13 November 2025
EMADOC-1700519818-2822399
Human Medicines Division

List of nationally authorised medicinal products

Active substance(s): codeine / paracetamol

EURD list No. PSUSA/00000851/202503

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Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000 An agency of the European Union

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
DAFALGAN CODEINE 500 mg/30 mg Filmtabletten	not available	BE205886	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg Filmtabletten	not available	BE205886	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, Brausetabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, Brausetabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, Brausetabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEÏNE 500 mg/30 mg, bruistabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEÏNE 500 mg/30 mg, bruistabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEÏNE 500 mg/30 mg, bruistabletten	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, comprimés effervescents	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, comprimés effervescents	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, comprimés effervescents	not available	BE137776	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, comprimés pelliculés	not available	BE205886	UPSA SAS	BE
DAFALGAN CODEINE 500 mg/30 mg, comprimés pelliculés	not available	BE205886	UPSA SAS	BE
DAFALGAN CODEÏNE 500 mg/30 mg, filmomhulde tabletten	not available	BE205886	UPSA SAS	BE
DAFALGAN CODEÏNE 500 mg/30 mg, filmomhulde tabletten	not available	BE205886	UPSA SAS	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA 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
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537751	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537724	TEVA 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
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000 mg/60 mg Filmtabletten	DK/H/2786/002	BE537751	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537751	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA 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
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg comprimés pelliculés	DK/H/2786/002	BE537751	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde	DK/H/2786/002	BE537724	TEVA 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
tabletten				
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537751	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde	DK/H/2786/002	BE537733	TEVA 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
tabletten				
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537724	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537733	TEVA B.V	BE
Paracetamol/Codeine Teva 1000mg/60mg filmomhulde tabletten	DK/H/2786/002	BE537751	TEVA B.V	BE
Gelonida Schmerztabletten Tabletten mit 500 mg Paracetamol und 30 mg Codeinphosphat-Hemihydrat Zur Anwendung bei Erwachsenen und Jugendlichen ab 12 Jahren	not available	13300.00.00	PFIZER PHARMA GMBH	DE
Gelonida Schmerztabletten Tabletten mit 500 mg Paracetamol und 30 mg Codeinphosphat-Hemihydrat Zur Anwendung bei Erwachsenen und Jugendlichen ab 12 Jahren	not available	13300.00.00	PFIZER PHARMA GMBH	DE
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE 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
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Kodipar, filmovertrukne tabletter	not available	12042	ORIFARM HEALTHCARE A/S	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	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
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000/60 mg	DK/H/2786/002	53786	TEVA B.V	DK
Pinex comp, filmovertrukne tabletter 1000+60 mg	DK/H/2786/002	53786	TEVA B.V	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
AnalgiPlus 500 mg/30 mg comprimidos recubiertos con película.	not available	63.118	FAES FARMA, S.A.	ES
AnalgiPlus 500 mg/30 mg comprimidos recubiertos con película.	not available	63.118	FAES FARMA, S.A.	ES
Cod-Efferalgan 500 mg/ 30 mg comprimidos efervescentes	not available	60.748	UPSA SAS	ES
Dolocatil Codeína 650 mg/30 mg comprimidos	not available	65.510	FERRER INTERNACIONAL, S.A.	ES
Panacod brustablett	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod brustablett	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod brustablett	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod brustablett	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod brustablett	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod poretabletti	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod poretabletti	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod poretabletti	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod poretabletti	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod poretabletti	not available	11050	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI

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Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tablett	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
Panacod tabletti	not available	10294	CNX THERAPEUTICS IRELAND LIMITED	FI
CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR
CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR
CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR
CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR
CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR

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CLARADOL CODEINE 500 mg/20 mg, comprimé	not available	333 046-5	TEOFARMA S.R.L.	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 928 9 5	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 792 7 0	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 927 2 7	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 332 207 5 4	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 793 3 1	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 791 0 2	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 928 9 5	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 792 7 0	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 927 2 7	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 332 207 5 4	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 793 3 1	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 791 0 2	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 928 9 5	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 792 7 0	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 560 927 2 7	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 332 207 5 4	OPELLA HEALTHCARE FRANCE SAS	FR
CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 793 3 1	OPELLA HEALTHCARE FRANCE SAS	FR

