VADEMÉCUM
SAVAL

Contenido exclusivo para Profesionales de la Salud habilitados para prescribir y dispensar medicamentos.

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SAVAL

Siempre junto a ti

Vademecum SAVAL

Indice de productos

AB

Bucco-pharyngeal antiseptic

Composition:

Each tablet contains:
Chlorexidine hydrochloride 5 mg
Excipients q.s.

Presentations:

12 tablet pack

Indications:

Mild mouth and throat infections, stomatitis, glossitis, gingivitis and pharyngitis. Dental inflammatory processes; pre and post dental surgery. Antiseptic for mouth and throat, oropharynge, upper buccal vestibule, tonsils, buccal mucous membrane, sublingual region, palate

AB ANTITUSIVO

Bucco-pharyngeal antiseptic / Antitussive

Composition:

Each tablet contains:
Chlorhexidne hydrochloride 5 mg
Noscapine 10 mg
Excipients q.s.

Presentations:

12 tablet pack

Indications:

Cough associated with: mild mouth and throat infections, stomatitis, glossitis, gingivitis and pharyngitis. Cough symptomatic treatment, in case of laryngitis, pharyngitis and tracheitis. Mouth and throat antiseptic and antitussive drug for relieving cough which is generally associated with laryngitis, pharyngitis and tracheitis.

ACTAN

Antidepressant

Composition:

Each ACTAN capsule contains:
Fluoxetine hydrochloride 20 mg
Excipients q.s.

Each ACTAN CD dispersable tablet contains:
Fluoxetine hydrochloride 20 mg
Excipients q.s.

Presentations:

30 or 60 capsule packs
30 dispersible scored tablet packs

Indications:

Fluoxetine hydrochloride is indicated for the acute and maintenance treatment of depression, associated or not with anxiety; for the treatment of obsessive-compulsive disorder in adults and children 8 years and above; for the treatment of premenstrual dysphoric disorder.

It is indicated as an adjuvant to psychotherapy for the reduction of binge-eating and purging activity in patients with bulimia nervosa.

ACTAN CD

Antidepressant

Composition:

Cada comprimido dispersable contiene:
Fluoxetina clorhidrato 20 mg.
Excipientes c.s.

Presentations:

Actan CD: Estuche con 30 comprimidos dispersables

Indications:

Fluoxetina clorhidrato está indicado para el tratamiento agudo y de mantención de la depresión asociada o no con ansiedad; para el tratamiento del trastorno obsesivo compulsivo en adultos y niños a partir de 8 años de edad; para el tratamiento del desorden disfórico premenstrual. Indicado como complemento a la psicoterapia para la reducción de la ingesta excesiva de alimentos, la provocación del vómito y el uso de laxantes en pacientes con bulimia nerviosa.

ADAX

Antianxiety

Composition:

Each ADAX tablet contains:
Alprazolam 0,25 mg
Excipients q.s.

Alprazolam 0,50 mg
Excipients q.s.

Alprazolam 1,0 mg
Excipients q.s.

Each ADAX RETARD sustained release tablet contains:
Alprazolam 0,5 mg
Excipients q.s.

Alprazolam 1,0 mg
Excipients q.s.

Alprazolam 2,0 mg
Excipients q.s.

Presentations:

ADAX: 30 scored tablet pack: 0.25 mg, 0.50 mg and 1 mg
ADAX RETARD: 30 sustained-release tablet pack: 0.5 and 1 mg

Indications:

Neurotic or reactive depression.

For the management of anxiety disorders as a coadjuvant treatment for anxiety associated with depression.

In the treatment of panic disorders with or without agoraphobia.

AERO-ITAN

Prokinetic / Antianxiety / Antiflatulent

Composition:

Each capsule contains:
Chlordiazepoxide hydrochloride 5 mg
Metoclopramide hydrochloride 5 mg
Dimethyl polysiloxane 100 mg
Excipients q.s.

Presentations:

20 capsule pack

Indications:

AEROITAN is indicated in stress-related dyspeptic conditions presenting with flatulence, postprandial fullness, aerophagy and meteorism. Gastric hypokinesia.

ALERTEX

Vaso-dilatant / Brain Stimulant

Composition:

Each tablet contains:
Modafinil 100 mg
Excipients q.s.

Modafinil 200 mg
Excipients q.s.

Presentations:

100 and 200 mg Alertex tablets: 30 tablet pack

Indications:

Modafinil is indicated to improve alertness in patients with excessive daytime sleepiness associated with narcolepsy.

Uses:
Modafinil is indicated to improve wakefulness in patients with excessive sleepiness associated with:
- Narcolepsy: a CNS disorder characterized by somnolence, often accompanied by sudden attacks of weakness (catalepsy), and nighttime sleep alteration. Occasional hypnagogic hallucinations (the state between waking and sleeping) and/or sleep paralysis before falling asleep and waking up. The disorder involves wakefulness and sleep deregulation.
- Obstructive sleep apnea/hypopnea syndrome. (OSAHS)
- Shift work sleep disorder (SWSD).

ALEXIA / ALEXIA FORTE

Antihistamine

Composition:

Each ALEXIA tablet contains:
Fexofenadine hydrochloride 120 mg
Excipients q.s.

Each ALEXIA FORTE tablet contains:
Fenofexadine hydrochloride 180 mg
Excipients q.s.

Presentations:

ALEXIA: 10 and 30 coated tablet packs
ALEXIA FORTE: 10 and 30 coated tablet packs

Indications:

Fexofenadine is indicated for the treatment of chronic idiopathic urticaria and seasonal allergic rhinitis.

ALEXIA-D

Antihistamine / Decongestant

Composition:

Each capsule contains:
Fexofenadine hydrochloride 60 mg
Pseudoephedrine sulfate 120 mg
Excipients q.s.

Presentations:

10 and 20 capsule packs

Indications:

For the relief of symptoms associated with seasonal allergic rhinitis in adults and children over 12 years of age.

ALIZON

Oral Contraceptive

Composition:

Each active, pink coated tablet contains:
Chlormadinone acetate 2 mg
Ethinyl estradiol 0.03 mg
Excipients q.s.

Each placebo, white coated tablet contains:
Excipients q.s.

Presentations:

Pack containing 28 coated tablets

Indications:

Prevention of pregnancy and is useful for treating signs of androgenization, such as acne.

ALIZON 20

Oral Contraceptive

Composition:

Each active, pink coated tablet contains:
Chlormadinone acetate 2 mg
Ethinyl estradiol 0.02 mg
Excipients q.s.

Each placebo, white coated tablet contains:
Excipients q.s.

Presentations:

Pack containing 28 coated tablets

Indications:

Hormonal contraception. Treatment of moderate acne in women who have been indicated this contraceptive.

ALTAZINC

Zinc Supplement

Composition:

Each ml of oral drops solution contains:
Zinc (as sulfate heptahydrate) 5 mg
Excipients q.s.

Each tablet contains:
Zinc (as sulfate heptahydrate) 10 mg
Excipients q.s.

Zinc (as sulfate heptahydrate) 15 mg
Excipients q.s.

Presentations:

Bottle with 30 ml oral solution for drops: 5 mg / ml (20 drops)
Pack with 40 score tablets of 10 and 15 mg

Indications:

Prophylaxis and treatment of zinc deficiency conditions caused by low contribution or poor intestinal absorption, and other conditions that interfere with the use thereof or that increase its body loss, with prior determination of zinc plasma levels.

Uses:
It is used in conditions requiring specific protection such as: child growth, skeletal development, surgical procedure, nutritional deficiency, and zinc deficiency. Coadjuvant in the therapy of hypogonadism and hypothyroidism.

AMINTA

Oral Contraceptive

Composition:

Each coated tablet contains:
Desogestrel 75 mcg
Excipients q.s.

Presentations:

Pack containing 28 coated tablets

Indications:

Selective oral contraceptive to be used during lactation or when estrogens are contraindicated.

AMOVAL

Antibiotic

Composition:

Each AMOVAL tablet contains:
Amoxicillin (as trihydrate) 500 mg
Excipients q.s.

Amoxicillin (as trihydrate) 750 mg
Excipients q.s.

Each 5 ml of AMOVAL oral suspension contains:
Amoxicillin (as trihydrate) 250 mg
Excipients q.s.

Amoxicillin (as trihydrate) 500 mg
Excipients q.s.

Presentations:

AMOVAL: 21 tablet pack
Oral Suspension: Packs with powder to prepare 100ml of suspension

Indications:

Treatment of infections of the upper and lower respiratory tract, infections of the skin and soft tissues, intra-abdominal sepsis, osteomyelitis, infections of the urinary tract caused by susceptible microorganisms, supported by an antibiogram.

AMOVAL 1 GRAMO

Antibiotic

Composition:

Each tablet contains:
Amoxicillin (As trihydrate) 1000 mg
Excipients q.s.

Presentations:

14 tablet pack
20 tablet pack

Indications:

Treatment of airway bacterial infections. Indicated in infections caused by sensitive germs in localized tissues such as gastrointestinal infections, urinary infections, gonorrhea, skin, soft tissue infections.

Indicated for the eradication of Helicobacter pylori in triple therapy with Clarithromycin and omeprazole or esomeprazole.

AMOVAL DUO / 400 / 800 / 1000

Antibiotic

Composition:

Each 5 ml of reconstituted suspension contains:
Amoxicillin (as trihydrate) 400 mg
Excipients q.s.

Amoxicillin (as trihydrate) 800 mg
Excipients q.s.

Amoxicillin (as trihydrate) 1000 mg
Excipients q.s.

Presentations:

Amoval Duo 400: packs to reconstitute 70 ml of suspension
Amoval Duo 800: packs to reconstitute 35 and 70 ml of suspension
Amoval Duo 1000: packs to reconstitute 50 and 90 ml of suspension

Indications:

Indicated in the treatment of airway infections caused by sensitive germs. Indicated also in infections of other localizations caused by sensitive germs, such as otitis media, skin and soft tissue infections, urinary infections, gonorrhea and gastrointestinal infections.

ANTALIN / ANTALIN FORTE

Antidepressant / Antianxiety

Composition:

Each ANTALIN tablet contains:
Amitriptyline hydrochloride 12,5 mg
Chlordiazepoxide (as hydrochloride) 5 mg
Excipients q.s.

Each ANTALIN FORTE tablet contains:
Amitriptyline hydrochloride 25 mg
Chlordiazepoxide (as hydrochloride) 10 mg
Excipients q.s.

Presentations:

30 tablet packs of ANTALIN or ANTALIN Forte

Indications:

ANTALIN and ANTALIN FORTE are indicated for the treatment of mild to moderate depressive disorders; accompanied by mild to moderate anxiety episodes.

ANTIAX

Antacid / Antiflatulent

Composition:

Each tablet contains:
Magaldrate 480 mg
Simethicone 100 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Magaldrate 480 mg
Simethicone 100 mg
Excipients q.s

Presentations:

6 and 24 tablet packs
180 and 360 ml suspension bottles

Indications:

Treatment of dyspepsia, flatulence, and heartburn

ARIANA

Oral Contraceptive

Composition:

Each active, white coated tablet contains:
Nomegestrol Acetate 2.5 mg
Estradiol 1.5 mg
Excipientes: q.s.

Each placebo, red coated tablet contains:
Excipients q.s.

Presentations:

Pack containing 28 coated tablets

Indications:

Oral contraception.
The decision to prescribe Ariana must take into account the current risk factors of the woman in particular, specifically those of venous thromboembolism, and how the risk of venous thromboembolism with Ariana compares with that of other combined hormonal contraceptives.

ATROPINA

Mydriatic-Cycloplegic

Composition:

Each ml of sterile ophthalmic solution contains:
Atropine sulfate 10 mg (1%)
Excipients q.s.

Atropine sulfate 5 mg (0,5%)
Excipients q.s.

Presentations:

Sealed bottle containing 10 ml of sterile ophthalmic solution

Indications:

The ophthalmic preparations of Atropina are used to produce mydriasis and cyclopegia during retinal and optic disc examination, together with the refractive error measurement.

Atropina is also used in the management of acute inflammatory processes of the iris (iridociclytis), and the uveal tract (uveitis). Atropina is also used for its cyclopegic effect for the treatment of amblyopia suppression by reducing visual acuity in the non-affected eye, while the affected eye forces fixation. It has also been used to treat patients with a functional excess of accommodation and convergence.

BEOF

Antiglaucoma

Composition:

Each ml of sterile ophthalmic solution contains:
Betaxolol hydrochloride 5 mg (0,5%)
Excipients q.s.

Presentations:

Sealed drop dispenser bottle with 5 ml of sterile ophthalmic solution

Indications:

BEOF is indicated for the treatment of ocular hypertension and open angle chronic glaucoma. It may be used alone or combined with other medications for glaucoma.

BEQUIUM

Antitussive / Antihistamine / Decongestant

Composition:

Each 5 ml of oral solution contains:
Chlorphenamine maleate 2 mg
Pseudoephedrine hydrochloride 30 mg
Codeine 10 mg
Excipients q.s.

Presentations:

120 ml bottle

Indications:

BEQUIUM is indicated for the symptomatic treatment of non-productive cough, allergic, irritative in origin, associated with bronchitis and/or cold involving nasal congestion.

BLAXITEC

Antihistamine

Composition:

Each tablet contains 20 mg of bilastine.

Presentations:

Package with 30 tablets

Indications:

Symptomatic treatment of allergic rhinoconjunctivitis (seasonal and perennial) and urticaria. Blaxitec 20 mg tablets are indicated in adults and adolescents (age equal to or greater than 12 years).

BLAXITEC ODT

Antihistamine

Composition:

Each orodispersible tablet contains:
10 mg of bilastine
Excipients q.s.

Presentations:

Package with 30 orodispersible tablets

Indications:

Symptomatic treatment of allergic rhinoconjunctivitis (seasonal and perennial) and urticaria. Blaxitec ODT 10 mg orodispersible tablets are indicated for children aged 2 to 11 years.

BLOX

Antihypertensive

Composition:

Each tablet contains:
Candersartan cilexetil 8 mg
Excipients q.s.

Candersartan cilexetil 16 mg
Excipients q.s.

Candersartan cilexetil 32 mg
Excipients q.s.

Presentations:

Blox 8 mg: package with 30 tablets
Blox 16 mg: package with 30 tablets
Blox 32 mg: package with 30 tablets

Indications:

It is indicated for:
Treatment of High Blood Pressure.
Treatment of patients with heart failure and left ventricular systolic dysfunction (ejection fraction less than or equal to 40%), as a complement of ACE inhibitors, or in case of intolerance to this kind of medications.

BLOX AM

Antihypertensive

Composition:

Each BLOX AM 8/5 capsule contains:
Candesartan cilexetil 8 mg
Amlodipine besilate 5 mg
Excipients q.s.

Each BLOX AM 16/5 capsule contains:
Candesartan cilexetil 16 mg
Amlodipine besilate 5 mg
Excipients q.s.

Each BLOX AM 16/10 capsule contains:
Candesartan cilexetil 16 mg
Amlodipine besilate 10 mg
Excipients q.s.

Presentations:

Each package of BLOX AM contains 30 capsules.

Indications:

BLOX AM is indicated as substitution therapy in adult patients with essential hypertension whose blood pressure is already adequately controlled on candesartan and amlodipine administered concomitantly at the same dose level.

BLOX-D

Antihypertensive / Diuretic

Composition:

Each tablet contains:
Candersartan cilexertil 8 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Candersartan cilexetil 16 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Candersartan cilexetil 32 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Presentations:

30 scored tablet packs of:
BLOX-D 8/12.5
BLOX-D 16/12.5
BLOX-D 32/ 12.5

Indications:

This product is indicated for the treatment of idiopathic arterial hypertension in patients not achieving an appropriate control thereof, with a monotherapy with candesartan cilexethyl or hydrochlorothiazide.

