

Form 2025 R2.0: Central Nervous System Tumor Pre-HSCT Data

Center:

CRID:

Key Fields	
Sequence Number: _____	
Date Received: ____-____-____	
CIBMTR Center Number: _____	
CIBMTR Recipient ID: _____	
Today's Date: ____-____-____	
Date of HSCT for which this form is being completed: ____-____-____	
HSCT type: (Check all that apply)	
<input type="checkbox"/>	Autologous
<input type="checkbox"/>	Allogeneic, unrelated
<input type="checkbox"/>	Allogeneic, related
<input type="checkbox"/>	Syngeneic (identical twin)
Product type: (Check all that apply)	
<input type="checkbox"/>	Marrow
<input type="checkbox"/>	PBSC
<input type="checkbox"/>	Cord blood
<input type="checkbox"/>	Other product
Specify: _____	
<input type="checkbox"/>	If this is a report of a second or subsequent transplant, check here and continue with question 72.

Disease Assessment at Diagnosis		Questions: 1 - 71
1	What was the date of pathologic diagnosis of the central nervous system (CNS) tumor? ____-____-____	
2	What was the primary disease for which the HSCT was performed? _____	
3	Specify other PNET: _____	
4	Specify other high-grade glial tumor: _____	
5	Specify pathologic diagnosis from Astrocytomas or Other tumors: _____	
6	What was the extent of the CNS tumor at diagnosis? _____	
7	What was the primary CNS tumor site at diagnosis?	
<input type="checkbox"/>	brainstem (medulla/pons/midbrain)	
<input type="checkbox"/>	cerebral hemisphere	
<input type="checkbox"/>	cerebellar hemisphere/vermis	
<input type="checkbox"/>	optic chiasma/hypothalamus/suprasellar area	
<input type="checkbox"/>	spinal cord	
<input type="checkbox"/>	thalamus/basal ganglia/corpus callosum	
<input type="checkbox"/>	extra-CNS primary site	
8	Specify site: _____	
9	Does the recipient have a history of co-existing phakomatosis?	
<input type="checkbox"/>	yes	<input type="checkbox"/> no <input type="checkbox"/> Unknown

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10 Specify the co-existing phakomatosis:

- ☐ cerebrometinal angiomatosis (Von Hippel-Lindau disease)
- ☐ encephalotrigeminal angiomatosis (Sturge-Weber syndrome)
- ☐ Neurofibromatosis type 1
- ☐ neurofibromatosis type 2
- ☐ tuberous sclerosis (Bourneville disease)
- ☐ other phakimatosi

11 Specify phakomatosis: \_\_\_\_\_

12 At the time of diagnosis, did the recipient have a family history of cancer in first degree relatives under 40 years of age?

- ☐ yes
- ☐ no
- ☐ Unknown

Specify the cancer(s) present in first degree relatives:

13 Basal cell carcinoma

- ☐ yes
- ☐ no

14 Brain tumors

- ☐ yes
- ☐ no

15 Breast cancer

- ☐ yes
- ☐ no

16 Colo-rectal carcinoma

- ☐ yes
- ☐ no

17 Malignant nerve sheath tumors

- ☐ yes
- ☐ no

18 Neurofibromas

- ☐ yes
- ☐ no

19 Soft tissue sarcoma

- ☐ yes
- ☐ no

20 Other cancer

- ☐ yes
- ☐ no

21 Specify cancer: \_\_\_\_\_

22 Was therapy given between diagnosis and the start of the preparative regimen?

- ☐ yes
- ☐ no

Pre-HSCT Treatment for CNS Tumor (1)

Questions: 23 - 71

Line of Therapy

23 Date therapy started: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_

24 Date therapy stopped: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_

25 Systemic Therapy:

- ☐ yes
- ☐ no

26 Number of cycles \_\_\_\_\_ ☐ Number of cycles unknown/not applicable

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## Treatment:

**27** bleomycin (BLM, Blenoxane)

☐ yes ☐ no

**28** carboplatin (Paraplatin)

☐ yes ☐ no

**29** Cisplatin (Platinol, CDDP)

☐ yes ☐ no

**30** Corticosteroids

☐ yes ☐ no

**31** Cyclophosphamide (Cytoxan)

☐ yes ☐ no

**32** Etoposide (VP-16, VePesid)

☐ yes ☐ no

**33** Ifosfamide (Ifex)

☐ yes ☐ no

**34** melphalan (L-PAM, Alkeran)

☐ yes ☐ no

**35** methotrexate (MTX, Folex)

☐ yes ☐ no

**36** nitrosourea (carmustine)

☐ yes ☐ no

**37** procarbazine (Matulane)

☐ yes ☐ no

**38** temozolomide (Temodar)

☐ yes ☐ no

**39** thiotepa (Thioplex)

☐ yes ☐ no

**40** topotecan (Hycamtin)

☐ yes ☐ no

**41** Vincristine (VCR, Oncovin)

☐ yes ☐ no

**42** other therapy

☐ yes ☐ no

**43** Specify other therapy: \_\_\_\_\_

**44** Hematopoietic growth factor?

☐ yes ☐ no

**45** Number of chemo cycles used with hematopoietic growth factor:

☐ <5 ☐ >=5 ☐ Unknown

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## 46 Radiation Therapy:

