

# RMP - Rangueil Midi-Pyrénées

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Last update : 08/12/2014 | Version : 1 | ID : 60172

| General  |   |
|--|---|
| Identification   |   |
| Detailed name  | Rangueil Midi-Pyrénées  |
| Sign or acronym  | RMP   |
| CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation | Accord CNIL (30/08/2003)  |
| General Aspects  |   |
| Medical area   | Rheumatology  |
| Health determinants  | Geography   |
| Others (details)   | Rheumatoid arthritis  |
| Keywords   | joint inflammatory activity, structural damage, Health episodes, progression, quality of life |
| Scientific investigator(s)<br>(Contact)  |   |
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| Unit   | UMR1043 ? Centre de Physiopathologie de Toulouse-Purpan                                       |
| Organization   | CHU de  |
| Name of the director   | Constantin  |

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| Unit   | U 558 INSERM   |
| Organization   | INSERM - Institut National de la Santé et de la Recherche          |
| Collaborations   |  |
| Funding  |  |
| Funding status   | Public   |
| Details  | CENTRE DE RHUMATOLOGIE DES HÔPITAUX DE TOULOUSE                    |
| Governance of the database   |  |
| Sponsor(s) or organisation(s) responsible                              | CHU de toulouse  |
| Organisation status  | Public   |
| Sponsor(s) or organisation(s) responsible                              | INSERM - Institut National de la Santé et de la Recherche Médicale |
| Organisation status  | Public   |
| Additional contact   |  |
| Main features  |  |
| Type of database   |  |
| Type of database   | Study databases  |
| Study databases (details)  | Cohort study   |
| 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.                     | Prospective Inclusion cut-off date: 01/12/2002                     |

| Database objective                           |  |
|--|--|
| Main objective                               | General objective: to define clinical, radiological and biological criteria that allow formal diagnosis of early cases of rheumatoid arthritis among other early cases of inflammatory rheumatism. Secondary objective: To identify prognostic factors, ahead of early rheumatoid arthritis, that allow the assessment of medium and long-term progression, in order to set the best adapted therapeutic strategy at an early stage. |
| Inclusion criteria                           | Adult patients with early inflammatory rheumatism, showing first clinical signs within the last year   |
| Population type                              |  |
| Age  | Adulthood (19 to 24 years)<br>Adulthood (25 to 44 years)<br>Adulthood (45 to 64 years)   |
| Population covered                           | Sick population  |
| Gender                                       | Male<br>Woman  |
| Geography area                               | Regional   |
| French regions covered by the database       | Languedoc-Roussillon Midi-Pyrénées   |
| Detail of the geography area                 | Midi-Pyrénées region   |
| Data collection                              |  |
| Dates  |  |
| Date of first collection (YYYY or MM/YYYY)   | 06/1992  |
| Date of last collection (YYYY or MM/YYYY)    | 12/2012  |
| Size of the database                         |  |
| Size of the database (number of individuals) | [500-1000[ individuals   |
| Details of the number of individuals         | 823  |
| Data   |  |

|                                       |   |
|---------------------------------------|---|
| Database activity                     | Data collection completed   |
| Type of data collected                | Clinical data<br>Declarative data<br>Paraclinical data<br>Biological data   |
| Clinical data (detail)                | Direct physical measures<br>Medical registration  |
| Declarative data (detail)             | Paper self-questionnaire<br>Face to face interview  |
| Paraclinical data (detail)            | Imaging   |
| Biological data (detail)              | Type of samples taken: ESR, CRP, FR, anti-keratin<br>AC (GIFOR, GBlot, AhFibA, Anti-CCP), acAN  |
| Presence of a biobank                 | Yes   |
| Contents of biobank                   | DNA   |
| Details of biobank content            | DNA bank  |
| Health parameters studied             | Health event/morbidity<br>Health event/mortality<br>Quality of life/health perception   |
| Procedures                            |   |
| Data collection method                | Self-administered questionnaire: entry from a paper questionnaire (manual input); Interviews: entry from a paper questionnaire (manual input); Clinical examination: handwritten (manual input) Biological analysis: handwritten (manual input) |
| Participant monitoring                | Yes   |
| Details on monitoring of participants | Patients in the RMP cohort are no longer monitored.   |
| Links to administrative sources       | No  |
| Promotion and access                  |   |
| Promotion                             |   |
| Link to the document                  | <a href="http://www.ncbi.nlm.nih.gov/pubmed/?term">http://www.ncbi.nlm.nih.gov/pubmed/?term</a>   |
| Link to the document                  | <a href="https://www.em-consulte.com/en/article/86987">https://www.em-consulte.com/en/article/86987</a>   |
| Access                                |   |

|   |   |
|---|---|
| Terms of data access (charter for data provision, format of data, availability delay) | To be decided if data may be used by academic teams To be decided if data may be used by industrial teams |
| Access to aggregated data   | Access on specific project only   |
| Access to individual data   | Access on specific project only   |