

## **Pediatric Thoracic - Lung Transplant Recipient Registration 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.

Paginiant Information	
Recipient Information	DOD.
Name:	DOB:
SSN:	Gender:
HIC:	Tx Date:
State of Permanent Residence: *	
Permanent Zip:★	-
Provider Information	
Recipient Center:	
Physician Name: *	
Physician NPI#:*	
Surgeon Name: *	
Surgeon NPI#: *	
Donor Information	
UNOS Donor ID #:	
Donor Type:	
Patient Status	
Primary Diagnosis:*	
Specify:	
Date: Last Seen, Retransplanted or Death★	
	LIVING
But the sky	
Patient Status: *	● DEAD
	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	6 6 6
the transplant admission:	O YES O NO UNK
	IN INTENSIVE CARE UNIT
Medical Condition:*	HOSPITALIZED NOT IN ICU
	NOT HOSPITALIZED
Patient on Life Support: *	© YES © NO
	Extra Corporeal Membrane Oxygenation
	☐ Intra Aortic Balloon Pump
	Prostacyclin Infusion
	Prostacyclin Inhalation
	☐ Intravenous Inotropes
	☐ Inhaled NO
	☐ Ventilator
	☐ Other Mechanism
Specify:	
Functional Status: *	
Functional Status:*	Definite Cognitive delay/impairment
Functional Status: *	© Definite Cognitive delay/impairment
	Probable Cognitive delay/impairment
Functional Status: *  Cognitive Development: *	<ul> <li>Probable Cognitive delay/impairment</li> <li>Questionable Cognitive delay/impairment</li> </ul>
	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment
	<ul> <li>Probable Cognitive delay/impairment</li> <li>Questionable Cognitive delay/impairment</li> </ul>
	<ul> <li>Probable Cognitive delay/impairment</li> <li>Questionable Cognitive delay/impairment</li> <li>No Cognitive delay/impairment</li> <li>Not Assessed</li> </ul>
	Probable Cognitive delay/impairment Questionable Cognitive delay/impairment No Cognitive delay/impairment Not Assessed  Definite Motor delay/impairment
Cognitive Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment
	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment
Cognitive Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment  No Motor delay/impairment
Cognitive Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment
Cognitive Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment  No Motor delay/impairment  No Motor delay/impairment
Cognitive Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment  No Motor delay/impairment  No Motor delay/impairment  Not Assessed  Within One Grade Level of Peers
Cognitive Development: *  Motor Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment  No Motor delay/impairment  No Motor delay/impairment  Not Assessed  Within One Grade Level of Peers  Delayed Grade Level
Cognitive Development: *	<ul> <li>Probable Cognitive delay/impairment</li> <li>Questionable Cognitive delay/impairment</li> <li>No Cognitive delay/impairment</li> <li>Not Assessed</li> <li>Definite Motor delay/impairment</li> <li>Probable Motor delay/impairment</li> <li>Questionable Motor delay/impairment</li> <li>No Motor delay/impairment</li> <li>Not Assessed</li> <li>Within One Grade Level of Peers</li> <li>Delayed Grade Level</li> <li>Special Education</li> </ul>
Cognitive Development: *  Motor Development: *	Probable Cognitive delay/impairment  Questionable Cognitive delay/impairment  No Cognitive delay/impairment  Not Assessed  Definite Motor delay/impairment  Probable Motor delay/impairment  Questionable Motor delay/impairment  No Motor delay/impairment  Not Assessed  Within One Grade Level of Peers  Delayed Grade Level

Academic Activity Level:*		oad in academics due to disease or condition ears old/ High School graduate
Source of Payment:		
Primary: <b>*</b>		
Specify:		
Secondary:		
Clinical Information : PRETRANSPLAN	Т	
Date of Measurement:	<u> </u>	
Height:★	ft. in.	cm ST=
Weight:**	lbs	kg ST=
BMI:	kg/m <sup>2</sup>	
Previous Transplants:		
Previous Transplant Organ	Previous Transplant Date	Previous Transplant Graft Fail Date
The three most recent transplants are listed h 4334 or by emailing unethelpdesk @unos.org.	nere. Please contact the UNet Help Desk to cont	firm more than three previous transplants by calling 800-978-
Viral Detection:		
	Positive	
HIV Serostatus: ★	Negative	
THV Selostatus.	Not Done	
	UNK/Cannot Disclose	e
	Positive	
CMV IgG:★	Not Done	
	UNK/Cannot Disclose	е
	Positive	
CMV IgM: <b>*</b>	Not Done	
	UNK/Cannot Disclose	e
	Positive	
HBV Core Antibody: ★		
The second secon		

