

Form 2143 R3.0: Multiple Sclerosis Post-HSCT Data

Center:

CRID:

Key Fields

Sequence Number: \_\_\_\_\_

Date Received: \_\_\_\_-\_\_\_\_-\_\_\_\_

CIBMTR Center Number: \_\_\_\_\_

CIBMTR Recipient ID: \_\_\_\_\_

EBMT Center Identification Code (CIC): \_\_\_\_\_

Today's Date: \_\_\_\_-\_\_\_\_-\_\_\_\_

Date of HSCT for which this form is being completed: \_\_\_\_-\_\_\_\_-\_\_\_\_

HSCT type: (check all that apply)

- ☐ Autologous
- ☐ Allogeneic, unrelated
- ☐ Allogeneic, related
- ☐ Syngeneic (identical twin)

Product type: (check all that apply)

- ☐ Marrow
- ☐ PBSC
- ☐ Cord blood
- ☐ Other product

Specify: \_\_\_\_\_

Visit:

- ☐ 100 day ☐ 6 months ☐ 1 year ☐ 2 years ☐ > 2 years,

Specify: \_\_\_\_\_

Multiple Sclerosis Post-HSCT Data

Questions: 1 - 58

1 Date of actual contact with the recipient to determine medical status for this follow-up report: \_\_\_\_-\_\_\_\_-\_\_\_\_

Disease Relapses or Progression of Disability Post-HSCT

2 Were there any relapses of multiple sclerosis (MS) since the date of the last report?

- ☐ yes ☐ no

3 Specify the date of first relapse since the date of the last report: \_\_\_\_-\_\_\_\_-\_\_\_\_

4 Specify the number of relapses since the date of the last report: \_\_\_\_\_ ☐ number unknown

5 Did the recipient experience continuous progression of MS since the date of the last report?

- ☐ yes ☐ no ☐ Unknown

6 Was a MRI scan of the brain performed since the date of the last report?

- ☐ yes ☐ no ☐ Unknown

7 Date of most recent MRI: \_\_\_\_-\_\_\_\_-\_\_\_\_ ☐ Date unknown

8 Was disease relapse / progression detected on the MRI findings?

- ☐ yes ☐ no

Specify which MRI findings indicated disease relapse / progression:

9 New gadolinium-enhancing lesions

- ☐ yes ☐ no ☐ Unknown

10 Enlarging T2 lesions

- ☐ yes ☐ no ☐ Unknown

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**11** Radiology report states worsening of MS associated findings

☐ yes ☐ no ☐ Unknown

**12** Were the Kurtzke Functional Systems Scores (FSS) assessed by a neurologist since the date of the last report?

☐ yes ☐ no ☐ Unknown

**13** Pyramidal score: \_\_\_\_\_ ☐ score unknown

**14** Cerebellar score: \_\_\_\_\_ ☐ score unknown

**15** Brainstem score: \_\_\_\_\_ ☐ score unknown

**16** Sensory score: \_\_\_\_\_ ☐ score unknown

**17** Bowel and bladder score: \_\_\_\_\_ ☐ score unknown

**18** Visual score: \_\_\_\_\_ ☐ score unknown

**19** Cerebral score: \_\_\_\_\_ ☐ score unknown

**20** Other function score: \_\_\_\_\_ ☐ score unknown

**21** Specify other function: \_\_\_\_\_

**22** Specify the Kurtzke Expanded Disability Status Scale (EDSS): \_\_\_\_\_ ☐ EDSS unknown

**23** Specify date of EDSS assessment: \_\_\_\_-\_\_\_\_-\_\_\_\_ ☐ Date unknown

## Post-HSCT Treatment for Multiple Sclerosis

**24** Did the recipient receive any treatment for MS since the date of the last report?

☐ yes ☐ no ☐ Unknown

**Specify the reason(s) for initiating post-HSCT therapy:**

**25** Continued therapy from pre-HSCT

☐ yes ☐ no

**26** Planned therapy per protocol

☐ yes ☐ no

**27** Disease relapse

☐ yes ☐ no

**28** Disease progression

☐ yes ☐ no

**29** Other reason

☐ yes ☐ no

**30** Specify other reason for initiating therapy: \_\_\_\_\_

**31** What was the date the first therapy started? \_\_\_\_-\_\_\_\_-\_\_\_\_

**Indicate all treatments for MS that the recipient received since the date of the last report:**

**32** Alemtuzumab (Campath)

☐ yes ☐ no ☐ Unknown

**33** Azathioprine (Azasan, Imuran)

☐ yes ☐ no ☐ Unknown

**34** Belimumab (LymphoStat-B)

☐ yes ☐ no ☐ Unknown

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## 35 Cladribine (2-CdA, Leustatin)

☐ yes ☐ no ☐ Unknown

## 36 Corticosteroids (chronic use, not to treat acute relapse)

☐ yes ☐ no ☐ Unknown

## 37 Corticosteroids (to treat acute relapse)

☐ yes ☐ no ☐ Unknown

## 38 Cyclophosphamide (CTX, Cytosan, Neosar)

☐ yes ☐ no ☐ Unknown

## 39 Daclizumab (Zenapax, anti-CD25)

☐ yes ☐ no ☐ Unknown

## 40 Fingolimod (FTY720)

☐ yes ☐ no ☐ Unknown

## 41 Fumarate (oral) (BG00012)

☐ yes ☐ no ☐ Unknown

## 42 Glatiramer acetate (Copaxone) [previously copolymer-1]

☐ yes ☐ no ☐ Unknown

## 43 Immune globulin (IVIG, Gamimune, Gammagard)

☐ yes ☐ no ☐ Unknown

## 44 Interferon beta-1a (Avonex, Rebif)

☐ yes ☐ no ☐ Unknown

## 45 Interferon beta-1b (Betaseron)

☐ yes ☐ no ☐ Unknown

## 46 Laquinimod

☐ yes ☐ no ☐ Unknown

## 47 Methotrexate (MTX, Folex)

☐ yes ☐ no ☐ Unknown

## 48 Mitoxantrone (Novantrone)

☐ yes ☐ no ☐ Unknown

## 49 Mycophenolate mofetil (MMF, Cellcept)

☐ yes ☐ no ☐ Unknown

## 50 Natalizumab (Tysabri, Antegren)

☐ yes ☐ no ☐ Unknown

## 51 Rituximab (anti-CD20, Rituxan, MabThera)

☐ yes ☐ no ☐ Unknown

## 52 Sirolimus (Rapamune)

☐ yes ☐ no ☐ Unknown

## 53 Tacrolimus (FK 506, Prograf)

☐ yes ☐ no ☐ Unknown

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## 54 Teriflunomide (oral) (HMR1726)

☐ yes ☐ no ☐ Unknown

## 55 Blinded randomized trial agent

☐ yes ☐ no ☐ Unknown

## 56 Specify trial agent: \_\_\_\_\_

## 57 Other treatment

☐ yes ☐ no ☐ Unknown

## 58 Specify other treatment: \_\_\_\_\_



If the person completing this form is a Neurologist, check here and continue with the signature lines below.

First Name: \_\_\_\_\_ Last Name: \_\_\_\_\_

Phone number: \_\_\_\_\_ Fax number: \_\_\_\_\_

E-mail address: \_\_\_\_\_