Company Led Medicinal Product Recall of Parallel Imported Nurofen® Zatoki

2.1.2014 | Circular Number P01/2014

Information on the recall of Nurofen® Zatoki

Ecosse Ltd. is voluntarily recalling the product Nurofen® Zatoki (Cold and Flu) to pharmacy level under the supervision of the Medicines Authority. While there are no quality concerns with the product itself, in order to prevent confusion to patients and suppliers of medicine, the labeling of the product will be modified to make it clear that this product is a Cold and Flu preparation.

Ecosse Ltd. has already contacted all affected customers and issued a recall notice. The product batches recalled were AK278, AM366, AM524 and AK829. The product will be back in stock when the necessary labeling modifications have been carried out.

Reporting Adverse Drug Reactions

Healthcare professionals and patients are encouraged to report any side-effects related to Nurofen Cold and Flu. ADR forms can be downloaded from **http://www.medicinesauthority.gov.mt/adrportal** and sent to 203, Level 3 Rue D'Argens Gzira, to **postlicensing.medicinesauthority@gov.mt** or to the marketing authorisation holder or their local representatives.

Dr John Joseph Borg
Post-licensing Director

Healthcare professionals and patients are encouraged to regularly check the Medicines Authority website for product safety updates as these are issued on an ongoing basis.

Medicines Authority
203 Level 3, Rue D'Argens, Gzira, GZR 1368 – Malta.
Tel. (+356) 23439000 Fax. (+356) 23439161 www.medicinesauthority.gov.mt