

SAM - Severe Drug Allergy Physiopathology and Characterisation of Novel Diagnostic and Therapeutic Targets

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General

Identification

Detailed name Severe Drug Allergy Physiopathology and Characterisation of Novel Diagnostic and Therapeutic Targets

Sign or acronym SAM

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

General Aspects

Medical area Study of allergies

Health determinants Genetic

Scientific investigator(s) (Contact)

Name of the director Nicolas

Surname Jean-François

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Unit Equipe 14 Inserm U1111- CIRI "Immunologie de l'allergie cutanée et vaccination"

Organization INSERM - Institut National de la Santé et de la Recherche

Name of the director Berard

Surname	Frédéric
Collaborations	
Funding	
Funding status	Mixed
Details	AO HCL Action incitative 2009 INSERM/DHOS 2009
Governance of the database	
Sponsor(s) or organisation(s) responsible	INSERM - Institut National de la Santé et de la Recherche Médicale
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. All patients who attended CCR2A and agreed to be included in cohort. No randomisation or selection.
Database objective	
Main objective	To improve pathophysiological understanding of cutaneous Adverse Drug Reaction (cADRs) by clarifying the number and functional status of T lymphocytes (LT) CD8 + CTLs specific to medication. We wish to: i) confirm the involvement of CD8+ cells for different diseases; ii) understand why the same cell (CD8+) is responsible for different clinical forms of severity.
Inclusion criteria	Patient presenting with maculopapular rash (MPR), Drug Hypersensitivity Syndrome (DRESS) or Lyell's Syndrome (toxic epidermal necrolysis) (SJS/TEN)

caused by medication, diagnosed by referring physicians in the Rhône-Alpes Auvergne Centre de Compétences (CCR2A)

Population type

Age
Adulthood (19 to 24 years)
Adulthood (25 to 44 years)
Adulthood (45 to 64 years)

Population covered
Sick population

Gender
Male
Woman

Geography area
Regional

French regions covered by the database
Auvergne Rhône-Alpes

Detail of the geography area
Rhône-Alpes-Auvergne

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)
2009

Size of the database

Size of the database (number of individuals)
< 500 individuals

Details of the number of individuals
102

Data

Database activity
Current data collection

Type of data collected
Clinical data

Clinical data (detail)
Direct physical measures
Medical registration

Presence of a biobank
Yes

Contents of biobank
Whole blood
Serum
Blood cells isolated
Fluids (saliva, urine, amniotic fluid, ?)

Tissues
Others

Details of biobank content Blood sample (30 mL heparin tube), Two skin biopsies (3 mm in diameter), one for damaged skin and one for healthy skin

Health parameters studied Health event/morbidity
Health event/mortality
Health care consumption and services

Care consumption (detail) Hospitalization
Medical/paramedical consultation
Medicines consumption

Procedures

Data collection method All cohort samples were collected by Allergobiotec (Headed up by Dr. Aurore Rozieres, INSERM U1111-CIRI/HCL).

Participant monitoring Yes

Details on monitoring of participants Systematic visit 3 months following reaction. Other follow-ups (6 months and 9 months) may be scheduled depending on the patient's condition and severity of reaction. A visit at 1 year is also systematic.

Links to administrative sources No

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay) Data from this cohort are accessible through scientific publications. If other researchers wish to access this data, a request must be made and data will be made available according to good clinical practice. Filtered "access request". No "open access".

Access to aggregated data Access on specific project only

Access to individual data Access on specific project only