Genetic Evaluation of Criollo Cattle and Their Crossbreds in Colombia

RESUMEN

Título: Evaluación genética del ganado criollo y sus cruces en Colombia.

Colombia posee siete razas de bovinos criollos adaptadas a las condiciones tropicales; el número total de animales puros es menor de 18,000, aunque si se consideran los animales criollos cruzados, dicho censo asciende a 50,000 animales aproximadamente. El Proyecto de Desarrollo Ganadero de 1986 sugirió aumentar su número y desarrollar planes de mejoramiento a nivel de finca. Estos objetivos se pueden lograr mediante un programa de evaluación genética nacional que incluya todas las razas criollas y sus cruces. El presente artículo presenta las estructuras básicas de las poblaciones y de los datos, a partir de los cuales es posible realizar una evaluación genética nacional, la cual se analiza en términos de la actual situación colombiana. Se discuten además, métodos lineales y procedimientos computacionales unirraciales y multirraciales para su manejo, y se sugiere el uso de programas de computación. Así mismo, se plantea una estrategia general para desarrollar dicho Programa Nacional de Evaluación Genética, enfatizando en algunos aspectos específicos desde el punto de vista colombiano. La implementación de dicho programa beneficiaría la conservación y el desarrollo de las razas criollas y de sus cruces, además de servir como modelo para desarrollar de manera sistemática la evaluación genética de los bovinos colombianos.

Palabras claves: ganado bovino, criollo, evaluación genética, cruzamientos, componentes de varianza.

INTRODUCTION

Colombia is currently pursuing a Livestock Modernization Plan with the collaboration of producers and governmental institutions. One of the specific objectives of this plan is to improve reproductive and productive traits through the utilization of cattle adapted to tropical conditions. As part of its strategy to achieve this goal, the Livestock Modernization Plan: 1) suggests the development of animal improvement programs at a farm level, and 2) encourages the specialization of researchers in areas of critical importance to allow access and use of advanced methodologies (Añanador, 1996). Colombia is in a good position to achieve these goals because: 1) the country has several Criollo breeds of cattle, although small in number, well adapted to their tropical environment, and 2) it has a sizable number of capable researchers in various research centers and universities. From a genetic improvement standpoint, what is needed is a coordinated effort among producers, government officials, and researchers to develop the infrastructure to genetically evaluate animals of the various Criollo breeds for their use in straightbreeding and crossbreeding. Thus, the objectives of this work are: 1) to present population and data structures that allow the prediction on production of the animal genetic effects in farm conditions, 2) to describe current genetic evaluation methods and computational procedures, and 3) to suggest a strategy for development in medium term, a national genetic evaluation program for the Criollo breeds and crossbreeds in Colombia.

Population and Data Structures for Genetic Predictions under Farm Conditions

Types of Populations

A population is defined as a group of animals (males and females) that interbreed. Populations can be unibreed and multibreed. Unibreed populations contain animals of a single breed. For example, seven unibreed Criollo populations exist in Colombia: Blanco Orejinegro...
(BON), Casanare (C), Chino Santanderiano (CS), Costeño Con Cuernos (CCC), Hartón del Valle (HDV), Romsusianuro (R) and Sam Martino (SM). Multibred populations are formed by animals of several breeds and crossbred groups. For example, a two-bred RZ (R x Zebu) multibred population was formed in the Turunapai Investigation Center (Córdoba region) in the 1980 s. This RZ multibred population was composed of purebred R and Z as well as RZ, 3/4R/1/4Z, 1/2R/1/2Z, 1/4R/3/4Z and several other crossbred groups of other R and Z fractions. This was a complex multibred population. A much simpler multibred population would be one formed by purebreds and first generation crossbreds only (e.g., R, Z, and 1/2R/1/2Z). Several simple multibred populations of this type were to be generated by the mating plans for purebred Criollo bulls of the seven Colombian breeds above mentioned and suggested in the 1986 Livestock Development Project. This plan called for the mating of purebred Criollo bulls to Criollo and to Zebu dams (Bejarano et al., 1986). Criollo cattle will, in all likelihood, continue to be used for intrabred and interf bred mating programs in Colombia in the foreseeable future. Thus, both types of populations will be considered here.

