

Patient information leaflet

Scheduling status
Schedule 0

Proprietary name, strength and pharmaceutical form

Reneel

Read all of this leaflet carefully because it contains important information for you.

Reneel is available without a doctor's prescription, for you to treat a mild illness. Nevertheless you still need to use Reneel carefully to get the best results from it.

- Keep this leaflet. You may need to read it again.
- Do not share Reneel with any other person.
- Ask your pharmacist if you need more information or advice.
- You must see a doctor if your symptoms worsen or do not improve.

1. What Reneel contains

1 tablet cont.:

The active substances are: Berberis vulgaris D2 15 mg, Acidum nitricum D4 30 mg, Cantharis D5 30 mg, Plumbum acetikum D6 30 mg, Pareira brava D3 30 mg, Sabal serrulatum D2 30 mg, Causticum Hahnemanni D4 60 mg, Aluminium oxydatum D12 75 mg.

The other ingredient is: Magnesium stearate.

1 tablet contains approx. 300 mg lactose.

2. What Reneel is used for

Pharmacological classification:

D. 33.2. Homeopathy.

Discipline of the medicine: Homeopathy

This medicine is prepared in accordance with homeopathic principles and is proposed for use in inflammatory diseases in the region of the urinary passages, with and without lithiasis.

3. Before you take Reneel

Do not take Reneel:

- if you are hypersensitive (allergic) to active substances or any of the other ingredients of Reneel.

Pregnancy and breastfeeding:

If you are pregnant or breastfeeding your baby while taking this medicine, please consult your doctor, pharmacist or other healthcare professional for advice.

Important information about some of the ingredients of Reneel:

This preparation contains natural lactose. Although the quantity of lactose present is probably not sufficient to cause discomfort, a health professional should be consulted in strong cases of lactose intolerance. A temporary aggravation of the existing symptoms is possible after taking a homeopathic preparation.

Taking other medicines with Reneel:

If you are taking other medicines on a regular basis, including complementary or traditional medicines, the use of Reneel with these medicines may cause undesirable interactions.

Please consult your doctor, pharmacist or other healthcare professional for advice.

4. How to take Reneel

Do not share medicines prescribed for you with any other person.

Always take Reneel exactly as your doctor has instructed you. You should check with your doctor or pharmacist if you are unsure.

The usual dose is:

Adults and children 12 years and older:

1 tablet 3x daily.

Children 6-11 years: 1 tablet 2x daily.

Children 2-5 years: 1 tablet 1-2x daily.

Infants up to 2 years: 1 tablet 1x daily.

Acute or initial dose:

Adults and children 12 years and older:

1 tablet every ½ to 1 hour, up to 12x daily, and then continue with usual dosage.

Children 6-11 years:

1 tablet every 1 to 2 hours, up to 8x daily, and then continue with usual dosage.

Children 2-5 years:

1 tablet every 1 to 2 hours, up to 6x daily, and then continue with usual dosage.

Infants up to 2 years:

1 tablet every 1 to 2 hours, up to 4x daily, and then continue with usual dosage.

Method of administration:

The tablets should be allowed to dissolve slowly in the mouth. For children and infants, it is recommended to crush the tablet and administer it dissolved in a little water.

It is recommended not to eat or drink 15 minutes before or after taking the medication.

If you take more Reneel than you should:

In the event of overdose, consult your doctor or pharmacist. If neither is available, seek help at the nearest hospital or poison control centre.

If you forget to take Reneel:

Do not take a double dose to make up for forgotten individual doses.

5. Possible side effects

Reneel can have side effects.

None known.

Not all side effects reported for Reneel are included in this leaflet. Should your general health worsen while taking this medicine, please consult your doctor, pharmacist or other healthcare professional for advice.

If you notice any side effects not mentioned in this leaflet, please inform your doctor or pharmacist.

6. Storing and disposing of Reneel

Keep all medicines out of the reach and sight of children.

- Store in a cool (below 25 °C) dry place.

7. Presentation of Reneel

Containers of 50 tablets.

8. Identification of Reneel

White to light yellow tablets.

9. Registration number / Reference number

U 5508 (Act 101/1965)

10. Name and business address of the holder of the certificate of registration

ModHomCo (Pty) Ltd
96 Amsterdam Street
Clubview, 0157 Centurion

11. Date of publication

August 2024

This unregistered medicine has not been evaluated by the SAHPRA for its quality, safety or intended use.



