10 September 2015
EMA/619753/2015
Procedure Management and Committees Support

List of nationally authorised medicinal products

Active substance: pseudoephedrine, triprolidine

Procedure no.: PSUSA/00003047/201412

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Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Actifed		018723092	Johnson & Johnson S.P.A.	IT
Actifed		8559013	Johnson & Johnson Lda	PT
Actifed		8559104	Johnson & Johnson Lda	PT
Actifed		AIC N. 018723080	Johnson & Johnson S.P.A.	IT
Actifed		MA167/00103	Glaxo Wellcome Ltd	MT
Actifed		MA167/00104	Glaxo Wellcome Ltd	MT
Actifed		MA167/00107	Glaxo Wellcome Ltd	MT
Actifed		PA 823/6/2	Mcneil Healthcare (Ireland) Limited	IE
Actifed		PA 823/6/3	Mcneil Healthcare (Ireland) Limited	IE
Multi-Action Actifed		PL 15513/0014	Mcneil Products Limited	UK
Rhinopront Kombi Tabletten		1184.00.00	Recordati Pharma Gmbh	DE