DNA Databases for Offender Identification in Europe — The Need for Technical, Legal and Political Harmonization

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In 1995, a national database has been established quite successfully in the U.K.. It is being used for the identification of suspects using short tandem repeat (STR) typing results from casework as well as from reference DNA samples obtained from suspected and convicted offenders. The introduction of multiplex PCR typing systems allowing the simultaneous analysis of ten independent loci or more greatly facilitates the rapid typing of samples and computer-based storage of results in large DNA profile databases. However, in order to introduce such a database as well at the European level, it must be recognized that the legal systems in the member states of the European Union are quite diverse and may not allow the storage of personal genetic data for the purpose of criminal investigation. At present, there is still a significant heterogeneity among the European countries already concerning the possibility to obtain DNA samples from suspects and the acceptance of DNA evidence in casework [for review, see ref. 1].

There is no generally agreed model regarding the organisational structure of a national DNA database. Therefore, Fig. 1 may serve as an example for such a database exhibiting typical features which should ascertain the efficient use in criminal investigations and at the same time provides a maximum of data protection and quality assurance for the DNA profiles entered. This model is divided into three separate organisational areas: the DNA database with profiling laboratory for typing and storage of anonymous DNA samples collected from offenders only for the purpose of database searches; an independent database only for storage of personal records and identification tags used to anonymize the DNA data-

base samples; the police carrying out routine casework investigations on crime scene samples.

Thus, the database is completely separated from casework investigations only serving as an intelligence tool for offender identification. The typing of reference samples from known offenders submitted to the database has to be subjected to rigorous internal quality control and quality assurance procedures to avoid storage of unconfirmed or erroneous typing results, as these could lead to a wrongful exclusion of a perpetrator. In a criminal investigation regarding the origin of an unknown crime scene sample, DNA typing would be carried out in a routine lab on behalf of the police, and the results would then be submitted to the database for a search against the profiles of known offenders (person-to-scene match) or against other samples from unsolved crimes (scene-to-scene match). If a match is found, the database lab can retrieve the stored reference sample for a confirmatory analysis before forwarding the respective ID code to the personal database. The police unit carrying out the case investigations will then be informed about the identity of the suspect. If arrested, a fresh DNA sample has to be obtained from the suspect for further investigations and to serve as evidence in court.

At present, national DNA databases are in operation in 4 European countries. Plans for a database are at different levels of preparation in 8 more countries. Only 4 countries do not plan to introduce a database in the near future (see Table 1). At the political level, a decision has been reached in 1997 between the members of the European Union to create a framework for a European DNA Data-

Table 1: DNA Databases in Europe

Database in operation	Date of introduction	Database in preparation	Date of legislation (date of planned operation)	Currently no plans for database
UK	April 1995	Belgium	September 1998	Ireland
Netherlands	1997	Denmark	?	Italy
Austria	October 1997	Finland	July 1997 (1.1.1999)	Greece
Germany	April 1998	France	end of 1998	Portugal
•	-	Norway	September 1997	-
		Spain	?	
		Sweden	January 1999	
		Switzerland	end of 1998	

base for offenders convicted for sexual abuse of children. To allow the exchange of DNA profiling data for this purpose, agreements have to be reached regarding the typing technology and the selection of standard DNA systems forming the core of the database. Recommendations have been made by the DNA Working Group of the European Network of Forensic Science Institutes (ENFSI – a network of police and government laboratories) and

by Interpol to use the following STR loci as core systems: TH01, vWA, FGA, D21S11. These loci have initially been recommended as suitable for standardization by the EDNAP (European DNA Profiling) Group (a working group of the International Society for Forensic Haemogenetics – ISFH) based on a series of collaborative exercises [2-4].