BRIMOF

Antiglaucoma

Composition:

Each 1 mL (28 drops)of ophthalmic solution contains:
Brimonidine Tartrate 2 mg
Excipients: q.s.

Each 100 ml of ophthalmic solution contains:
Brimonidine Tartrate 0,2 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophtalmic solution

Indications:

BRIMOF is indicated for the reduction of elevated intraocular pressure in patients with open-angle glaucoma and ocular hypertension.

BRINZOF-T

Antiglaucoma

Composition:

Each 1 mL (27 drops) of ophthalmic suspension contains:
Brinzolamide 10mg
Timolol (as maleate) 5 mg
Excipients q.s.

Each 100 mL of ophthalmic suspension contains:
Brinzolamide 1.0 g
Timolol (as maleate) 0.5 g
Excipients q.s.

Presentations:

Sealed drop bottle with 5 mL of sterile ophthalmic suspension

Indications:

Lowering of intraocular pressure (IOP) in adult patients with open-angle glaucoma or ocular hypertension for whom monotherapy does not provide sufficient IOP reduction.

BRONCATOX

Antitussive

Composition:

Each 10 ml of oral solution contains:
Levodropropizine 60 mg
Excipients q.s.

Presentations:

120 ml bottle

Indications:

Symptomatic treatment of non-productive cough

BUXON

Antidepressant / Smoking Cessation Agent

Composition:

Each tablet contains:
Anfebutamona hydrochloride (Bupropion) 150 mg
Excipients q.s.

Presentations:

Aluminum blister/aluminum packs containing 30 and 60 coated sustained-release tablets

Indications:

Indicated for the treatment of depression.
Indicated as a coadjuvant in the treatment of smoking cessation.

CALDEVAL

Vitamins and minerals supplement

Composition:

Each chewable tablet contains:
Calcium carbonate 1250 mg
(Equivalent to 50 mg of ion calcium)
Vitamin D3 400 I.U
Excipients q.s.

Presentations:

Blister pack with 30 and 60 pleasant-tasting flavor, scored tablets

Indications:

Treatment and prophylaxis of calcium deficiency conditions associated with vitamin D deficit. Indicated as stimulant of fracture consolidation processes.
Uses:
Osteoporosis, osteomalacia, rickets, in all forms thereof, hypocalemia associated with hypoparathyroidism. Breastfeeding, pregnancy, pre- and post-menopausic demineralization and old age.

CEFIRAX

Antibiotic

Composition:

Each tablet contains:
Cefpodoxime 200 mg
Excipients q.s.

Each 5 mL of oral suspension contains:
Cefpodoxime 100 mg
Excipients q.s.

Presentations:

Tablets: 10 and 20 tablet pack
Suspension: Bottle containing 75 mL of oral suspension

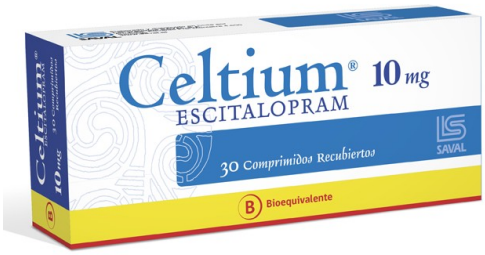
Indications:

Cefpodoxime proxetil is indicated in the oral treatment of respiratory tract infections caused by susceptible microorganisms.
• Acute exacerbations of chronic bronchitis.
• Community-acquired pneumonia.
• Acute maxillary sinusitis.
• Acute otitis media.
• Pharyngitis and tonsillitis caused by streptococcus pyogenes.

It is also indicated in infections caused by susceptible microorganisms:
• Non-complicated gonorrhea.
• Non-complicated skin or skin structure infections.
• Non-complicated urinary tract infections.

CELTIUM

Antidepressant



Composition:

Each coated tablet contains:
Escitalopram (as oxalate) 10 mg
Excipients q.s.

Escitalopram (as oxalate) 20 mg
Excipients q.s.

Presentations:

30 tablet packs of 10 or 20 mg
60 tablet packs of 10 mg

Indications:

CELTIUM contains escitalopram, an antidepressant, indicated for:
• Acute treatment and maintenance of major depressive disorder in adults and adolescents 12 years of age.
• Treatment of panic attacks with or without agoraphobia.
• Treatment of social anxiety disorders (social phobia).
• Treatment of generalized anxiety disorders.
• Treatment of obsessive-compulsive disorders.

CIPRODEX

Ophthalmic Antibiotic / Corticosteroid

Composition:

Each 100 g of ophthalmic ointment contains:
Ciprofloxacin (as hydrochloride monohydrate) 0,3 g
Dexamethasone 0,1 g
Excipients q.s.

Each 100 ml of ophthalmic suspension contains:
Ciprofloxacin (as hydrochloride monohydrate) 0,3 g
Dexamethasone 0,1 g
Excipients q.s.

Presentations:

Ophthalmic suspension: sealed eye drop bottle with 5 mL of sterile ophthalmic suspension
Ophthalmic ointment: ointment tubes with 3.5 g

Indications:

Ocular infections caused by susceptible microorganisms when the anti-inflammatory action of dexamethasone is needed. Blepharitis, blepharoconjunctivitis and conjunctivitis caused by sensitive germs, including Staphylococcus aureus, Staphylococcus epidermidis and Streptococcus pneumoniae. Prophylaxis in ocular surgery and post-surgery.

CIPRODEX OTICO

Otic Antibiotic / Corticosteroid

Composition:

Each ml of otic suspension contains:
Ciprofloxacin (as hydrochloride monohydrate) 3 mg
Dexamethasone 1 mg
Excipients q.s.

Presentations:

7.5 ml sterile suspension bottle

Indications:

Otic ciprodex is indicated in:
• Acute otitis media with tympanostomy due to an infection caused by microorganisms sensitive to Ciprofloxacin children older than 6 months.
• Acute external otitis due to an infection caused by microorganisms sensitive to Ciprofloxacin.

CIPROVAL

Antibiotic

Composition:

Each CIPROVAL tablet contains:
Ciprofloxacin (as hydrochloride monohydrate) 250 mg
Excipients q.s.

Ciprofloxacin (as hydrochloride monohydrate) 500 mg
Excipients q.s.

Presentations:

Pack with 14 tablets of 250 mg
Pack with 10 and 20 tablets of 500 mg

Indications:

This medication is for the treatment of single or mixed infections caused by two or more sensitive microorganisms, for example, respiratory tract infections, urinary infections, gastrointestinal infections, gonorrhea. Additionally, it is an alternative agent against infections caused by antibiotic resistance mechanisms.

CIPROVAL OFTÁLMICO

Ophthalmic Antibiotic

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Ciprofloxacin (as hydrochloride) 0,3 g
Excipients q.s.

Each 100 g of sterile ophthalmic ointment contains:
Ciprofloxacin (as hydrochloride) 0,3 g
Excipients q.s.

Presentations:

Sealed bottle with 5 ml sterile ophthalmic solution.
Ointment container with 3.5 g sterile ophthalmic ointment.

Indications:

CIPROVAL® Ophthalmic contains an antibiotic (ciprofloxacin) that belongs to a group of drugs called fluoroquinolones, which acts against certain microorganisms that cause eye infections.

CIPROVAL® ophthalmic solution: is indicated for the treatment of the following infections in adults, neonates (0-27 days), infants (28 days to 23 months), children (2-11 years) and adolescents (12 to less than 18 years): corneal ulcers, keratitis, corneal abscesses and purulent bacterial conjunctivitis, caused by sensitive bacteria.

CIPROVAL® ophthalmic ointment: is indicated for the treatment of the following infections in adults and children from 1 year of age: purulent bacterial conjunctivitis and blepharitis caused by sensitive bacteria.

Official recommendations on the proper use of antibacterial agents should be taken into account.

CIPROVAL ÓTICO

Otic Antibiotic

Composition:

Each 100 ml of sterile solution for ear drops contains:
Ciprofloxacin (as hydrochloride) 0,3 g
Excipients q.s.

Presentations:

Pack with 5 ml plastic bottle.

Indications:

CIPROVAL Otic Solution is an antimicrobial quinolone indicated in the treatment of otic infections caused by germs sensitive to ciprofloxacin.

CLAVINEX

Antibiotic

Composition:

Each tablet contains:
Clavulanic acid (as potassium clavulanate) 125 mg
Amoxicillin (as trihydrate) 500 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Clavulanic acid (as potassium clavulanate) 62,5 mg
Amoxicillin (as trihydrate) 250 mg
Excipients q.s.

Presentations:

Tablets: 21 tablet pack
Suspension: bottle with powder to prepare 60 ml of 250 mg/5ml suspension

Indications:

Treatment of infections of the upper and lower respiratory tract, urinary tract, skin and soft tissue, intraabdomianl sepsis, osteomyelitis, caused by sensitive microorganisms y beta-lactamase producers supported by an antibiogram.

Clinical uses:
• Acute bacterial sinusitis (properly diagnosed)
• Acute otitis media, tonsillitis
• Acute exacerbation of chronic bronchitis (properly diagnosed)
• Community acquired pneumonia
• Cystitis, urethritis
• Pyelonephritis
• Skin and soft tissue infections, particularly cellulitis, animal bites, severe dental abscesses with spreading cellulitis.
• Bone and joint infections, particularly osteomyelitis.

CLAVINEX DUO CD

Antibiotic

Composition:

Each dispersable tablet contains:
Amoxicillin 875 mg
Clavulanic acid 125 mg
Excipients q.s.

Presentations:

14 and 20 dispersible tablets pack

Indications:

CLAVINEX DUO CD is indicated for the treatment of Beta-lactamase-producing bacteria infections, of the respiratory tract (including ear and nose), urinary tract, skin and soft tissues, intraabdominal sepsis and osteomyelitis caused by organisms sensitive to the association.

CLAVINEX DUO y CLAVINEX DUO FORTE

Antibiotic

Composition:

Each CLAVINEX DUO tablet contains:
Amoxicillin (as trihydrate) 875 mg
Clavulanic acid (as potassium clavulanate) 125 mg
Excipients q.s.

Each 5 ml of CLAVINEX DUO recontituted suspension contains:
Amoxicillin (as trihydrate) 400 mg
Clavulanic acid (as potassium clavulanate) 57 mg
Excipients q.s.

Each 5 ml of CLAVINEX DUO FORTE reconstituted suspension contains:
Amoxicillin (as trihydrate) 800 mg
Clavulanic acid (as potassium clavulanate) 57 mg
Excipients q.s.

Presentations:

CLAVINEX Duo
14 tablet and 20 tablet Packs.
Pack containing 1 bottle with powder to reconstitute 35 ml or 70 ml suspension plus 1 bottle with solvent and dosing syringe

CLAVINEX Duo FORTE
Pack containing 1 bottle with powder to reconstitute 35 ml or 70 ml suspension plus 1 bottle with solvent and dosing syringe

Indications:

Treatment of infections of the upper and lower respiratory tract, urinary tract, skin and soft tissue, intraabdomianl sepsis, osteomyelitis, caused by sensitive microorganisms y beta-lactamase producers supported by an antibiogram.

Used in infections caused by microorganisms sensitive to Amoxicillin and infections caused by microorganisms sensitive to Amoxicillin and beta-lactamase producers such as:

• Infections of the upper respiratory tract: e.g.: tonsillitis, sinusitis, and otitis media.
• Infections of the lower respiratory tract: e.g.: acute chronic bronchitis with exacerbations, bronchopneumonia.
• Infections of the genitourinary tract: cystitis, urethritis, pyelonephritis, female genital infections.
• Skin and soft tissue infections.
• Bone and joint infections: e.g. osteomyelitis
• Other infections: e.g.: septic abortion, puerperal sepsis, intraabdominal sepsis, septicemia, peritonitis, post-operative infections.

• Prophylaxis against infections associated with major surgical procedures such as gastrointestinal, pelvis, head and neck, heart, kidney, bone replacement, and biliary tract surgeries.

CLIMAVAL

Hormone replacement therapy

Composition:

Each tablet contains:
Tibolone 2,5 mg
Excipients q.s.

Presentations:

Pack containing 30 tablets

Indications:

Climaval is indicated in:
• Treatment of estrogen deficiency symptoms in postmenopausal women, more than a year after menopause.
• Prevention of osteoporosis in postmenopausal women at high risk of fractures in the future who cannot tolerate or have contraindications for other approved medicinal products for the prevention of osteoporosis.

NOTE
For all women, the decision to prescribe Climaval should be based on an assessment of the general risks of each patient and, in particular in those over 60 years of age, should include consideration of the risk of stroke.

CLONEX CD

Antianxiety / Anticonvulsivant

Composition:

Each dispersable tablet contains:
Clonazepam 0,5 mg
Excipients q.s.

Clonazepam 1,0 mg
Excipients q.s.

Clonazepam 2,0 mg
Excipients q.s.

Presentations:

CLONEX 0.5 mg, 1.0 mg and 2.0 mg in 30 dispersible tablet packs

Indications:

It is indicated in clinical epilepsy alone or as adjuvant in the treatment of petit mal seizure (absence) and Lennox-Gastaut syndrome. Also used in akinetic and atonic seizures, mioclonic seizure, primary or secondary generalized tonic-clonic seizure, panic attack and panic disorders with or without agoraphobia (fear to be alone in open spaces and public areas).

CLORANFENICOL

Ophthalmic Antibiotic

Composition:

Each ml of ophthalmic solution contains:
Chloranphenicol 5 mg
Excipients q.s.

Each g of ophthalmic ointment contains:
Chloranphenicol 10 mg
Excipients q.s.

Presentations:

10 ml sealed eye drop applicator with sterile ophthalmic solution
3.5 g ointment tube with ophthalmic ointment

Indications:

Superficial eye infections such as: acute microbial conjunctivitis, microbial keratitis of lacrimal ducts, blepharitis or other infections caused by sensitive bacteria.

DEPRAX

Antidepressant

Composition:

Each film coated tablet contains:
Sertraline hydrochloride 50 mg
Excipients q.s.

Each film coated tablet contains:
Sertraline hydrochloride 100 mg
Excipients q.s.

Presentations:

30 and 60 coated scored tablet pack of Deprax 50 mg
30 coated scored tablet pack of Deprax 100 mg

Indications:

Sertraline is indicated for the treatment of depression, including major depressive disorder and relapse prevention, obsessive-compulsive disorder, in adults and pediatric patients aged 6 to 17 years, panic disorder with or without agoraphobia, posttraumatic stress disorder, premenstrual dysphoric disorder, and for the treatment of social anxiety disorder.

DERMABIOTICO

Antiseptic external use

Composition:

Each g of ointment contains:
Polymyxin B (as sulfate) 10000 I.U.
Bacitracin (as Bacitracin zinc) 500 I.U.
Excipients q.s.

Presentations:

Topical ointment, 15 g tube.

Indications:

Antibiotic combination for external use. It is used to prevent infections in cuts, abrasions, scratches and minor burns, bacterial dermatoses sensitive to antibiotics, and it is an essential component in first aid kit.

DERMOSONA

Topical Corticosteroid

Composition:

Each g of cream contains:
Mometasone furoate 1 mg (0,1%)
Excipients q.s.

Presentations:

Tubes of 10 and 15 g.

Indications:

Used to relieve inflammatory and itching manifestations of corticosteroid-sensitive dermatoses, such as atopic dermatitis, allergic contact dermatitis and psoriasis.

DESPEVAL

Antihistamine

Composition:

Each coated tablet contains:
Desloratadine 5 mg
Excipients q.s.

Each 5 ml of oral solution contains:
Desloratadine 2,5 mg
Excipients q.s.

