

"AZF" Santé Cohort - Cohort of Toulouse Conurbation Workers

Head :Buisson Catherine, DÉPARTEMENT SANTÉ TRAVAIL (DST)

Diène Eloi, Département santé travail Institut de veille sanitaire 94415 Saint Maurice Cedex

Last update : 12/01/2019 | Version : 1 | ID : 60005

General

Identification

Detailed name	Cohort of Toulouse Conurbation Workers
Sign or acronym	"AZF" Santé Cohort
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CCTIRS (number 03.143), CNIL (number 904129).

General Aspects

Medical area	Otolaryngology or ENT Psychology and psychiatry
Study in connection with Covid-19	No
Health determinants	Medicine Occupation Social and psychosocial factors
Keywords	Major health events observed: symptoms of post-traumatic stress, depressive symptoms, hearing disorders. Main cohort topics: mental health, auditory health, working life.

Scientific investigator(s) (Contact)

Name of the director	Buisson
Surname	Catherine
Address	94410 saint Maurice
Phone	+33 (0)1 41 79 67 00
Email	c.buisson@invs.sante.fr
Unit	DÉPARTEMENT SANTÉ TRAVAIL (DST)

Organization	Institut de Veille
Name of the director	Diène
Surname	Eloi
Address	ARS Midi Pyrénées, 10 chemin du raisin, 31050 Toulouse Cedex
Phone	+33 (0)5 34 30 26 81
Email	eloi.diene@ars.sante.fr
Unit	Département santé travail Institut de veille sanitaire 94415 Saint Maurice Cedex
Organization	InVS
Collaborations	
Others	Other related cohorts: COSET: the AZF cohort is a procedure test for COSET and CONSTANCES cohorts.
Funding	
Funding status	Public
Details	Partnership agreement between InVS, CPAM Toulouse (CES) and INSERM.
Governance of the database	
Sponsor(s) or organisation(s) responsible	Institut de veille sanitaire
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Additional information regarding sample selection.	Inclusion method: Prospective. Other bodies active in creating this cohort: - CPAM-TOULOUSE health examination clinics - INSERM U1018 CESP. Closing date for inclusion: 12/2009. All volunteers from the first cross-sectional study were included in the cohort.

Database objective

Main objective

General objectives:

- To investigate the medium-term health effects from the explosion at the AZF plant that occurred in September 2001 in the population of workers;
- To test access procedures for medical-administrative data (ERASMUS) for monitoring large prospective cohorts.

Inclusion criteria

Working within the Toulouse conurbation and agreeing to participate in the cohort during the cross-sectional study, carried out between September 2002 and April 2003.

Population type

Age

Adulthood (19 to 24 years)
Adulthood (25 to 44 years)
Adulthood (45 to 64 years)
Elderly (65 to 79 years)

Population covered

General population

Gender

Male
Woman

Geography area

Local

Detail of the geography area

French study. Geographical area covered: workers in the Toulouse conurbation.

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)

05/2003

Date of last collection (YYYY or MM/YYYY)

12/2009

Size of the database

Size of the database (number of individuals)

[1000-10 000[individuals

Details of the number of individuals

3,000 participants.

Data

Database activity	Data collection completed
Type of data collected	Clinical data Declarative data Paraclinical data Biological data Administrative data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Clinical and paraclinical examination: at baseline and final cohort assessment. Information collected is that generated during periodic examinations conducted in the primary Health Insurance Fund examination clinics. Psychiatric diagnosis by MINI.
Declarative data (detail)	Paper self-questionnaire Face to face interview
Details of collected declarative data	Self-administered questionnaire on occupation and health: at baseline and during annual follow-up. Self-administered questionnaire time span: 1 (year). Information collected by self-administered questionnaire: *health events and occupational events *mental health scales - psychological unease (GHQ) - depressive symptoms (CES-D) - symptoms of post-traumatic stress (IES-R). Psychiatric diagnosis by MINI - questionnaire is conducted face-to-face in a cohort sub-sample.
Paraclinical data (detail)	Radiology, pulmonary function tests, audiograms.
Biological data (detail)	Biology, biochemistry.
Administrative data (detail)	ERASMUS database.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health care consumption and services Quality of life/health perception
Care consumption (detail)	Medical/paramedical consultation Medicines consumption

Procedures

Data collection method	Annual collection.
Quality procedure(s) used	Self-administered questionnaire: optical input from a paper questionnaire (automated data reading) from second year of follow-up. Missing data

management: systematic search of individuals' addresses where post was returned undelivered. Subject reminder for follow-up visits? Yes. Patients are informed about the use of their data.

Participant monitoring Yes

Details on monitoring of participants Follow-up duration: 5 years.

Links to administrative sources Yes

Linked administrative sources (detail) Database(s) used: ERASMUS.

Promotion and access

Promotion

Link to the document <http://www.ncbi.nlm.nih.gov/pubmed/19252759>

Link to the document <http://www.ncbi.nlm.nih.gov/pubmed/22020864>

Link to the document <http://opac.invs.sante.fr/index.php?lvl>

Access

Terms of data access (charter for data provision, format of data, availability delay) - Can data be used by academic teams? Yes: access subject to written request according to InVS data access procedure.