DECISION

no. 26/11.10.2013
on approval of change of classification for release for Telfast 120 mg, film-coated tablets, box containing 1 blister x 10 film-coated tablets

The Scientific Council of the National Agency for Medicines and Medical Devices (NAMMD), established based on Order of the Minister of Health No. 158/18.02.2013, reunited on summons of the NAMMD President in the ordinary meeting of 11.10.2013, in accordance with Article 12 (5) of Emergency Government Decision no. 734/2010 on the organisation and operation of the National Agency for Medicines and Medical Devices, hereby adopts the following

DECISION

Sole article. - The change of classification for release from release based on medical prescription to release without medical prescription is approved for Telfast 120 mg, film-coated tablets, Marketing Authorisation Holder: Sanofi – Aventis Romania SRL, packaging size: box containing 1 blister x 10 film-coated tablets.

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