## Records ?

## Pediatric Kidney Transplant Recipient Follow-Up Worksheet

FORM APPROVED: O.M.B. NO. 0915-0157 Expiration Date: 10/31/2010

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:	Gender:
HIC:	Tx Date:
Previous Follow-Up:	Previous Px Stat Date:
Transplant Discharge Date:	
State of Permanent Residence:*	
Zip Code: *	
Provider Information	
Recipient Center:	
Followup Center:	
Physician Name: <b>*</b> NPI#: <b>*</b>	
Follow-up Care Provided By: *	<ul> <li>Transplant Center</li> <li>Non Transplant Center Specialty Physician</li> <li>Primary Care Physician</li> <li>Other Specify</li> </ul>
Specify:	
Donor Information	
UNOS Donor ID #:	
Donor Type:	
Patient Status	
Date: Last Seen, Retransplanted or Death $st$	

Patient Status:\*

O DEAD

Primary Cause of Death:	
Specify:	
Contributory Cause of Death:	
Specify:	
Contributory Cause of Death:	
Specify:	
Hospitalizations:	
Has the patient been hospitalized since the last patient status date: *	C YES C NO C UNK
Number of Hospitalizations:	ST=
	Disease Recurrence:
	No recurrence
TRR	Suspected recurrence (not confirmed or unknown is confirmed by biopsy)
Diagnosis:	Biopsy confirmed recurrence
	C Unknown
Noncompliance:	
Was there evidence of noncompliance with immunosuppression medication during this follow-up period that compromised the patient's recovery:	YES NO CUNK
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
	Probable Motor delay/impairment
Motor Development: *	Questionable Motor delay/impairment
	No Motor delay/impairment
	Not Assessed
	Within One Grade Level of Peers
	C Delayed Grade Level
Academic Progress: *	Special Education
	Not Applicable < 5 years old/ High School graduate or GED
	C Status Unknown
	Full academic load
	Reduced academic load
Academic Activity Level:*	Unable to participate in academics due to disease or condition
	Not Applicable < 5 years old/ High School graduate or GED
	G Status Unknown
Primary Insurance at Follow-up: *	
Specify:	
Clinical Information	
Date of Measurement:	
Height: *	ft in cm ST=
Weight: *	lbs. kg ST=
BMI: kg/m <sup>2</sup>	
Urine Protein Found By Any Method:	VES NO UNK
Diabetes onset during the follow-up period: $ imes$	C YES C NO C UNK
If yes, insulin dependent:	CYES NO CUNK
Graft Status: *	Functioning Failed

If Functioning, Most Recent Serum Creatinin	e:ST=
Date of Failure:	
Primary Cause of Graft Failure:	
Other, Specify:	
Contributory causes of graft failure:	
Acute Rejection	🦷 yes 🖷 no 🥌 unk
Chronic Rejection	C YES C NO C UNK
Graft Thrombosis	C YES C NO C UNK
Infection	CYES CNO CUNK
Urological Complications	CYES CNO CUNK
Patient Noncompliance	YES 🦳 NO 🧖 UNK
Recurrent Disease	CYES CNO CUNK
BK (Polyoma) Virus	🧉 YES 🌀 NO 🥌 UNK
Other, Specify:	
	© NO
	YES, RESUMED MAINTENANCE DIALYSIS
Dialysis Since Last Follow-Up: *	YES, NO MAINTENANCE RESUMPTION
	YES, MAINTENANCE RESUMPTION UNKNOWN
	C UNKNOWN
Date Maintenance Dialysis Resumed:	
Select a Dialysis Provider:	
Provider #:	
Provider Name:	

