**Key Fields**

- **Sequence Number:**
- **Date Received:** __ __ __ __ - __ __ __ __
- **CIBMTR Center Number:**
- **CIBMTR Research ID:**
- **Event date:** __ __ __ __ - __ __ __ __

### Inotuzumab Ozogamicin (Besponsa™)

#### Questions: 1 - 14

1. Did the recipient receive more than one cycle of Inotuzumab ozogamicin? (Besponsa™) (1 cycle = 3 doses)
   - Yes
   - No

2. Number of cycles: __________________________

#### Cycle(s) (1)

Questions: 3 - 14

3. Date of first dose for cycle
   - Known
   - Unknown

4. Date of first dose for cycle: __ __ __ __ - __ __ __ __

5. Date of last dose for cycle
   - Known
   - Unknown

6. Date of last dose for cycle: __ __ __ __ - __ __ __ __

7. Combined dose per cycle (e.g., if patient received 3 doses in cycle 1 at 0.8 mg in day 1, 0.5 in days 8 and 14 then total dose is 1.8 mg)
   - Known
   - Unknown

8. Dose: __________________________ mg/m²

9. Were three doses given in this cycle?
   - Yes
   - No

10. Best response to this cycle of therapy
    - Complete remission (CR) - All of the following response criteria without progression for at least four weeks: < 5% blasts in the bone marrow, no blasts with Auer rods, no extramedullary disease (e.g., central nervous system or soft tissue involvement), ANC of ≥ 1,000/µL, Platelets ≥ 100,000/µL.
    - Complete remission with incomplete hematologic recovery (CRi) - All CR criteria except for residual neutropenia (< 1000/µL) and/or thrombocytopenia (< 100,000/µL)
    - No complete remission

11. Was recipient MRD negative following this cycle of therapy?
    - Yes
    - No
    - Not done

12. MRD method of detection
    - Known
    - Unknown

    13. Minimal residual disease (MRD) testing method
        - Flow cytometry
        - Next generation sequencing (NGS)
        - Polymerase chain reaction (PCR)

14. MRD
    - Positive
    - Negative

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First Name: __________________________

Last Name: __________________________

E-mail address: __________________________

Date: __ __ __ __ - __ __ __ __

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