## **Pediatric 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<sup>®</sup> 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<sup>®</sup> 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:		
Solidi Type.		
Patient Status		
Primary Diagnosis: *		
Specify:		
Date: Last Seen, Retransplanted or Death*		
Patient Status: *	OLIVING	
	ODEAD	
	ORETRANSPLANTE	D
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: *		
Cognitive Development: *	Operate Committee	
<u>cognitive beveropment.</u>		e delay/impairment ve delay/impairment
		gnitive delay/impairment
	No Cognitive dela	ny/impairment
	Not Assessed	
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motor Development: *		Operinite Motor delay	/impairment	
		Probable Motor delay	y/impairment	
		Questionable Motor of	delay/impairment	
		No Motor delay/impa	airment	
		Not Assessed		
Academic Progress:*		Within One Grade Lev	vel of Peers	
		<b>Delayed Grade Level</b>		
		Special Education		
		Not Applicable, too y	oung for school/ High Sch	nool graduate or GED
		Status Unknown		
Academic Activity Level: *		Full academic load		
		Reduced academic lo	pad	
		Ounable to participate	in academics due to dise	ase or condition
		Unable to participate	regularly due to dialysis	
		Not Applicable, too y	oung for school/ High Sch	nool graduate or GED
		Status Unknown		
Source of Payment: Primary: *				
Specify:				
. ,				
Height Measurement Date:				
Height: *		ft. in.	cm	ST=
Weight Measurement Date:				
Weight: *		lbs	kg	ST=
BMI:	kg/m <sup>2</sup>			
Previous Transplants:				
Previous Transplant Organ	Previous Transp	plant Date	Previous Transplant G	raft Fail Date
Previous Transplant Organ  The three most recent transplants are listed here emailing unethelpdesk@unos.org.				
The three most recent transplants are listed here				
The three most recent transplants are listed here emailing unethelpdesk@unos.org.  Pretransplant Dialysis: *  If Yes, Date of Most Recent Initiation of Chro	e. Please contact t	he UNet Help Desk to confirr		
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The three most recent transplants are listed here emailing unethelpdesk@unos.org.  Pretransplant Dialysis: *  If Yes, Date of Most Recent Initiation of Chro Dialysis:  Average Daily Insulin Units: *	e. Please contact t	he UNet Help Desk to confirm  YES NO UNK	m more than three previous t	stansplants by calling 800-978-4334 or by  ST=
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HBV Core Antibody: *	○ Positive
	Negative
	ONot Done
	UNK/Cannot Disclose
HBV Surface Antigen: *	
riby Surface Artigett. *	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
HCV Serostatus: <b>*</b>	Positive
	Negative
	ONot Done
	UNK/Cannot Disclose
EBV Serostatus: *	Positive
	O Negative
	Not Done
	OUNK/Cannot Disclose
Vaccination Status:	
Did the recipient receive Hepatitis B vaccines prior to transplant?: $\pmb{\ast}$	YES NO OUNK
Reason not vaccinated:	Immunity
	Medical precaution
	Time constraints
	Patient objection
	Product out of stock
	Other, specify
Specify:	, ,
NAT Results:	
HIV NAT:*	O- W
TIV NAL.*	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
HBV NAT: ∗	Positive
	Negative
	Not Done
	UNK/Cannot Disclose
HCV NAT: ∗	Positive
	ONegative
	Not Done
	OUNK/Cannot Disclose
	Sinty cullist biscost
Malignancies between listing and transplant: *	○YES ○NO ○UNK
This question is NOT applicable for patients receiving living do	nor transplants who were never on the waiting list.

