

**Extracorporeal Photopheresis (ECP) for the Management of Progressive Bronchiolitis Obliterans Syndrome (BOS) in Medicare-Eligible Recipients of Lung Allografts**

**DEMOGRAPHICS/MEDICAL HISTORY - Case Report Form (CRF)**

**Please include the following types of source documents:**

1. History and Physical Or Consultation Note
2. Operative Report of Transplant Procedure
3. Pulmonary Function Test Reports (for each FEV-1 submitted)

Source Document Type	Document Name	Submission Date
Select Source Document Type	<input type="text" value="History and Physical or Consultation Note"/>	<input type="text"/>
Attach Source Document:	<input type="text"/>	<input type="button" value="Browse..."/> <input type="button" value="Upload"/>
<input type="button" value="Save"/>	<input type="button" value="Submit"/>	

**Date of Participant Enrollment in Study:**

**SECTION A. Demographic Information**

1. Age:

2. Gender: ☐ Male ☐ Female

3. Race:

- ☐ White
- ☐ Black or African-American
- ☐ Asian
- ☐ American Indian
- ☐ Alaska Native
- ☐ Native Hawaiian or Other Pacific Islander
- ☐ Other
- ☐ No Response


4. Are you of Hispanic or Latino origin: ☐ YES ☐ NO

## **SECTION B. Previous Medical History and Pulmonary Function Testing Results**

5. Please provide a description of the underlying disease necessitating lung transplantation:

- ☐ Chronic Obstructive Pulmonary Disease (COPD) including Emphysema
- ☐ Interstitial Lung Disease
- ☐ Cystic Fibrosis
- ☐ Pulmonary Hypertension
- ☐ Alpha 1-Antitrypsin Deficiency Emphysema
- ☐ Replacing previously transplanted lung that failed
- ☐ Other

If other, please describe:

6. Date of lung transplantation:  

7. Operation performed:

- ☐ Single
- ☐ Bilateral
- ☐ Heart-lung
- ☐ Other

If other, please describe:

8. Weight at time of transplant:  kilograms

9. Height:  Centimeters

10. Medical History - Co-Morbid Conditions at any time prior to enrollment:

- |                           |                          |  |
|---------------------------|--------------------------|--|
| <input type="radio"/> YES | <input type="radio"/> NO | Hypertension   |
| <input type="radio"/> YES | <input type="radio"/> NO | Diabetes   |
| <input type="radio"/> YES | <input type="radio"/> NO | GERD - if yes, treatment: <input type="radio"/> medical therapy <input type="radio"/> fundoplication       |
| <input type="radio"/> YES | <input type="radio"/> NO | High Cholesterol   |
| <input type="radio"/> YES | <input type="radio"/> NO | Current Smoker   |
| <input type="radio"/> YES | <input type="radio"/> NO | Previous Smoker  |
| <input type="radio"/> YES | <input type="radio"/> NO | Coronary Artery Disease - if yes: <input type="radio"/> angina <input type="radio"/> myocardial infarction |
| <input type="radio"/> YES | <input type="radio"/> NO | Congestive Heart Failure   |
| <input type="radio"/> YES | <input type="radio"/> NO | Chronic Kidney Disease   |
| <input type="radio"/> YES | <input type="radio"/> NO | Stroke   |
| <input type="radio"/> YES | <input type="radio"/> NO | Neurologic Disorder  |
| <input type="radio"/> YES | <input type="radio"/> NO | Other Active Conditions  |

If other active conditions, please describe:

11. Has the patient received any anti-platelet drug(s) within the last six months:

- ☐ YES    ☐ NO    Anti-Thrombotic
- ☐ YES    ☐ NO    Anti-Platelet Agent

12. Check all drugs that were previously used as maintenance immunosuppression and/or BOS prevention in this participant:

