

RDPLF - Registre de Dialyse Péritonéale de Langue Française

Head :Verger Christian

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General

Identification

Detailed name Registre de Dialyse Péritonéale de Langue Française

Sign or acronym RDPLF

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

General Aspects

Medical area Urology, andrology and nephrology

Health determinants Nutrition

Others (details) peritoneal dialysis, peritonitis, management of the anemia

Keywords dialysis, peritoneal dialysis, peritonitis, anemia, co-morbidities, quality of dialysis, nurse aspects, survival, nutrition

Scientific investigator(s) (Contact)

Name of the director Verger

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Organization RDPLF

Collaborations

Funding

Funding status	Private
Details	Amgen, Baxter, Fresenius, Genzyme, HAC Pharma, Roche, SHIRE,
Governance of the database	
Sponsor(s) or organisation(s) responsible	Association RDPLF
Organisation status	Private
Additional contact	
Main features	
Type of database	
Type of database	Morbidity registers
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is made on the basis of:	Another treatment or procedure
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	All patients with renal insufficiency treated via peritoneal dialysis in voluntary centers for participation; about 85% of French patients treated via peritoneal dialysis are followed
Database objective	
Main objective	Continuous analysis of the survival rates and complications of patients treated via peritoneal dialysis, assistance for participating centers in order to evaluate and compare themselves, prevent complications, manage the quality of dialysis, nutrition, anemia, surgical aspects linked to the peritoneal approach, assistance in publication. Information for patients.
Inclusion criteria	All patients treated via peritoneal dialysis in French-speaking centers
Population type	
Age	Infant (28 days to 2 years) Early childhood (2 to 5 years)

Childhood (6 to 13 years)
Adolescence (13 to 18 years)
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 International

Detail of the geography area France, Belgium, Switzerland, Tunisia, Morocco, Algeria

Data collection

Dates

Date of first collection (YYYY or MM/YYYY) 1986

Size of the database

Size of the database (number of individuals) Greater than 20 000 individuals

Details of the number of individuals France métropolitaine : 27108 DOM TOM : 1373

Data

Database activity Current data collection

Type of data collected Clinical data
Paraclinical data
Biological data

Clinical data (detail) Direct physical measures
Medical registration

Paraclinical data (detail) date of birth, date of treatments, therapeutic methods

Biological data (detail) urea, creatinine, CBC, albumin, bacteriology, ionogram,

Presence of a biobank No

Health parameters studied

Health event/morbidity
Health event/mortality

Procedures

Data collection method

collection on paper forms, updated at least every 3 months

Participant monitoring

Yes

Details on monitoring of participants

reminders by telephone and mail as soon as the delay in updating exceeds 3 months

Links to administrative sources

No

Promotion and access

Promotion

Link to the document

<http://www.rdplf.org/publications.html>

Link to the document

Description

Link to the document

Description

Access

Terms of data access (charter for data provision, format of data, availability delay)

send back to the centers the statistical analysis of their own data, publication of the annual epidemiological data for the site, annual report can be downloaded from the site. Possibility open to everyone to use a portion of the database using pre-established personal queries on the site for depersonalized export of the database each evening.
The personalized publications and studies can be accessed by everyone, subject to approval by a committee of 12 nurses and doctors elected from within the centers (Commission on Good Practices). Organization of a congress every 2 years bringing together 300 nephrologists and nurses

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only