

Form 2024 R2.0: Sarcoma 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)

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

Product Type (check all that apply)

- Marrow
- PBSC
- Cord blood
- Other product

Specify:

If this is a report of a second or subsequent transplant, check here and continue with 115.

Disease Assessment at Diagnosis Questions: 1 - 76

- 1 What was the date of diagnosis of bone or soft tissue Sarcoma?
- 2 On the CIBMTR Form 2000 - Recipient Baseline Data, was the primary disease for which the HSCT was performed (question9) either "bone sarcoma (excluding Ewing family tumors)" (solid tumors option 10) or "soft tissue sarcoma (excluding Ewing family tumors)" (solid tumors option 21)?
- yes no
- 3 Specify bone or soft tissue sarcoma:

- 4 Primary site of sarcoma at diagnosis: (check only one)

- calcaneus
- femur
- fibula
- humerus
- metacarpal
- metatarsal
- multifocal
- patella
- pelvis
- radius
- rib
- scapula
- skull

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- ☐ sternum
- ☐ tibia
- ☐ ulna
- ☐ vertebra
- ☐ other bone location
- ☐ abdominal wall
- ☐ buttock
- ☐ Chest wall
- ☐ foot
- ☐ gastrointestinal
- ☐ genitourinary
- ☐ great vessels
- ☐ gynecologic
- ☐ hand
- ☐ head and neck
- ☐ heart
- ☐ lower arm
- ☐ lower leg
- ☐ lung / pleura
- ☐ Mediastinum
- ☐ retroperitoneum
- ☐ upper arm
- ☐ upper leg
- ☐ other viscera
- ☐ other soft tissue location

- 5 Specify multifocal: _____
- 6 Specify other bone location: _____
- 7 Specify other viscera: _____
- 8 Specify other soft tissue location: _____

9 What were the two largest dimensions of tumor mass at _____ X _____ cm
diagnosis?

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10 Tumor mass was assessed by:

- ☐ apparent by palpation
- ☐ apparent by visualization
- ☐ plain film / x-ray without contrast
- ☐ plain film / x-ray with contrast
- ☐ CT scan
- ☐ MRI scan
- ☐ radioisotope scan
- ☐ ultrasound
- ☐ Other method
- ☐ Unknown

11 Specify assessment method: _____

12 (For soft-tissue sarcoma only) What was the soft-tissue sarcoma grade at diagnosis?

- ☐ low
- ☐ intermediate
- ☐ high
- ☐ Unknown

13 Were metastases present at diagnosis?

- ☐ yes
- ☐ no
- ☐ Unknown

Specify the site(s) of metastases at diagnosis:

14 Abdominal - diffuse

- ☐ yes
- ☐ no
- ☐ Unknown

15 Bone marrow

- ☐ yes
- ☐ no
- ☐ Unknown

16 Central nervous system (CNS)

- ☐ yes
- ☐ no
- ☐ Unknown

17 Liver

- ☐ yes
- ☐ no
- ☐ Unknown

18 Lungs

- ☐ yes
- ☐ no
- ☐ Unknown

19 Lymph nodes - distant

- ☐ yes
- ☐ no
- ☐ Unknown

20 Lymph nodes - regional

- ☐ yes
- ☐ no
- ☐ Unknown

21 Skin

- ☐ yes
- ☐ no
- ☐ Unknown

22 Other site:

- ☐ yes
- ☐ no
- ☐ Unknown

23 Specify site: _____

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24 On the CIBMTR Form 2000 - Recipient Baseline Data, was there a history of malignancy other than the primary disease for which this HSCT is being performed (question 22, answered "yes")?

☐ yes ☐ no ☐ Unknown

Specify any treatment(s) given for the other malignancy:

25 Chemotherapy

☐ yes ☐ no ☐ Unknown

26 Radiation

☐ yes ☐ no ☐ Unknown

27 Other treatment

☐ yes ☐ no ☐ Unknown

28 Specify treatment _____

29 Was a cytogenetic analysis of the tumor mass performed at any time?

☐ yes ☐ no ☐ Unknown

30 Results of tests at diagnosis:

☐ Yes abnormalities identified

☐ No evaluable metaphases

☐ No abnormalities

Specify Cytogenetic abnormalities identified at diagnosis:

Translocation

31 t(1;13)

☐ yes ☐ no

32 t(1;16)

☐ yes ☐ no

33 t(2;13)

☐ yes ☐ no

34 t(7;16)

☐ yes ☐ no

35 t(7;22)

☐ yes ☐ no

36 t(11;22)

☐ yes ☐ no

37 t(12;14)

☐ yes ☐ no

38 t(12;15)

☐ yes ☐ no

39 t(12;16)

☐ yes ☐ no

40 t(12;19)

☐ yes ☐ no

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41 t(12;22)



yes



no

42 t(13;22)



yes



no

43 t(17;22)



yes



no

44 t(21;22)



yes



no

45 t(X;17)



yes



no

46 t(X;18)



yes



no

Deletion

47 del(16q) / 16q-



yes



no

48 del (17q) / 17q-



yes



no

Insertion

49 ins(19p) / 19p+



yes



no

Other

50 complex (>=3 distinct abnormalities)



yes



no

51 other abnormality



yes



no

52 specify other abnormality: _____

53 Results of tests after diagnosis to prior to the preparative regimen:



Yes abnormalities identified



No evaluable metaphases



No abnormalities on any tests after diagnosis and before the preparative regimen

Specify cytogenetic abnormalities identified at any test result between diagnosis and preparative regimen

