02 October 2025
EMADOC-1700519818-2421448
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): cyclophosphamide

EURD list No. PSUSA/00000901/202501

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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
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE

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
oplossing voor injectie/infusie				
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662893	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE

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mg/ml solution à diluer pour solution injectable/pour perfusion				
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662894	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662895	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE
Cyclofosfamide Accord 200 mg/ml concentraat voor oplossing voor injectie/infusie	FI/H/1194/001	BE662896	ACCORD HEALTHCARE B.V.	BE

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
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions-	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK

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/infusionsvæske, opløsning				
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide "Accord", koncentrat til injektions- /infusionsvæske, opløsning	FI/H/1194/001	69146	ACCORD HEALTHCARE B.V.	DK
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE

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mg/ml süste-/infusioonilahuse kontsentraat				
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml süste-/infusioonilahuse kontsentraat	FI/H/1194/001	1157124	ACCORD HEALTHCARE B.V.	EE
Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio-	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI

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/infuusiokonsentraatti, liuosta varten				
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio- /infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI

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Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord 200 mg/ml injektio-/infuusiokonsentraatti, liuosta varten	FI/H/1194/001	42091	ACCORD HEALTHCARE B.V.	FI
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/001	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/002	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/003	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/004	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/005	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/006	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/007	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml	FI/H/1194/001	LT/1/24/5497/008	ACCORD HEALTHCARE B.V.	LT

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
koncentratas injekciniam ar infuziniam tirpalui				
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/009	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/010	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/011	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/012	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/013	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/014	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/015	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord Healthcare 200 mg/ml koncentratas injekciniam ar infuziniam tirpalui	FI/H/1194/001	LT/1/24/5497/016	ACCORD HEALTHCARE B.V.	LT
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV

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Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV

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injekciju/infūziju šķīduma pagatavošanai				
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml koncentrāts injekciju/infūziju šķīduma pagatavošanai	FI/H/1194/001	24-0116	ACCORD HEALTHCARE B.V.	LV
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons- /infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons- /infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons- /infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons- /infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons- /infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO

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mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning				
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml konsentrat til injeksjons-/infusjonsvæske, oppløsning	FI/H/1194/001	23-15299	ACCORD HEALTHCARE B.V.	NO
Cyclophosphamide Accord 200 mg/ml koncentrat till injektions-/infusionsvätska, lösning	FI/H/1194/001	64398	ACCORD HEALTHCARE B.V.	SE
Cyclophosphamide Accord 200 mg/ml koncentrat till injektions-/infusionsvätska, lösning	FI/H/1194/001	64398	ACCORD HEALTHCARE B.V.	SE

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
Cyclophosphamide Accord 200 mg/ml koncentrat till injektions-/infusionsvätska, lösning	FI/H/1194/001	64398	ACCORD HEALTHCARE B.V.	SE
Cyclophosphamide Accord 200 mg/ml koncentrat till injektions-/infusionsvätska, lösning	FI/H/1194/001	64398	ACCORD HEALTHCARE B.V.	SE
CYCLOPHOSPHAMIDE ACCORD 200 mg/mL, solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	34009 302 943 8 3	ACCORD HEALTHCARE FRANCE SAS	FR
CYCLOPHOSPHAMIDE ACCORD 200 mg/mL, solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	34009 302 943 9 0	ACCORD HEALTHCARE FRANCE SAS	FR
CYCLOPHOSPHAMIDE ACCORD 200 mg/mL, solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	34009 302 944 0 6	ACCORD HEALTHCARE FRANCE SAS	FR
CYCLOPHOSPHAMIDE ACCORD 200 mg/mL, solution à diluer pour solution injectable/pour perfusion	FI/H/1194/001	34009 302 944 2 0	ACCORD HEALTHCARE FRANCE SAS	FR
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG

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
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG

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
инжекционен/инфузионен разтвор				
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Циклофосфамид Акорд 200 mg/ml концентрат за инжекционен/инфузионен разтвор	FI/H/1194/001	20240251	ACCORD HEALTHCARE POLSKA SP. Z O.O.	BG
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE	CZ

