DECISION
no. 27/03.07.2015
on approval of switch from classification for release of Lomexin 20 mg/gram cream (fenticonazole)

The Scientific Council of the National Agency for Medicines and Medical Devices (NAMMD), established based on Order of the Minister of Health No. 374/02.04.2014, reunited on summons of the President of the National Agency for Medicines and Medical Devices in the ordinary meeting of 03.07.2015, in accordance with Article 12(5) of Government Decision no. 734/2010 on establishment, organization and operation of the National Agency for Medicines and Medical Devices, as amended, adopts the following :

DECISION

Sole article - The switch of classification for release from release on medical prescription to over the counter release of **Lomexin 20 mg/gram cream (fenticonazole),** marketing authorisation holder: Recordati S.p.A., Italy, packaging size: 30 g Aluminum Ointment Tube, is approved.

PRESIDENT
of the Scientific Council
of the National Agency for Medicines and Medical Devices,
Acad. Prof. Dr. Leonida Gherasim