Did patient have any acute rejection episodes during the follow-up period: ≭	<ul> <li>Yes, none treated with additional anti-rejection agent</li> <li>No</li> <li>Unknown</li> </ul>
Was biopsy done to confirm acute rejection:	<ul> <li>Biopsy not done</li> <li>Yes, rejection confirmed</li> <li>Yes, rejection not confirmed</li> <li>Unknown</li> </ul>
Viral Detection:	
CMV lgG:	<ul> <li>Positive</li> <li>Negative</li> <li>Not Done</li> <li>UNK/Cannot Disclose</li> </ul>
CMV IgM:	<ul> <li>Positive</li> <li>Negative</li> <li>Not Done</li> <li>UNK/Cannot Disclose</li> </ul>
Is growth hormone therapy used during this followup period: <del>*</del>	C YES C NO C UNK
Post Transplant Malignancy: *	YES NO UNK
Donor Related:	C YES C NO C UNK
Recurrence of Pre-Tx Tumor:	C YES C NO C UNK
Post Tx De Novo Solid Tumor:	C YES C NO C UNK
De Novo Lymphoproliferative disease and Lymphoma:	C YES C NO C UNK
Bone Disease:	

Fracture in the past year (or since last follow-

up):*	C YES C NO C UNK		
	Spine-compression fracture:	# of fractures:	
Specify Location and number of fractures: $st$	Extremity:	# of fractures:	
	Other:	# of fractures:	
AVN (avascular necrosis):*	🦷 yes 🔍 no 🔍 unk		

Treatment	
Biological or Anti-viral therapy:	YES NO Unknown/Cannot disclose
	Acyclovir (Zovirax)
	Cytogam (CMV)
	Gamimune
	Gammagard
	Ganciclovir (Cytovene)
If Yes, check all that apply:	Valgancyclovir (Valcyte)
	HBIG (Hepatitis B Immune Globulin)
	Flu Vaccine (Influenza Virus)
	Lamivudine (Epivir) (for treatment of Hepatitis B)
	Valacyclovir (Valtrex)
	Other, Specify
Specify: *	
Specify:	
Treatment for BK (polyoma) virus:	C YES C NO
	Yes, Immunosuppression reduction
	Yes, Cidofovir
If Yes, check all that apply:	Yes, IVIG
	Yes, Type Unknown
	Yes, Other, Specify

Specify: *	
Other therapies:	YES NO
	Photopheresis
If Yes, check all that apply:	Plasmapheresis
	Total Lymphoid Irradiation (TLI)

Immunosuppressive Information	
Previous Validated Maintenance Follow-Up Medications:	
Previous Validated Maintenance Follow-Up Medications:	
	Yes, same as validated TRR form
Were any medications given during the follow-	Yes, same as previous validated report
up period for maintenance:	Yes, but different than previous validated report
	None given
Did the physician discontinue all maintenance immunosuppressive medications:	SYES NO
Did the patient participate in any clinical research protocol for immunosuppressive medications:	YES NO
Specify: *	

Immunosuppressive Medications View Immunosuppressive Medications

## Definitions Of Immunosuppressive Follow-Up Medications

For each of the immunosuppressant medications listed, check **Previous Maintenance (Prev Maint)**, **Current Maintenance (Curr Maint)** or **Anti-rejection (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 <u>should not</u> be listed under AR immunosuppression, but <u>should be</u> 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.** 

	Prev Maint	Curr Maint	AR
Steroids (Prednisone,Methylprednisolone,Solumedrol,Medrol,Decadron)			
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)			
Modified Release Tacrolimus FK506E (MR4)			
Sirolimus (RAPA, Rapamycin, Rapamune)			
Myfortic (Mycophenolate Sodium)			

Other Immunosuppressive Medications			
	Prev Maint	Curr Maint	AR
Campath - Alemtuzumab (anti-CD52)			
Cyclophosphamide (Cytoxan)			
Leflunomide (LFL, Arava)			

Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)		
Other Immunosuppressive Medication, Specify		
Rituximab		

Investigational Immunosuppressive Medications			
	Prev Maint	Curr Maint	AR
Everolimus (RAD, Certican)			
Other Immunosuppressive Medication, Specify			

UNOS View Only	
Comments:	