

COPANFLU - Cohort of 1,000 Households for H1N1 Pandemic Follow-Up

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General

Identification

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|--|---|
| Detailed name | Cohort of 1,000 Households for H1N1 Pandemic Follow-Up |
| Sign or acronym | COPANFLU |
| CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation | CPP (IDF 1) : Réf. 09-12074 (08/09/2009); Afssaps : Réf. 2009-B90987-60 (10/09/2009); CNI: MR001 - projet qualifié en RBM |

General Aspects

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| Medical area | Infectious diseases |
| Health determinants | Intoxication Social and psychosocial factors |
| Keywords | influenza A (H1N1), infectious respiratory syndromes, Health episodes, infections, complications, infection |

Scientific investigator(s) (Contact)

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| Unit | ECOLE DES HAUTES ETUDES EN SANTÉ PUBLIQUE |
| Organization | EHESP |

Collaborations

Participation in projects, networks and consortia Yes

Funding

Funding status Public

Details EHESP, IRD

Governance of the database

Sponsor(s) or organisation(s) responsible ECOLE DES HAUTES ETUDES EN SANTÉ PUBLIQUE - EHESP

Organisation status Public

Additional contact

Main features

Type of database

Type of database Study databases

Study databases (details) Cohort study

Database recruitment is carried out by an intermediary A population file

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Prospective Other bodies active in creating this cohort: INSEE Inclusion cut-off date: 01/10/2009

Database objective

Main objective

General objective: To identify individual epidemiological, environmental, immunological, social, genetic and virological risk determinants of infection by influenza A (H1N1) swine-like-(SWL) virus
Secondary Objectives: - to describe the impact on public health in terms of morbidity, complications and health care resources for A (H1N1)-SWL infection; - to characterise the ? natural? history of the infection and its variability, to determine how it is transmitted; - to study the individual and collective determinants of the variable clinical expressions of the infection; - to study behavioural changes induced by pandemic and assess the level of risk perception and its evolution over time; - to evaluate penetrance, application and effectiveness of control measures in place (treatment, vaccination, barriers); - to characterise the homeotypic and heterotypic immunity of participants, according to age ? evaluate herd immunity and to study innate and acquired cellular immunity subsequent to infection by this virus; - To characterise viral diversity and evolutionary mechanisms of virus A (H1N1)-SWL; to study the onset of drug resistance - to build and test anticipation models in real time on the impact of pandemic on the population and to simulate different control scenarios.

Inclusion criteria

Volunteer households in the metropolitan area whose members have all provided validated consent to participate in this cohort (Parental consent as well as children's is required, optional if they are over 7 years old and obligatory if they are over 13 years); covered by health insurance; can write and speak French and who have a land-line telephone.

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

General population

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| Gender | Male Woman |
| Geography area | International |
| Detail of the geography area | International Study involving: Bolivia, Laos, Mali, Cameroon (not limited) Geographical area covered: National (France protocol) |

Data collection

Dates

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|--|---------|
| Date of first collection (YYYY or MM/YYYY) | 08/2009 |
| Date of last collection (YYYY or MM/YYYY) | 12/2012 |

Size of the database

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

Details of the number of individuals 1451

Data

Database activity Current data collection

Type of data collected Declarative data
Biological data

Declarative data (detail) Paper self-questionnaire
Face to face interview
Phone interview

Biological data (detail) Type of samples taken: cells samples and and serum At baseline, after possible vaccination and twice a year

Presence of a biobank Yes

Contents of biobank Serum
DNA
DNAC/RNAm

Details of biobank content Serum bank, DNA bank

Health parameters studied Health event/morbidity
Health event/mortality

Procedures

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|------------------------|---|
| Data collection method | Self-administered questionnaire: input from a paper questionnaire with double data entry Interview: input from a paper questionnaire with double data entry Biological analysis: direct input |
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| Participant monitoring | Yes |
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| Details on monitoring of participants | Follow-up duration: 2 years |
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| Links to administrative sources | Yes |
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| Linked administrative sources (detail) | IRIS-2000 |
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Promotion and access

Promotion

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| Link to the document | http://www.hal.inserm.fr/COPANFLU |
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|-------------|-----------------------------|
| Description | List of publications in HAL |
|-------------|-----------------------------|

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|----------------------|---|
| Link to the document | http://www.ncbi.nlm.nih.gov/pubmed/?term=copanflu |
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| Description | List of publications in Pubmed |
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Access

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| Terms of data access (charter for data provision, format of data, availability delay) | According to IRESP Charter and INSERM/EHESP convention Contact Prof. Fabrice Carrat |
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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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