**Number of Animals**

The number of purebred and crossbred Criollo cattle in Colombia is small. The 1986 livestock development plan of Criollo breeds (Bejarano et al., 1986) reported the existence of about 15,000 purebred and upgraded dams and 3,000 purebred bulls of the seven Colombian Criollo breeds (BON, C, CS, CCC, HDV, R, and SM). When crossbred Criollo cattle were included, the total number increased to approximately 50,000 animals. Probably a smaller number of purebred and crossbred Criollo cattle exist today. These population numbers are too small to obtain accurate predictions of genetic values for a large number of parents. It would be desirable to increase the number of animals per Criollo breed as much as economically feasible for their use in straightbreeding and crossbreeding programs. On the other hand, the current number of Criollo cattle per breed should not be a deterrent to conducting the necessary research for development national genetic evaluation programs for these populations. Research in genetic evaluation procedures under Colombia conditions would help determine the distribution and amount of data that could be feasibly collected under field conditions, and the accuracies to be expected from such data sets.

Plans to increase the number of cattle per Criollo breed have existed in Colombia for at least 10 years. The 1986 Livestock Development Project based on Criollo breeds suggested the use of multiplication herds and upgrading to increase the numbers of the seven Colombian Criollo breeds (Bejarano et al., 1986). Unfortunately, this plan has yet to materialize. These two options, upgrading and multiplication herds, may still be the most feasible options to increase the numbers of Criollo cattle under current Colombian conditions. In particular, upgrading would be greatly helped by the use of artificial insemination. A third option that might be considered, provided that sufficient funds and trained personnel were available, would be superovulation and embryo transfer. A joint research project between CORPOICA, the University of Antioquia, and Texas A&M University will preserve 400 embryos per breed of all the Colombian Criollo breeds (Estrada, 1996).

**Contemporary Groups**

A contemporary group is a group of animals that allows a fair comparison among animals of interest. For example, contemporary groups for birth weight (BW), weaning weight (WW), and postweaning gain (PG) are usually defined as follows. Contemporary groups for BW are formed by all calves that: 1) were born in the same farm and the same season, 2) are of the same sex, and 3) came from dams in the same management group. Contemporary groups for WW include all calves that: 1) belong to the same BW contemporary group, 2) were in the same preweaning feeding and management group, and 3) were weaned on the same date. Contemporary groups for PG consider calves that: 1) were in the same WW contemporary group, and 2) were in the same postweaning feeding and management group. It should be noted that PG contemporary groups are nested within WW contemporary groups, which in turn are nested within BW contemporary groups.

The only difference between contemporary groups in unibred and in multibred populations is that unibred contemporary groups contain calves of a single breed only, whereas multibred contemporary groups include calves of various breeds and crossbred groups.

**Connected Data Sets**

A data set with linked contemporary groups that permits the comparison of any pair of animals is a connected data set. A connected data set is the basis of a genetic evaluation system. Connections across contemporary groups are created by using reference animals (e.g., sires) in several contemporary groups. Reference animals need not be in all comparison groups, but they must create enough connections across contemporary groups such that the comparison between any two evaluated animals is possible. Artificial insemination would be a very important tool to quickly achieve a high degree of connectedness across contemporary groups.

Records in a connected data set should contain the following information: 1) identification of the calf, its sire and dam; if available, its maternal grandsire; 2) the breed composition of each calf and its ancestors; 3) the codes for each fixed effect (e.g., contemporary group, sex of calf, age of dam), and 4) the measurement of each trait being evaluated (adjusted if required).

