

## Please remember BLINCYTO may cause:

1. Neurological events, including immune effector cell-associated neurotoxicity syndrome (ICANS). Signs and symptoms of this side effect may include:
  - » Uncontrolled shaking (tremor)
  - » Difficulty in communicating (aphasia) and/or writing (dysgraphia)
  - » Confusion
  - » Disturbances of brain function (encephalopathy)
  - » Seizure (convulsion)
  - » Feeling less alert
2. Cytokine release syndrome including:
  - » Fever
  - » Swelling
  - » Chills
  - » Dizziness or lightheadedness



**Please seek urgent attention from your doctor or nurse immediately or seek emergency help if you experience any of these symptoms**



**Please travel home safely and do not drive or operate moving vehicles/heavy machinery, or engage in hazardous activities whilst receiving BLINCYTO**

Version 4.0 Approved April 2026

This Patient Card fulfills the conditions of the marketing authorization and has been approved by the competent authority

# Patient Card



**Please show this card to all emergency and healthcare providers**

**Information about  
BLINCYTO®▼(blinatumomab)**

My name is \_\_\_\_\_

In Case of Emergency (ICE) contact

I am being treated with BLINCYTO,  
a treatment for B-precursor acute  
lymphoblastic leukaemia, which can  
lower my immune system.

I started treatment on \_\_\_\_\_

**Before providing any treatment, please call my prescribing physician at the number below.** If any medical evaluations are undertaken, please provide copies of all medical records, including any treatments and/or test results, to the doctor(s) named below.

|                   | Name | Hospital | City | Phone Number |
|-------------------|------|----------|------|--------------|
| Haematologist     |      |          |      |              |
| Oncologist        |      |          |      |              |
| Haematology Nurse |      |          |      |              |

▼ This medicinal product is subject to additional monitoring. This will allow quick identification of new safety information. If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this card. Side effects can be reported directly to the Health Products Regulatory Authority (HPRA) using the available methods via [www.hpra.ie](http://www.hpra.ie). By reporting side effects you can help provide more information on the safety of this medicine. Side effects should also be reported to Amgen Limited on +44 (0) 1223 436441 or Freephone 1800 535 160.