Fast and accurate relatedness estimation from high-throughput sequencing data in the presence of inbreeding

Hanghøj, Kristian; Moltke, Ida; Andersen, Philip Alstrup; Manica, Andrea; Korneliussen, Thorfinn Sand

Published in:
GigaScience

DOI:
10.1093/gigascience/giz034

Publication date:
2019

Document version
Publisher's PDF, also known as Version of record

Document license:
CC BY

Citation for published version (APA):
TECHNICAL NOTE

Fast and accurate relatedness estimation from high-throughput sequencing data in the presence of inbreeding

Kristian Hanghøj 1,2,*, Ida Moltke 3, Phillip Alstrup Andersen 3, Andrea Manica 4 and Thorfinn Sand Korneliussen 1,4,*

1Centre for GeoGenetics, Natural History Museum of Denmark, University of Copenhagen, 1350 Copenhagen K, Denmark; 2Université de Toulouse, University Paul Sabatier (UPS), Laboratoire AMIS, CNRS UMR 5288, Toulouse, France; 3Department of Biology, University of Copenhagen, Ole Maaløes Vej 5, 2200 Copenhagen, Denmark; and 4Department of Zoology, University of Cambridge, Downing Street, Cambridge CB2 3EJ, UK

*Correspondence address. Kristian Hanghøj, E-mail: k.hanghoej@snm.ku.dk http://orcid.org/0000-0003-1941-5495; and Thorfinn Sand Korneliussen, E-mail: tskorneliussen@snm.ku.dk http://orcid.org/0000-0001-7576-5380

Abstract

Background: The estimation of relatedness between pairs of possibly inbred individuals from high-throughput sequencing (HTS) data has previously not been possible for samples where we cannot obtain reliable genotype calls, as in the case of low-coverage data. Results: We introduce ngsRelateV2, a major revision of ngsRelateV1, a program that originally allowed for estimation of relatedness from HTS data among non-inbred individuals only. The new revised version takes into account the possibility of individuals being inbred by estimating the 9 condensed Jacquard coefficients along with various other relatedness statistics. The program is threaded and scales linearly with the number of cores allocated to the process. Conclusion: The program is available as an open source C/C++ program under the GPL license and hosted at https://github.com/ANGSD/ngsRelate. To facilitate easy analysis, the program is able to work directly on the most commonly used container formats for raw sequence (BAM/CRAM) and summary data (VCF/BCF).

Keywords: relatedness estimation; inbreeding; Jacquard coefficients; high-throughput sequencing data; genotype likelihood; next-generation sequencing; threading; population genetics

Introduction

Being able to estimate how closely related 2 individuals are and whether they are inbred is important in several different fields ranging from conservation genetics to medical genetics. For this purpose, numerous coefficients, such as the kinship coefficient and inbreeding coefficients, have been defined and many programs for estimating these coefficients have been proposed.

Notably, the genetic relationship between 2 individuals can be quantified by the extent to which the 2 individuals share their alleles via identity by descent (IBD), i.e., are identical as a result of recent common ancestry. More specifically, for 2 diploid individuals, and thus 4 alleles, there are 15 distinct possible IBD sharing patterns at any given site (detailed identity states). If we ignore the maternal or paternal origin of the alleles, these 15 detailed states can be collapsed into 9 condensed states [1] (here denoted \(j_1, j_2, \ldots, j_9\)), and their corresponding frequencies in the genome of 2 individuals are called the condensed Jacquard coefficients (here denoted \(J_1, J_2, \ldots, J_9\)). These condensed coefficients provide a comprehensive description of the common ancestry between 2 individuals that can be used to infer their familial relationship. Furthermore, many other commonly used...
coefficients, such as the kinship coefficient and inbreeding coefficients, can be expressed as linear combinations of the 9 condensed Jacquard coefficients.

In the specific case in which neither individual is inbred, only 3 of the condensed Jacquard coefficients can be positive, namely, $J_J, J_J$, and $J_J$, which are often also denoted $k_2, k_1$, and $k_0$, respectively, and known as Cotterman coefficients [2]. Numerous approaches, based on either method of moments (e.g., [3]) or maximum-likelihood estimation (e.g., [4]), have been devised to estimate these 3 quantities assuming that the rest are zero and thus that the individuals are not inbred. This includes commonly used methods such as PLINK and KING [5-7]. Importantly, these methods can lead to wrong estimates and conclusions if applied to inbred individuals because the assumption that only $J_J, J_J$, and $J_J$ can be positive is violated. Hence, in the presence of inbreeding one needs to estimate all 9 coefficients. Several methods for doing this have been proposed [6-8]. However, few current tools allow the user to infer all 9 Jacquard coefficients and thus allows for inference of relatedness in the presence of inbreeding.