Table 2: European Countries with DNA Databases in Operation

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U.K.	Netherlands	Austria	Germany			
Custodian /Location of Database						
Forensic Science Service,	Dutch Forensic Science	Central DNA Typing	Bundeskriminalamt (BKA),			
central database lab in	Laboratory, Rijswijk	Laboratory, Institute of	Wiesbaden			
Birmingham		Legal Medicine, Innsbruck	(Federal Criminal Office)			
Samples stored and entry						
DNA profiles and refer-	<u>DNA profiles</u> only of:	<u>DNA profiles</u> and <u>DNA</u>	<u>DNA profiles</u> only of:			
ence samples of:	 convicted offenders 	reference samples of:	- suspects			
- suspects	- unknown samples	- suspects	- convicted offenders			
 convicted offenders 	for serious crimes with 2	- convicted offenders	- unknown samples,			
- unknown samples	years imprisonment or	- unknown samples,	for serious crimes with one			
for "any recordable	more after court order	for crimes against life and	year imprisonment or more,			
offense"		health, sexual abuse,	sexual abuse and other seri-			
		robbery, theft, arson,	ous crimes,			
		blackmail, drug-related	at present only for results			
		and other serious crimes	obtained from routine case-			
			work when DNA typing was			
			ordered by a judge			
Anonymization requirement	ents					
anonymous storage of	anonymous storage of	anonymous storage of	open storage of DNA profiles			
reference samples and	DNA profiles only, sepa-	reference samples and	together with personal data,			
DNA profiles, separate	rate register for personal	DNA profiles, separate	typing of anonymized per-			
register for personal	records	register for personal	sonal and crime scene sam-			
records	(crime samples can be	records outside the central	ples in police and university			
	stored)	DNA lab	laboratories			
Removal of entries			_			
Acquitted suspects only	offenders: after 30 years	acquitted suspects only	routine controls for samples			
	samples: after 18 years		to be removed every 5 years			
No. of entries (June 1998))					
263,000	offenders: 200	4,500	no statistics available yet			
	unknown samples: 400		•			
DNA systems used (see als	so Table 4)					
- Quadruplex	- Quadruplex	SGM	4 European core systems			
- SGM	- SGM		+ SE33			
- TGM						
Remarks						
	Change of legislation		Additional legislation pro-			
	planned to allow entry of		posed to obtain samples from			
	offender profiles without		convicted offenders in cases			
	court order		where no DNA typing was			
			carried out during investiga-			
			tion			

Table 3: European Countries with Databases in Preparation (if no information is given, the respective issue is still under discussion).

(if no information is given, the respective issue is still under discussion)					
Database custodian	Entry criteria	Sample storage and removal periods	DNA systems used for typing		
Belgium National Institute of Criminalistics, Brussels	Convicted criminals with court order for crimes with 3 years of imprisonment or more	DNA profiles only from - convicted offenders - unknown samples, removal after 30 years	4 European core systems + at least 3 additional STR systems (not yet defined)		
Denmark University Institute of Forensic Genetics, Copenhagen	no details available yet, a co submitted to the parliament	ommission report has been			
Finland Crime Laboratory, National Bureau of Inves- tigation, Vantaa	Suspects for crimes with 1 year of imprisonment or more, for offenders con- victed before 1.7.97 also retrospectively if still held in prison	DNA profiles and DNA reference samples from - suspects - convicted offenders - unknown samples, removal after 1 year if suspect is acquitted, legal limit for data storage 10 years (law may be changed for DNA profiles)	Promega or ABI multiplex STR systems, no final decision yet		
France	Sexual assault on children				
Norway University Institute of Legal Medicine, Oslo	Convicted criminals with court order for sexual abuse, crimes against life and health, crimes posing danger to the public (e.g. arson), blackmail and robbery	DNA profiles only from - convicted offenders - unknown samples, no removal except after death or proven innocence	ABI SGM Plus likely, no final decision yet		
Spain	Legislation had been proposed in 1995 and was rejected. It will be presented again in a few months.				
Sweden SKL – National Institute of Forensic Science, Linkøping	Convicted criminals for crimes with 2 years of imprisonment or more	DNA profiles only from - convicted offenders - unknown samples, removal 10 years after release from prison (without further offense)	ABI Profiler		
Switzerland University Institute of Legal Medicine		DNA profiles and DNA reference samples may be stored, removal periods are under discussion	ABI Profiler (Plus) likely, no final decision yet		

The surveys from Table 2 (databases in operation) and Table 3 (databases in preparation) represent the situation of DNA database projects in Europe in June 1998. In a number of countries, no final decisions have been made yet, or changes may still be possible to the information given here.

Regarding the system standardization, most countries are using or planning to use either the SGM (second generation multiplex) developed and used by the Forensic Science Service (FSS) for the U.K. National DNA Database, or multiplex PCR systems offered by commercial companies like Promega or Applied Biosystems (for

Tuble 4. Composition and properties of STR multiplexes selected for databases				
Multiplex kit/loci	STR system composition	Chance for a random match		
SGM	TH01 ⁺ , vWA ⁺ , FGA ⁺ , D8S1179, D18S51,	1 in 50 Million		
	D21S11 ⁺ , AMG			
ABI Profiler*	TH01 ⁺ , vWA ⁺ , FGA ⁺ , TPOX, CSF1PO,	1 in 3.5 Billion		
	D3S1358, D5S818, D7S820, D13S317,			
	AMG			
ABI SGM Plus	TH01 ⁺ , vWA ⁺ , FGA ⁺ , D2S1338, D6S477,	more than 1 in 100 Billion		

