11 January 2018
EMA/27324/2018
Human Medicines Evaluation Division

# List of nationally authorised medicinal products

Active substance: risperidone

Procedure no.: PSUSA/00002649/201705

30 Churchill Place ● Canary Wharf ● London E14 5EU ● United Kingdom
**Telephone** +44 (0)20 3660 6000 **Facsimile** +44 (0)20 3660 5525
**Send a question via our website** www.ema.europa.eu/contact

| 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 |
|---|---|---|---|---|
| Risperidona Farmalider 0,5 mg comprimidos recubiertos con película | not available | 66804 | FARMALIDER, S.A. | ES |
| Eperon 0,5 mg επικαλυμμένα με λεπτό υμένιο δισκία | DK/H/0980/001 | 20177 | MEDOCHEMIE LTD. | CY |
| Eperon 6 mg επικαλυμμένα με λεπτό υμένιο δισκία | DK/H/0980/006 | 20182 | MEDOCHEMIE LTD. | CY |
| Eperon 8 mg επικαλυμμένα με λεπτό υμένιο δισκία | DK/H/0980/007 | 20183 | MEDOCHEMIE LTD. | CY |
| RISPERDAL 0,5 mg filmomhulde tabletten | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg filmomhulde tabletten | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg filmomhulde tabletten | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | BE |
| Risperdal 4 mg Filmtabletten | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 50 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/015 | BE254615 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 2 mg Schmelztabletten | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg Filmtabletten | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| Risperdal 3 mg Filmtabletten | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 0,5 mg Schmelztabletten | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| Risperdal 0,5 mg Filmtabletten | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 25 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/013 | BE254597 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 1 mg Schmelztabletten | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 37,5 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/014 | BE254606 | JANSSEN-CILAG NV | BE |
| Risperdal 6 mg Filmtabletten | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | BE |
| Risperdal 1 mg/ml Lösung zum Einnehmen | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg filmdragerade tabletter | DE/H/2184/004 | 11479 | JANSSEN-CILAG OY | FI |
| Risperdal 3 mg filmdragerade tabletter | DE/H/2184/005 | 11480 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 37,5 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/014 | 16895 | JANSSEN-CILAG OY | FI |
| Risperdal 6 mg kalvopäällysteiset tabletit | DE/H/2184/007 | 12302 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 25 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/013 | 16894 | JANSSEN-CILAG OY | FI |

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|---|---|---|---|---|
| Risperdal Instasolv 0,5 mg munsönderfallande tabletter | DE/H/2184/016 | 17263 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 3 mg munsönderfallande tabletter | DE/H/2184/011 | 21348 | JANSSEN-CILAG OY | FI |
| Risperdal 4 mg kalvopäällysteiset tabletit | DE/H/2184/006/MR | 11481 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 1 mg munsönderfallande tabletter | DE/H/2184/009 | 17264 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 2 mg munsönderfallande tabletter | DE/H/2184/010 | 17265 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg kalvopäällysteiset tabletit | DE/H/2184/003 | 11478 | JANSSEN-CILAG OY | FI |
| Risperdal 0,5 mg filmdragerade tabletter | DE/H/2184/002 | 13364 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg/ml oral lösning | DE/H/2184/008 | 12113 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 4 mg munsönderfallande tabletter | DE/H/2184/012 | 21349 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 50 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/015 | 16896 | JANSSEN-CILAG OY | FI |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681609 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3389493 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3432645 5 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637442 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681561 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681584 1 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 982 5 4 | JANSSEN-CILAG | FR |

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|---|---|---|---|---|
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3432639 4 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637407 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637413 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637459 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637465 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34009 566 553 7 3 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3389470 2 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 980 2 5 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637502 4 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637399 0 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637494 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681555 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681590 2 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 984 8 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 5665572 4 | JANSSEN-CILAG | FR |
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3389530 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34009 566 554 3 4 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 5665566 3 | JANSSEN-CILAG | FR |

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|---|---|---|---|---|
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 983 1 5 | JANSSEN-CILAG | FR |
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3389547 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637488 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681549 0 | JANSSEN-CILAG | FR |
| Risperdal 0,5 mg Filmtabletten | DE/H/2184/002 | 2001040022 | JANSSEN-CILAG NV | LU |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | 2004058265 | JANSSEN-CILAG NV | LU |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | 2004058266 | JANSSEN-CILAG NV | LU |
| Risperdal 3 mg Filmtabletten | DE/H/2184/005 | 2004058267 | JANSSEN-CILAG NV | LU |
| Risperdal 6 mg Filmtabletten | DE/H/2184/007 | 2008089903 | JANSSEN-CILAG NV | LU |
| Risperdal 4 mg Filmtabletten | DE/H/2184/006 | 2004058268 | JANSSEN-CILAG NV | LU |
| Risperdal Consta 50 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/015 | 2004010028 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 2 mg Schmelztabletten | DE/H/2184/010 | 2003050032 | JANSSEN-CILAG NV | LU |
| Risperdal Consta 37,5 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/014 | 2004010027 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 0,5 mg Schmelztabletten | DE/H/2184/016 | 2003050030 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 1 mg Schmelztabletten | DE/H/2184/009 | 2003050031 | JANSSEN-CILAG NV | LU |

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|---|---|---|---|---|
| Risperdal Consta 25 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/013 | 2004010026 | JANSSEN-CILAG NV | LU |
| RISPERDAL 2 mg filmsko obložene tablete | DE/H/2184/004 | H/96/01358/014 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 50 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/015 | H/96/01358/003 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 1 mg filmsko obložene tablete | DE/H/2184/003 | H/96/01358/008 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 25 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/013 | H/96/01358/001 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL® 1 mg/ml peroralna raztopina | DE/H/2184/008 | H/96/01358/004 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 37,5 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/014 | H/96/01358/002 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 4 mg filmsko obložene tablete | DE/H/2184/006 | H/96/01358/024 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 3 mg filmsko obložene tablete | DE/H/2184/005 | H/96/01358/020 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |

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|---|---|---|---|---|
| RISPOLEPT CONSTA 25 mg pulbere si solvent pentru suspensie injectabila cu eliberare prelungita | DE/H/2184/013 | 6876/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT 1mg/ml soluţie orală | DE/H/2184/008 | 6875/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT CONSTA 50 mg pulbere şi solvent pentru suspensie injectabilă cu eliberare prelungită | DE/H/2184/015 | 6878/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT CONSTA 37,5 mg pulbere si solvent pentru suspensie injectabila cu eliberare prelungita | DE/H/2184/014 | 6877/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT 1mg/ml soluţie orală | DE/H/2184/008 | 6875/2014/02 | JANSSEN PHARMACEUTICA NV | RO |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | PL 00242/0376 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | PL 00242/0375 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | PL 00242/0377 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL CONSTA 50 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/015 | 68/070/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 4 mg, potahované tablety | DE/H/2184/006 | 68/185/95-D/C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 1 mg potahované tablety | DE/H/2184/003 | 68/185/95-A/C | JANSSEN-CILAG S.R.O | CZ |

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|---|---|---|---|---|
| RISPERDAL, 1 mg/ml, perorální roztok | DE/H/2184/008 | 68/298/99-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL CONSTA 37,5 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/014 | 68/069/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 3 mg, potahované tablety | DE/H/2184/005 | 68/185/95-C/C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL CONSTA 25 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/013 | 68/068/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 2 mg, potahované tablety | DE/H/2184/004 | 68/185/95-B/C | JANSSEN-CILAG S.R.O | CZ |
| Risperdal Consta 25 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/013 | 17868 | JANSSEN-CILAG AB | SE |
| Risperdal Consta 50 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/015 | 01-9860 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 37,5 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/014 | 01-9859 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 50 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/015 | 17870 | JANSSEN-CILAG AB | SE |
| Risperdal Consta 37,5 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/014 | IS/1/02/132/02 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 37,5 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/014 | 16895 | JANSSEN-CILAG OY | FI |

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|---|---|---|---|---|
| Risperdal Consta 25 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/013 | IS/1/02/132/01 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 25 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/013 | 16894 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 25 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/013 | 01-9858 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 50 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/015 | IS/1/02/132/03 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 50 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/015 | 16896 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 37,5 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/014 | 17869 | JANSSEN-CILAG AB | SE |
| RISPERDAL 1 mg film-coated tablets | DE/H/2184/003 | MA018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | MA018/00203 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | MA018/00203 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 1 mg film-coated tablets | DE/H/2184/003 | 018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 1mg/ml oral solution | DE/H/2184/008 | MA018/00204 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 3 mg film-coated tablets | DE/H/2184/005 | 018/00202 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 3 mg film-coated tablets | DE/H/2184/005 | MA018/00202 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 1 mg film-coated tablets | DE/H/2184/003 | 018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |

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|---|---|---|---|---|
| RISPERDAL 1 mg, filmomhulde tabletten | DE/H/2184/003 | RVG 16096 | JANSSEN-CILAG BV | NL |
| RISPERDAL 4 mg, filmomhulde tabletten | DE/H/2184/006 | RVG 16099 | JANSSEN-CILAG BV | NL |
| RISPERDAL 3 mg, filmomhulde tabletten | DE/H/2184/005 | RVG 16098 | JANSSEN-CILAG BV | NL |
| RISPERDAL 1 mg/ml drank | DE/H/2184/008 | RVG 19127 | JANSSEN-CILAG BV | NL |
| RISPERDAL 0,5 mg, filmomhulde tabletten | DE/H/2184/002 | RVG 22714 | JANSSEN-CILAG BV | NL |
| RISPERDAL 2 mg, filmomhulde tabletten | DE/H/2184/004 | RVG 16097 | JANSSEN-CILAG BV | NL |
| RISPERDAL 1 mg/ml oral solution | DE/H/2184/008 | PL 00242/0199 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 0.5mg orodispersible tablets | DE/H/2184/016 | PL 00242/0378 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL QUICKLET 4 mg, orodispergeerbare tabletten | DE/H/2184/012 | RVG 31776 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 25 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/013 | RVG 27178 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 1 mg, orodispergeerbare tabletten | DE/H/2184/009 | RVG 27791 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 2 mg, orodispergeerbare tabletten | DE/H/2184/010 | RVG 27792 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 50 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/015 | RVG 27180 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 3 mg, orodispergeerbare tabletten | DE/H/2184/011 | RVG 31775 | JANSSEN-CILAG BV | NL |

