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, severy, care patterns.

Scientific investigator(s) (Contact)

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Unit	Service de Dermatologie
Organization	Centre Hospitalier Victor Dupouy

Collaborations

Funding

Funding status	Public
Details	CHU

Governance of the database

Sponsor(s) or organisation(s) responsible	Centre Hospitalier Argenteuil
Organisation status	Public

Additional contact

Main features

Type of database

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.

Database objective

Main objective	Disadvantaged socioeconomic level is associated with a delay in cancer and cardiovacsular 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

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	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.
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	01/2013
Date of last collection (YYYY or MM/YYYY)	05/2013
Size of the database	
Size of the database (number of individuals)	[500-1000[individuals
Details of the number of individuals	683
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Declarative data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Care patterns, psoriasis severity, rheumatisms, comorbidity, smoking, body mass index, depression.
Declarative data (detail)	Face to face interview
Details of collected declarative data	Care patterns, psoriasis severity, rheumatisms, comorbidity, smoking, body mass index, depression.
Presence of a biobank	No
Health parameters studied	Health event/morbidity
Procedures	
Participant monitoring	No

Links to administrative sources	No
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.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only