



Caring for science.
Caring for you.

PRODUCT LIST 2025



PAIN & FEVER
COUGH & THROAT
ANTI-INFLAMMATORY
GASTROENTEROLOGY
ANTIVIRAL
ANTIBACTERIAL
CARDIOLOGY
CENTRAL NERVOUS SYSTEM
GLUCOCORTICOID
WARTS TREATMENT
CALCIUM FOLINATE
ANTIHISTAMINIC
ANTIDISLIPIDEMIC
BENIGN PROSTATIC HYPERPLASIA
VITAMINS & MINERALS
HAIR LOSS
HAIR CARE
ANTI-SEBORRHEIC
ACNE
CICATRIZATION
DRY & PSORIATIC SKIN
BABY CARE
BATH & HYDRATION
ANIMAL HEALTH

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PRODUCT LIST



MEDINFAR is a fully integrated Pharmaceutical Company with headquarters in Lisbon, **Portugal**. Founded in 1970, Medinfar is focused in R&D, Production, Distribution, Marketing and Sales, being currently the N°1 Portuguese Company in Consumer Health and Dermatology. In Portugal, besides our own brands, we market licensed products in partnerships with top international pharmaceutical companies.

Medinfar out license a wide range of products: Prescription medicines, Consumer Health, Dermocosmetics, Food Supplements and Veterinary products, **in more than 70 countries.**

Through our **R&D - GMP center** and our **EU-GMP manufacturing** site Medinfar Manufacturing, we offer tailor made turnkey solutions, from galenic development to contract manufacturing.

We offer

LICENSING OUT
CONTRACT MANUFACTURING
CONTRACT DEVELOPMENT



human health

LABORATÓRIO MEDINFAR

Marketing and Sales of Prescription
Branded Products

MEDINFAR CONSUMER HEALTH

Marketing and Sales of OTC and Other
Consumer Health Products

GP GENÉRICOS PORTUGUESES

Marketing and Sales of Prescription
Generic Products

manufacturing

MEDINFAR MANUFACTURING

EU • Manufacturing Site

animal health

MEDINFAR SOROLÓGICO

Marketing and Sales of Products for Veterinary Use





CMO
CDMO



CONTRACT MANUFACTURING

Strict compliance with international requirements and certifications:

EU GMP

NP EN ISO 9001:2015
NP EN ISO 14001:2015
NP ISO 45001:2019
EN ISO 22716:2007

Tech Transfer - Integrated Project Management Model With Regulatory Support.

SOLID DOSAGE FORMS

Tablets, Coated Tablets, Pellets, Granules and Capsules

SEMISOLID DOSAGE FORMS

Creams, Ointments, Gels and Suppositories

LIQUID DOSAGE FORMS

Syrups, Solutions and Suspensions

INSTALLED CAPACITY:

130
MILLION
UNITS/YEAR

MORE THAN:

40
CORPORATE
COSTUMERS

GROWING BEYOND:

300
QUALIFIED
PROFESSIONALS

EU GMP SITE

FORMULATION DEVELOPMENT

SCALE-UP

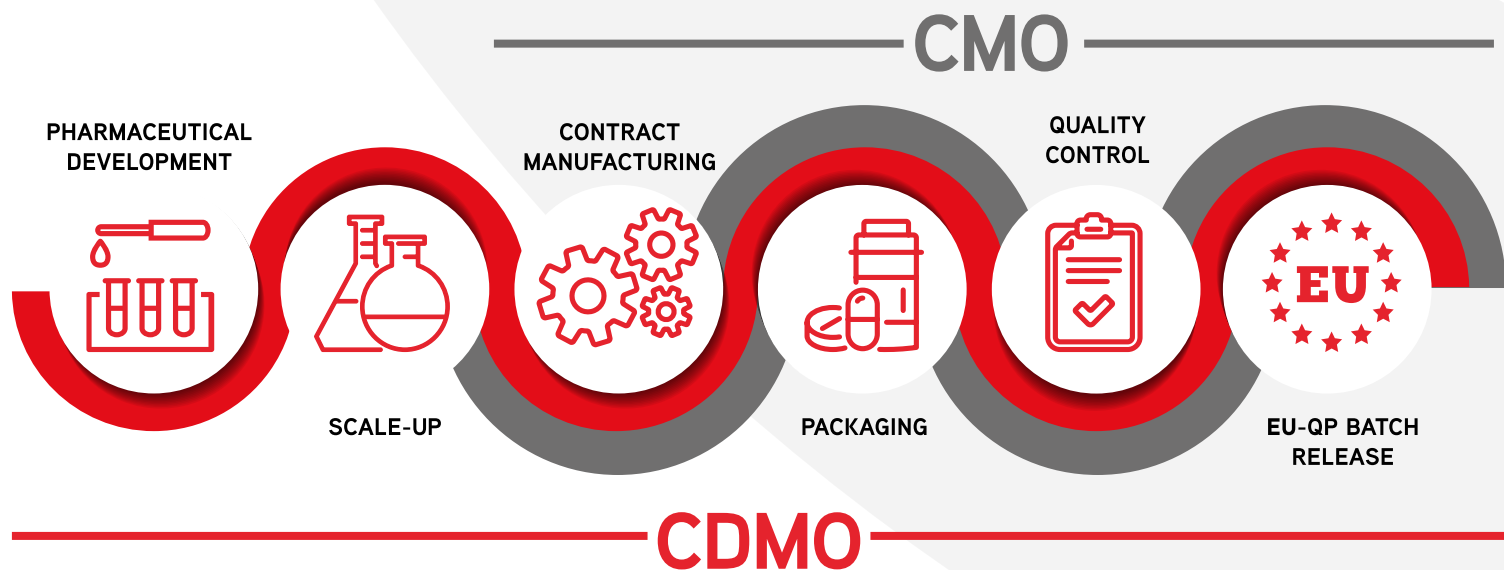
MANUFACTURING PROCESS DEVELOPMENT & VALIDATION

