

# REGATE - Longitudinal Study on Patients with Rheumatoid Arthritis and Study on Tolerance and Efficacy of Tocilizumab (REGistry-RoAcTEmra)

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## General

### Identification

Detailed name Longitudinal Study on Patients with Rheumatoid Arthritis and Study on Tolerance and Efficacy of Tocilizumab (REGistry-RoAcTEmra)

Sign or acronym REGATE

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL REGistry-RoAcTEmra

### General Aspects

Medical area Geriatrics  
Rheumatology

Health determinants Medicine

Keywords RA, TCZ, tolerance, comorbidity, efficacy

### Scientific investigator(s) (Contact)

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## Collaborations

## Funding

Funding status Private

Details French Rheumatology Society

## Governance of the database

Sponsor(s) or organisation(s) responsible Société Française de Rhumatologie

Organisation status Private

Sponsor(s) or organisation(s) responsible Club Rhumatiques et Inflammation

Organisation status Private

## Additional contact

## Main features

## Type of database

Type of database Study databases

Study databases (details) Longitudinal study (except cohorts)

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 No

study

## Database objective

Main objective The main objective of the REGATE study is to identify the tolerance and efficacy of Tocilizumab currently used for Rheumatoid Arthritis (RA).

Inclusion criteria - aged 18 or over  
- patient with rheumatoid arthritis

## 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)

Population covered Sick population

Gender Male  
Woman

Geography area National

Detail of the geography area France

## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY) 01/2011

Date of last collection (YYYY or MM/YYYY) 10/2012

### Size of the database

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

Details of the number of individuals 1,500

### Data

Database activity Data collection completed

Type of data collected Clinical data  
Biological data

Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	History, drug intake, DAS28 score, comorbidities such as: Severe infection, cancer, infusion reactions, pregnancy, gastrointestinal perforation, cardiovascular events, cytopenia, cytolysis.
Biological data (detail)	Rheumatoid factor, ACPA
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health care consumption and services
Care consumption (detail)	Medicines consumption
<b>Procedures</b>	
Participant monitoring	Yes
Details on monitoring of participants	at 3 and 6 months, then every 6 months for 5 years.
Links to administrative sources	No
<b>Promotion and access</b>	
<b>Promotion</b>	
Link to the document	<a href="http://www.ncbi.nlm.nih.gov/pubmed/?term=regate+AND+%28Morel[author]+OR+Sibilia[author]%29">http://www.ncbi.nlm.nih.gov/pubmed/?term=regate+AND+%28Morel[author]+OR+Sibilia[author]%29</a>
Description	List of publications in Pubmed
<b>Access</b>	
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