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CODOLIPRANE ADULTES 400 mg/20 mg, comprimé sécable	not available	34009 326 791 0 2	OPELLA HEALTHCARE FRANCE SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 333 167 7 8	UPSA SAS	FR
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 126 9 8	UPSA 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
DAFALGAN CODEINE, comprimé effervescent sécable	not available	34009 556 125 2 0	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 332 758 1 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 581 219 7 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 557 156 9 6	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 332 758 1 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 581 219 7 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 557 156 9 6	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 332 758 1 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 581 219 7 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 557 156 9 6	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 332 758 1 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 581 219 7 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 557 156 9 6	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 332 758 1 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 581 219 7 5	UPSA SAS	FR
DAFALGAN CODEINE, comprimé pelliculé	not available	34009 557 156 9 6	UPSA SAS	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	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**
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 331 808 5 0	LABORATOIRES GRÜNENTHAL S.A.S.	FR
LINDILANE 400 mg/25 mg, comprimé	not available	34009 358 825 8 5	LABORATOIRES GRÜNENTHAL S.A.S.	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 356 156 1 9	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 357 486 5 2	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 336 104 6 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 356 156 1 9	PIERRE FABRE MEDICAMENT	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
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 357 486 5 2	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 336 104 6 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 356 156 1 9	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 357 486 5 2	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 300 mg/25 mg, comprimé	not available	34009 336 104 6 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 558 349 5 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 556 954 9 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 357 487 1 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 558 349 5 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 556 954 9 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 357 487 1 3	PIERRE FABRE MEDICAMENT	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
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 558 349 5 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 556 954 9 3	PIERRE FABRE MEDICAMENT	FR
PARACETAMOL/CODEINE PIERRE FABRE 600 mg/50 mg, comprimé	not available	34009 357 487 1 3	PIERRE FABRE MEDICAMENT	FR
Lonalgal (500+30) mg δισκία	not available	1950101	LAVIPHARM S.A	GR
Lonalgal (500+30) mg δισκία	not available	1950101	LAVIPHARM S.A	GR
Kapake 15mg/500mg Tablets.	IE/H/0720/001	PA22680/2/1	GALEN PHARMA IRELAND LIMITED	IE
Panadeine Extra Strength Tablets Paracetamol 500 mg Codeine Phosphate hemihydrate 12.8 mg	not available	PA0678/026/003	HALEON IRELAND LIMITED	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol 500 mg/30 mg Effervescent Tablets	not available	PA1113/027/002	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE

Product Name (in authorisation country)	MRP/DCP Authorisation number	National Authorisation Number	MAH of product in the member state	Member State where product is authorised
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Solpadol Caplets 500 mg/30 mg Tablets	not available	PA1113/027/001	PHOENIX LABS UNLIMITED COMPANY	IE
Tylex 30 mg / 500 mg Hard Capsules	not available	PA 891/14/1	UCB (PHARMA) IRELAND LIMITED	IE
Tylex 30 mg / 500 mg Hard Capsules	not available	PA 891/14/1	UCB (PHARMA) IRELAND LIMITED	IE
Tylex 30 mg / 500 mg Hard Capsules	not available	PA 891/14/1	UCB (PHARMA) IRELAND LIMITED	IE
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	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
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín 500 mg/10 mg filmuhúðaðar töflur	not available	IS/1/17/023/01	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	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
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Parkódín forte 1000 mg/60 mg filmuhúðaðar töflur	DK/H/2786/002	IS/1/15/052/02	TEVA B.V	IS
Co Efferalgan 500 mg + 30 mg compresse effervescenti	not available	027989019	UPSA SAS	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
Co Efferalgan 500 mg + 30 mg compresse rivestite con film	not available	027989033	UPSA SAS	IT
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg poudre pour solution buvable.	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
Algocod 500 mg/30 mg Pulver zur Herstellung einer Lösung zum Einnehmen	not available	2010040003	LABORATOIRES SMB S.A.	LU
DAFALGAN CODEINE 500 mg/30 mg Filmtabletten	not available	2006028443	UPSA SAS	LU
DAFALGAN CODEINE 500 mg/30 mg, Brausetabletten	not available	2006028442	UPSA SAS	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
DAFALGAN CODEINE 500 mg/30 mg, Brausetabletten	not available	2006028442	UPSA SAS	LU
DAFALGAN CODEINE 500 mg/30 mg, comprimés effervescents	not available	2006028442	UPSA SAS	LU
DAFALGAN CODEINE 500 mg/30 mg, comprimés effervescents	not available	2006028442	UPSA SAS	LU
DAFALGAN CODEINE 500 mg/30 mg, comprimés pelliculés	not available	2006028443	UPSA SAS	LU
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte stikkpiller "Karo Pharma"	not available	6139	KARO PHARMA AS	NO
Paralgin forte tablett "Karo Pharma".	not available	4088	KARO PHARMA AS	NO
Paralgin forte tablett "Karo Pharma".	not available	4088	KARO PHARMA AS	NO
Paralgin major stikkpiller "Karo Pharma"	not available	7735	KARO PHARMA AS	NO
Paralgin major stikkpiller "Karo Pharma"	not available	7735	KARO PHARMA AS	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO

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
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Forte brusetabletter	not available	7522	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO

