Letter to the Editor

PROBABILITY OF NONPATERNITY DETERMINED BY MULTIPLE ALLELE CODOMINANT SYSTEMS

To the Editor: The average probability of nonpaternity is usually defined as the likelihood a specific genetic system will exclude a falsely accused male in a paternity action. A number of published reports explore the chances of proving nonpaternity employing evidence from genetic analysis. Weiner [1] briefly reviews the history of these calculations for the major blood group systems. The exact formulas for many of these calculations depend on the specific system being considered (e.g., ABO [2], Rh [3], and MNSs [4]).

For codominant systems where all genotypes are detectable, it is possible to express the probability of nonpaternity in a general form. If

$$p_1, p_2, \ldots, p_k (\sum_{i=1}^k p_i = 1.0)$$

represent the gene frequencies associated with a codominant system with k alleles, then P (nonpaternity) =

$$\sum_{i=1}^{k} \{ p_i [1-p_i] \}^2 + \sum_{i>j} \sum_{i>j} p_i p_j \{ [1-p_i]^3 + [1-p_j]^3 + [p_i+p_j]^3 + [p_i+p_j] [1-(p_i+p_j)]^2 \} \},$$

where the assumption is made that all individuals involved in the paternity case come from a large random mating population at equilibrium (random with respect to the genetic system employed).

The following justifies the above expression for the average probability of paternity for a k allele system. Consider two specific alleles symbolized by A and B with gene frequencies p_i and p_j . Four possible exhaustive and mutually exclusive mother-child pairs arise, namely, (1) AX-AA, (2) AY-AB, (3) BZ-AB, and (4) AB-AB. X represents the presence of any one of the k alleles, Y represents the presence of any one of the k alleles not equal to B (frequency, $1 - p_j$), and Z represents the presence of any one of the k alleles not equal to A (frequency, $1 - p_i$). Therefore, the population frequencies of the four mother-child pairs in a large random mating population at Hardy-Weinberg equilibrium are: (1) AX-AA = p_i^2 , (2) AY-AB = p_ip_j ($1 - p_j$), (3) BZ-AB = p_ip_j ($1 - p_i$), and (4) AB-AB = p_ip_j ($p_i + p_j$). These mother-child pairs each exclude specific paternal genotypes. These genotypes and their associated probability of exclusion are: (1) the AX-AA pair excludes all men who do not possess an A allele. The population frequency of these men is $(1 - p_i)^2$; (2) AY-AB pair excludes all men who do not possess a B allele. The population frequency of these men is $(1 - p_j)^2$; (3) the BZ-AB pair excludes all men who do not possess an A allele. The population frequency

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of these men is $(1 - p_i)^2$; and (4) the *AB*-*AB* pair excludes all men who possess neither an *A* nor a *B* allele. The population frequency of these men is $(1 - p_i - p_j)^2$. The probability that a randomly selected man possess allele *A* or *B* or both is $p_i (2 - p_i) + p_j (2 - p_j) - 2p_i p_j$, giving the complementary probability $1 - [p_i (2 - p_i) + p_j (2 - p_j) - 2p_i p_j] = (1 - p_i - p_j)^2$ of possessing neither alleles *A* nor *B*.

Combining the exclusion probability for each mother-child pair with the frequency of the mother-child pairs in the population gives the probability of exclusion for a specific system from evidence supplied by the genes A and B as $p_i^2[(1 - p_i)^2] + p_j^2[(1 - p_j)^2] + p_ip_j(1 - p_j)[(1 - p_j)^2] + p_ip_j(1 - p_i)[(1 - p_i)^2] + p_ip_j(p_i + p_j)[(1 - p_i - p_j)^2]$. The sum over all k alleles yields the expression given above for the average probability of paternity.

Special cases of the probability of nonpaternity expression are well known. For example, a codominant system with two alleles (k = 2) yields a probability of nonpaternity of p_1p_2 $(1 - p_1p_2)$, which appears in several textbooks [5]. For three alleles (k = 3), the general expression given above agrees with the results obtained by Weiner [1], which are somewhat more compact.

The maximum value for the probability of nonpaternity for k allele codominant systems is achieved when all gene frequencies are equal. That is, the maximum P (nonpaternity) is $(k - 1) (k^3 - k^2 - 2k + 3)/k^4$, which occurs when $p_1 = p_2 = \ldots = p_k = 1/k$, and can be demonstrated by applying Lagrange's method of maximization [6]. Maximum probability of nonpaternity values are given in table 1 for genetic systems with up to 10 alleles $(k = 2, 3, \ldots, 10)$. Also noteworthy is that the probability of nonpaternity increases as the number of alleles within a system increases.

The probability of detecting nonpaternity plays a fundamental role in judging the efficacy of various genetic systems for detecting nonpaternity. For example, the erythrocyte acid phosphatase (EAP) system among whites $(p_1 = .327, p_2 = .612, \text{ and } p_3 = .061)$ gives P(nonpaternity) = .234, whereas among blacks $(p_1 = .223, p_2 = .223, p_2 = .223)$

TABLE 1

MAXIMUM PROBABILITY OF NONPATERNITY FOR CODOMINANT GENETIC SYSTEMS OF & ALLELES WHEN ALL GENOTYPES ARE DETECTABLE

| k | Maximum* | | |
|-----------------------|---|---|--|
| 2 3 4 5 6 | 3/16 30/81 43/256 372/625 855/1296 | = .188 = .370 = .504 = .595 = .660 | |
| 7 8 9 10 | 1698/2401 3045/4096 5064/6561 7447/10000 | $\begin{array}{rcl} & .707 \\ = & .743 \\ = & .772 \\ = & .795 \end{array}$ | |

* Maximum = $(k - 1)(k^3 - k^2 - 2k + 3)/k^4$.

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.771, and $p_3 = .006$), the same probability is .151. Both cases show that the EAP system is fairly effective for detecting nonpaternity when compared with the maximum possible value of .370.

STEVE SELVIN

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