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
mg/ml koncentrát pro injekční/infuzní roztok			POLSKA SP. Z O.O.	
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Cyclophosphamide Accord 200 mg/ml koncentrát pro injekční/infuzní roztok	FI/H/1194/001	44/015/23-C	ACCORD HEALTHCARE POLSKA SP. Z O.O.	CZ
Ciklofosfamid Accord 200 mg/ml koncentrat za otopinu za injekciju/infuziju	FI/H/1194/001	HR-H-872342996	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HR
Ciklofosfamid Accord 200 mg/ml koncentrat za otopinu za injekciju/infuziju	FI/H/1194/001	HR-H-872342996	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HR
Ciklofosfamid Accord 200 mg/ml koncentrat za otopinu za injekciju/infuziju	FI/H/1194/001	HR-H-872342996	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HR
Ciklofosfamid Accord 200 mg/ml koncentrat za otopinu za injekciju/infuziju	FI/H/1194/001	HR-H-872342996	ACCORD HEALTHCARE POLSKA SP. Z O.O.	HR
Cyclophosphamide Accord, 200 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	FI/H/1194/001	28911	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Cyclophosphamide Accord, 200 mg/mL, koncentrat do	FI/H/1194/001	28911	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL

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
sporządzania roztworu do wstrzykiwań / do infuzji				
Cyclophosphamide Accord, 200 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	FI/H/1194/001	28911	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Cyclophosphamide Accord, 200 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	FI/H/1194/001	28911	ACCORD HEALTHCARE POLSKA SP. Z O.O.	PL
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/01	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/02	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/03	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/04	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/05	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/06	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/07	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/08	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/09	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200	FI/H/1194/001	15486/2024/10	ACCORD HEALTHCARE	RO

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
mg/ml concentrat pentru soluție injectabilă/perfuzabilă			POLSKA SP. Z O.O.	
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/11	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/12	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/13	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/14	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/15	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Ciclofosfamidă Accord 200 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	FI/H/1194/001	15486/2024/16	ACCORD HEALTHCARE POLSKA SP. Z O.O.	RO
Cyclophosphamide Accord 200 mg/ml πυκνό σκεύασμα για παρασκευή ενεσίμου διαλύματος ή διαλύματος προς έγχυση	FI/H/1194/001	024027	ACCORD HEALTHCARE S.L.U.	CY
Cyclophosphamide Accord 200 mg/ml πυκνό σκεύασμα για παρασκευή ενεσίμου διαλύματος ή διαλύματος προς έγχυση	FI/H/1194/001	024027	ACCORD HEALTHCARE S.L.U.	CY
Cyclophosphamide Accord 200 mg/ml πυκνό σκεύασμα για παρασκευή ενεσίμου διαλύματος ή διαλύματος προς έγχυση	FI/H/1194/001	024027	ACCORD HEALTHCARE S.L.U.	CY
Cyclophosphamide Accord 200 mg/ml πυκνό σκεύασμα για παρασκευή ενεσίμου διαλύματος ή διαλύματος προς έγχυση	FI/H/1194/001	024027	ACCORD HEALTHCARE S.L.U.	CY
Ciclofosfamida Accord 200 mg/ml concentrado para	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES

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
solución inyectable y para perfusion				
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES

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
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamida Accord 200 mg/ml concentrado para solución inyectable y para perfusion	FI/H/1194/001	89.739	ACCORD HEALTHCARE S.L.U.	ES
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209017	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209029	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209031	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per	FI/H/1194/001	051209043	ACCORD HEALTHCARE S.L.U.	IT

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
soluzione iniettabile/per infusione				
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209056	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209068	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209070	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209082	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209094	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209106	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209118	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209120	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209132	ACCORD HEALTHCARE S.L.U.	IT

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
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209144	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209157	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamide Accord 200 mg/mL concentrato per soluzione iniettabile/per infusione	FI/H/1194/001	051209169	ACCORD HEALTHCARE S.L.U.	IT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877956	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877964	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877972	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	FI/H/1194/001	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877956	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877956	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877956	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877964	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução	FI/H/1194/001	5877964	ACCORD HEALTHCARE S.L.U.	PT

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
injetável ou para perfusão				
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877964	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877972	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877972	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	5877972	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	FI/H/1194/001	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	FI/H/1194/001	ACCORD HEALTHCARE S.L.U.	PT
Ciclofosfamida Accord 200 mg/ml concentrado para solução injetável ou para perfusão	FI/H/1194/001	FI/H/1194/001	ACCORD HEALTHCARE S.L.U.	PT
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI

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
concentrate for solution for injection/infusion				
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Cyclophosphamide 200 mg/ml concentrate for solution for injection/infusion	FI/H/1194/001	PL 0142/1329	ACCORD-UK LIMITED	XI
Sendoxan, pulver til injektionsvæske, opløsning	not available	09746	BAXTER A/S	DK
Sendoxan, pulver til injektionsvæske, opløsning	not available	04482	BAXTER A/S	DK
Sendoxan, pulver til injektionsvæske, opløsning	not available	09748	BAXTER A/S	DK

