Pediatric Kidney-Pancreas Transplant Recipient Registration Worksheet

Note: These worksheets are provided to function as a guide to what data will be required in the online TIEDI® application. Currently in the worksheet, a red asterisk is displayed by fields that are required, independent of what other data may be provided. Based on data provided through the online TIEDI® application, additional fields that are dependent on responses provided in these required fields may become required as well. However, since those fields are not required in every case, they are not marked with a red asterisk.

Recipient Information		
Name:		DOB:
SSN:		Birth sex:
HIC:		Transplant Date and Time:
State of Permanent Residence: *		
Permanent Zip: *	-	
*		
Provider Information		
Recipient Center:		
Surgeon Name: *		
NPI#:*		
Donor Information UNOS Donor ID #:		
Recovering OPO:		
Donor Type:		
Patient Status		
Kidney Primary Diagnosis: *		
Specify:		7
Pancreas Primary Diagnosis:*		
Specify:		
Date: Last Seen, Retransplanted or Death*		7
Patient Status: *	OL TATALIC	
ration status.	CLIVING	
	ODEAD	
	RETRANSPLANTE	
Retransplanted organ:		
		s OKidney/Pancreas
Primary Cause of Death:		
Primary Cause of Death:		
Primary Cause of Death: Specify:		
Primary Cause of Death: Specify: Contributory Cause of Death: Specify:		
Primary Cause of Death: Specify: Contributory Cause of Death:		
Primary Cause of Death: Specify: Contributory Cause of Death: Specify: Contributory Cause of Death: Specify:		
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Primary Cause of Death: Specify: Contributory Cause of Death: Specify: Contributory Cause of Death: Specify: Transplant Hospitalization: Date of Admission to Tx Center:*		
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Primary Cause of Death: Specify: Contributory Cause of Death: Specify: Contributory Cause of Death: Specify: Transplant Hospitalization: Date of Admission to Tx Center: * Date of Discharge from Tx Center: Clinical Information : PRETRANSPLANT	Okidney Pancrea	s Kidney/Pancreas delay/impairment
Primary Cause of Death: Specify: Contributory Cause of Death: Specify: Contributory Cause of Death: Specify: Transplant Hospitalization: Date of Admission to Tx Center: * Date of Discharge from Tx Center: Clinical Information : PRETRANSPLANT Functional Status: *	Continue Cognitive Probable Cognitive	delay/impairment
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Probable Motor delay/impairment Questionable Motor delay/impairment No Motor delay/impairment Not Assessed Academic Progress:* Within One Grade Level of Peers Delayed Grade Level Special Education Not Applicable, too young for school/ High School graduate or GED Status Unknown Academic Activity Level:* Full academic load Reduced academic load Unable to participate in academics due to disease or condition Unable to participate regularly in academics due to dialysis Not Applicable, too young for school/ High School graduate or GED Status Unknown Kidney Source of Payment: Primary:* Specify: Pancreas Source of Payment: Primary:* Specify: Pancreas Source of Payment: Primary:* Specify: Pancreas Source of Payment: Primary:* Specify: Belght Measurement Date: Weight Measurement Date: Weight Measurement Date: Weight Measurement Date: Weight Measurement Date: Previous Transplants: Previous Transplant Graft Fail Date Previous Transplant Graft Fail Date Previous Transplants by calling 800-978-4334 or by manifering interlegiptosk@unco.cgn.
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emailing unethelpdesk@unos.org.
If Yes, Date of Most Recent Initiation of Chronic Maintenance Dialysis: ST=
Average Daily Insulin Units: * units/kg/day ST=
Serum Creatinine at Time of Tx: * mg/dl ST=
Viral Detection:
HIV Serostatus:* Positive
Negative
Not Done
UNK/Cannot Disclose
CMV Status* Positive
○ Negative
Not Done
OUNK/Cannot Disclose
HBV Surface Antibody Total *
O Negative
○Negative ○Not Done ○UNK/Cannot Disclose