To simulate data with $L$ sites and $N$ diploid individuals, we first sampled $L$ allele frequencies from a uniform distribution with a minor-allele frequency (MAF) filter on 0.05 and 0.1. For each site for each of the $N$ individuals, we sample 2 alleles using independent Bernoulli trials with the probability of success equal to the allele frequency for the given site, implying that the data are generated under the assumption of Hardy-Weinberg equilibrium. The outcome of these 2 trials represents the genotype. Gametes of these individuals are subsequently generated by sampling either of the 2 alleles from the 2 haplotypes for every site with equal probability. We assume that each site is independent; thus, linkage disequilibrium (LD) is not modeled. Allosomes are disregarded as well.

For the $N$ founder individuals, we simulate offspring to generate 3 different pedigrees. From these pedigrees, we have analyzed pairs of individuals with the expected Jacquard coefficients as presented in Table 3.

We then proceed by calculating genotype likelihoods by assuming different sequencing depths $d = \{1x, 2x, 4x, 8x, 16x\}$, error rate $\epsilon = 0.001$, and number of sites $s = \{10,000, 30,000, 50,000\}$, for the individuals of interest. The per-site-per-individual sequencing depth is given by sampling the depth from a Poisson distribution with parameter $d$ and using the binomial density distribution with $\epsilon$. This approach is similar to the previous approach in Korneliussen and Moltke [11], which...
Figure 1: Scenario 1: 100 independent simulations of 2 outbred cousins across variable sequencing depth and informative sites with a MAF cutoff on 10%. \(J_1\) to \(J_9\) refer to the 9 Jacquard coefficients, \(R\) is the relatedness, and \(F_1\) and \(F_2\) refer to the individual inbreeding coefficients. Simulation (green) are the true values against which we compare ngsRelateV1 (red) and the new program ngsRelateV2 (blue).

Table 1: Probabilities for various allelic states, given modes of IDB from Table 1 in Anderson and Weir [8], with triallelic sites disregarded

<table>
<thead>
<tr>
<th>Allelic state</th>
<th>(J_1)</th>
<th>(J_2)</th>
<th>(J_3)</th>
<th>(J_4)</th>
<th>(J_5)</th>
<th>(J_6)</th>
<th>(J_7)</th>
<th>(J_8)</th>
<th>(J_9)</th>
</tr>
</thead>
<tbody>
<tr>
<td>(A,A,A,A)</td>
<td>(p_1)</td>
<td>(p_2)</td>
<td>(p_3)</td>
<td>(p_4)</td>
<td>(p_5)</td>
<td>(p_6)</td>
<td>(p_7)</td>
<td>(p_8)</td>
<td>(p_9)</td>
</tr>
<tr>
<td>(A,A,A,A)</td>
<td>0</td>
<td>(p_1p_2)</td>
<td>0</td>
<td>(p_1p_3)</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>0</td>
<td>0</td>
<td>(p_2p_3)</td>
</tr>
<tr>
<td>(A,A,A,A)</td>
<td>0</td>
<td>0</td>
<td>(p_1p_2)</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>0</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>(p_2p_3)</td>
</tr>
<tr>
<td>(A,A,A,A)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>(p_1p_2)</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>(p_2p_3)</td>
</tr>
<tr>
<td>(A,A,A,A)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>(p_1p_2)</td>
<td>0</td>
<td>(p_2p_3)</td>
<td>(p_2p_3)</td>
<td>(p_2p_3)</td>
</tr>
</tbody>
</table>

does not model the spatial properties of true recombination and LD.

Results

To test the performance of ngsRelateV2, we use 3 simulated scenarios (see Simulations section) and compare it with ngsRelateV1 [11]. For every scenario, we generate 100 independent simulations for every combination of sequencing effort and number of segregating sites. In Scenario 1, we compare 2 outbred cousins (Fig. 1). As expected, both versions of ngsRelate find not only the correct level of relatedness but also the correct estimates of the 3 relevant Jacquard coefficients \((J_1, J_3, J_5)\). Scenario 2 also includes 2 cousins, but this time we have introduced inbreeding in 1 of the individuals. The parents of the inbred individual are related equivalent to a parent-child relation. In this scenario, even at low sequencing effort and only 10,000 sites, ngsRelateV2 correctly estimates the coefficients of relat-
Relatedness estimation from high-throughput sequencing data in the presence of inbreeding

Figure 2: Scenario 2: 100 independent simulations of 2 cousins, with 1 individual being inbred, across variable sequencing depth and segregating sites with a MAF cutoff on 10%. J9 to J1 refer to the 9 Jacquard coefficients, R is the relatedness, and F1 and F2 refer to the individual inbreeding coefficients. Simulation (green) are the true values against which we compare ngsRelateV1 (red) and the new program ngsRelateV2 (blue).

