

Form 2126 R2.0: Neuroblastoma Post-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: _____

Visit: ☐ 100 day ☐ 6 months ☐ 1 year ☐ 2 years ☐ > 2 years,

Specify: _____

Disease Assessment at the Time of Best Response to HSCT Questions: 1 - 159

1 Compared to the disease status prior to the preparative regimen, what was the best response since the date of the last report? (Include response to any post-HSCT treatment planned as of Day 0.)

☐ complete response - no primary tumor, no metasrarcic sites, catecholamines normal; included continued complete response

☐ very good partial response - primary tumor decreased by 90-99%, no metastatic sites, catecholamines normal; residual ⁹⁹Tc bone changes allowed

☐ partial response - primary tumor decreased by > 50%, all measurable metastatic sites decreased by > 50%, number of positive bone sites decreased by > 50%, no more than 1 positive bone marrow site allowed, 1 positve marrow aspirate or biopsy allowed if this represents a decrease from the number of postive sites at diagnosis

☐ minimal response - no new lesion; > 50% reduction of any measurable lesion (primary or metastases) with < 50% reduction in any other; < 25% increase in any existing lesion

☐ no response - no new lesions; < 50% reduction but < 25% increase in any existing lesion

☐ progressive disease - any new lesions; increase of any measurable lesion by > 25%; previous negative marrow positive for tumor

☐ Not assessed

☐ unknown / not tested

Specify the site(s) of persistent tumor:

2 Adrenal gland

☐ yes ☐ no

3 Bone

☐ yes ☐ no

4 Bone marrow

☐ yes ☐ no

Form 2126 R2.0: Neuroblastoma Post-HSCT Data

Center: CRID:

5

Cerebellum

yesno

6

Cerebrospinal fluid (CSF)

yesno

7

Cerebrum

yesno

8

Cranial nerves

yesno

9

Liver

yesno

10

Lymph node

yesno

11

Mediastinum

yesno

12

Paraspinal ganglion

yesno

13

Retro-orbital area

yesno

14

Skin / subcutaneous tissue

yesno

15

Elevated catecholemines

yesno

16

Other site:

yesno

17

Specify other site: _____

18

Specify the date best response was determined: _____ - ____ - ____

Date previously reported

19

Were tumor markers evaluated for the best response post-HSCT determination?

yesno

Specify the following tumor marker analyses performed:

20

Homovanilic acid (HVA):

KnownNot known

21

_____ <!--[endif]-->µg/mg creatinine

22

Date of best response determination: _____ - ____ - ____

23

Neuron specific enolase:

KnownNot known

24

_____ ng/mL

25

Date of best response determination: _____ - ____ - ____

26

Serum ferritin:

KnownNot known

Form 2126 R2.0: Neuroblastoma Post-HSCT Data

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CRID:

27 _____ ng/mg or µg/L

28 Date of best response determination: ____ - ____ - ____

29 Vanilmandelic acid (VMA):

☐ Known ☐ Not known

30 _____ µg/mg creatinine

31 Date of best response determination: ____ - ____ - ____

32 Other tumor marker analysis:

☐ Known ☐ Not known

33 Specify other analysis: _____

34 Specify level and units: _____

35 Was the recipient given planned therapy per protocol post-HSCT treatment for neuroblastoma?

☐ yes ☐ no

36 Was radiotherapy given?

☐ yes ☐ no

Specify the site(s) of radiotherapy:

37 Bone metastases

☐ yes ☐ no

38 Primary tumor

☐ yes ☐ no

39 Other site:

☐ yes ☐ no

40 Specify: _____

41 Specify the date radiotherapy was started: ____ - ____ - ____

42 Number of fractions given: _____

43 Dose per fraction: _____ <!--[endif]-->cGy (rads)

44 Was MIBG given?

☐ yes ☐ no

Specify the radioisotope given:

45 ¹³¹I-MIBG

☐ yes ☐ no

46 Other

☐ yes ☐ no

47 Specify: _____

48 Specify the date MIBG treatment was performed: ____ - ____ - ____

49 Were retinoids given?

☐ yes ☐ no

Specify the retinoids given:

50 Isotretinoin

☐ yes ☐ no

Form 2126 R2.0: Neuroblastoma Post-HSCT Data

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51 Other

☐ yes ☐ no

52 Specify: _____

53 Specify the date retinoid therapy was started: ____-____-____

54 Was immunotherapy given?

☐ yes ☐ no

Specify the drug (s) given:

55 α -interferon

☐ yes ☐ no

56 Anti-GD2 anitbody CH14.18

☐ yes ☐ no

57 Interleukin-2 (IL-2)

☐ yes ☐ no

58 Other

☐ yes ☐ no

59 Specify: _____

60 Specify the date immunotherapy was started: ____-____-____

61 Was chemotherapy given?

☐ yes ☐ no

62 Adriamycin

☐ yes ☐ no

63 Cisplatin

☐ yes ☐ no

64 Cyclophosphamide

☐ yes ☐ no

65 Dacarbazine (DTIC)

☐ yes ☐ no

66 Etoposide (VP-16)

☐ yes ☐ no

67 Isosfamide

☐ yes ☐ no

68 Melphalan (L-PAM)

☐ yes ☐ no

69 Teniposide (VM26)

☐ yes ☐ no

70 Vincristine

☐ yes ☐ no

71 Other

☐ yes ☐ no

72 Specify: _____

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73 Specify the date chemotherapy was started: ____ - ____ - ____

