed reference and target populations under study.

277

As populations become increasingly mobile, the probability of admixture is greater and the extension of these and future computational tools to more genetically complex populations across the world is vital and, as we have demonstrated, is possible. The conclusions of this study will be increasingly relevant and generalizable.

283

284

### 285 Acknowledgements

286

We thank Dr. Brenna Henn and Dr. Elizabeth Atkinson for their assistance and support of this research. We thank the study participants of the projects cited here without their contribution, this research would not be possible.

290

291

## 292 Author Contributions

293

294 CU designed the study, wrote the first draft of the manuscript and performed the 295 computational analyses. MM and EH helped to develop the research and edited the 296 manuscript.

297

298	
299	Funding
300	
301	This research was funded (partially or fully) by the South African government through
302	the South African Medical Research Council and the National Research Foundation.
303	CU was supported by a fellowship from the Claude Leon Foundation.
304	
305	Conflict of Interest
306	
307	The authors declare that they have no conflict of interest.
308	
309	
310	Ethical Approval and Informed Consent
311	
312	All procedures performed in studies involving human participants were in accordance
313	with the ethical standards of the institutional and/or national research committee and
314	with the 1964 Helsinki declration and its later amendments or comparable ethical
315	standards. Informed consent was obtained from all individual participants included in
316	
	the study. This study was approved by the Stellenbosch University Health Research
317	the study. This study was approved by the Stellenbosch University Health Research Ethics Committee (Reference #N11/07/210).
317 318	the study. This study was approved by the Stellenbosch University Health Research Ethics Committee (Reference #N11/07/210).
317 318 319	the study. This study was approved by the Stellenbosch University Health Research Ethics Committee (Reference #N11/07/210).
317 318 319 320	the study. This study was approved by the Stellenbosch University Health Research Ethics Committee (Reference #N11/07/210). References

- 1000 Genomes Project Consortium (2010), 2010 A map of human genome variation
   from population-scale sequencing. Nature 467: 1061–1073.
- 1000 Genomes Project Consortium, G. R. Abecasis, A. Auton, L. D. Brooks, M. A.
- 325 DePristo *et al.*, 2012 An integrated map of genetic variation from 1,092 human 326 genomes. Nature 491: 56–65.
- Alexander, D. H., J. Novembre, and K. Lange, 2009 Fast model-based estimation of
   ancestry in unrelated individuals. Genome Res. 19: 1655–1664.
- Atkinson, E., 2018 Calculations of accuracy comparing Williams lab simulations to
- 330 *RFmix runs: eatkinson/LAlaccuracy.*
- Brown, R., and B. Pasaniuc, 2014 Enhanced methods for local ancestry assignment

in sequenced admixed individuals. PLoS Comput. Biol. 10: e1003555.

333 Cheng, C. Y., W. H. Kao, N. Patterson, A. Tandon, C. A. Haiman et al., 2009

Admixture mapping of 15,280 African Americans identifies obesity

susceptibility loci on chromosomes 5 and X. PLoS.Genet 5: e1000490.

- 336 Cheng, J. Y., T. Mailund, and R. Nielsen, 2017 Fast admixture analysis and
- population tree estimation for SNP and NGS data. Bioinformatics 33: 2148–
  2155.

Chimusa, E. R., M. Daya, M. Möller, R. Ramesar, B. M. Henn et al., 2013

340 Determining ancestry proportions in complex admixture scenarios in South

- Africa using a novel proxy ancestry selection method. PLoS ONE 8: e73971.
- 342 Daya, M., L. van der Merwe, U. Galal, M. Möller, M. Salie et al., 2013 A Panel of
- 343 Ancestry Informative Markers for the Complex 5-way Admixed South African
- Coloured Population. PLoS ONE 8: e82224.

