



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

10 December 2020
EMA/33992/2021
Human Medicines Evaluation Division

List of nationally authorised medicinal products

Active substance: teicoplanin

Procedure no.: EMEA/H/N/PSR/S/0025

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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
TARGOCID 100 mg, poudre et solvant pour solution injectable/pour perfusion ou solution buvable	DE/H/3916/001	NL 15047	SANOFI-AVENTIS FRANCE	France
TARGOCID 100 MG	DE/H/3916/001	27257.00.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	Germany
Targocid 200 mg Trockenstechampullen mit Lösungsmittel	DE/H/3916/002	1-19652	SANOFI-AVENTIS GMBH OSTERREICH	Austria
TARGOCID 200 MG, POEDER EN OPLOSMIDDEL VOOR OPLOSSING VOOR INJECTIE	DE/H/3916/002	BE146876	SANOFI BELGIUM	Belgium
Targocid 200 mg prašak i otapalo za otopinu za injekciju/infuziju ili oralnu otopinu	DE/H/3916/002	HR-H-903402079	SANOFI-AVENTIS CROATIA D.O.O.	Croatia
TARGOCID 200 MG	DE/H/3916/002	15/216/93-A/C	SANOFI-AVENTIS SRO	Czech Republic
TARGOCID 200 mg, poudre et solvant pour solution injectable/pour perfusion ou solution buvable	DE/H/3916/002	NL 14997	SANOFI-AVENTIS FRANCE	France

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
TARGOCID 200 MG	DE/H/3916/002	27257.01.00	SANOFI-AVENTIS DEUTSCHLAND GMBH	Germany
TARGOCID	DE/H/3916/002	1204/07-09-12	VIANEX S.A.	Greece
Targocid 200 mg powder and solvent for solution for injection/infusion or oral solution	DE/H/3916/002	540/21/1	SANOFI-AVENTIS IRELAND Ltd. T/A SANOFI	Ireland
TARGOSID 200 mg polvere e solvente per soluzione iniettabile/infusione o soluzione orale	DE/H/3916/002	026458012	SANOFI SPA	Italy
Targocid 200 mg poudre et solvant pour solution injectable/pour perfusion ou solution buvable	DE/H/3916/002	0006/10010671	SANOFI BELGIUM	Luxembourg
Targocid 200 mg powder and solvent for solution for injection/infusion or oral solution	DE/H/3916/002	082/03401	Sanofi Malta Ltd	Malta