74 Was other treatment given?

☐ yes ☐ no

75 Specify other treatment: _____

76 Specify the date other treatment was started: ____ - ____ - ____

77 Did the neuroblastoma recur or progress since the date of the last report?

☐ yes ☐ no

Specify the known site(s) of disease progression / recurrence:

78 Adrenal gland

☐ yes ☐ no

79 Date determined: ____ - ____ - ____

80 Bone

☐ yes ☐ no

81 Date determined: ____ - ____ - ____

82 Bone marrow

☐ yes ☐ no

83 Date determined: ____ - ____ - ____

84 Cerebellum

☐ yes ☐ no

85 Date determined: ____ - ____ - ____

86 Cerebrospinal
fluid (CSF)

☐ yes ☐ no

87 Date determined: ____ - ____ - ____

88 Cerebrum

☐ yes ☐ no

89 Date determined: ____ - ____ - ____

90 Cranial nerves

☐ yes ☐ no

91 Date determined: ____ - ____ - ____

92 Liver

☐ yes ☐ no

93 Date determined: ____ - ____ - ____

94 Lymph nodes

☐ yes ☐ no

95 Date determined: ____ - ____ - ____

96 Mediastinum

☐ yes ☐ no

97 Date determined: ____ - ____ - ____

98 Paraspinal ganglion

☐ yes ☐ no

99 Date determined: ____ - ____ - ____

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100 Retro-orbital area

☐ yes ☐ no

101 Date determined: ____ - ____ - ____

102 Skin/subcutaneous tissue

☐ yes ☐ no

103 Date determined: ____ - ____ - ____

104 Other site:

☐ yes ☐ no

105 Date determined: ____ - ____ - ____

106 Specify other site: _____

Specify the methods used to examine sites of disease recurrence / persistence / progression:

107 Biopsy

☐ yes ☐ no

108 Specify disease status:

☐ Positive ☐ Negative

109 Bone scan

☐ yes ☐ no

110 Specify disease status:

☐ Positive ☐ Negative

111 Radiology

☐ yes ☐ no

112 Specify disease status:

☐ Positive ☐ Negative

113 Other method

☐ yes ☐ no

114 Specify disease status:

☐ Positive ☐ Negative

115 Specify other method: _____

116 Was the recipient given treatment for post-HSCT persistent, progressive or recurrent disease since the date of the last report?

☐ yes ☐ no ☐ Unknown

117 Was radiotherapy given?

☐ yes ☐ no

Specify the site(s) of radiotherapy:

118 Bone metastases

☐ yes ☐ no

119 Primary tumor

☐ yes ☐ no

120 Other ☐ yes ☐ no

121 Specify: _____

122 Specify the date radiotherapy was started: ____ - ____ - ____

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123 Number of fractions given: _____

124 Dose per fraction: _____ <!--[endif]-->cGy (rads)

125 Was MIBG given?

☐ yes ☐ no

Specify the radioisotope given:

126 ¹³¹I-MIBG

☐ yes ☐ no

127 Other

☐ yes ☐ no

128 Specify: _____

129 Specify the date MIBG treatment was performed: ____ - ____ - ____

130 Were retinoids given?

☐ yes ☐ no

Specify the retinoids given:

131 Isoretinoin

☐ yes ☐ no

132 Other

☐ yes ☐ no

133 Specify: _____

134 Specify the date retinoid therapy was started: ____ - ____ - ____

135 Was immunotherapy given?

☐ yes ☐ no

Specify the drug(s) given:

136 α -interferon

☐ yes ☐ no

137 Anti-GD2 antibody CH 14.18

☐ yes ☐ no

138 Interleuken-2 (IL-2)

☐ yes ☐ no

139 Other

☐ yes ☐ no

140 Specify: _____

141 Specify the date immunotherapy was started: ____ - ____ - ____

142 Was chemotherapy given?

☐ yes ☐ no

Specify the treatment(s) given:

143 Adriamycin

☐ yes ☐ no

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144 Cisplatin

☐ yes ☐ no

145 Cyclophosphamide

☐ yes ☐ no

146 Dacarbazine (DTIC)

☐ yes ☐ no

147 Etoposide (VP-16)

☐ yes ☐ no

148 Ifosfamide

☐ yes ☐ no

149 Melphalan (L-PAM)

☐ yes ☐ no

150 Tenoposide (VM26)

☐ yes ☐ no

151 Vincristine

☐ yes ☐ no

152 Other

☐ yes ☐ no

153 Specify: _____

154 Specify the date chemotherapy was started: ____ - ____ - ____

155 Was other treatment given?

☐ yes ☐ no

156 Specify other treatment: _____

157 Specify the date other treatment was started: ____ - ____ - ____

158 What is the current disease status?

☐ complete remission

☐ Not in complete remission

159 Specify the date the current disease status was determined: ____ - ____ - ____

First Name: _____ Last Name: _____

Phone number: _____ Fax number: _____

E-mail address: _____