

**KURZPROTOKOLL**  
**20150136**

<b>Öffentlicher Titel</b>	Beobachtungsstudie zu Blinatumomab bei B-ALL
<b>Wissenschaftl. Titel</b>	An Observational Study of Blinatumomab Safety and Effectiveness, Utilization, and Treatment Practices
<b>Kurztitel</b>	20150136
<b>Studienart</b>	multizentrisch, Anwendungsbeobachtung, prospektiv, offen/unverblindet, einarmig, nicht-interventionelle Studie
<b>Studienphase</b>	nicht zutreffend
<b>Erkrankung</b>	Blut: Akute lymphatische Leukämie (ALL): Rezidiert/refraktär
<b>Einschlusskriterien</b>	<ul style="list-style-type: none"><li>- Medical records of patients initiating Blincyto after country-specific reimbursement in routine clinical practice will be eligible for abstraction.</li></ul>
<b>Ausschlusskriterien</b>	<ul style="list-style-type: none"><li>- Medical records of patients who have participated in Blincyto clinical trials will be excluded since their treatment will be prescribed by the study protocol unless the patient is receiving new Blincyto treatment outside the clinical trial.</li><li>- Medical records of patients participating in other Amgen non-interventional prospective studies in which safety endpoints are collected will be excluded</li><li>- Medical records of patients who have received Blincyto via an expanded access/compassionate use program will be excluded</li><li>- In countries where patient informed consent is required for access to their medical records, any patient who does not provide informed consent will be excluded.</li></ul>
<b>Alter</b>	18 Jahre und älter
<b>Prüfzentren</b>	<b>Universitätsklinikum Frankfurt</b> (Nachbeobachtung) Medizinische Klinik II, Hämatologie/Onkologie Theodor-Stern-Kai 7 60590 Frankfurt am Main Roswitha Kotthoff Tel: 069 6301-87146 Fax: 069 6301-7463 <a href="mailto:r.kotthoff@med.uni-frankfurt.de">r.kotthoff@med.uni-frankfurt.de</a>
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<b>Registrierung in anderen Studienregistern</b>	ClinicalTrials.gov NCT03117621
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