EPIDERMAS: epidemiological survey on seborrheic dermatitis risk factors - Case-control study of patients attending a dermatologist: risk factors for seborrheic dermatitis

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General

Identification

Detailed name	Case-control study of patients attending a dermatologist: risk factors for seborrheic dermatitis
Sign or acronym	EPIDERMAS: epidemiological survey on seborrheic dermatitis risk factors

General Aspects

Medical area	Dermatology, venereology
Health determinants	Genetic Lifestyle and behavior
Keywords	Risk factors, impact, stress, alcohol, tobacco

Scientific investigator(s) (Contact)

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Unit	CIC1430 CIC HENRI MONDORÉquipe/activité : Module plurithématique
Organization	AP-HP

Collaborations

Funding

Funding status	Public

Details	APHP

Governance of the database

Sponsor(s) or organisation(s) responsible	AP-HP
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Case control study
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.	Cases with sporadic or non-sporadic seborrheic dermatitis are selected and controls are matched according to age (+/- 5 years) and gender that are attending consultation for another reason (except psoriasis and eczema).

Database objective

Main objective	To analyse risk factors for seborrheic dermatitis and risk factors for a flare-ups. To assess the impact of seborrheic dermatitis on quality of life.
Inclusion criteria	- male and female - adult - patient attending a dermatologist

Population type

Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years)
Population covered	General population
Gender	Male Woman

Geography area	National
Detail of the geography area	France

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	10/2005
Date of last collection (YYYY or MM/YYYY)	12/2011

Size of the database

Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	380: - 190 cas/cases - 190 témoins/controls

Data

Database activity	Data collection completed
Type of data collected	Clinical data Declarative data
Clinical data (detail)	Medical registration
Details of collected clinical data	Consultation with dermatologist.
Declarative data (detail)	Paper self-questionnaire
Details of collected declarative data	Sociodemographic characteristics, disease parameters and exposure to risk factors in the months preceding consultation with dermatologist.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Quality of life/health perception

Procedures

Data collection method	Data are collected with questionnaires: the questionnaire is completed by the physician and the self-administered questionnaire is completed by the patient.
Participant monitoring	No

Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://www.sciencedirect.com/science/article/pii/S0151963814008746
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only