

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 | <p>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.</p> <p>Secondary: The secondary objectives are:</p> <ul style="list-style-type: none"> - 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. |

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|--------------------|---|
| Inclusion criteria | <p>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:</p> <ul style="list-style-type: none"> - 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. |
|--------------------|---|

Population type

| | |
|-----|--|
| Age | <p>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)</p> |
|-----|--|

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|--------------------|-----------------|
| Population covered | Sick population |
|--------------------|-----------------|

| | |
|--------|------------------------|
| Gender | <p>Male Woman</p> |
|--------|------------------------|

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|----------------|----------|
| Geography area | National |
|----------------|----------|

| | |
|------------------------------|---------------------|
| Detail of the geography area | Metropolitan France |
|------------------------------|---------------------|

Data collection

Dates

| | |
|--|------|
| Date of first collection (YYYY or MM/YYYY) | 2009 |
|--|------|

| | |
|---|------|
| Date of last collection (YYYY or MM/YYYY) | 2009 |
|---|------|

Size of the database

| | |
|--|------------------------|
| Size of the database (number of individuals) | [500-1000[individuals |
|--|------------------------|

| | |
|--------------------------------------|-----|
| Details of the number of individuals | 965 |
|--------------------------------------|-----|

Data

| | |
|-------------------|---------------------------|
| Database activity | Data collection completed |
|-------------------|---------------------------|

| | |
|------------------------|---------------|
| Type of data collected | Clinical data |
|------------------------|---------------|

| | |
|------------------------|--------------------------|
| Clinical data (detail) | Direct physical measures |
|------------------------|--------------------------|

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 |