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|---|---|---|---|---|
| RISPERDAL CONSTA 37,5 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/014 | RVG 27179 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | 018/00205 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | MA018/00205-7 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | MA018/00205-7 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg filmomhulde tabletten | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | BE |
| RISPERDAL 6 mg comprimés pelliculés | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | BE |
| RISPERDAL 3 mg comprimés pelliculés | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg filmomhulde tabletten | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg filmomhulde tabletten | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| RISPERDAL 4 mg comprimés pelliculés | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| RISPERDAL 4 mg filmomhulde tabletten | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| RISPERDAL 6 mg filmomhulde tabletten | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | BE |
| RISPERDAL 3 mg filmomhulde tabletten | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 0,5 mg orodispergeerbare tabletten | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1mg/ml solution buvable | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 2 mg orodispergeerbare tabletten | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg/ml drank | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 37,5 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/014 | BE254606 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 2 mg comprimés orodispersibles | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 50 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/015 | BE254615 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg orodispergeerbare tabletten | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 25 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/013 | BE254597 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg comprimés orodispersibles | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg comprimés orodispersibles | DE/H/2184/009 | 2003050031 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| RISPERDAL 2 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/004 | 14397 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/013 | 33091 | JANSSEN-CILAG A/S | DK |
| RISPERDAL CONSTA 37,5 mg prášok a disperzné prostredie na intramuskulárnu injekčnú suspenziu s predĺženým uvoľňovaním | DE/H/2184/014 | 68/0106/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL CONSTA 25 mg prášok a disperzné prostredie na intramuskulárnu injekcnú suspenziu s predlženým uvolnovaním | DE/H/2184/013 | 68/0105/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL 3 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/005 | 14398 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL 4 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/006 | 14399 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/014 | 33092 | JANSSEN-CILAG A/S | DK |
| RISPERDAL 1 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/003 | 14396 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL 1 mg/ml πόσιμο διάλυμα | DE/H/2184/008 | 17844 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/015 | 33093 | JANSSEN-CILAG A/S | DK |
| RISPERDAL INSTASOLV 0,5 mg comprimés orodispersibles | DE/H/2184/016 | 2003050030 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA50 mg prášok a disperzné prostredie na intramuskulárnu injekčnú suspenziu s predĺženým uvoľňovaním | DE/H/2184/015 | 68/0107/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL INSTASOLV 2 mg comprimés orodispersibles | DE/H/2184/010 | 2003050032 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA 37,5 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/014 | 2004010027 | JANSSEN-CILAG NV | LU |
| RISPERDAL 4 mg comprimés pelliculés | DE/H/2184/006 | 2004058268 | JANSSEN-CILAG NV | LU |
| RISPERDAL 1mg/ml solution buvable | DE/H/2184/008 | 2008089904 | JANSSEN-CILAG NV | LU |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | 2004058265 | JANSSEN-CILAG NV | LU |
| RISPERDAL 3 mg comprimés pelliculés | DE/H/2184/005 | 2004058267 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA 25 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/013 | 2004010026 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA 50 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/015 | 2004010028 | JANSSEN-CILAG NV | LU |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | 2001040022 | JANSSEN-CILAG NV | LU |
| RISPERDAL 6 mg comprimés pelliculés | DE/H/2184/007 | 2008089903 | JANSSEN-CILAG NV | LU |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | 2004058266 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| RISPERDAL 4 mg smeltetabletter | DE/H/2184/012 | 04-3141 | JANSSEN-CILAG A/S | NO |
| Risperdal 4 mg frystorkade tabletter | DE/H/2184/012 | 21886 | JANSSEN-CILAG AB | SE |
| RISPERDAL INSTASOLV 0,5 mg tabletti, suussa hajoava | DE/H/2184/016 | 17263 | JANSSEN-CILAG OY | FI |
| Risperdal 0,5 mg munndreifitöflur | DE/H/2184/016 | IS/1/03/119/01 | JANSSEN-CILAG AB | IS |
| Risperdal 3 mg frystorkade tabletter | DE/H/2184/011 | 21885 | JANSSEN-CILAG AB | SE |
| RISPERDAL QUICKLET 1 mg, Schmelztabletten | DE/H/2184/009 | 28754.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 4 mg, Schmelztabletten | DE/H/2184/012 | 28754.03.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL Lösung 1 mg/ml, Lösung zum Einnehmen | DE/H/2184/008 | 35950.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 3 mg smeltetabletter | DE/H/2184/011 | 04-3140 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg/ml oral lösning | DE/H/2184/008 | 12677 | JANSSEN-CILAG AB | SE |
| RISPERDAL 1 mg/ml belsőleges oldat | DE/H/2184/008 | OGYI-T-8812/07 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL 1 mg/ml oral solution | DE/H/2184/008 | PA 0748/003/001 | JANSSEN-CILAG LIMITED | IE |
| Rispolept, 2 mg, tabletki powlekane | not available | R/6705 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 1 mg munndreifitöflur | DE/H/2184/009 | IS/1/03/119/02 | JANSSEN-CILAG AB | IS |
| RISPERDAL INSTASOLV 1 mg tabletti, suussa hajoava | DE/H/2184/009 | 17264 | JANSSEN-CILAG OY | FI |
| Rispolept, 4 mg, tabletki powlekane | not available | R/6707 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT, 1 mg/ml suukaudne lahus | DE/H/2184/008 | 273799 | UAB JOHNSON & JOHNSON | EE |
| Risperdal 1 mg/ml soluzione orale | DE/H/2184/008 | 028752145 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| Risperdal 2 mg munndreifitöflur | DE/H/2184/010 | IS/1/03/119/03 | JANSSEN-CILAG AB | IS |
| RISPERDAL 1 mg/ml oraaliliuos | DE/H/2184/008 | 12113 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 2 mg tabletti, suussa hajoava | DE/H/2184/010 | 17265 | JANSSEN-CILAG OY | FI |
| RISPERDAL QUICKLET 2 mg, Schmelztabletten | DE/H/2184/010 | 28754.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 0,5 mg, Schmelztabletten | DE/H/2184/016 | 28754.04.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 3 mg, Schmelztabletten | DE/H/2184/011 | 28754.02.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept 1 mg/ml geriamasis tirpalas | DE/H/2184/008 | LT/1/98/0215/001 | UAB JOHNSON & JOHNSON | LT |
| Risperdal 1 mg/ml soluzione orale | DE/H/2184/008 | 028752095 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 1 mg/ml mikstur, oppløsning | DE/H/2184/008 | 95-0684 | JANSSEN-CILAG A/S | NO |
| RISPERDAL FLAS 4 mg comprimidos bucodispersables | DE/H/2184/012 | 67.051 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL FLAS 2 mg comprimidos bucodispersables | DE/H/2184/010 | 65.698 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL FLAS 3 mg comprimidos bucodispersables | DE/H/2184/011 | 67.052 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL FLAS 0,5 mg comprimidos bucodispersables | DE/H/2184/016 | 65.696 | JANSSEN-CILAG S.A. | ES |
| Rispolept 1 mg/ml šķīdums iekšķīgai lietošanai | DE/H/2184/008 | 01-0372 | UAB JOHNSON & JOHNSON | LV |
| RISPERDAL FLAS 1 mg comprimidos bucodispersables | DE/H/2184/009 | 65.697 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 1 mg/ml solución oral | DE/H/2184/008 | 62.096 | JANSSEN-CILAG S.A. | 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 |
|---|---|---|---|---|
| Risperdal, smeltetabletter | DE/H/2184/009 | 33697 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/002 | 30230 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/004 | 14977 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/003 | 14976 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/006 | 14979 | JANSSEN-CILAG A/S | DK |
| Risperdal, oral opløsning | DE/H/2184/008 | 17570 | JANSSEN-CILAG A/S | DK |
| Risperdal | DE/H/2184/016 | 33696 | JANSSEN-CILAG A/S | DK |
| Risperdal, smeltetabletter | DE/H/2184/010 | 33698 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/005 | 14978 | JANSSEN-CILAG A/S | DK |
| Risperdal 2 mg tabletti, kalvopäällysteinen | DE/H/2184/004 | 11479 | JANSSEN-CILAG OY | FI |
| Risperdal 0,5 mg tabletti, kalvopäällysteinen | DE/H/2184/002 | 13364 | JANSSEN-CILAG OY | FI |
| Risperdal 4 mg tabletti, kalvopäällysteinen | DE/H/2184/006 | 11481 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg tabletti, kalvopäällysteinen | DE/H/2184/003 | 11478 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 4 mg tabletti, suussa hajoava | DE/H/2184/012 | 21349 | JANSSEN-CILAG OY | FI |
| Risperdal 6 mg tabletti, kalvopäällysteinen | DE/H/2184/007 | 12302 | JANSSEN-CILAG OY | FI |
| Risperdal 3 mg tabletti, kalvopäällysteinen | DE/H/2184/005 | 11480 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 3 mg tabletti, suussa hajoava | DE/H/2184/011 | 21348 | JANSSEN-CILAG OY | FI |
| Rispolept, 1 mg, tabletki powlekane | DE\H\2184\003\R\001 | R/6704 | JANSSEN-CILAG INTERNATIONAL NV | 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 |
|---|---|---|---|---|
| RISPOLEPT 4 mg, õhukese polümeerikattega tabletid | DE/H/2184/006 | 179797 | UAB JOHNSON & JOHNSON | EE |
| Rispolept, 1 mg/ml, roztwór doustny | DE/H/2184/008 | 4238 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 1 mg/ml mixtúra, lausn | DE/H/2184/008 | 950155 | JANSSEN-CILAG AB | IS |
| RISPERDAL CONSTA 25 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/013 | 65.213 | JANSSEN-CILAG S.A. | ES |
| RISPOLEPT 2 mg, õhukese polümeerikattega tabletid | DE/H/2184/004 | 179597 | UAB JOHNSON & JOHNSON | EE |
| RISPOLEPT 3 mg, õhukese polümeerikattega tabletid | DE/H/2184/005 | 179697 | UAB JOHNSON & JOHNSON | EE |