**Pedigree Data Set**

The pedigree data set must include identification numbers of all evaluated animals and their immediate ancestors (sire, dam, maternal grandsire), their birth dates, and for multibred populations, the breed composition of each animal and its ancestors. The animals and ancestors included in the pedigree file will depend on the model chosen for the analysis. For example, if an Animal Model (AM, see definition page 19) is used, all calves, sires, and dams must be included in the pedigree file, but only sires, maternal grandsires, and their male ancestors are needed for a Sire-Maternal Grand sire Model (SMM, see definition page 19).

The pedigree file will be used to construct the inverse of the covariance matrix of additive genetic effects (unibred population; Henderson, 1975 a,b, and 1976 a,b; Quaas, 1976), and the covariance matrices of additive and nonadditive genetic effects (multibred population; Ello, 1996 and 1990 a,b), using sets of computational rules.

**Traits and Genetic Effects**

The traits considered in the evaluation will depend on the objectives of the evaluation, and the capability of the software. For sire summaries in general, the main limitation is the set of traits that can be recorded in the field. Thus, the usual reproduction and survival traits (e.g., age at first calving, calving interval, survival after 3 days, survival to weaning and to yearling), growth traits (BW, WW, PG), and carcass
Figure 1. Hato de Chino Santanderiano

Figure 2. Vaca Blanco Orejinegro

Figure 3. Toro Casanareño
Figure 5a. Torete Costeño con cuernos

Figure 4. Chino Santandereano

Figure 6. Vaca Costeña con cuernos
Figure 7. Novilla Hartón del Valle

Figure 8. Toro Romosinuano

Figure 9. Toro Sanmartinero
traits (e.g., hot carcass weight, fat over the longissimus dorsi, marbling score, shear force) might be the first batch of traits that can be evaluated. As the evaluation system matures over time, either more traits, or a different set of traits that better reflect the prevailing economic interests at that time, might be considered.

The types of genetic effects considered per trait will vary according to the type and the level of inheritance of a trait and the type of population animals belong to. For traits that are highly inheritable, both additive and nonadditive effects need to be considered, particularly interbreed nonadditive genetic effects in multibreed populations. Further, if the dam affects the outcome of a trait (e.g., WW), then maternal effects will be considered in addition to direct genetic effects. Direct genetic effects are the result of an animal's own genes. Maternal effects are caused by the influence of the dam over the calf (e.g., the effect of maternal milk over WW). From a statistical standpoint, each effect within a trait will be treated as a different trait. Thus, a trait influenced by direct and maternal effects (e.g., WW) in a multibreed population may need to consider up to four embedded traits (direct and maternal additive and nonadditive genetic effects).

Assuming that a suitable software package is available, the genetic effects for each trait considered in the model will depend on: (1) the previous knowledge about the inheritance of a trait; for example, a model for a highly inheritable post-weaning trait (e.g., PG) or a carcass trait (e.g., hot carcass weight) will probably consider direct genetic effects, but no maternal effects, and (2) the type of population where animals are being evaluated; for instance, a model for traits of medium to low heritability (e.g., BW, WW) in a multibreed population may include nonadditive genetic effects, in addition to additive genetic effects, if the structure of the data set permits such predictions.

**Genetic Evaluation Methods and Computational Procedures**

**Genetic Evaluation Methods**

The genetic evaluation tests currently used to evaluate animals in the field can be broadly grouped in two categories: linear and nonlinear methods. Linear methods are better suited to predict genetic values of animals for traits that show continuous responses (e.g., BW, WW, PG) than for traits that have categorical responses (e.g., calving ease, calf survival). If traits have categorical responses, then nonlinear methods are preferred to linear ones. The method of choice to predict genetic values of animals for continuous traits is called Best Linear Unbiased Prediction (BLUP; Henderson et al., 1959; Henderson, 1973; Pollak and Quaas, 1980; Elzo, 1983; Elzo and Fumula, 1985; Arnold et al., 1995), whereas Bayesian methods are preferred for nonlinear traits (Foulley et al., 1983; Gianola and Foulley, 1988; Gianola and Fernando, 1986; Hoeschele et al., 1995). Detailed description of nonlinear models and procedures is beyond the scope of this paper. Thus, only linear models will be discussed.

