

CEMC-Auvergne - Registry of Congenital Malformations in Auvergne (Certified Registry 2012-2015)

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General

Identification

Detailed name Registry of Congenital Malformations in Auvergne (Certified Registry 2012-2015)

Sign or acronym CEMC-Auvergne

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation 1387396

General Aspects

Medical area Disability/handicap
Pediatrics
Rare diseases

Health determinants Addictions
Genetic
Iatrogenic
Lifestyle and behavior
Medicine

Keywords prenatal diagnosis

Scientific investigator(s) (Contact)

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Organization CEMC-Auvergne

Collaborations	
Funding	
Funding status	Mixed
Details	Regional Health Agency - ARS - French Institute for Public Health Surveillance - InVS -French National Institute of Health and Medical Research - INSERM, private donations.
Governance of the database	
Sponsor(s) or organisation(s) responsible	CHU de Clermont-Ferrand
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Morbidity registers
Additional information regarding sample selection.	Selection of subjects having the required inclusion criteria:- Public and private maternity wards in the Auvergne region- Multidisciplinary center for prenatal diagnosis- Medical and surgical pediatric departments- Cardiology departments- Maxillo-facial surgery- Pediatric urology- Public and private cytogenetics laboratories- Anatomopathology laboratory- Mother and child protection centers (PMIs)- Medical information departments (DIMs)
Database objective	
Main objective	The registry's primary objective is to conduct epidemiological surveillance of congenital malformations in Auvergne, with a role of sounding the alert when any environmental teratogenic agents become evident. With the development of prevention measures - particularly in the field of prenatal diagnosis - this surveillance-alert mission has gradually been extended to include an essential role in evaluating the impact of public health policies with regards to perinatal care on the population. The registry also helps to improve knowledge in the field of malformations by collaborating on studies with specialist clinical

departments (particularly genetic and foetal medicine departments) and by being involved in the Multidisciplinary Center for Prenatal Diagnosis in Clermont-Ferrand (the registry's data provides food for thought from obstetricians on the meaning and action to take when certain malformations are observed during prenatal diagnosis). Lastly, the registry's data informs healthcare stakeholders and the public authorities about congenital malformations and chromosomal abnormalities.

Inclusion criteria

All malformed children are listed, whether stillborn or a live birth after a pregnancy of at least 22 weeks of amenorrhoea, or in the event of therapeutic abortion due to foetal malformation, regardless of term. For children who are born alive, the malformation diagnosis must have been made before the end of their first year of life (infants may be recorded up to 12 months after their birth). All types of malformation are included, whether these concern isolated or polymalformative syndromes, whether or not identified, with normal or abnormal karyotype. The only types to be excluded are inborn metabolic errors and minor malformations or deformations (snapping hips without real dislocation, deformed feet, small superficial naevi or angiomas less than 4cm², inguinal hernia, umbilical hernia not requiring surgery, isolated persistent ductus arteriosus in premature infants weighing less than 2.5kg, hypertrophic pyloric stenosis, single umbilical artery, uni- or bilateral ectopic testis).

Population type

Age	Newborns (birth to 28 days) Infant (28 days to 2 years)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	Regional
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French regions covered by the database	Auvergne Rhône-Alpes
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Detail of the geography area	Auvergne
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Data collection

Dates

Date of first collection (YYYY or MM/YYYY) 1983

Size of the database

Size of the database (number of individuals) [10 000-20 000[individuals

Details of the number of individuals 10 145 patients (1983-2011)

Data

Database activity Current data collection

Type of data collected
Clinical data
Paraclinical data
Biological data
Administrative data

Clinical data (detail) Direct physical measures

Paraclinical data (detail) MRI, scan, radiographies, ultrasounds (prenatal, foetal echocardiography, in post-natal stages: urinary tracts, cardio, etc.).

Biological data (detail) cytogenetic examinations, maternal serum markers.

Administrative data (detail) Identification data.

Presence of a biobank No

Health parameters studied
Health event/morbidity
Health event/mortality

Procedures

Data collection method primarily active +/- passive.

Classifications used ICD 10, ATC.

Participant monitoring Yes

Details on monitoring of participants
Vital status: The vital status is reported systematically when each case is being registered. The following are distinguished: - children who are alive when the report is made to the registry - therapeutic abortions before 22 weeks of amenorrhoea- therapeutic abortions after 22 weeks of amenorrhoea- fetal deaths in utero- infants who

have died (in the first 8 days of life, between 8 and 28 days or after 28 days)

Links to administrative sources	No
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Promotion and access

Promotion

Link to the document	http://tinyurl.com/PUBMED-CEMC
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Description	Liste des publications dans Pubmed
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Link to the document	http://tinyurl.com/HAL-CEMC
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Description	Liste des publications dans HAL
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Access

Terms of data access (charter for data provision, format of data, availability delay)	By request to the scientist in charge.
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Access to aggregated data	Access on specific project only
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Access to individual data	Access on specific project only
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