Table 2: Various relatedness statistics estimated by ngsRelateV2 and the summary statistics on which they are based

<table>
<thead>
<tr>
<th>Statistic</th>
<th>Formula</th>
<th>Summary statistic</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>$r_{AB}$</td>
<td>$J_1 + J_2 + 0.75(J_3 + J_5) + 0.5J_8$</td>
<td>IBD</td>
<td>[12]</td>
</tr>
<tr>
<td>$F_A$</td>
<td>$J_1 + J_2 + J_3$</td>
<td>IBD</td>
<td>[19]</td>
</tr>
<tr>
<td>$F_S$</td>
<td>$J_1 + J_2 + J_5 + J_6$</td>
<td>IBD</td>
<td>[19]</td>
</tr>
<tr>
<td>$\theta$</td>
<td>$J_1 + 0.5(J_3 + J_5 + J_6)$</td>
<td>IBD</td>
<td>[19]</td>
</tr>
<tr>
<td>$F_{12}$</td>
<td>$J_1 + 0.5J_3$</td>
<td>IBD</td>
<td>[12]</td>
</tr>
<tr>
<td>$F_{31}$</td>
<td>$J_1 + 0.5J_6$</td>
<td>IBD</td>
<td>[12]</td>
</tr>
<tr>
<td>Fraternity</td>
<td>$J_2 + J_3$</td>
<td>IBD</td>
<td>[20]</td>
</tr>
<tr>
<td>Identity</td>
<td>$J_1$</td>
<td>IBD</td>
<td>[20]</td>
</tr>
<tr>
<td>Zygosity</td>
<td>$J_1 + J_2 + J_3$</td>
<td>IBD</td>
<td>[20]</td>
</tr>
<tr>
<td>2-3-IBD</td>
<td>$J_1 + J_2 + J_3 + J_5 + J_6 + 0.5(J_4 + J_5 + J_6 + J_8)$</td>
<td>IBD</td>
<td>[16]</td>
</tr>
<tr>
<td>$F_{eff}$</td>
<td>$0.5(J_4 - J_8)$</td>
<td>IBD</td>
<td>[16]</td>
</tr>
<tr>
<td>R0</td>
<td>$(C + G)/E$</td>
<td>IBS</td>
<td>[17]</td>
</tr>
<tr>
<td>R1</td>
<td>$E/(B + D + H + F + G)$</td>
<td>IBS</td>
<td>[17]</td>
</tr>
<tr>
<td>King</td>
<td>$[E - 2(C + G)]/(B + D + H + F + 2F)$</td>
<td>IBS</td>
<td>[17]</td>
</tr>
</tbody>
</table>

IBD: identity by descent; IBS: identity by state.
Figure 3: Scenario 3: 100 independent simulations of 2 cousins, both being inbred, across variable sequencing depth and segregating sites with a MAF cutoff on 10%. $J_i$ to $J_9$ refer to the 9 Jacquard coefficients, $R$ is the relatedness, and $F_1$ and $F_2$ refer to the individual inbreeding coefficients. Simulation (green) are the true values against which we compare ngsRelateV1 (red) and the new program ngsRelateV2 (blue).

Table 3: Expected Jacquard coefficients, relatedness, and inbreeding coefficients for 3 simulated scenarios

