

Pediatric Kidney Transplant Recipient Registration Worksheet

FORM APPROVED: O.M.B. NO. 0915-0157 Expiration Date: 12/31/2011

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.

Name:	DOB:
SSN:	Gender:
HIC:	Tx Date:
State of Permanent Residence: *	
Permanent Zip: *	
Termanent Zip.	
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Recipient Center:	
Surgeon Name: *	
NPI#: *	
UNOS Donor ID #: Donor Type:	
Donor Type.	
Primary Diagnosis: **	
Specify:	
Date: Last Seen, Retransplanted or Death ★	
Date. Last Seen, Retransplanted of Death ***	
	C
al-	0
Patient Status: **	DEAD
	RETRANSPLANTED
	ACTIVITIES ENTRES
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:	
Was patient hospitalized during the last 90 days prior to the transplant admission:	C YES C NO C UNK
	0
	IN INTENSIVE CARE UNIT
Medical Condition at time of transplant: ★	O HOSPITALIZED NOT IN ICU
	NOT HOSPITALIZED
Functional Status: *	
	Definite Cognitive delay/impairment
	Probable Cognitive delay/impairment
Cognitive Development: *	Questionable Cognitive delay/impairment
	No Cognitive delay/impairment
	Not Assessed
	Definite Motor delay/impairment
Motor Development: *	Probable Motor delay/impairment
	Questionable Motor delay/impairment

	0	No Motor delay/impairment
	0	Not Assessed
	0	Within One Grade Level of Peers
	0	
Academic Progress: ★	0	Delayed Grade Level
Academic Progress. The		Special Education
	0	Not Applicable < 5 years old/ High School graduate or GED
	0	Status Unknown
	0	Full academic load
	0	Reduced academic load
	0	
Academic Activity Level: *	0	Unable to participate in academics due to disease or condition
	0	Unable to participate regularly in academics due to dialysis
		Not Applicable < 5 years old/ High School graduate or GED
	0	Status Unknown
Source of Payment:		
Primary: *		
Specify:		
Secondary:	<u> </u>	
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Previous Transplants:)
Previous Transplant Organ	Previous Transplant	Date Previous Transplant Graft Fail Date
The three most recent transplants are listed 978-4334 or by emailing unethelpdesk@uno		UNet Help Desk to confirm more than three previous transplants by calling 800-
Pretransplant Dialysis: *	0	YES NO UNK
If Yes, Date of Most Recent Initiation of Chro Maintenance Dialysis:	onic	ST=

Serum Creatinine at Time of Tx: *		mg/dl ST=
Viral Detection:		
HIV Serostatus: ★	0 0 0	Positive Negative Not Done UNK/Cannot Disclose
CMV lgG: *	0 0 0	Positive Negative Not Done UNK/Cannot Disclose
CMV IgM: ★	0 0 0	Positive Negative Not Done UNK/Cannot Disclose
HBV Core Antibody: ★	0 0 0	Positive Negative Not Done UNK/Cannot Disclose
HBV Surface Antigen: ★	0 0 0	Positive Negative Not Done UNK/Cannot Disclose
HCV Serostatus: ★	0	Positive

	0	Negative
	0	Not Done
	0	UNK/Cannot Disclose
	0	Positive
	0	Negative
EBV Serostatus: *	0	
	0	Not Done
		UNK/Cannot Disclose
Was preimplantation kidney biopsy performed at the transplant center:	0	YES NO
Did patient receive any pretransplant blood transfusions:	0	YES NO UNK
Any tolerance induction technique used:	0	YES NO UNK
	_	
	0	NO PREVIOUS PREGNANCY
	0	1 PREVIOUS PREGNANCY
	0	2 PREVIOUS PREGNANCIES
	0	3 PREVIOUS PREGNANCIES
Previous Pregnancies:	0	4 PREVIOUS PREGNANCIES
	0	5 PREVIOUS PREGNANCIES
	0	MORE THAN 5 PREVIOUS PREGNANCIES
	0	NOT APPLICABLE: < 10 years old
	0	UNKNOWN
Malignancies between listing and transplant: ★	0	YES NO C UNK

This question is NOT applicable for patients receiving living donor transplants who were never on the waiting list.

