Amsterdam, 02 September 2021
EMA/520175/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: sodium iron gluconate (parenteral preparations)

Procedure no.: PSUSA/00010867/202101

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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
FERLIXIT 62,5 mg/5 ml soluzione per uso orale e uso endovenoso	not available	021455023	SANOFI S.P.A	IT
Ferrlecit 12,5 mg /ml injekční roztok	not available	12/174/73/-C	SANOFI-AVENTIS SRO	CZ
Ferrlecit 12,5 mg /ml injekční roztok	not available	12/174/73/-C	SANOFI-AVENTIS SRO	CZ
Ferrlecit 12,5 mg/ml oldatos injekció	not available	OGYI-T-438/01	SANOFI-AVENTIS ZRT	HU