		0	UNK	/Can	not	Disc	close					
		0	Posit	ive								
LIDUO C. A. C. Mc		0	Nega	itive								
HBV Surface Antigen: <del>**</del>		0	Not E	Done	•							
		0	UNK	/Can	not	Disc	close					
		0	Posit	ive								
		0	Nega	itive								
HCV Serostatus: <b>*</b>		0	Not [	Oone	•							
		0	UNK	/Can	not	Disc	close					
		0	Posit	ive								
		0	Nega	itive								
EBV Serostatus: *		0	Not E	Oone	•							
		0	UNK	/Can	not	Disc	close					
Most Recent Hemodynamics:									Inotre	nos/\	/aso	dilators:
								ST=				
PA (sys)mm/Hg: *									0	YES	0	NO
PA(dia) mm/Hg: *								ST=	0	YES	0	NO
PA(mean) mm/Hg:≭								ST=		YES		NO
1 Allically mining.								[		ILS		NO
PCW(mean) mm/Hg:*								ST=	0	YES	0	NO
CO L/min:*								ST=	0	YES	0	NO
Most Recent Serum Creatinine: ★							mg/dl	ST=				
Most Recent Total Bilirubin: ★							mg/dl	ST=				
Oxygen Requirement at Rest:*							L/min	ST=				
Chronic Steroid Use:*		0	YES	0	NO	0	UNK					
Pulmonary Status (Give most recent value):												
FVC:*							%predicted:	ST=				
FeV1:*							%predicted:	ST=				
pCO2: *							mm/Hg:	ST=				
Events occurring between listing and transplant:												
Transfusions: <b>*</b>		0	YES	0	NO	0	UNK					
Pulmonary Embolism:*		0	YES	0	NO	0	UNK					
Infection Requiring IV Therapy within 2 wks prior to	o Tx:*						UNK					
					-							

Cerebrovascular Event:	O YES O NO O UNK
Dialysis:*	O YES O NO O UNK
Implantable Defibrillator:	O YES O NO O UNK
Episode of Ventilatory Support: *	O YES O NO O UNK
If yes, indicate most recent timeframe:	C At time of transplant Within 3 months of transplant C >3 months prior to transplant
Tracheostomy:*	O YES O NO O UNK
Prior Thoracic Surgery other than prior transplant:*	O YES O NO O UNK
If yes, number of prior sternotomies:	<ul> <li>Unknown if there were prior sternotomies</li> <li>0</li> <li>1</li> <li>2</li> <li>3</li> <li>4</li> </ul>

	© 5+					
	Unknown number of prior sternotomies					
	Unknown if there were prior thoracotomies					
	O 0					
	© 1					
	C 2					
If yes, number of prior thoracotomies:	<b>○</b> 3					
	€ 4					
	Unknown number of prior thoracotomies					
Prior congenital cardiac surgery:	© YES © NO © UNK					
If yes, palliative surgery:	O YES O NO O UNK					
If yes, corrective surgery:	O YES O NO UNK					
If yes, single ventricular physiology:	O YES O NO O UNK					
	O NO PREVIOUS PREGNANCY					
	1 PREVIOUS PREGNANCY					
	2 PREVIOUS PREGNANCIES					
	3 PREVIOUS PREGNANCIES					
Previous Pregnancies:	4 PREVIOUS PREGNANCIES					
	5 PREVIOUS PREGNANCIES					
	MORE THAN 5 PREVIOUS PREGNANCIES					
	NOT APPLICABLE: < 10 years old					
	UNKNOWN					
Malignancies between listing and transplant: *	C YES C NO C UNK					
This question is NOT applicable for patients receiving living dono	r transplants who were never on the waiting list.					
	Skin Melanoma					
	Skin Non-Melanoma					
	☐ CNS Tumor					
	☐ Genitourinary					
If yes, specify type:	☐ Breast					
	☐ Thyroid					
	☐ Tongue/Throat/Larynx					
	Lung					