HBV Core Antibody: ∗	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
HBV Surface Antigen: ★	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
HCV Serostatus: *	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
EBV Serostatus: *	Positive
EBV Scrosulus. W	Negative
	Not Done
	UNK/Cannot Disclose
Vaccination Status	O OTRY Carriet Disclose
Vaccination Status: Did the recipient receive Hepatitis B vaccines prior to	YES NO UNK
transplant?: *	YES NO UNK
Reason not vaccinated:	○Immunity
	OMedical precaution
	OTime constraints
	Patient objection
	Product out of stock
	Other, specify
Specify:	
NAT Results:	
HIV NAT: ∗	Positive
	○ Negative
	Not Done
	OUNK/Cannot Disclose
HBV NAT: ∗	Positive
	ONegative
	ONot Done
	OUNK/Cannot Disclose
HCV NAT: ∗	Positive
	Negative
	Not Done
	OUNK/Cannot Disclose
Previous Pregnancies:	YES
	ONO
	ONOT APPLICABLE: < 10 years old
Malignancies between listing and transplant: $\!$	○YES ○NO
This question is NOT applicable for patients receiving living dono	or transplants who were never on the waiting list.
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If yes, specify type:	□Skin Melanoma
	□Skin Non-Melanoma
	□CNS Tumor
	Genitourinary
	□Breast
	☐Thyroid
	☐Tongue/Throat/Larynx
	□Lung
	□Leukemia/Lymphoma
	Liver
	Other, specify
Specify:	
Bone Disease:	
Fracture in the past year (or since last follow-up): *	YES NO UNK
rracture in the past year (or since last follow-up).	YES NO UNK
Specify Location and number of fractures: *	☐ Spine-compression fracture: # of fractures:
	Extremity: # of fractures:
	Other: # of fractures:
AVN (avascular necrosis): ∗	YES NO UNK
Clinical Information : TRANSPLANT PROCE	DURE
Multiple Organ Recipient	
Were extra vessels used in the transplant procedure:	
Procedure Type:	
Surgical Information:	
Graft Placement: *	INTRA-PERITONEAL
	ORETRO-PERITONEAL
	OPARTIAL INTRA/RETRO-PERITONEAL
Operative Technique: *	Simultaneous Kidney-Pancreas
	Cluster
	Multi-Organ Non-Cluster
Duct Management: *	○ENTERIC W/ROUX-EN-Y
	ENTERIC W/O ROUX-EN-Y
	Суутоутому
	DUCT INJECTION IMMEDIATE
	DUCT INJECTION DELAYED
	OTHER SPECIFY
	OTHER SPECIAL
Specify:	
Venous Vascular Management: *	SYSTEMIC SYSTEM (ILIAC:CAVA)
	PORTAL SYSTEM (PORTAL OR TRIBUTARIES)
	NA/Multi-organ cluster
Arterial Reconstruction:*	CELIAC WITH PANCREAS
	Y-GRAFT TO SPA & SMA
	SPA TO SMA DIRECT
	SPA TO SMA WITH INTERPOSITION
	SPA ALONE
	OTHER SPECIFY
Specify:	
Venous Extension Graft: *	YES NO
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Kidney and Pancreas Preservation Information:	
Total Cold ischemia Time Right KI(OR EN-BLOC): (if pumped, include pump time):	hrs ST=
Total Cold Ischemia Time Left KI (If pumped, include pump time):	hrs ST=
Total Pancreas Preservation Time (include Cold, Warm, Anastomotic time): $\ensuremath{\ast}$	hrs ST=
Kidney(s) received on:∗	○Ice
	Pump
	O _{N/A}
Received on ice:	Stayed on ice
	Put on pump
Received on pump:	
received on pump.	Stayed on pump
	OPut on ice
If put on pump or stayed on pump:	
Right Kidney Final resistance at transplant:	ST=
Right Kidney Final flow rate at transplant:	ST=
Left Kidney Final resistance at transplant:	ST=
Left Kidney Final flow rate at transplant:	ST=
Organ Check-in Information:	
Pancreas Check- Date: Time: In Date and Time:	Military time Time Zone: ST=
Left Kidney Date: Time: Check-In Date and Time:	Military time Time Zone: ST=
Right Kidney Date: Time: Time: and Time:	Military time Time Zone: ST=
En Bloc Kidneys Date: Time: Check-In Date and Time:	Military time Time Zone: ST=
Clinical Information: POST TRANSPLANT Kidney Graft Status: *	Functioning Failed
If death is indicated for the recipient, and the death was a result of	f some other factor unrelated to graft failure, select Functioning.
Resumed Maintenance Dialysis:	OYES ONO
Date Maintenance Dialysis Resumed:	
Kidney Date of Graft Failure:	
Kidney Primary Cause of Graft Failure:	OHYPERACUTE REJECTION
	ACUTE REJECTION
	PRIMARY NON-FUNCTION (GRAFT NEVER FUNCTIONED POST-TRANSPLANT)
	GRAFT THROMBOSIS
	OINFECTION
	SURGICAL COMPLICATIONS
	Ourological complications
	CRECURRENT DISEASE
	OTHER SPECIFY CAUSE
Specify:	
Did patient have any acute kidney rejection episodes	Voc. at least one enjoyed treated with auti-valuation
between transplant and discharge: *	Yes, at least one episode treated with anti-rejection agent
	Yes, none treated with additional anti-rejection agent
	○No
Is growth hormone therapy used between listing and transplant: *	OYES ONO OUNK
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Most Recent Serum Creatinine Prior to Discharge: \ast	mg/dl ST :	=
Patient Need Dialysis within First Week: *	OYES ONO	
Pancreas Graft Status: *	○Functioning ○Failed	
If death is indicated for the recipient, report graft status up until the	instance of death.	
Patient using either oral medication or diet for blood sugar control: $\!$	YES NO UNK	
Patient on oral medication to control blood sugar?*	YES NO UNK	
Date of medications resumed: ∗	ST=	
Patient using diet to control blood sugar:*	YES ONO OUNK	
Patient on insulin?*	YES ONO OUNK	
Date insulin resumed: ∗		ST=
Average total insulin dosage per day: *	units/kg/day	ST=
Insulin duration of use: *	days	ST=
C-peptide value:	ng/mL ST=	
HbA1c:	% ST=	
Pancreas Date of Graft Failure:		
Pancreas Primary Cause of Graft Failure:		
Pancreas Primary Cause of Graft Failure/Specify:		
Contributory causes of graft failure:		
Pancreas Graft/Vascular Thrombosis:	YES NO UNK	
Pancreas Infection:	YES NO UNK	
Bleeding:	YES NO UNK	
Anastomotic Leak:	YES ONO OUNK	
Hyperacute Rejection:	YES NO UNK	
Pancreas Acute Rejection:	OYES ONO OUNK	
Biopsy Proven Isletitis:	YES NO UNK	
Pancreatitis:	YES NO UNK	
Other, Specify:		
Did patient have any acute pancreas rejection episodes between transplant and discharge: *	Yes, at least one episode treated with anti-rejec	tion agent
<u>-</u>	\bigcirc Yes, none treated with additional anti-rejection	agent
	○No	
Pancreas Transplant Complications:		
(Not leading to graft failure.)		
Pancreatitis:*	YES NO UNK	
Anastomotic Leak: *	OYES ONO OUNK	
Abscess or Local Infection: *	OYES ONO OUNK	
Other:		
Weight Post Transplant: *	lbs. kg ST=	
Immunosuppressive Information Are any medications given currently for maintenance or anti-rejection: *	○YES ○NO	
Immunosuppressive Medications		
View Immunosuppressive Medications		
Definitions Of Immunosuppressive Medications		
For each of the immunosuppressive medications listed, select Ind prescribed for the recipient during the initial transplant hospitalizat		

