

## **Pediatric Liver Transplant Recipient Registration Worksheet**

The revised worksheet sample is for reference purposes only and is pending OMB approval.

Note: These worksheets are provided to function as a guide to what data will be required in the online TIEDI<sup>B</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>B</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:
0.7 (0	
State of Permanent Residence: *	
Permanent Zip: *	-
Provider Information	
Recipient Center:	
Surgeon Name: *	
NPI: *	
Donor Information	
UNOS Donor ID #:	
Donor Type:	
Patient Status	
Primary Diagnosis: *	
Specify:	
Date: Last Seen, Retransplanted or Death *	
	LIVING
Patient Status: *	© DEAD
	© RETRANSPLANTED
	RETRANSPLANTED
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:	© YES © NO © UNK
Medical Condition at time of transplant: ★	<ul><li>IN INTENSIVE CARE UNIT</li><li>HOSPITALIZED NOT IN ICU</li><li>NOT HOSPITALIZED</li></ul>
Patient on Life Support: *	✓ YES ✓ NO  Ventilator  Artificial Liver
Specify:	Other Mechanism, Specify
Functional Status: *	
Cognitive Development: *	Definite Cognitive delay/impairment (verified by IQ score <70 or unambiguous behavioral observation)  Probable Cognitive delay/impairment (not verified or unambiguous but more likely than not, based on behavioral observation or other evidence)  Questionable Cognitive delay/impairment (not judged to be more likely than not, but with some indication of cognitive delay/impairment such as expressive/receptive language and/or learning difficulties)  No Cognitive delay/impairment (no obvious indicators of cognitive delay/impairment)  Not Assessed
Motor Development: *	<ul> <li>Definite Motor delay/impairment (verified by physical exam or unambiguous behavioral observation)</li> <li>Probable Motor delay/impairment (not verified or unambiguous but more likely than not, based on behavioral observation or other evidence)</li> <li>Questionable Motor delay/impairment (not judged to be more likely than not, but with some indications of motor delay/impairment)</li> <li>No Motor delay/impairment (no obvious indicators of motor delay/impairment)</li> <li>Not Assessed</li> </ul>
	Within One Grade Level of Peers

Academic Progress: *	C Delayed Grade Level C Special Education C Not Applicable < 5 years old C Status Unknown		
Academic Activity Level: *	<ul> <li>Full academic load</li> <li>Reduced academic load</li> <li>Unable to participate in academics due to disease or condition</li> <li>Not Applicable &lt; 5 years old/ High School graduate</li> <li>Status Unknown</li> </ul>		
Source of Payment:			
Primary: *			
Specify:			
Secondary:			
Clinical Information : PRETRANSPLA	NT		
Date of Measurement: *			
Height: *	ft. in. cm %ile ST=		
Weight: *	lbs kg %ile ST=		
BMI:	kg/m <sup>2</sup> %ile		
Previous Transplants:			
Previous Transplant Organ	Previous Transplant Date Previous Transplant Graft Fail Date		
The three most recent transplants are listed here. Please contact the UNet Help Desk to confirm more than three previous transplants by calling 800-978-4334 or by emailing unethelpdesk@unos.org.			
Viral Detection:			
	Positive		
HIV Serostatus: ★	Negative		
	Not Done		
	O UNK/Cannot Disclose		
	C Positive		
	Negative		
CMV IgG: *	○ Not Done		
	UNK/Cannot Disclose		

CMV IgM: <b>≭</b>	C Positive
	Negative
	Not Done
	C UNK/Cannot Disclose
	C Positive
	C Negative
HBV Core Antibody: ★	C Not Done
	C UNK/Cannot Disclose
	C Positive
HBV Surface Antigen: ★	Negative
nov Surface Antigen.	Not Done
	C UNK/Cannot Disclose
	C Positive
LIGITO A STATE OF THE STATE OF	C Negative
HCV Serostatus: *	Not Done
	C UNK/Cannot Disclose
	C Positive
EBV Serostatus: *	C Negative
EBV Gelosiands.	Not Done
	C UNK/Cannot Disclose
Any tolerance induction technique used:	C YES C NO C UNK
Pretransplant Lab Date:	
SGPT/ALT:	U/L ST=
Malignancies between listing and transplant: *	C YES C NO C UNK
This question is NOT applicable for patients receiving living de	onor transplants who were never on the waiting list.
	Skin Melanoma
	Skin Non-Melanoma
	CNS Tumor
	Genitourinary
	☐ Breast

I and the second				
	☐ Thyroid			
If yes, specify type:	Tongue/Throat/Larynx			
	Lung			
	Leukemia/Lymphoma			
	☐ Hepatoblastoma			
	Hepatocellular Carcinoma			
	Liver			
	Other, specify			
Specify:				
Clinical Information : TRANSPLANT PROCEDURE				
Multiple Organ Recipient				
Were extra vessels used in the transplant procedure:				
Vessel Donor ID:				
	G entretene			
Surgical Procedure:	ORTHOTOPIC HETEROTOPIC			
	HETEROTOPIC			
	Mhole Liver			
	Partial Liver, remainder not Tx or Living Transplant			
Procedure Type:	Split Liver			
Troccuire Type.	Whole Liver with Pancreas (Technical Reasons)			
	Partial Liver with Pancreas (Technical Reasons)			
	Split Liver with Pancreas (Technical Reasons)			
Split Type:				
- Special Control of the Control of				
Preservation Information:				
Warm Ischemia Time (include anastomotic time):	min ST=			
Total Cold Ischemia Time (if pumped, include pump time): *	hrs ST=			
Risk Factors:				
Did Patient receive 5 or more units of packed red blood cells within 48 hours prior to transplantation due to spontaneous portal hypertensive bleeding:	C YES ONO UNK			
Spontaneous Bacterial Peritonitis:	C YES ONO UNK			
Previous Abdominal Surgery: *	C YES NO UNK			
Portal Vein Thrombosis: *				

