



**HAL**  
open science

## Survey of European programmes for the epidemiological surveillance of congenital toxoplasmosis.

Antoine Bénard, Eskild Petersen, Roger Salamon, Geneviève Chêne, Ruth Gilbert, Louis Rachid Salmi

► **To cite this version:**

Antoine Bénard, Eskild Petersen, Roger Salamon, Geneviève Chêne, Ruth Gilbert, et al.. Survey of European programmes for the epidemiological surveillance of congenital toxoplasmosis.. *Euro surveillance: bulletin européen sur les maladies transmissibles = European communicable disease bulletin*, 2008, 13 (15). inserm-00405336

**HAL Id: inserm-00405336**

**<https://www.hal.inserm.fr/inserm-00405336>**

Submitted on 20 Jul 2009

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# Survey of European programmes for the epidemiological surveillance of congenital toxoplasmosis

Bénard Antoine <sup>1 4 \*</sup>, Petersen Eskild <sup>2</sup>, Salamon Roger <sup>1 4</sup>, Chêne Geneviève <sup>1 4</sup>, Gilbert Ruth <sup>3</sup>, Salmi Louis Rachid <sup>1 4</sup>, for the European Toxo Prevention Study Group (EUROTOXO)

<sup>1</sup> Centre épidémiologie et biostatistique INSERM : U897, Université Victor Segalen - Bordeaux II, FR

<sup>2</sup> Statens Serum Institut Statens Serum Institut, DK

<sup>3</sup> Centre for Paediatric Epidemiology and Biostatistics Institute of Child Health, University College London, London, GB

<sup>4</sup> Centre Hospitalier Universitaire de Bordeaux, FR

\* Correspondence should be addressed to: Antoine Bénard <antoine.benard@isped.u-bordeaux2.fr>

## Abstract

The objective of this investigation was to describe systems for the epidemiological surveillance of congenital toxoplasmosis implemented in European countries. In September 2004, a questionnaire, adapted from the evaluation criteria published by the United States Centers for Disease Control and Prevention, was sent to a panel of national correspondents in 35 countries in the European geographical area with knowledge of the epidemiological surveillance systems implemented in their countries. Where necessary, we updated the information until July 2007. Responses were received from 28 countries. Some 16 countries reported routine surveillance for toxoplasmosis. In 12 countries (Bulgaria, Cyprus, Czech Republic, England and Wales, Estonia, Ireland, Latvia, Lithuania, Malta, Poland, Scotland and Slovakia), surveillance was designed to detect only symptomatic toxoplasmosis, whether congenital or not. Four countries reported surveillance of congenital toxoplasmosis, on a regional basis in Italy and on a national basis in Denmark, France and Germany. In conclusion, epidemiological surveillance of congenital toxoplasmosis needs to be improved in order to determine the true burden of disease and to assess the effectiveness of and the need for existing prevention programmes.

**MESH Keywords** Disease Notification ; methods ; statistics & numerical data ; Disease Outbreaks ; statistics & numerical data ; Europe ; epidemiology ; Humans ; Incidence ; Population Surveillance ; methods ; Risk Assessment ; methods ; Risk Factors ; Toxoplasmosis, Congenital ; epidemiology

## Introduction

Toxoplasmosis is caused by a protozoan parasite (*Toxoplasma gondii*). While toxoplasmosis infection is often benign, congenital toxoplasmosis (transmission to the foetus when a pregnant woman acquires toxoplasma infection for the first time during pregnancy) can lead to severe sequelae for the foetus and the newborn with visual or neurological impairment or death.

It is important to evaluate the burden of toxoplasma infection in the general population, as well as in pregnant women, foetuses, newborns and children, because this contributes to the rationale behind the different screening programmes currently performed (none, prenatal or postnatal) [1–3]. Frequency and severity of a disease are the basic measurements used to assess its burden, and data on this can be collected in specific studies or surveillance systems. The value of epidemiological surveillance is that it can be used to monitor trends over time. Public health strategies to prevent congenital toxoplasmosis differ between European countries. It is still being debated which are the best methods for controlling congenital toxoplasmosis, and the debate is not always based on accurate information.

The EUROTOXO project (<http://eurotox.isped.u-bordeaux2.fr>) is a European consensus initiative aimed at defining the implications of current scientific knowledge for a research agenda and for policy decisions on how best to prevent congenital toxoplasmosis and its consequences. The project has reviewed the state of the knowledge concerning the burden of toxoplasma infection in Europe. This article presents a systematic review of the systems implemented in European countries for the epidemiological surveillance of toxoplasmosis.

