DECISION

No. 10/09.03.2007

on approval of change of classification for release of certain medicinal products

The Scientific Council of the National Medicines Agency,
set up based on Minister of Public Health Order no. 485/09.05.2005, as amended through Minister of Public Health Order no. 159/22.02.2006, no. 1599/12.12.2006 and no. 395/27.02.2007, reunited on summons of the National Medicines Agency President in the ordinary meeting of 09.03.2007, in accord 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, as further amended, agrees on the following

DECISION

Single article. - Approval of change of classification for release of certain medicinal products, 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

Medicinal products approved by the Scientific Council of the National Medicines Agency for change of classification for release

1. **ESCAPELLE 1.5 mg tablets** (levonorgestrel), box containing one blister of PVC/Al of one tablet, Marketing Authorisation Holder GEDEON RICHTER LTD., Hungary, is approved for transfer from the status of medicinal product subject to medical prescription (P-RF) to OTC status. The medicinal product shall be released without medical prescription only for patients aged over 18 years old.
2. **POSTINOR-2, 0.75 mg tablets** (levonorgestrel), box containing one blister of PVC/Al of 2 tablets, Marketing Authorisation Holder GEDEON RICHTER LTD., Hungary, is approved for transfer from the status of medicinal product subject to medical prescription (P-RF) to OTC status. The medicinal product shall be released without medical prescription only for patients aged over 18 years old.