

ELFE - French longitudinal study from childhood

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Général

Identification

Nom détaillé French longitudinal study from childhood

Sigle ou acronyme ELFE

Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.) CNIL n°910504 - CNIS n°2011X716AU - CPP n°IDF IX-11-024 (13/05/2011)

Thématiques générales

Domaine médical Biology
Dermatology, venereology
Disability/handicap
Endocrinology and metabolism
Gynecology/ obstetrics
Infectious diseases
Ophthalmology
Otolaryngology or ENT
Pediatrics
Pneumology
Psychology and psychiatry
Study of allergies
Traumatology

Etude en lien avec la Covid-19

Yes

Déterminants de santé

Geography
Healthcare system and access to health care services
Lifestyle and behavior
Nutrition
Occupation
Pollution
Social and psychosocial factors

Mots-clés

family, pregnancy, growth, development, pollution, environment, nutrition, living environment, schooling, socialisation, children

Responsable(s) scientifique(s)

Nom du responsable

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Laboratoire	?Elfe? INED-INSERM joint unit
Organisme	University of Picardie
Collaborations	
Participation à des projets, des réseaux, des consortiums	Yes
Précisions	European Lifecycle project
Financements	
Financements	Public
Précisions	Ministry for Higher Education and Research, the Ministry for Labour, Employment and Health, and the Ministry for Ecology Sustainable Development, Transport and Housing, + RE-CO-NAI is the recipient of an ANR "Investing in the Future" grant
Gouvernance de la base de données	
Organisation(s) responsable(s) ou promoteur	French National Institute for Demographic Studies (INED)
Statut de l'organisation	Secteur Public

Organisation(s) responsable(s) ou promoteur	French National Institute for Health and Medical Research (INSERM)
Statut de l'organisation	Secteur Public
Existence de comités scientifique ou de pilotage	Yes
Contact(s) supplémentaire(s)	
Caractéristiques	
Type de base de données	
Type de base de données	Study databases
Base de données issues d'enquêtes, précisions	Cohort study
Origine du recrutement des participants	
Le recrutement dans la base de données s'effectue dans le cadre d'une étude interventionnelle	No
Informations complémentaires concernant la constitution de l'échantillon	random sample of maternity units
Objectif de la base de données	
Objectif principal	<ul style="list-style-type: none"> - Track changes in the child's living conditions (family, social environment, etc.) and measure their impact on his or her physical and mental development. - Study the interactions between these living conditions and the child's school career - Measure the impact of events during pregnancy and the child's health at birth on the latter's subsequent development - Observe eating habits and their effects on growth (overweight, obesity) - Assess the child's exposure to chemical, physical or environmental pollutants - Measure the incidence and prevalence of childhood pathologies at different stages in growth - Study associations between pathologies and exposure to environmental pollutants <p>Elfe is one of the two cohorts making up the RECONAI research platform, the other being Epipage</p> <p>2. The general aim of this platform is to set up an</p>

infrastructure to support the collection, highly secure storage and distribution of data on pregnancy, birth and children. The platform will use the information yielded by the child cohorts, both followed from birth, to address key issues in the areas of children's health, development and socialization from an overarching, multidisciplinary perspective. It will also ensure that the cohorts are given a high profile in the world of academic research (French and international), as well as among bodies, organizations and manufacturers with an interest in children. This will ensure optimum use of the data collected and promote their dissemination.

Critères d'inclusion

- Children born after 33 weeks of pregnancy
- single or twin pregnancy
- in mainland France
- mother aged 18 years and over

Type de population

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)

Population concernée

General population

Pathologie

Sexe

Male
Woman

Champ géographique

National

Détail du champ géographique

Mainland France

Collecte

Dates

Année du premier recueil

2011

Taille de la base de données

Taille de la base de données (en nombre d'individus)

[10 000-20 000] individuals

Détail du nombre d'individus

18300

Données

Activité de la base	Current data collection
Type de données recueillies	Clinical data Declarative data Paraclinical data Biological data Administrative data
Données cliniques, précisions	Direct physical measures Medical registration
Détail des données cliniques recueillies	> Data available in the medical records at the time of birth: neonatal characteristics (weight, height, gestational age, congenital anomalies, circumstances of birth, infections, etc.) > Medical examinations: vaccination, hearing, eyesight, weight/height, asthma, allergies, genital organs and neuropsychomotor tests at different ages
Données déclaratives, précisions	Paper self-questionnaire Internet self-questionnaire Face to face interview Phone interview
Détail des données déclaratives recueillies	Exposure: medication, pollution, occupational, tobacco use, alcohol, diet, chemical and physical agents, social status. > Results: congenital anomalies, infections, growth/obesity, neurological development, asthma/allergies, pubescent development, sleep, mental health > Use of healthcare resources
Données paracliniques, précisions	pregnancy monitoring tests, medical imaging
Données biologiques, précisions	At birth, urine, venous blood and maternal hair post-partum: Breast milk, meconium, stools (at D3); urine, hair, infant stools at 3.5 years
Données administratives, précisions	pairing with SNIIR-AM data
Existence d'une biothèque	Yes
Contenu de la biothèque	Whole blood Serum Plasma Cord blood Fluids (saliva, urine, amniotic fluid, ?) Tissues DNA DNAc/RNAm

	Others
Détail des éléments conservés	plasma, serum, whole blood, red blood cells, buffy coat, urine, hair, milk, stools and meconium, RNA (sub-sample)
Paramètres de santé étudiés	Health event/morbidity Health care consumption and services Quality of life/health perception
Modalités	
Mode de recueil des données	- Self-administered questionnaire on diet and exposure during pregnancy completed by the mother in the maternity unit.- notes from obstetrics record- Face-to-face meeting, at home- Telephone surveys- Self-administered questionnaires on infant diet between
Nomenclatures employées	ICD-10
Suivi des participants	Yes
Modalités de suivi des participants	Monitoring by contact with the participant (mail, e-mail, telephone etc.) Monitoring by crossing with a medical-administrative database
Détail du suivi	annual follow-up for up to 3 years, then every 2-3 years
Appariement avec des sources administratives	Yes
Sources administratives appariées, précisions	SNIIR-AM database
Valorisation et accès	
Valorisation et accès	
Lien vers le document	http://tinyurl.com/HAL-ELFE
Description	List of publications in HAL
Lien vers le document	http://tinyurl.com/Pubmed-ELFE
Description	List of publications in Pubmed

Lien vers le document

[Elfe.pdf](#)

Accès

Site internet dédié

<https://www.elfe-france.fr/>

Existence d'un document qui répertorie les variables et les modalités de codage

Yes

Charte d'accès aux données (convention de mise à disposition, format de données et délais de mise à disposition)

Data access platform: access via the website: www.elfe-france.fr/acces-donnees or direct access: https://pandora.vjf.inserm.fr/public/

The proposals should be submitted to an ELFE data access committee which will determine the relevance of the project and the suitability of the data requested.

Accès aux données agrégées

Access on specific project only

Accès aux données individuelles

Access on specific project only