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PROCEEDINGS

SIMPLE MATERNITY SEARCH WITH BAYESIAN NETWORKS

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Abstract. DNA profiles are commonly used in forensic identification problems. It was in England, in 1995, that the first DNA profiles databases were built. They gave rise to new challenges concerning the forensic identification. In this paper it is intended to exemplify how to use the analysis of DNA profiles to solve the problem of simple maternity search. For it it is necessary to make use of a probabilistic expert system (PES), in this case an object-oriented Bayesian network (OOBN).

Keywords: Bayesian networks, DNA profiles, paternity dispute problems

Mathematics Subject Classification: 62F15

1 Introduction

The use of networks that "carry" probabilities began with the geneticist Sewall Wright in 1921. Then they were used in several forms in various areas as social sciences and economics. The models used in these sciences are, in general, linear. Examples are the Path Diagrams or Structural Equation Models (SEM). Non linear models, called Bayesian networks or Probabilistic Expert Systems (PES), are usually used in artificial intelligence.

This work target is the simple maternity problem approach: the recognizing if a woman is the mother of a child – in general a dead child. To solve the real problems data analysis is performed through object-oriented Bayesian networks, which are a PES example, using Hugin¹ software.

2 Simple Maternity Search

From Jornal PÚBLICO 19.01.2012 (translated):

According to Francisco Corte-Real, INML², last year were carried out 5709 kinship biological research exams, concerning 1217 judicial processes. In the previous year (2010) had been carried out 5595 exams in the context of 1379 examinations processes.

The specialist stated that through these tests, paternities and maternities are looked for, being confirmed that the first are the most frequent. In the case of maternity examinations, these are

www.hugin.com – OOBN a resource available in the Hugin 6.4 software.

² INML – Instituto Nacional de Medicina Legal (National Legal Medicine Institute)

requested in cases like crimes of infanticide when a newborn is found dead and seeks to identify the mother, but, according to Francisco Corte-Real, are much less frequent.

According to this charge, approximately 90% of examinations are referring to doubts raised by parents about the paternity of the children and the remaining 10% are made in their private capacity. In the INML took place, only last year, a total of 4405 forensic examinations for biological processes involving criminal cases as 1165 rapes and murders. In 2010, had been carried out 4795 concerning 1105 examinations processes.

Under 60 processes were conducted, for individual examinations of genetic identification, 203 (142 in 2010, to 88 processes), usually requested by prosecutors and for identification of corpses or parts. *End of citation*³.

The news above shows how often, and in what context, the analysis of DNA profiles is used for forensic proposes in Portugal. In this paper it will be analyzed the situation of maternity examinations cases, much less usual than the paternity ones, particularly the requested in cases like crimes of infanticide, when a newborn is found dead and it is imperative to identify the mother. It is supposed that are available the genotypes from the putative mother (*pmgt*) and from the child (*chgt*). This is generally called the simple maternity search.

So the hypotheses are:

 H_P : The putative mother is the true mother of the child

VS.

 H_D : The true mother of the child is another individual chosen randomly from the population, not related with the putative mother.

The evidence is E = (chgt, pmgt) – the child genotype and the putative mother genotype. The posterior odds is, as in (Ferreira and Andrade, 2009),

$$\frac{P(H_P|E)}{P(H_D|E)} = \frac{P(E|H_P)}{P(E|H_D)} * \frac{P(H_P)}{P(H_D)}$$
(1)

and assuming $P(H_P) = P(H_D)$ as usual,

$$\frac{P(H_P|E)}{P(H_D|E)} = \frac{P(E|H_P)}{P(E|H_D)}.$$
 (2)

As only a single likelihood ratio is being considered it may be computed using a Bayesian network, Fig. 1, where the where the nodes **pg** and **mg** are of class *founder* (single node network which states are the observed alleles with the observed population frequencies). Nodes **pmgt** and **chgt** are of class *genotype* (representing the individuals genotypes). Nodes **tpg** and **tmg** specify if the corresponding allele belongs or not to the putative mother. If **ch_match_pm?** is true the child allele is identical to the one of the putative mother, if not it is chosen randomly in the population. Node **chmg** defines the Mendel inheritance being the allele of the individual chose randomly after the

³ http://www.publico.pt/Sociedade/mais-de-5700-testes-de-paternidade-realizados-em-2011-1529758

ancestral alleles. Node **chpg** is the other element of the child genotype pair chosen randomly from the population.

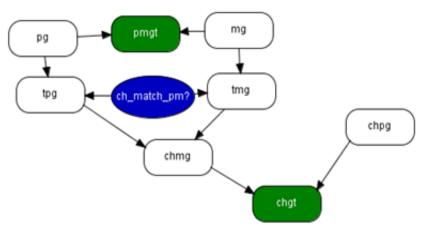


Figure 1. Simple maternity network.

3 Data and Results

To exemplify the application of the described tools, three markers (FGA, D21S11 and PENTA D) were chosen. In Table 1 are presented the genotypes and the allelic frequencies in the population.

Marker		Allele Fre	chgt, pmgt			
FGA	p_{20}	p_{21}	p_{24}	p_{25}	(20, 24); (24,25)	
	0.1421	0.1768	0.1325	0.0718	(20, 24), (24,23)	
D21S11	p_{27}	p_{29}	p_{30}	<i>p</i> _{31.2}		
	0.0246	0.2136	0.2437	0.1138	(27,31.2); (29,31.2)	
PENTA D	p_9	p_{11}	p_{13}	p_{15}		
	0.1984	0.1777	0.2066	0.0250	(11, 13); (9,11)	

Table 1. Genetic profiles and population frequencies for the chosen markers

The obtained results are presented in Table 2. Among many possible conclusions it must be highlighted that when there is a share of an allele with low frequencies in the population the probability of the yes hypothesis is very high, as it is expected.

Marker	FGA	D21S11	PENTA D	Rescaled	
ch_match_pm	yes	0.7848	0.6872	0.5845	0.9185
en_maien_pm	no	0.2152	0.3128	0.4155	0.0815

Table 2. Analysis results

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