Addendum to Multiphor II Electrophoresis System User Manual

About this addendum

This document is an addendum to *Multiphor II Electrophoresis System User Manual 18-1103-43 AK*. This addendum contains additional regulatory information to comply with the Eurasian Customs Union technical regulations.

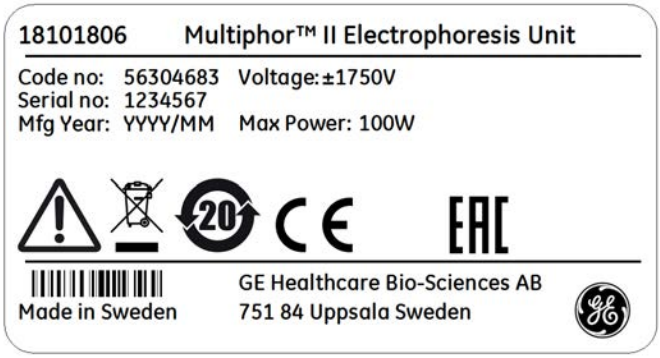
Manufacturing information

The table below summarizes the required manufacturing information. For further information, see the EU Declaration of Conformity (DoC) document.

Requirement	Content
Name and address of manufacturer	GE Healthcare Bio-Sciences AB, Björkgatan 30, SE 751 84 Uppsala, Sweden Telephone: + 46 771 400 600
Importer and/or company for obtaining information about importer	GE Healthcare LLC GE Healthcare Life Sciences Presnenskaya nab., 10C, 12th floor RU-123 317 Moscow, Russian Federation Telephone 1: + 7 495 411 9714 Fax nr: + 7 495 739 6932 Email: LSrus@ge.com

System label

Note: *The specific data shown on this system label is only an example. Actual data is specific for each individual system and may vary from system to system.*



Label	Meaning
EAC	Eurasian Conformity mark: the single conformity mark indicates that the product is approved for circulation on the markets of the member states of the Eurasian Customs Union.

For information regarding shelf life please contact your local GE representative. For information regarding manufacturing date, see year and month of production on the product.

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For local office contact information, visit
www.gelifesciences.com/contact

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