## Methods

### Source of information

We identified contacts for national surveillance programmes in 30 European countries (Table 1) from the following sources:

- the members of the Eurosurveillance Editorial Board listed on the Eurosurveillance website at the time;
- the Inventory of Resources for Infectious Diseases in Europe (IRIDE) (<http://iride.cineca.org/public/invcountries.html>);
- and the European Programme for Intervention Epidemiology Training (EPIET) network (<http://www.epiet.org/>).

Contacts for six other European countries (Albania, Bosnia-Herzegovina, Bulgaria, Croatia, Macedonia, and Serbia-Montenegro) were identified by Google search.

We did not find correspondents for Andorra, Monaco or Northern Ireland. The list of correspondents is shown in Table 1. All contacts were sent emails in September 2004 and those who did not respond were sent three further emails in January/February, April, and July 2005. We maintained contact with our correspondents in each participating countries until July 2007 and updated the data when a change in the surveillance systems was signalled. This was the case for France (implementation of a new surveillance system) and Denmark (surveillance system stopped).

### Data collection and interpretation

We developed a comprehensive questionnaire, based on the criteria published by the United States' (US) Centers for Disease Control and Prevention (CDC) for the evaluation of epidemiological surveillance systems [4]. Epidemiological surveillance was defined as ongoing and systematic collection, analysis, and interpretation of health data in the process of describing and monitoring a health event. The survey included questions about the objective of the surveillance system, the description of the health event under surveillance (case definition), the population under surveillance, the period of data collection, who was responsible for case reporting (sources of information) and a flow chart describing the system. We also asked how often the data were analysed and fed back to the reporting sources, and for the estimated costs of the toxoplasmosis surveillance system.

The usefulness of a given surveillance system was evaluated according to the following criteria:

- simplicity (ease of operation), flexibility (adaptability to changing information needs or operating conditions) and acceptability (cooperation of people on whom the system depends) based on the number and qualification of the reporting sources;
- sensitivity (proportion of cases detected by the system) and representativeness (the ability to describe the distribution of cases over time and in the population) based on the qualification of the reporting sources and on the figures available from the surveillance systems;
- timeliness (delay between steps in the system) based on the frequency of analysis and reports distribution.

These criteria are described in the US CDC's guidelines (<http://www.cdc.gov/mmwr/preview/mmwrhtml/00001769.htm>) [4,5].

## Results

We received responses from 28 of 35 countries. Seven countries (Albania, Luxembourg, Croatia, Hungary, the Former Yugoslav Republic of Macedonia, Serbia-Montenegro and Spain) did not send a response at all. Information on Denmark and France was updated in July 2007.

Of the 28 countries that responded, 12 did not have a surveillance system for toxoplasmosis (congenital or not). The 16 countries that did report to have a system for the epidemiological surveillance of toxoplasmosis in place, are almost all situated in central or eastern Europe (Figure) (Table 2). Poland has the oldest surveillance system (dating from 1966), while the most recent systems are in Cyprus, Ireland and Malta (dating from 2004).

Only four countries operate surveillance specifically for congenital toxoplasmosis: Denmark, France, Italy and Germany.

In **Denmark**, a nationwide neonatal screening programme based on neonatal Guthrie card testing for toxoplasma-specific IgM was implemented in 1999 but discontinued on 31 July, 2007 (Petersen E; personal communication). The Danish National Health Board found insufficient evidence that treatment for toxoplasmosis was effective, neither in preventing later attacks of ocular toxoplasmosis in children born without ocular lesions nor in preventing further attacks in children born with ocular lesions [6]. In case of a positive Guthrie result, peripheral blood samples were taken from the newborn and the mother and analysed for IgM, IgA and IgG profiles. The epidemiological surveillance system was based on this screening programme and therefore included all infants with congenital toxoplasmosis, whether or not they had clinical manifestations. Surveillance and all laboratory analyses were coordinated by Statens Serum Institut in Copenhagen.

In **France**, a surveillance system for congenital toxoplasmosis was initiated in May 2007 which lies in the area of responsibility of the French National Institute of Public Health (Institut national de Veille Sanitaire; InVS) and the National Reference Centre for Toxoplasmosis (CNR toxoplasme). The surveillance includes fetuses, newborns and children until the age of one year. Congenital toxoplasmosis cases are notifiable and defined as:

- Detection of *T. gondii* in body tissues or fluids by polymerase chain reaction (PCR), inoculation of mice, cell culture or immunocytochemistry;
- Detection of specific IgM or IgA antibodies;

- Neosynthesis of specific IgG, IgM or IgA antibodies;
- Stable specific IgG titres until after the age of one month;
- Persistently stable specific IgG titres until the age of one year.