Presentations:

10, 30 and 40 coated tablet packs.
100 ml syrup bottle.

Indications:

For the relief of symptoms associated with allergic rhinitis. Relief of symptoms of urticaria such as the relief of itching and reduction of wheals size.

DIAREN

Antidiarrheal

Composition:

Each tablet contains:
Nifuroxazide 200 mg
Activated attalpugite 350 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Nifuroxazide 200 mg
Activated attalpugite 500 mg
Excipients q.s.

Presentations:

Tablets: 20 tablet packs
Suspension: 100 ml bottle

Indications:

DIAREN is used for acute bacterial diarrheas in children and adults, and chronic diarrheas.

DIGENIL

Antiflatulent / Digestant

Composition:

Each capsule contains:
Pancreatin x 6NF 230 mg
(Equivalent to:5500 FIP U. Amylase, 6500 FIP U. Lipase, 400 FIP U. Protease)
Simethicone 80 mg
Excipients q.s.

Presentations:

Presentation: 10 capsules packs

Indications:

Symptomatic relief of digestive disorders characterized by gastrointestinal flatulence.

DOLODROPS

Nonsteroidal anti-inflammatory

Composition:

Each ml (25 drops) contains:
Diclofenac resinate 46,44 mg
Excipients q.s.

Presentations:

25 ml bottle

Indications:

Inflammatory rheumatism in children. Post-operative and post-traumatic painful and inflammatory conditions. Adjuvant in severe painful infections of the ear, nose and throat infections.

DOLOVERINA

Antispasmodic-Musculotropic

Composition:

Each sustained-release tablet contains:
Mebeverine Hydrocloride: 200 mg
Excipients q.s.

Presentations:

20 sustained-release tablet pack

Indications:

Symptomatic treatment of pain and intestinal discomfort due to functional alterations in the gastrointestinal tract and the bile duct.
Clinical uses:
Irritable bowel syndrome, persistent abdominal pain and cramping, non specific diarrhea (with or without alternating constipation), feeling of fullness in the stomach. Treatment of gastrointestinal spasms secondary to organic disorders. Enteritis.

DRONAVAL

Antiresortive

Composition:

Each coated tablet contains:
150 mg Ibandronic Acid (as Ibandronate monosodium monohydrate)
Excipients c.s.

Presentations:

3 and 1 tablet packs

Indications:

It is indicated for the treatment and prevention of postmenopausal osteoporosis

DRONAVAL PACK

Antiresortive

Composition:

Each coated tablet contains:
Ibandronic Acid (as Ibandronate monosodium monohydrate) 150 mg
Excipients q.s.

Each chewable tablet contains:
Calcium carbonate 1250 mg
(Equivalent to 500 mg of ion calcium)
Vitamin D3 400 I.U
Excipients q.s.

Presentations:

Pack containing: 1 coated tablet of 150 mg Ibandronic Acid and 60 chewable tablets of 500 mg calcium + 400 IU Vitamin D3

Indications:

It is indicated for the treatment and prevention of osteoporosis
Treatment and prevention of postmenopausal osteoporosis

DUALTEN

Antihypertensive

Composition:

Each tablet contains:
Carvedilol 6,25 mg
Excipients q.s.

Carvedilol 12,5 mg
Excipients q.s.

Carvedilol 25 mg
Excipients q.s.

Presentations:

Dualten 6,25 mg: 30 tablet packs
Dualten 12,5 mg: 30 tablet packs
Dualten 25 mg: 30 tablet packs

Indications:

Treatment of the essential Hypertension. Treatment of stable angina. Treatment of symptomatic congestive heart failure.

DUOVAL

Antihistamine / Decongestant

Composition:

Each ml (24 drops) of drop solution contains:
Chlorphenamine maleate 1 mg
Pseudoephedrine hydrochloride 30 mg
Excipients q.s.

Each 5 ml of oral solution contains:
Chlorphenamine maleate 2 mg
Pseudoephedrine hydrochloride 30 mg
Excipients q.s.

Presentations:

Syrup: plastic bottle containing 120 ml solution
Drops: plastic bottle containing 15 ml solution

Indications:

Congestion of the upper respiratory tract (including the paranasal sinuses and the Eustachian tubeS), common cold, hey fever, vasomotor and allergic rhinitis.

ELIXINE

Antiasthmatic

Composition:

Each 15 ml of oral solution contains:
Theophylline 80 mg
Excipients q.s.

Each lentocaps (sustained release capsule) contains:
Theophylline 125 mg
Excipients q.s.

Theophylline 250 mg
Excipients q.s.

Theophylline 300 mg
Excipients q.s.

Presentations:

Oral solution: 250 ml bottle
Pack containing 20 lentocaps of 125 mg
Pack containing 20 lentocaps of 250 mg
Pack containing 20 lentocaps of 300 mg

Indications:

Indicated as bronchodilator in the symptomatic treatment of asthma and reversible bronchospasm associated with chronic bronchitis and emphysema. Indicated for the symptomatic treatment of chronic obstructive pulmonary disease.

ENALTEN

Antihypertensive

Composition:

Each tablet contains:
Enalapril maleate 5 mg
Excipients q.s.

Enalapril maleate 10 mg
Excipients q.s.

Enalapril maleate 20 mg
Excipients q.s.

Presentations:

Pack containing 30 scored tablets of 5 mg, 10 mg and 20 mg, in a double laminated hermetic blister

Indications:

ENALTEN is indicated for the treatment of arterial hypertension and for the treatment of congestive heart failure, concomitantly with other treatments.

ENALTEN-D / ENALTEN-DN

Antihypertensive / Diuretic

Composition:

Each ENALTEN D tablet contains:
Enalapril maleate 10 mg
Hydrochlorotiazide 25 mg
Excipients q.s.

Each ENALTEN DN tablet contains:
Enalapril maleate 20 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Presentations:

ENALTEN-DN 20/12.5: Double laminated hermetic blister pack containing 30 scored tablets
ENALTEN-D (10/25): Double laminated hermetic blister pack containing 30 scored tablets

Indications:

ENALTEN D / ENALTER DN is indicated for the treatment of essential arterial hypertension, resistant to monotherapy.

ETEROVAL

Analgesic / Anti-Inflammatory NSAID

Composition:

Each coated tablet of Eteroval 60 mg contains:
Etoricoxib 60 mg
Excipients q.s.

Each coated tablet of Eteroval 90 mg contains:
Etoricoxib 90 mg
Excipients q.s.

Each coated tablet of Eteroval 120 mg contains:
Etoricoxib 120 mg
Excipients q.s.

Presentations:

Eteroval 60 mg: package with 14 coated tablets
Eteroval 90 mg: package with 14 coated tablets
Eteroval 120 mg: package with 7 coated tablets

Indications:

ETEROVAL is indicated in adults and adolescents over 16 years of age for:

Eteroval 60 mg:
• Symptomatic relief of osteoarthritis and rheumatoid arthritis (RA)
• Treatment of ankylosing spondylitis (AS)
• Treatment of primary dysmenorrhea
• Short-term treatment of moderate pain associated with dental surgery
• Pain and signs of inflammation associated with acute gouty arthritis

Eteroval 90 mg:
• Symptomatic relief of rheumatoid arthritis (RA)
• Treatment of ankylosing spondylitis (AS)
• Short-term treatment of moderate pain associated with dental surgery
• Moderate to severe treatment of acute postoperative pain associated with abdominal gynecological surgery

Eteroval 120 mg:
• Treatment of acute gouty arthritis
• Relief of acute pain
• Treatment of primary dysmenorrhea

The decision to prescribe a selective COX-2 inhibitor should be based on an individual assessment of the patient's overall risks.

EUROCOR

Antihypertensive

Composition:

Each coated tablet of Eurocor 1,25 mg contains:
Bisoprolol Fumarate 1,25 mg
Excipients q.s.

Each coated tablet of Eurocor 2,5 mg contains:
Bisoprolol Fumarate 2,5 mg
Excipients q.s.

Each coated tablet of Eurocor 5 mg contains:
Bisoprolol Fumarate 5 mg
Excipients q.s.

Each coated tablet of Eurocor 10 mg contains:
Bisoprolol Fumarate 10 mg
Excipients q.s.

Presentations:

EUROCOR 1,25 mg: 14 tablet pack
EUROCOR 2,5 mg: 35 tablet pack
EUROCOR 5 mg: 35 tablet pack
EUROCOR 10 mg: 35 tablet pack

Indications:

Bisoprolol fumarate is indicated in:
- Treatment of arterial hypertension
- Treatment of coronary heart disease
- Treatment of stable chronic heart failure concomitantly with ACE inhibitors, diuretics, and optionally cardiac glycosides

EUROCOR AM

Antihypertensive

Composition:

Each coated tablet of EUROCOR AM 5/5 mg contains:
Bisoprolol Fumarate 5 mg
Amlodipine (as besylate) 5 mg
Excipients q.s.

Each coated tablet of EUROCOR AM 5/10 mg contains:
Bisoprolol Fumarate 5 mg
Amlodipine (as besylate) 10 mg
Excipients q.s.

Presentations:

EUROCOR AM 5/5 mg: 35 coated tablets pack
EUROCOR AM 5/10 mg: 35 coated tablets pack

Indications:

EUROCOR AM is indicated in the treatment of arterial hypertension, as a sustitution therapy, in patients properly controlled with the individual products administered concomitantly at the same dose level as in the combination, but as separate tablets.

EUROCOR D

Antihypertensive / Diuretic

Composition:

Each coated tablet of Eurocor-D 2,5/6,25 contains:
Bisoprolol Fumarate 2,5 mg
Hydrochlorotiazide 6,25 mg
Excipients q.s.

Each coated tablet of Eurocor-D 5/6,25 contains:
Bisoprolol Fumarate 5 mg
Hydrochlorotiazide 6,25 mg
Excipients q.s.

Presentations:

Eurocor D 2,5/6,25: 35 coated tablet pack
Eurocor D 5/6,25: 35 coated tablet pack

Indications:

Eurocor-D (Bisoprolol Fumarate and hydrochlorotiazide) is indicated for the treatment of High Blood Pressure

EUROGESIC

Analgesic / Antipyretic / Anti-Inflammatory NSAID / Decongestant

Composition:

Each EUROGESIC ADULTO tablet contains:
Naproxen sodium 275 mg
Excipients q.s.

Each EUROGESIC INFANTIL tablet contains:
Naproxen sodium 100 mg
Excipients q.s.

Each EUROGESIC INFANTIL suppository contains:
Naproxen sodium 50 mg
Excipients q.s.

Each ml of EUROGESIC INFANTIL oral suspension contains:
Naproxen sodium 25 mg
Excipients q.s.

Presentations:

Pack containing 10 adult tablets
Pack containing 10 pediatric tablets
Pack containing 6 suppositories
60 ml oral suspension bottle

Indications:

As analgesic and anti-inflammatory in acute and chronic, painful and inflammatory processes of soft tissues and skeletal muscle. As analgesic and anti-inflammatory for bursitis, tendinitis, synovitis, tenosynovitis and lumbago. For analgesic action after dislocation, sprain, orthopedic manipulations, surgery and dental extraction. For uterine relaxation and post delivery analgesic for the mother who does not breastfeed. As analgesic in dysmenorrhea and after IUD insertion. As anti-inflammatory and analgesic drug in rheumatoid arthritis, osteoarthritis (degenerative arthritis), ankylosing spondylitis and gout.

EUROGESIC FORTE

Analgesic / Antipyretic / Anti-Inflammatory NSAID / Decongestant

Composition:

Each tablet contains:
Naproxen sodium 550 mg
Excipients q.s.

Presentations:

10 tablet pack

Indications:

As analgesic and anti-inflammatory in acute and chronic, painful and inflammatory processes of soft tissues and skeletal muscle. As analgesic and anti-inflammatory for bursitis, tendinitis, synovitis, tenosynovitis and lumbago. For analgesic action after dislocation, sprain, orthopedic manipulations, surgery and dental extraction. For uterine relaxation and post delivery analgesic for the mother who does not breastfeed. As analgesic in dysmenorrhea and after IUD insertion. As anti-inflammatory and analgesic drug in rheumatoid arthritis, osteoarthritis (degenerative arthritis), ankylosing spondylitis and gout.

EUROGREL

Antithrombotic

Composition:

Each coated tablet contains:
Clopidogrel (as bisulfate) 75 mg
Excipients q.s.

Presentations:

Hermetically sealed double aluminum blister pack with 35 tablets

Indications:

EUROGREL is used to prevent vascular events (acute myocardial infarction, ischemic stroke) in patients with a history of symptomatic atherosclerosis. Also used to reduce atherosclerotic events in: myocardial infarction, acute coronary syndrome, ischemic stroke, peripheral arterial disease. Acute coronary syndrome (unstable angina/ non-Q-wave MI).

EUROMICINA

Antibiotic

Composition:

Each tablet contains:
Clarithromycin 250 mg
Excipients q.s.

Clarithromycin 500 mg
Excipients q.s.

Each 5 ml of reconstituted suspension contains:
Clarithromycin 125 mg
Excipients q.s.

Clarithromycin 250 mg
Excipients q.s.

Presentations:

Packs with 14 and 20 tablets of 250 or 500 mg.
80 ml bottle of reconstituted suspension with 125 or 250 mg/5ml EUROMICINA, including solvent for reconstitution.

Indications:

Clarithromycin is indicated for the treatment of infections caused by sensitive organisms. These infections include:
• Infection of the lower respiratory tract
• Infection of the upper respiratory tract.
• Skin and subcutaneous tissue infections.
• Infections caused or spread by Mycobacterium avium or intracellulare.
• Localized infections due to Mycobacterium fortuitum or mycobacterium kanassi.
Indicated together with Amoxicillin and omeprazol for the treatment of eradication of Helicobacter pylori, resulting in a reduction of duodenal ulcer recurrence.

EUROVIR

Antiviral

Composition:

Each tablet contains:
Acyclovir 400 mg
Excipients q.s.

Acyclovir 800 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Acyclovir 200 mg
Excipients q.s.

Each 5 ml of EUROVIR FORTE oral suspension contains:
Acyclovir 400 mg
Excipients q.s.

Presentations:

EUROVIR 400 (Packs with 3 and 15 tablets of 400 mg)
EUROVIR 800 (Packs with 5 and 35 tablets of 800 mg)
EUROVIR Suspension (100 ml Bottle with 200 mg/5ml)
EUROVIR Suspension Forte: 100 ml bottle with 400 mg/5 ml

Indications:

EUROVIR 400 is used in the treatment of herpes labialis and genital herpes, and in some forms of prophylaxis in immunocompromised patients.

EUROVIR 800 is used in: herpes zoster and suppression of recurrent genital herpes.

EUROVIR Suspension is specially used in children with: zoster-varicella, herpetic gingivostomatitis, severe herpes labialis, prophylaxis for herpes simplex infections in immunocompromised patients, and for suppression of recurrent herpes simplex infections in immunocompetent patients.

EUROVIR suspension Forte is used in the treatment of viral infections caused by herpes simplex virus types 1 and 2, herpes zoster and varicella-herpes zoster in immunodepressed patients.

EUROVIR CREMA

Antiviral

Composition:

Each gram of cream contains:
Acyclovir 50 mg
Excipients q.s.

Presentations:

EUROVIR Cream (5 and 15 gr. tubes)

Indications:

EUROVIR cream is used in: herpes labialis. It is also used as an oral coadjuvant therapy for other cutaneous herpetic infections

EVOCAZ CD

Cholinesterase Inhibitor

Composition:

Each dispersible tablet contains:

Donepezil Hydrochloride 5 mg
Excipients q.s

Donepezil Hydrochloride 10 mg
Excipients q.s.