**Best Linear Unbiased Predictor (BLUP) and Mixed Model Equations**

The BLUP predictor minimizes the error variance of a prediction within the group of linear unbiased predictors. BLUP computing using the original formula is not an easy task. Thus, C. R. Henderson et al. (1959) came up with a set of mixed model equations (MME) that yielded not only BLUP of random effects, but also generalized least squares (GLS) of fixed effects in the model. The MME became, and still are, the procedure of choice for computing BLUP.

**Genetic Models**

From a practical standpoint, the purpose of a genetic evaluation model is to account genetic and environmental effects in a population as thoroughly as possible such that the predictions of the genetic animal values are the best that can be computed given the available data. This purpose applies to any kind of model and any type of population.

The genetic evaluation models can be classified according to the type of animal included in the model as follows:

1) **Animal Model (AM):** a model that contains all animals in a vector of random genetic effects. The AM predicts expected breeding values (EBV) and it accounts for all relationships among animals in the data set. These EBV must subsequently be divided by two to compute the expected progeny differences (EPD) that currently are published in sire summaries.

2) **Reduced Animal Model (RAM):** a model that explains nonparents in terms of their parental genetic effects and their random Mendelian factors, thus leaving only parents in vector of random genetic effects. The RAM also predicts EBV, and it accounts for all relationships. The RAM yields the same predictions as the AM. The purpose of the RAM is to save the storage space when computing EBV in large data sets.

3) **Sire-Dam Model (SDM):** a model that contains only parents in the vector of random genetic effects. The SDM predicts EPD, and it accounts for relationships among parents only.

4) **Sire-Maternal Grand sire Model (SMM):** a model that contains only sires and maternal grandsires in the vector of random genetic effects. The SMM predicts EPD, and it accounts for relationships among sires and maternal grandsires only.

5) **Sire Model (SM):** a prediction model that contains only sires in the vector of random genetic effects. The SM predicts EPD, and it accounts for relationships among sires only.

The models SDM, SMM, and SM are approximations to the AM. Their predictions are less accurate than those of the AM (or RAM) because less information is available than the AM. Although it may be desirable to use any kind of model to perform an AM in order to predict genetic values, computing constraints may force the utilization of an approximation model.

All these genetic models can be either unibreed or multibreed depending on the type of population animals they are going to be applied. By definition, unibreed models include additive and nonadditive genetic effects from a single breed only. Multibreed models are an extension of unibreed models in that they consider both intrabreed and interbreed additive and nonadditive genetic effects.

**Genetic Evaluation Model and Assumptions**

A genetic evaluation model could be represented, in general form, as follows:

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  RECORD
  = CONTEMPORARY GROUP (FIXED)
  + OTHER FIXED ENVIRONMENTAL EFFECTS (e.g., age of dam)
  + GROUP (FIXED) GENETIC EFFECTS
  + RANDOM GENETIC EFFECTS
  + PERMANENT ENVIRONMENTAL EFFECTS
  + RESIDUAL
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The assumptions of this model are as follows:

1) The expected value of a record is equal to the sum of the fixed effects in the model: contemporary group (assumed here to include an overall mean) + other fixed environmental effects + group genetic effects.

2) The variance of a record is equal to the variance of the random genetic effects + the variance of the residual. The random genetic effects, the permanent environmental effects and the residual are assumed to be uncorrelated.

This general model applies to all unibreed and multibreed models described above (AM, RAM, SDM, SMM, SM).