<table>
<thead>
<tr>
<th>Scenario</th>
<th>$J_1$</th>
<th>$J_2$</th>
<th>$J_3$</th>
<th>$J_4$</th>
<th>$J_5$</th>
<th>$J_6$</th>
<th>$J_7$</th>
<th>$J_8$</th>
<th>$J_9$</th>
<th>$R$</th>
<th>$F_1$</th>
<th>$F_2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0.25</td>
<td>0.75</td>
<td>0.13</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0.06</td>
<td>0.19</td>
<td>0</td>
<td>0.38</td>
<td>0.38</td>
<td>0.23</td>
<td>0</td>
<td>0.25</td>
</tr>
<tr>
<td>3</td>
<td>0.02</td>
<td>0.02</td>
<td>0.09</td>
<td>0.12</td>
<td>0.06</td>
<td>0.06</td>
<td>0.06</td>
<td>0.38</td>
<td>0.22</td>
<td>0.38</td>
<td>0.25</td>
<td>0.13</td>
</tr>
</tbody>
</table>

edness and inbreeding; however, the estimates of the 9 Jacquard coefficients are somewhat noisy, and $\geq 50,000$ segregating sites are needed to increase the accuracy (Fig. 2). Scenario 3, being the most complex, includes the inbred individual from Scenario 2 and another inbred cousin whose parents are related equivalent to a grandparent-grandchild relationship. Interestingly, with such a complex pedigree, ngsRelateV2 still manages to recover the exact estimates for relatedness and individual inbreeding coefficients, even with only 10,000 segregating sites and a low sequencing depth (Fig. 3). Similarly to the results from Scenario 2, accurate estimates of the 9 Jacquard coefficients required increasing the number of informative sites and/or the sequencing effort. We also applied ngsRelateV2 to these 3 scenarios using a MAF cutoff on 0.05 (Supplementary Figs 1–3). We find that ngsRelateV2 recovers comparable accuracy with a MAF filter on 0.05.

We also applied ngsRelateV2 to real HTS data and compared the estimates with those obtained with ngsRelateV1. We used 6 pairwise related genomes, sequenced to low coverage (~4×), from the Luhya in Webuye, Kenya (LWK), population generated as part of the 1000 Genomes Project [21]. We calculated genotype likelihoods of the related individuals, using ANGSD [18], at
Relatedness estimation from high-throughput sequencing data in the presence of inbreeding

Figure 4: Estimated Jacquard coefficients from 6 pairs of related individuals. The estimates are based on low-depth next-generation sequencing data from the 1000 Genomes Project using ngsRelateV1 and ngsRelateV2. \( J \) refers to the 9 Jacquard coefficients, \( R \) is the relatedness, and \( F_1 \) and \( F_2 \) refer to the individual inbreeding coefficients. CO: cousins; HS: half siblings; PC: parent-child; AV: avuncular; FS: full siblings.

Figure 4: Estimated Jacquard coefficients from 6 pairs of related individuals. The estimates are based on low-depth next-generation sequencing data from the 1000 Genomes Project using ngsRelateV1 and ngsRelateV2. \( J \) refers to the 9 Jacquard coefficients, \( R \) is the relatedness, and \( F_1 \) and \( F_2 \) refer to the individual inbreeding coefficients. CO: cousins; HS: half siblings; PC: parent-child; AV: avuncular; FS: full siblings.

genomic sites with MAF in the LWK population on 0.05, summing up to 4.6 million segregating sites. We not only show that ngsRelateV2 obtains relatedness estimates comparable to those obtained by ngsRelateV1, with this novel software, we also show that all the tested individuals show an inbreeding coefficient <1% (Fig. 4).

In extremely complicated pedigrees with symmetric inbreeding, such as multiple generations of full sibling mating, we find multiple global maxima where several combinations of the 9 Jacquard coefficients, including the expected coefficients, are equally likely. Albeit observing such identifiability challenges, we, importantly, still find accurate relatedness estimates and individual inbreeding coefficients by summing the relevant Jacquard coefficients.

For every pair of individuals, ngsRelateV2 generates and outputs estimates of the 9 Jacquard coefficients, the relatedness, the individual inbreeding coefficients as described above but also other combinations of the 9 Jacquard coefficients: the kinship coefficient, fraternity, and the 3 summary statistics inbred relatedness, identity, and zygosity, suggested by Ackerman and colleagues [20]. It also produces the King statistic [22] based on the 2D site frequency spectrum of pairs of individuals following the methodology of Waples et al. [17]. The latter statistics do not require population allele frequencies.
Computational speed and memory requirements

To take advantage of the increasing number of cores of available on modern computers we employ a multilevel threading approach by parallelizing both the file reading and the actual analysis. In Fig. 5 we analyzed a semi-random dataset consisting of 135 samples mainly from de Barros et al. [23]. The input for the program was a 34-GB BCF file generated with standard bcftools with a liberal 164 million number of single-nucleotide polymorphism sites. We timed the actual runtime (wall clock) and the CPU time for a varying number of cores (1, 2, 4, 8, 16, 32) and noted the memory usage for each run because allocating more cores for the process requires additional internal datastructures and therefore also increases the memory requirements as reported in Table 4. From both the table and figure we observe a near linear correlation between the number of cores and the runtime, with the CPU time remaining almost constant.

