

Adult Thoracic - Lung 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: ** | |
| | |
| Recipient Center: | |
| Physician Name: ** | |
| Physician NPI#: * | |
| Surgeon Name: * | |
| Surgeon NPI#: ★ | |
| | |
| UNOS Donor ID #: | |
| Donor Type: | |
| | |
| Primary Diagnosis: * | |
| Specify: | |
| Date: Last Seen, Retransplanted or Death ** | |
| | C LIVING |
| Patient Status: * | C DEAD |
| | C 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: | C YES C NO C UNK |
| | IN INTENSIVE CARE UNIT |
| Medical Condition: ★ | C HOSPITALIZED NOT IN ICU |
| | NOT HOSPITALIZED |
| Patient on Life Support: * | C YES NO |
| | |
| | Extra Corporeal Membrane Oxygenation |
| | Intra Aortic Balloon Pump |
| | Prostacyclin Infusion |
| | Prostacyclin Inhalation |
| | Inhaled NO |
| | Ventilator |
| | Other Mechanism |

| Specify: | | |
|----------------------------|--|--|
| Functional Status: * | | |
| Physical Capacity: | No Limitations Limited Mobility Wheelchair bound or more limited Not Applicable (< 1 year old or hospitalized) Unknown | |
| Working for income: ★ | C YES C NO C UNK | |
| If No, Not Working Due To: | | |
| If Yes: | 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: | Within One Grade Level of Peers Delayed Grade Level Special Education Not Applicable < 5 years old/ High School graduate or GED Status Unknown | |

| 0 | | e in academics due to disea ears old/ High School gradu | |
|--------------------------------------|-----------------------|---|---|
| 0 | Not Applicable < 5 ye | | |
| | | ears old/ High School gradu | uate or GED |
| ° | Status Unknown | | |
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| | Tt. III. | cm | ST= |
| | lbs | kg | ST= |
| kg/m ⁻ | | | |
| | | | |
| Previous Transplant I | Date | Previous Transplant G | raft Fail Date |
| ed here. Please contact the nos.org. | UNet Help Desk to co | onfirm more than three prev | ious transplants by calling 800 |
| 0 | Positive | | |
| 0 | Negative | | |
| 0 | Not Done | | |
| 0 | UNK/Cannot Disclos | e | |
| 0 | Positive | | |
| | | | |
| | contact the nos.org. | Previous Transplant Date ad here. Please contact the UNet Help Desk to conso.org. Positive Negative Not Done UNK/Cannot Disclos | Previous Transplant Date Previous Transplant G ad here. Please contact the UNet Help Desk to confirm more than three previous.org. Positive Negative Not Done UNK/Cannot Disclose |

| | 0 | Not Done |
|---------------------------|---|-------------------------|
| | 0 | UNK/Cannot Disclose |
| | 0 | Positive |
| 000U W W | 0 | Negative |
| CMV IgM: ₩ | 0 | Not Done |
| | 0 | UNK/Cannot Disclose |
| | 0 | Positive |
| HBV Core Antibody: ★ | 0 | Negative |
| TIBV GOTE ATTRIBUTY. | 0 | Not Done |
| | 0 | UNK/Cannot Disclose |
| | 0 | Positive |
| HBV Surface Antigen: ★ | 0 | Negative |
| | 0 | Not Done |
| | 0 | UNK/Cannot Disclose |
| | 0 | Positive |
| HCV Serostatus: ★ | 0 | Negative |
| | 0 | Not Done |
| | 0 | UNK/Cannot Disclose |
| | 0 | Positive |
| EBV Serostatus: * | 0 | Negative |
| | 0 | Not Done |
| | 0 | UNK/Cannot Disclose |
| Most Recent Hemodynamics: | | Inotropes/Vasodilators: |
| PA (sys)mm/Hg: * | | ST= C YES NO |

| PA(dia) mm/Hg: * | | ST= | YES NO |
|--|------------------|-----|------------|
| PA(mean) mm/Hg: ★ | | ST= | C YES C NO |
| PCW(mean) mm/Hg: * | | ST= | C YES C NO |
| CO L/min: * | | ST= | C YES C 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: ** | C YES C NO C UNK | | |
| Pulmonary Status (Give most recent value): | | | |
| FVC: * | %predicted: | ST= | |
| FeV1: * | %predicted: | ST= | |
| pCO2: ** | mm/Hg: | ST= | |
| Events occurring between listing and transplant: | | | |
| Transfusions: ★ | O YES O NO O UNK | | |
| Pulmonary Embolism: | O YES O NO UNK | | |
| Infection Requiring IV Therapy within 2 wks prior to Tx: * | C YES C NO UNK | | |
| Cerebrovascular Event: | O YES O NO UNK | | |
| Dialysis: * | C YES C NO C UNK | | |
| Implantable Defibrillator: | C YES C NO UNK | | |
| Prior Cardiac Surgery (non-transplant): ★ | C YES O NO UNK | | |
| If yes, check all that apply: | | | |