- ☐ YES    ☐ NO    Tacrolimus
- ☐ YES    ☐ NO    Prednisone
- ☐ YES    ☐ NO    Alemtuzumab
- ☐ YES    ☐ NO    Sirolimus (Rapamycin)
- ☐ YES    ☐ NO    Everolimus
- ☐ YES    ☐ NO    Cyclosporine A
- ☐ YES    ☐ NO    Methotrexate
- ☐ YES    ☐ NO    Macrolide Antibiotic, Azithromycin
- ☐ YES    ☐ NO    Mycophenolate Mofetil (Cellcept or Myfortic)
- ☐ YES    ☐ NO    Anti-Thymocyte Globulin - ATG (Thymoglobulin or Atgam)

13. Check all drugs that were previously used as active treatment of BOS in this participant:

- ☐ YES    ☐ NO    Tacrolimus
- ☐ YES    ☐ NO    Prednisone
- ☐ YES    ☐ NO    Alemtuzumab
- ☐ YES    ☐ NO    Sirolimus (Rapamycin)
- ☐ YES    ☐ NO    Everolimus
- ☐ YES    ☐ NO    Cyclosporine A
- ☐ YES    ☐ NO    Methotrexate
- ☐ YES    ☐ NO    Macrolide Antibiotic, Azithromycin
- ☐ YES    ☐ NO    Mycophenolate Mofetil (Cellcept or Myfortic)
- ☐ YES    ☐ NO    Anti-Thymocyte Globulin - ATG (Thymoglobulin or Atgam)
- ☐ YES    ☐ NO    Total Lymphoid Irradiation


14. Has the participant received prednisone therapy within the last 6 months? ☐ YES ☐ NO

If yes, daily starting dose:  mg

15. During the last 6 months, has the patient required a prednisone dose escalation for any period of greater than 5 days? ☐ YES ☐ NO

If yes, to what daily dose:  mg

Estimate average daily dose over 6 months:  mg

16. Date of diagnosis of post-transplantation BOS:  

17. Post-transplant BOS stage at diagnosis: ☐ 0 ☐ 0-p ☐ 1 ☐ 2 ☐ 3

Stage	Definition
BOS Stage 0	FEV1 > 90% and FEF 25%-75% > 75% of baseline
BOS Stage 0-p	FEV1 = 81% to 90% of baseline and or FEF 25%-75% 75% of baseline
BOS Stage 1	FEV1 = 66% to 80% of baseline
BOS Stage 2	FEV1 = 51% to 65% of baseline
BOS Stage 3	FEV1 less than or equal to 50% of baseline

18. Please provide the baseline pulmonary function testing results using the ISHLT definition = the average of the two highest FEV1 measurements obtained 3 weeks apart after transplantation. Also please provide the results of the two component PFT assessments used for this calculation:

a. Baseline FEV1 (pre-bronchodilator)(mean FEV1 values below):  liters


b. First component assessment:

i. Date:  

ii. FEV1 (pre-bronchodilator):  liters

iii. FVC (pre-bronchodilator):  liters

c. Second component assessment:

i. Date:  

ii. FEV1 (pre-bronchodilator):  liters

iii. FVC (pre-bronchodilator):  liters

## **SECTION C. Clinical Status at or Within One Week of Enrollment**

19. Donor specific antibody at time of study enrollment? ☐ YES ☐ NO

20. Is the participant on any anticoagulant or anti-platelet drugs? ☐ YES ☐ NO

If yes, list drugs:

Name drug 1:

Name drug 2:

Name drug 3:

21. Date when the following baseline vital signs were obtained (values should be obtained within one week prior to enrollment):  

22. Weight:  kilograms

23. Blood pressure: systolic  mmHg diastolic  mmHg

24. Heart rate:  beats per minute

25. Respiratory rate:  breaths per minute

26. Resting oxygen saturation:  %

27. Is the participant receiving supplemental oxygen? ☐ YES ☐ NO

If yes, how much?  liters/minute

28. Comments:

Save

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