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
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Pinex Major 1000 mg/60 mg filmdrasjerte tabletter	DK/H/2786/002	14/9970	TEVA B.V	NO
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Efferalgan Codeine, 500 mg + 30 mg, tabletki musujące	not available	R/6780	UPSA SAS	PL
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE 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**
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 40 mg/ml + 1 mg/ml Xarope	not available	3/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 500 mg + 20 mg Comprimidos	not available	2/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 500 mg + 20 mg Comprimidos	not available	2/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 500 mg + 20 mg Comprimidos	not available	2/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron 500 mg + 20 mg Comprimidos	not available	2/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE 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
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 1000 mg + 60 mg Supositórios	not available	6/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE FARMACÊUTICA, LDA.	PT
Dol-u-ron Forte 500 mg + 30 mg Cápsulas	not available	5/93	BENE 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**
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg brustabletter	not available	10151	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon 500 mg/30 mg, tabletter	not available	9777	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	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
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Citodon forte 1 g/60 mg, suppositorier	not available	11529	EVOLAN PHARMA AB	SE
Co-codamol 15/500mg Film-coated Tablets	not available	PL 0142/0921	ACCORD-UK LIMITED	XI
Co-codamol 15/500mg Film-coated Tablets	not available	PL 0142/0921	ACCORD-UK LIMITED	XI
Co-codamol 15/500mg Film-coated Tablets	not available	PL 0142/0921	ACCORD-UK LIMITED	XI
Co-codamol 15mg/500mg Film-coated Tablets	not available	PL 49565/0094	RUDIPHARM LIMITED	XI
Co-codamol 15mg/500mg Film-coated Tablets	not available	PL 49565/0094	RUDIPHARM LIMITED	XI
Co-codamol 15mg/500mg Film-coated Tablets	not available	PL 49565/0094	RUDIPHARM LIMITED	XI
Co-codamol 15mg/500mg Tablets	IE/H/0720/001	PL 27827/0011	GALEN LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Effervescent Tablets	not available	PL 17780/0046	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA 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
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500 Tablets.	not available	PL 17780/0044	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 30/500mg Effervescent Tablets	not available	PL 51463/0016	KENT PHARMA UK LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 30mg/500mg Tablets	not available	PL 17907/0479	BRISTOL LABORATORIES LIMITED	XI
Co-codamol 8 mg/500 mg effervescent tablets	not available	PL 0142/1258	ACCORD-UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA 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
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Effervescent Tablets	not available	PL 17780/0511	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500 Tablets	not available	PL 17780/0510	ZENTIVA PHARMA UK LIMITED	XI
Co-codamol 8/500mg Capsules	not available	PL 33446/0002	CUSTOM HEALTHCARE LIMITED	XI
Co-codamol 8/500mg Capsules	not available	PL 33446/0002	CUSTOM HEALTHCARE LIMITED	XI
Co-Codamol 8/500mg Capsules.	not available	PL 33446/0001	CUSTOM HEALTHCARE LIMITED	XI
Co-Codamol 8/500mg Capsules.	not available	PL 33446/0001	CUSTOM HEALTHCARE LIMITED	XI
CO-CODAMOL 8/500mg TABLETS	not available	PL 20075/0702	ACCORD HEALTHCARE LIMITED	XI
Co-codamol 8mg/500mg tablets	not available	PL 25298/0427	BROWN & BURK UK LIMITED	XI
Co-codamol 8mg/500mg tablets	not available	PL 25298/0427	BROWN & BURK 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
Co-codamol 8mg/500mg tablets	not available	PL 25298/0427	BROWN & BURK UK LIMITED	XI
Co-codamol 8mg/500mg tablets	not available	PL 25298/0427	BROWN & BURK UK LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 20416/0739	CRESCENT PHARMA LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0477	BRISTOL LABORATORIES LIMITED	XI
Co-Codamol 8mg/500mg Tablets	not available	PL 17907/0478	BRISTOL LABORATORIES LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA 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
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0018	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Co-codamol Effervescent Tablets 8/500mg	not available	PL 51463/0036	KENT PHARMA UK LIMITED	XI
Codeine/Paracetamol 60mg/1000mg film-coated tablets	not available	PL 36687/0141	TORRENT PHARMA (UK) LTD.	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING LIMITED	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING LIMITED	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING LIMITED	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING LIMITED	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING LIMITED	XI
Panadol Ultra	not available	PL 44673/0010	HALEON UK TRADING 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
Paracetamol & Codeine 500mg/8mg Effervescent Tablets	not available	PL 0142/1258	ACCORD-UK LIMITED	XI
Solpadeine Max Tablets	not available	PL 02855/0074	OMEGA PHARMA LTD	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
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Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Caplets	not available	PL 35104/0052	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	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**
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
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Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
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Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	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
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
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Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	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**
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
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Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Solpadol 30mg/500mg Effervescent Tablets	not available	PL 35104/0053	PHOENIX LABS UNLIMITED COMPANY	XI
Zipamol PLUS 8 mg/500 mg effervescent tablets	not available	PL 0142/1258	ACCORD-UK LIMITED	XI