		Skin Melanoma		
		Skin Non-Melanoma		
		CNS Tumor		
		Genitourinary		
		Breast		
If yes, specify type:		Thyroid		
		Tongue/Throat/Larynx		
		Lung		
		Leukemia/Lymphoma		
		Liver		
		Other, specify		
Specify:				
Bone Disease:				
Fracture in the past year (or since last follow-up): *	0	YES O NO UNK		
	fractur	Spine-compression re:	# of fractures:	
Specify Location and number of fractures: *		Extremity:	# of fractures:	
		Other:	# of fractures:	
AVN (avascular necrosis): ★	0	YES O NO UNK		
AVN (avascular necrosis): *	0	YES NO UNK		
AVN (avascular necrosis): * Multiple Organ Recipient	0	YES NO UNK		
	0	YES NO UNK		

Kidney Preservation Information:

Total Cold ischemia Time Right KI(OR EN-BLOC): (if pumped, include pump time):	hrs	ST=			
Total Warm Ischemia Time Right KI (OR EN-BLOC): (Include Anastomotic time):	min	ST=			
Total Cold ischemia Time Left KI (if pumped, include pump time):	hrs	ST=			
Total Warm ischemia Time Left KI (include Anastomotic time):	min	ST=			
	C				
Kidney(s) received on: ★	Pump				
	C _{N/A}				
	C Stayed on ice				
Received on ice:	C Put on pump				
	C Stayed on pump				
Received on pump:	Put on ice				
If put on pump or stayed on pump:					
Final resistance at transplant:		ST=			
Final flow rate at transplant:		ST=			
Incidental Tumor found at time of Transplant:	C YES C NO UNK				
	Oncocytoma				
	Renal Cell Carcinoma				
	Carcinoid				
If yes, specify tumor type:	Adenoma				
	Transitional Cell Carcinoma				
	Other Primary Kidney Tumor, Specify.				
Specify:					

Graft Status: ★	C Functioning Failed
If death is indicated for the recipient, and the death was a re-	sult of some other factor unrelated to graft failure, select Functioning.
Resumed Maintenance Dialysis:	C YES NO
Date Maintenance Dialysis Resumed:	
Select a Dialysis Provider:	
Provider #:	
Provider Name:	
Date of Graft Failure:	
	HYPERACUTE REJECTION
	ACUTE REJECTION
	PRIMARY FAILURE
D: 0 (0 (5)	GRAFT THROMBOSIS
Primary Cause of Graft Failure:	INFECTION
	SURGICAL COMPLICATIONS
	UROLOGICAL COMPLICATIONS
	RECURRENT DISEASE
	OTHER SPECIFY CAUSE
Specify:	
Contributory causes of graft failure:	
Acute Rejection:	C YES C NO C UNK
Graft Thrombosis:	C YES C NO C UNK
Infection:	C YES C NO C UNK
Surgical Complications:	C YES NO O UNK
Urological Complications:	C YES NO C UNK

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Recurrent Disease:	C YES O NO UNK
Other, Specify:	
Most Recent Serum Creatinine Prior to Discharge: **	mg/dl ST=
Kidney Produced > 40ml of Urine in First 24 Hours:	C YES C NO
Patient Need Dialysis within First Week: *	C YES C NO
Creatinine decline by 25% or more in first 24 hours on 2 separate samples:	C YES C NO
	Yes, at least one episode treated with anti-rejection agent
Did patient have any acute rejection episodes between transplant and discharge: **	Yes, none treated with additional anti-rejection agent
	C _{No}
	Biopsy not done
Was biopsy done to confirm acute rejection:	Yes, rejection confirmed
	Yes, rejection not confirmed
Is growth hormone therapy used between listing and transplant: **	C YES C NO C UNK
Date of Measurement:	
Height: *	ft. in. cm ST=
Weight: ★	lbs sT=
BMI:	kg/m ²
Biological or Anti-viral Therapy:	C YES NO Unknown/Cannot disclose
If Yes, check all that apply:	Acyclovir (Zovirax)

