



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

30 January 2025
EMA/82944/2025
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): phenylpropanolamine

Procedure No. PSUSA/00010483/202406



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
Rinexin 25 mg depottabletter	not available	6882	VIATRIS AS	NO
Rinexin 25 mg depottabletter	not available	6882	VIATRIS AS	NO
Rinexin 25 mg depottabletter	not available	9407	VIATRIS AB	SE
Rinexin 25 mg depottabletter	not available	9407	VIATRIS AB	SE
Rinexin 50 mg depottabletter	not available	6312	VIATRIS AS	NO
Rinexin 50 mg depottabletter	not available	6312	VIATRIS AS	NO
Rinexin 50 mg depottabletter	not available	9250	VIATRIS AB	SE
Rinexin 50 mg depottabletter	not available	9250	VIATRIS AB	SE