Cases are notified by laboratories qualified for antenatal or postnatal diagnosis.

In **Germany**, congenital toxoplasmosis cases have been notifiable since 2001, when a nationwide surveillance system was implemented under the Protection Against Infection Act. The case definition of congenital toxoplasmosis is based on at least one of the following criteria:

- Demonstration of *Toxoplasma gondii* in body tissues or fluids;
- Detection of specific IgM or IgA antibodies;
- Persistently stable specific IgG titres or a single elevated specific IgG-titre.

Laboratories report anonymised cases to the Robert Koch institute in Berlin. Part of the data can be accessed at <http://www3.rki.de/SurvStat/QueryForm.aspx>. Quarterly summaries and yearly reports are also published [7].

In **Italy**, surveillance is confined to a regional programme in the Campania region, which has been running since 1997. The population under surveillance are living newborn babies. A case of congenital toxoplasmosis is defined as the persistence of specific IgG antibodies until the age of one year. Cases are reported by social workers, paediatricians and neonatologists. Information about toxoplasmosis primary infection among pregnant women is collected retrospectively on medical records, and information about congenital toxoplasmosis and complications among congenitally infected children are collected prospectively. The creation of a nationwide surveillance system is still being debated.

In the 12 other countries (Bulgaria, Cyprus, Czech Republic, England and Wales, Estonia, Ireland, Latvia, Lithuania, Malta, Poland, Scotland, and Slovakia, see Table 2), the health event under surveillance is toxoplasmosis (congenital or not), as defined by the European Union (symptomatic toxoplasmosis cases serologically confirmed) [8]. It is considered a notifiable disease and subject to continuous data collection (Table 2). Cases are reported by physicians, epidemiologists, or laboratories. Several sources of reporting contribute to the systems, except in Cyprus, Lithuania, Poland and Slovakia, where the physicians are the only health professionals to report cases, and in the Czech Republic and Scotland, where cases are declared only by epidemiologists and laboratories, respectively.

All 16 surveillance systems analyse the data regularly (from daily to annually). The reports are sent to the health authorities weekly to annually.

Only two countries were able to provide data about the costs of the system. In Italy, the global cost of the regional pilot programme is estimated to be 68,000 Euros a year for 67,000 to 70,000 live births. In Denmark, the cost of the nationwide surveillance system was estimated to be 600,000 Euros a year.

## Discussion

Our study provides detailed, up-to-date information on systems implemented for the surveillance of toxoplasmosis (congenital or not) in 28 European countries. We have identified a high degree of heterogeneity.

12 countries do not have any surveillance system for toxoplasmosis in place. In 12 countries, the event under surveillance was symptomatic toxoplasmosis. Five of those countries did not provide details about the qualification of the physicians who reported the information. In the field of toxoplasmosis, gynaecologists, ophthalmologists, paediatricians or neurologists are able to diagnose toxoplasmosis at different stages of the disease. Therefore, it is important that all those specialists take part in the surveillance process. However, toxoplasmosis is a notifiable disease in all those countries, and we assume that all registered medical practitioners are involved in the surveillance system.

Denmark, France, Germany, and Italy (the latter only at regional level), are the only participating European countries who have implemented a surveillance system that is specifically dedicated to congenital toxoplasmosis and that is able to detect symptomatic as well as asymptomatic cases. Systems which survey symptomatic toxoplasmosis in the general population are of least interest because it is impossible to distinguish congenital from acquired toxoplasmosis without data on the serological status during pregnancy or at birth [9]. Furthermore, the vast majority of acquired toxoplasmosis infections in healthy individuals are benign and the proportion of asymptomatic cases is estimated to be 70% [10–13]. Differences in the structure of these four specific surveillance systems may be responsible for

differences in their usefulness. We consider the surveillance system in Denmark to be simpler than those in Italy, Germany, and France. Centralised analysis like in Denmark and France also increases the acceptability as the system relies on professionals specifically dedicated to the system, contrary to the systems in Italy and Germany where the tasks are divided between health professionals and laboratories. The Danish surveillance system could also be considered the most flexible, because of its centralised approach, which allows for changes to be implemented in only one place, should they become necessary.

In Denmark, the surveillance system was linked to a nationwide systematic neonatal screening [14]. The sensitivity and the representativeness of this system could thus be considered higher than in Germany where the surveillance system is suffering from an underestimation of the number of congenital toxoplasmosis cases. Data on the number of congenital toxoplasmosis cases detected by the two surveillance systems were available for 2001 and 2002. In Germany, 38 cases were reported 2001 and 18 in 2002 (<http://www3.rki.de/SurvStat/QueryForm.aspx>) among a population of 82 million inhabitants, compared to 19 cases in 2001 and 13 in 2002 in Denmark (5.4 million inhabitants) [14]. According to these data, the estimated frequency of congenital toxoplasmosis is ten-fold lower in Germany than in Denmark. Based on what is known about the geographical variation of the burden of congenital toxoplasmosis, this is unlikely.

