



Cellular Therapy Infusion

Registry Use Only

Sequence Number:

Date Received:

CIBMTR Center Number: _____

CIBMTR Research ID: _____

Event date: ____-____-____
 YYYY MM DD

Product Infusion

1. Specify the identifiers associated with this cell product (*check all that apply*)

- ☐ Cell product ID – ***Go to question 2***
- ☐ Batch number– ***Go to question 3***
- ☐ Lot number – ***Go to question 4***

2. Cell product ID: _____

3. Batch number: _____

4. Lot number: _____

5. Date of this product infusion: _____ — _____ — _____
 YYYY MM DD

6. Was the entire volume of product infused?

- ☐ Yes – **Go to question 9**
- ☐ No– **Go to question 7**

7. Specify what happened to the reserved portion (*check all that apply*)

- ☐ Discarded – **Go to question 9**
- ☐ Cryopreserved for future use – **Go to question 9**
- ☐ Research – **Go to question 9**
- ☐ Training or Quality Control – **Go to question 9**
- ☐ Other fate – **Go to question 8**

8. Specify other fate:

9. Specify the route of product infusion

- ☐ Intravenous – ***Go to question 13***
- ☐ Intramedullary – ***Go to question 13***
- ☐ Intraperitoneal – ***Go to question 13***
- ☐ Intra arterial - ***Go to question 13***
- ☐ Intramuscular - ***Go to question 13***
- ☐ Intrathecal / Intra-ventricular - ***Go to question 13***
- ☐ Intraorgan - ***Go to question 11***
- ☐ Locally in the tissue - ***Go to question 13***
- ☐ Other route of infusion - ***Go to question 10***

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10. Specify other route of infusion: _____ - **Go to question 13**

11. Specify the site of intraorgan administration of cells

- ☐ Bone - **Go to question 13**
- ☐ Heart - **Go to question 13**
- ☐ Liver - **Go to question 13**
- ☐ Pancreas - **Go to question 13**
- ☐ Kidney - **Go to question 13**
- ☐ Brain - **Go to question 13**
- ☐ Lung - **Go to question 13**
- ☐ Other site - **Go to question 12**

12. Specify other site: _____

Report the total number of cells (not cells per kilogram) contained in the product administered.

13. Total number of cells administered

- ☐ Known— **Go to question 14**
- ☐ Unknown— **Go to question 15**

14. Total number of cells administered: _____ • _____ x 10 _____

15. Specify the cell type(s) administered (*check all that apply*)

Lymphocytes

- ☐ Lymphocytes (unselected) – **Go to question 16**
- ☐ CD4+ lymphocytes – **Go to question 17**
- ☐ CD8+ lymphocytes – **Go to question 18**
- ☐ Regulatory T-cells (TREG) – **Go to question 19**

Other Cells

- ☐ Mesenchymal stromal stem cells (MSCs) – **Go to question 20**
- ☐ Unspecified mononuclear cells – **Go to question 21**
- ☐ Other cell type – **Go to question 22**

16. Total number of lymphocytes (unselected): _____ • _____ x 10 _____

17. Total number of CD4+ lymphocytes: _____ • _____ x 10 _____

18. Total number of CD8+ lymphocytes: _____ • _____ x 10 _____

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19. Total number of TREG cells: _____ • _____ x 10 _____
20. Total number of MSCs: _____ • _____ x 10 _____
21. Total number of unspecified mononuclear cells: _____ • _____ x 10 _____
22. Specify other cell type: _____
23. Total number of other cells: _____ • _____ x 10 _____

Concomitant Therapy

24. Did the recipient receive concomitant therapy?

- ☐ Yes – **Go to question 25**
- ☐ No – **Go to End of Form**

Copy and complete questions 25 – 27 to report each concomitant therapy received.

25. Specify start date: _____

YYYY MM DD

26. Specify drug

- ☐ Atezolizumab (Tecentriq®) – **Go to End of Form**
- ☐ Avelumab (Bavencio®) – **Go to End of Form**
- ☐ Durvalumab – **Go to End of Form**
- ☐ IL-2 – **Go to End of Form**
- ☐ IL-15 – **Go to End of Form**
- ☐ Ipilimumab (Yervoy®) – **Go to End of Form**
- ☐ Lenalidomide (Revlimid®) – **Go to End of Form**
- ☐ Nivolumab (Opdivo®) – **Go to End of Form**
- ☐ Pembrolizumab (Keytruda) – **Go to End of Form**
- ☐ Pomalidomide – **Go to End of Form**
- ☐ Other - **Go to question 27**

27. Specify other drug: _____

Copy and complete questions 25 – 27 to report each concomitant therapy received.