	Leukemia/Lymphoma							
	Liver							
	Other, specify							
Specify:								
Clinical Information : TRANSPLANT PROCEDURE								
Multiple Organ Recipient								
Were extra vessels used in the transplant procedure:								
	SINGLE LEFT LUNG							
	SINGLE RIGHT LUNG							
Procedure Type:	BILATERAL SEQUENTIAL LUNG							
Trocedure Type.	EN-BLOC DOUBLE LUNG							
	C LOBE, RIGHT							
	C LOBE, LEFT							
Was this a retransplant due to failure of a previous thoracic graft:	YES NO							
Total Organ Ischemia Time (include cold, warm and anast	comotic time):							
Left Lung:	min ST=							
Right Lung (OR EN-BLOC):	min ST=							
Incidental Tumor found at time of Transplant:	€ YES € NO € UNK							
	Adenoma							
	Carcinoma							
If yes, specify tumor type:	Carcinoid							
ii yes, specily tumor type:	C Lymphoma							
	C Harmartoma							
	Other Primary Lung Tumor, Specify							
Specify:								
Clinical Information : POST TRANSPLANT								
Graft Status:*	Functioning Failed							
If death is indicated for the recipient, and the death was a resu	ult of some other factor unrelated to graft failure, select Functioning.							
Date of Graft Failure:								
	Primary Non-Function							
Primary Cause of Graft Failure:	Acute Rejection							
	•							

	Chronic Rejection/Atherosclerosis					
	C Other, Specify					
Specify:						
Events Prior to Discharge:						
Any Drug Treated Infection:	C YES NO C UNK					
Stroke: *	C YES NO C UNK					
Dialysis:*	C YES NO C UNK					
Cardiac Re-Operation:	C YES NO UNK					
Other Surgical Procedures:	C YES NO C UNK					
	© No					
	C Ventilator support for <= 48 hours					
	C Ventilator support for >48 hours but < 5 days					
Ventilator Support: *	C Ventilator support >= 5 days					
	C Ventilator support, duration unknown					
	C Unknown Status					
Reintubated: *	© YES © NO © UNK					
Permanent Pacemaker:*	C YES NO UNK					
Chest drain >2 weeks:	C YES NO UNK					
Airway Dehiscence:★	C YES C NO C UNK					
	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					
	○ No					
	Biopsy not done					
Was biopsy done to confirm acute rejection:	Yes, rejection confirmed					
	Yes, rejection not confirmed					
Treatment						
Biological or Anti-viral Therapy:	C YES NO Unknown/Cannot disclose					
	☐ Acyclovir (Zovirax)					
	Cytogam (CMV)					
	Gamimune					

	☐ Gammagard						
	☐ Ganciclovir (Cytovene)						
	☐ Valgancyclovir (Valcyte)						
	☐ HBIG (Hepatitis B Immune Globulin)						
If Yes, check all that apply:	☐ Flu Vaccine (Influenza Virus)						
	☐ Lamivudine (Epivir) (for treatment of Hepatitis B)						
	☐ Other, Specify						
	☐ Valacyclovir (Valtrex)						
Specify:							
Specify:							
1, 1, 2							
Other therapies:	€ YES € 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:	© YES © NO						
If Yes, Specify:							
Immunosuppressive Medications							
View Immunosuppressive Medications							
Definitions Of Immunosuppressive Medications							
	nd (Induction), Maint (Maintenance) or AR (Anti-rejection) to indicate all medications that spitalization period, and for what reason. If a medication was not given, leave the associated						
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.							
long-term or intermediate term with a tapering of the dosage un	ions given before, during or after transplant for varying periods of time which may be either till the drug is either eliminated or replaced by another long-term maintenance drug (example: Azathioprine, or Rapamycin). This does not include any immunosuppressive medications						
initial post-transplant period or during a specific follow-up period Atgam, OKT3, or Thymoglobulin). When switching maintenance	uppressive medications given for the purpose of treating an acute rejection episode during the d, usually up to 30 days after the diagnosis of acute rejection (example: Methylprednisolone, e drugs (example: from Tacrolimus to Cyclosporine; or from Mycophenolate Mofetil to ed under AR immunosuppression, but should be listed under maintenance						
	being administered (e.g., new monoclonal antibodies), select Ind, Maint, or AR next to Other fithe medication in the space provided. <b>Do not list non-immunosuppressive medications.</b>						
	Ind. Days ST 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			
	Ind. Days	ST	Maint
Campath - Alemtuzumab (anti-CD52)			
Cyclophophomida (Cytayan)	П		
Cyclophosphamide (Cytoxan)			

Methotrexate (Folex, PFS, Mexate-AQ, Rheumatrex)	Г				
Other Immunosuppressive Medication, Specify					
Rituximab	Г				
Investigational Immunosuppressive Medications					
	Ind	. Days	ST	Maint	AR
Everolimus (RAD, Certican)	Г				
Other Immunosuppressive Medication, Specify					
UNOS View Only					
Comments:					