| Rispolept, 3 mg, tabletki powlekane | not available | R/6706 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPERDAL CONSTA 50 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/015 | 65.214 | JANSSEN-CILAG S.A. | ES |
| RISPOLEPT 1 mg, õhukese polümeerikattega tabletid | DE/H/2184/003 | 179497 | UAB JOHNSON & JOHNSON | EE |
| RISPERDAL CONSTA 37,5 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/014 | 65.215 | JANSSEN-CILAG S.A. | ES |
| Risperdal 2 mg compresse rivestite con film | DE/H/2184/004 | 028752020 | JANSSEN-CILAG SPA | IT |
| Risperdal 1 mg compresse rivestite con film | DE/H/2184/003 | 028752057 | JANSSEN-CILAG SPA | IT |
| Risperdal 2 mg compresse rivestite con film | DE/H/2184/004 | 028752069 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 4 mg compresse rivestite con film | DE/H/2184/006 | AIC 028752083 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| Risperdal 3 mg compresse rivestite con film | DE/H/2184/005 | 028752071 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 1 mg, Filmtabletten | DE/H/2184/003 | 28758.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 4 mg, Filmtabletten | DE/H/2184/006 | 28758.03.00 | JANSSEN-CILAG GMBH | DE |
| Risperdal 1 mg compresse rivestite con film | DE/H/2184/003 | 028752018 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 4 mg compresse rivestite con film | DE/H/2184/006 | AIC 028752044 | JANSSEN-CILAG SPA | IT |
| Rispolept 3 mg plėvele dengtos tabletės | DE/H/2184/005 | LT/1/96/0215/004 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 2 mg apvalkotās tabletes | DE/H/2184/004 | 98-0100 | UAB JOHNSON & JOHNSON | LV |
| Risperdal 3 mg compresse rivestite con film | DE/H/2184/005 | 028752032 | JANSSEN-CILAG SPA | IT |
| Rispolept 4 mg plėvele dengtos tabletės | DE/H/2184/006 | LT/1/96/0215/005 | UAB JOHNSON & JOHNSON | LT |
| RISPERDAL 6 mg, Filmtabletten | DE/H/2184/007 | 37961.00.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept 4 mg apvalkotās tabletes | DE/H/2184/006 | 98-0102 | UAB JOHNSON & JOHNSON | LV |
| Rispolept 2 mg plėvele dengtos tabletės | DE/H/2184/004 | LT/1/96/0215/003 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 1 mg plėvele dengtos tabletės | DE/H/2184/003 | LT/1/96/0215/002 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 3 mg apvalkotās tabletes | DE/H/2184/005 | 98-0101 | UAB JOHNSON & JOHNSON | LV |
| Rispolept 1 mg apvalkotās tabletes | DE/H/2184/003 | 98-0099 | UAB JOHNSON & JOHNSON | LV |
| RISPERDAL 0,5 mg, Filmtabletten | DE/H/2184/002 | 43776.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 2 mg, Filmtabletten | DE/H/2184/004 | 28758.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 3 mg, Filmtabletten | DE/H/2184/005 | 28758.02.00 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| Risperdal 2 mg filmdragerade tabletter | DE/H/2184/004 | 11993 | JANSSEN-CILAG AB | SE |
| Risperdal 1 mg filmdragerade tabletter | DE/H/2184/003 | 11992 | JANSSEN-CILAG AB | SE |
| Risperdal 0,5 mg filmdragerade tabletter | DE/H/2184/002 | 14318 | JANSSEN-CILAG AB | SE |
| Risperdal 4 mg filmdragerade tabletter | DE/H/2184/006 | 11995 | JANSSEN-CILAG AB | SE |
| Risperdal 3 mg filmdragerade tabletter | DE/H/2184/005 | 11994 | JANSSEN-CILAG AB | SE |
| RISPERDAL 6 mg film-coated tablets | DE/H/2184/007 | PA 0748/003/002 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0,5 mg tabletter, filmdrasjerte | DE/H/2184/002 | 98-4328 | JANSSEN-CILAG A/S | NO |
| RISPERDAL 2 mg tabletter, filmdrasjerte | DE/H/2184/004/MR | 8038 | JANSSEN-CILAG A/S | NO |
| RISPERDAL 1 mg Filmtabletten | DE/H/2184/003 | 1-20297 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 4 mg filmuhúðaðar töflur | DE/H/2184/006 | 920126 | JANSSEN-CILAG AB | IS |
| RISPERDAL 2 mg Filmtabletten | DE/H/2184/004 | 1-20301 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 2 mg filmuhúðaðar töflur | DE/H/2184/004 | 920124 | JANSSEN-CILAG AB | IS |
| RISPERDAL 3 mg tabletter, filmdrasjerte | DE/H/2184/005 | 8039 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg filmuhúðaðar töflur | DE/H/2184/003 | 920123 | JANSSEN-CILAG AB | IS |
| RISPERDAL 3 mg Filmtabletten | DE/H/2184/005 | 1-20302 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 3 mg filmuhúðaðar töflur | DE/H/2184/005 | 920125 | JANSSEN-CILAG AB | IS |
| RISPERDAL 1 mg/ml Lösung zum Einnehmen | DE/H/2184/008 | 1-21466 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 0,5 mg filmuhúðaðar töflur | DE/H/2184/002 | 980135 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| RISPERDAL 4 mg Filmtabletten | DE/H/2184/006 | 1-20303 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 6 mg Filmtabletten | DE/H/2184/007 | 1-22846 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 1 mg tabletter, filmdrasjerte | DE/H/2184/003 | 8037 | JANSSEN-CILAG A/S | NO |
| RISPERDAL 4 mg tabletter, filmdrasjerte | DE/H/2184/006 | 8040 | JANSSEN-CILAG A/S | NO |
| RISPERDALCONSTA L.P. 50 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/015 | 34009 362 494 2 4 | JANSSEN-CILAG | FR |
| RISPERDALCONSTA L.P. 37,5 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/014 | 34 00 9 3624936 3 | JANSSEN-CILAG | FR |
| RISPERDALCONSTA L.P. 25 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/013 | 34 00 9 3624913 4 | JANSSEN-CILAG | FR |
| RISPERDAL CONSTA 50 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/015 | 1-24630 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL CONSTA 25 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/013 | 1-24628 | JANSSEN-CILAG PHARMA 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 |
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| RISPERDAL CONSTA 37,5 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/014 | 1-24629 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 3 mg comprimidos recubiertos con película | DE/H/2184/005 | 60.335 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 1 mg comprimidos recubiertos con película | DE/H/2184/003 | 60.336 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 6 mg comprimidos recubiertos con película | DE/H/2184/007 | 62.803 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL CONSTA50 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/015 | 19589 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL CONSTA 25 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/013 | 52995.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL CONSTA 50 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/015 | 52995.02.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept Consta, 50 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu | DE/H/2184/015 | 10580 | JANSSEN-CILAG INTERNATIONAL NV | 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 |
|---|---|---|---|---|
| RISPOLEPT CONSTA 37,5 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/014 | 03-0097 | UAB JOHNSON & JOHNSON | LV |
| RISPOLEPT CONSTA 37,5 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/014 | LT/1/03/3651/002 | UAB JOHNSON & JOHNSON | LT |
| Rispolept Consta 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | 20030321 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | BG |
| RISPOLEPT CONSTA 50 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/015 | LT/1/03/3651/003 | UAB JOHNSON & JOHNSON | LT |
| Rispolept Consta, 25 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu. | DE/H/2184/013 | 10582 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 4 mg munndreifitöflur | DE/H/2184/012 | IS/1/06/139/02 | JANSSEN-CILAG AB | IS |
| Rispolept Consta, 37,5 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu. | DE/H/2184/014 | 10581 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT CONSTA 25 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/013 | 03-0096 | UAB JOHNSON & JOHNSON | LV |
| RISPOLEPT CONSTA 25 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/013 | LT/1/03/3651/001 | UAB JOHNSON & JOHNSON | 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 |
|---|---|---|---|---|
| RISPOLEPT CONSTA 25 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/013 | 410603 | UAB JOHNSON & JOHNSON | EE |
| RISPOLEPT CONSTA 50 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/015 | 410803 | UAB JOHNSON & JOHNSON | EE |
| RISPERDAL CONSTA 37,5 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/014 | 52995.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPOLEPT CONSTA 37,5 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/014 | 410703 | UAB JOHNSON & JOHNSON | EE |
| RISPOLEPT CONSTA 50 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/015 | 03-0098 | UAB JOHNSON & JOHNSON | LV |
| Risperdal 25 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/013 | 028752172 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 50 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/015 | AIC N° 028752196 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 37,5 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/014 | AIC N° 028752184 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA 25 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/013 | 67148/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperdal Consta 37,5 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/014 | OGYI-T-8812/04 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL CONSTA 50 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/015 | 67150/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperdal Consta 50 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/015 | OGYI-T-8812/06 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL Quicklet 0,5 mg Schmelztabletten | DE/H/2184/016 | 1-25155 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL Quicklet 4 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/012 | 67146/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL CONSTA 37,5 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/014 | 67149/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperdal Consta 25 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/013 | OGYI-T-8812/02 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3389487 0 | JANSSEN-CILAG | 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 |
|---|---|---|---|---|
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3442738 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3389501 3 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 981 9 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637471 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681532 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637436 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637382 2 | JANSSEN-CILAG | FR |
