Development of a model webserver for breed identification using microsatellite DNA marker

Mir Asif Iquebal1, Sarika1, Sandeep Kumar Dhanda2, Vasu Arora1, Sat Pal Dixit3, Gajendra PS Raghava2, Anil Rai1 and Dinesh Kumar1*

Abstract

Background: Identification of true to breed type animal for conservation purpose is imperative. Breed dilution is one of the major problems in sustainability except cases of commercial crossbreeding under controlled condition. Breed descriptor has been developed to identify breed but such descriptors cover only “pure breed” or true to the breed type animals excluding undefined or admixture population. Moreover, in case of semen, ova, embryo and breed product, the breed cannot be identified due to lack of visible phenotypic descriptors. Advent of molecular markers like microsatellite and SNP have revolutionized breed identification from even small biological tissue or germplasm. Microsatellite DNA marker based breed assignments has been reported in various domestic animals. Such methods have limitations viz. non-availability of allele data in public domain, thus each time all reference breed has to be genotyped which is neither logical nor economical. Even if such data is available but computational methods need expertise of data analysis and interpretation.

Results: We found Bayesian Networks as best classifier with highest accuracy of 98.7% using 51850 reference allele data generated by 25 microsatellite loci on 22 goat breed population of India. The FST values in the study were seen to be low ranging from 0.051 to 0.297 and overall genetic differentiation of 13.8%, suggesting more number of loci needed for higher accuracy. We report here world’s first model webserver for breed identification using microsatellite DNA markers freely accessible at http://cabin.iasri.res.in/gomi/.

Conclusion: Higher number of loci is required due to less differentiable population and large number of breeds taken in this study. This server will reduce the cost with computational ease. This methodology can be a model for various other domestic animal species as a valuable tool for conservation and breed improvement programmes.

Keywords: Bayesian network, Breed, Goat, Microsatellite, Prediction, Webserver

Background

Breed of a given species are known to emerge over years during evolution within a specific ecological niche. Each breed is a unique combination of gene in a given gene pool and over the period of time with selection for survival as well as also for productivity due to human intervention. Except cases of commercial crossbreeding under controlled condition, the breed dilution is one of the major problems in sustainability of the breed. The identification of true to breed type animal for conservation purpose is imperative. If we conserve crossbred or admixture breed, its long term sustenance is compromised as breed is not well adapted over period of time to its native ecological niche. Cross breeding of native goats with exotic breeds of goats (Alpine, Saanen and Boer) has shown poor reproductive performance and high mortality rate in higher grade crosses thus selective breeding of true to the breed type animals is desirable with maintained diversity level for successful conservation and long term sustainability of breed [1]. Such identification tool is also needed to establish breed product’s origin in today’s global market [2].

Though breed descriptor has been developed in India to identify breed but such descriptors cover only “pure breed” type animals which excludes more than 2/3rd of population which are either undefined or admixture [3-5]. In case of close resemblance of phenotype it becomes subjective to identify the breed. Moreover, when
degree of admixture is not so conspicuously visible then it is hard to differentiate between true to breed type and “admixed breed”. Advent of molecular tools like microsatellite and SNP have revolutionized the breed identification even from small samples of biological tissue or germplasm without having ova and semen. In case of semen, ova or embryo the breed cannot be identified as there are no visible breed descriptors.

Microsatellite DNA marker based breed identification has been reported in various domestic animals like cattle [6,7], sheep [8,9], goat [10,11], pig [12], horse [13], dog [14] poultry and rabbit [15]. Such methods have limitations namely, non-availability of allele data in public domain, thus each time all reference breed has to be genotyped which is neither logical nor economical. Even if such data is available but computational methods needs expertise of data analysis and interpretation.

The present work aims at development of a model web server for breed identification where one need not do genotyping of all referral breeds each time increasing the cost of molecular level identification. In order to achieve this, we have used 51850 allelic data of microsatellite marker obtained from DNA fingerprinting of 22 goat breeds on 25 loci across India. This methodology demonstrates that it can be used as model for other domestic animal species and breed for identification and conservation for long term sustainability endeavor.

