



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

16 May 2019
EMA/401589/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: indoramin

Procedure no.: PSUSA/00001740/201809

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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
Indoramin 20 mg Tablets	not available	PL 17736/0089	CHEMIDEX PHARMA LIMITED	UK
Doralese Tiltab Tablets 20 mg	not available	PL 17736/0089	CHEMIDEX PHARMA LIMITED	UK
Indoramin 20 mg Film-coated Tablets	not available	PA 1161/4/1	CHEMIDEX PHARMA LIMITED	IE