		Cytogam (CMV)
		Gamimune
		Gammagard
		Ganciclovir (Cytovene)
		Valgancyclovir (Valcyte)
		HBIG (Hepatitis B Immune Globulin)
		Flu Vaccine (Influenza Virus)
		Lamivudine (Epivir) (for treatment of Hepatitis B)
		Other, Specify
		Valacyclovir (Valtrex)
Specify:		
Specify:		
Other therapies:	0	YES NO
		Photopheresis
If Yes, check all that apply:		Plasmapheresis
		Total Lymphoid Irradiation (TLI)
<u> </u>		
Are any medications given currently for maintenance or	_	
Are any medications given currently for maintenance or anti-rejection: **		YES NO
Did the patient participate in any clinical research protocol for immunosuppressive medications:	О	YES NO
If Yes, Specify:		
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View Immunosuppressive Medications		

For each of the immunosuppressant medications listed, check **Previous Maintenance (Prev Maint)**, **Current Maintenance (Curr Maint)** or **Antirejection (AR)** to indicate all medications that were prescribed for the recipient during this follow-up period, and for what reason. If a medication was not given, leave the associated box(es) blank.

Previous Maintenance (Prev Maint) includes all immunosuppressive medications given during the report period, which covers the period from the last clinic visit to the current clinic visit, for varying periods of time which may be either long-term or intermediate term with a tapering of the dosage until the drug is either eliminated or replaced by another long-term maintenance drug (example: Prednisone, Cyclosporine, Tacrolimus, Mycophenolate Mofetil, Azathioprine, or Rapamycin). This does not include any immunosuppressive medications given to treat rejection episodes.

Current Maintenance (Curr Maint) includes all immunosuppressive medications given at the current clinic visit to begin in the next report for varying periods of time which may be either long-term or intermediate term with a tapering of the dosage until the drug is either eliminated or replaced by another long-term maintenance drug (example: Prednisone, Cyclosporine, Tacrolimus, Mycophenolate Mofetil, Azathioprine, or Rapamycin). This does not include any immunosuppressive medications given to treat rejection episodes.

Anti-rejection (AR) immunosuppression includes all immunosuppressive medications given for the purpose of treating an acute rejection episode since the last clinic visit (example: Methylprednisolone, Atgam, OKT3, 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.

Note: The Anti-rejection field refers to any anti-rejection medications since the last clinic visit, not just at the time of the current clinic visit.

If an immunosuppressive medication other than those listed is being administered (e.g., new monoclonal antibodies), select Previous Maint, or Current 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.**

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, Atgam, Thymoglobulin, OKT3, Simulect, or Zenapax). Some of these drugs might be used for another finite period for rejection therapy and would be recorded as 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 or Zenapax 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 for varying periods of time which may be either long-term or intermediate term with a tapering of the dosage until the drug is either eliminated or replaced by another long-term maintenance drug (example: Prednisone, Cyclosporine, Tacrolimus, Mycophenolate Mofetil, Azathioprine, or Rapamycin). 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, Atgam, OKT3, 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.

Steroids (Prednisone, Methylprednisolone, Solumedrol, Medrol, Deca dron)	Ind.	Days	ST	Maint	AR
Atgam (ATG)					
OKT3 (Orthoclone, Muromonab)					
Thymoglobulin					
Simulect - Basiliximab					
Zenapax - Daclizumab					

Azathioprine (AZA, Imuran)				
EON (Generic Cyclosporine)				
Gengraf (Abbott Cyclosporine)				
Other generic Cyclosporine, specify brand:				
Neoral (CyA-NOF)				
Sandimmune (Cyclosporine A)				
CellCept (Mycophenolate Mofetil; MMF)				
Generic MMF (Generic CellCept)				
Prograf (Tacrolimus, FK506)				
Generic Tacrolimus (Generic Prograf)				
Advagraf (Tacrolimus Extended or Modified Release)				
Nulojix (Belatacept)				
Sirolimus (RAPA, Rapamycin, Rapamune)				
Myfortic (Mycophenolate Sodium)				
Compath Name (Ind. Day	s S	laint .	_
Campath - Alemtuzumab (anti-CD52)				
Cyclophosphamide (Cytoxan)				
Leflunomide (LFL, Arava)				
Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)				
Other Immunosuppressive Medication, Specify				
Rituximab				

	Ind.	Days	ST	Maint	AR
Zortress (Everolimus)					
Other Immunosuppressive Medication, Specify					