Implementation
Genomic DNA isolation and creation of data set
Blood samples were collected from a total of 1037 unrelated animals belonging to twenty two different Indian goat breeds. The breeds selected were from diverse geographical regions and climatic conditions with varying utilities and body sizes. Genomic DNA was isolated from the blood samples by using SDS-Proteinase-K method [16,17]. The quality and quantity of the DNA extracted was assessed by Nanodrop 1000 (Thermo Scientific, USA) before further use. A total of 51850 allelic data generated by 25 microsatellite (details can be seen at http://cabin.iasri.res.in/gomi/tutorial.html) loci based DNA fingerprinting of 22 goat breeds on 25 loci across India. This methodology demonstrates that it can be used as model for other domestic animal species and breed for identification and conservation for long term sustainability endeavor.

Data Generation by allele detection and genotyping
PCR products were mixed in ratio of 1:1.5:2.2 of FAM (blue), VIC (green), NED (yellow) and PET (red) labelled respectively after determining the optimal pooling ratio and dilution ratio for a set of primers. In order to ensure size calibration of alleles 0.5 μL of this mixture was combined with 0.3 μL of Liz 500 as internal lane standard (Applied Biosystems) and 9.20 μL of Hi-Di Formamide per sample. The resulting mixture was denatured by incubation for 5 min at 95°C to run on automated DNA sequencer of Applied Biosystems (ABI 3100 Avant). The electropherograms were drawn through Gene Scan and used to extract DNA fragment sizing details using Gene Mapper software (version 3.0) (Applied Biosystems). Generated data is numeric in terms of base pair which is size of each allele along with genotype (combination of allele at every diploid locus). The protocol has been described at http://cabin.iasri.res.in/gomi/tutorial.html. The obtained allelic data were further analysed using FSTAT software (http://www2.unil.ch/popgen/softwares/fstat.htm) to compute relative locus differentiation of each breed in the entire dataset.

Bayesian networks as classifiers
Classification is a technique to identify class labels for instances based on a set of features (attributes). Building accurate classifiers from pre-classified data is a very active research topic of machine learning and data mining. In last two decades, many classification algorithms have been proposed including Naïve-Bayes, Neural Network (Multilayer Perceptron), Random Forest and Bayesian Network based classifiers.

Naïve-Bayes, an effective classifier is easy to construct as the structure is given a priori i.e., no structure learning procedure is required. It assumes that features are independent of each other. Although this assumption is not realistic, Naïve-Bayes has surprisingly outperformed many sophisticated classifiers over a large number of
datasets, especially where the features are not strongly correlated [18]. Bayesian Network (BN) is a kind of un-restricted classifier. A common feature of Naive Bayes is that the class node is treated as a special node: the parent of all the feature nodes. However, BN treats the class nodes as an ordinary node, it is not necessary a parent of all the feature nodes. The learning methods and the performance of BN for classification are well described by Friedman et al. in 1999 [19]. It has powerful probabilistic representation for classification. A Bayesian network B which is a graphical model that encodes a probability distribution \( P_B(A_1, A_2, \ldots, A_n|C) \) from a given training set. The resulting model can be used so that, given a set of attributes \( a_1, a_2, \ldots, a_n \), the classifier based on \( B \) returns the label/class \( c \) which maximizes the posterior probability, i.e.

\[
P_B(c|a_1, a_2, \ldots, a_n)
\]

Let \( D = \{u_i, u_2, \ldots, u_n\} \) denotes the training data set. Here, each \( u_i \) is a tuple of the form \( \langle a_1^i, a_2^i, \ldots, a_n^i, c^i \rangle \) which assigns values to the attributes \( A_1, A_2, \ldots, A_n \) to the class variable \( C \). The log likelihood function, which measures the quality of learned model, can be written as

\[
LL(B|D) = \sum_{i=1}^{N} \log P_B(c^i|a_1^i, a_2^i, \ldots, a_n^i) + \sum_{i=1}^{N} \log P_B(a_1^i, a_2^i, \ldots, a_n^i)
\]