☐ yes ☐ no

### 47 Whole brain

☐ yes ☐ no

48 Specify total dose: \_\_\_\_\_ cGy (rads)

### 49 Local cranial

☐ yes ☐ no

50 Specify total dose: \_\_\_\_\_ cGy (rads)

### 51 Craniospinal

☐ yes ☐ no

52 Specify total dose: \_\_\_\_\_ cGy (rads)

### 53 Gamma knife / radiosurgery

☐ yes ☐ no

54 Specify total dose: \_\_\_\_\_ cGy (rads)

### 55 Interstitial irradiation / brachytherapy

☐ yes ☐ no

56 Specify total dose: \_\_\_\_\_ cGy (rads)

### 57 Radioactive instillation

☐ yes ☐ no

58 Specify total dose: \_\_\_\_\_ cGy (rads)

### 59 Local spinal

☐ yes ☐ no

60 Specify total dose: \_\_\_\_\_ cGy (rads)

### 61 Other site:

☐ yes ☐ no

62 Specify other site: \_\_\_\_\_

63 Specify total dose: \_\_\_\_\_ cGy (rads)

### 64 Fractionation schedule

☐ single ☐ single daily ☐ multiple daily ☐ other schedule

## 65 Surgical Biopsy/Resection:

☐ yes ☐ no

66 Type of surgery \_\_\_\_\_

67 Size of residual tumor after surgery (see codes below) \_\_\_\_\_

### 68 Was this line of therapy given for stem cell priming?

☐ yes ☐ no

69 Best Response to Line of Therapy \_\_\_\_\_

70 Date response evaluated: \_\_\_\_ - \_\_\_\_ - \_\_\_\_

### 71 Did patient relapse/progress following this line of therapy?

☐ yes ☐ no

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Disease Involvement Between Diagnosis and the Preparative Regimen

Questions: 72 - 86

Specify all sites of disease involvement between diagnosis and the start of the preparative regimen:

72 Cerebrospinal fluid

yes no Unknown

73 Extraneural

yes no Unknown

74 Distant intracranial parenchymal

yes no Unknown

75 Intracranial leptomeningeal

yes no Unknown

76 Spinal leptomeningeal

yes no Unknown

77 Local primary site

yes no Unknown

78 Other site:

yes no Unknown

79 Specify other site:

80 Was CNS tumor present in the recipient's bone at any time between diagnosis and the preparative regimen?

yes no

81 Bone scan

yes no

82 Date of bone scan: - - - - - Date bone scan unknown

83 Was the bone scan positive for CNS tumor?

yes no

84 MRI

yes no

85 Date of MRI: - - - - - Date of MRI unknown

86 Was the MRI postive for CNS tumor?

yes no

Bone Marrow Aspirate / Biopsy Performed

Questions: 87 - 102

87 Was a bone marrow aspirate / biopsy performed with in 30 days of the preparative regimen?

Yes No Not known

88 Specify the date the bone marrow biopsy was performed: - - - - - Date the bone marrow biopsy performed unknown

89 Was any tumor present in the biopsy?

yes no

Specify the test(s) used:

90 Cytogenetics

yes no

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91 Specify cytogenetic results:

- ☐ positive for tumor involvement
- ☐ Negative

92 Immunohistochemistry

- ☐ yes
- ☐ no

93 Specify immunohistochemistry results:

- ☐ positive for tumor involvement
- ☐ Negative

94 Routine histopathology

- ☐ yes
- ☐ no

95 Specify histopathology results:

- ☐ positive for tumor involvement
- ☐ Negative

96 Other test method

- ☐ yes
- ☐ no

97 Specify other test: \_\_\_\_\_

98 Specify test results:

- ☐ positive for tumor involvement
- ☐ Negative

99 Was a bone scan performed within 30 days of the preparative regimen (other than that reported at question 80)?

- ☐ yes
- ☐ no
- ☐ Unknown

100 Date of bone scan: \_\_\_\_-\_\_\_\_-\_\_\_\_ e Date unknown

101 Was the bone scan positive for CNS tumor?

- ☐ yes
- ☐ no

102 What was the sensitivity of the CNS tumor to chemotherapy prior to the preparative regimen?(Report response to last chemotherapy given prior to HSCT; chemotherapy must be >= two cycles of treatment given < 6 months prior to the preparative regimen.)

- ☐ sensitive: ≥ 50% reduction in bidimensional diameter of all disease sites with no new sites of disease (CR, CRU, PR)
- ☐ resistant: <50% reduction in diameter of all disease sites or development of new disease sites (NR, PD)
- ☐ untreated, or treated > 6 months prior to transplant
- ☐ not assessed, or chemotherapy < 2 cycles

Disease Status at the Last Assessment Prior to the Preparative Regimen

Questions: 103 - 112

103 What was the disease status immediately prior to the preparative regimen? (see CNS disease status definitions on page 4) \_\_\_\_\_

Specify all sites of residual disease:

104 Cerebrospinal fluid (CSF)

- ☐ yes
- ☐ no

105 Extraneural

- ☐ yes
- ☐ no

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106 Distant intracranial parenchymal

☐ yes ☐ no

107 Intracranial leptomeningeal

☐ yes ☐ no

108 Spinal leptomeningeal

☐ yes ☐ no

109 Local primary site

☐ yes ☐ no

110 Other site:

☐ yes ☐ no

111 Specify other site: \_\_\_\_\_

112 Date of the most recent assessment for disease status prior to the preparative regimen: \_\_\_\_ - \_\_\_\_ - \_\_\_\_

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

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

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