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | PA 0748/003/012 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | PA 0748/003/011 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 4219887 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 4219689 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 4 mg comprimido orodispersível | DE/H/2184/012 | 5827589 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 4219788 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1mg comprimidos revestidos por película | DE/H/2184/003 | 2305985 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 2 mg comprimidos revestidos por película | DE/H/2184/004 | 2306082 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg/ml solução oral | DE/H/2184/008 | 2527984 | JANSSEN FARMACÊUTICA PORTUGAL, 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 |
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| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 4219986 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220182 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220083 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 3 mg comprimido orodispersível | DE/H/2184/011 | 5827381 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1mg comprimidos revestidos por película | DE/H/2184/003 | 2305787 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 4 mg comprimido orodispersível | DE/H/2184/012 | 5827480 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 3 mg comprimido orodispersível | DE/H/2184/011 | 5827282 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 50 mg pó e veículo para suspensão de libertação prolongada para injecção intramuscular | DE/H/2184/015 | 4753686 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681578 0 | JANSSEN-CILAG | FR |
| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 5701289 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 25 mg pó e veículo para suspensão de libertação prolongada para injeção intramuscular | DE/H/2184/013 | 4753588 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL Quicklet 3 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/011 | 67145/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 1 mg/ml solução oral | DE/H/2184/008 | 2715381 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 5701180 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL Quicklet 2 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/010 | 67144/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | 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 |
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| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220083 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 37,5 mg pó e veículo para suspensão de libertação prolongada para injeção intramuscular | DE/H/2184/014 | 4753687 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA37,5 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/014 | 19588 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL CONSTA 25 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/013 | 19587 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL Quicklet 1 mg Schmelztabletten | DE/H/2184/009 | 1-25156 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 3 mg munndreifitöflur | DE/H/2184/011 | IS/1/06/139/01 | JANSSEN-CILAG AB | IS |
| RISPERDAL Quicklet 2 mg Schmelztabletten | DE/H/2184/010 | 1-25157 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 2 mg comprimidos revestidos por película | DE/H/2184/004 | 2306181 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg comprimidos revestidos por película | DE/H/2184/003 | 2305886 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 2 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/004 | 67138/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 4 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/006 | 67140/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | 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 |
|---|---|---|---|---|
| RISPERDAL 4 mg comprimidos revestidos por película | DE/H/2184/006 | 2306587 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/003 | 67137/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 4 mg comprimidos revestidos por película | DE/H/2184/006 | 2306488 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 3 mg comprimidos revestidos por película | DE/H/2184/005 | 2306389 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg/ml πόσιμο διάλυμα | DE/H/2184/008 | 67147/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 3 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/005 | 67139/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 6 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/007 | 67140/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 0,5 mg comprimidos revestidos por película | DE/H/2184/002 | 3219888 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 3 mg comprimidos revestidos por película | DE/H/2184/005 | 2306280 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | PA 0748/003/010 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 3 mg film-coated tablets | DE/H/2184/005 | PA 0748/003/006 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 2 mg smeltetabletter | DE/H/2184/010 | 02-1026 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg compresse orodispersibili | DE/H/2184/009 | 028752222 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| Risperdal 1 mg frystorkade tabletter | DE/H/2184/009 | 18385 | JANSSEN-CILAG AB | SE |
| RISPERDAL 1mg film-coated tablets | DE/H/2184/003 | PL 00242/0186 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 4 mg film-coated tablets | DE/H/2184/006 | PA 0748/003/007 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL Quicklet 1 mg orodispersible tablets | DE/H/2184/009 | PA 0748/003/014 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0.5 mg film-coated tablets | DE/H/2184/002 | PA 0748/003/009 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | PA 0748/003/005 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL Quicklet 3 mg orodispersible tablets | DE/H/2184/011 | PA 0748/003/016 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 3mg film-coated tablets | DE/H/2184/005 | PL 00242/0188 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 1 mg smeltetabletter | DE/H/2184/009 | 02-1025 | JANSSEN-CILAG A/S | NO |
| RISPERDAL Quicklet 4 mg orodispersible tablets | DE/H/2184/012 | PL 0242/0408 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 1 mg orodispersible tablets | DE/H/2184/009 | PL 0242/0379 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 2mg orodispersible tablets | DE/H/2184/010 | PL 0242/0380 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 2mg film-coated tablets | DE/H/2184/004 | PL 00242/0187 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 6mg film-coated tablets | DE/H/2184/007 | PL 00242/0317 | JANSSEN-CILAG LIMITED | UK |
| Risperdal 0,5 mg frystorkade tabletter | DE/H/2184/016 | 18384 | JANSSEN-CILAG AB | SE |
| RISPERDAL 2 mg compresse orodispersibili | DE/H/2184/010 | 028752259 | JANSSEN-CILAG SPA | IT |
| RISPERDAL Quicklet 4 mg orodispersible tablets | DE/H/2184/012 | PA 0748/003/017 | JANSSEN-CILAG LIMITED | IE |
| Risperdal 2 mg frystorkade tabletter | DE/H/2184/010 | 18386 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| RISPERDAL 0.5mg film-coated tablets | DE/H/2184/002 | PL 00242/0347 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 0.5mg orodispersible tablets | DE/H/2184/016 | PA0748/003/013 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0,5 mg smeltetabletter | DE/H/2184/016 | 02-1024 | JANSSEN-CILAG A/S | NO |
| RISPERDAL Quicklet 3 mg orodispersible tablets | DE/H/2184/011 | PL 0242/0407 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 1 mg compresse orodispersibili | DE/H/2184/009 | 028752234 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 2 mg compresse orodispersibili | DE/H/2184/010 | 028752246 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 4mg film-coated tablets | DE/H/2184/006 | PL 00242/0189 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 2 mg orodispersible tablets | DE/H/2184/010 | PA 0748/003/015 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 1 mg film-coated tablets | DE/H/2184/003 | PA 0748/003/04 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL INSTASOLV 0,5 mg comprimés orodispersibles | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| RISPERDAL Quicklet 0,5 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/016 | 67142/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL Quicklet 1 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/009 | 67143/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperidone EG 8 mg comprimés pelliculés | not available | 0019/09110045 | EUROGENERICS N.V./S.A. | LU |
| Risperidone EG 8 mg filmomhulde tabletten | not available | BE329813 | EUROGENERICS N.V./S.A. | BE |
| Risperidone EG 8 mg comprimés pelliculés | not available | BE329813 | EUROGENERICS N.V./S.A. | BE |
| Risperidone EG 8 mg Filmtabletten | not available | BE329813 | EUROGENERICS N.V./S.A. | BE |
| Risperidon - 1 A Pharma 0,25 mg Filmtabletten | DE/H/0923/001 | 67593.00.00 | 1 A PHARMA 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 |
|---|---|---|---|---|
| Risperidon HEXAL 0,25 mg Filmtabletten | DE/H/0922/001 | 67607.00.00 | HEXAL AG | DE |
| Risperidon Sandoz 0,25 mg Filmtabletten | DE/H/0924/001 | 67727.00.00 | HEXAL AG | DE |
| Risperidon - 1 A Pharma 6 mg Filmtabletten | DE/H/0923/002 | 67594.00.00 | 1 A PHARMA GMBH | DE |
| Risperidon Sandoz 6 mg Filmtabletten | DE/H/0924/002 | 67728.00.00 | HEXAL AG | DE |
| Risperidon HEXAL 6 mg Filmtabletten | DE/H/0922/002 | 67608.00.00 | HEXAL AG | DE |
| Risperidon HEXAL 0,5 mg Filmtabletten | DE/H/0792/001 | 66671.00.00 | HEXAL AG | DE |
| Risperidon - 1 A Pharma 0,5 mg Filmtabletten | DE/H/0793/001 | 66672.00.00 | 1 A PHARMA GMBH | DE |
| Risperidon Sandoz 0,5 mg Filmtabletten | DE/H/0794/001 | 66673.00.00 | HEXAL AG | DE |
| Risperanne, filmovertrukne tabletter | DE/H/0794/001 | 39628 | SANDOZ A/S | DK |
| Risperidon Sandoz 0,25 mg - Filmtabletten | DE/H/0924/001 | 1-27295 | SANDOZ GMBH | AT |
| Risperidon Sandoz 0,5 mg - Filmtabletten | DE/H/0794/001 | 1-27042 | SANDOZ GMBH | AT |
| Risperidon Hexal 0,5 mg – Filmtabletten | DE/H/0792/001 | 1-27039 | HEXAL PHARMA GMBH | AT |
| Risperidon 1A Pharma 0,5 mg – Filmtabletten | DE/H/0793/001 | 1-27040 | 1A PHARMA GMBH | AT |
| Risperidon Sandoz 0,5 mg tabletti, kalvopäällysteinen | DE/H/0794/001 | 22442 | SANDOZ A/S | FI |
| Risperidon Sandoz 0,5 mg filmdragerade tabletter | DE/H/0794/001 | 24184 | SANDOZ A/S | SE |
| Risperidon Sandoz 0,25 mg filmdragerad tablett | DE/H/0924/001 | 42117 | SANDOZ A/S | SE |
| Risperidone 4mg Tablets | FI/H/0373/004 | PL 04416/0665 | SANDOZ LTD | UK |
| Risperidone 1mg Tablets | FI/H/0373/001 | PL 04416/0662 | SANDOZ LTD | UK |