**Group genetic effects.** Group genetic effects are usually constructed when animals being evaluated have incomplete pedigree information. In a multibreed model group genetic effects could be formed for both additive and nonadditive genetic effects. The preferred grouping strategy is the accumulated grouping which incorporates information of ancestor groups by exploiting the structure of the additive relationship matrix (Thompson, 1979; Quaas, 1988; Westell et al., 1988). In models with group genetic effects, random genetic effects are computed as deviations from genetic group effects. Group genetic effects are more likely to be needed in field data sets where parents come from many different sources where the level of accuracy of record keeping varies widely. If only a few herds with well-kept pedigree records supply most what bulls needed for straight-breeding and crossbreeding, then genetic groups may not be needed because most (if not all) of the important relationships among evaluated animals will be accounted for. This is the situation with some of the Colombian Criollo breeds (e.g., R, CCC). It is also the case for some of the available multibreed data sets. For example, the SMM model used for the genetic evaluation of the multibreed RZ data set (1980-1992) of Turipaná did not require the use of genetic groups (Elzo et al., 1996) because all relevant relationships among sires and maternal grandsires were accounted for.

**Random genetic effects.** Depending on the trait, additive direct and maternal additive and nonadditive genetic effects can be considered in the model. By the usual assumptions of the infinitesimal theory of quantitative inheritance, additive and nonadditive genetic effects are independent (Kempthorne, 1955). Direct and maternal effects, on the other hand, will usually be correlated. Thus, the structure of the covariance matrix for each animal is block diagonal with two blocks: one for additive direct and maternal genetic covariances among all the traits being evaluated, and another block for the nonadditive direct and maternal covariances among the traits in the analysis. For example, consider the data set of the RZ multibreed population of Turipaná, and assume that, i) a multiple-trait AM for two traits: BW and WW will be used in the evaluation, and ii) only additive and nonadditive (interbreed [RZ] intralocus interactions) are important. Here, i) both the additive and the nonadditive genetic covariance blocks of each animal in the model will be 2 x 2, and an animal's complete block-diagonal covariance matrix will be 4 x 4, and 2) the size of covariance matrix of all random effects will be 4n x 4n, where n is the number of animals in the evaluation.

**Permanent and residual environmental effects.** Permanent environmental effects are considered when there are several records per animal, where all these records can be assumed to be repeated measures of the same trait. There could be direct and maternal permanent. Permanent environmental effects are usually assumed to be uncorrelated to genetic effects; they are also uncorrelated across animals. Thus, the covariance matrix of permanent environmental effects is block diagonal where each block contains covariances among permanent environmental effects for each animal.

Residual effects contain genetic and/or environmental random effects not explicitly accounted for in the model. Residual effects are assumed to be uncorrelated among animals. Thus, the covariance matrix of residual effects is block diagonal, with one block per animal.

**Genetic Base.**

If genetic predictions are obtained for national genetic evaluation summaries, then a factor that needs to be considered is a genetic base. The choice of an appropriate genetic base will be particularly important for genetic evaluations in multibreed populations because of the potential economic implications of comparing animals of different breeds and crossbred groups.

A genetic base is formed by a group of animals whose mean predicted genetic value is the mean from which the predicted genetic values of all animals in the population are deviated. A genetic base can be floating or fixed. A floating genetic base is determined automatically by the genetic evaluation procedure; thus, its composition changes each time animals or data are added to or deleted from a genetic evaluation. A fixed genetic base, on the other hand, is formed by a specific group of animals, and it remains unchanged for a specific period of time (e.g., 10 years).

The criteria used to define the group of animals for a fixed genetic base will depend on the type of population. In unibreed populations, a time factor (e.g., year of entry into the stud) may be sufficient. In multibreed populations, breed composition and time may be needed. For example, in a RZ multibreed population composed of straightbred R and Z, and crossbred groups of these two breeds, a fixed genetic base could be defined by all R animals that entered the stud the first year data were recorded.