Table 4: Run statistics for 34-GB BCF file as a function of number of cores

<table>
<thead>
<tr>
<th>Cores</th>
<th>Memory usage (GB)</th>
<th>Wall clock time (h)</th>
<th>CPU time (h)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>45</td>
<td>85.59</td>
<td>87.14</td>
</tr>
<tr>
<td>2</td>
<td>46.9</td>
<td>41.38</td>
<td>81.39</td>
</tr>
<tr>
<td>4</td>
<td>49.3</td>
<td>23.36</td>
<td>89.03</td>
</tr>
<tr>
<td>8</td>
<td>54.2</td>
<td>11.88</td>
<td>88.69</td>
</tr>
<tr>
<td>16</td>
<td>63.5</td>
<td>6.30</td>
<td>89.73</td>
</tr>
<tr>
<td>32</td>
<td>83.2</td>
<td>3.39</td>
<td>87.81</td>
</tr>
</tbody>
</table>

Presented are the memory requirement, wall clock time (actual runtime), and CPU time; see also Fig. 5.

Conclusion

The tool presented in this Technical Note allows researchers to perform relatedness analysis for inbred individuals in a statistical framework that is especially suited for low-coverage sequence data. The results show that the method performs well for estimating all 9 coefficients, at least when the underlying pedigrees are not extremely complex. And even when the underlying pedigree is very complex, compound summaries of the output, such as relatedness and inbreeding coefficients, will still be correct. The implementation is a fast multi-threaded C++
program that can be directly applied to the most commonly used data files used for HTS data.

**Implementation Details**

The program is implemented in a fast multi-threaded C++ program and takes as input either genotype likelihood files and frequencies, BCF/VCF files as produced from standard tools such as GATK [24] or SAMtools [25], or binary-format PLINK files [3]. We also include an R implementation that we used for simulating data. Of note, the simulations generated in this study do not account for LD. In case of LD between genetic variants, the likelihood function becomes a composite likelihood function. The maximum-likelihood estimate of such a function is consistent with that found with a likelihood function of independent sites [26].

The optimization follows the approach described in Korneliussen and Moltke [11]. The optimization is an accelerated expectation maximization following the squared iterative approach in S3 in Varadhan and Roland [27] and is initialized with a random start point within the parameter space. The borders of the parameter space are manually examined after convergence. Because the expectation maximization algorithm is only guaranteed to find a local optimum, it is recommended to rerun with multiple different seeds although we note that we did not find an issue with multiple local optima in our examples.

**Availability of source code and requirements**

- Project name: ngsRelateV2
- Project home page: http://github.com/ANGSD/ngsRelate
- Operating system(s): platform independent
- Programming language: C++
- Other requirements: htslib (only for parsing VCF/BCF files)
- License: GNU GPL (version 3)
- RRID: SCR_016588
- GigaDB: Snapshots of the code and other supporting data are available in the GigaScience repository [28]

**Additional files**

Supplementary Methods and Results are available via the additional file associated with this article.

**Abbreviations**

HTS: high-throughput sequencing; IBD: identity by descent; IBS: identity by state; LD: linkage disequilibrium; LWK: Luhya in Webuye, Kenya; MAF: minor-allele frequency.

**Competing interests**

The authors declare that they have no competing interests.

**Funding**

K.H. is funded by the Danish National Research Foundation (DNRF94) and the Initiative d’Excellence Chaires d’attractivité, Université de Toulouse (OURASIL); T.S.K. by a grant from the Carlsberg Foundation (CF16-0913), Danish National Research Foundation Centre for GeoGenetics Funding, (DNRF 0094), Lundbeck Foundation GeoGenetics Centre for Brain, Disease, & Evolution grant No. R302-2018-2155; I.M. by Independent Research Fund Denmark (DFF - 4090-00244); A.M. by an ERC Consolidator Grant LocalAdaptation 647787.

**Authors’ contributions**

T.S.K. devised the model. It was first prototyped by P.A.A. as part of a Master’s project under the supervision of T.S.K. and I.M. K.H. implemented and ran all analyses. I.M. and A.M. devised test scenarios and improved early versions of the method. All authors wrote the article.

**Acknowledgements**

We thank the reviewers for their helpful comments.

**References**

18. Korneliussen TS, Albrechtsen A, Nielsen R. ANGSD: Analy-