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If yes, specify type:	Skin Melanoma
	Skin Non-Melanoma
	□CNS Tumor
	Breast
	☐Thyroid
	☐Tongue/Throat/Larynx
	□Leukemia/Lymphoma
	Cliver
	Other, specify
Specify:	
Clinical Information : TRANSPLANT PROCE	DURE
Multiple Organ Recipient	
Were extra vessels used in the transplant procedure:	
Procedure Type:	
Surgical Information:	
Graft Placement: *	OINTRA-PERITONEAL
	ORETRO-PERITONEAL
	PARTIAL INTRA/RETRO-PERITONEAL
Operative Technique: *	PANCREAS ALONE
operative recilinques x	PANCREAS AFTER KIDNEY
	CLUSTER  MULTI-ORGAN NON-CLUSTER
	PANCREAS WITH KIDNEY DIFFERENT DONOR
Duct Management: *	CENTERIC W/ROUX-EN-Y
	CENTERIC W/O ROUX-EN-Y
	Ссуѕтоѕтому
	ODUCT INJECTION IMMEDIATE
	ODUCT INJECTION DELAYED
	OTHER SPECIFY
Specify:	
Venous Vascular Management: *	SYSTEMIC SYSTEM (ILIAC:CAVA)
	OPORTAL SYSTEM (PORTAL OR TRIBUTARIES)
	ONA/Multi-organ cluster
Arterial Reconstruction:*	CELIAC WITH PANCREAS
	CY-GRAFT TO SPA & SMA
	SPA TO SMA DIRECT
	SPA TO SMA WITH INTERPOSITION
	SPA ALONE
	OTHER SPECIFY
Specify:	
Venous Extension Graft: *	YES NO
·	-, EU -NU
Preservation Information:	
Total Pancreas Preservation Time (include Cold, Warm, Anastomotic time): *	hrs ST=

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Organ Check-in Information:	
Pancreas Check- Date: Time: In Date and Time:	Military time Time Zone:
Clinical Information : POST TRANSPLANT	
Pancreas Graft Status: *	Functioning Failed
If death is indicated for the recipient, report graft status up until t	he 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 ○NO ○UNK
Patient on insulin?*	YES NO UNK

Clinical Information : POST TRANSPLANT					
Pancreas Graft Status: *	Functioning Faile	d			
If death is indicated for the recipient, report graft status up until the	ne 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?*	OYES ONO OUNK				
Date of medications resumed:∗			S	T=	
Patient using diet to control blood sugar:*	OYES ONO OUNK				
Patient on insulin?*	OYES 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=		
Date of Graft Failure:					
Pancreas Primary Cause of Graft Failure:					
Specify:					
Contributory causes of graft failure:					
Pancreas Graft/Vascular Thrombosis:	OYES ONO OUNK				
Pancreas Infection:	OYES ONO OUNK				
Bleeding:	YES NO UNK				
Anastomotic Leak:	OYES ONO OUNK				
Hyperacute Rejection:	OYES ONO OUNK				
Pancreas Acute Rejection:	OYES ONO OUNK				
Biopsy Proven Isletitis:	OYES ONO OUNK				
Pancreatitis:	OYES ONO OUNK				
Other, Specify:					
Pancreas Transplant Complications:					
(Not leading to graft failure.)					
Pancreatitis:*	YES NO UNK				
Anastomotic Leak: *	OYES ONO OUNK				
Abscess or Local Infection: *	OYES ONO OUNK				
Pancreas Transplant Complications: Other					
Did patient have any acute rejection episodes between transplant and discharge: *	Yes, at least one ep	pisode treate	d with anti	-rejection agent	
	Yes, none treated v	with addition	al anti-reje	ection agent	
	○No				
Immunosuppressive Information					
Are any medications given currently for maintenance or anti-rejection: *	YES ONO				

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** (Induction), **Maint** (Maintenance) or **AR** (Anti-rejection) to indicate all medications that were prescribed for the recipient during the initial transplant hospitalization period, and for what reason. If a medication was not given, leave the associated box(es) blank

Induction (Ind) immunosuppression includes all medications given for a short finite period 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 will not 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 total number of days the drug was actually administered 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.** 

Dwg used for industion, partie 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					
brugs 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:	21101	Days	5.	rianic	AIC
- 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:					
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- Envarsus XR (tacrolimus XR)	
- Prograf (tacrolimus)	
- Generic tacrolimus (generic Prograf)	
Othor drugs	
Other drugs  Ind. Days ST Maint	AR
Other immunosuppressive medication, specify:	
Other immunosuppressive medication, specify:	

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