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
Sendoxan, overtrukne tabletter	not available	02705	BAXTER A/S	DK
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 500 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 750 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 1000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 2000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN omhulde tablet, omhulde tabletten, 50 mg	not available	RVG 01155	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 200 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 500 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 750 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 1000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 2000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN omhulde tablet, omhulde tabletten, 50 mg	not available	RVG 01155	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 200 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 500 mg	not available	RVG 08058	BAXTER B.V.	NL

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
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 750 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 1000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 2000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN omhulde tablet, omhulde tabletten, 50 mg	not available	RVG 01155	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 200 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 500 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 750 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 1000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 2000 mg	not available	RVG 08058	BAXTER B.V.	NL
ENDOXAN omhulde tablet, omhulde tabletten, 50 mg	not available	RVG 01155	BAXTER B.V.	NL
ENDOXAN I.V., poeder voor oplossing voor injectie (lyofilisaat) 200 mg	not available	RVG 08058	BAXTER B.V.	NL
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND	DE

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
			GMBH	
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.03.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.03.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND	DE

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
			GMBH	
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.02.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.00.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.01.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan	not available	6035903.03.00	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan®	not available	6035903.00.01	BAXTER DEUTSCHLAND GMBH	DE
Endoxan® „Baxter" 50 mg - Dragees	not available	10.824	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 200 mg – Trockenstechampulle	not available	10.823	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 1 g –	not available	17.948	BAXTER HEALTHCARE GMBH	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
Trockenstechampulle				
Endoxan® „Baxter" 500 mg – Trockenstechampulle	not available	13.546	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 200 mg – Trockenstechampulle	not available	10.823	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 500 mg – Trockenstechampulle	not available	13.546	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 1 g – Trockenstechampulle	not available	17.948	BAXTER HEALTHCARE GMBH	AT
Endoxan® „Baxter" 50 mg - Dragees	not available	10.824	BAXTER HEALTHCARE GMBH	AT
Cyclophosphamide Tablets 50 mg	not available	PL 00116/0389	BAXTER HEALTHCARE LTD.	XI
Cyclophosphamide Injection 500 mg.	not available	PL 00116/0387	BAXTER HEALTHCARE LTD.	XI
Cyclophosphamide Injection 1 g	not available	PL 00116/0388	BAXTER HEALTHCARE LTD.	XI
Cyclophosphamide Tablets 50 mg	not available	PL 00116/0389	BAXTER HEALTHCARE LTD.	XI
Cyclophosphamide Tablets 50 mg	not available	PL 00116/0389	BAXTER HEALTHCARE LTD.	XI
Cyclophosphamide Tablets 50 mg	not available	PL 00116/0389	BAXTER HEALTHCARE LTD.	XI
Endoxan 50 mg drajeuri	not available	9095/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 50 mg drajeuri	not available	9095/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 50 mg drajeuri	not available	9095/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 200 mg pulbere pentru solutie perfuzabila/injectabila	not available	9092/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 500 mg pulbere pentru soluţie perfuzabilă/injectabilă	not available	9093/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 1 g pulbere pentru solutie perfuzabila/injectabila	not available	9094/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan 50 mg drajeuri	not available	9095/2016/01	BAXTER HEALTHCARE SRL	RO
Endoxan® - Επικαλυμμένο δισκίο	not available	36818/10/18-03-2011	BAXTER HELLAS	GR
Endoxan® - Επικαλυμμένο δισκίο	not available	36818/10/18-03-2011	BAXTER HELLAS	GR
Endoxan® Κόνις για ενέσιμο	not available	36816/10/18-03-2011	BAXTER HELLAS	GR

