

# R-ens 2013 - Cross-sectional study on psoriasis severity at a first consultation in dermatology and patient's socioeconomic status

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## General

### Identification

Detailed name	Cross-sectional study on psoriasis severity at a first consultation in dermatology and patient's socioeconomic status
Sign or acronym	R-ens 2013

### General Aspects

Health determinants	Occupation Social and psychosocial factors
Keywords	socioeconomic characteristics, severity, care patterns.

### Scientific investigator(s) (Contact)

Name of the director	Mahé
Surname	Emmanuel
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Unit	Service de Dermatologie
Organization	Centre Hospitalier Victor Dupouy

### Collaborations

### Funding

Funding status	Public
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Details	CHU
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### Governance of the database

Sponsor(s) or organisation(s) responsible	Centre Hospitalier Argenteuil
Organisation status	Public
<b>Additional contact</b>	
<b>Main features</b>	
<b>Type of database</b>	
Type of database	Study databases
Study databases (details)	Not-repeated cross-sectional studies (except case control studies)
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.	Any adult consulting for the first time for psoriasis in one of the participating centers was included in the study.
<b>Database objective</b>	
Main objective	Disadvantaged socioeconomic level is associated with a delay in cancer and cardiovascular diseases care. No data is available on psoriasis. The objective of this study is to highlight a potential correlation between patient's socioeconomic profile and psoriasis severity at the first consultation in dermatology.
Inclusion criteria	- age superior or equal to 18 - psoriasis diagnosis
<b>Population type</b>	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years)
Population covered	Sick population
Gender	Male Woman
Geography area	National

Detail of the geography area	France : Argenteuil, Saint-Mandé, Marseille, Reims, Clamart, Orléans, Martigues, Amiens, Auxerres, Le Mans, Paris, Vienne, Nancy, Montpellier, Pontoise, Boulogne-Billancourt.
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## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY)	01/2013
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Date of last collection (YYYY or MM/YYYY)	05/2013
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### Size of the database

Size of the database (number of individuals)	[500-1000[ individuals
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Details of the number of individuals	683
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### Data

Database activity	Data collection completed
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Type of data collected	Clinical data Declarative data
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Clinical data (detail)	Direct physical measures Medical registration
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Details of collected clinical data	Care patterns, psoriasis severity, rheumatism, comorbidity, smoking, body mass index, depression.
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Declarative data (detail)	Face to face interview
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Details of collected declarative data	Care patterns, psoriasis severity, rheumatism, comorbidity, smoking, body mass index, depression.
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Presence of a biobank	No
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Health parameters studied	Health event/morbidity
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### Procedures

Participant monitoring	No
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Links to administrative sources	No
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## Promotion and access

### Promotion

### Access

Terms of data access (charter for data provision, format of data, availability delay)	Via publications. For more information, contact the scientific investigator.
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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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