

## **Adult Kidney Transplant Recipient Follow-Up 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:
Previous Follow-Up:	Previous Px Stat Date:
Transplant Discharge Date:	
State of Permanent Residence: *	
Zip Code: ★	
Recipient Center:	
Followup Center:	
Physician Name: **	
NPI#: *	
	C Transplant Center
Follow-up Care Provided By: **	Non Transplant Center Specialty Physician
	Primary Care Physician
	Other Specify
Specify:	
UNOS Donor ID #:	
Donor Type:	
Date: Last Seen, Retransplanted or Death *	

Patient Status: *	C LIVING DEAD	
	RETRANSPLANTED	
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 sta	NC C	) )
Number of Hospitalizations:	ST=	NK.
	Disease Recurrence:	
	No recurrence	
TRR Diagnosis:	Suspected recurrence (not confirmed or unknown is confirmed by biopsy)	
	Biopsy confirmed recurrence	
	C Unknown	
Noncompliance:		
Was there evidence of noncompliance with immunosuppres recovery:	ssion medication during this follow-up period that compromised the patient's  YE	) ES

0
NO
0
UNK

Functional Status: *		
Physical Capacity:	0 0 0	No Limitations  Limited Mobility  Wheelchair bound or more limited  Not Applicable (< 1 year old or hospitalized)  Unknown
Working for income: *	0	YES NO UNK
If No, Not Working Due To:		
If Yes:	0000000	Working Full Time  Working Part Time due to Demands of Treatment  Working Part Time due to Disability  Working Part Time due to Insurance Conflict  Working Part Time due to Inability to Find Full Time Work  Working Part Time due to Patient Choice  Working Part Time Reason Unknown  Working, Part Time vs. Full Time Unknown
Academic Progress:	0	Within One Grade Level of Peers  Delayed Grade Level  Special Education

	Not Applicable < 5 years old/ 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
	Not Applicable < 5 years old/ High School graduate or GED  Status Unknown
Primary Insurance at Follow-up: *	
Specify:	
Height: Weight: BMI:	ft. in. cm ST= ST= ST= ST=
Urine Protein Found By Any Method:	C YES C NO C UNK
Diabetes onset during the follow-up period: *	C YES C NO C UNK
If yes, insulin dependent:	C YES C NO C UNK
Graft Status: *	C Functioning Failed
If death is indicated for the recipient, and the death was	s a result of some other factor unrelated to graft failure, select Functioning.
If Functioning, Most Recent Serum Creatinine:	mg/dl ST=
Date of Failure:	
Primary Cause of Graft Failure:	
Other, Specify:	
Contributory causes of graft failure:	

Acute Rejection		0	YES O	NO
Chronic Rejection		0	YES O	NO
Graft Thrombosis		0	YES O	NO
Infection		0	YES O	NO
Urological Complications		0	YES O	NO
Patient Noncompliance		0	YES O	NO
Recurrent Disease		0	YES O	NO
BK (Polyoma) Virus		0	YES O	NO
Other, Specify:				
Dialysis Since Last Follow-Up: <b>米</b>	O NO O YES, RESUMED MAINTENANCE DIALYSIS O YES, NO MAINTENANCE RESUMPTION O YES, MAINTENANCE RESUMPTION UNKNOWN O UNKNOWN			
Date Maintenance Dialysis Resumed:				
Select a Dialysis Provider:				

Provider #:	
Provider Name:	
Did patient have any acute rejection episodes during the follow-up period: ★	Yes, at least one episode treated with anti-rejection agent Yes, none treated with additional anti-rejection agent No Unknown
Was biopsy done to confirm acute rejection:	Biopsy not done  Yes, rejection confirmed  Yes, rejection not confirmed  Unknown
Viral Detection:	
CMV IgG:	Positive Negative Not Done UNK/Cannot Disclose
CMV IgM:	Positive Negative Not Done UNK/Cannot Disclose
Post Transplant Malignancy: *	C YES C NO C UNK
Donor Related:	C YES C NO 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
Biological or Anti-viral therapy:	C YES NO C 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 NO
	Yes, Immunosuppression reduction
If Yes, check all that apply:	Yes, Cidofovir
	Yes, IVIG

	Yes, Type Unknown
	Yes, Other, Specify
Specify: *	
Other therapies:	C <sub>YES</sub> C <sub>NO</sub>
	Photopheresis
If Yes, check all that apply:	Plasmapheresis
	Total Lymphoid Irradiation (TLI)
Previous Validated Maintenance Follow-Up Medications:	·
Previous Validated Maintenance Follow-Up Medications:	
Were any medications given during the follow-up period for	Yes, same as validated TRR form Yes, same as previous validated report
maintenance:	Yes, but different than previous validated report
	None given
Did the physician discontinue all maintenance immunosuppressive medications:	C <sub>YES</sub> C <sub>NO</sub>
Did the patient participate in any clinical research protocol for immunosuppressive medications:	C YES C NO
Specify: *	
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 **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 <a href="maintenance immunosuppression">should be listed under AR immunosuppression</a>, but <a href="maintenance immunosuppression">should be listed under AR immunosuppression</a>.

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. <u>Do not list non-immunosuppressive</u> 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)			
Advagraf (Tacrolimus Extended or Modified Release)			
Nulojix (Belatacept)			
Sirolimus (RAPA, Rapamycin, Rapamune)			
Myfortic (Mycophenolate Sodium)			
	Prev Maint	Curr Maint	AR
Campath - Alemtuzumab (anti-CD52)	Prev Maint	Curr Maint	AR
Campath - Alemtuzumab (anti-CD52)  Cyclophosphamide (Cytoxan)	Prev Maint		_
Cyclophosphamide (Cytoxan)			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)  Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)  Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)  Other Immunosuppressive Medication, Specify			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)  Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)  Other Immunosuppressive Medication, Specify			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)  Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)  Other Immunosuppressive Medication, Specify			
Cyclophosphamide (Cytoxan)  Leflunomide (LFL, Arava)  Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)  Other Immunosuppressive Medication, Specify			