



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

31 October 2019
EMA/597965/2019
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: Nicorandil

Procedure no.: PSUSA-00002152-201902

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An agency of the European Union



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
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
ADANCOR 10 mg, comprimé sécable	FR/H/0596/001	34009 335 483 3 9	MERCK SANTÉ S.A.S.	FR
ADANCOR 20 mg, comprimé	FR/H/0596/002	34009 335 485 6 8	MERCK SANTÉ S.A.S.	FR
ANGICOR	FR/H/0595/002	14453	SANOFI A/S	DK
Dancor 10 mg comprimidos	FR/H/0596/001	2414886	MERCK, S.A.	PT
Dancor 10 mg comprimidos	FR/H/0596/001	2414985	MERCK, S.A.	PT
Dancor 10 mg Tabletten	FR/H/0596/001	1-20773	MERCK GESELLSCHAFT MBH	AT
Dancor 20 mg comprimidos	FR/H/0596/002	2415081	MERCK, S.A.	PT
Dancor 20 mg Tabletten	FR/H/0596/002	1-20770	MERCK GESELLSCHAFT MBH	AT

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
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
Ikorel 10 mg Tablets	FR/H/0595/001	PL 04425/0327	AVENTIS PHARMA LTD	UK
IKOREL 10 mg, comprimé sécable	FR/H/0595/001	34009 335 481 0 0	SANOFI-AVENTIS FRANCE	FR
IKOREL 10, TABLETTEN 10 MG	FR/H/0595/001	RVG 15221	SANOFI-AVENTIS NETHERLANDS B.V.	NL
Ikorel 10mg tablets	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE

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
Ikorel 10mg tablets	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Ikorel 10mg tablets	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
Ikorel 20 mg Tablets	FR/H/0595/002	PL 04425/0328	AVENTIS PHARMA LTD	UK
IKOREL 20 MG, COMPRIME	FR/H/0595/002	335 482-7	SANOFI-AVENTIS FRANCE	FR

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
Ikorel 20mg Tablets	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Ikorel 20mg Tablets	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
Ikorel 20mg Tablets	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 10 MG	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 10 MG	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 10 MG	FR/H/0595/001	PA 540/102/1	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 20 MG	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 20 MG	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE
IKOREL TABLETS 20 MG	FR/H/0595/002	PA 540/102/2	SANOFI-AVENTIS IRELAND LTD. T/A SANOFI	IE

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NICORANDIL ZENTIVA 10 mg, comprimé sécable	not available	34009 378 035 2 6	ZENTIVA FRANCE	FR
NICORANDIL ZENTIVA 20 mg, comprimé	not available	34009 378 036 9 4	ZENTIVA FRANCE	FR