In Italy, congenital toxoplasmosis cases are declared by social workers, paediatricians and neonatologists. It is well known that passive reporting by physicians only captures a fraction of cases, most often only the most serious ones [15,16].

Overall, we consider the epidemiological surveillance system that was implemented in Denmark be the most useful. However, it was discontinued in July 2007.

A European survey was conducted within the EUROTOXO initiative to describe the national public health policies and routine programmes to prevent congenital Toxoplasmosis [17]. One of the fundamental criteria to evaluate the efficiency of such programmes is the frequency of the disease in question. Some countries did not define congenital toxoplasmosis as a public health issue and consequently have not implemented a prevention programme or surveillance system.

Several countries that do not have a congenital toxoplasmosis prevention policy have nevertheless defined congenital toxoplasmosis as a public health issue and implemented a surveillance system. But of these countries only Germany has implemented a system specifically dedicated to congenital toxoplasmosis.

Austria, Denmark, France, Italy, Lithuania and Slovenia have defined congenital toxoplasmosis as a public health issue and implemented a national systematic prevention programme [17]. Among these six countries, Denmark and France are the only countries where a specific and exhaustive surveillance system of congenital toxoplasmosis was implemented. However, screening and surveillance in Denmark were stopped in July 2007 and in France has only existed since May 2007, 29 years after the implementation of the national screening programme.

In the absence of a dedicated surveillance system, data on the burden of a disease can be obtained only through ad hoc epidemiological surveys. A systematic review of the published data on the burden of congenital toxoplasmosis was conducted by the EUROTOXO study group in 2005 [18]. The main results of this review were the following: Firstly, the prevalence of toxoplasmosis among pregnant women (the reservoir of congenital toxoplasmosis) decreases over the years, as previously reported. Due to limited available data, other epidemiological parameters such as incidence of seroconversion in susceptible pregnant women or incidence of complications among congenitally infected children cannot be analysed in detail. Such accurate data on the trends of diseases can only be obtained through continuous data collection such as surveillance systems.

Secondly, published data on the burden of congenital toxoplasmosis in Europe are limited, in terms of both quantity and quality. In fact, the vast majority of surveys evaluated by the group were not representative, in particular with respect to rare events such as the incidence of complications among congenitally infected children. For these estimates to be sufficiently precise, children were recruited in specialised centres. Such representative estimates could be improved by systematic data collection, for example as part of a surveillance system.

Nevertheless, periodic snapshot surveys based on consistent reporting definitions can also be an effective way of determining the burden of congenital toxoplasmosis. This is the approach used in the United Kingdom for symptomatic toxoplasmosis in children through the British Paediatric Surveillance Unit and the British Ophthalmic Surveillance Unit [9].

Few countries in Europe have implemented specific surveillance systems in accordance with their prevention policies regarding congenital toxoplasmosis. The epidemiological surveillance of congenital toxoplasmosis needs to be improved in order to determine the true burden of disease and assess the need for and effectiveness of existing prevention programmes.

## **Acknowledgements:**

The authors would like to thank Alain Moren (EPIET training programme coordinator) and H el ene Therre (Eurosurveillance Monthly editor) for their help in contacting the European correspondents.

EUROTOXO is a joint initiative of the Institute of Child Health (London, UK), the Staten Serum Institute (Copenhagen, Denmark), and the Institute of Public Health, Epidemiology and Development (Bordeaux, France). The EUROTOXO Group was financed by the European Commission (Contract No. QL4G-CT-2002-30262) and worked from 2002–2005.