**Computational Procedures and Computer Programs.**

Currently all computational algorithms use the MMREML to obtain the BLUP of the random effects in the model either by direct or by iterative procedures. Direct procedures usually involve some kind of factorization (Golub and Van Loan, 1983; Van Vleck and Dwyer, 1985a,b). Iterations may or may not require constructing the LHS. The preferred algorithm to solve very large sets of MMREML is one that iterates on the data provided (Schaeffer and Kennedy, 1986), and it does not require the construction of the LHS.

The best two free-of-charge unibreed computer packages, that can be run on personal computers, are: i) MTDFREML (Boldman et al., 1993), and ii) DMU (Jensen and Madsen, 1993). Both programs can estimate covariance components in addition to predicting genetic values under a great variety of unibreed genetic models. There is a research computer system for multibreed populations resulting from two base breeds (LREM; Elzo, 1996b) that predict genetic values and estimate covariance components for several traits and effects using various multibreed genetic models.
The basic steps involved in the computation of genetic predictions using one of the above packages are as follows:

1) Preparation of input animal files. Users need to prepare a data file and a pedigree file. The data file contains the identification of animals with records, the identification number of each environmental effect in the model, the identification number of the ancestors of the animal with records, and the animal’s records for all traits. The pedigree file contains the number of each animal and its ancestors. In addition to this information, the LREM3 multiregion program requires the breed composition of each animal and its ancestors in the data file and in the pedigree file.

2) Preparation of input parameter files. These files specify the type of model to be run (e.g., AM), effects to be considered per trait, variances and covariances among all traits and effects in the model, and output files.

3) Execution of the programs needed for the desired objective. The set of programs to be executed may depend on whether the objective is genetic evaluation only, estimation of covariance components only, or estimation of covariance components and genetic evaluation. These programs can be executed separately, or sequentially through a script file. A script file is an executable file that contains commands that will be executed by operating system in the order they were specified.

If one of the objectives of the study were to estimate covariances, this would be specified in the parameter files; then, the specified covariances would be taken as initial values by the iterative procedures used to estimate these covariances. The MTDREML and the DMU packages use a derivative-free algorithm to estimate covariance components within a breed (Smith and Graser, 1986). The LREM3 system uses an EM algorithm (Dempster et al., 1977; Elzo, 1996b) to estimate multiregion covariance components.

Strategy to Develop a National Genetic Evaluation Program for the Colombian Criollo Breeds

The development of a national genetic evaluation program for the Criollo breeds in Colombia should be an integral part of the Livestock Modernization Plan. A task force formed by breeders and producers, government officials, and genetic researchers would probably need to be created to develop the necessary policies and to organize the research and development work. A genetic evaluation center should be created in order to provide stability and continuity to the genetic evaluation effort.

General Strategy

A strategy to develop a genetic evaluation program for the Colombian Criollo breeds could consider the following sequential phases:

A. Research Phase:
1) Obtain a current inventory of the animals and historical data sets available.
2) Determine the level of connectedness of various data sets within across Criollo breeds and crossbred groups among Criollo breeds and between Criollo breeds and other breeds (e.g., RZ crossbred groups in Turipana).
3) Agree on a set of traits of economic importance whose measurements are feasible to be collected in the field.
4) Agree on a uniform set of data recording formats for all traits of interest and for all breeds.
5) Agree on a uniform set of adjustment formulas for traits that require additional adjustments prior to their use in genetic evaluation.
6) Define a tentative mating plan for each breed (intrasubbreed only, intrasubbreed and interbreed), and the subpopulations (unibreed and multiregion) created by these mating schemes.
7) Agree on genetic models and procedures to be used in the genetic evaluation.
8) Agree on a set of reference bulls to ensure that the defined unibreed and multiregion subpopulations are connected.
9) Agree on a genetic base from which all genetic evaluations will be deviated.
10) Define the computer resources (hardware and software) for: i) the editing and preparation of the data files; ii) the computation of the genetic predictions; and, iii) the publication of the genetic evaluation summaries.