Pasiënt-inligtingstuk

052742/5007/ZA

Skeduleringstatus
Skedule 0

Handelsnaam, sterkte en farmaseutiese vorm

Tablets / Tablette

Lees die hele inhoud van hierdie inligtingstuk noukeurig aangesien dit belangrike inligting bevat.

Reneel is sonder 'n doktersvoorskrif beskikbaar om 'n ligte siekte mee te behandel. Nogtans moet u Reneel versigtig gebruik om die beste resultate te verkry.

- Hou hierdie inligtingstuk. U mag dit weer moet lees.
- Moenie Reneel met enige ander persoon deel nie.
- Vra u apteker indien u meer inligting of advies benodig.
- U moet 'n geneesheer raadpleeg indien u simptome vererger of nie verbeter nie.

1. Wat Reneel bevat

1 tablet bevat:

Die aktiewe bestanddele is: Berberis vulgaris D2 15 mg, Acidum nitricum D4 30 mg, Cantharis D5 30 mg, Plumbum acetikum D6 30 mg, Pareira brava D3 30 mg, Sabal serrulatum D2 30 mg, Causticum Hahnemanni D4 60 mg, Aluminium oxydatum D12 75 mg.

Die ander bestanddeel is: Magnesiumstearaat.

1 tablet bevat ong. 300 mg laktose.

2. Waarvoor Reneel gebruik word

Farmakologiese klassifikasie:

D. 33.2. Homeopatie.

Dissipline van die medisyne: Homeopatie

Hierdie medisyne is in ooreenstemming met homeopatiëse beginsels voorberei en word voorgestel vir gebruik in inflammatoriese siektes van die urieneweg, met of sonder steenvorming (litiase).

3. Voordat u Reneel neem

Moenie Reneel neem:

- indien u hipersensitief (allergies) is vir die aktiewe bestanddele of enige van die ander bestanddele van Reneel.

Swangerskap en borsvoeding:

Indien u swanger is of u baba borsvoed terwyl u hierdie medisyne neem, raadpleeg asseblief u geneesheer, apteker of ander professionele gesondheidswerker vir advies.

Belangrike inligting omtrent sommige van die bestanddele van Reneel:

Hierdie preparaat bevat natuurlike laktose. Hoewel die hoeveelheid laktose aanwesig waarskynlik nie genoegsaam is om ongemak te veroorsaak nie, moet 'n professionele gesondheidswerker geraadpleeg word in sterk gevalle van laktose-intoleransie. 'n Tydelike verergering van die bestaande simptome is moontlik na die inname van 'n homeopatiëse preparaat.

Die neem van ander medisyne saam met Reneel:

Indien u op 'n gereelde basis ander medisyne neem, insluitend komplementêre of tradisionele medisyne, mag die gebruik van Reneel saam met hierdie medisyne ongewenste interaksies tot gevolg hê. Raadpleeg asseblief u geneesheer, apteker of ander professionele gesondheidswerker vir advies.

4. Hoe om Reneel te neem

Moenie medisyne wat vir u voorgeskryf is met enige ander persoon deel nie.

Neem Reneel altyd presies soos wat dit vir u voorgeskryf is. U moet u geneesheer of apteker raadpleeg indien u onseker is.

Die gewone dosis is:

Volwassenes en kinders 12 jaar en ouer:

1 tablet 3x daaglik.

Kinders 6-11 jaar: 1 tablet 2x daaglik.

Kinders 2-5 jaar: 1 tablet 1-2x daaglik.

Babas tot 2 jaar oud: 1 tablet 1x daaglik.

Akute of aanvanklike dosis:

Volwassenes en kinders 12 jaar en ouer:

1 tablet elke ½ tot 1 uur, tot 12x daaglik, en gaan dan voort met die gewone dosis.

Kinders 6-11 jaar:

1 tablet elke 1 tot 2 uur, tot 8x daaglik, en gaan dan voort met die gewone dosis.

Kinders 2-5 jaar:

1 tablet elke 1 tot 2 uur, tot 6x daaglik, en gaan dan voort met die gewone dosis.

Babas tot 2 jaar oud:

1 tablet elke 1 tot 2 uur, tot 4x daaglik, en gaan dan voort met die gewone dosis.

Metode van administrasie:

Die tablette moet stadig in die mond opgelos word. Vir kinders en babas word dit aanbeveel om die tablet fyn te maak en in 'n bietjie water opgelos, toe te dien.

Dit word aanbeveel dat daar vir 15 minute voordat en nadat die medikasie geneem word, nie geëet of gedrink word nie.