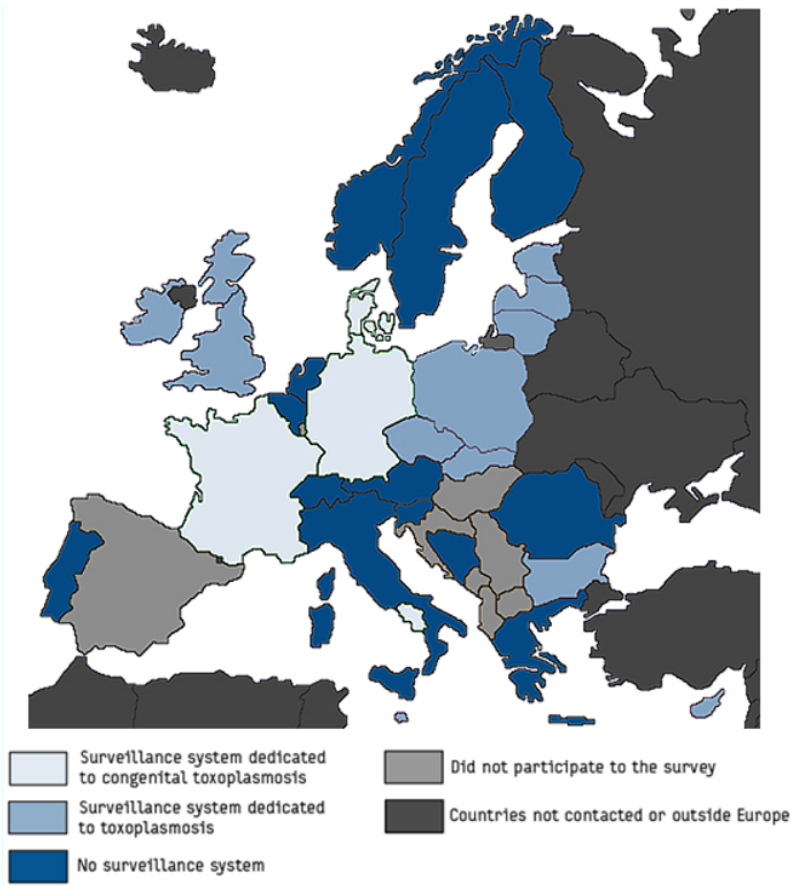
The EUROTOXO Group was composed as follows (by alphabetical order, except for chairs): Steering committee: Roger Salamon, Institute of Public Health, Epidemiology and Development (ISPED), Bordeaux, France (Chair): Leiv S. Bakketeig (University of Southern Denmark, Denmark), G erard Br art (INSERM U-149, Paris, France), Wilma Buffolano (Universita Federico II, Naples, Italy), Genevi ve Ch ene (INSERM U-593, Bordeaux, France), Birgitta Evengard (Karolinska Institute, Huddinge Hospital, Sweden), Ruth Gilbert (Institute of Child Health, London, UK), Michael Hayde (University Children's Hospital, Vienna, Austria), Eskild Petersen (Staten Serum Institute, Copenhagen, Denmark), Fran ois Peyron (H opital de la Croix-Rousse, Lyon, France), Aniki Rothova (Academisch Ziekenhuis Utrecht, Utrecht, The Netherlands), L. Rachid Salmi (ISPED, Bordeaux, France), Babill Stray-Pedersen (Institute of General Practice and Community Medicine, University of Oslo, Oslo, Norway), Philippe Thulliez (Institut de Pu riculture, Paris, France). Scientific secretariat: L. Rachid Salmi, ISPED, Bordeaux, France (Chair), Antoine B nard (INSERM U-593, Bordeaux, France), Ruth Gilbert (Institute of Child Health, Londres, UK), Val eriane Leroy (INSERM U-593, Bordeaux, France), Eskild Petersen (Staten Serum Institute, Copenhagen, Denmark), Rodolphe Thi baut (INSERM U-593 and U-875, Bordeaux, France). Scientific Secretariat collaborators: H el ene Bricout (INSERM U593, Bordeaux, France), Sabrina Di-Costanzo (INSERM U593, Bordeaux, France), Sandy Leproust (INSERM U593, Bordeaux, France). Panel 1: L. Rachid Salmi, ISPED, Bordeaux, France (Chair). Antoine B nard (France), Christine Binquet (France), Michel Cot (France), Catherine De Vigan (France), Marianne Forsgren (Sweden), Justus Garweg (Switzerland), Dorota Nowakowska (Poland), Stefania Salmaso (Italy), Alessandra Sensini (Italy), Rudiger Von Kries (Germany), Anton Van Loon (The Netherlands). Panel 2: Rodolphe Thi baut, INSERM U-593 and U-875, Bordeaux, France (Chair), Ole Andersen (Denmark), Alain Berrebi (France), Antoine Br zin (France), Bruno Charpiat (France), Francis Derouin (France), Uwe Gross (Germany), Fran ois Kieffer (France), Philippe Lepage (Belgium), Laurent Mandelbrot (France), Valeria Meroni (Italy), Maarten Postma (The Netherlands), Miles Stanford (UK). Panel 3: Val eriane Leroy, INSERM U-593, Bordeaux, France (Chair), Antony Ades (UK), Marie-Laure Dard  (France), Anne Eskild (Norway), Christos Hadjichristodoulou (Greece), Pal Jenum (Norway), Monique Kaminski (France), Babak Khoshnood (France), Jean-Fran ois Korobelnik (France), Bogumila Milewska-Bobula (Poland), Pierre-Alain Raeber (Switzerland), Christoph Rudin (Switzerland), Isabelle Villena (France). External experts: Am elie Daveluy, Annie Fourrier-R glat, Fran ois Goffinet, J r me Harambat, Fran oise Haramburu, Paul Perez, Martine Wallon. Secretariat, documentation and editorial support: Erica L. Gollub, Evelyne Mouillet, Karine Surlanne, Hooi-Kuan Tan.