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
διάλυμα				
Endoxan® Κόνις για ενέσιμο διάλυμα	not available	36817/10/18-03-2011	BAXTER HELLAS	GR
Endoxana Coated Tablets 50 mg	not available	PA2299/027/003	BAXTER HOLDING B.V.	IE
Endoxana Injection 500 mg Powder for Solution for Injection	not available	PA2299/027/001	BAXTER HOLDING B.V.	IE
Endoxana Injection 1000 mg Powder for Solution for Injection	not available	PA2299/027/002	BAXTER HOLDING B.V.	IE
Endoxana Coated Tablets 50 mg	not available	PA2299/027/003	BAXTER HOLDING B.V.	IE
Endoxana Coated Tablets 50 mg	not available	PA2299/027/003	BAXTER HOLDING B.V.	IE
Endoxana Injection 500 mg Powder for Solution for Injection	not available	MA1277/00801	BAXTER HOLDING B.V.	MT
Endoxana Injection 1000 mg Powder for Solution for Injection	not available	MA1277/00802	BAXTER HOLDING B.V.	MT
Endoxana Coated Tablets 50 mg	not available	MA1277/00803	BAXTER HOLDING B.V.	MT
Endoxana Injection 500 mg Powder for Solution for Injection	not available	MA1277/00801	BAXTER HOLDING B.V.	MT
Endoxana Injection 1000 mg Powder for Solution for Injection	not available	MA1277/00802	BAXTER HOLDING B.V.	MT
Endoxana Injection 500 mg Powder for Solution for Injection	not available	MA1277/00801	BAXTER HOLDING B.V.	MT
Endoxana Injection 1000 mg Powder for Solution for Injection	not available	MA1277/00802	BAXTER HOLDING B.V.	MT
Endoxan 500 mg por oldatos injekcióhoz	not available	OGYI-T-4534/01	BAXTER HUNGARY KFT.	HU
Endoxan 1 g por oldatos injekcióhoz	not available	OGYI-T-4534/02	BAXTER HUNGARY KFT.	HU
Endoxan bevont tabletta	not available	OGYI-T-4534/03	BAXTER HUNGARY KFT.	HU
Endoxan bevont tabletta	not available	OGYI-T-4534/04	BAXTER HUNGARY KFT.	HU
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV

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
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV

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
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 200 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0498	BAXTER LATVIA SIA	LV
ENDOXAN 1 g pulveris injekciju škiduma pagatavošanai	not available	96-0500	BAXTER LATVIA SIA	LV
ENDOXAN 500 mg pulveris injekciju šķīduma pagatavošanai	not available	96-0499	BAXTER LATVIA SIA	LV
ENDOXAN 50 mg apvalkotas tabletes	not available	96-0501	BAXTER LATVIA SIA	LV
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS

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
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 50 mg húðaðar töflur	not available	860028	BAXTER MEDICAL AB	IS
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 50 mg húðaðar töflur	not available	860028	BAXTER MEDICAL AB	IS
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 50 mg húðaðar töflur	not available	860028	BAXTER MEDICAL AB	IS
Sendoxan 50 mg húðaðar töflur	not available	860028	BAXTER MEDICAL AB	IS

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
Sendoxan 200 mg, stungulyfsstofn, lausn	not available	650708 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 500 mg, stungulyfsstofn, lausn	not available	711396 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 1 g (1000 mg), stungulyfsstofn, lausn	not available	920275 (IS)	BAXTER MEDICAL AB	IS
Sendoxan 2 g (2000 mg), stungulyfsstofn, lausn	not available	IS/1/04/157/01	BAXTER MEDICAL AB	IS
Sendoxan 500 mg pulver til injeksjonsvæske, oppløsning	not available	13-9570	BAXTER MEDICAL AB	NO
Sendoxan 1000 mg pulver til injeksjonsvæske, oppløsning	not available	13-9571	BAXTER MEDICAL AB	NO
Sendoxan 200 mg pulver til injeksjonsvæske, oppløsning	not available	3861	BAXTER MEDICAL AB	NO
Sendoxan 50 mg, tabletter drasjerte	not available	3862	BAXTER MEDICAL AB	NO
Sendoxan 500 mg pulver til injeksjonsvæske, oppløsning	not available	13-9570	BAXTER MEDICAL AB	NO
Sendoxan 1000 mg pulver til injeksjonsvæske, oppløsning	not available	13-9571	BAXTER MEDICAL AB	NO
Sendoxan 200 mg pulver til injeksjonsvæske, oppløsning	not available	3861	BAXTER MEDICAL AB	NO
Sendoxan pulver till injektionsvätska, lösning	not available	5866	BAXTER MEDICAL AB	SE
Sendoxan pulver till injektionsvätska, lösning	not available	5866	BAXTER MEDICAL AB	SE
Sendoxan pulver till injektionsvätska, lösning	not available	5866	BAXTER MEDICAL AB	SE
Sendoxan pulver till injektionsvätska, lösning	not available	5866	BAXTER MEDICAL AB	SE
Sendoxan 50 mg dragerad tablett	not available	10402	BAXTER MEDICAL AB	SE
Endoxan pó para solução injectável 500 mg	not available	4408381	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	4408480	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	8024570	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT

