

ABSINTHE - Observational study of the therapeutic strategy in the management of acute sinusitis in primary care

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Général

Identification

Nom détaillé Observational study of the therapeutic strategy in the management of acute sinusitis in primary care

Sigle ou acronyme ABSINTHE

Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.) CCTI-RS 04.342, CNIL 904410

Thématiques générales

Domaine médical Otolaryngology or ENT

Déterminants de santé Iatrogenic

Mots-clés telithromycin, antibiotic, ENT, Ketek, management, effectiveness, therapeutic strategies, recommendations, pharmacoepidemiology, cohort, Department of Pharmacology, Bordeaux

Responsable(s) scientifique(s)

Nom du responsable Moore

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Laboratoire Service de Pharmacologie, CIC-P 0005-INSERM U657- Université Bordeaux Segalen

Organisme Université Bordeaux

Collaborations

Financements

Financements Mixed

Précisions Laboratoire Sanofi-Aventis (soutien inconditionnel) - Sanofi-Aventis (unconditional support)

Gouvernance de la base de données

Organisation(s) responsable(s) ou promoteur INSERM

Statut de l'organisation Secteur Public

Contact(s) supplémentaire(s)

Caractéristiques

Type de base de données

Type de base de données Study databases

Base de données issues d'enquêtes, précisions Longitudinal study (except cohorts)

Origine du recrutement des participants A selection of health care professionals

Le recrutement dans la base de données s'effectue dans le cadre d'une étude interventionnelle No

Informations complémentaires concernant la constitution de l'échantillon A database of general practitioners and non-hospital ENT specialists provided by Wanadoo was used for selection of physicians. General practitioners and ENT specialists were to include prospectively over a period of 4 weeks patients with an episode of acute sinusitis diagnosed according to criteria of the physician. To avoid conducting the study over a period with marked ecological features such as a period of epidemic influenza or respiratory syncytial virus that can interfere with diagnosis and / or treatment of acute sinusitis, the study was conducted over two four-week periods: March / April and September / October. Each physician was to include the first patients meeting the study inclusion criteria (5 during the first wave of inclusion and 6 in the second).

Objectif de la base de données

Objectif principal	The study objectives were to describe the management of acute sinusitis in real-life conditions of prescription and to evaluate the effectiveness of the initial therapeutic strategy.
Critères d'inclusion	Patient consulting for acute sinusitis diagnosed according to criteria of the practitioner; Patient without a previous episode of acute sinusitis during the two months preceding diagnosis; Patient aged 18 years and over; Patient agreeing to participate in the study; Patient not included in a clinical trial (Huriet-Sérusclat); Patient who may be followed for two months; Patient showing no severe active disease (life-threatening in the next three months).
Type de population	
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)
Population concernée	Sick population
Sexe	Male Woman
Champ géographique	National
Détail du champ géographique	General practitioners and ENT specialists in metropolitan France
Collecte	
Dates	
Année du premier recueil	2005
Année du dernier recueil	2005
Taille de la base de données	
Taille de la base de données (en nombre d'individus)	[1000-10 000[individuals
Détail du nombre d'individus	5693
Données	
Activité de la base	Data collection completed
Type de données recueillies	Clinical data

Declarative data

Données cliniques, précisions Direct physical measures

Données déclaratives, précisions Paper self-questionnaire

Existence d'une bibliothèque No

Paramètres de santé étudiés Health event/morbidity
Health event/mortality
Health care consumption and services

Consommation de soins, précisions Medicines consumption

Modalités

Mode de recueil des données Patients included were the subject of collection of indirectly personal medical data (medical questionnaire completed by the physician at inclusion and 10 days and 2 months after inclusion) and they also had to complete a self-administered within 10 days following inclusion. Participating physicians were also asked to identify and register all patients with an episode of acute sinusitis (with a maximum of 20 patients).

Suivi des participants Yes

Détail du suivi Patients were followed for 2 months with an evaluation point at 10 days and 2 months after diagnosis. These assessment points concerned the evolution of sinusitis and associated care (antibiotics, drainage, other).

Appariement avec des sources administratives No

Valorisation et accès

Valorisation et accès

Lien vers le document <http://tinyurl.com/Pubmed-ABSINTHE>

Description List of publications in Pubmed

Accès

Charte d'accès aux données (convention de mise à disposition, format de données et délais de mise à disposition) A final study report was submitted to the funder. The final study report and scientific communications (posters, papers, ...) are validated by the study Scientific Committee. Ownership of

study data is the subject of an agreement between the University of Bordeaux Segalen and the funder. Terms for third-party access to the database are to be defined.

Accès aux données agrégées

Access on specific project only

Accès aux données individuelles

Access on specific project only