Form 2541 R1.0: Inotuzumab Ozogamicin (Besponsa™) Supplemental Data Collection Form Center **Key Fields** Sequence Number: 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^{T M}) (1 cycle = 3 doses) C Yes C No 2 Number of cycles: Cycle(s) (1) Questions: 3 - 14 Report the start / stop date of each cycle of Inotuzumab ozogamicin (BesponsaTM) 3 Date of first dose for cycle C Known C Unknown 4 Date of first dose for cycle: 5 Date of last dose for cycle C Known C 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) C Known C Unknown 8 Dose: mg/m² **9** Were three doses given in this cycle? C Yes C No 10 Best response to this cycle of therapy Complete - 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 remission extramedullary disease (e.g., central nervous system or soft tissue involvement), ANC of ≥ 1,000/µL, Platelets ≥ 100,000/µL. (CR) 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? C Yes C No C Not done 12 MRD method of detection C Known C Unknown 13 Minimal residual disease (MRD) testing method Flow cytometry Next generation sequencing (NGS) Polymerase chain reaction (PCR) **14** MRD

Positive Negative

First Name: _____ Last Name: ____ E-mail address: