

## Blood and Marrow Transplant Clinical Trials Network

### Acute GVHD Form (GVH)

Web Version: 1.0; 10.14; 12-09-16

Segment (PROTSEG): A

Visit Number (VISNO):

1. Date of staging:(STAGEDT)  (mm/dd/yyyy)  
 Start of GVHD Assessment Period: (GVASSTDY)  (mm/dd/yyyy)  
 End of GVHD Assessment Period:(GVASENDT)  (mm/dd/yyyy)

*The assessment for which you are entering data must have taken place within the above dates. If the patient was not seen during the assessment period specified above, please exit the form and request an exception for this form.*

2. Immunosuppressant (prophylaxis) received:(IMMUNORC) 
  - 0 - Prednisone
  - 1 - Cyclosporine
  - 2 - Tacrolimus
  - 3 - Not taken during assessment
3. Record most recent blood level of immunosuppressant (prophylaxis):  
(TROUGHLV)  (xxx.x) ng/mL
4. Record date blood sample obtained:(TROUGHDT)  (mm/dd/yyyy)

**Record the highest level of organ abnormalities, the etiologies contributing to the abnormalities and any biopsy results during the assessment period.**

5. Skin abnormalities:(GVHSKINA) 
  - 0 - No Rash
  - 1 - Maculopapular Rash, <25% of Body Surface
  - 2 - Maculopapular Rash, 25-50% of Body Surface
  - 3 - Generalized Erythroderma
  - 4 - Generalized Erythroderma with Bullus Formation and Desquamation

6. Skin etiologies:

<b>GVHD</b>	<b>Drug Reaction</b>	<b>Conditioning Regimen Toxicity</b>
(SETGVHD) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(SETDRGRX) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(SETCRTOX) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No
<b>Infection</b>	<b>Other</b>	
(SETINFCT) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(SETOTHER) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	

Specify other skin etiologies:(GVHSKNSP)

7. Skin biopsy for GVHD:(GVHSKINB) 
  - 1 - Positive
  - 2 - Negative
  - 3 - Equivocal
  - 4 - Not Done

8. Upper GI abnormalities:(GVHUPGIA) 
  - 0 - No Protracted Nausea and Vomiting
  - 1 - Persistent Nausea, Vomiting or Anorexia

9. Upper intestinal tract etiologies:

<b>GVHD</b>	<b>Drug Reaction</b>	<b>Conditioning Regimen Toxicity</b>
(UGIETGVH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(UGIETDRG) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(UGIETCON) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No
<b>TPN</b>	<b>Infection</b>	<b>Other</b>
(UGIETTPN) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(UGIETINF) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(UGIETOTH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No

Specify other upper intestinal tract etiologies:(UGIETSPC)

10. Upper intestinal tract biopsy for GVHD:(UGBIORS)

- 1 - Positive
- 2 - Negative
- 3 - Equivocal
- 4 - Not Done

11. Lower GI abnormalities:(GVHINTA)

- 0 - No Diarrhea
- 1 - Diarrhea Less Than or Equal to 500 mL/day or <280 mL/m<sup>2</sup>
- 2 - Diarrhea >500 but Less Than or Equal to 1000 mL/day or 280-555 mL/m<sup>2</sup>
- 3 - Diarrhea >1000 but Less Than or Equal to 1500 mL/day or 556-833 mL/m<sup>2</sup>
- 4 - Diarrhea >1500 mL/day or >833 mL/m<sup>2</sup>
- \*Additional Options Listed Below

Use mL/day for adult patients and mL/m<sup>2</sup> for pediatric patients

12. Lower intestinal tract etiologies:

<b>GVHD</b>	<b>Drug Reaction</b>	<b>Conditioning Regimen Toxicity</b>
(LGIETGVH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LGIETDRG) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LGIETCON) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No
<b>TPN</b>	<b>Infection</b>	<b>Other</b>
(LGIETTPN) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LGIETINF) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LGIETOTH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No

Specify other lower intestinal tract etiologies:(LGIETSPC)

13. Lower intestinal tract biopsy for GVHD:(LGIBIORS)

- 1 - Positive
- 2 - Negative
- 3 - Equivocal
- 4 - Not Done

14. Liver abnormalities:(GVHLIVRA)

- 0 - Bilirubin <2.0 mg/dL
- 1 - Bilirubin 2.0-3.0 mg/dL
- 2 - Bilirubin 3.1-6.0 mg/dL
- 3 - Bilirubin 6.1-15.0 mg/dL
- 4 - Bilirubin >15.0 mg/dL

15. Liver etiologies:

<b>GVHD</b>	<b>Drug Reaction</b>	<b>Conditioning Regimen Toxicity</b>	<b>TPN</b>
(LIVETGVH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LIVETDRG) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LIVETCND) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LIVETTPN) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No
<b>Infection</b>	<b>VOD</b>	<b>Other</b>	
(LIVETINF) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LIVETVOD) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	(LIVETOTH) <input type="checkbox"/> 1 - Yes <input type="checkbox"/> 2 - No	

Specify other liver etiologies:(GVHLIVRS)

16. Liver biopsy for GVHD:(GVHLIVRB)

- 1 - Positive
- 2 - Negative
- 3 - Equivocal
- 4 - Not Done

17. Was any treatment of GVHD modified during this assessment period?  
(GVHTHERP)

- 1 - Yes
- 2 - No

*This only applies to TREATMENT for GVHD. If GVHD prophylaxis was the only modification during this assessment period, this question should be answered "2 - No".*

18. If yes, specify agent name:(GVHAGENT)

- 1 - CSA
- 2 - FK506
- 3 - Topical Steroids
- 4 - Prednisone
- 5 - ATG
- \*Additional Options Listed Below

Specify other agent:(GVHAGNSP)

19. Indicate treatment modification:(GVHTRMOD)

- 1 - Started
- 2 - Stopped
- 4 - Tapered
- 5 - Increased

Comments:(GVHCOMM)

## Additional Selection Options for GVH

### Lower GI abnormalities:

5 - Severe Abdominal Pain with or without Ileus, or Stool with Frank Blood or Melena

### If yes, specify agent name:

6 - MMF

7 - Daclizumab

8 - Methylprednisolone

9 - Other