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
Endoxan pó para solução injectável 1000 mg	not available	8024588	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan 50 mg comprimido revestido	not available	8024604	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	4408381	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	4408480	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	8024570	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	8024588	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan 50 mg comprimido revestido	not available	8024604	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	4408381	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	4408480	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	8024570	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	8024588	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan 50 mg comprimido revestido	not available	8024604	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	4408381	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	4408480	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 500 mg	not available	8024570	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan pó para solução injectável 1000 mg	not available	8024588	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Endoxan 50 mg comprimido revestido	not available	8024604	BAXTER MÉDICO-FARMACÊUTICA, LDA.	PT
Sendoxan 50 mg tabletti, päällystetty	not available	9277	BAXTER OY	FI
Sendoxan 50 mg dragerad tablett	not available	9277	BAXTER OY	FI

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
Sendoxan 50 mg tabletti, päällystetty	not available	9277	BAXTER OY	FI
Sendoxan 50 mg dragerad tablett	not available	9277	BAXTER OY	FI
Sendoxan 2 000 mg injektiokuiva-aine, liuosta varten	not available	9276	BAXTER OY	FI
Sendoxan 1 000 mg injektiokuiva-aine, liuosta varten	not available	9276	BAXTER OY	FI
Sendoxan 500 mg injektiokuiva-aine, liuosta varten	not available	9276	BAXTER OY	FI
Sendoxan 200 mg injektiokuiva-aine, liuosta varten	not available	9276	BAXTER OY	FI
Sendoxan 2 000 mg pulver till injektionsvätska, lösning	not available	9276	BAXTER OY	FI
Sendoxan 1 000 mg pulver till injektionsvätska, lösning	not available	9276	BAXTER OY	FI
Sendoxan 500 mg pulver till injektionsvätska, lösning	not available	9276	BAXTER OY	FI
Sendoxan 200 mg pulver till injektionsvätska, lösning	not available	9276	BAXTER OY	FI
Sendoxan 50 mg tabletti, päällystetty	not available	9277	BAXTER OY	FI
Sendoxan 50 mg dragerad tablett	not available	9277	BAXTER OY	FI
Sendoxan 50 mg tabletti, päällystetty	not available	9277	BAXTER OY	FI
Sendoxan 50 mg dragerad tablett	not available	9277	BAXTER OY	FI
Sendoxan 50 mg tabletti, päällystetty	not available	9277	BAXTER OY	FI
Sendoxan 50 mg dragerad tablett	not available	9277	BAXTER OY	FI
ENDOXAN, 200 mg, proszek do sporzadzania roztworu do wstrzykiwan	not available	R/2409	BAXTER POLSKA SP Z O.O.	PL
ENDOXAN, 200 mg, proszek do sporzadzania roztworu do wstrzykiwan	not available	R/2409	BAXTER POLSKA SP Z O.O.	PL

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
ENDOXAN, 1g, proszek do sporzadzania roztworu do wstrzykiwan	not available	R/2410	BAXTER POLSKA SP Z O.O.	PL
ENDOXAN, 50mg, tabletki drażowane	not available	R/2408	BAXTER POLSKA SP Z O.O.	PL
Endoxan Baxter 50 mg Compresse rivestite	not available	015628 011	BAXTER S.P.A.	IT
Endoxan Baxter 200 mg Polvere per soluzione iniettabile	not available	015628 062	BAXTER S.P.A.	IT
Endoxan Baxter 500 mg Polvere per soluzione iniettabile	not available	015628 074	BAXTER S.P.A.	IT
Endoxan Baxter 1 g Polvere per soluzione iniettabile	not available	015628 086	BAXTER S.P.A.	IT
Endoxan 100 mg, Pulver zur Herstellung einer Injektionslösung	not available	BE 000095	BAXTER SA	BE
Endoxan 200 mg, Pulver zur Herstellung einer Injektionslösung Cyclophosphamid	not available	BE 000122	BAXTER SA	BE
Endoxan 500 mg, Pulver zur Herstellung einer Injektionslösung	not available	BE 000682	BAXTER SA	BE
Endoxan 1000 mg, Pulver zur Herstellung einer Injektionslösung	not available	BE 151207	BAXTER SA	BE
Endoxan 50 mg, Überzogene Tabletten	not available	BE 000131	BAXTER SA	BE
Endoxan 200 mg, Lyophilisat zur Herstellung einer Injektionslösung	not available	BE 158855	BAXTER SA	BE
Endoxan 500 mg, Lyophilisat zur Herstellung einer Injektionslösung	not available	BE 158864	BAXTER SA	BE
Endoxan 1000 mg, Lyophilisat zur Herstellung einer Injektionslösung	not available	BE 158873	BAXTER SA	BE
Endoxan 50 mg, omhulde	not available	BE 000131	BAXTER SA	BE

