

## Reference Standards & Logistics Department

# ORIGIN OF GOODS

### Golimumab BRP

The **European Pharmacopoeia** (being elaborated by the Council of Europe – European Directorate for the Quality of Medicines & HealthCare or EDQM) is a single reference work for the **quality control of medicines** in the signatory states of the Convention on its elaboration.

The official standards published within provide a legal and scientific basis for quality control during the development, production and marketing processes.

They concern the qualitative and quantitative tests and assays to be carried out on medicines and/or on the raw materials used in production of medicines. All producers of medicines and/or substances for pharmaceutical use must therefore apply these quality standards in order to market their products in the signatory states of the Convention.

To apply these tests and assays, users must use these official reference standards.

Code catalogue	Batch number	Material origin	Country of non-preferential origin for components
Y0002502	1	Human ( <i>500 IU of tumor necrosis factor neutralising activity + 500 IU of TNF binding activity + 500 IU of Fcγ<sub>3</sub> binding activity + 500 IU of ADCC activity</i> )	Ireland

\*Information applies to batch number and sub-batches.

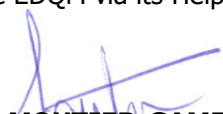
- The material is not of animal origin;
- The goods are in quantities of no greater than 20mL or 20g for each individually packaged unit;
- The above material does not contain any animal materials or cell animal culture derived products or animal additives such as albumin or serum.

**EDQM Reference Standards are supplied for laboratory test and assay only as described in the European Pharmacopoeia.**

**They are not intended for use in humans or animals as drugs, medical devices, dietary supplement or food.**

The goods transported shall be considered of "diplomatic origin" with special status and the Council of Europe EDQM shall be considered as manufacturer of these goods, although its components may originate from different countries in the world. The Council of Europe cannot provide the Certificate EUR1 / movement certificate / certificate of origin. The special status of the transferred goods does not prevent the customs authority from granting the importer a release only after it has itself settled and paid the duties and taxes.

In case of question, please contact the EDQM via its HelpDesk available from its website at [www.edqm.eu](http://www.edqm.eu).



**Dr Fanny MOUTIER-GAME**  
Head of Reception & Distribution Section  
Reference Standards and Logistics Department