## References:

1. Salmi LR , Mathoulin S , Perez P , Lawson-Ayayi S Screening and early detection in blood transfusion: when are they indicated?. *Transfus Clin Biol.* 1997; 4: (4) 417- 27
2. Wilson JMG , Jungner G Principles and practice of screening for disease. *Public Health Papers.* 34: World Health Organization; Geneva 1968;
3. Gilbert R , Dezateux C Newborn screening for congenital toxoplasmosis: feasible, but benefits are not established. *Arch Dis Child.* 2006; 91: (8) 629- 31
4. Centers for Disease Control and Prevention Guidelines for evaluating surveillance systems. *MMWR Morb Mortal Wkly Rep.* 1988; 37: (S-5) 1- 18
5. Churchill R , Teutsch S Principles and practice of public health surveillance. New York Oxford University Press; 2000;
6. SYROCOT (Systematic Review on Congenital Toxoplasmosis) study group Thi baut R , Leproust S , Ch ene G , Gilbert R Effectiveness of prenatal treatment for congenital toxoplasmosis: a meta-analysis of individual patients' data. *Lancet.* 2007; 369: (9556) 115- 22
7. Robert Koch Institute Infektionsepidemiologisches Jahrbuch meldepflichtiger Krankheiten.
8. European Union. Commission decision of 19 March 2002 laying down case for reporting communicable diseases to the Community network under Decision No 2119/98/EC of the European Parliament and of the Council. *OJEC L86/58, 3.4.2002. Official Journal of the European Communities.* 2002. Available from: [http://eur-lex.europa.eu/LexUriServ/site/en/oj/2002/l\\_086/l\\_08620020403en00440062.pdf](http://eur-lex.europa.eu/LexUriServ/site/en/oj/2002/l_086/l_08620020403en00440062.pdf)
9. Gilbert R , Tan HK , Cliffe S , Guy E , Stanford M Symptomatic toxoplasmosis in childhood in the UK. *Arch Dis Child.* 2006; 91: (6) 495- 8
10. Mawhorter SD , Effron D , Blinkhorn R , Spagnuolo PJ Cutaneous manifestations of toxoplasmosis. *Clin Infect Dis.* 1992; 14: (5) 1084- 8
11. Montoya JG , Liesenfeld O Toxoplasmosis. *Lancet.* 2004; 363: (9425) 1965- 76
12. McAuley J , Boyer KM , Patel D , Mets M , Swisher C , Roizen N Early and longitudinal evaluations of treated infants and children and untreated historical patients with congenital toxoplasmosis: the Chicago Collaborative Treatment Trial. *Clin Infect Dis.* 1994; 18: (1) 38- 72
13. Binquet C , Wallon M , Quantin C , Kodjikian L , Garweg J , Fleury J Prognostic factors for the long-term development of ocular lesions in 327 children with congenital toxoplasmosis. *Epidemiol Infect.* 2003; 131: (3) 1157- 68
14. Schmidt DR , Hogh B , Andersen O , Fuchs J , Fledelius H , Petersen E The national neonatal screening programme for congenital toxoplasmosis in Denmark: results from the initial four years, 1999–2002. *Arch Dis Child.* 2006; 91: (8) 661- 5
15. Vogt RL , LaRue D , Klaucke DN , Jillson DA Comparison of an active and passive surveillance system of primary care providers for hepatitis, measles, rubella, and salmonellosis in Vermont. *Am J Public Health.* 1983; 73: (7) 795- 7
16. Marier R The reporting of communicable diseases. *Am J Epidemiol.* 1977; 105: (6) 587- 90
17. Leroy V , Raeber PA , Petersen E , Salmi LR , Kaminski M , Villena I for the Eurotox Group (Panel 3) National public health policies and routines programs to prevent congenital Toxoplasmosis, Europe, 2005 [Unpublished report]. Bordeaux (France) The Eurotox Group; 2005;
18. B nard A , Salmi LR for Panel 1 of the Eurotox Group Systematic review on the burden of congenital toxoplasmosis in Europe [Unpublished report]. Bordeaux (France) The Eurotox Group; 2005;

