

e-TUD - Transversal study on medical management of glaucoma /ocular hypertension

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General

Identification

Detailed name Transversal study on medical management of glaucoma /ocular hypertension

Sign or acronym e-TUD

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation N° CNIL : 1340473

General Aspects

Medical area Ophthalmology

Health determinants Social and psychosocial factors

Others (details) glaucoma

Keywords caring for glaucoma, ocular hypertension

Scientific investigator(s) (Contact)

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Unit Pfizer

Collaborations

Funding

Funding status	Private
Details	Pfizer
Governance of the database	
Sponsor(s) or organisation(s) responsible	pfizer
Organisation status	Private
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.	Recruitment through participating doctors
Database objective	
Main objective	Primary: The main objective of this survey is to describe the diagnostic and therapeutic care for glaucoma and/or ocular hypertensions (defined by an intraocular pressure ≥ 21 mm Hg) examined on the day of the survey. Secondary: The secondary objectives are: - The description of the satisfaction and of the observance perceived by the doctors according to the treatments, - The diagnostic techniques used: visual fields, ocular imaging and gonioscopy, - The risk factors for patients in routine practice, - The exercise mode for ophthalmologists in caring for glaucoma and/or ocular hypertensions. - Putting the data into relation with the data from the survey conducted in 2003.

Inclusion criteria	All of the patients having glaucoma or an OHT coming to a consultation in ophthalmology on the given day were included in the survey, if they met the following conditions: - at least 18 years of age, - diagnosis of glaucoma, or OHT already established at the time of the consultation. - were informed about the survey and gave oral agreement for the use of the data concerning them.
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Population type

Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years) Great age (80 years and more)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	National
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Detail of the geography area	Metropolitan France
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Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	2009
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Date of last collection (YYYY or MM/YYYY)	2009
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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	965
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Data

Database activity	Data collection completed
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Type of data collected	Clinical data
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Clinical data (detail)	Direct physical measures
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Medical registration

Presence of a biobank	No
Health parameters studied	Health care consumption and services
Care consumption (detail)	Medical/paramedical consultation
Procedures	
Data collection method	Electronic CRF
Classifications used	Who-Drug
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)	Methods for accessing the database are currently being defined
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only