Presentations:

EVOCAZ CD 5 mg: 30 dispersible tablets pack
EVOCAZ CD 10 mg: 30 dispersible tablets pack

Indications:

EVOCAZ CD is indicated for the adjuvant therapy of Alzheimer-type dementia, mild to moderate nature.

FIBROLOW LIDOSE

Antihyperlipidemic

Composition:

Each capsule contains:
Fenofibrate 200 mg
Excipients q.s.

Presentations:

30 lidose capsule pack

Indications:

FIBROLOW Lidose is indicated as a complement to diet and non-pharmacological management (as exercise, weight reduction) in the next cases:

- Treatment of severe hypertriglyceridemia with or without low levels of HDL cholesterol.
- Mixed hyperlipidemia when statins are contraindicated or not tolerated by the patient.
- Mixed hyperlipidemia together with statin in patients with high cardiovascular risk, in whom triglycerides and HDL cholesterol have not been adequately controlled.

FIBROTINA LIDOSE

Antihyperlipidemic

Composition:

Each capsule contains:
Fenofibrate 160 mg
Pravastatin sodium 40 mg
Excipients q.s.

Presentations:

30 lidose capsule pack

Indications:

FIBROTINA LIDOSE is indicated as a supplement for diets and other non-pharmacological treatments (eg, exercise, weight reduction) for the treatment of mixed hyperlipidemia in adult patients with a high cardiovascular risk, to reduce triglycerides and increase the level of HDL cholesterol when LDL cholesterol values are sufficiently controlled when treated with pravastatin 40 mg as monotherapy.

FIBROX XR

Skeletal Muscle Relaxant

Composition:

Each controlled release coated tablet contains:
Cyclobenzaprine hydrochloride 10 mg
Excipients q.s.

Cyclobenzaprine hydrochloride 20 mg
Excipients q.s.

Presentations:

10 and 20 tablet pack

Indications:

Indicated for symptomatic treatment and as a coadjuvant treatment in rest and physical therapy for the relief of muscle spasms associated with acute musculoskeletal contractures.

FINEX

Antifungal

Composition:

Each tablet contains:
Terbinafine (as hydrochloride) 250 mg
Excipients q.s.

Each 100 g of cream contains:
Terbinafine hydrochloride 1 g (1%)
Excipients q.s.

Presentations:

FINEX tablets: packs with 14 and 28 tablets of 250 mg
FINEX cream: 15 g tube

Indications:

FINEX tablets are indicated in fungal infections, where oral therapy is considered appropriate owing to the site, severity or extent of the infection.

FINEX is effective in the treatment of skin, hair and nails infections caused by dermatophytes such as Trichophyton (e.g., T. rubrum, T. mentagrophytes, T. verrucosum, T. tonsurans, T. violaceum), Microsporum canis and Epidermophyton floccosum.

Oral FINEX is indicated in the treatment of: tinea corporis, tinea cruris and tinea pedis, and cutaneous infections caused by yeast of the genus Candida (for example, Candida albicans).Onychomycosis caused by dermatophyte fungi. Oral terbinafine is not effective in pityriasis versicolor and vaginal candidiasis.

Topically applied terbinafine has a rapid onset of action and can be effective with a short duration of treatment. Less than 5% of the dose is absorbed after topical application to humans.

FOSVAL

Antiresortive

Composition:

Each tablet contains:
Alendronic acid (as alendronato sodium trihydrate) 70 mg
Excipients q.s.

Presentations:

FOSVAL 70 mg: 12 tablet blister calendar packs

Indications:

FOSVAL is indicated in the treatment and prevention of Osteoporosis in postmenopausal women.
For the prevention of fractures, including hip and spine.
Indicated for men and women in the treatment and prevention of glucocorticoid-induced osteoporosis.

FREMAVAL

Nasal corticosteroid

Composition:

Each 100 mL of suspension contains:
Fluticasone furoate 0.055 g
Excipients q.s.

Presentations:

Bottle with 120 doses

Indications:

Fremaval Nasal Spray Suspension 27.5 mcg/dose is indicated for the treatment of symptoms of seasonal and perennial allergic rhinitis in adult patients and children 2 years of age and older.

FUNGIUM CREMA

Antifungal

Composition:

Each 100 g of cream contains:
Ketoconazole 2 g (2%)
Excipients q.s.

Presentations:

Tube with 15 g of dermal cream

Indications:

Topical treatment of skin infections: tiena corporis, tinea cruris, tinea manus and tinea pedis caused by Trichophyton sp., Mocrosporum sp., and Epidermophyton sp. Topical treatment of seborrheic dermatitis associated to infections caused by Pityrosporum ovale. Topical treatment of cutaneous candidiasis caused by Candida albicans and Tinea versicolor caused by malassezia furfur.

GESIX

Nonsteroidal anti-inflammatory

Composition:

Each capsule contains:
Celecoxib 200 mg
Excipients q.s.

Presentations:

10 and 30 capsule pack

Indications:

Gesix is indicated for:
Symptomatic treatment of osteoarthritis and rheumatoid arthritis
Relief of the signs and symptoms of ankylosing spondylitis
Acute pain management
Treatment of primary dysmenorrhea

GINTOL

Nutritional Suplement

Composition:

Each capsule for oral administration contains:

Información nutricional	Por 100 g	1 porción	1 porción % *
Vitamina C (mg)	4168	40,00	67
Vitamina E (mg)	1250	12,00	60
Vitamina B1 (mg)	153	1,470	105
Vitamina B2 (mg)	178	1,710	107
Niacina (mg)	1667	16,00	89
Ácido pantonénico (mg)	625	6,00	60
Vitamina B6 (mg)	146	1,40	70
Biotina (mcg)	7918	76	25
Ácido fólico (mcg)	41675	400	200
Vitamina B12 (mcg)	260	2,5	250
Cobre (mg)	104	1,00	50
Hierro (mg)	1459	14,00	100
Zinc (mg)	1042	10,00	67
Manganeso (mg)	104	1,00	50
Selenio (mcg)	5730	55	79
Yodo (mcg)	15628	150	100
Cromo (mcg)	2084	20	57

* Percentage according to recommended daily dose

Presentations:

Pack with 30 soft capsules

HEMOVAL

Iron supplement

Composition:

Each chewable tablet contains:
Iron (as Iron III hydroxide polymallose complex) 100 mg
Excipients q.s.

Each ml of drop solution contains:
Iron (as Iron III hydroxide polymallose complex) 50 mg
Excipients q.s.

Presentations:

Chewable tablets: package of 42 tablets
Oral drops: 30 ml Bottle

Indications:

Prophylaxis and treatment of patients with iron-deficiency due to iron low intake, or poor absorption.
Uses
Iron deficiency states, such as anemia, pregnancy, and different growth stages.

HIALOF HIDRA

Eye lubricant SP

Composition:

Hialof Hidra does not contain preservatives.

Each 1 mL (20 drops) of sterile ophthalmic solution contains:
Sodium hyaluronate: 2 mg
Excipients q.s.

Each 100 mL of of sterile ophthalmic solution contains:
Sodium hyaluronate: 0,2 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 mL ophthalmic solution

Indications:

It is indicated as an ocular lubricant, for the temporary relief of burning, foreign body sensation, stinging and dryness due to dry keratoconjunctivitis (external inflammatory conditions of the eye).

HIDRIUM

Diuretic

Composition:

Each tablet contains:
Furosemide 40 mg
Amiloride hydrochloride 5 mg
Excipients q.s.

Presentations:

20 scored tablet pack

Indications:

HIDRIUM is indicated for the treatment of arterial hypertension, congestive heart failure, edematous states associated with heart failure, heart failure, nephrosis, corticosteroid or estrogen therapies, and cirrhosis-related ascites.

IDON

Antiemetic - Prokinetic

Composition:

Each capsule contains:
Domperidone 10 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Domperidone 5 mg
Excipients q.s.

Each ml of drop solution contains:
Domperidone 10 mg
Excipients q.s.

Presentations:

30 capsule pack
60 capsule pack
Bottle with 15 ml drops
Bottle with 100 ml of pediatric suspension

Indications:

Adults:
• Relief of symptoms such as nausea and vomiting. Stomach burn sensation.
Children:
• Relief of symptoms of nausea and vomiting

IPSON / IPSON FORTE

Analgesic / Antipyretic

Composition:

Each 5ml of IPSON oral suspension contains:
Ibuprofen 100 mg
Excipients q.s.

Each 5 ml of IPSON FORTE oral suspension contains:
Ibuprofen 200 mg
Excipients q.s.

Presentations:

IPSON: PET bottle with 120 ml of oral suspension
IPSON FORTE: PET bottle with 120 ml of oral suspension

Indications:

IPSON is indicated for the treatment of inflammatory and painful processes of soft tissue and for febrile syndrome in patients over 6 months of age. Juvenile rheumatoid arthritis.

IPSON-D / IPSON-D FORTE

Analgesic / Antipyretic / Anti-Inflammatory NSAID

Composition:

Each 5 ml of IPSON D oral suspension contains:
Ibuprofen 100 mg
Pseudoephedrine hydrochloride 15 mg
Excipients q.s.

Each 5 ml of IPSON D FORTE oral suspension contains:
Ibuprofen 200 mg
Pseudoephedrine hydrochloride 30 mg
Excipients q.s.

Each IPSON D FORTE tablet contains:
Ibuprofen 400 mg
Pseudoephedrine hydrochloride 60 mg
Excipients q.s.

Presentations:

Oral suspension: Bottle with 120 ml
Tablets: 10 tablet pack.

Indications:

Ipson-D and Ipson-D Forte are indicated for transient relief of nasal congestion, headache and fever associated with flu or common cold. They are also useful in nasal congestion caused by infection or inflammation as well as in cases of sinusitis.

KEVAL

Antimigraine

Composition:

Each coated tablet contains:
Eletriptan (as hydrobromide) 40 mg
Excipients q.s.

Presentations:

Package with 2 or 6 coated tablets

Indications:

KEVAL® is indicated for the acute treatment of headache phase of migraine attacks with or without aura, for adults.

KI-CAB

Potassium competitive acid blocker

Composition:

Each coated tablet contains:
Tegoprazan 50.0 mg
Excipients: q.s.

Presentations:

Package with 30 coated tablets

Indications:

KI-CAB is indicated for:
- Treatment of erosive gastroesophageal reflux disease
- Treatment of non-erosive gastroesophageal reflux disease
- Treatment of gastric ulcer
- Combination antibiotic therapy for the eradication of Helicobacter pylori in patients with peptic ulcer and/or chronic atrophic gastric disease.

LATOF

Antiglaucoma

Composition:

Each 100 ml of ophthalmic solution contains:
Latanoprost 0,005 g
Excipients q.s.

Presentations:

Sealed drop bottle with 2.5 ml of sterile ophtalmic solution, with a controlled drop dispenser which aids in releasing drops making application easy and preventing drop loss and ensuring dose uniformity.

Indications:

Reduction of elevated intraocular pressure in patients with open-angle glaucoma and ocular hypertension.

LATOF SP

Antiglaucoma

Composition:

Latof SP does not contain preservants.

Each 1 mL (32 drops) of ophthalmic solution of LATOF SP contains:
Latanoprost 0,05 mg
Excipients q.s.

Each 100 ml of ophthalmic solution of LATOF SP contains:
Latanoprost 0,005 g
Excipients q.s.

Presentations:

Sealed drop bottle with 2.5 ml of sterile ophtalmic solution

Indications:

Reduction of elevated intraocular pressure in patients with open angle glaucoma and ocular hypertension in patients with intolerance to other drugs that decrease intraocular pressure.

Reduction of elevated intraocular pressure in pediatric patients with elevated intraocular pressure and pediatric glaucoma.

LATOF-T

Antiglaucoma

Composition:

Each 100 ml of ophthalmic solution contains:
Latanoprost 0,005 g
Timolol maleate 0,5 g
Excipients q.s.

Presentations:

Sealed drop bottle with 2.5 ml of sterile ophtalmic solution, with a controlled drop dispenser which aids in releasing drops making application easy and preventing drop loss and ensuring dose uniformity

Indications:

Reduction of elevated intraocular pressure in patients with open-angle glaucoma and ocular hypertension.

LATOF-T SP

Antiglaucoma

Composition:

LATOF T SP does not contain preservants.

Each 1 mL (33 drops) of ophthalmic solution of LATOF T SP contains:
Latanoprost 0,05 mg
Timolol: 5,00 mg
Excipients q.s.

Each 100 ml of ophthalmic solution of LATOF T SP contains:
Latanoprost 0,005 g
Timolol: 0,500 g
Excipients q.s.

Presentations:

Sealed drop bottle with 2.5 ml of sterile ophtalmic solution

Indications:

It is indicated for the reduction of elevated intraocular pressure (IOP) in patients with open angle glaucoma or ocular hypertension who respond insufficiently to topical agents that reduce IOP.

LEOVAL

Antiepileptic

Composition:

Each coated tablet of LEOVAL 500 mg contains:
Levetiracetam 500 mg
Excipients q.s.

Each coated tablet of LEOVAL 1000 mg contains:
Levetiracetam 1000 mg
Excipients q.s.

Presentations:

LEOVAL 500 mg: package with 30 coated tablets
LEOVAL 1000 mg: package with 30 coated tablets

Indications:

LEOVAL is indicated as monotherapy in the treatment of partial onset seizures, with or without secondary generalization in adults and adolescents over 16 years of age with a new diagnosis of epilepsy.

LEOVAL is indicated as concomitant therapy:
- In the treatment of partial onset seizures with or without secondary generalization in adults, adolescents and children over 4 years of age with epilepsy.
-In the treatment of myoclonic seizures in adults and adolescents over 12 years of age with Juvenile Myoclonic Epilepsy.
-In the treatment of primary generalized tonic-clonic seizures in adults and adolescents over 12 years of age with Generalized Idiopathic Epilepsy.

LEOVAL INYECTABLE

Antiepileptic

Composition:

Each vial of 5 mL of solution contains:
Levetiracetam 500 mg
Excipients q.s.

Presentations:

Pack containing 10 vials of 5 mL

Indications:

LEOVAL is indicated as monotherapy in the treatment of partial onset seizures, with or without secondary generalization in adults and adolescents over 16 years of age with a new diagnosis of epilepsy.

LEOVAL is indicated as concomitant therapy:
- In the treatment of partial onset seizures with or without secondary generalization in adults, adolescents and children over 4 years of age with epilepsy.
- In the treatment of myoclonic seizures in adults and adolescents over 12 years of age with Juvenile Myoclonic Epilepsy.
- In the treatment of primary generalized tonic-clonic seizures in adults and adolescents over 12 years of age with Generalized Idiopathic Epilepsy.

Levetiracetam concentrate is an alternative for patients in whom oral administration is temporarily not feasible.

LEOVAL SOLUCION

Antiepileptic

Composition:

Each mL of oral solution contains:
Levetiracetam 100 mg
Excipients q.s.

Presentations:

Bottle with 300 mL of oral solution

Indications:

LEOVAL is indicated as monotherapy in the treatment of partial onset seizures, with or without secondary generalization in adults and adolescents over 16 years of age with a new diagnosis of epilepsy.

LEOVAL is indicated as concomitant therapy:
- In the treatment of partial onset seizures with or without secondary generalization in adults, adolescents and children over 4 years of age with epilepsy.
-In the treatment of myoclonic seizures in adults and adolescents over 12 years of age with Juvenile Myoclonic Epilepsy.
-In the treatment of primary generalized tonic-clonic seizures in adults and adolescents over 12 years of age with Generalized Idiopathic Epilepsy.

LIFTER

Erectile Dysfunction Agent

Composition:

Each tablet contains:
Sildenafil (as citrate) 50 mg
Excipients q.s.