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
tabletten				
Endoxan 100 mg, poeder voor oplossing voor injectie	not available	BE 000095	BAXTER SA	BE
Endoxan 100 mg poudre pour solution injectable	not available	BE 000095	BAXTER SA	BE
Endoxan 200 mg,poudre pour solution injectable	not available	BE 000122	BAXTER SA	BE
Endoxan 200 mg, poeder voor oplossing voor injectie	not available	BE 000122	BAXTER SA	BE
Endoxan 50 mg, comprimés enrobés	not available	BE 000131	BAXTER SA	BE
Endoxan 500 mg - poudre pour solution injectable	not available	BE 000682	BAXTER SA	BE
Endoxan 500 mg, poeder voor oplossing voor injectie	not available	BE 000682	BAXTER SA	BE
Endoxan 1000 mg, poeder voor oplossing voor injectie	not available	BE 151207	BAXTER SA	BE
Endoxan 200 mg - lyofilisaat voor oplossing voor injectie	not available	BE 158855	BAXTER SA	BE
Endoxan 200 mg lyophilisat pour solution injectable	not available	BE 158855	BAXTER SA	BE
Endoxan 500 mg - lyofilisaat voor oplossing voor injectie	not available	BE 158864	BAXTER SA	BE
Endoxan 500 mg - lyophilisat pour solution injectable	not available	BE 158864	BAXTER SA	BE
Endoxan 1000 mg lyofilisaat voor oplossing voor injectie	not available	BE 158873	BAXTER SA	BE
Endoxan 1000 mg lyophilisat pour solution injectable.	not available	BE 158873	BAXTER SA	BE
Endoxan 1000 mg poudre pour solution injectable	not available	BE 151207	BAXTER SA	BE
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur	not available	0039277	BAXTER SA	LU

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
Herstellung einer Injektions-/Infusionslösung				
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour	not available	0039280	BAXTER SA	LU

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
solution injectable /pour perfusion				
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer	not available	0039294	BAXTER SA	LU

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
Injektionslösung				
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-	not available	0039263	BAXTER SA	LU

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
/Infusionslösung				
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU

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
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg Pulver zur Herstellung einer Injektionslösung	not available	0039294	BAXTER SA	LU
Endoxan 200 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039280	BAXTER SA	LU
Endoxan 100 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039277	BAXTER SA	LU
Endoxan 50 mg, überzogene Tabletten	not available	0039246	BAXTER SA	LU
Endoxan 1000 mg, Pulver zur Herstellung einer Injektions-/Infusionslösung	not available	0039263	BAXTER SA	LU
Endoxan 100 mg, poudre pour solution injectable / pour perfusion	not available	0039277	BAXTER SA	LU
Endoxan 200 mg, poudre pour solution injectable /pour perfusion	not available	0039280	BAXTER SA	LU
Endoxan 50 mg, comprimés enrobés	not available	0039246	BAXTER SA	LU
Endoxan 500 mg - poudre pour solution injectable	not available	0039294	BAXTER SA	LU
Endoxan 1000 mg, poudre pour solution injectable /pour perfusion	not available	0039263	BAXTER SA	LU
Endoxan 500 mg, poudre pour solution injectable	not available	34009 321 195 0 9	BAXTER SAS	FR
Endoxan 500 mg, poudre pour solution injectable	not available	34009 321 196 7 7	BAXTER SAS	FR
Endoxan 500 mg, poudre pour solution injectable	not available	34009 321 197 3 8	BAXTER SAS	FR
Endoxan 500 mg, poudre pour solution injectable	not available	34009 315 820 4 5	BAXTER SAS	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
Endoxan 500 mg, poudre pour solution injectable	not available	34009 321 193 8 7	BAXTER SAS	FR
Endoxan 500 mg, poudre pour solution injectable	not available	34009 321 194 4 8	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 558 370 4 6	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 558 371 0 7	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 558 372 7 5	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 558 373 3 6	BAXTER SAS	FR
ENDOXAN 500 mg, poudre pour solution injectable	not available	34009 550 287 8 9	BAXTER SAS	FR
ENDOXAN 500 mg, poudre pour solution injectable	not available	34009 550 287 9 6	BAXTER SAS	FR
ENDOXAN 500 mg, poudre pour solution injectable	not available	34009 550 288 0 2	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 550 286 5 9	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 550 286 6 6	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 550 286 7 3	BAXTER SAS	FR
ENDOXAN 1000 mg, poudre pour solution injectable	not available	34009 550 286 8 0	BAXTER SAS	FR
ENDOXAN 50 mg, comprimé enrobé	not available	34009 303 589 0 0	BAXTER SAS	FR
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK

