Contact Information for Healthcare Professionals

Please contact the patient's haematologist (details overleaf) for more information.

Please consult the Summary of Product Characteristics for Columvi (glofitamab) available at: **www.medicines.org.uk** if based in Great Britain or **www.emcmedicines.com/en-gb/northernireland** if based in Northern Ireland; or contact Roche Medical Information (tel: 0800 328 1629, email:**medinfo.uk@roche.com**).

▼This medicine is subject to additional monitoring. This will allow quick identification of new safety information. You can help by reporting any side effects you may get.

Important Information for Patients

If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in the patient card. You should also report side effects to Roche Products Ltd by emailing the Roche Drug Safety Centre at **welwyn.uk_dsc@roche.com** or calling +44 (0)1707 367554. You can also report any side effects via the Yellow Card Scheme at: **www.yellowcard.mhra.gov.uk**. By reporting side effects you can help provide more information on the safety of this medicine.

Date of Columvi initiation:

Name of haematologist:

Contact Number:

After-hours contact number:

My name:

My contact number:

Emergency contact:

Emergency contact number:

Roche

Important Safety Information for Patients Receiving Columvi®▼(glofitamab):

Patient Card

Columvi is used to treat patients for the following indication:

Columvi as monotherapy for the treatment of adult patients with relapsed or refractory diffuse large B-cell lymphoma (DLBCL), after two or more lines of systemic therapy.

This educational material is provided by Roche Products Limited and mandatory as a condition of the Marketing Authorisation in order to further minimise important selected risks.

M-GB-00013178 Date of preparation: July 2023
Version 1.0 MHRA Approval Date: September 2023

Please carry this card with you at all times while you are receiving Columvi (glofitamab).

Show this card to ALL Healthcare Professionals (including doctors, nurses, pharmacists) involved in your treatment and at any visits to the hospital.

Important Information for the Patient

Contact your Doctor or get emergency help right away if you have any of these symptoms. Do not attempt to diagnose and treat these symptoms yourself.

- Fever (38°C or higher)
- Fast or irregular heartbeat
- Chills or shaking chills
- Confusion
- Severe fatigue or weakness
- Difficulty breathing
- Dizziness or light-headedness
- Fainting or blurred vision
- Cold or pale clammy skin
- Headache
- Nausea
- Vomiting
- Diarrhoea

Tell your doctor if these symptoms persist or worsen.

Experiencing any of these symptoms could be due to **Cytokine Release Syndrome**, which requires immediate evaluation by a Doctor.

Cytokine Release Syndrome (CRS)

- is a group of symptoms caused by small proteins called cytokines, released in your body during inflammation.
- may involve any organ systems.
- may be caused by receiving Columvi.

Please refer to the Patient Information Leaflet (PIL) for Columvi (glofitamab) for more information, which is available at: www.medicines.org.uk if based in Great Britain or www.emcmedicines.com/en-gb/northernireland if based in Northern Ireland.

Important Information for Healthcare Professionals

This patient has received Columvi (glofitamab) - which may cause Cytokine Release Syndrome (CRS).

- Evaluate the patient immediately for signs and symptoms of CRS and treat symptoms accordingly.
- CRS may involve any organ systems and can be serious and life-threatening.
- If CRS is suspected, please refer to the latest product information for Columvi (glofitamab) for comprehensive instructions on CRS management.
- **Contact the prescribing doctor immediately for further information** – they may need to modify the next infusion of Columvi (glofitamab).