Indien u meer Reneel neem as wat u moet:

In die geval van 'n oordosering, raadpleeg u geneesheer of apteker. Indien nie een van hulle beskikbaar is nie, soek hulp by die naaste hospitaal of gifbeheersentrum.

Indien u vergeet om Reneel te neem:

Moenie 'n dubbeldosis neem om op te maak vir vergeete individuele dosisse nie.

5. Moontlike nuwe-effekte

Reneel kan nuwe-effekte hê.

Geen bekend nie.

Nie alle nuwe-effekte wat vir Reneel gerapporteer is word in hierdie inligtingstuk ingesluit nie. Indien u algemene gesondheid vererger terwyl u hierdie medisyne neem, raadpleeg asseblief u geneesheer, apteker of ander professionele gesondheidswerker vir advies.

Indien u enige nuwe-effekte opmerk wat nie in hierdie inligtingstuk genoem word nie, stel asseblief u geneesheer of apteker in kennis.

6. Opberging van en beskikking oor Reneel

Bêre alle medisyne buite die bereik en sig van kinders.

- Bêre in 'n koel (onder 25 °C) droë plek.

7. Aanbieding van Reneel

Houers van 50 tablette.

8. Identifikasie van Reneel

Wit tot liggeel tablette.

9. Registrasienommer / Verwysingsnommer

U 5508 (Wet 101/1965)

10. Naam en besigheidsadres van die houer van die registrasiesertifikaat

ModHomCo (Edms) Bpk
Amsterdamstraat 96
Clubview, 0157 Centurion

11. Datum van publikasie

Augustus 2024

Hierdie ongeregistreerde medisyne is nie deur die SAHPRA vir kwaliteit, veiligheid of beoogde gebruik geëvalueer nie.

-Heel



Reneel

1. Scheduling status

Schedule 0

2. Proprietary name and dosage form

Reneel Tablets

3. Composition

1 tablet cont.: Berberis vulgaris D2 15 mg, Acidum nitricum D4 30 mg, Cantharis D5 30 mg, Plumbum aceticum D6 30 mg, Pareira brava D3 30 mg, Sabal serrulatum D2 30 mg, Causticum Hahnemanni D4 60 mg, Aluminium oxydatum D12 75 mg. Excipient: Magnesium stearate. 1 tablet contains approx. 300 mg lactose.

4. Pharmacological classification

D. 33.2. Homeopathy.

5. Pharmacological action

Action based on homeopathic principles.

6. Indications

This medicine is prepared in accordance with homeopathic principles and is proposed for use in inflammatory diseases in the region of the urinary passages, with and without lithiasis.

7. Contraindications

Hypersensitivity to any of the ingredients, including excipients.

8. Warnings

This preparation contains natural lactose. Although the quantity of lactose present is probably not sufficient to cause discomfort, a health professional should be consulted in strong cases of lactose intolerance. A temporary aggravation of the existing symptoms is possible after taking a homeopathic preparation.

9. Interactions

No interactions studies have been performed.

10. Pregnancy and lactation

Safety and/or efficacy has not been established.

11. Dosage and directions for use

The usual dose is:
Adults and children 12 years and older: 1 tablet 3x daily.
Children 6-11 years: 1 tablet 2x daily.
Children 2-5 years: 1 tablet 1-2x daily.
Infants up to 2 years: 1 tablet 1x daily.

Acute or initial dose:
Adults and children 12 years and older: 1 tablet every ½ to 1 hour, up to 12x daily, and then continue with usual dosage.
Children 6-11 years: 1 tablet every 1 to 2 hours, up to 8x daily, and then continue with usual dosage.
Children 2-5 years: 1 tablet every 1 to 2 hours, up to 6x daily, and then continue with usual dosage.
Infants up to 2 years: 1 tablet every 1 to 2 hours, up to 4x daily, and then continue with usual dosage.

Method of administration:
The tablets should be allowed to dissolve slowly in the mouth. For children and infants, it is recommended to crush the tablet and administer it dissolved in a little water.

It is recommended not to eat or drink 15 minutes before or after taking the medication.

12. Side effects and special precautions

12.1 Side effects

None known.

12.2 Special precautions

None known.

12.3 Effects on ability to drive and use machines

13. Known symptoms of overdose and particulars of its treatment

None known.

14. Identification

White to light yellow tablets.

15. Presentation

Containers of 50 tablets.

16. Storage instructions

Store in a cool (below 25 °C) dry place beyond the reach of children.

17. Registration number

U 5508 (Act 101/1965)

18. Name and business address of the holder of the certificate of registration

ModHomCo (Pty) Ltd
96 Amsterdam Street
Clubview, 0157 Centurion
Manufactured in Germany.