| 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 |
|---|---|---|---|---|
| Risperidone 2mg Tablets | FI/H/0373/002 | PL 04416/0663 | SANDOZ LTD | UK |
| Risperidone 3mg Tablets | FI/H/0373/003 | PL 04416/0664 | SANDOZ LTD | UK |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/001 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/002 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/003 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/004 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/005 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/006 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/007 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/008 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/009 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/010 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 0,5 mg orodisperzibilni filmi | DE/H/2441/001 | H/10/01968/011 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/012 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/013 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/014 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/015 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/016 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |

| 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 |
|---|---|---|---|---|
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/017 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/018 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/019 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/020 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/021 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 1 mg orodisperzibilni filmi | DE/H/2441/002 | H/10/01968/022 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/023 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/024 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/025 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/026 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/027 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/028 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/029 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/030 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/031 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/032 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 2 mg orodisperzibilni filmi | DE/H/2441/003 | H/10/01968/033 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/034 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |

| 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 |
|---|---|---|---|---|
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/035 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/036 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/037 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/038 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/039 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/040 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/041 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/042 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/043 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| Rispolux Neo 3 mg orodisperzibilni filmi | DE/H/2441/004 | H/10/01968/044 | LEK PHARMACEUTICALS D.D. LJUBLJANA | SI |
| RISPERIDONE EG 1 mg/ml, solution buvable | not available | NL32169 | EG LABO LABORATOIRES EUROGENERICS | FR |
| Risperidona Farmalider 4 mg comprimidos recubiertos con película | not available | 66806 | FARMALIDER, S.A. | ES |
| RISPERDAL 0,5 mg filmomhulde tabletten | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg filmomhulde tabletten | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg filmomhulde tabletten | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| Risperdal 4 mg Filmtabletten | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 50 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/015 | BE254615 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 2 mg Schmelztabletten | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg Filmtabletten | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | BE449617 | JANSSEN-CILAG NV | BE |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | BE |
| Risperdal 3 mg Filmtabletten | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 0,5 mg Schmelztabletten | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| Risperdal 0,5 mg Filmtabletten | DE/H/2184/002 | BE449600 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 25 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/013 | BE254597 | JANSSEN-CILAG NV | BE |
| Risperdal Instasolv 1 mg Schmelztabletten | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| Risperdal Consta 37,5 mg, Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/014 | BE254606 | JANSSEN-CILAG NV | BE |
| Risperdal 6 mg Filmtabletten | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| Risperdal 1 mg/ml Lösung zum Einnehmen | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | BE449626 | JANSSEN-CILAG NV | BE |
| Risperdal 2 mg filmdragerade tabletter | DE/H/2184/004 | 11479 | JANSSEN-CILAG OY | FI |
| Risperdal 3 mg filmdragerade tabletter | DE/H/2184/005 | 11480 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 37,5 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/014 | 16895 | JANSSEN-CILAG OY | FI |
| Risperdal 6 mg kalvopäällysteiset tabletit | DE/H/2184/007 | 12302 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 25 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/013 | 16894 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 0,5 mg munsönderfallande tabletter | DE/H/2184/016 | 17263 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 3 mg munsönderfallande tabletter | DE/H/2184/011 | 21348 | JANSSEN-CILAG OY | FI |
| Risperdal 4 mg kalvopäällysteiset tabletit | DE/H/2184/006/MR | 11481 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 1 mg munsönderfallande tabletter | DE/H/2184/009 | 17264 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 2 mg munsönderfallande tabletter | DE/H/2184/010 | 17265 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg kalvopäällysteiset tabletit | DE/H/2184/003 | 11478 | JANSSEN-CILAG OY | FI |
| Risperdal 0,5 mg filmdragerade tabletter | DE/H/2184/002 | 13364 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg/ml oral lösning | DE/H/2184/008 | 12113 | JANSSEN-CILAG OY | FI |
| Risperdal Instasolv 4 mg munsönderfallande tabletter | DE/H/2184/012 | 21349 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| Risperdal Consta 50 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/015 | 16896 | JANSSEN-CILAG OY | FI |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681609 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3389493 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3432645 5 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637442 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681561 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681584 1 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 982 5 4 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3432639 4 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637407 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637413 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637459 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637465 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34009 566 553 7 3 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3389470 2 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 980 2 5 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637502 4 | JANSSEN-CILAG | 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 |
|---|---|---|---|---|
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637399 0 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637494 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681555 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681590 2 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 984 8 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 5665572 4 | JANSSEN-CILAG | FR |
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3389530 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34009 566 554 3 4 | JANSSEN-CILAG | FR |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 5665566 3 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 983 1 5 | JANSSEN-CILAG | FR |
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3389547 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637488 1 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681549 0 | JANSSEN-CILAG | FR |
| Risperdal 0,5 mg Filmtabletten | DE/H/2184/002 | 2001040022 | JANSSEN-CILAG NV | LU |
| Risperdal 1 mg Filmtabletten | DE/H/2184/003 | 2004058265 | JANSSEN-CILAG NV | LU |
| Risperdal 2 mg Filmtabletten | DE/H/2184/004 | 2004058266 | JANSSEN-CILAG NV | LU |
| Risperdal 3 mg Filmtabletten | DE/H/2184/005 | 2004058267 | JANSSEN-CILAG NV | LU |
| Risperdal 6 mg Filmtabletten | DE/H/2184/007 | 2008089903 | JANSSEN-CILAG NV | LU |
| Risperdal 4 mg Filmtabletten | DE/H/2184/006 | 2004058268 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| Risperdal Consta 50 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/015 | 2004010028 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 2 mg Schmelztabletten | DE/H/2184/010 | 2003050032 | JANSSEN-CILAG NV | LU |
| Risperdal Consta 37,5 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/014 | 2004010027 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 0,5 mg Schmelztabletten | DE/H/2184/016 | 2003050030 | JANSSEN-CILAG NV | LU |
| Risperdal Instasolv 1 mg Schmelztabletten | DE/H/2184/009 | 2003050031 | JANSSEN-CILAG NV | LU |
| Risperdal Consta 25 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionsuspension | DE/H/2184/013 | 2004010026 | JANSSEN-CILAG NV | LU |
| RISPERDAL 2 mg filmsko obložene tablete | DE/H/2184/004 | H/96/01358/014 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 50 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/015 | H/96/01358/003 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 1 mg filmsko obložene tablete | DE/H/2184/003 | H/96/01358/008 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 25 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/013 | H/96/01358/001 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |

| 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 |
|---|---|---|---|---|
| RISPERDAL® 1 mg/ml peroralna raztopina | DE/H/2184/008 | H/96/01358/004 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL CONSTA® 37,5 mg prašek in vehikel za suspenzijo za injiciranje s podaljšanim sproščanjem | DE/H/2184/014 | H/96/01358/002 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 4 mg filmsko obložene tablete | DE/H/2184/006 | H/96/01358/024 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPERDAL 3 mg filmsko obložene tablete | DE/H/2184/005 | H/96/01358/020 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | SI |
| RISPOLEPT CONSTA 25 mg pulbere si solvent pentru suspensie injectabila cu eliberare prelungita | DE/H/2184/013 | 6876/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT 1mg/ml soluţie orală | DE/H/2184/008 | 6875/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT CONSTA 50 mg pulbere şi solvent pentru suspensie injectabilă cu eliberare prelungită | DE/H/2184/015 | 6878/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT CONSTA 37,5 mg pulbere si solvent pentru suspensie injectabila cu eliberare prelungita | DE/H/2184/014 | 6877/2014/01 | JANSSEN PHARMACEUTICA NV | RO |
| RISPOLEPT 1mg/ml soluţie orală | DE/H/2184/008 | 6875/2014/02 | JANSSEN PHARMACEUTICA NV | RO |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | PL 00242/0376 | JANSSEN-CILAG LIMITED | UK |

| 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | PL 00242/0375 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | PL 00242/0377 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL CONSTA 50 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/015 | 68/070/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 4 mg, potahované tablety | DE/H/2184/006 | 68/185/95-D/C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 1 mg potahované tablety | DE/H/2184/003 | 68/185/95-A/C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL, 1 mg/ml, perorální roztok | DE/H/2184/008 | 68/298/99-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL CONSTA 37,5 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/014 | 68/069/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 3 mg, potahované tablety | DE/H/2184/005 | 68/185/95-C/C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL CONSTA 25 mg, prášek a rozpouštědlo pro injekční suspenzi s prodlouženým uvolňováním | DE/H/2184/013 | 68/068/03-C | JANSSEN-CILAG S.R.O | CZ |
| RISPERDAL 2 mg, potahované tablety | DE/H/2184/004 | 68/185/95-B/C | JANSSEN-CILAG S.R.O | CZ |
| Risperdal Consta 25 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/013 | 17868 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| Risperdal Consta 50 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/015 | 01-9860 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 37,5 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/014 | 01-9859 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 50 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/015 | 17870 | JANSSEN-CILAG AB | SE |
| Risperdal Consta 37,5 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/014 | IS/1/02/132/02 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 37,5 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/014 | 16895 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 25 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/013 | IS/1/02/132/01 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 25 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/013 | 16894 | JANSSEN-CILAG OY | FI |
| Risperdal Consta 25 mg pulver og væske til depotinjeksjonsvæske, suspensjon | DE/H/2184/013 | 01-9858 | JANSSEN-CILAG A/S | NO |
| Risperdal Consta 50 mg stungulyfsstofn og leysir, forðadreifa | DE/H/2184/015 | IS/1/02/132/03 | JANSSEN-CILAG AB | IS |
| Risperdal Consta 50 mg injektiokuiva-aine ja liuotin, depotsuspensiota varten | DE/H/2184/015 | 16896 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| Risperdal Consta 37,5 mg pulver och vätska till injektionsvätska, depotsuspension | DE/H/2184/014 | 17869 | JANSSEN-CILAG AB | SE |
| RISPERDAL 1 mg film-coated tablets | DE/H/2184/003 | MA018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | MA018/00203 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | MA018/00203 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 1 mg film-coated tablets | DE/H/2184/003 | 018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 1mg/ml oral solution | DE/H/2184/008 | MA018/00204 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 3 mg film-coated tablets | DE/H/2184/005 | 018/00202 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 3 mg film-coated tablets | DE/H/2184/005 | MA018/00202 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL® 1 mg film-coated tablets | DE/H/2184/003 | 018/00201 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 1 mg, filmomhulde tabletten | DE/H/2184/003 | RVG 16096 | JANSSEN-CILAG BV | NL |
| RISPERDAL 4 mg, filmomhulde tabletten | DE/H/2184/006 | RVG 16099 | JANSSEN-CILAG BV | NL |
| RISPERDAL 3 mg, filmomhulde tabletten | DE/H/2184/005 | RVG 16098 | JANSSEN-CILAG BV | NL |
| RISPERDAL 1 mg/ml drank | DE/H/2184/008 | RVG 19127 | JANSSEN-CILAG BV | NL |
| RISPERDAL 0,5 mg, filmomhulde tabletten | DE/H/2184/002 | RVG 22714 | JANSSEN-CILAG BV | NL |
| RISPERDAL 2 mg, filmomhulde tabletten | DE/H/2184/004 | RVG 16097 | JANSSEN-CILAG BV | NL |
| RISPERDAL 1 mg/ml oral solution | DE/H/2184/008 | PL 00242/0199 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 0.5mg orodispersible tablets | DE/H/2184/016 | PL 00242/0378 | JANSSEN-CILAG LIMITED | UK |

| 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 |
|---|---|---|---|---|
| RISPERDAL QUICKLET 4 mg, orodispergeerbare tabletten | DE/H/2184/012 | RVG 31776 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 25 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/013 | RVG 27178 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 1 mg, orodispergeerbare tabletten | DE/H/2184/009 | RVG 27791 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 2 mg, orodispergeerbare tabletten | DE/H/2184/010 | RVG 27792 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 50 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/015 | RVG 27180 | JANSSEN-CILAG BV | NL |
| RISPERDAL QUICKLET 3 mg, orodispergeerbare tabletten | DE/H/2184/011 | RVG 31775 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 37,5 mg, poeder en oplosmiddel voor suspensie voor intramusculaire injectie met verlengde afgifte | DE/H/2184/014 | RVG 27179 | JANSSEN-CILAG BV | NL |
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | 018/00205 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | MA018/00205-7 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | MA018/00205-7 | JANSSEN-CILAG INTERNATIONAL NV | MT |
| RISPERDAL 2 mg filmomhulde tabletten | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | 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 |
|---|---|---|---|---|
| RISPERDAL 6 mg comprimés pelliculés | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | BE |
| RISPERDAL 3 mg comprimés pelliculés | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | BE |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | BE165697 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg filmomhulde tabletten | DE/H/2184/003 | BE165681 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg filmomhulde tabletten | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| RISPERDAL 4 mg comprimés pelliculés | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | BE |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | BE219983 | JANSSEN-CILAG NV | BE |
| RISPERDAL 4 mg filmomhulde tabletten | DE/H/2184/006 | BE165715 | JANSSEN-CILAG NV | BE |
| RISPERDAL 6 mg filmomhulde tabletten | DE/H/2184/007 | BE183881 | JANSSEN-CILAG NV | BE |
| RISPERDAL 3 mg filmomhulde tabletten | DE/H/2184/005 | BE165706 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 0,5 mg orodispergeerbare tabletten | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1mg/ml solution buvable | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 2 mg orodispergeerbare tabletten | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL 1 mg/ml drank | DE/H/2184/008 | BE183897 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 37,5 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/014 | BE254606 | JANSSEN-CILAG NV | 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 |
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| RISPERDAL INSTASOLV 2 mg comprimés orodispersibles | DE/H/2184/010 | BE250162 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 50 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/015 | BE254615 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg orodispergeerbare tabletten | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| RISPERDAL CONSTA 25 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/013 | BE254597 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg comprimés orodispersibles | DE/H/2184/009 | BE250153 | JANSSEN-CILAG NV | BE |
| RISPERDAL INSTASOLV 1 mg comprimés orodispersibles | DE/H/2184/009 | 2003050031 | JANSSEN-CILAG NV | LU |
| RISPERDAL 2 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/004 | 14397 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/013 | 33091 | JANSSEN-CILAG A/S | DK |
| RISPERDAL CONSTA 37,5 mg prášok a disperzné prostredie na intramuskulárnu injekčnú suspenziu s predĺženým uvoľňovaním | DE/H/2184/014 | 68/0106/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL CONSTA 25 mg prášok a disperzné prostredie na intramuskulárnu injekcnú suspenziu s predlženým uvolnovaním | DE/H/2184/013 | 68/0105/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL 3 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/005 | 14398 | JANSSEN-CILAG INTERNATIONAL NV | CY |

