Interfant 99.

No registrations found.

Ethical review	Positive opinion
Status	Recruitment stopped
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON21662

Source
NTR

Brief title
Interfant 99

Health condition

Acute lymphoblastic leukemia.

Sponsors and support

Primary sponsor: Interfant Collaborative Group.
Source(s) of monetary or material Support: N/A

Intervention

Outcome measures

Primary outcome

Event Free Survival.

Secondary outcome

N/A

Study description

Background summary

Outcome for infant ALL is relatively poor. The Interfant-99 protocol has 3 aims:

1. assess`the outcome of a hybrid therapy schedule including AML elements on an ALL backbone.

2. assess the value of a late intensification course (VIMARAM) including high-dose araC.

3. determine which clinical and biological factors have independent prognostic relevance.

Study objective

A late intensification course (VIMARAM) improves the outcome of infants with acute lymphoblastic leukemia.

Study design

N/A

Intervention

Intensification course VIMARAM.

Contacts

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Eligibility criteria

Inclusion criteria

1. Age < 366 days;

2. Acute lymphoblastic leukemia.

Exclusion criteria

Prior therapy for leukemia (except emergency treatment).

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Randomized controlled trial
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL

Recruitment status:	Recruitment stopped
Start date (anticipated):	01-01-1999
Enrollment:	500
Type:	Actual

Ethics review

Positive opinion

Date:	05-09-2005
Application type:	First submission

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL147
NTR-old	NTR182
Other	: N/A
ISRCTN	ISRCTN24251487

Study results

Summary results
Lancet. 2007 Jul 21;370(9583):240-50.