The first term in above equation measures efficiency of network \( B \) to estimate the probability of a class given set of attribute values. The second term measures how well network \( B \) estimates the joint distribution of the attributes. Since the classification is determined based on \( P_B(C|A_1, A_2, \ldots, A_n) \), only the first term is related to the score of the network as a classifier i.e., its predictive accuracy. This term is dominated by the second term, when there are many observations. As \( n \) grows larger, the probability of each particular assignment to \( A_1, A_2, \ldots, A_n \) becomes smaller, since the number of possible assignments grows exponentially in \( n \). In our study,
number of feature \((n)\) are the number of alleles (two alleles per locus) \textit{i.e.} 50 and the total number of samples is 1037 which includes 22 breeds (classes). Prediction performance of a Bayesian network has also been compared with Multilayer Perceptron [20] and Random forest algorithm [21].

In this study, WEKA machine learning workbench with extensive collection of machine learning algorithms and data pre-processing methods was used for classification and prediction [22].

### Assessment of the prediction accuracy

The best model was selected using various statistical measures \textit{viz.} sensitivity, specificity, precision or positive predictive value (PPV), negative predictive value (NPV), accuracy, false discovery rate (FDR) and Mathew's correlation coefficient (MCC). Accuracy estimate was obtained using five-fold cross-validation technique [23]. For five-fold cross validation technique, the total observations were divided into five parts. Training was done with four sets of observations and testing with one set. The same was repeated such that each set got the opportunity to fall under the test set. Accuracy for each was recorded and the averages of all these five accuracies were reported. The measures are defined as follows:

#### Table 2 Performance of different classifiers

| Method            | Sensitivity | Specificity | Accuracy | MCC    | FDR    |
|-------------------|-------------|-------------|----------|--------|--------|
| Bayes NET         | 0.858       | 0.993       | 0.987    | 0.851  | 0.142  |
| Naive Bayes       | 0.404       | 0.972       | 0.946    | 0.376  | 0.596  |
| Multilayer-Perceptron | 0.450 | 0.974       | 0.950    | 0.424  | 0.550  |
| Random Forest     | 0.682       | 0.985       | 0.971    | 0.667  | 0.318  |

The best performing classifier is represented in bold.

![Figure 1](http://www.biomedcentral.com/1471-2156/14/S2/1) Confusion matrix to show prediction power of BayesNet for each goat breed.
where \( TP \) = True Positive, \( TN \) = True Negative, \( FP \) = False Positive, \( FN \) = False Negative.

Web implementation
The server is developed using CGI-Perl script, Hyper Text Markup Language (HTML) and Java Scripts to make it more user-friendly and launched using open source web server software program, Apache. Other models like Random Forest, Multiple Layer Perceptron were logically excluded in web implementation ensuring objectivity of identification accuracy. The user needs to submit the microsatellite allelic data having numeric values in base pairs at http://cabin.iasri.res.in/gomi/gomi.html. The data can also be uploaded either using .csv or .txt format or direct entry in the submission form. The server has tutorial for the users for easy understanding with a sample data at http://cabin.iasri.res.in/gomi/tutorial.html.

Results and discussion
In order to evaluate the performance of Bayesian Network classifier with respect to other popular classifiers such as Naïve Bayes, Multilayer Perceptron and Random Forest, were trained and tested using five-fold cross validation and prediction performance measures were averaged over five test sets. These classifiers were applied over the 51850 allelic/microsatellite data of Indian goat breeds and it has been observed that Bayes Network outperformed other methods (viz. Naïve Bayes, Multilayer Perceptron and Random Forest method) with sensitivity (TP Rate), specificity, PPV, NPV, accuracy and MCC values as 0.858, 0.993, 0.860, 0.993, 0.987 and 0.851. The performance of these classifiers is shown in Table 2. Confusion matrix to show prediction power of Bayesian Network for each goat breed is represented in Figure 1. Graphical representation of various evaluation measures (sensitivity or TP Rate, accuracy and ROC area) over all the 22 breeds of goat gives clear picture of the result obtained (Figure 2). The area under ROC (total area equals 1) represents the quality of classification. Higher the value better is the classification which is also evident from our result.