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
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Endoxan 50 mg obalené tablety	not available	44/0298/97-S	BAXTER SLOVAKIA S.R.O.	SK
Genoxal 50 mg comprimidos recubiertos	not available	33214	BAXTER, S.L.	ES
Genoxal 200 mg polvo para solución inyectable y para perfusión	not available	33411	BAXTER, S.L.	ES
Genoxal 1.000 mg polvo para solución inyectable y para perfusión	not available	48972	BAXTER, S.L.	ES
Cyclophosphamid beta 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	2205439.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/002	2205440.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	2205441.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	2205439.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	2205439.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	2205439.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-	DE/H/6587/002	2205440.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE

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
/Infusionslösung				
Cyclophosphamid beta 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/002	2205440.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	2205441.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/002	2205440.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	2205441.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Cyclophosphamid beta 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	2205441.00.00	BETAPHARM ARZNEIMITTEL GMBH	DE
Ciclofosfamidă Dr. Reddy’s 500 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/001	14350/2022/01	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy’s 500 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/001	14350/2022/02	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy’s 500 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/001	14350/2022/03	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy’s 500 mg/ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/001	14350/2022/04	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy’s 1000 mg/2 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/002	14351/2022/01	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy’s 1000 mg/2 ml concentrat pentru	DE/H/6587/002	14351/2022/02	DR. REDDY'S LABORATORIES ROMANIA	RO

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
soluție injectabilă/perfuzabilă			SRL	
Ciclofosfamidă Dr. Reddy's 1000 mg/2 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/002	14351/2022/03	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy's 1000 mg/2 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/002	14351/2022/04	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy's 2000 mg/4 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/003	14352/2022/01	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy's 2000 mg/4 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/003	14352/2022/02	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy's 2000 mg/4 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/003	14352/2022/03	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ciclofosfamidă Dr. Reddy's 2000 mg/4 ml concentrat pentru soluție injectabilă/perfuzabilă	DE/H/6587/003	14352/2022/04	DR. REDDY'S LABORATORIES ROMANIA SRL	RO
Ендоксан 50 mg обвити таблетки	not available	20020260	ECOPHARM GROUP	BG
Ендоксан 50 mg обвити таблетки	not available	20020260	ECOPHARM GROUP	BG
Ендоксан 50 mg обвити таблетки	not available	20020260	ECOPHARM GROUP	BG
Ендоксан 200 mg, прах за инжекционен разтвор	not available	20020298	ECOPHARM GROUP	BG
Ендоксан 500 mg, прах за инжекционен разтвор	not available	20020299	ECOPHARM GROUP	BG
Ендоксан 1 g, прах за инжекционен разтвор	not available	20020300	ECOPHARM GROUP	BG
Ендоксан 50 mg обвити таблетки	not available	20020260	ECOPHARM GROUP	BG
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/2977/003	90401.00.00	HEXAL AG	DE
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer	NL/H/2977/003	90401.00.00	HEXAL AG	DE

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
Injektions-/Infusionslösung				
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/2977/003	90401.00.00	HEXAL AG	DE
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/2977/003	90401.00.00	HEXAL AG	DE
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/2977/003	90401.00.00	HEXAL AG	DE
Cyclophosphamid HEXAL 2000 mg Pulver zur Herstellung einer Injektions-/Infusionslösung	NL/H/2977/003	90401.00.00	HEXAL AG	DE
ENDOXAN 200 mg, süstelahuse pulber	not available	145896	OÜ BAXTER ESTONIA	EE
ENDOXAN 200 mg, süstelahuse pulber	not available	145896	OÜ BAXTER ESTONIA	EE
ENDOXAN 500 mg, süstelahuse pulber	not available	145996	OÜ BAXTER ESTONIA	EE
ENDOXAN 1 g, süstelahuse pulber	not available	146096	OÜ BAXTER ESTONIA	EE
ENDOXAN 50 mg, kaetud tabletid	not available	182197	OÜ BAXTER ESTONIA	EE
ENDOXAN 200 mg, süstelahuse pulber	not available	145896	OÜ BAXTER ESTONIA	EE
ENDOXAN 200 mg, süstelahuse pulber	not available	145896	OÜ BAXTER ESTONIA	EE
ENDOXAN 500 mg, süstelahuse pulber	not available	145996	OÜ BAXTER ESTONIA	EE
ENDOXAN 1 g, süstelahuse pulber	not available	146096	OÜ BAXTER ESTONIA	EE
ENDOXAN 50 mg, kaetud tabletid	not available	182197	OÜ BAXTER ESTONIA	EE
Cyclophosphamid Reddy 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	140968	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 1000	DE/H/6587/002	140969	REDDY HOLDING GMBH	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
mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung				
Cyclophosphamid Reddy 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	140968	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	140968	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/002	140969	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/002	140969	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	140970	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	140970	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/003	140970	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 500 mg/ml Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DE/H/6587/001	140968	REDDY HOLDING GMBH	AT
Cyclophosphamid Reddy 1000 mg/2 ml Konzentrat zur Herstellung einer Injektions-	DE/H/6587/002	140969	REDDY HOLDING GMBH	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
/Infusionslösung				
Cyclophosphamid Reddy 2000 mg/4 ml Konzentrat zur Herstellung einer Injektions- /Infusionslösung	DE/H/6587/003	140970	REDDY HOLDING GMBH	AT
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500	DE/H/6587/001	86548	REDDY PHARMA IBERIA,	ES