Translocation

54 t(1;13)



yes



no

55 t(1;16)



yes



no

56 t(2;13)



yes



no

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57 t(7;16)
☐ yes ☐ no

58 t(7;22)
☐ yes ☐ no

59 t(11;22)
☐ yes ☐ no

60 t(12;14)
☐ yes ☐ no

61 t(12;15)
☐ yes ☐ no

62 t(12;16)
☐ yes ☐ no

63 t(12;19)
☐ yes ☐ no

64 t(12;22)
☐ yes ☐ no

65 t(13;22)
☐ yes ☐ no

66 t(17;22)
☐ yes ☐ no

67 t(21;22)
☐ yes ☐ no

68 t(X;17)
☐ yes ☐ no

69 t(X;18)
☐ yes ☐ no

Deletion

70 del(16q) / 16q-
☐ yes ☐ no

71 del(17q) / 17q-
☐ yes ☐ no

Insertion

72 ins(19p) / 19p+
☐ yes ☐ no

Other

73 complex (>=3 distinct abnormalities)
☐ yes ☐ no

74 other abnormality
☐ yes ☐ no

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75 specify other abnormality: _____

76 Is a copy of the cytogenetic or FISH report attached?

☐ yes ☐ no

Pre-HSCT Treatment for Sarcoma

Questions: 77 - 114

77 Was therapy given (including surgery and neo-adjuvant or adjuvant therapy) between diagnosis and the start of the preparative regimen?

☐ yes ☐ no

Lines of Therapy (1)

Questions: 78 - 114

Line of Therapy

78 Systemic Therapy:

☐ yes ☐ no

79 Date therapy started: ____-____-____

80 Date therapy stopped: ____-____-____

81 Number of cycles _____ ☐ Unknown/not applicable

82 Cisplatin (Platinol, CDDP)

☐ yes ☐ no

83 cyclophosphamide (Cytoxan)

☐ yes ☐ no

84 dactinomycin (Actinomycin D)

☐ yes ☐ no

85 dacarbazine (DTIC)

☐ yes ☐ no

86 Doxorubicin (Adriamycin)

☐ yes ☐ no

87 etoposide (VP-16, VePesid)

☐ yes ☐ no

88 Ifosfamide (Ifex)

☐ yes ☐ no

89 imatinib (Gleevec)

☐ yes ☐ no

90 melphalan (L-PAM, Alkeran)

☐ yes ☐ no

91 sunitinib (Sutent, SU11248)

☐ yes ☐ no

92 topotecan (Hycamtin)

☐ yes ☐ no

93 Vincristine (VCR, Oncovin)

☐ yes ☐ no

94 Other systemic therapy

☐ yes ☐ no

95 Specify other therapy: _____

96 Radiation Therapy:

☐ yes ☐ no

97 Date therapy started: ____ - ____ - ____

98 Date therapy stopped: ____ - ____ - ____

99 Local / regional

☐ yes ☐ no

100 Specify total dose _____ cGy (rads)

101 Sites of non-contiguous metastases

☐ yes ☐ no

102 Specify total dose _____ cGy (rads)

103 Other radiation therapy site

☐ yes ☐ no

104 Specify other radiation site _____

105 Specify total dose _____ cGy (rads)

106 Surgical Biopsy/Resection:

☐ yes ☐ no

107 Date of surgery: ____ - ____ - ____

108 Type of surgery

109 Specify other surgery _____

110 Site of surgery:

☐ primary lesion ☐ metastatic lesion ☐ Both

111 Best Response to Line of Therapy: (see definitions on page 5)

- ☐ CR (CR) - disappearance of all target lesions for a period of at least one month
- ☐ CRU (CRU) - complete response with persistent imaging abnormalities of unknown significance
- ☐ PR (PR) - at least 30% decrease in the sum of the longest diameter of measured lesions (target lesions) taking as reference the baseline sum of longest diameters
- ☐ SD (SD) - neither sufficient shrinkage to qualify for PR nor sufficient increase to qualify for PD, taking as reference the smallest sum of the longest diameters since the treatment started
- ☐ PD (PD) - at least a 20% increase in the sum of the longest diameter of measured lesions (target lesions), taking as reference the smallest sum of the longest diameters recorded since the treatment started or the appearance of one or more new lesions
- ☐ NA (Not assessed)

112 Date response evaluated: ____ - ____ - ____

113 Did disease relapse/progress following this line of therapy?

☐ yes ☐ no

114 Date of relapse/progression: ____ - ____ - ____

Laboratory Studies Prior to the Start of the Preparative Regimen

Questions: 115 - 117

115 Serum alkaline phosphatase:

☐ Known ☐ Not known

116 _____ ☐ IU/L ☐ μ kat/L

117 Upper limit of normal for alkaline phosphatase: _____

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| Disease Status at the Last Assessment Prior to the Start of the Preparative Regimen | | Questions: 118 - 120 |
|---|--|--------------------------------------|
| 118 | What was the disease status at the last evaluation prior to the preparative regimen? (see definitions above) | |
| <div><div><input type="radio"/></div>CR</div> | | <div><input type="radio"/></div> CRU |
| <div><input type="radio"/></div> PR | | <div><input type="radio"/></div> SD |
| <div><input type="radio"/></div> PD | | <div><input type="radio"/></div> NA |
| <div><input type="radio"/></div> Unknown | | |
| 119 | Specify reason: _____ | |
| 120 | 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: _____ | | |