Induction (Ind) immunosuppression includes all medications given for a <u>short finite period</u> in the perioperative period for the purpose of preventing acute rejection. Though the drugs may be continued after discharge for the first 30 days after transplant, it <u>will not</u> be used long-term for immunosuppressive maintenance. Induction agents are usually polyclonal, monoclonal, or IL-2 receptor antibodies (example: methylprednisolone, Campath, Thymoglobulin, or Simulect). Some of these drugs might be used for another finite period for rejection therapy and would be recorded as anti-rejection therapy if used for this reason. For each induction medication indicated, write the <u>total number of days the drug was actually administered</u> in the space provided. For example, if Simulect was given in 2 doses a week apart then the total number of days would be 2, even if the second dose was given after the patient was discharged.

Maintenance (Maint) includes all immunosuppressive medications given before, during or after transplant with the intention to maintain them <u>long-term</u> (example: prednisone, cyclosporine, tacrolimus, mycophenolate mofetil, azathioprine, or Rapamune). This does not include any immunosuppressive medications given to treat rejection episodes, or for induction.

Anti-rejection (AR) immunosuppression includes all immunosuppressive medications given for the purpose of treating an acute rejection episode during the initial post-transplant period or during a specific follow-up period, usually up to 30 days after the diagnosis of acute rejection (example: methylprednisolone, or Thymoglobulin). When switching maintenance drugs (example: from tacrolimus to cyclosporine; or from mycophenolate mofetil to azathioprine) because of rejection, the drugs should not be listed under AR immunosuppression, but should be listed under maintenance immunosuppression.

If an immunosuppressive medication other than those listed is being administered (e.g., new monoclonal antibodies), select Ind, Maint, or AR next to Other Immunosuppressive Medication field, and enter the full name of the medication in the space provided. **Do not list non-immunosuppressive medications.**

Drug used for induction, acute rejection, or maintenance					
Drug used for induction, acute rejection, or maintenance	Ind.	Days	ST	Maint	AR
Steroids (prednisone, methylprednisolone, Solumedrol, Medrol)					
Drugs used for induction or acute rejection					
Drugs used for induction of acute rejection	Ind.	Days	ST	Maint	AR
Atgam					
Campath (alemtuzumab)					
Cytoxan (cyclophosphamide)					
Methotrexate (Folex PFS, Mexate-AQ, Rheumatrex)					
Rituxan (rituximab)					
Simulect (basiliximab)					
Thymoglobulin					
Drugs primarily used for maintenance	Ind.	Days	ST	Maint	AR
Cyclosporine, select from the following:					
- Gengraf					
- Neoral					
- Sandimmune					
- Generic cyclosporine					
Imuran (azathioprine, AZA)					
Leflunomide (LFL)					
Mycophenolic acid, select from the following:					
- CellCept (MMF)					
- Generic MMF (generic CellCept)					
- Myfortic (mycophenolic acid)					
- Generic Myfortic (generic mycophenolic acid)					
mTOR inhibitors, select from the following:					
- Rapamune (sirolimus)					
- Generic sirolimus					
- Zortress (everolimus)					
Nulojix (belatacept)					
Tacrolimus, select from the following:					
- Astagraf XL (extended release tacrolimus)					
- Envarsus XR (tacrolimus XR)					
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- Prograf (tacrolimus)					
- Generic tacrolimus (generic Prograf)					
Other drugs	T., d	D	CT	Maint	4.0
Other immunosuppressive medication, specify:	Ind.	Days	ST	Maint	AR
Other immunosuppressive medication, specify:					

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