

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