**Figure**

Different surveillance systems for toxoplasmosis in Europe. Eurotoxox, 2007



**Table 1**

European correspondents contacted to participate in the survey on the epidemiological surveillance of toxoplasmosis

	Countries that participated in the survey
Austria	Reinhild Strauss, Federal Ministry for Health, Family and Youth, General Directorate of Public Health, Vienna
Belgium	Germaine Hanquet, Scientific Institute for Public Health, Unit of Epidemiology, Ministry of Social Affairs, Public Health and Environment, Brussels
Bosnia and Herzegovina	and Semra Cavaljuga, Institute for Epidemiology and Biostatistics, School of Medicine, University of Sarajevo, Sarajevo
Bulgaria	Mira Kojuharova, National Centre of Infectious and Parasitic Diseases, Department of Epidemiology, Sofia
Cyprus	Olga Kalakouta, on behalf of Dr. Chrystalla Hadjianastassiou, Chief medical Officer, Medical and Public Health Services, Ministry of Health, Nicosia
Czech Republic	Petr Kodym, National Reference Laboratory for Toxoplasmosis, National Institute of Public Health, Prague
Denmark	Henrik Vedel Nielsen, Unit for Mycology and Parasitology, Statens Serum Institut, Copenhagen
England and Wales	Robert Smith, Public Health Laboratory, Service of Communicable Disease, Surveillance Centre Wales, Cardiff
Estonia	Kuulo Kutsar, Department of Communicable Diseases, Health Protectorate Inspectorate, Tallinn
Finland	Maija Lappalainnen, Department of Virology, Hospital District of Helsinki and Uusimaa, Helsinki
France	Isabelle Villena, National Reference centre for toxoplasmosis, Reims; Véronique Goulet, Department of Infectious Diseases, Institut de Veille Sanitaire, Saint-Maurice
Germany	Katharina Alpers, Department for Infectious Disease Epidemiology, Gastrointestinal Infections, Zoonoses and Tropical Infections, Robert Koch Institute, Berlin
Greece	Yanis Tselentis, Laboratory of Clinical Bacteriology, Parasitology, Zoonoses and Geographical Medicine, University of Crete, Faculty of Medicine, Heraklion
Ireland	Darina O'Flanagan, HSE-Health Protection Surveillance Centre, Dublin
Italy	Wilma Buffolano, Perinatal Infection Unit, Department of Paediatrics, Federico II University of Naples; Maria Grazia Pompa, Communicable Disease Unit, DG Health Prevention, Ministry of Health, Rome
Latvia	Irina Lucenko, Division of Epidemiology of Infectious Diseases, State Public Health Agency, Riga
Lithuania	Bronius Morkunas, Centre for Communicable Disease Prevention and Control, Vilnius
Malta	Tanya Melillo Fenech, Disease Surveillance Unit, Department of Public Health, Ministry of Health, Msida
Netherlands	Laetitia M. Kortbeek, Diagnostic Laboratory for Infectious Diseases and Perinatal Screening (LIS) National Institute of Public Health and the Environment, Bilthoven
Norway	Hans Blystad, Department of Infectious Disease Epidemiology, Norwegian Institute of Public Health, Oslo
Poland	Malgorzata Sadkowska-Todys, Laboratory of Zoonoses, Department of Epidemiology, National Institute of Hygiene, Warsaw
Portugal	Judite Catarino, General Health Directorate, Lisbon
Romania	Adriana Pistol, General Department of Public Health, Service of Prevention and Control of Communicable Diseases, Ministry of Health and Family, Bucharest
Scotland	Lynda Browning, Zoonoses Section, Health Protection Scotland, Glasgow
Slovakia	Maria Avdicova, Department of Epidemiology, Regional Institute of Public Health, Baska Bystrica
Slovenia	Jernej Logar, Institute of Microbiology, Medical Faculty, Ljubljana
Sweden	Johan Lindh, Department of Parasitology, Mycology and Water, Swedish Institute for Infectious Disease Control, Solna
Switzerland	Karim Boubaker, Infectious Diseases Section, Division of Communicable Diseases, Swiss Federal Office of Public Health, Public Health Direction, Bern
	Countries that were asked but did not participate in the survey
Albania	Eduard Kakarriqi, Department of Epidemiology, Institute of Public Health, Rruga
Croatia	Ira Gjenero-Margan, Croatian Public Health Institute, Department of Epidemiology of Infectious Diseases, Zagreb
Hungary	Márta Melles, 'Johan Béla' National Centre for Epidemiology, Budapest
Luxembourg	Robert Hemmer, National service of Infectious Diseases, Centre Hospitalier de Luxembourg, Luxembourg
Macedonia	Kristin Vasilevska, Medical Faculty Skopje, University 'Sv. Kiril i Metodij' Institute of Epidemiology, Biostatistics and Medical Informatics, Skopje
Spain	Luiza Sanchez Serrano, Sección de Sistema de Información Microbiológica, Vigilancia de Salud Pública, National Centre of Edpidemiology, Hospital Carlos III, Madrid
Serbia-Montenegro	Danica Masanovic, Sanitary Inspection of the Ministry of Health of Montenegro, Podgorica