	C YES C NO C UNK
Transjugular Intrahepatic Portacaval Stint Shunt: *	C YES O NO UNK
Incidental Tumor found at time of Transplant:	C YES C NO C UNK
	C Hepatocellular Adenoma
	Hemangioma
If yes, specify tumor type:	Hemangioendothelioma
	<ul><li>Angiomyolipoma</li></ul>
	Bile Duct Cystadenocarcinoma
ii yes, specify tuffor type.	Cholangiocarcinoma
	Hepatocellular Carcinoma
	Hepatoblastoma
	Angiosarcoma
	Other Primary Liver Tumor, Specify
Specify:	
Clinical Information : POST TRANSPLANT	
Juliou iniviliation i Tool Titalioi Eali	

Clinical Information : POST TRANSPLANT			
Pathology Conf. Liver Diag. of Hospital Discharge: *			
Specify:			
Graft Status: *	C Functioning C Failed		
If death is indicated for the recipient, and the death was a result of some other factor unrelated to graft failure, select Functioning.			
Date of Graft Failure:			
Causes of graft failure:			
Primary Graft Failure	G YES G NO G UNK		
Vascular Thrombosis	C YES NO UNK		
Hepatic arterial thrombosis:	C YES NO UNK		
Hepatic outflow obstruction:	YES NO UNK		
Portal vein thrombosis:	YES NO UNK		
Biliary Tract Complication	YES NO UNK		
Hepatitis: DeNovo	YES NO UNK		
Hepatitis: Recurrent	C YES NO UNK		

Recurrent Disease (non-Hepatitis)	C YES ONO UNK			
Acute Rejection	YES NO UNK			
Infection	C YES C NO C UNK			
Other, Specify:				
Discharge Lab Date:				
Total Bilirubin:	mg/dl ST=			
SGPT/ALT:	U/L ST=			
Serum Albumin:	g/dl ST=			
Serum Creatinine:	mg/dl ST=			
INR:	ST=			
Did patient have any acute rejection episodes between transplant and discharge: *	Yes, at least one episode treated with anti-rejection agent Yes, none treated with additional anti-rejection agent No			
Was biopsy done to confirm acute rejection:	<ul><li>Biopsy not done</li><li>Yes, rejection confirmed</li><li>Yes, rejection not confirmed</li></ul>			
Treatment				
Biological or Anti-viral Therapy:				
	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)			
	☐ Other, Specify			
	☐ Valacyclovir (Valtrex)			
Specify:				

Specify:			
Other therapies:	C YES O NO		
	_		
	Photopheresis		
If Yes, check all that apply:	Plasmapheresis		
	☐ Total Lymphoid Irradiation (TLI)		
Immunosuppressive Information			
Are any medications given currently for maintenance or anti-rejection: *	© YES © NO		
Did the patient participate in any clinical research protocol for immunosuppressive medications:	C YES NO		
If Yes, Specify:			
Immunosuppressive Medications			
View Immunosuppressive Medications			
2 5 10 20			
Definitions Of Immunosuppressive Medications			
For each of the immunosuppressive medications listed, select <b>Ind</b> (Induction), <b>Maint</b> (Maintenance) or <b>AR</b> (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. <b>Do not list non-immunosuppressive medications.</b>			
	Ind. Days ST		
Steroids (Prednisone, Methylprednisolone, Solumedrol, Medrol, Decadro			
Atgam (ATG)			
OKT3 (Orthoclone, Muromonab)			

Thymoglobulin			
Simulect - Basiliximab			
Zenapax - Daclizumab			
			1
Azathioprine (AZA, Imuran)			
EON (Generic Cyclosporine)			
Gengraf (Abbott Cyclosporine)			
Other generic Cyclosporine, specify brand:		п	
Neoral (CyA-NOF)			
Sandimmune (Cyclosporine A)		П	
Mycophenolate Mofetil (MMF, Cellcept, RS61443)		П	
Tacrolimus (Prograf, FK506)		п	
Modified Release Tacrolimus FK506E (MR4)			
Sirolimus (RAPA, Rapamycin, Rapamune)			
Myfortic (Mycophenolate Sodium)			
Other Immunosuppressive Medications			
	Ind. Days	ST	Maint AR
Campath - Alemtuzumab (anti-CD52)			
Cyclophosphamide (Cytoxan)			
Leflunomide (LFL, Arava)			
Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)			
Other Immunosuppressive Medication, Specify			
Other Immunosuppressive Medication, Specify			
Rituximab			

Investigational Immunosuppressive Medications			
	Ind. Days	ST	Maint AR
Everolimus (RAD, Certican)			
FTY 720			
Other, Specify			
UNOS View Only			
Comments:			