Similar case of microsatellite data based breed identification using Bayesian method has been found with much higher accuracy for example 99.63% accuracy in five Spanish sheep breed viz. Churra, Latxa, Castellana, Rasa-Aragonesa and Merino using 18 microsatellite markers [4]. Similar works have been reported in cattle [24], camel [25] and dog [26].

The novel approach and methodology developed in this study gives higher accuracy which is in similar range of earlier studies in cattle [27]. In some reported cases number of loci needed for breed identification ranged much lower like 3-10 [26,28]. For our study, all the 25 loci were needed which is due to poor differentiation of loci in the breeds. Populations having higher \( F_{ST} \) values always needed minimum loci. Contrary to this, population having low \( F_{ST} \) needs more number of loci and still the accuracy is compromised. For example, Murciana and Granadina populations with 25 microsatellites of low \( F_{ST} \) value (0.0432) have been reported with just 80%
accuracy \[29\]. Contrary to this, in case of horse, where \( \text{F}_{ST} \) was having a range of 0.2 to 0.259, the accuracy has been high up to 95%, even with minimum of 3 loci \[28\].

In case of very low \( \text{F}_{ST} \) like 0.009, the breed identification accuracy has been reported as low as 39-48% in four breeds. The poor success in correct breed assignment is due to weak genetic differentiation and gene flow between populations \[29\]. In our study, the \( \text{F}_{ST} \) values were calculated and were seen to be low ranging from 0.051 at 5th locus to 0.297 at 10th locus and overall genetic differentiation of 13.8%, suggesting more number of loci needed for higher accuracy and we found the expected result in our study (Figure 3). In our observation when loci number was increased this low \( \text{F}_{ST} \) was compensated for identification accuracy. The relationship between locus differentiation \( \text{F}_{ST} \) and accuracy of prediction is proportionate. If \( \text{F}_{ST} \) value in a given population of locus selected are higher (> 0.10) then number of locus needed is relatively less. If \( \text{F}_{ST} \) value of loci in a given population is low (<0.05) then more number of loci is required to achieve accuracy \[26\].

Poor \( \text{F}_{ST} \) in Indian goat population is already reported in many studies related to goat breeds of India \[16,30,31\]. This is happening due to unplanned and indiscriminate mating prevalent in breeding region leads to small effective population size or mating between relatives and consequent genetic drift. The general practice of breeding here is to allow few bucks for the whole village/flock \[30\]. For conservation, proper breeding strategies must be designed by rotating the bucks in their flock since the male:female sex ratio is too low. We found Jhakrana, Marwari and Sirohi having lower sensitivity and MCC (Table 3) which is due to overlapping native breeds of tract where mixing of population prevails in Western India. The low MCC of Jamunapari and Marwari population are obviously expected as lot of