Sildenafil (as citrate) 100 mg
Excipients q.s.

Presentations:

50 mg tablets: packs with 1, 5 and 10 tablets
100 mg tablets: pack with 5 tablets

Indications:

For the treatment of erectile dysfunction.

LOMEX IV

Antiulcer

Composition:

Each vial of lyophilized contains:
Omeprazole (as sodium salt) 40 mg
Excipients q.s.

Each ampoule contains 10 ml of reconstitution solvent.

Presentations:

LOMEX 40 mg I.V. for injection:
Combined pack containing one vial with omeprazole and one ampoule with 10 ml of solvent.

Indications:

LOMEX I.V.: prophylaxis of acid aspiration, prophylaxis of digestive hemorrhage in critical patients, upper gastrointestinal hemorrhage, Zollinger-Ellison syndrome and ulcer-related gastric retention.

LUDIUM 20 MG

Phosphodiesterase Type 5 Inhibitor

Composition:

Each coated tablet contains:
Tadalafil 20 mg
Excipients q.s.

Presentations:

Package with 1 and 4 coated tablets

Indications:

LUDIUM 20 mg is indicated in:

- The treatment of erectile dysfunction in adult men. In order for LUDIUM to be effective, sexual stimulation is necessary.

The use of LUDIUM is not indicated in women.

LUDIUM 5 MG

Phosphodiesterase Type 5 Inhibitor

Composition:

Each coated tablet contains:
Tadalafil 5 mg
Excipients q.s.

Presentations:

Package with 30 coated tablets

Indications:

LUDIUM 5 mg is indicated in:

- Treatment of erectile dysfunction in adult men. In order for LUDIUM to be effective, sexual stimulation is necessary.

- Treatment of the signs and symptoms of benign prostatic hyperplasia in adult men.

The use of LUDIUM is not indicated in women.

LUKANEX

Antiasthmatic

Composition:

Each chewable tablet contains:
Montelukast sodium: 4 mg
Excipients q.s.

Montelukast sodium: 5 mg
Excipients q.s.

Each coated tablet contains:
Montelukast: 10 mg
Excipients q.s.

Each sachet contains:
Montelukast: 4 mg
Excipients q.s.

Presentations:

4 mg granulated Lukanex: 40 sachets pack
4 mg Lukanex chewable tablets: 40 chewable tablet pack
5 mg Lukanex chewable tablets: 40 chewable tablet pack
10 mg Lukanex coated tablets: 40 coated tablets pack

Indications:

Montelukast is indicated in:
- Prophylaxis and chronic treatment of asthma in adults and pediatric patients from 12 months of age
- Prevention of exercise-induced bronchoconstriction in adult and adolescent patients older than 15 years of age
- Relief of symptoms of seasonal allergic rhinitis in adult and adolescent patients from 2 years of age
- Relief of symptoms of perennial allergic rhinitis in adult patients and children form 6 months of age

MENTANIA

Vaso-dilatant / Brain Stimulant

Composition:

Each soft capsule contains:
Standardized dry extract of Gingko biloba leaves 60 mg
(Equivalent to 14,4 mg of flavonglycosides)
Standardized dry extract of Panax ginseng root C.A.Meyer 100 mg
(Equivalent to 4 mg of glycosides)
Excipients q.s.

Presentations:

30 soft capsule packs

Indications:

Treatment of cerebral vascular insufficiency and some its manifestations such as cognitive deficit, mood swings and the like.

MIGRAX

Antijaquecoso

Composition:

Each coated tablet contains:
Acetaminophen 450 mg
Caffeine 40 mg
Dihydroergotamine mesylate 1 mg
Excipients q.s.

Presentations:

10 tablet packs

Indications:

MIGRAX is indicated for the symptomatic treatment of vascular headaches.

MIGTAL

Antimigraine

Composition:

Each coated tablet contains:
Naratriptan (as hydrochloride) 2,5 mg
Excipients q.s.

Presentations:

2, 6 and 12 tablet Pack

Indications:

For the treatment of headache – migraine: Indicated for the treatment of acute migraine attacks with or without aura in adults.

MOXAVAL

Antibiotic

Composition:

Each coated tablet contains:
Moxifloxacin (as hydrochloride) 400 mg
Excipients q.s.

Presentations:

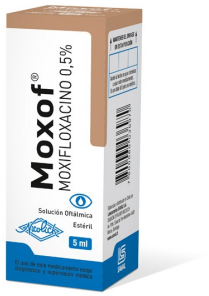
7 and 10 coated tablet packs

Indications:

Moxifloxacin is indicated for the treatment of adults over 18 years of age with upper and lower respiratory tract infections such as acute sinusitis; acute exacerbation of chronic bronchitis; extra hospital pneumonia. In addition, it is indicated for cutaneous infections and soft tissue infections.

MOXOF

Ophthalmic Antibiotic



Composition:

Each 100 ml of ophthalmic solution contains:
Moxifloxacin (as hydrochloride) 0,5 g
Excipients q.s.

Presentations:

Sealed drop-dispenser bottle with 5 mL of sterile ophthalmic solution

Indications:

Moxof is indicated for the treatment of bacterial conjunctivitis caused by susceptible microorganism strains.

MUXELIX

Mucolitic

Composition:

Each 100 ml of oral solution contains:
Hedera helix L. dry extract (5-7,5 : 1) 0,7 g
(Equivalent to no less than 70 mh hederacoside C)
Excipients q.s.

Presentations:

120 ml syrup bottle

Indications:

Treatment of inflammatory bronchial disease symptoms associated with cough.

MUXOL

Mucolitic

Composition:

Each tablet contains:
Ambroxol hydrochloride 30 mg
Excipients q.s.

Each 5 ml of adult oral solution contains:
Ambroxol hydrochloride 30 mg
Excipients q.s.

Each 5 ml of pediatric oral solution contains:
Ambroxol hydrochloride 15 mg
Excipients q.s.

Presentations:

20 tablet pack
100 ml adult syrup bottle
100 ml pediatric syrup bottle

Indications:

Ambroxol is indicated as mucolitic expectorant agent for the relief of productive cough caused by flu states.

NEUROVAL CD

Antianxiety

Composition:

Each dispersable tablet contains:
Clotiazepam 5 mg
Excipients q.s.

Each dispersable tablet contains:
Clotiazepam 10 mg
Excipients q.s.

Presentations:

NEUROVAL®CD 5 mg: 30 dispersable tablet packs
NEUROVAL®CD 10 mg: 30 dispersable tablet packs

Indications:

Treatment of generalized anxiety

NEURUM CÁPSULAS

Antiepileptic

Composition:

Each capsule contains:
Pregabalin 50 mg
Excipients q.s.

Presentations:

Neurum 50 mg: Package with 30 capsules

Indications:

Pregabalin is indicated for the treatment of:
Treatment of neuropathic pain in adults, adjunctive therapy of partial seizures, with or without secondary generalization, in patients 12 years of age and older. Management of fibromyalgia syndrome. Treatment of generalized anxiety disorder.

NEURUM COMPRIMIDOS

Antiepileptic

Composition:

Each tablet contains:
Pregabalin 75 or 150 mg
Excipients q.s.

Presentations:

Neurum 75 mg: Package with 30 scored tablets
Neurum 150 mg: Package with 30 scored tablets

Indications:

Neuropathic pain in adults:
Pregabalin is indicated for the treatment of peripheral and central neuropathic pain in adults.

Epilepsy:
Pregabalin is indicated in adults for the combined treatment of partial seizures with or without secondary generalization in patients from 12 years of age.

Generalized anxiety disorder:
Pregabalin is indicated for the treatment of generalized anxiety disorder (GAD) in adults.

fibromyalgia:
Pregabalin is indicated for the management of fibromyalgia syndrome.

NICODROPS

Ophthalmic Decongestant Drops

Composition:

Each 100 ml of ophthalmic solution contains:
Naphazoline hydrochloride 0,025 g
Dextran-70 0,100 g
Hypromellose 0,300 g
Excipients q.s.

Presentations:

10 ml sterile ophthalmic solution bottle

Indications:

Transient relief of eye redness due to minor ocular irritation. It is indicated in all conditions in which relief of irritation, burning, pruritus and/or ocular congestion is required, for example: dust, smoke, air pollution, sun exposure, wind, contact lenses, allergy, cold or swimming.

NICOL

Oral Contraceptive

Composition:

Each active, white coated tablet contains:
Dienogest 2.00 mg
Ethinyl estradiol 0.03 mg
Excipients q.s.

Each placebo, green coated tablet contains:
Excipients q.s.

Presentations:

Pack containing 28 coated tablets

Indications:

Hormonal contraception.
Treatment of moderately severe acne, which is refractory to local treatment in women in whom the administration of oral contraceptives is not contraindicated.
The decision to prescribe Nicol should take into account the current risk factors of each individual woman, specifically those of venous thromboembolism and how the risk of venous thromboembolism compares with that of other combined hormonal contraceptives.

NICOTEARS

Ophthalmic Drops substitute

Composition:

Each ml of sterile ophthalmic solution contains:
Hypromellose 3 mg
Dextran-70 1 mg
Excipients q.s.

Presentations:

Sealed drop-dispenser bottle with 20 ml sterile ophthalmic solution

Indications:

NICOTEARS is indicated as a lubricating and moistening agent for the eye in the symptomatic treatment of dry eye. It is also indicated in the irritation caused by contact lenses; in Keratitis caused by exposure to irritants; decreased corneal sensitivity; burning; photophobia; conjunctival hyperemia, and generally in recurrent corneal and/or conjunctival erosions and irritations.

NICOTEARS GEL

Substitute for tears

Composition:

Each g of sterile ophthalmic gel contains:
Carbomer 940 1,5 mg
Manitol 46 mg
Excipients q.s.

Presentations:

Sealed tube with 5 g of sterile ophthalmic gel.

Indications:

Relief of the dry eye due to environmental agents such as wind, cigarette smoke, pollution, and solar radiation, or additionally, by physiological conditions manifested by a deficit in tear production. It is also indicated in irritation caused by hard contact lenses; exposure Keratitis, burning, photophobia; conjunctival hyperemia and generally in superior corneal irritations.

NIRVAN

Nonbenzodiazepine Hypnotic

Composition:

Each coated tablet contains:
Eszopiclone 2 mg
Excipients q.s.

Eszopiclone 3 mg
Excipients q.s.

Presentations:

40 scored coated tablets pack

Indications:

Indicated for the treatment of temporary or chronic insomnia

NOVOTEARS

Ophthalmic decongestant

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Naphazoline hydrochloride 0,012 g
Hypromellose 0,3 g
Dextran-70 0,1 g
Excipients q.s.

Presentations:

Sealed drop-dispenser bottle with 10 ml sterile ophthalmic solution

Indications:

Transient relief of eye redness due to minor ocular irritations such as burning sensation, itching, and/or ocular congestion, due to dust, smoke, air pollution, sun exposure, wind, contact lenses, allergy or swimming, winds.

NYSKIN

Antiacne

Composition:

Each 100 g of dermal gel 0,3% contains:
Adapalene 0,3 g
Excipients q.s.

Each 100 g of dermal gel 0,1% contains:
Adapalene 0,1 g
Excipients q.s.

Each 100 g of dermal cream 0,1% contains:
Adapalene 0,1 g
Excipients q.s.

Presentations:

35 g cream or gel tubes

Indications:

Adapalene is indicated for the topical treatment of acne vulgaris.

OBEXOL

Anorexigenic Agent

Composition:

Each capsule contains:
Phentermine Hydrochloride 18,75 or 37,5 mg
Excipients q.s.

Presentations:

30 capsule pack

Indications:

OBEXOL is indicated for short-term treatment of exogenous obesity together with diet to reduce body weight, based on exercise, modification of eating habits and calorie restriction. The treatment is suitable for patients with a body mass index (BMI) greater than or equal to 30 kg / m2 or greater than or equal to 27 kg / m2 in the presence of risk factors such as hypertension, diabetes and hyperlipidemia.

OFTABIÓTICO

Ophthalmic Antibiotic

Composition:

Each ml of sterile ophthalmic solution contains:
Polymyxin B (as sulfate) 5000 I.U.
Neomycin (as sulfate) 1,7 mg
Gramicidin 0,025 mg
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution

Indications:

OFTABIOTICO is indicated for short-term treatment of superficial external ocular infections caused by sensitive organisms.

OFTABIÓTICO UNGÜENTO

Ophthalmic Antibiotic

Composition:

Each g of sterile ophthalmic ointment contains:
Polymyxin B (as sulfate) 6000 I.U.
Neomycin (as sulfate) 3,5 mg
Bacitracin 400 I.U.
Excipients q.s.

Presentations:

3.5 g ointment tubes.

Indications:

OFTABIOTICO ointment is indicated for short-term treatment of superficial external ocular infections caused by sensitive organisms.

OFTAFILM

Eye lubricant

Composition:

Each 100 ml of ophthalmic solution contains:
Sodium hyaluronate 0,400 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution

Indications:

It is indicated as ocular lubricant for the transient relief of burning sensation, foreign body sensation, itching and dryness due to keratoconjunctivitis sicca.

Uses: keratoconjunctivitis sicca, exposure keratitis, neuroparalitic keratitis, mild ocular irritation by sun rays, dust, air, chlorinated waters, weak chemical agents, weak chemical agents, and exposure to strong light.

OFTAFILM SP

Eye lubricant SP

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Sodium hyaluronate 0,400 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 mL ophthalmic solution

Indications:

It is indicated as ocular lubricant for the transient relief of burning sensation, foreign body sensation, itching and dryness due to dry keratoconjunctivitis.

Pediatric use: Do not use, safety and efficacy of this product has not been established in pediatric patients.

Geriatric use: No significant differences were observed in clinical safety and efficacy in elderly patients compared to younger people.

OFTAGEN

Ophthalmic Antibiotic

Composition:

Each ml of sterile ophthalmic solution contains:
Gentamicin (as sulfate) 3 mg
Excipients q.s.

Each g of ophthalmic ointment contains:
Gentamicin (as sulfate) 3 mg
Excipients q.s.

Presentations:

Sealed drop bottle with 5 ml of sterile ophthalmic solution.
3.5g ointment tube.

Indications:

OFTAGEN is indicated for the treatment of bacterial infections of the surface structure of the eye caused by microorganisms sensitive to gentamicin, for example, conjunctivitis, keratitis, keratoconjunctivitis, corneal ulcer, blepharitis, blepharoconjunctivitis, meibomianitis, and dacriocystitis.
It is also indicated in the prophylaxis of postoperative ocular infections.

OFTAGEN COMPUESTO

Ophthalmic Antibiotic / Corticosteroid

Composition:

Each ml of sterile ophthalmic solution contains:
Gentamicin (as sulfate) 3 mg
Bethamethasone disodium fosfate 1 mg
Excipients q.s.

Each g of ophthalmic ointment contains:
Gentamicin (as sulfate) 3 mg
Bethamethasone disodium phosphate 1 mg
Excipients q.s.

Presentations:

Sealed drop bottle with 5 ml of sterile ophthalmic solution
3.5 g ointment tube

Indications:

OFTAGEN is indicated for the treatment of bacterial infections of the surface structure of the eye caused by microorganisms sensitive to gentamicin; associated with inflammatory and allergic conditions such as conjunctivitis, keratitis, and keratoconjunctivitis. Indicated as prophylaxis of postoperative ocular infections of the eye.

OFTALER

Ophthalmic Antihistamine

Composition:

Each 100 ml of ophthalmic solution contains:
Ketotifen (as fumarate) 0,025 g (0,025%)
Excipients q.s.

Ketotifen (as fumarate) 0,050 g (0,050%)
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution 0.05% and 0.025%.