345	Daya, M., L. van der Merwe, C. R. Gignoux, P. D. van Helden, M. Möller et al., 2014
346	Using multi-way admixture mapping to elucidate TB susceptibility in the South
347	African Coloured population. BMC Genomics 15: 1021.
348	Freedman, M. L., C. A. Haiman, N. Patterson, G. J. McDonald, A. Tandon et al.,
349	2006 Admixture mapping identifies 8q24 as a prostate cancer risk locus in
350	African-American men. Proc.Natl.Acad.Sci.U.S.A 103: 14068–14073.
351	Gurdasani, D., T. Carstensen, F. Tekola-Ayele, L. Pagani, I. Tachmazidou et al.,
352	2015 The African Genome Variation Project shapes medical genetics in
353	Africa. Nature 517: 327–332.
354	Henn, B. M., C. R. Gignoux, M. Jobin, J. M. Granka, J. M. Macpherson <i>et al.</i> , 2011
355	Hunter-gatherer genomic diversity suggests a southern African origin for
356	modern humans. Proc. Natl. Acad. Sci. U.S.A. 108: 5154–5162.
357	Manichaikul, A., J. C. Mychaleckyj, S. S. Rich, K. Daly, M. Sale et al., 2010 Robust
358	relationship inference in genome-wide association studies. Bioinformatics 26:
359	2867–2873.
360	Maples, B. K., S. Gravel, E. E. Kenny, and C. D. Bustamante, 2013 RFMix: a
361	discriminative modeling approach for rapid and robust local-ancestry
362	inference. Am. J. Hum. Genet. 93: 278–288.
363	Martin, A. R., M. Lin, J. M. Granka, J. W. Myrick, X. Liu et al., 2017 An Unexpectedly
364	Complex Architecture for Skin Pigmentation in Africans. Cell 171: 1340-
365	1353.e14.
366	O'Connell, J., D. Gurdasani, O. Delaneau, N. Pirastu, S. Ulivi <i>et al.</i> , 2014 A General
367	Approach for Haplotype Phasing across the Full Spectrum of Relatedness.
368	PLOS Genet 10: e1004234.

- Padhukasahasram, B., 2014 Inferring ancestry from population genomic data and its
   applications. Front. Genet. 5:.
- Petersen, D. C., O. Libiger, E. A. Tindall, R.-A. Hardie, L. I. Hannick et al., 2013
- 372 Complex Patterns of Genomic Admixture within Southern Africa. PLOS Genet373 9: e1003309.
- The International HapMap Consortium, 2007 A second generation human haplotype map of over 3.1 million SNPs. Nature 449: 851–861.
- Uren, C., M. Kim, A. R. Martin, D. Bobo, C. R. Gignoux et al., 2016 Fine-Scale
- 377 Human Population Structure in Southern Africa Reflects Ecogeographic
- Boundaries. Genetics 204: 303–314.
- Williams, A., 2016 admix-simu: admix-simu: program to simulate admixture between
  multiple populations.
- de Wit, E., W. Delport, C. E. Rugamika, A. Meintjes, M. Moller et al., 2010 Genome-
- wide analysis of the structure of the South African Coloured Population in the
  Western Cape. Hum.Genet. 128: 145–153.
- Wojcik, G., M. Graff, K. K. Nishimura, R. Tao, J. Haessler et al., 2018 The PAGE
- 385 Study: How Genetic Diversity Improves Our Understanding of the Architecture
  386 of Complex Traits. bioRxiv 188094.
- 387
- 388 Figure Legends:
- **Figure 1:** Computational workflow
- 390

391 The number (n) of individuals included in each dataset, over all ancestral 392 populations. For details, please see the methods section.

393

Figure 2: Comparison between observed global ancestry proportions and "true"
proportions showing RFMix performs more accurately than ADMIXTURE in ancestry
determination.

397

398 Admixture proportions calculated by ADMIXTURE are in black and RFMix in red.

399 Root Mean Square Errors for every comparison are shown.

400

401

402 Figure 3: Boxplot showing the accuracy with which RFMix assigns an ancestral403 origin to a genetic region, stratified by reference population.

404

The median (bold horizontal line) and the upper and lower quartiles are shown. Data faliing outside this range are plotted as outliers. The differences in accuracies across ancestries were assessed using a Wilcoxon non-parametric test. All statistically significant p values (< 0.01) are shown.

# Table 1: Population characteristics of the final merged dataset

Population	Number of individuals included
KhoeSan (Nama and ≠Khomani	284
San)	
European (British)	79
African (Yoruba and Luhya)	35
East Asian (Han)	50
South East Asian (Gujarati)	103

# Table 2: Average admixture proportions

	Previously Reported (Uren <i>et al.</i> 2016) (%)	Simulation (%)	ADMIXTURE (%)	RFMix (%)
Bantu-speaking African	32	27	33	26
KhoeSan	30	33	25	33
European	19	22	26	23
East Asian	7	6	7	6
South East Asian	12	12	9	12





Actual Ancestry Proportions



KhoeSan