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| RISPERDAL 4 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/006 | 14399 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/014 | 33092 | JANSSEN-CILAG A/S | DK |
| RISPERDAL 1 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/003 | 14396 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL 1 mg/ml πόσιμο διάλυμα | DE/H/2184/008 | 17844 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| Risperdal Consta, depotinjektionsvæske, pulver og solvens til suspension | DE/H/2184/015 | 33093 | JANSSEN-CILAG A/S | DK |
| RISPERDAL INSTASOLV 0,5 mg comprimés orodispersibles | DE/H/2184/016 | 2003050030 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA50 mg prášok a disperzné prostredie na intramuskulárnu injekčnú suspenziu s predĺženým uvoľňovaním | DE/H/2184/015 | 68/0107/03-S | JOHNSON & JOHNSON, S.R.O | SK |
| RISPERDAL INSTASOLV 2 mg comprimés orodispersibles | DE/H/2184/010 | 2003050032 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA 37,5 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/014 | 2004010027 | JANSSEN-CILAG NV | LU |
| RISPERDAL 4 mg comprimés pelliculés | DE/H/2184/006 | 2004058268 | JANSSEN-CILAG NV | LU |
| RISPERDAL 1mg/ml solution buvable | DE/H/2184/008 | 2008089904 | JANSSEN-CILAG NV | LU |
| RISPERDAL 1 mg comprimés pelliculés | DE/H/2184/003 | 2004058265 | JANSSEN-CILAG NV | LU |
| RISPERDAL 3 mg comprimés pelliculés | DE/H/2184/005 | 2004058267 | JANSSEN-CILAG NV | LU |