Indications:

Relief of the signs and symptoms of allergic conjunctivitis.

OFTALER FORTE

Ophthalmic Antihistamine

Composition:

OFTALER FORTE
Cada 1 mL (30 gotas) de solución oftálmica contiene:
Ketotifeno (como fumarato) 0,5 mg
Excipientes c.s.

Cada 100 mL de solución oftálmica contiene:
Ketotifeno (como fumarato) 0,05 g
Excipientes: Conforme a la última fórmula autorizada en el registro sanitario.

Presentations:

OFTALER FORTE: Frasco gotario con 10 ml de solución oftálmica estéril

Indications:

Alivio de los síntomas y signos de la conjuntivitis alérgica.

OFTALIRIO

Ophthalmic Decongestant / Antihistamine

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Naphazoline hydrochloride 0,05 g
Antazoline fosfate 0,50 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution.

Indications:

The onset of action with OFTALIRIO can be noted within few minutes after topical application and its effect lasts for 2 -3 hours. Rapid relief of pruritus and ocular congestive symptoms associated with irritating, allergic and inflammatory conditions.

OFTASONA-N

Ophthalmic Antibiotic / Corticosteroid

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Bethamethasone disodium phosphate 0,1 g
Neomycin (as sulfate) 0,35 g
Excipients q.s.

Each 100 g of ophthalmic ointment contains:
Bethamethasone disodium phosphate 0,1 g
Neomycin (as sulfate) 0,35 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution
3 g ointment tube.

Indications:

OFTASONA-N is indicated for the treatment of infectious processes caused by germs sensitive to neomycin accompanied by an inflammatory component.

OFTASONA-P

Ophthalmic Corticosteroid

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Bethamethasone disodium phosphate 0,1 g
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution.

Indications:

Indicated for the treatment of allergic conjunctivitis, inflammatory conditions of the anterior segment of the eye, caused by thermal, chemical, postoperative, and radiation factors.

OFTAVIR

Ophthalmic Antiviral

Composition:

Each 100 g of ophthalmic ointment contains:
Acyclovir 3 g
Excipients q.s.

Presentations:

3.5 g tube with sterile ophthalmic ointment

Indications:

OFTAVIR is indicated in the selective treatment of Herpes simplex keratitis. Coadjuvant therapy in systemic treatment of Herpes Zoster with ocular involvement.

OFTAVITA

Polivitamin with minerals

Composition:

Each coated tablet contains:

Vitamin A (As beta carotene)	3300 IU	(2,00 mg)
Vitamin C (As ascorbic acid)	200,00 mg	
Vitamin E (As DL- Tocopheryl acetate)	75,00 IU	(75,00 mg)
Vitamin B2	5,00 mg	
Zinc (As acetate dihydrate)	30,00 mg	
Selenium (As amino acid complex)	0,02 mg	
Copper (As glycinate chelate)	2,00 mg	
Manganese (As glycinate chelate)	5,00 mg	
Lutein / zeaxanthin	2,00 mg	
Excipients q.s.		

Presentations:

OFTAVITA®: 30 coated tablets pack

Indications:

Vitamin and mineral deficiency states

OFTIC

Ophthalmic Analgesic - Anti-Inflammatory NSAID

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Diclofenac sodium 0,1 g
Excipients q.s.

Presentations:

10 ml sealed pack with sterile ophthalmic solution.

Indications:

OFTIC is used in the treatment of inflammation in patients undergoing cataract surgery.

OFTOL / OFTOL FORTE

Ophthalmic Corticosteroid

Composition:

100 mL of OFTOL ophthalmic suspension contain:
0.200 g Loteprednol Etabonate
Excipients c.s.

100 mL of OFTOL FORTE ophthalmic suspension contain:
0.500 g Loteprednol Etabonate
Excipients c.s.

Presentations:

Sealed drop bottle containing 5 mL of sterile ophthalmic suspension.

Indications:

The physician will indicate the concentration and dose of the drug to be used according to the nature and intensity of the clinical condition. Generally, OFTOL FORTE is used in the treatment of inflammatory conditions of the palpebral and bulbar conjunctiva, cornea and anterior segment of the globe that respond to steroids such as allergic conjunctivitis, acne rosacea, superficial punctate keratitis, herpes zoster keratitis, iritis, cyclitis, selected infective conjunctivititis, when the potential damage of steroid use has been accepted, in order to obtain a decrease in edema and inflammation. It is also used in the treatment of inflammation after ocular surgery. OFTOL is usually indicated for transient relief of signs and symptoms of seasonal allergic conjunctivitis.

OFTOL PLUS

Ophthalmic Antibiotic / Corticosteroid

Composition:

Each 100 ml of Ophthalmic Solution contains:
Loteprednol Etabonate 0,5 g
Tobramycin 0,3 g
Excipients q.s.

Presentations:

Sealed drop bottle with 5 mL of sterile ophthalmic solution

Indications:

The association is indicated for the treatment of inflammatory eye conditions who present superficial bacterial infection or when there is a risk of bacterial ocular infection.

Corticosteroids are indicated for inflammation of the conjunctiva of eyelids and bulbar conjunctiva, cornea, and anterior segment of the eye including allergic conjunctivitis, acne rosacea, superficial punctate keratitis, herpes zoster keratitis, iritis, cyclitis.

It is also used in cases of infectious conjunctivitis when it is beneficial to reduce edema and inflammation and at the same time when the risk of using topical corticosteroids agents is justified.

It is used to treat chronic anterior uveitis and cornea damage due to chemicals components, radiation, burns, or penetration of foreign bodies.

The use of Loteprednol combined with tobramycin (antibiotic) is indicated where the risk of superficial ocular infection is high or where there is a potential risk of bacterial growth in the eye.

OLOF

Decongestant / Antihistamine

Composition:

Each 1 mL (28 drops) of ophthalmic solution contains:
Olopatadine (Hydrochloride) 2 mg
Excipients: q.s.

Each 100 mL of ophthalmic solution contains:
Olopatadine (Hydrochloride) 0,2 g
Excipients: q.s.

Presentations:

Sealed drop bottle containing 5 mL of sterile ophthalmic suspension.

Indications:

Indicated for the treatment of ocular itching associated with allergic conjunctivitis.

OTICUM

Otic Antibiotic / Corticosteroid

Composition:

Each 100 ml of otic solution contains:
Polymyxin B (as sulfate) 1000000 I.U.
Neomycin (as sulfate) 0,35 g
Bethamethasona disodium phosphate 0,1 g
Lidocaine hydrochloride 2 g
Excipients q.s.

Presentations:

5 ml sterile solution pack

Indications:

OTICUM is indicated for the treatment of external otitis accompanied by pain and inflammation, caused by sensitive bacteria. Complementary treatment of acute and chronic otitis media.

Indicated as a coadjuvant in the parenteral antibiotic treatment and other therapeutic measures, and in cavity protection following mastoidectomy and fenestration.

PAXON

Antianxiety

Composition:

Each tablet contains:
Buspirone hydrochloride 5 mg
Excipients q.s.

Buspirone hydrochloride 10 mg
Excipients q.s.

Presentations:

Packs with 20 scored tablets of 5 mg or 10 mg

Indications:

PAXON is indicated for the treatment of acute and chronic anxiety disorders with the following symptoms: anxiety, unsettledness, internal restlessness, and tension states.

PERTIUM

Antihypertensive

Composition:

Each tablet contains:
Nebivolol (as hydrochloride) 5 mg
Excipients q.s.

Presentations:

Package of 42 tablets

Indications:

Treatment of essential hypertension alone or in combination with other antihypertensive agents. Treatment of stable, mild, moderate, and severe heart failure in addition to standard therapies (eg diuretics, ACE inhibitors and angiotensin II antagonists) in elderly patients > 70 years of age.

PILOCARPINA

Antiglaucoma / Direct Acting Miotic

Composition:

Each ml of sterile ophthalmic solution contains:

Pilocarpine hydrochloride 20 mg (2%)
Excipients q.s.

Pilocarpine hydrochloride 40 mg (4%)
Excipients q.s.

Presentations:

Sealed drop bottle with 10 ml of sterile ophthalmic solution 2% or 4%.

Indications:

Antagonist of the mydriatic and cycloplegic effects of atropine. Open angle glaucoma. Narrow angle glaucoma.

PROSTOP-D

Alpha adrenergic receptor antagonist

Composition:

Each capsule contains:
Dutasteride 0.5 mg
Tamsulosin (as hydrochloride) 0.4 mg
Excipients q.s.

Presentations:

Pack with 30 capsules

Indications:

For the treatment of the symptoms of benign prostatic hyperplasia (BPH) in men with an enlarged prostate.

PROTIUM - F

Nutritional Suplement

Composition:

Each capsule contains:
Lactobacillus acidophilus La-14® (4x109 UFC)
Lactobacillus rhamnosus HN001® (1x109 UFC)
Excipients q.s.

Presentations:

30 capsules pack

PROTIUM - I

Nutritional Suplement

Composition:

Each capsule contains:
Lactobacillus acidophilus NCFM® + *Bifidobacterium lactis* Bi-07® (1*10¹º UFC)
Excipients q.s.

Presentations:

30 capsules pack

PROTIUM - T

Nutritional Suplement

Composition:

Each capsule contains:
Bifidobacterium lactis HN019® + *Lactobacillus acidophilus* NCFM® ($1*10^{10}$ UFC)
Excipients q.s.

Presentations:

30 capsules pack

PRUVAL

Other drugs for constipation

Composition:

Each coated tablet 1 mg PRUVAL contains:
Prucalopride (as succinate) 1 mg
Excipients q.s.

Each coated tablet 2 mg PRUVAL contains:
Prucalopride (as succinate) 2 mg
Excipients q.s.

Presentations:

Pruval 1 mg: 30 tablet packs
Pruval 2 mg: 30 tablet packs

Indications:

PRUVAL is indicated for the symptomatic treatment of chronic constipation in women for whom laxatives have not provided adequate relief.

QUETIUM

Antipsychotic

Composition:

Each coated tablet of QUETIUM® 25 mg contains:
Quetiapine (fumarate) 25 mg
Excipients q.s.

Each coated tablet of QUETIUM® 100 mg contains:
Quetiapine (fumarate) 100 mg
Excipients q.s.

Each coated tablet of QUETIUM® 200 mg contains:
Quetiapine (fumarate) 200 mg
Excipients q.s.

Each coated tablet of QUETIUM® 300 mg contains:
Quetiapine (fumarate) 300 mg
Excipients q.s.

Presentations:

QUETIUM® 25 mg: package of 30 coated tablets
QUETIUM® 100 mg: package of 30 coated tablets
QUETIUM® 200 mg: package of 30 coated tablets
QUETIUM® 300 mg: package of 30 coated tablets

Indications:

QUETIUM® is indicated in:
- Schizophrenia
- Manic episodes associated with bipolar disorder
- Depressive episodes associated with bipolar disorder
- Preventing recurrences in the treatment of maintenance of bipolar disorder (manic, mixed or depressive episodes) in combination with lithium or valproate

REALTA

Antidepressant

Composition:

Each capsule with enteric-coated granules of Realta 30 mg contains:
Duloxetine: 30 mg
Excipients q.s.

Each capsule with enetric-coated granules of Realta 60 mg contains:
Duloxetine: 60 mg
Excipients q.s.

Presentations:

Realta 30 mg: 30 capsule pack
Realta 60 mg: 30 capsule pack

Indications:

Realta is indicated in:
- Major Depressive Disorder
- Generalized Anxiety Disorder
- Diabetic Peripheral Neurophatic Pain
- Fibromyalgia
- Musculoskeletal chronic pain

REMSIMA

Tumor necrosis factor inhibitor

Composition:

One vial contains 100 mg of infliximab. After reconstitution each mL contains 10 mg of infliximab.

Presentations:

1 vial of 100 mg

Indications:

Rheumatoid arthritis:
Remsima, in combination with methotrexate, is indicated for the reduction of signs and symptoms as well as the improvement in physical function in:
• adult patients with active disease when the response to disease-modifying antirheumatic drugs (DMARDs), including methotrexate, has been inadequate.
• adult patients with severe, active and progressive disease not previously treated with methotrexate or other DMARDs.

In these patient populations, a reduction in the rate of the progression of joint damage, as measured by X-ray, has been demonstrated.

Adult Crohn's disease:
Remsima is indicated for:
• treatment of moderately to severely active Crohn's disease, in adult patients who have not responded despite a full and adequate course of therapy with a corticosteroid and/or an immunosuppressant; or who are intolerant to or have medical contraindications for such therapies.
• treatment of fistulising, active Crohn's disease, in adult patients who have not responded despite a full and adequate course of therapy with conventional treatment (including antibiotics, drainage and immunosuppressive therapy).

Paediatric Crohn's disease:
Remsima is indicated for treatment of severe, active Crohn's disease in children and adolescents aged 6 to 17 years, who have not responded to conventional therapy including a corticosteroid, an immunomodulator and primary nutrition therapy; or who are intolerant to or have contraindications for such therapies. Infliximab has been studied only in combination with conventional immunosuppressive therapy.

Ulcerative colitis:
Remsima is indicated for treatment of moderately to severely active ulcerative colitis in adult patients who have had an inadequate response to conventional therapy including corticosteroids and 6-mercaptopurine (6-MP) or azathioprine (AZA), or who are intolerant to or have medical contraindications for such therapies.

Ankylosing spondylitis:
Remsima is indicated for treatment of severe, active ankylosing spondylitis, in adult patients who have responded inadequately to conventional therapy.

Psoriatic arthritis:
Remsima is indicated for treatment of active and progressive psoriatic arthritis in adult patients when the response to previous DMARD therapy has been inadequate.
Remsima should be administered:
• in combination with methotrexate
• or alone in patients who show intolerance to methotrexate or for whom methotrexate is contraindicated.

Infliximab has been shown to improve physical function in patients with psoriatic arthritis, and to reduce the rate of progression of peripheral joint damage as measured by X-ray in patients with polyarticular symmetrical subtypes of the disease.

Psoriasis:
Remsima is indicated for treatment of moderate to severe plaque psoriasis in adult patients who failed to respond to, or who have a contraindication to, or are intolerant to other systemic therapy including cyclosporine, methotrexate or psoralen ultra-violet A (PUVA).

RINOVAL

Nasal corticosteroid

Composition:

Each dose contains:
Mometasone furoate (as monohydrate) 50 mcg
Excipients q.s.

Presentations:

Nasal spray bottle with 120 doses of mometasone furoate 50 mcg/dose.

Indications:

• Treatment of the symptoms of seasonal and perennial rhinitis in adults and pediatric patients from 2 years of age.
• Prophylaxis of the nasal symptoms of allergic rhinitis in adults and children older than 12 years of age, 2 to 4 weeks prior to the commencement of the exposure to allergens.
• Indicated in adults and patients over 12 years of age as adjuvant to the antibiotic therapy in the treatment of acute sinusitis episodes.
• Treatment of nasal polyps in patients from 18 years of age.

RIVOXA 10 mg

Platelet Aggregation Inhibitor

Composition:

Each Rivoxa film-coated tablet contains:
Rivaroxaban 10 mg
Excipients: q.s.

Presentations:

Rivoxa 10 mg: Envase con 10 comprimidos recubiertos

Bioequivalente: este producto ha demostrado su equivalencia terapéutica.

Indications:

Prevention of venous thromboembolism (VTE) in patients undergoing major orthopedic surgery of the lower extremities.
Treatment of deep vein thrombosis (DVT) and pulmonary embolism (PE), and prevention of recurrences of DVT and PE in adults.

RIVOXA 15 mg y 20 mg

Platelet Aggregation Inhibitor

Composition:

Each Rivoxa film-coated tablet contains:
Rivaroxaban 15 or 20 mg
Excipients: q.s.