B. Developmental Phase:
1) Organize a reliable in-farm data recording scheme.
2) Assign responsibilities for the various tasks to be accomplished during the genetic evaluation process to the various organizations and individuals involved. Examples of tasks are: i) gathering and preliminary editing of the data; ii) final editing of the data, construction of the evaluation data and pedigree file; iii) conducting the genetic evaluation; and, iv) publishing the summaries of predicted genetic values.

3) Conduct research to improve the genetic evaluation procedures and the efficiency of the genetic evaluation program.

Specific Considerations

The main areas involved in the general strategy are discussed in more detail below.

Inventory of animals and historical data sets. This inventory should cover all types of data sets where Criollo cattle were used intrasubbreed, crossbred, experimental, field data sets kept by breeders and commercial producers, and pedigree information. This inventory: i) will be used to create the inventory data and pedigree files used to determine connectedness within and across breed groups (purebred and crossbred), and to identify traits in common across the component individual data sets, ii) will create a knowledge base upon which future decisions will be based upon (e.g., common recording formats, adjustment procedures, mating plans), and iii) will be the first step in the creation of a national data base for Criollo breeds and their crossbred groups.

Traits. A national genetic evaluation should include all traits (reproductive, growth, carcass) of economic importance that were recorded in the field data sets. The initial inventory files will probably have only a limited set of traits that were measured in all contemporary groups. The traits most likely to be available will be growth traits (e.g., BW, WW). There could also be some reproductive traits that could be either directly available or that could be recovered from the existing files (e.g., survival from birth to weaning, age at first calving, calving interval). The least recorded traits are likely to be carcass traits (e.g., hot carcass weight, area of the muscle longissimus dorsi, marbling score, tenderness or sicles force); and, their only source might be experimental data sets.

Thus, the first set of traits to be considered for a national genetic evaluation of Criollo breeds is a set of growth traits. Reproduction traits would be considered next. The evaluation of sires (and dams) for carcass traits will probably need to wait until a national carcass recording scheme is in place.

Mating Plans and Subpopulations. Mating decisions are made by breeders and commercial producers. The role of the participating animal breeders will largely be the identification of unibreed and multiregion subpopulations created by these mating decisions, and the identification of bulls to help connect contempo-
ratory groups for genetic evaluation purposes. The most important factor when identifying these unibreed and multibreed subpopulations is the identification of contemporary groups. The key factor that determines whether a contemporary group is unibreed or multibreed is the breed composition of the animals in them. Multibreed contemporary groups must have animals representing several breeds and (or) crossbred groups. It is not necessary to have all base breeds and crossbred groups in every multibreed contemporary group. However, it would be desirable that reference sires of several breed compositions were represented in multibreed contemporary groups. The connected unibreed and multibreed subpopulations found in the inventory data file will reflect past mating decisions based on a great variety of traits ranging from purely experimental goals to purely commercial goals. Mating plans will probably be narrower if only breeder and commercial producer goals are agreed upon. These mating plans will be the ones that define the first set of subpopulations whose data sets will be used for national genetic evaluations.

Genetic Models and Procedures. Several models will need to be explored during the research phase. The type of models tested will primarily depend on the defined subpopulation types (i.e., unibreed or multibreed) and on the pattern of missing data (some effects may be completely confounded). Regardless of the shortcomings that inventory files may have, a clearer vision should emerge of the environmental and genetic effects to be included in models for the defined unibreed and multibreed subpopulations, and the additional data and links across contemporary groups needed. The model chosen for each subpopulation will surely establish an important compromise between accuracy of prediction and practicality. However, this should not be a major concern. These initial models will evolve into more complete (and complicated) ones as the data sets expand in future years.