| Breed          | Sensitivity | Specificity | Accuracy (*) | MCC     | FDR     |
|----------------|-------------|-------------|--------------|---------|---------|
| Blackbengal    | 0.958       | 0.998       | 0.996 (0.005) | 0.956   | 0.042   |
| Ganjam         | 0.958       | 0.997       | 0.995 (0.005) | 0.946   | 0.061   |
| Gohilwari      | 0.958       | 0.998       | 0.996 (0.005) | 0.956   | 0.042   |
| Jharkhandblack | 0.833       | 0.994       | 0.986 (0.006) | 0.844   | 0.130   |
| Attapaddy      | 0.854       | 0.997       | 0.990 (0.006) | 0.887   | 0.068   |
| Changthangi    | 0.979       | 0.998       | 0.997 (0.003) | 0.968   | 0.041   |
| Kutchi         | 0.978       | 0.999       | 0.998 (0.005) | 0.977   | 0.022   |
| Mehsana        | 0.854       | 0.996       | 0.989 (0.006) | 0.877   | 0.089   |
| Sirohi         | 1.000       | 1.000       | 1.000 (0.000) | 1.000   | 0.000   |
| Malabari       | 0.917       | 0.993       | 0.989 (0.006) | 0.884   | 0.137   |
| Jamunapari     | 0.458       | 0.980       | 0.956 (0.003) | 0.467   | 0.476   |
| Jhakarana      | 0.625       | 0.990       | 0.973 (0.011) | 0.671   | 0.250   |
| Surti          | 0.750       | 0.995       | 0.984 (0.008) | 0.803   | 0.122   |
| Gaddi          | 0.917       | 0.986       | 0.983 (0.005) | 0.825   | 0.241   |
| Marwari        | 0.667       | 0.976       | 0.961 (0.021) | 0.597   | 0.429   |
| Barbari        | 0.729       | 0.995       | 0.983 (0.009) | 0.790   | 0.125   |
| Beetal         | 0.792       | 0.991       | 0.982 (0.007) | 0.790   | 0.191   |
| Kanniadu       | 0.979       | 0.986       | 0.986 (0.011) | 0.862   | 0.230   |
| Sangamnari     | 0.938       | 0.999       | 0.996 (0.002) | 0.956   | 0.022   |
| Osmanabadhi    | 0.979       | 0.996       | 0.995 (0.006) | 0.947   | 0.078   |
| Zalawari       | 1.000       | 1.000       | 1.000 (0.000) | 1.000   | 0.000   |
| Cheguhu        | 0.791       | 0.989       | 0.981 (0.017) | 0.763   | 0.244   |

*The values in parenthesis are the respective standard deviations computed from 5-fold cross validation.

Data in bold represent the weighted average, where weights are the sample sizes for each breed.
allele are getting introduced through immigrant goat breeds in the respective population [30,31].

**Conclusion**

Through the present study, we are reporting first web server for breed prediction with accuracy of more than 98% using 22 goat breeds of India. The number of loci needed is relatively high due to less differentiable population and large number of breeds taken in this study. The web server can be used for other domestic species thus relevant for global use. Further studies are warranted to look for new algorithm to reduce the number of loci in prevailing conditions of large number of breeds and with lower differentiation especially prevailing in “breed melting pot” regions like India and other major diversity regions of the world. This server will reduce the cost with computational ease. This methodology would become a model for all flora and fauna for variety and breed identification required in improvement, conservation, sovereignty issues in trans-border germplasm movement and management.

**Availability and requirements**

Webserver can be accessed freely at http://cabin.iasri.res.in/gomi/

**Server Name:** http://cabin.iasri.res.in/

**Project home page:** http://cabin.iasri.res.in/gomi/

**Operating system(s):** e.g. Platform independent

**Programming language:** PERL, Java, PHP

**Other requirements:** Internet connectivity

**License:** No restrictions on non-commercial/Research use

**Any restrictions to use by non-academics:** Non-academics may contact corresponding author

**Competing interest**

The authors declare that they have no competing interests.

**Authors’ contribution**

DK, GPSR and AR conceived this study. SPD participated in sample collection and data generation. MAI, S, SKD & VA created the work-flow, web application and performed data analyses. MAI and DK drafted the manuscript. All authors read and approved the manuscript.

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**Author details**

1. Centre for Agricultural Bioinformatics, Indian Agricultural Statistics Research Institute, Library Avenue, PUSA, New Delhi 110012, India. 2. Bioinformatics Centre, CSIR-Institute of Microbial Technology, Sector 39A, Chandigarh 160036, India. 3. National Bureau of Animal Genetic Resources, Karnal, Haryana 132 001, India.

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