# EMU - Cross-sectional Study on Migraine Patients in Emergency Service Departments: Diagnosis and Treatment

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## General

### Identification

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| Detailed name | Cross-sectional Study on Migraine Patients in Emergency Service Departments: Diagnosis and Treatment |
| Sign or acronym | EMU |
| CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation | CNIL no. 906286 |

### General Aspects

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| Medical area | Emergency medicine<br>Neurology |
| Keywords | emergency services |

### Scientific investigator(s) (Contact)

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| Unit | Pfizer |

### Collaborations

### Funding

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| Funding status | Private |

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| 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 by participating doctors |
| **Database objective** | |
| Main objective | Primary:<br>- To estimate the frequency of patients admitted with headaches and to estimate the frequency of patients diagnosed with migraine.<br>- To characterise patients diagnosed with migraines attending emergency services based on demographic data and migraine history (age, severity).<br>- To describe the treatment of patients diagnosed with a migraine during emergency service consultation.<br>Secondary:<br>- To describe the treatment of patients diagnosed with a migraine according to the type of migraine.<br>- To describe the frequency of patients diagnosed with a migraine and their treatment by emergency structure type.<br>- To describe treatment follow-up. |
| Inclusion criteria | - Patients aged 18 or older.<br>- Patients deemed capable of answering the |

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| | questions.<br>- Patients agreeing to participate in the survey.<br>- Patients admitted to emergency department |
| **Population type** | |
| Age | Adulthood (19 to 24 years)<br>Adulthood (25 to 44 years)<br>Adulthood (45 to 64 years)<br>Elderly (65 to 79 years)<br>Great age (80 years and more) |
| Population covered | Sick population |
| Gender | Male<br>Woman |
| Geography area | National |
| Detail of the geography area | Metropolitan France |
| **Data collection** | |
| **Dates** | |
| Date of first collection (YYYY or MM/YYYY) | 2006 |
| Date of last collection (YYYY or MM/YYYY) | 2008 |
| **Size of the database** | |
| Size of the database (number of individuals) | < 500 individuals |
| Details of the number of individuals | 479 |
| **Data** | |
| Database activity | Data collection completed |
| Type of data collected | Clinical data<br>Declarative data |
| Clinical data (detail) | Direct physical measures<br>Medical registration |
| Declarative data (detail) | Face to face interview |
| Presence of a biobank | No |

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| Health parameters studied | Health event/morbidity |
| **Procedures** | |
| Data collection method | Paper CRF |
| Participant monitoring | Yes |
| Details on monitoring of participants | 6-8 weeks |
| Links to administrative sources | No |
| **Promotion and access** | |
| **Promotion** | |
| **Access** | |
| Terms of data access (charter for data provision, format of data, availability delay) | Submission of project to the Pfizer scientific team. |
| Access to aggregated data | Access on specific project only |
| Access to individual data | Access on specific project only |