Genetic Bases. There will be a different genetic base (floating or fixed) for each separate unibreed and multibreed subpopulation. A choice of a genetic base should be made during the research phase. Perhaps all subpopulations should choose the same type of genetic base (floating or fixed). The advantage of floating genetic base is that the computer program automatically defines the base. However, the chosen genetic base may not be the most suitable for the publication of the genetic predictions, particularly for the multibreed subpopulations. Additional computations will be necessary to deviate the predicted genetic values from the mean genetic value of specific group of animals, effectively turning this floating genetic base into a fixed genetic base. A major advantage of a fixed genetic base is that it provides stability for comparisons over a period of time. In addition, genetic predictions of animals that were born, or that became parents, in subsequent years will give breeders and producers information on genetic trends in their herds.

Computer Hardware. A dedicated set of computers is essential for the implementation of national genetic evaluation programs. Various types of computers and printers, and a redundant disk storage system will be needed. Microcomputers could be used for data entry and editing, whereas powerful microcomputers or UNIX workstations will be required for the computation of genetic predictions. The speed and storage capabilities of the microcomputers have increased dramatically in recent years. Perhaps powerful microcomputers (e.g., a 200 megahertz Pentium Pro) with appropriate amounts of memory (e.g., 128 megabytes of RAM), and disk space (e.g., 10 gigabytes) may be enough for the computation of genetic predictions during the research phase. On the other hand, if these machines were also used for the evaluation of the larger Zebu population in Colombia, then perhaps more powerful UNIX workstations (e.g., IBM RS6000) with more memory and disk space would be needed. Server-type microcomputers and UNIX workstations with a redundant disk storage system will minimize data loss due to an hardware failure and should be considered as an option. Impact printers for production runs and laser printers for high-quality printouts would be needed. In general, current and future computing needs and hardware update schedules should play a major role in hardware purchases.

Computer Software. Various types of software will be needed for specific stages of the production of genetic prediction summaries.

1) Data entry and initial editing of data files. Perhaps standard spreadsheet and database management programs could be used here. To eliminate those data file formatting problems, it would be most advisable to choose a single software system that can create standard ASCII files for the next stage of the genetic evaluation process.

2) Final editing of data files, and construction of input data and pedigree files for the genetic prediction programs. These programs are very specific to the populations being evaluated. Thus, specialized software will need to be written (e.g., in FORTRAN) and the necessary compilers will need to be purchased. These computer programs will be the ones that define contemporary groups and all other environmental fixed effect subclasses, adjust trait measurements, and assign sequential numbers to animals in the evaluation.

3) Computation of genetic predictions. Computer programs for this stage can be general or specialized. General programs are more flexible but less efficient than specialized programs. The unibreed computer packages MTDFREML and DMU are general computer programs. The DMU program is more flexible and complete, but more difficult to learn than the MTDFREML. The multibreed computer system LREM3 can currently handle a variety of genetic models for two-breed multibreed populations only. This type of research system is continuously evolving. An earlier version of this system (RLREM) was used to compute covariance components and genetic predictions of sires used between 1980 and 1992 in the RZ multibreed herd at Turipán (Elzo et al., 1996b). The structure of the multibreed subpopulations were defined in the research phase will determine if changes and additions to the LREM3 system will need to be made.

4) Publication of genetic prediction summaries. Distribution of the genetic prediction summaries can be done electronically or in print. Electronic distribution of genetic evaluation files could be done by one (or more) of the following means: diskettes, downloading files from a dedicated computer in the genetic evaluation center, and (or) accessing these files directly in a national genetic evaluation homepage. These last two options would require an excellent telephone or network communications. The software required would be Web communication and editing programs. Printed distribution of genetic evaluations will probably be more expensive than electronic distribution. A desktop publishing software package and a laser printer could be used to produce high quality genetic prediction summaries.

In-Farm Recording Scheme. An in-farm recording program is the core of a national genetic evaluation program. Thus, the cooperation of breeders and producers as well as geneticists in participating experi-
The development of a national genetic evaluation program involving the seven Criollo breeds and their crossbreds will not only achieve this goal, but it will create the infrastructure for a national sire evaluations for all cattle breeds and their crossbreds in Colombia.

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