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
mg/ml concentrado para solución inyectable y para perfusión			S.A.	
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
Ciclofosfamida Dr. Reddy 500 mg/ml concentrado para solución inyectable y para perfusión	DE/H/6587/001	86548	REDDY PHARMA IBERIA, S.A.	ES
CYCLOPHOSPHAMIDE REDDY PHARMA 1000 mg/2 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/002	34009 302 430 5 3	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 1000 mg/2 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/002	34009 550 860 7 9	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 1000 mg/2 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/002	34009 550 860 8 6	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 1000 mg/2 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/002	34009 550 860 9 3	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 2000 mg/4 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/003	34009 302 430 6 0	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 2000 mg/4 mL, solution à diluer pour solution	DE/H/6587/003	34009 550 861 0 9	REDDY PHARMA SAS	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
injectable/pour perfusion				
CYCLOPHOSPHAMIDE REDDY PHARMA 2000 mg/4 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/003	34009 550 861 1 6	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 2000 mg/4 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/003	34009 550 861 2 3	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 500 mg/1 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/001	34009 302 430 4 6	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 500 mg/1 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/001	34009 550 860 4 8	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 500 mg/1 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/001	34009 550 860 5 5	REDDY PHARMA SAS	FR
CYCLOPHOSPHAMIDE REDDY PHARMA 500 mg/1 mL, solution à diluer pour solution injectable/pour perfusion	DE/H/6587/001	34009 550 860 6 2	REDDY PHARMA SAS	FR
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz	DK/H/3394/001	68206	SANDOZ A/S	DK
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE

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
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamide Sandoz 100 mg/ml koncentrat till injektions-/infusionsvätska, lösning	DK/H/3394/001	63696	SANDOZ A/S	SE
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-/Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions-	DK/H/3394/001	141977	SANDOZ GMBH	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
/Infusionslösung				
Cyclophosphamid Sandoz 100 mg/ml – Konzentrat zur Herstellung einer Injektions- /Infusionslösung	DK/H/3394/001	141977	SANDOZ GMBH	AT
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Cyclophosphamide 2000 mg Powder for Solution for Injection or Infusion	NL/H/2977/003	PL 04416/1395	SANDOZ LTD	XI
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен разтвор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен разтвор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен разтвор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG

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
раствор				
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен раствор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен раствор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Циклофосфамид Сандоз 100 mg/ml концентрат за инжекционен/инфузионен раствор	DK/H/3394/001	20240053	SANDOZ PHARMACEUTICALS D.D.	BG
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100 mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL
Cyclophosphamide Sandoz, 100	DK/H/3394/001	28340	SANDOZ POLSKA SP. Z O.O.	PL

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
mg/mL, koncentrat do sporządzania roztworu do wstrzykiwań / do infuzji				
Endoxan 200 mg milteliai injekciniam tirpalui	not available	LT/1/97/3087/002	UAB "BAXTER LITHUANIA"	LT
Endoxan 500 mg milteliai injekciniam tirpalui	not available	LT/1/97/3087/003	UAB "BAXTER LITHUANIA"	LT
Endoxan 1000 mg milteliai injekciniam tirpalui	not available	LT/1/97/3087/004	UAB "BAXTER LITHUANIA"	LT
Endoxan 50 mg dengtos tabletės	not available	LT/1/97/3087/001	UAB "BAXTER LITHUANIA"	LT