19. Date of publication of the professional information

August 2024

This unregistered medicine has not been evaluated by the SAHPRA for its quality, safety or intended use.

Tablets / Tablette

1. Skeduleringstatus

Skedule 0

2. Handelsnaam en doseervorm

Reneel Tablette

3. Samestelling

1 tablet bevat: Berberis vulgaris D2 15 mg, Acidum nitricum D4 30 mg, Cantharis D5 30 mg, Plumbum aceticum D6 30 mg, Pareira brava D3 30 mg, Sabal serrulatum D2 30 mg, Causticum Hahnemanni D4 60 mg, Aluminium oxydatum D12 75 mg. Bymiddel: Magnesiumstearaat. 1 tablet bevat ong. 300 mg laktose.

4. Farmakologiese klassifikasie

D. 33.2. Homeopatie.

5. Farmakologiese werking

Werking gebaseer op homeopatiese beginsels.

6. Indikasies

Hierdie medisyne is volgens homeopatiese beginsels voorberei en word voorgestel in gebruik in inflammatoriese siektes in die omgewing van die urinerweg, met of sonder steenvorming (litiase).

7. Kontra-indikasies

Hipersensitiwiteit vir enige van die bestanddele, insluitend die bymiddels.

8. Waarskuwings

Hierdie preparaat bevat natuurlike laktose. Hoewel die hoeveelheid laktose aanwesig waarskynlik nie genoegsaam is om ongemak te veroorsaak nie, moet 'n professionele gesondheidswerker geraadpleeg word in sterk gevalle van laktose-intoleransie. 'n Tydelike verergering van die bestaande simptome is moontlik na die inname van 'n homeopatiese preparaat.

9. Interaksies

Geen interaksie-studies is uitgevoer nie.

10. Swangerskap en borsvoeding

Veiligheid en/of doeltreffendheid is nie vasgestel nie.

11. Dosering en gebruiksaanwysings

Die gewone dosis is:
Volwassenes en kinders 12 jaar en ouer: 1 tablet 3x daaglik.
Kinders 6-11 jaar: 1 tablet 2x daaglik.
Kinders 2-5 jaar: 1 tablet 1-2x daaglik.
Babas tot 2 jaar oud: 1 tablet 1x daaglik.

Akute of aanvanklike dosis:
Volwassenes en kinders 12 jaar en ouer: 1 tablet elke ½ tot 1 uur, tot 12x daaglik, en gaan dan voort met die gewone dosis.
Kinders 6-11 jaar: 1 tablet elke 1 tot 2 uur, tot 8x daaglik, en gaan dan voort met die gewone dosis.
Kinders 2-5 jaar: 1 tablet elke 1 tot 2 uur, tot 6x daaglik, en gaan dan voort met die gewone dosis.
Babas tot 2 jaar oud: 1 tablet elke 1 tot 2 uur, tot 4x daaglik, en gaan dan voort met die gewone dosis.

Metode van administrasie:
Die tablette moet stadig in die mond opgelos word. Vir kinders en babas word dit aanbeveel om die tablet fyn te maak en in 'n bietjie water opgelos, toe te dien.

Dit word aanbeveel dat daar vir 15 minute voordat en nadat die medikasie geneem word, nie geëet of gedrink word nie.

12. Nuwe-effekte en spesiale voorsorgmaatreëls

12.1 Nuwe-effekte

Geen bekend nie.

12.2 Spesiale voorsorgmaatreëls

Geen bekend nie.

12.3 Effekte op die vermoë om motorvoertuie te bestuur en masjinerie te gebruik

13. Bekende simptome van oordosering en besonderhede van die behandeling daarvan

Geen bekend nie.

14. Identifikasie

Wit tot liggeel tablette.

15. Aanbieding

Houers van 50 tablette.

16. Opbergingsinstruksies

Bêre in 'n koel (onder 25 °C) droë plek buite die bereik van kinders.

17. Registrasienuommer

U 5508 (Wet 101/1965)

18. Naam en besigheids-adres van die houër van die registrasie-sertifikaat

ModHomCo (Edms) Bpk
Amsterdamstraat 96
Clubview, 0157 Centurion
Vervaardig in Duitsland.

19. Datum van publikasie van hierdie professionele inligting

Augustus 2024

Hierdie ongeregisteerde medisyne is nie deur die SAHPRA vir kwaliteit, veiligheid of beoogde gebruik geëvalueer nie.