Presentations:

Rivoxa 15 mg: Package with 30 coated tablets
Rivoxa 20 mg: Package with 30 coated tablets

Bioequivalent: This product has demonstrated its therapeutic equivalence

Indications:

Prevention of cerebrovascular accident and systemic embolism in adult patients with nonvalvular atrial fibrillation, with one or more risk factors, such as congestive heart failure, hypertension, age ≥ 75 years, diabetes mellitus, previous cerebrovascular accident or transient ischemic attack.

Treatment of deep vein thrombosis (DVT) and pulmonary embolism (PE), and prevention of recurrences of DVT and PE in adults.

RULOXAN

Antihistamine

Composition:

Each tablet contains:
Rupatadine (as fumarate) 10 mg
Excipients q.s.

Presentations:

Package with 30 tablets

Indications:

Symptomatic treatment of allergic rhinitis and chronic idiopathic urticaria in adults over 18 years of age.
Symptomatic treatment of allergic rhinitis and urticaria in adults and adolescents (over 12 years of age).

RULOXAN SOLUCIÓN ORAL

Antihistamine

Composition:

Each 2.5 ml of oral solution contains:
Rupatadine (as fumarate) 2.5 mg
Excipients q.s.

Each 5 ml of oral solution contains:
Rupatadine (as fumarate) 5 mg
Excipients q.s.:

Presentations:

120 mL bottle

Indications:

Ruloxan oral solution 1 mg/mL is indicated for the symptomatic treatment of:
- Allergic rhinitis (including persistent allergic rhinitis) in children aged 2 to 11 years
- Urticaria in children aged 2 to 11 years

RUX

Antihyperlipidemic

Composition:

Each film coated tablet contains:

Rosuvastatin (as calcium salt) 5 mg
Excipiets q.s.

Rosuvastatin (as calcium salt) 10 mg
Excipiets q.s.

Rosuvastatin (as calcium salt) 20 mg
Excipients q.s.

Presentations:

RUX 5 mg: 30 coated tablets pack
RUX 10 mg: 30 and 60 coated tablets packs
RUX 20 mg: 30 coated tablets pack

Indications:

Rosuvastatin is indicated for the treatment of:

Adults:

1. Primary hyperlipidemia and mixed dyslipidemia

Rosuvastatin is indicated as adjunctive therapy to diet to reduce elevated total cholesterol (TC), LDL-C, ApoB, non-HDL cholesterol and triglycerides and to increase HDL-C in adult patients with primary hyperlipidemia or mixed dyslipidemia.

2. Hypertriglyceridemia

Rosuvastatin is indicated as adjunctive therapy to diet for the treatment of patients with hypertriglyceridemia.

3. Type III familial hyperlipoproteinemia.

Rosuvastatin is indicated as adjunctive therapy to diet for the treatment of patients with Type III familial hyperlipoproteinemia (a genetic disease characterized by hypertriglyceridemia, hypercholesterolemia, and the presence of cholesterol-rich, very low density lipoproteins in plasma).

4. Homozygous familial hypercholesterolemia

Rosuvastatin is indicated as adjunctive therapy to other lipid-lowering treatments (e.g., LDL apheresis) or if such treatments are unavailable to reduce LDL-C, total-C, and ApoB in adult patients with homozygous familial

hypercholesterolemia.

5. Generalized atherosclerosis

Rosuvastatin is indicated as adjunctive therapy to diet to slow atherosclerosis progression in adult patients with the aim of reducing total-C and LDL-C.

Pediatrics

1. Treatment of heterozygous familial hypercholesterolemia (HeFH)

It is indicated in 10 to 17-year-old pediatric patients for the treatment of heterozygous familial hypercholesterolemia (HeFH) as a supplement to diet to reduce Total-C, LDL-C and ApoB levels in male and female adolescents (at least one year after menarche). Drug treatment should be started after failing to achieve expected results with a diet therapy, and where laboratory tests show: LDL-C >190 mg/dL, or >160 mg/dL plus family history of early cardiovascular disease (CVD) or two or more cardiovascular risk factors.

RUX EZ

Antihyperlipidemic

Composition:

Each RUX EZ 10/10 coated tablet contains:
Rosuvastatin (as calcium salt) 10 mg
Ezetimibe 10 mg
Excipients q.s.

Each RUX EZ 20/10 coated tablet contains:
Rosuvastatin (as calcium salt) 20 mg
Ezetimibe 10 mg
Excipients q.s.

Presentations:

Package with 30 coated tablets

Indications:

Primary Hypercholesterolemia:
RUX EZ is indicated as an adjunct to diet for the treatment of primary hypercholesterolemia as replacement therapy in adult patients adequately controlled with the individual products, given simultaneously at the same dose level as in the fixed-dose combination, but as separate products.

Prevention of Cardiovascular Events:
RUX EZ is indicated to reduce the risk of cardiovascular events as replacement therapy in patients with coronary artery disease and a history of acute coronary syndrome, who have been adequately controlled using the same active ingredients and doses, but separately as those included in the fixed-dose combination, but as separate medicinal products.

SALCAL

Anorexigenic Agent

Composition:

Each tablet contains:
Femproporex (as hydrochloride) 10 mg
Excipients q.s.

Presentations:

30 tablet pack

Indications:

SALCAL is indicated in: obesity by overfeeding, obesity by sedentarism, obesity in the elderly. And in partial adeposidities: Madelung's disease, Dercum's disease, Barraquer –Simons' progressive lipodystrophy.

SOMNO

Nonbenzodiazepine Hypnotic

Composition:

Each coated tablet contains:
Zolpidem hemitartrate 5 mg
Excipients q.s.

Zolpidem hemitartrate 10 mg
Excipientsq.s.

Presentations:

Packs containing 30 scored tablets of 5 or 10 mg

Indications:

Treatment of short-term insomnia, either sleep-onset insomnia, early morning awakening, or increased number of awakenings at night, for a 2 – 3 week therapy.

SOMNO XR

Nonbenzodiazepine Hypnotic

Composition:

Each coated sustained release tablet contains:
Zolpidem hemitartrate 12,5 mg
Excipients q.s.

Presentations:

Packs containing 30 coated bilayer tablets

Indications:

Somno XR is indicated for the treatment of insomnia characterized by the inability to fall asleep and/or to maintain sleep (measured as awakenings after falling asleep) for a 2 – 3 week therapy.

SUMAVAL

Antineoplastic agent

Composition:

Cada cápsula de SUMAVAL 12,5 mg contiene:
Sunitinib (como malato) 12,5 mg
Excipientes c.s.

Cada cápsula de SUMAVAL 25 mg contiene:
Sunitinib (como malato) 25 mg
Excipientes c.s.

Cada cápsula de SUMAVAL 50 mg contiene:
Sunitinib (como malato) 50 mg
Excipientes c.s.

BIOEQUIVALENCIA: Este producto farmacéutico ha demostrado equivalencia terapéutica.

Presentations:

Package with 30 capsules

Indications:

Tumores del estroma gastrointestinal (GIST):
Sunitinib está indicado en el tratamiento de los tumores del estroma gastrointestinal (GIST), en adultos después del fracaso del tratamiento con mesilato de imatinib debido a resistencia o intolerancia. La experiencia con Sunitinib como tratamiento de primera línea es limitada.

Carcinoma metastásico de células renales (CMCR):
Sunitinib está indicado en el tratamiento del carcinoma metastásico avanzado de células renales (CMCR) en adultos.

Tumor pancreático neuroendrocrino (pNET):
Sunitinib está indicado para el tratamiento de tumores pancreáticos neuroendocrinos (pNET) bien diferenciados, no extraíbles por cirugía o metastáticos, con progresión de la enfermedad en adultos.

Adyuvante en Carcinoma de Células Renales (CCR):
Sunitinib está indicado para el tratamiento adyuvante de pacientes adultos con alto riesgo de carcinoma de células renales (CCR) recurrente después de una nefrectomía.

TELLMI

Antihypertensive

Composition:

Each tablet contains:
Telmisartan 40 mg
Excipients q.s

Telmisartan 80 mg
Excipients q.s

Presentations:

TELLMI 40 mg: 30 tablet pack
TELLMI 80 mg: 30 tablet pack

Indications:

Treatment of essential arterial hypertension. Prevention of cardiovascular morbidity and mortality in patients 55 years of age and older with a high risk of cardiovascular disease.

TELLMI AM

Antihypertensive

Composition:

Each TELLMI AM 40/5 tablet contains:
Telmisartan 40 mg
Amlodipine (as besylate) 5 mg
Excipients: q.s.

Each TELLMI AM 80/5 tablet contains:
Telmisartan 80 mg
Amlodipine (as besylate) 5 mg
Excipients: q.s.

Each TELLMI AM 80/10 tablet contains:
Telmisartan 80 mg
Amlodipine (as besylate) 10 mg
Excipients: q.s.

Presentations:

Package with 30 tablets

Indications:

Indicated for the treatment of hypertension, alone or with other antihypertensive agents.
It may also be used as initial therapy in patients who are likely to require multiple medications to achieve their blood pressure goals. The choice of TELLMI AM tablets as initial therapy for hypertension is based on an assessment of the potential benefits and risks, including whether the patient is able to tolerate the initial dose of this product.

TELLMI D

Antihypertensive / Diuretic

Composition:

Each tablet of 40/12,5 contains:

Telmisartan	40 mg
Hydrochlorothiazide	12,5 mg
Excipients q.s	

Each tablet of 80/12,5 contains:

Telmisartan	80 mg
Hydrochlorothiazide	12,5 mg
Excipients q.s	

Presentations:

TELLMI-D 40/12,5 mg: 30 tablet pack
TELLMI-D 80/12,5 mg: 30 tablet pack

Indications:

Treatment of essential hypertension.

TELLMI-D, in fixed-dose combination (40 mg telmisartan / 12.5 mg hydrochlorothiazide and 80 mg telmisartan / 12.5 mg hydrochlorothiazide), is indicated in adults whose blood pressure can not be adequately controlled with telmisartan alone.

TIOF

Antiglaucoma

Composition:

Each ml of sterile ophthalmic solution contains:
Timolol (as maleate) 2,5 mg (0,25%)
Excipients q.s.

Timolol (as maleate) 5,0 mg (0,50%)
Excipients q.s.

Presentations:

Sealed drop bottle containing 10 ml of TIOF sterile ophthalmic solution 0.25 % and 0.5 %.

Indications:

Treatment of elevated intraocular pressure in patients with ocular hypertension, open-angle glaucoma, or aphakic glaucoma.

TIOF PLUS

Antiglaucoma

Composition:

Each 100 ml of sterile ophtalmic solution contains:
Dorzolamide (as Clorhidrate) 2 g
Timolol (as Maleate) 0,5 g
Excipients q.s.

Presentations:

Sealed drop bottle containing 6 mL of sterile ophtalmic solution
Sealed drop bottle containing 10 mL of sterile ophtalmic solution

Indications:

In the treatment of elevated intraocular pressure in patients with ocular hypertension, open-angle glaucoma, pseudoexfoliative glaucoma or other secondary open-angle glaucoma when concomitant therapy is approved.

TOL 12 FORTE INYECTABLE

Vitamins supplement

Composition:

Each ampoule of TOL 12 FORTE (5000) contains:
Vitamin B1 (Thiamine hydrochloride) 200 mg
Vitamin B6 (Pyridoxine hydrochloride) 100 mg
Vitamin B12 (Hydroxocobalamin) 5000 mcg
Excipients q.s.

Each ampoule of TOL 12 FORTE (10000) contains:
Vitamin B1 (Thiamine hydrochloride) 200 mg
Vitamin B6 (Pyridoxine hydrochloride) 100 mg
Vitamin B12 (hydroxocobalamina) 10000 mcg
Excipients q.s.

Presentations:

Pack containing 3 ampoules of 5,000 or 10,000

Indications:

Indicated in those cases where administration is required due to deficiency conditions and/or requiring a higher intake of complex B vitamins.
Uses:
Neuralgic and neuritis syndromes; intercostal neuralgia; lumbosciatic pain; shoulder-arm syndrome, trigeminal neuralgia; facial paralysis; neuritis, radiculitis, Herpes zoster; viral, deficiency, alcoholic, diabetic polyneuritis and polyneuritis in pregnancy. Sprains; luxations; fractures.
General stimulant in patients with anorexia, during convalescence and in geriatric patients.

TOL 12 ORAL / TOL 12 FORTE

Orexigenic

Composition:

Each 5 ml of oral solution contains:
Vitamin B1 15 mg
Vitamin B6 15 mg
Vitamin B12 50 mcg
Excipients q.s.

Each capsule contains:
Vitamin B1 (Thiamine hydrochloride) 200 mg
Vitamin B6 (Pyridoxine hydrochloride) 200 mg
Vitamin B12 (Hydroxocobalamin acetate) 1000 mcg
Excipients q.s.

Presentations:

250 ml bottle
20 capsule pack

Indications:

Indicated in those cases where administration is required due to deficiency conditions and/or requiring a higher intake of complex B vitamins.
Uses:
Neuralgic and neuritis syndromes; intercostal neuralgia; lumbosciatic pain; shoulder-arm syndrome, trigeminal neuralgia; facial paralysis; neuritis, radiculitis, Herpes zoster; viral, deficiency, alcoholic, diabetic polyneuritis and polyneuritis in pregnancy. Sprains; luxations; fractures.
General stimulant in patients with anorexia, during convalescence and in geriatric patients.

TOL TOTAL JARABE

Vitamins supplement

Composition:

Each 100 ml of oral solution contains:
Vitamin A palmitate 50000 I.U.
Vitamin B1 hydrochloride 18,0 mg
Bitamin B2 phosphate sodium 20,0 mg
Vitamin B6 hydrochloride 28,0 mg
Vitamin B12 0,10 mg
Vitamin C 900,0 mg
Vitamin D3 8000 I.U.
Vitamin E acetate 196,68 mg
Biotin 1,70 mg
Dexpanthenol 70,0 mg
Nicotinamide 220,0 mg
Excipients q.s.

Presentations:

100 ml bottle

Indications:

Treatment and prevention of vitamin deficiency conditions.

TRAVOF SP

Antiglaucoma

Composition:

Each 100 ml of ophthalmic solution contains:
Travoprost 0,004 g
Excipients

Presentations:

Sealed drop bottle with 2,5 mL ophthalmic solution

Indications:

Travof® SP is indicated for the reduction of elevated intraocular pressure in adult patients with ocular hypertension or open-angle glaucoma.
Travof® SP is indicated for the reduction of elevated intraocular pressure in pediatric patients aged 2 months to <18 years with ocular hypertension or pediatric glaucoma.

TRAVOF T SP

Antiglaucoma

Composition:

Each 100 mL of ophthalmic solution contains:
Travoprost 0.004g
Timolol (as maleate) 0.5 g
Excipients q.s.

Presentations:

Sealed drop bottle with 2.5 mL ophthalmic solution

Indications:

Travof T SP is indicated for the reduction of elevated intraocular pressure in patients with ocular hypertension or open-angle glaucoma, in whom treatment with a single drug only results in insufficient reduction of intraocular pressure.

TREX / TREX FORTE

Antibiotic

Composition:

Each TREX tablet contains:
Azithromycin (as dihydrate) 500 mg
Excipients q.s.

Each 5 ml of TREX oral suspension contains:
Azithromycin (as dihydrate) 200 mg
Excipients q.s.

Each 5 ml of TREX FORTE oral suspension contains:
Azithromycin (as dihydrate) 400 mg
Excipients q.s.