| | Valve Replacement/Repair |
|---|-------------------------------|
| | Congenital |
| | Left Ventricular Remodeling |
| | Other, specify |
| Specify: | |
| Prior Lung Surgery (non-transplant): ★ | C YES C NO UNK |
| | Pneumoreduction |
| | Pneumothorax Surgery-Nodule |
| | Pneumothorax Decortication |
| If you about all that apply | Lobectomy |
| If yes, check all that apply: | Pneumonectomy |
| | Left Thoracotomy |
| | Right Thoracotomy |
| | Other, specify |
| Specify: | |
| Episode of Ventilatory Support: ** | C YES C NO C UNK |
| | At time of transplant |
| If yes, indicate most recent timeframe: | Within 3 months of transplant |
| | >3 months prior to transplant |
| Tracheostomy: * | C YES C NO C UNK |
| | С |
| | NO PREVIOUS PREGNANCY |
| Previous Pregnancies: | 1 PREVIOUS PREGNANCY |
| | 2 PREVIOUS PREGNANCIES |

| | 0 | 3 PREVIOUS PREGNANCIES |
|--|----------|---|
| | 0 | 4 PREVIOUS PREGNANCIES |
| | 0 | 5 PREVIOUS PREGNANCIES |
| | 0 | MORE THAN 5 PREVIOUS PREGNANCIES |
| | 0 | |
| | 0 | NOT APPLICABLE: < 10 years old |
| | | UNKNOWN |
| | | |
| Malignancies between listing and transplant: * | 0 | YES NO UNK |
| This question is NOT applicable for patients receiving living d | onor tra | ansplants 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 |
| | П | Leukemia/Lymphoma |
| | П | Liver |
| | | Other, specify |
| Specify: | | |
| 1 | | |
| Multiple Organ Recipient | | |
| Ware outra vessels used in the transplant precedure: | | |
| Were extra vessels used in the transplant procedure: Vessel Donor ID: | | |

| | SINGLE LEFT LUNG |
|--|---|
| | SINGLE RIGHT LUNG |
| Procedure Type: | © BILATERAL SEQUENTIAL LUNG |
| | EN-BLOC DOUBLE LUNG |
| | C LOBE, RIGHT |
| | C LOBE, LEFT |
| Was this a retransplant due to failure of a previous thoracic graft: | C _{YES} C _{NO} |
| Total Organ Ischemia Time (include cold, warm and anas | stomotic time): |
| Left Lung: | min ST= |
| Right Lung (OR EN-BLOC): | min ST= |
| Incidental Tumor found at time of Transplant: | C YES C NO UNK |
| | C Adenoma |
| | Carcinoma |
| If yes, specify tumor type: | Carcinoid |
| | Lymphoma |
| | C Harmartoma |
| | Other Primary Lung Tumor, Specify |
| Specify: | |
| | |
| Graft Status: ★ | C Functioning C Failed |
| If death is indicated for the recipient, and the death was a res Date of Graft Failure: | sult of some other factor unrelated to graft failure, select Functioning. |
| Date of Statt Latitute. | |

| | О | Primary Non-Function |
|---------------------------------|---|---|
| Primary Cause of Graft Failure: | 0 | Acute Rejection |
| Timaly dauge of Grant andre. | О | Chronic Rejection/Atherosclerosis |
| | 0 | Other, Specify |
| Specify: | | |
| | | |
| Events Prior to Discharge: | | |
| Any Drug Treated Infection: | 0 | YES O NO UNK |
| Stroke: * | 0 | YES NO UNK |
| Dialysis: * | 0 | YES NO UNK |
| Cardiac Re-Operation: | 0 | YES NO UNK |
| Other Surgical Procedures: | 0 | YES NO UNK |
| | 0 | No |
| | 0 | Ventilator support for <= 48 hours |
| Ventilator Support: ★ | 0 | Ventilator support for >48 hours but < 5 days |
| ventilator Support. | 0 | Ventilator support >= 5 days |
| | 0 | Ventilator support, duration unknown |
| | 0 | Unknown Status |
| Reintubated: * | 0 | YES NO UNK |
| Permanent Pacemaker: ★ | 0 | YES O NO UNK |
| Chest drain >2 weeks: | 0 | YES O NO UNK |
| Airway Dehiscence: * | 0 | YES O NO UNK |

| | O | Yes, at least one episode treated with anti-rejection agent |
|--|---|---|
| Did patient have any acute rejection episodes between transplant and discharge: ** | 0 | Yes, none treated with additional anti-rejection agent |
| | 0 | No |
| | 0 | Biopsy not done |
| Was biopsy done to confirm acute rejection: | 0 | Yes, rejection confirmed |
| | 0 | Yes, rejection not confirmed |
| - | | |
| | | |
| Biological or Anti-viral Therapy: | 0 | 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) |
| | | Other, Specify |
| | | Valacyclovir (Valtrex) |
| Specify: | | |
| Specify: | | |
| | | |
| Other therapies: | 0 | YES NO |

| | | Photopheresis |
|--|---|----------------------------------|
| If Yes, check all that apply: | | Plasmapheresis |
| | | Total Lymphoid Irradiation (TLI) |
| | | |
| Are any medications given currently for maintenance or anti-rejection: ** | 0 | YES NO |
| Did the patient participate in any clinical research protocol for immunosuppressive medications: | 0 | YES NO |
| If Yes, Specify: | | |
| | | |
| View Immunosuppressive Medications | | |
| Definitions Of 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. <u>Do not list non-immunosuppressive medications.</u>

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,Decadron) | 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) | I | | | | |
| | | | | | |
| Campath - Alemtuzumab (anti-CD52) | Ind. | Days | ST | Maint | AR |
| 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 | | | | | |

С