DECISION

No. 9/29.02.2008

on approval of changing the classification for the supply of medicinal products

The Scientific Council of the National Medicines Agency,
set up based on Order of the Minister of Health No. 485/09.05.2005, as amended, reunited on summons of the National Medicines Agency President in the ordinary meeting of 29.02.2008 in accordance with Article 10 of Government Ordinance No. 125/1998 related to the set up, organisation and functioning of the National Medicines Agency, approved as amended through Law No. 594/2002, with further changes and completions, agrees on the following

DECISION

Single article. – The changing of the classification for the supply of medicinal products is approved, according to the Annex which is integral part of this Decision.

PRESIDENT
of the Scientific Council
of the National Medicines Agency

Acad. Prof. Dr. Victor Voicu

ANNEX

List of medicinal products for which the National medicines Agency's Scientific Council has approved the change of classification concerning their release

1. CORNEREGEL 50 mg/g ophthalmic gel (dexpanthenol), box with aluminium tube, provided with an applicator, containing 5 g or 10 g ophthalmic gel, Marketing Authorisation Holder: Dr. Gerhard Mann, Chem. – Pharm. Fabrik GmbH, GERMANY, is approved for transfer from the status of medicinal product released via medical prescription (P-RF) to the OTC status.

2. DERMAZIN 10 mg/g cream (silver sulfadiazine), box with a tube, containing 50 g cream, Marketing Authorisation Holder: Lek Pharmaceuticals d.d., Slovenia, is approved for transfer from the status of medicinal product released via medical prescription (P-RF) to the OTC status.