Presentations:

Pack with 3 and 6 tablets of 500 mg
Bottle containing 15 mL of Suspension (200 mg/5mL)
Bottle containing 30 mL of suspension (200 mg/5 mL)
Trex suspension Forte: Bottle containing 20 ml of suspension (400 mg / 5 ml). A solvent to prepare the suspension and a dosing syringe for better dose adjustment and administration in children are included

Indications:

Azithromycin is indicated for the treatment of infections of the upper and lower respiratory tract, urinary tract, skin and soft tissues caused by sensitive microorganisms, supported by an antibiogram. Treatment of uncomplicated genital infections due to Chlamydia trachomatis. It is also indicated in the treatment of uncomplicated genital infections due to non-multiresistant Neisseria gonorrhoeae.
Clinical uses:
Infection of the lower respiratory tract, including bronchitis and pneumonia. Exacerbation of pulmonary obstructive chronic disease due to H. Influenzae, M. Catarrhalis and S. Pneumoniae.
Mild or moderate infection of the upper respiratory tract including laryngitis, pharyngitis, tonsillitis, sinusitis caused by S. Pyogenes.
Skin and soft tissue infections.
Azithromycin is indicated for the treatment of infections of the uncomplicated genitourinary tract or sexually transmitted infections (urethritis, endocervicitis) caused by Clamydia trachomatis. Likewise, it is indicated in genital infections caused by non-multiresistant Neisseria gonorrhoeae and Ureaplasma urealyticum.

TRIM 300

Digestive Motility Regulator

Composition:

Each sustained release coated tablet contains:
Trimebutine maleate 300 mg
Excipients q.s.

Presentations:

Packs with 10 and 30 sustained-release tablets

Indications:

TRIM 300 is used in irritable bowel syndrome: functional colonic diseases, inflammatory bowel disease, irritable colon, spastic colon and its manifestations such as intestinal spasms, colonic dysfunction, alternating diarrhea and constipation, meteorism, abdominal pain and distention, constipation, aftermath of gastrectomy.

TRIOFENO

Anti-Flu

Composition:

Each coated tablet contains:
Ibuprofen: 200 mg
Pseudoephedrine Hydrocloride: 30 mg
Chlorpheniramine Maleate: 2 mg
Excipients q.s.

Presentations:

Pack with 20 score tablets

Indications:

TRIOFENO is indicated for the transient relief of nasal congestion, headache, and fever caused by flu states or common cold. TRIOFENO is useful in the management of nasal congestion caused by airway infections or inflammation and also in cases of sinusitis.

TRIOVAL

Anti-Flu

Composition:

Each tablet contains:
Acetaminophen 500 mg
Pseudoephedrine hydrochloride 60 mg
Chlorphenamine maleate 4 mg
Excipients q.s.

Each 5 ml of oral suspension contains:
Acetaminophen 125 mg
Pseudoephedrine hydrochloride 30 mg
Chlorphenamine maleate 2 mg
Excipients q.s.

Each ml of drops solution contains:
Acetaminophen 120 mg
Pseudoephedrine hydrochloride 7,5 mg
Chlorphenamine maleate 0,75 mg
Excipients q.s.

Presentations:

Tablets: packs with 10 and 80 tablets
Suspension: 100 ml suspension bottle
Oral drops: 15 ml drop bottle

Indications:

Relief of the symptoms of flu and common cold.

TRIOVAL DÍA y NOCHE

Anti-Flu

Composition:

Tablets:

Each white tablet (DAY) contains:
Acetaminophen 500 mg
Pseudoephedrine hydrochloride 60 mg
Excipients q.s.

Each blue tablet (NIGHT) contains:
Acetaminophen 500 mg
Pseudoephedrine hydrochloride 60 mg
Chlorphenamine maleate 4 mg
Excipients q.s

Sachets:
Each sachet day for oral solution contains:
Acetominophen: 500 mg
Pseudoephedrine Hydrocloride: 60 mg
Excipients q.s.

Each sachet night for oral solution contains:
Acetominophen: 500 mg
Pseudoephedrine Hydrocloride: 60 mg
Chlorpheniramine Maleate: 4 mg
Excipients q.s.

Presentations:

Tablets: Packs with 15 white tablets (day) and 5 blue tablets (night)
Sachets: Packs with 3 sachets

Indications:

Relief of the symptoms of flu and common cold.

TRUXIMA

Monoclonal Antibody

Composition:

Each mL of concentrate contains 10 mg of rituximab.

Each vial of 50 mL contains 500 mg of rituximab.
Each vial of 10 mL contains 100 mg of rituximab.

Presentations:

- Truxima 500 mg concentrate for solution for infusion: Pack of 1 vial
- Truxima 100 mg concentrate for solution for infusion: Pack of 2 vials

Indications:

The registered formulation of Rituximab is 10mg/ml concentrate for solution for intravenous (IV) infusion, and it is indicated for:

• Non-Hodgkin's Lymphoma (NHL): Rituximab is indicated for the treatment of previously untreated patients with stage III-IV follicular lymphoma (FL) along with chemotherapy or in patients who are chemo-resistant or are in their second and subsequent relapse after chemotherapy. It is also given in patients with CD20 positive large B cell NHL in combination with cyclophosphamide, doxorubicin, vincristine, prednisolone chemotherapy. Additionally, maintenance therapy is indicated for the treatment of FL individuals responding to induction therapy.
• Chronic Lymphocytic Leukaemia (CLL): Rituximab is indicated for the treatment of patients with previously untreated and relapsed CLL in combination with chemotherapy.
• Rheumatoid Arthritis (RA): Rituximab in combination with methotrexate is given to the adult patient with severe active RA who had insufficient response or intolerance to other disease-modifying anti-rheumatic drugs (DMARD) including one or more tumour necrosis factor inhibitor (TNFi) therapies.
• Granulomatosis with Polyangiitis and Microscopic Polyangiitis (GPA/MPA): In combination with glucocorticoids, Rituximab is given for the induction of remission in adult patients with severe, active GPA/MPA.

UNDERAN

Antiseptic external use

Composition:

Each 100 g of ointment contains:
Mupirocin 2 g (2%)
Excipients q.s.

Presentations:

Aluminum tube containing 15 g of UNDERAN.

Indications:

UNDERAN is indicated for the topical treatment of skin infections caused by susceptible pathogens: impetigo caused by Staphylococcus aureus and group A ß-hemolytic Streptococcus (Streptococcus pyogenes), ecthyma, pyoderma, folliculitis, superinfected eczema, furunculosis, epidermolysis bullosa, superinfected dermatosis and infected traumatic skin lesions, burns. Prophylactic treatment for preventing bacterial infections and promoting rapid healing of lesions. As an alternative treatment for perianal streptococcal cellulitis.

VALAX

Antihypertensive

Composition:

Each coated tablet contains:
Valsartan 80 mg
Excipients q.s.

Valsartan 160 mg
Excipients q.s.

Presentations:

35 tablet pack

Indications:

Arterial hypertension:
Indicated for the treatment of arterial hypertension. It may be used alone or concomitantly with other antihypertensive agents.
Heart Failure:
Treatment of heart failure stages II-IV (New York Heart Association) in patient on continuous treatment, for example, with diuretics, digitalics, or beta-blockers. The use of all these conventional treatments is not mandatory.
Post-myocardial infarction:
Valsartan is indicated to improve survival following myocardial infarction in clinically stable patients and with signs and symptoms or radiologic signs of left ventricular failure, or with left ventricular systolic dysfunction.

VALAX D

Antihypertensive / Diuretic

Composition:

Each coated tablet contains:
Valsartan 80 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Valsartan 160 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Valsartan 160 mg
Hydrochlorotiazide 25 mg
Excipients q.s.

Presentations:

35 coated tablet pack

Indications:

Treatment of hypertension in patients whose blood pressure is not adequately controlled with monotherapy

VALAXAM

Antihypertensive

Composition:

Each coated tablet contains:
Valsartan 80 mg
Amlodipine (as besylate) 5 mg
Excipients q.s.

Valsartan 160 mg
Amlodipine (as besylate) 5 mg
Excipients q.s.

Valsartan 160 mg
Amlodipine (as besylate) 10 mg
Excipients q.s.

Presentations:

35 tablet pack

Indications:

Treatment of arterial hypertension in patients whose blood pressure does not respond to therapy with amlodipine or valsartan as monotherapy.

VALAXAM D

Antihypertensive / Diuretic

Composition:

Valaxam D 5/160/12,5: Each coated tablet contains:
Valsartan 160 mg
Amlodipine Besylate 5 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Valaxam D 10/160/12,5: Each coated tablet contains:
Valsartan 160 mg
Amlodipine Besylate 10 mg
Hydrochlorotiazide 12,5 mg
Excipients q.s.

Valaxam D 160/10/25: Each coated tablet contains:
Valsartan 160 mg
Amlodipine Besylate 10 mg
Hydrochlorotiazide 25 mg
Excipients q.s.

Valaxam D 160/10/25: Each coated tablet contains:
Valsartan 160 mg
Amlodipine Besylate 10 mg
Hydrochlorotiazide 25 mg
Excipients q.s.

Valaxam D 320/10/25: Each coated tablet contains:
Valsartan 320 mg
Amlodipine Besylate 10 mg
Hydrochlorotiazide 25 mg
Excipients q.s.

Presentations:

35 tablets pack

Indications:

Treatment of essential arterial hypertension. This fixed-dose combination is not indicated for the initial treatment of hypertension.

VENLAX

Antidepressant

Composition:

Each tablet contains:
Venlafaxine (as hydrochloride) 50 mg
Excipients q.s.

Venlafaxine (as hydrochloride) 75 mg
Excipients q.s.

Presentations:

Packs with 30 scored tablets (50 mg and 75 mg)

Indications:

VENLAX is indicated for the treatment of all kinds of depression, alone or with comorbid anxiety. It is also indicated for the treatment of generalized anxiety disorders.

VERTIFIN

Antivertigo

Composition:

Each tablet contains:
Betahistine dihydrochloride 16 mg
Excipient q.s.

Each tablet contains:
Betahistine dihydrochloride 24 mg
Excipient q.s.

Bioequivalence: This pharmaceutical product has demonstrated therapeutic equivalence.

Presentations:

Package with 30 tablets

Indications:

Pathological disorders due to microcirculatory deficit in the labyrinth: vertigo, tinnitus, hearing loss associated with Meniere's syndrome and correlated vertigo conditions.

VERTIUM

Antivertigo

Composition:

Each tablet contains:
Diphenidol Hydrochloride 25 mg
Excipients q.s.

Presentations:

10 tablet pack
40 tablet pack

Indications:

Vertium is indicated in prevention and symptomatic treatment of peripheral vertigo associated with nausea and vomiting in conditions such as Meniere's disease or inner or middle ear surgery
It is also indicated for prevention and treatment of nausea and vomiting associated with postoperative states, malignant neoplasms, labyrinthitis, and therapies with antineoplastic agents, radiotherapy and infectious diseases

VITAYDE-C

Vitamins supplement

Composition:

Each ml of drops solution contains:
Vitamin A 5000 I.U.
Vitamin C 75 mg
Vitamin D 1000 I.U.
Excipients q.s.

Presentations:

30 ml bottle

Indications:

Used to prevent and treat vitamin A, C and D deficiency in unweaned babies and children.

XOLOF-D

Ophthalmic Antibiotic / Corticosteroid

Composition:

Each 100 ml of sterile ophthalmic solution contains:
Tobramycin 0,3 g
Dexamethasone 0,1 g
Excipients q.s.

Each 100 g of sterile ophthalmic ointment contains:
Tobramycin 0,3 g
Dexamethasone 0,1 g
Excipients q.s.

Presentations:

Sealed drop bottle with 5 ml of sterile ophthalmic suspension
Tubes with 3.5 g of sterile ophthalmic ointment

Indications:

Treatment of steroid-responsive inflammatory ocular conditions accompanied by a bacterial ocular infection, or risk thereof. Used in inflammatory conditions of the palpebral and bulbar conjunctiva, cornea and anterior segment of the eye, in chronic anterior uveitis and corneal injury from chemical or thermal burns, radiation or penetration of foreign bodies.

ZAFIN

Analgesic

Composition:

Each coated tablet contains:
Tramadol hydrochloride 37,5 mg
Acetaminophen 325,0 mg
Excipients q.s.

Presentations:

14 or 28 tablet pack

Indications:

Indicated for the treatment of moderate to severe, acute and chronic pain.

ZIVAL

Antihistamine

Composition:

Each film coated tablet contains:
Levocetirizine dihydrochloride 5 mg
Excipients q.s.

Presentations:

40 tablet pack

Indications:

Indicated in prevention and treatment of skin and airway allergy syndromes, such as, seasonal and perennial allergic rhinitis; and for the treatment of idiopathic chronic urticaria.

ZIVAL SOLUCIÓN

Antihistamine

Composition:

Each 5 ml of oral solution contains:
Levocetirizine dihydrochloride 2,5 mg
Excipients q.s.

Each ml (24 drops) of drops solution contains:
Levocetirizine dihydrochloride 5 mg
Excipients q.s.

Presentations:

Oral solution: Bottle of 120 ml
Oral drops: Dropper bottle of 20 ml

Indications:

Indicated in prevention and treatment of skin and airway allergy syndromes, such as, seasonal and perennial allergic rhinitis; and for the treatment of idiopathic chronic urticaria.

ZOMEL

Antiulcer

Composition:

Each enteric coated tablet contains:
Esomeprazole (as magnesium salt) 20 mg
Excipients q.s.

Each enteric coated tablet contains:
Esomeprazole (as magnesium salt) 40 mg
Excipients q.s.

Bioequivalence: This pharmaceutical product has demonstrated therapeutic equivalence.

Presentations:

Zomel 20 mg: Package with 30 enteric-coated tablets
Zomel 40 mg: Package with 30 enteric-coated tablets

Indications:

Zomel contains a medicine called esomeprazole which belongs to a group of medicines called 'proton pump inhibitors'. These work by reducing the amount of acid the stomach produces.

Zomel is used to treat the following disorders:
Gastroesophageal reflux disease (GERD):
- Treatment of erosive reflux esophagitis
- Long-term preventive treatment of patients with healed esophagitis to prevent relapse
- Symptomatic treatment of gastroesophageal reflux disease (GERD).
- Use in adolescents from 12 years of age in the short-term treatment of gastroesophageal reflux.

In patients who need continuous treatment with non-steroidal anti-inflammatory drugs (NSAIDs):
- Indicated to reduce the incidence of gastric ulcers associated with continuous therapy with NSAIDs in patients over 18 years of age at risk of developing gastric ulcers. Patients at risk are considered to be those over 60 years of age and/or with a documented history of gastric ulcer.

For patients who require continuous treatment with low-dose aspirin (75-325 mg):
- Prevention of gastric and/or duodenal ulcers associated with low-dose aspirin treatment in patients at risk.

Following treatment with esomeprazole IV for maintenance of hemostasis and prevention of rebleeding of gastric and duodenal ulcers.
In combination with an appropriate antibacterial therapeutic regimen to eradicate Helicobacter pylori and to:
- Scarring of duodenal ulcer associated with Helicobacter pylori and
- Prevention of relapses of peptic ulcers associated with Helicobacter pylori.

ZOMEL HP

Antiulcer, Tritherapy

Composition:

Each dispersible tablet contains:
Amoxicillin (As Trihydrate) 1 g
Excipients q.s.

Each coated tablet contains:
Clarithromycin 500 mg
Excipients q.s.

Each capsule with enteric coated microgranules contains:
Esomeprazole 20 mg
Excipients q.s.

Presentations:

Package with 20 doses for 10 days treatment
Package with 28 doses for 14 days treatment

Indications:

Combined therapy indicated for eradication of infections caused by Helicobacter pylori in patients with gastric and/or duodenal ulcer.