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 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 compares 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