Forms Now Available in FormsNet3

Audience: International and Domestic Data Managers and Medical Directors

Revised Forms
The CIBMTR is excited to announce a new set of revised forms have been released and are now in production as Friday, January 22, 2021. The forms include:

<table>
<thead>
<tr>
<th>Forms</th>
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<tbody>
<tr>
<td>2400 R8 – Pre-Transplant Essential Data</td>
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<tr>
<td>4000 R7 – Cellular Therapy Essential Data Pre-Infusion</td>
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<tr>
<td>4003 R4 – Cellular Therapy Product</td>
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<tr>
<td>4006 R5 – Cellular Therapy Infusion</td>
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<tr>
<td>4100 R6 – Cellular Therapy Essential Data Follow-Up</td>
</tr>
<tr>
<td>3501 R2 – Pregnancy Form</td>
</tr>
<tr>
<td>2801 R5 – Request for Recipient Transfer</td>
</tr>
</tbody>
</table>

Release Information
These forms have been revised to ensure that the forms are capturing currently relevant data, particularly where new definitions or tests have been accepted for use in the field.

The blank forms are available on the Data Collection Forms page and the summary for the Release Highlights are available on the Upcoming Releases page. The corresponding Forms Instruction Manual sections for these forms are updated and available in the CIBMTR Forms Instruction Manual. If your center has feedback regarding the forms or manuals, please contact CIBMTR Center Support.

Registry Donor ID Update
Each of the forms listed below has the field for “Non-NMDP unrelated donor ID” updated to “Registry donor ID” per WMDA (World Marrow Donor Association) guidelines. There are no changes to the intent or how we collect the data, only a name change.

- 2400 R8 – Pre-Transplant Essential Data
- *2100 R6 – Post-HCT Follow-up Data
- *2004 R5 – Infectious Disease Markers
- *2005 R7 – Confirmation of HLA Typing
- *2006 R5 – HCT Infusion
- *2450 R5 – Post-TED
- 4000 R7 – Cellular Therapy Essential Data Pre-Infusion
- 4003 R4 – Cellular Therapy Product
- *3001 R3 – Adverse Event Form
- *3010 R3 – Product Complaint Form

*indicates no form revision required
Fillable Request for Recipient Transfer Form (2801 R5)
The form is now a fillable, interactive PDF document that allows for simplified completion by eliminating the need to print or scan the document. To complete the fillable document for transfer scenarios, follow the steps listed below:

1. Navigate to the Data Collection Forms page here.

2. Select the “2801 (PDF-RF).

3. Download the document.

4. Further instructions on how to complete the form are included on the first page of the document.

Form Pairings
During the revision process, changes were made to the forms with the intent that they would be completed and paired together. The following updates were made on the center’s behalf to maintain data integrity and quality. **No action is required by the center.**

Forms in DUE status:
- Cellular Therapy Product (4003) – will remain as R3 and **not** update to the new R4
- Cellular Therapy Essential Infusion (4006) – will remain as R4 and **not** update to the new R5

Consent Tool
FormsNet3 now captures CIBMTR research database consent for recipients within a new tool. Forms 2400 and 4000 no longer include questions capturing research database consent. Outcomes expected from this change are to:
- Support capturing consent and recipient contact information pre-infusion, when possible.
- Allow the CIBMTR Survey Research Group (SRG) to approach patients for pre-infusion patient-reported outcomes (PRO) surveys for clinical research.

After a CRID has been created for the recipient, a new link to the consent tool will be available to report the recipient’s consent. If a consent response ("yes", "no", or "not approached") has not been reported for a recipient, users will not be able to complete the forms 2400/2402 or 4000 until a consent is reported. For existing recipients, the recipient’s latest consent was moved to the consent tool when possible. There are some circumstances where consent will not be moved (e.g., the forms 2400/2402 are in SVD status, “not applicable” was reported on the form 4000, etc.). If you find the edit form icon for forms 2400/2402 or 4000 appears disabled after the release, please report consent within the consent tool. The edit icon will then enable and allow for form completion. Instructions with an FAQ for the consent tool are located in the Data Management Guide.

If you have any questions regarding the upcoming release, please contact CIBMTR Center Support.