DIVAT - Données Informatisées et VAlidées en Transplantation

Head :Giral Magali, Institut de Transplantation Urologie Néphrologie, RTRS ?Centaure?, UMR Inserm 1064Centre d'Investigation Clinique et de Biothérapie

Last update : 11/27/2014 | Version : 3 | ID : 21204

General

Identification

Detailed name	Données Informatisées et VAlidées en Transplantation
Sign or acronym	DIVAT
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL (17/09/2004) n°891735, Network DIVAT : 10.16.618

General Aspects

Medical area	Urology, andrology and nephrology
Keywords	Transplantation

Scientific investigator(s) (Contact)

Name of the director	Giral
Surname	Magali
Address	30, bd Jean Monnet, 44093 Nantes Cedex 01
Phone	+33(0)2.40.08.74.43
Email	mgiral@chu-nantes.fr
Unit	Institut de Transplantation Urologie Néphrologie, RTRS ?Centaure?, UMR Inserm 1064Centre d'Investigation Clinique et de Biothérapie
Organization	Centre Hospitalier Universitaire Hôtel Dieu

Collaborations

Funding

Funding status	Mixed

Details	Each center supports its own costs of creation, follow-up and development.We occasionaly receive a financial support from private pharmaceutical lab.

Governance of the database

Sponsor(s) or organisation(s) responsible	Institut de Transplantation Urologie Néphrologie, RTRS ?Centaure?, UMR Inserm 1064
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Morbidity registers
Additional information regarding sample selection.	All recipients of a kidney and/or a pancreas transplant are included in the cohort.

Database objective

Main objective	The eight centers established between them collaboration in order to be able to collect and exchange information, to conduct clinical and epidemiological studies all together, to communicate and exchange data and results with other academic researchers, to construct partnership with pharmaceutical industries and to create a biocollection database linked to clinical data (for 3 of the 8 centers).
Inclusion criteria	Kidney and/or pancreas transplant recipients

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	Nantes, Nancy, Montpellier, Toulouse, Necker

	(Paris), Lyon, Saint-Louis (Paris) et Nice

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1994

Size of the database

Size of the database (number of individuals)	[10 000-20 000[individuals
Details of the number of individuals	11000

Data

Database activity	Current data collection
Type of data collected	Clinical data Paraclinical data Biological data Administrative data
Clinical data (detail)	Direct physical measures Medical registration
Paraclinical data (detail)	Biopsies of the transplant at pre- or post-transplantation. Identification of bacterial, viral, parasitic or mycotic infectious agents.
Biological data (detail)	At pre-transplantation : viral status, immunization and only for pancreas transplantations (C-peptide, HbA1C, serum creatinin, serum proteins, serum albumin, LDL, HDL, cholesterol, triglycerides). At post-transplantation : serum creatinin, proteinuria, hemoglobin, LDL, triglycerides, immunization and only for pancreas transplantations (glycemia, C-peptide, HbA1C, insuline, serum amylase, serum lipase, cholesterol).
Administrative data (detail)	File number, Cristal number (registration on waiting list), patient contact details and name of referring doctor.
Presence of a biobank	Yes
Contents of biobank	Whole blood Serum Plasma Blood cells isolated

	Fluids (saliva, urine, amniotic fluid, ?)
Details of biobank content	DMSO CELLS, SERUM, WHOLE BLOOD, TRIZOL CELLS, URINE, PAXGENE, PLASMA
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Data are collected at pre-transplantation and prospectively at each folow-up visit.
Participant monitoring	Yes
Details on monitoring of participants	Follow-up parameters are collected at 3 months, 6 months, 1 year and then every year.
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://tinyurl.com/PUBMED-DIVAT
Description	Liste des publications dans Pubmed
Link to the document	http://tinyurl.com/HAL-DIVAT
Description	Liste des publications dans HAL

Access

Terms of data access (charter for data provision, format of data, availability delay)	The Divat network encourage collaborations and enables interested researchers to submit a research project. Guidelines for requesting data analyses from the registry will be soon available on the DIVAT website.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only