DECISION
No. 11 /24.10.2017
on repeal of application for change of classification for release of Robitussin junior 3.75 mg/ 5 ml, oral solution and Robitussin antitussicum 7.5 mg / 5 ml oral solution (dextromethorphanum)

The Scientific Council of the National Agency for Medicines and Medical Devices (NAMMD), established based on Order of the Minister of Health no. 104/09.02.2017, convened on summons by the NAMMD President in the ordinary session of 24.10.2017, in accordance with provisions of Article 12 (5) of Decision of the Romanian Government no. 734/2010 on organisation and operation of the National Agency for Medicines and Medical Devices, as amended, hereby adopts the following

DECISION

Sole Article – The application for change of classification for release of Robitussin junior 3.75 mg/ 5 ml, oral solution and Robitussin antitussicum 7.5 mg / 5 ml oral solution (dextromethorphanum) – Marketing Authorisation Holder: PFIZER CORPORATION AUSTRIA GMBH – AUSTRIA – is repealed.

PRESIDENT
of the Scientific Council
of the National Agency for Medicines and Medical Devices,

Prof. Dr. Anca-Dana Buzoianu