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| RISPERDAL CONSTA 25 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/013 | 2004010026 | JANSSEN-CILAG NV | LU |
| RISPERDAL CONSTA 50 mg poudre et solvant pour suspension à libération prolongée destinée à l'injection intramusculaire | DE/H/2184/015 | 2004010028 | JANSSEN-CILAG NV | LU |
| RISPERDAL 0,5 mg comprimés pelliculés | DE/H/2184/002 | 2001040022 | JANSSEN-CILAG NV | LU |
| RISPERDAL 6 mg comprimés pelliculés | DE/H/2184/007 | 2008089903 | JANSSEN-CILAG NV | LU |
| RISPERDAL 2 mg comprimés pelliculés | DE/H/2184/004 | 2004058266 | JANSSEN-CILAG NV | LU |
| RISPERDAL 4 mg smeltetabletter | DE/H/2184/012 | 04-3141 | JANSSEN-CILAG A/S | NO |
| Risperdal 4 mg frystorkade tabletter | DE/H/2184/012 | 21886 | JANSSEN-CILAG AB | SE |
| RISPERDAL INSTASOLV 0,5 mg tabletti, suussa hajoava | DE/H/2184/016 | 17263 | JANSSEN-CILAG OY | FI |
| Risperdal 0,5 mg munndreifitöflur | DE/H/2184/016 | IS/1/03/119/01 | JANSSEN-CILAG AB | IS |
| Risperdal 3 mg frystorkade tabletter | DE/H/2184/011 | 21885 | JANSSEN-CILAG AB | SE |
| RISPERDAL QUICKLET 1 mg, Schmelztabletten | DE/H/2184/009 | 28754.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 4 mg, Schmelztabletten | DE/H/2184/012 | 28754.03.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL Lösung 1 mg/ml, Lösung zum Einnehmen | DE/H/2184/008 | 35950.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 3 mg smeltetabletter | DE/H/2184/011 | 04-3140 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg/ml oral lösning | DE/H/2184/008 | 12677 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| RISPERDAL 1 mg/ml belsőleges oldat | DE/H/2184/008 | OGYI-T-8812/07 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL 1 mg/ml oral solution | DE/H/2184/008 | PA 0748/003/001 | JANSSEN-CILAG LIMITED | IE |
| Rispolept, 2 mg, tabletki powlekane | not available | R/6705 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 1 mg munndreifitöflur | DE/H/2184/009 | IS/1/03/119/02 | JANSSEN-CILAG AB | IS |
| RISPERDAL INSTASOLV 1 mg tabletti, suussa hajoava | DE/H/2184/009 | 17264 | JANSSEN-CILAG OY | FI |
| Rispolept, 4 mg, tabletki powlekane | not available | R/6707 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT, 1 mg/ml suukaudne lahus | DE/H/2184/008 | 273799 | UAB JOHNSON & JOHNSON | EE |
| Risperdal 1 mg/ml soluzione orale | DE/H/2184/008 | 028752145 | JANSSEN-CILAG SPA | IT |
| Risperdal 2 mg munndreifitöflur | DE/H/2184/010 | IS/1/03/119/03 | JANSSEN-CILAG AB | IS |
| RISPERDAL 1 mg/ml oraaliliuos | DE/H/2184/008 | 12113 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 2 mg tabletti, suussa hajoava | DE/H/2184/010 | 17265 | JANSSEN-CILAG OY | FI |
| RISPERDAL QUICKLET 2 mg, Schmelztabletten | DE/H/2184/010 | 28754.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 0,5 mg, Schmelztabletten | DE/H/2184/016 | 28754.04.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL QUICKLET 3 mg, Schmelztabletten | DE/H/2184/011 | 28754.02.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept 1 mg/ml geriamasis tirpalas | DE/H/2184/008 | LT/1/98/0215/001 | UAB JOHNSON & JOHNSON | LT |
| Risperdal 1 mg/ml soluzione orale | DE/H/2184/008 | 028752095 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 1 mg/ml mikstur, oppløsning | DE/H/2184/008 | 95-0684 | JANSSEN-CILAG A/S | NO |
| RISPERDAL FLAS 4 mg comprimidos bucodispersables | DE/H/2184/012 | 67.051 | JANSSEN-CILAG S.A. | 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 |
|---|---|---|---|---|
| RISPERDAL FLAS 2 mg comprimidos bucodispersables | DE/H/2184/010 | 65.698 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL FLAS 3 mg comprimidos bucodispersables | DE/H/2184/011 | 67.052 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL FLAS 0,5 mg comprimidos bucodispersables | DE/H/2184/016 | 65.696 | JANSSEN-CILAG S.A. | ES |
| Rispolept 1 mg/ml šķīdums iekšķīgai lietošanai | DE/H/2184/008 | 01-0372 | UAB JOHNSON & JOHNSON | LV |
| RISPERDAL FLAS 1 mg comprimidos bucodispersables | DE/H/2184/009 | 65.697 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 1 mg/ml solución oral | DE/H/2184/008 | 62.096 | JANSSEN-CILAG S.A. | ES |
| Risperdal, smeltetabletter | DE/H/2184/009 | 33697 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/002 | 30230 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/004 | 14977 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/003 | 14976 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/006 | 14979 | JANSSEN-CILAG A/S | DK |
| Risperdal, oral opløsning | DE/H/2184/008 | 17570 | JANSSEN-CILAG A/S | DK |
| Risperdal | DE/H/2184/016 | 33696 | JANSSEN-CILAG A/S | DK |
| Risperdal, smeltetabletter | DE/H/2184/010 | 33698 | JANSSEN-CILAG A/S | DK |
| Risperdal, filmovertrukne tabletter | DE/H/2184/005 | 14978 | JANSSEN-CILAG A/S | DK |
| Risperdal 2 mg tabletti, kalvopäällysteinen | DE/H/2184/004 | 11479 | JANSSEN-CILAG 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 |
|---|---|---|---|---|
| Risperdal 0,5 mg tabletti, kalvopäällysteinen | DE/H/2184/002 | 13364 | JANSSEN-CILAG OY | FI |
| Risperdal 4 mg tabletti, kalvopäällysteinen | DE/H/2184/006 | 11481 | JANSSEN-CILAG OY | FI |
| Risperdal 1 mg tabletti, kalvopäällysteinen | DE/H/2184/003 | 11478 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 4 mg tabletti, suussa hajoava | DE/H/2184/012 | 21349 | JANSSEN-CILAG OY | FI |
| Risperdal 6 mg tabletti, kalvopäällysteinen | DE/H/2184/007 | 12302 | JANSSEN-CILAG OY | FI |
| Risperdal 3 mg tabletti, kalvopäällysteinen | DE/H/2184/005 | 11480 | JANSSEN-CILAG OY | FI |
| RISPERDAL INSTASOLV 3 mg tabletti, suussa hajoava | DE/H/2184/011 | 21348 | JANSSEN-CILAG OY | FI |
| Rispolept, 1 mg, tabletki powlekane | DE\H\2184\003\R\001 | R/6704 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT 4 mg, õhukese polümeerikattega tabletid | DE/H/2184/006 | 179797 | UAB JOHNSON & JOHNSON | EE |
| Rispolept, 1 mg/ml, roztwór doustny | DE/H/2184/008 | 4238 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 1 mg/ml mixtúra, lausn | DE/H/2184/008 | 950155 | JANSSEN-CILAG AB | IS |
| RISPERDAL CONSTA 25 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/013 | 65.213 | JANSSEN-CILAG S.A. | ES |
| RISPOLEPT 2 mg, õhukese polümeerikattega tabletid | DE/H/2184/004 | 179597 | UAB JOHNSON & JOHNSON | EE |
| RISPOLEPT 3 mg, õhukese polümeerikattega tabletid | DE/H/2184/005 | 179697 | UAB JOHNSON & JOHNSON | EE |
| Rispolept, 3 mg, tabletki powlekane | not available | R/6706 | JANSSEN-CILAG INTERNATIONAL NV | 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA 50 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/015 | 65.214 | JANSSEN-CILAG S.A. | ES |
| RISPOLEPT 1 mg, õhukese polümeerikattega tabletid | DE/H/2184/003 | 179497 | UAB JOHNSON & JOHNSON | EE |
| RISPERDAL CONSTA 37,5 mg polvo y disolvente para suspensión de liberación prolongada para inyección intramuscular | DE/H/2184/014 | 65.215 | JANSSEN-CILAG S.A. | ES |
| Risperdal 2 mg compresse rivestite con film | DE/H/2184/004 | 028752020 | JANSSEN-CILAG SPA | IT |
| Risperdal 1 mg compresse rivestite con film | DE/H/2184/003 | 028752057 | JANSSEN-CILAG SPA | IT |
| Risperdal 2 mg compresse rivestite con film | DE/H/2184/004 | 028752069 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 4 mg compresse rivestite con film | DE/H/2184/006 | AIC 028752083 | JANSSEN-CILAG SPA | IT |
| Risperdal 3 mg compresse rivestite con film | DE/H/2184/005 | 028752071 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 1 mg, Filmtabletten | DE/H/2184/003 | 28758.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 4 mg, Filmtabletten | DE/H/2184/006 | 28758.03.00 | JANSSEN-CILAG GMBH | DE |
| Risperdal 1 mg compresse rivestite con film | DE/H/2184/003 | 028752018 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 4 mg compresse rivestite con film | DE/H/2184/006 | AIC 028752044 | JANSSEN-CILAG SPA | IT |
| Rispolept 3 mg plėvele dengtos tabletės | DE/H/2184/005 | LT/1/96/0215/004 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 2 mg apvalkotās tabletes | DE/H/2184/004 | 98-0100 | UAB JOHNSON & JOHNSON | LV |
| Risperdal 3 mg compresse rivestite con film | DE/H/2184/005 | 028752032 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| Rispolept 4 mg plėvele dengtos tabletės | DE/H/2184/006 | LT/1/96/0215/005 | UAB JOHNSON & JOHNSON | LT |
| RISPERDAL 6 mg, Filmtabletten | DE/H/2184/007 | 37961.00.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept 4 mg apvalkotās tabletes | DE/H/2184/006 | 98-0102 | UAB JOHNSON & JOHNSON | LV |
| Rispolept 2 mg plėvele dengtos tabletės | DE/H/2184/004 | LT/1/96/0215/003 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 1 mg plėvele dengtos tabletės | DE/H/2184/003 | LT/1/96/0215/002 | UAB JOHNSON & JOHNSON | LT |
| Rispolept 3 mg apvalkotās tabletes | DE/H/2184/005 | 98-0101 | UAB JOHNSON & JOHNSON | LV |
| Rispolept 1 mg apvalkotās tabletes | DE/H/2184/003 | 98-0099 | UAB JOHNSON & JOHNSON | LV |
| RISPERDAL 0,5 mg, Filmtabletten | DE/H/2184/002 | 43776.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 2 mg, Filmtabletten | DE/H/2184/004 | 28758.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL 3 mg, Filmtabletten | DE/H/2184/005 | 28758.02.00 | JANSSEN-CILAG GMBH | DE |
| Risperdal 2 mg filmdragerade tabletter | DE/H/2184/004 | 11993 | JANSSEN-CILAG AB | SE |
| Risperdal 1 mg filmdragerade tabletter | DE/H/2184/003 | 11992 | JANSSEN-CILAG AB | SE |
| Risperdal 0,5 mg filmdragerade tabletter | DE/H/2184/002 | 14318 | JANSSEN-CILAG AB | SE |
| Risperdal 4 mg filmdragerade tabletter | DE/H/2184/006 | 11995 | JANSSEN-CILAG AB | SE |
| Risperdal 3 mg filmdragerade tabletter | DE/H/2184/005 | 11994 | JANSSEN-CILAG AB | SE |
| RISPERDAL 6 mg film-coated tablets | DE/H/2184/007 | PA 0748/003/002 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0,5 mg tabletter, filmdrasjerte | DE/H/2184/002 | 98-4328 | JANSSEN-CILAG A/S | NO |
| RISPERDAL 2 mg tabletter, filmdrasjerte | DE/H/2184/004/MR | 8038 | JANSSEN-CILAG A/S | 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 |
|---|---|---|---|---|
| RISPERDAL 1 mg Filmtabletten | DE/H/2184/003 | 1-20297 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 4 mg filmuhúðaðar töflur | DE/H/2184/006 | 920126 | JANSSEN-CILAG AB | IS |
| RISPERDAL 2 mg Filmtabletten | DE/H/2184/004 | 1-20301 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 2 mg filmuhúðaðar töflur | DE/H/2184/004 | 920124 | JANSSEN-CILAG AB | IS |
| RISPERDAL 3 mg tabletter, filmdrasjerte | DE/H/2184/005 | 8039 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg filmuhúðaðar töflur | DE/H/2184/003 | 920123 | JANSSEN-CILAG AB | IS |
| RISPERDAL 3 mg Filmtabletten | DE/H/2184/005 | 1-20302 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 3 mg filmuhúðaðar töflur | DE/H/2184/005 | 920125 | JANSSEN-CILAG AB | IS |
| RISPERDAL 1 mg/ml Lösung zum Einnehmen | DE/H/2184/008 | 1-21466 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 0,5 mg filmuhúðaðar töflur | DE/H/2184/002 | 980135 | JANSSEN-CILAG AB | IS |
| RISPERDAL 4 mg Filmtabletten | DE/H/2184/006 | 1-20303 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 6 mg Filmtabletten | DE/H/2184/007 | 1-22846 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 1 mg tabletter, filmdrasjerte | DE/H/2184/003 | 8037 | JANSSEN-CILAG A/S | NO |
| RISPERDAL 4 mg tabletter, filmdrasjerte | DE/H/2184/006 | 8040 | JANSSEN-CILAG A/S | NO |
| RISPERDALCONSTA L.P. 50 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/015 | 34009 362 494 2 4 | JANSSEN-CILAG | 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 |
|---|---|---|---|---|
| RISPERDALCONSTA L.P. 37,5 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/014 | 34 00 9 3624936 3 | JANSSEN-CILAG | FR |
| RISPERDALCONSTA L.P. 25 mg/2 ml, poudre et solvant pour suspension injectable à libération prolongée en seringue préremplie | DE/H/2184/013 | 34 00 9 3624913 4 | JANSSEN-CILAG | FR |
| RISPERDAL CONSTA 50 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/015 | 1-24630 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL CONSTA 25 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/013 | 1-24628 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL CONSTA 37,5 mg Pulver und Lösungsmittel zur Herstellung einer verzögert freisetzenden Suspension zur intramuskulären Injektion | DE/H/2184/014 | 1-24629 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 3 mg comprimidos recubiertos con película | DE/H/2184/005 | 60.335 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 1 mg comprimidos recubiertos con película | DE/H/2184/003 | 60.336 | JANSSEN-CILAG S.A. | ES |
| RISPERDAL 6 mg comprimidos recubiertos con película | DE/H/2184/007 | 62.803 | JANSSEN-CILAG S.A. | 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA50 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/015 | 19589 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL CONSTA 25 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/013 | 52995.00.00 | JANSSEN-CILAG GMBH | DE |
| RISPERDAL CONSTA 50 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/015 | 52995.02.00 | JANSSEN-CILAG GMBH | DE |
| Rispolept Consta, 50 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu | DE/H/2184/015 | 10580 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT CONSTA 37,5 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/014 | 03-0097 | UAB JOHNSON & JOHNSON | LV |
| RISPOLEPT CONSTA 37,5 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/014 | LT/1/03/3651/002 | UAB JOHNSON & JOHNSON | LT |
| Rispolept Consta 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | 20030321 | JOHNSON & JOHNSON PRODAJA MEDICINSKIH IN FARMACEVTSKIH IZDELKOV, D.O.O. | BG |
| RISPOLEPT CONSTA 50 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/015 | LT/1/03/3651/003 | UAB JOHNSON & JOHNSON | 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 |
|---|---|---|---|---|
| Rispolept Consta, 25 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu. | DE/H/2184/013 | 10582 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| Risperdal 4 mg munndreifitöflur | DE/H/2184/012 | IS/1/06/139/02 | JANSSEN-CILAG AB | IS |
| Rispolept Consta, 37,5 mg, proszek i rozpuszczalnik do sporządzania zawiesiny do wstrzykiwań domięśniowych o przedłużonym uwalnianiu. | DE/H/2184/014 | 10581 | JANSSEN-CILAG INTERNATIONAL NV | PL |
| RISPOLEPT CONSTA 25 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/013 | 03-0096 | UAB JOHNSON & JOHNSON | LV |
| RISPOLEPT CONSTA 25 mg milteliai ir tirpiklis pailginto atpalaidavimo injekcinei suspensijai | DE/H/2184/013 | LT/1/03/3651/001 | UAB JOHNSON & JOHNSON | LT |
| RISPOLEPT CONSTA 25 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/013 | 410603 | UAB JOHNSON & JOHNSON | EE |
| RISPOLEPT CONSTA 50 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/015 | 410803 | UAB JOHNSON & JOHNSON | EE |
| RISPERDAL CONSTA 37,5 mg Pulver und Lösungsmittel zur Herstellung einer intramuskulären Depot-Injektionssuspension | DE/H/2184/014 | 52995.01.00 | JANSSEN-CILAG GMBH | DE |
| RISPOLEPT CONSTA 37,5 mg, toimeainet prolongeeritult vabastava süstesuspensiooni pulber ja lahusti | DE/H/2184/014 | 410703 | UAB JOHNSON & JOHNSON | EE |