**Table 2**

Characteristics of European epidemiological surveillance programmes for toxoplasmosis. Eurotoxo, 2007

Country	Year started	Population under surveillance	Case definition	Reporting sources	Frequency of analysis	Frequency of surveillance reports	Surveillance report distribution
<b>Surveillance systems specifically dedicated to congenital toxoplasmosis</b>							
<b>Denmark</b>	1999	Live newborns	Detection of IgM on blood filter sample confirmed by IgA, IgM and IgG profile in newborn and mother	Statens Serum Institut	NA	Annually	Healthcare authorities
<b>France</b>	2007	Foetuses, newborns and infants	PCR, mouse inoculation, cell culture or immunocytochemistry on body tissues or fluids; detection of specific IgM or IgA antibodies; neosynthesis of specific IgG, IgM or IgA for antenatal or postnatal diagnosis	Laboratories qualified for antenatal or postnatal diagnosis	NA	NA	Healthcare authorities
<b>Germany</b>	NA	Live newborns and infants	Detection of <i>Toxoplasma gondii</i> in body tissues or fluids; detection of specific IgM or IgA antibodies; persistently stable specific IgG titres or a single elevated specific IgG titre	Laboratories	Continuous	Quarterly and annually	Free access on the Website of the Robert Koch Institute
<b>Italy *</b>	1997	Live newborns	Persistence of IgG until one year of age	Social workers Paediatricians Neonatalogists	Annually	Annually	National Health Institute, physicians, Regional Public Health Surveillance on Infectious Diseases
<b>Surveillance systems dedicated to toxoplasmosis (congenital or not)</b>							
<b>Bulgaria</b>	NA	All	EU (notifiable disease)	Physicians Laboratories Epidemiologists	Annually	Monthly and annually	Ministry of Health National Centre of Health Information National Centre of Infectious and Parasitic Diseases
<b>Cyprus</b>	2004	All	EU (notifiable disease)	All registered medical practitioners	Weekly	Twice a year	Physicians
<b>Czech Republic</b>	1970	All	EU (notifiable disease)	Epidemiologists	Monthly	Monthly and annually	Epidemiologists, Physicians Laboratories
<b>England and Wales</b>	1975	All	EU (notifiable disease)	Toxoplasma Reference Unit Swansea	Monthly	Quarterly and annually	Electronic distribution (and annually <a href="http://www.hpa.org.uk/hpr">http://www.hpa.org.uk/hpr</a> )
<b>Estonia</b>	1997	All	EU (notifiable disease)	General practitioners Laboratories	Monthly	Monthly and annually	Health protection Inspectorate and Ministry of Social Affairs
<b>Ireland</b>	2004	All	EU (notifiable disease)	All registered medical practitioners Laboratories	Weekly and annually	Weekly and annually	Physicians, Public health departments and population (Website)
<b>Latvia</b>	1995	All	EU (notifiable disease)	Physicians Epidemiologists	Annually	Monthly and annually	Ministry of Health, Physicians
<b>Lithuania</b>	1992	All	EU (notifiable disease)	All registered medical practitioners	Monthly and annually	Monthly and annually	Territorial healthcare institutions, Ministry of Health, European surveillance networks, WHO
<b>Malta</b>	2004	All	EU (notifiable disease)	Physicians Laboratories	Continuous	Weekly, monthly and annually	Physicians, Ministry of Health, WHO

<b>Poland</b>	1966 <sup>♦</sup>	All	EU (notifiable disease)	Physicians	Occasionally and annually	Quarterly Public administrations, research institutions and sanitary stations
<b>Scotland</b>	1988	All	EU (notifiable disease)	Laboratories	Continuous	Available on Free access for all (on demand) Website
<b>Slovakia</b>	1975	All	EU (notifiable disease)	Physicians	Monthly and annually	Monthly and Physicians, Ministry of Health annually

NA: Not available; WHO: World Health Organization;  
as defined by the European Union<sup>4</sup>

\* Regional surveillance system (Campania country)

Distinction between acquired and congenital toxoplasmosis since 1999

<sup>♦</sup> Distinction between acquired and congenital toxoplasmosis since 1997.