Table 4: Composition and properties of STR multiplexes selected for databases

TH01⁺, vWA⁺, FGA⁺, D21S11⁺

TH01⁺, vWA⁺, FGA⁺, D21S11⁺, SE33

D21S11⁺, AMG

D8S1179, D16S539, D18S51, D19S433,

further details, see other contributions to this volume). As these multiplexes comprise a number of common and different loci, efforts are being made to include at least the four European core systems in all multiplexes offered. All commercially available kits also contain the XYchromosomal Amelogenin locus (AMG) suitable for male/female detection. Nevertheless, the discrimination power of the four core loci is much less compared to the systems selected in national database projects (see Table 4). This may limit the future use of some of the national databases at the European level. The concept of "uniqueness" of a DNA profile in a database which was the basis of decision for selecting 13 STR loci in the United States for the national CODIS database, has not been adopted yet by most of the European countries.

German database loci

European core loci

In all European countries, specific legislation was required for the creation of national DNA databases, as the existing laws either prohibited the taking of a blood or saliva sample from suspects without consent or outside police investigations only for the purpose of a database, or the use of DNA profiling in criminal casework, and the storage of DNA profiles in computerized databases.

The protection of privacy rights at different levels has led to two different database models: in a number of countries, DNA profiles as well as reference DNA samples from suspects and/or convicted criminals may be stored anonymously in a central database facility, which enables a rigorous quality control of typing procedures and results, as well as further internal controls of a matching sample identified in a database search before the information about a match is being disclosed to the police. The storage of reference samples allows also to update database entries for future improvements in typing technology. In contrast, several other countries have decided that these reference samples (but not the crime scene samples) must be destroyed after completion of the typing procedure to prevent any illegal analyses of the genomic DNA samples. In Germany, DNA profiles may therefore

be stored without anonymization in a central police database, but the DNA laboratory responsible for the typing the (anonymized) casework samples has no access to the (non-anonymized) database records to verify the correctness of the entries. In these countries, the current typing technology has to be maintained over the next decades without the possibility of future enhancements for the existing records (except after having obtained a fresh sample again from casework).

1 in 10 Million 1 in 100,000

Further heterogeneity is observed regarding the crimes which may lead to a DNA database entry, the selection of persons, the basis of decision, as well as the storage periods. Criteria for a database entries may be as follows:

- all suspects or convicted offenders only (with or without a court order),
- retrospectively also for convicted offenders already serving prison sentences,
- for any recordable offense,
- sexual abuse (all cases or children only),
- crimes typically associated with stain evidence (e.g. serial theft, robbery, blackmail),
- severe crimes depending on a minimum period of imprisonment (typically 1-3 years),
- crimes against health and life,
- serious crime (e.g. organized crime),
- crimes causing danger to the public (e.g. arson).

The storage periods are either indefinite (except for acquitted suspects, or convicted offenders with proven innocence in a later trial), or limited to explicit periods between 10 and 30 years starting either from the date of database entry or from the date of release from prison.

This survey emphasizes the need for harmonization of these technical and legal issues at the European level in spite of considerable heterogeneities of the cultural, political and legal conditions among the European countries, which are based on historical developments and a

⁺ European core systems; * a different composition will be made available which also includes D21S11

different national heritage in each country. Nevertheless, the current developments regarding DNA databases represent a significant change in most countries. To further improve the usefulness of this powerful method in criminal investigations, and to respect and protect individual privacy rights at the same time, a continued collaborative effort of scientific and legal experts will be necessary.

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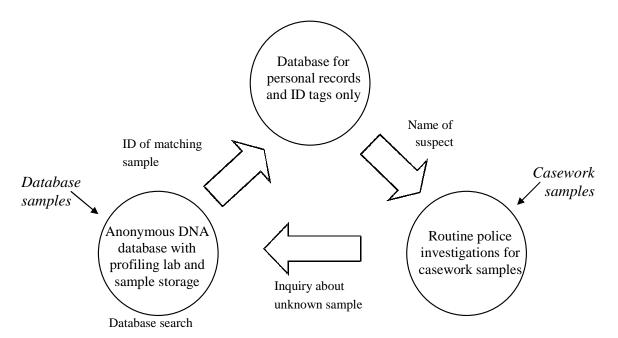


Figure 1: A database model