| 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 |
|---|---|---|---|---|
| RISPOLEPT CONSTA 50 mg pulveris un šķīdinātājs ilgstošas darbības injekciju suspensijas pagatavošanai | DE/H/2184/015 | 03-0098 | UAB JOHNSON & JOHNSON | LV |
| Risperdal 25 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/013 | 028752172 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 50 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/015 | AIC N° 028752196 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 37,5 mg/2 ml polvere e solvente per sospensione iniettabile a rilascio prolungato per uso intramuscolare | DE/H/2184/014 | AIC N° 028752184 | JANSSEN-CILAG SPA | IT |
| RISPERDAL CONSTA 25 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/013 | 67148/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperdal Consta 37,5 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/014 | OGYI-T-8812/04 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL CONSTA 50 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/015 | 67150/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | 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 |
|---|---|---|---|---|
| Risperdal Consta 50 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/015 | OGYI-T-8812/06 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL Quicklet 0,5 mg Schmelztabletten | DE/H/2184/016 | 1-25155 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL Quicklet 4 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/012 | 67146/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL CONSTA 37,5 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/014 | 67149/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperdal Consta 25 mg por és oldószer retard szuszpenziós injekcióhoz intramuszkuláris célra | DE/H/2184/013 | OGYI-T-8812/02 | JANSSEN-CILAG KFT. | HU |
| RISPERDAL 1 mg, comprimé pelliculé | DE/H/2184/003 | 34 00 9 3389487 0 | JANSSEN-CILAG | FR |
| RISPERDAL 4 mg, comprimé pelliculé | DE/H/2184/006 | 34 00 9 3442738 1 | JANSSEN-CILAG | FR |
| RISPERDAL 2 mg, comprimé pelliculé | DE/H/2184/004 | 34 00 9 3389501 3 | JANSSEN-CILAG | FR |
| RISPERDAL 1 mg/ml, solution buvable | DE/H/2184/008 | 34009 343 981 9 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 2 mg, comprimé orodispersible | DE/H/2184/010 | 34 00 9 3637471 3 | JANSSEN-CILAG | FR |
| RISPERDALORO 3 mg, comprimé orodispersible | DE/H/2184/011 | 34 00 9 3681532 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 1 mg, comprimé orodispersible | DE/H/2184/009 | 34 00 9 3637436 2 | JANSSEN-CILAG | FR |
| RISPERDALORO 0,5 mg, comprimé orodispersible | DE/H/2184/016 | 34 00 9 3637382 2 | JANSSEN-CILAG | 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA 50 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/015 | PA 0748/003/012 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL CONSTA 37.5 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/014 | PA 0748/003/011 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 4219887 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 4219689 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 4 mg comprimido orodispersível | DE/H/2184/012 | 5827589 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 4219788 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1mg comprimidos revestidos por película | DE/H/2184/003 | 2305985 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 2 mg comprimidos revestidos por película | DE/H/2184/004 | 2306082 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg/ml solução oral | DE/H/2184/008 | 2527984 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 4219986 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220182 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220083 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 3 mg comprimido orodispersível | DE/H/2184/011 | 5827381 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1mg comprimidos revestidos por película | DE/H/2184/003 | 2305787 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 4 mg comprimido orodispersível | DE/H/2184/012 | 5827480 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 3 mg comprimido orodispersível | DE/H/2184/011 | 5827282 | JANSSEN FARMACÊUTICA PORTUGAL, 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 |
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| RISPERDAL CONSTA 50 mg pó e veículo para suspensão de libertação prolongada para injecção intramuscular | DE/H/2184/015 | 4753686 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDALORO 4 mg, comprimé orodispersible | DE/H/2184/012 | 34 00 9 3681578 0 | JANSSEN-CILAG | FR |
| RISPERDAL QUICKLET 1 mg comprimido orodispersível | DE/H/2184/009 | 5701289 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 25 mg pó e veículo para suspensão de libertação prolongada para injeção intramuscular | DE/H/2184/013 | 4753588 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL Quicklet 3 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/011 | 67145/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 1 mg/ml solução oral | DE/H/2184/008 | 2715381 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL QUICKLET 0,5 mg comprimido orodispersível | DE/H/2184/016 | 5701180 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL Quicklet 2 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/010 | 67144/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL QUICKLET 2 mg comprimido orodispersível | DE/H/2184/010 | 4220083 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 37,5 mg pó e veículo para suspensão de libertação prolongada para injeção intramuscular | DE/H/2184/014 | 4753687 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA37,5 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/014 | 19588 | JANSSEN-CILAG INTERNATIONAL NV | CY |

| 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 |
|---|---|---|---|---|
| RISPERDAL CONSTA 25 mg κόνις και διαλύτης για παρασκευή ενέσιμου εναιωρήματος παρατεταμένης αποδέσμευσης για ενδομυϊκή χρήση | DE/H/2184/013 | 19587 | JANSSEN-CILAG INTERNATIONAL NV | CY |
| RISPERDAL Quicklet 1 mg Schmelztabletten | DE/H/2184/009 | 1-25156 | JANSSEN-CILAG PHARMA GMBH | AT |
| Risperdal 3 mg munndreifitöflur | DE/H/2184/011 | IS/1/06/139/01 | JANSSEN-CILAG AB | IS |
| RISPERDAL Quicklet 2 mg Schmelztabletten | DE/H/2184/010 | 1-25157 | JANSSEN-CILAG PHARMA GMBH | AT |
| RISPERDAL 2 mg comprimidos revestidos por película | DE/H/2184/004 | 2306181 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg comprimidos revestidos por película | DE/H/2184/003 | 2305886 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 2 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/004 | 67138/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 4 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/006 | 67140/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 4 mg comprimidos revestidos por película | DE/H/2184/006 | 2306587 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 1 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/003 | 67137/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 4 mg comprimidos revestidos por película | DE/H/2184/006 | 2306488 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 3 mg comprimidos revestidos por película | DE/H/2184/005 | 2306389 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |

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| RISPERDAL 1 mg/ml πόσιμο διάλυμα | DE/H/2184/008 | 67147/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 3 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/005 | 67139/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 6 mg επικαλυμμένα με λεπτό υμένιο δισκία | DE/H/2184/007 | 67140/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL 0,5 mg comprimidos revestidos por película | DE/H/2184/002 | 3219888 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL 3 mg comprimidos revestidos por película | DE/H/2184/005 | 2306280 | JANSSEN FARMACÊUTICA PORTUGAL, LDA | PT |
| RISPERDAL CONSTA 25 mg powder and solvent for prolonged-release suspension for intramuscular injection | DE/H/2184/013 | PA 0748/003/010 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 3 mg film-coated tablets | DE/H/2184/005 | PA 0748/003/006 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 2 mg smeltetabletter | DE/H/2184/010 | 02-1026 | JANSSEN-CILAG A/S | NO |
| Risperdal 1 mg compresse orodispersibili | DE/H/2184/009 | 028752222 | JANSSEN-CILAG SPA | IT |
| Risperdal 1 mg frystorkade tabletter | DE/H/2184/009 | 18385 | JANSSEN-CILAG AB | SE |
| RISPERDAL 1mg film-coated tablets | DE/H/2184/003 | PL 00242/0186 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 4 mg film-coated tablets | DE/H/2184/006 | PA 0748/003/007 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL Quicklet 1 mg orodispersible tablets | DE/H/2184/009 | PA 0748/003/014 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0.5 mg film-coated tablets | DE/H/2184/002 | PA 0748/003/009 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 2 mg film-coated tablets | DE/H/2184/004 | PA 0748/003/005 | JANSSEN-CILAG LIMITED | 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 |
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| RISPERDAL Quicklet 3 mg orodispersible tablets | DE/H/2184/011 | PA 0748/003/016 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 3mg film-coated tablets | DE/H/2184/005 | PL 00242/0188 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 1 mg smeltetabletter | DE/H/2184/009 | 02-1025 | JANSSEN-CILAG A/S | NO |
| RISPERDAL Quicklet 4 mg orodispersible tablets | DE/H/2184/012 | PL 0242/0408 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 1 mg orodispersible tablets | DE/H/2184/009 | PL 0242/0379 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 2mg orodispersible tablets | DE/H/2184/010 | PL 0242/0380 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 2mg film-coated tablets | DE/H/2184/004 | PL 00242/0187 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 6mg film-coated tablets | DE/H/2184/007 | PL 00242/0317 | JANSSEN-CILAG LIMITED | UK |
| Risperdal 0,5 mg frystorkade tabletter | DE/H/2184/016 | 18384 | JANSSEN-CILAG AB | SE |
| RISPERDAL 2 mg compresse orodispersibili | DE/H/2184/010 | 028752259 | JANSSEN-CILAG SPA | IT |
| RISPERDAL Quicklet 4 mg orodispersible tablets | DE/H/2184/012 | PA 0748/003/017 | JANSSEN-CILAG LIMITED | IE |
| Risperdal 2 mg frystorkade tabletter | DE/H/2184/010 | 18386 | JANSSEN-CILAG AB | SE |
| RISPERDAL 0.5mg film-coated tablets | DE/H/2184/002 | PL 00242/0347 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 0.5mg orodispersible tablets | DE/H/2184/016 | PA0748/003/013 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 0,5 mg smeltetabletter | DE/H/2184/016 | 02-1024 | JANSSEN-CILAG A/S | NO |
| RISPERDAL Quicklet 3 mg orodispersible tablets | DE/H/2184/011 | PL 0242/0407 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL 1 mg compresse orodispersibili | DE/H/2184/009 | 028752234 | JANSSEN-CILAG SPA | IT |
| RISPERDAL 2 mg compresse orodispersibili | DE/H/2184/010 | 028752246 | JANSSEN-CILAG SPA | 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 |
|---|---|---|---|---|
| RISPERDAL 4mg film-coated tablets | DE/H/2184/006 | PL 00242/0189 | JANSSEN-CILAG LIMITED | UK |
| RISPERDAL Quicklet 2 mg orodispersible tablets | DE/H/2184/010 | PA 0748/003/015 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL 1 mg film-coated tablets | DE/H/2184/003 | PA 0748/003/04 | JANSSEN-CILAG LIMITED | IE |
| RISPERDAL INSTASOLV 0,5 mg comprimés orodispersibles | DE/H/2184/016 | BE250144 | JANSSEN-CILAG NV | BE |
| RISPERDAL Quicklet 0,5 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/016 | 67142/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| RISPERDAL Quicklet 1 mg δισκία διασπειρόμενα στο στόμα | DE/H/2184/009 | 67143/12.09.2016 | JANSSEN-CILAG PHARMACEUTICAL S.A.C.I. | GR |
| Risperidon Aurobindo 6 mg Filmtabletten | NL/H/1957/006 | 80755.00.00 | AUROBINDO PHARMA GMBH | DE |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078014 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078026 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078038 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078040 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078053 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078065 | AUROBINDO PHARMA (ITALIA) S.R.L. | 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 |
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| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078077 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078089 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078091 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078103 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078584 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 0,5 mg compresse rivestite con film | NL/H/1957/001 | 040078596 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078519 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078521 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078533 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078545 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078558 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078560 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidone Aurobindo 6 mg compresse rivestite con film | NL/H/1957/006 | 040078572 | AUROBINDO PHARMA (ITALIA) S.R.L. | IT |
| Risperidon Aurobindo 0,5 mg, filmomhulde tabletten | NL/H/1957/001 | RVG 33495 | AUROBINDO PHARMA B.V. | NL |
| Risperidona Farmalider 2 mg comprimidos recubiertos con película | not available | 66805 | FARMALIDER, S.A. | ES |