

CONGENITAL HYPOTHYROIDISM - Newborn screening for congenital hypothyroidism - Development of young adults treated for congenital hypothyroidism since neonatal period: health, fertility, pregnancy outcomes and children's health

Head :Leger Juliane, CENTRE DE RÉFÉRENCE MALADIES ENDOCRINIENNES RARES DE LA CROISSANCE
HÔPITAL ROBERT DEBRÉ AP-HP

Last update : 09/05/2017 | Version : 3 | ID : 5664

General

Identification

Detailed name	Newborn screening for congenital hypothyroidism - Development of young adults treated for congenital hypothyroidism since neonatal period: health, fertility, pregnancy outcomes and children's health
---------------	--

Sign or acronym	CONGENITAL HYPOTHYROIDISM
-----------------	---------------------------

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	Accord CNIL n°906178 (31/07/2006)
--	-----------------------------------

General Aspects

Medical area	Endocrinology and metabolism
--------------	------------------------------

Health determinants	Genetic
---------------------	---------

Keywords	vital status, socio-professional insertion, fertility, children, Health episodes, treatment, quality of life, morbidity, pregnancy, environment
----------	---

Scientific investigator(s) (Contact)

Name of the director	Leger
----------------------	-------

Surname	Juliane
---------	---------

Address	48 boulevard Sérurier 75935 Paris Cedex 19
---------	---

Phone	+ 33 (0)1 40 03 23 54
-------	-----------------------

Email	juliane.leger@rdb.aphp.fr
-------	---------------------------

Unit	CENTRE DE RÉFÉRENCE MALADIES
------	------------------------------

ENDOCRINIENNES RARES DE LA CROISSANCE
HÔPITAL ROBERT DEBRÉ AP-HP

Organization	AP-HP
Collaborations	
Funding	
Funding status	Public
Details	PHRC 2005, Fondation WYETH
Governance of the database	
Sponsor(s) or organisation(s) responsible	Hôpital Robert Debré
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Longitudinal study (except cohorts)
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Prospective Inclusion cut-off date: 01/12/1988
Database objective	
Main objective	General objective: To analyse the development into adulthood for individuals treated for congenital hypothyroidism (CH) since neonatal period, thanks to routine screening established in France since 1978. Secondary objectives: - to assess health at adult age, quality of life and socio-professional insertion and fertility as well as children's health i.e. children of subjects treated for CH. To research thyroid developmental anomaly and general development during first year of life. - to identify

sub-groups of candidate patients according to phenotype for genetic study. The analysis of pregnancy progress and outcome as well as thyroxine requirements during pregnancy is included in the study of women treated for CH. - To improve treatment strategies (children, adolescents, adults, pregnant women).

Inclusion criteria Hypothyroidism detected during neonatal period between 1978 and 1988

Population type

Age Newborns (birth to 28 days)
 Infant (28 days to 2 years)
 Early childhood (2 to 5 years)
 Childhood (6 to 13 years)
 Adolescence (13 to 18 years)
 Adulthood (19 to 24 years)
 Adulthood (25 to 44 years)
 Adulthood (45 to 64 years)
 Elderly (65 to 79 years)
 Great age (80 years and more)

Population covered Sick population

Gender Woman

Geography area National

Detail of the geography area Multicentric cohort throughout metropolitan France (20 centres):

Data collection

Dates

Date of first collection (YYYY or MM/YYYY) 01/1978

Size of the database

Size of the database (number of individuals) [1000-10 000[individuals

Details of the number of individuals 1804

Data

Database activity Data collection completed

Type of data collected Declarative data

Declarative data (detail)	Paper self-questionnaire
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Quality of life/health perception
Procedures	
Data collection method	Self-administered questionnaire: manual input
Participant monitoring	Yes
Details on monitoring of participants	(Indefinite duration)
Links to administrative sources	Yes
Linked administrative sources (detail)	RNIAM (JO 11 JUNE 20007) French association for the detection and prevention of childhood disabilities
Promotion and access	
Promotion	
Link to the document	http://tinyurl.com/Pubmed-HYPOTHYROIDIE
Description	List of publications in Pubmed
Access	
Terms of data access (charter for data provision, format of data, availability delay)	To be decided if data may be used by academic teams Data may not be used by industrial teams
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only