## GO-PRACTICE - Study of the conditions of use of golimumab and its impact, in current practice, in patients with chronic inflammatory rheumatism.

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Funding status

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General	
Identification	
Detailed name	Study of the conditions of use of golimumab and its impact, in current practice, in patients with chronic inflammatory rheumatism.
Sign or acronym	GO-PRACTICE
General Aspects	
Medical area	Rheumatology
Pathology (details)	Rheumatoid arthritis, ankylosing spondylitis and rheumatic psoriasis
Scientific investigator(s) (Contact)	
Name of the director	LEVY-BACHELOT
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Name of the director	Gouyette
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Unit	MSD France
Collaborations	
Funding	

Private

Details MSD France

Governance of the database	
Sponsor(s) or organisation(s) responsible	MSD France
Organisation status	Private
Presence of scientific or steering committees	Yes
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Cohort study
Database objective	
Main objective	To evaluate the maintenance of golimumab therapy 24 months after initial prescription in adult patients with chronic inflammatory rheumatism in clinical practice in France The primary endpoint is the total percentage of patients who maintained treatment with golimumab two years after the initial prescription.
Inclusion criteria	Inclusion: 1. Patients aged 18 years or older. 2. Patients who gave oral consent to participate after receiving oral and written information about the study. 3. Patients with a diagnosis of chronic inflammatory rheumatism. 4. Patients with an initial hospital prescription of golimumab but not yet having initiated treatment with golimumab. 5. Patients able to understand and complete the self-evaluation questionnaires  Non-inclusion: 1. Patients previously treated with golimumab and discontinuing treatment prior to inclusion. 2. Patients who participated in previous golimumab

trials

3. Patients who had already begun treatment with golimumab prior to inclusion and were receiving

treatment at the time of inclusion.

4. Patients with pathologies or conditions that, according to the investigator, would limit the patient's ability to participate fully in the study or to meet all the requirements of the study;

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
Pathology	M05-M14 - Inflammatory polyarthropathies
Gender	Male Woman
Geography area	National
Detail of the geography area	No details available
Data collection	
Dates	
Size of the database	
Size of the database Size of the database (number of individuals)	[500-1000[ individuals
Size of the database (number of	[500-1000[ individuals 750 included
Size of the database (number of individuals)  Details of the number of	
Size of the database (number of individuals)  Details of the number of individuals	
Size of the database (number of individuals)  Details of the number of individuals  Data	750 included
Size of the database (number of individuals)  Details of the number of individuals  Data  Database activity	750 included  Current data collection  Clinical data Declarative data Paraclinical data
Size of the database (number of individuals)  Details of the number of individuals  Data  Data  Database activity  Type of data collected	750 included  Current data collection  Clinical data Declarative data Paraclinical data Biological data  Direct physical measures

Procedures	
Participant monitoring	Yes
Monitoring procedures	Monitoring by contact with the participant (mail, e-mail, telephone etc.) Monitoring by contact with the referring doctor
Links to administrative sources	No
Promotion and access	
Promotion	
Promotion Access	
	No data access charter
Access  Terms of data access (charter for data provision, format of	No data access charter  Access not yet planned