

Form 2011 R5.0: Acute Lymphoblastic Leukemia (ALL) Pre-Infusion Data

Center: CRID:

Key Fields

Sequence Number:
Date Received:
CIBMTR Center Number:
CIBMTR Research ID:
Event date:

Subsequent Transplant or Cellular Therapy

Is this the report of a second or subsequent transplant for the same disease?
yes no

Laboratory Studies at Diagnosis

Questions: 1 - 19

1 WBC
Known Unknown
2
x 10^9/L (x 10^3/mm^3)
x 10^6/L
3 Date sample collected:
4 Blasts in blood
Known Unknown
5 %
6 Date sample collected:
7 Blasts in bone marrow
Known Unknown
8 %
9 Date sample collected:
10 Was extramedullary disease present?
yes no Unknown

Specify site(s) of disease:

11 Central nervous system
yes no
12 Cerebrospinal fluid (CSF)
Yes No
13 Parenchyma (brain)
Yes No
14 Mediastinum
yes no
15 Skin
yes no
16 Soft tissue (soft tissue mass / granulocytic sarcoma)
yes no
17 Testes / ovaries
Yes No
18 Other site
yes no
19 Specify other site:

Pre-HCT or Pre-Infusion Therapy

Questions: 20 - 63

20 Was central nervous system prophylaxis given?
yes no Unknown
Specify prophylaxis:
21 Cranial irradiation
yes no
22 Craniospinal irradiation
Yes No
23 High-dose methotrexate
yes no

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Center:

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24 Intrathecal therapy (chemotherapy)

☐ yes ☐ no

25 Other prophylaxis

☐ yes ☐ no

26 Specify prophylaxis: _____

27 Was therapy given?

☐ yes ☐ no

Line of Therapy (1)

Questions: 28 - 63

28 Purpose of therapy

- ☐ Induction
☐ Consolidation
☐ Maintenance
☐ treatment for disease relapse

29 Intrathecal therapy

☐ yes ☐ no

30 Systemic therapy

☐ yes ☐ no

31 Date therapy started

☐ Known ☐ Unknown

32 Date started: ____ - ____ - ____

33 Date therapy stopped

☐ Known ☐ Unknown

34 Date stopped: ____ - ____ - ____

35 Number of cycles

☐ Known ☐ Unknown

36 Number of cycles: _____

37 Specify systemic therapy: (check all that apply for this line of therapy)

- ☐ Blinatumomab (Blincyto)
☐ Chemotherapy
☐ Dasatinib (Sprycel)
☐ Imatinib (Gleevec)
☐ Inotuzumab
☐ Nilotinib (AMN107, Tasigna)
☐ Ponatinib (Iclusig)
☐ Rituximab (Rituxan, MabThera)
☐ Other systemic therapy

38 Specify other systemic therapy: _____

39 Radiation therapy

☐ yes ☐ no

40 Date therapy started

☐ Known ☐ Unknown

41 Date started: ____ - ____ - ____

42 Date therapy stopped

☐ Known ☐ Unknown

43 Date stopped: ____ - ____ - ____

Specify site(s) of radiation therapy:

44 Central nervous system

☐ yes ☐ no

Specify CNS irradiation:

45 Cranial

☐ yes ☐ no

46 Craniospinal

☐ Yes ☐ No

47 Other site

☐ yes ☐ no

48 Specify other site: _____

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49 Cellular therapy (e.g. CAR T-cell)

☐ yes ☐ no

50 Best response to line of therapy

- ☐ Complete remission (CR) – 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 extramedullary disease (e.g., central nervous system or soft tissue involvement), ANC of $\geq 1,000/\mu\text{L}$, Platelets $\geq 100,000/\mu\text{L}$
- ☐ Complete remission with incomplete hematologic recovery (CRi) - All CR criteria except for residual neutropenia ($<1000/\mu\text{L}$) and/or thrombocytopenia ($<100,000/\mu\text{L}$)
- ☐ No complete remission

51 Date assessed: ____-____-____

52 Was the recipient MRD negative following this line of therapy?

☐ Yes ☐ No

53 Did the recipient relapse following this line of therapy?

☐ yes ☐ no

54 Date of relapse: ____-____-____

Specify sites of disease relapse:

55 Central nervous system

☐ yes ☐ no

56 Cerebrospinal fluid (CSF)

☐ Yes ☐ No

57 Parenchyma (brain)

☐ Yes ☐ No

58 Mediastinum

☐ Yes ☐ No

59 Skin

☐ yes ☐ no

60 Soft tissue (soft tissue mass / granulocytic sarcoma)

☐ yes ☐ no

61 Testes / ovaries

☐ Yes ☐ No

62 Other site

☐ yes ☐ no

63 Specify other site: _____

Laboratory Studies at Last Evaluation Prior to the Start of the Preparative Regimen / Infusion

Questions: 64 - 91

64 WBC

☐ Known ☐ Unknown

65 _____ ☐ $\times 10^9/\text{L}$ ($\times 10^3/\text{mm}^3$)
☐ $\times 10^6/\text{L}$

66 Date sample collected: ____-____-____

67 Blasts in blood

☐ Known ☐ Unknown

68 _____%

69 Date sample collected: ____-____-____

70 Blasts in bone marrow

☐ Known ☐ Unknown

71 _____%

72 Date sample collected: ____-____-____

73 Was flow cytometry performed?

☐ yes ☐ no ☐ Unknown

Specify tissue and results at last evaluation prior to the start of the preparative regimen:

74 Blood

☐ yes ☐ no

75 Date sample collected: ____-____-____

76 Was disease detected?

☐ yes ☐ no

77 Specify percent disease detected: _____%

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78 Bone marrow

☐ yes ☐ no

79 Date sample collected: ____ - ____ - ____

80 Was disease detected?

☐ yes ☐ no

81 Specify percent disease detected: _____ %

82 Was extramedullary disease present?

☐ yes ☐ no ☐ Unknown

Specify site(s) of disease:

83 Central nervous system

☐ yes ☐ no

84 Cerebrospinal fluid (CSF)

☐ Yes ☐ No

85 Parenchyma (brain)

☐ Yes ☐ No

86 Mediastinum

☐ yes ☐ no

87 Skin

☐ yes ☐ no

88 Soft tissue (soft tissue mass / granulocytic sarcoma)

☐ yes ☐ no

89 Testes / ovaries

☐ Yes ☐ No

90 Other site

☐ yes ☐ no

91 Specify other site: _____

First Name: _____

Last Name: _____

E-mail address: _____

Date: ____ - ____ - ____