ICH STABILITY

ANALYTICAL METHOD DEVELOPMENT

ANALYTICAL METHOD VALIDATION

CONTRACT DEVELOPMENT & LAB SERVICES





PAIN & FEVER



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Paracetamol	PARAMOLAN	500 mg	Tablet	20 tabs / 500 tabs	Fever, cold or flu symptoms and pain
Paracetamol	PARACETAMOL GP	1 g	Tablet	20 tabs	Fever, cold or flu symptoms and pain
Paracetamol	PARAMOLAN	125 mg 250 mg	Suppository	5 supp	Fever, cold or flu symptoms and pain
Paracetamol	PARAMOLAN	24 mg/ml	Oral solution	100 ml / 200 ml bottle	Fever, cold or flu symptoms and pain
Paracetamol+Vitamin C	PARAMOLAN C	500 mg + 250 mg	Powder for oral solution	20 sachets	Fever, cold or flu symptoms
Paracetamol+Caffeine	PARAMOLAN CAF	500 mg + 65 mg	Tablet	20 tabs	Fever, cold or flu symptoms and mild to moderate pain



PAIN & FEVER



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Ibuprofen	TRIFENE 200	200 mg	Film coated tablet	20 tabs / 60 tabs	Mild to moderate pain, fever and cold or flu symptoms
Ibuprofen	TRIFENE 400	400 mg	Film coated tablet	20 tabs / 60 tabs	Mild to moderate pain, fever and cold or flu symptoms
Ibuprofen	TRIFENE 600	600 mg	Film coated tablet	20 tabs / 60 tabs	Anti-rheumatic, analgesic, antipyretic
Ibuprofen	TRIFENE	20 mg/ml	Oral suspension	100 ml / 200 ml bottle	Mild to moderate pain and fever

PARAMOLAN® *Caf*

Migraine and headache relief



ESSENTIAL INFORMATION COMPATIBLE WITH THE SUMMARY OF PRODUCT CHARACTERISTICS Name of the medicinal product: Paramolan Caf 500 mg/65 mg tablets Qualitative and quantitative composition: Each tablet contains 500 mg of paracetamol and 65 mg of caffeine. Excipients with known effect: sodium (less than 23 mg per tablet) Pharmaceutical form: Tablet. White oblong tablets. Therapeutic indications: Paramolan Caf is indicated in adults and children aged 12 years and over in the treatment of mild to moderate pain and in febrile states. The situations of mild to moderate pain include: Symptoms associated with flu-like syndromes and colds; Headache, including migraine; Dental pain and pain arising from dental interventions; Odynophagia; Dysmenorrhoea; Muscle aches; Post-traumatic pain. If you do not feel better or if you feel worse after 3 days (in case of fever) or, in case of pain, after 5 days, you must talk to a doctor. Posology and method of administration: Posology: UAdults (including the elderly) and children aged 15 years and over 50 kg: U1-2 tablets, 3-4 times daily (every 6 - 8 hours), as required. Maximum daily dose: 3000 mg/390 mg (paracetamol/caffeine) (6 tablets in 24 hours). U Children 12-15 years (< 50 kg): 1 tablet, 3-4 times daily (every 6 - 8 hours), as required. Maximum daily dose: 2000 mg/260 mg (paracetamol/caffeine) (4 tablets in 24 hours). Maximum daily dose: The dose of 3 g of paracetamol daily, i.e. 6 tablets per day should not be exceeded. An interval of 4 hours (at least) should always be observed between administrations. Paediatric population: Paramolan Caf is not recommended for children under 12 years. Renal impairment: In the event of moderate or severe renal impairment, the dose should be reduced, as follows: Creatinine clearance between 50 - 10 ml/min: 1 tablet, every 6 hours (up to 4 times/day). Creatinine clearance <10 ml/min: 1 tablet, every 8 hours (up to 3 times/day). Hepatic impairment: Patients with mild to moderate hepatic impairment or with Gilbert's syndrome should take this medicine with caution. The dose should be reduced, or the dosing intervals extended. Patients with severe liver impairment should not take this medicinal product. U Method of administration: Oral route. Swallow the tablets with sufficient amount of water. Contraindications: Hypersensitivity to the active substances or to any of the excipients; Severe hepatic impairment; Acute hepatitis; Children under the age of 12 years old. Undesirable effects Adverse reactions have been ranked in frequency using the following convention: very common ($\geq 1/10$); common ($\geq 1/100$ to $< 1/10$); uncommon ($\geq 1/1,000$ to $< 1/100$); rare ($\geq 1/10,000$ to $< 1/1,000$); very rare ($< 1/10,000$); not known (cannot be estimated from the available data). Paracetamol: Blood and lymphatic system disorders - Very rare: haematopoiesis disorders (thrombocytopenia, leukopenia, isolated cases of agranulocytosis, pancytopenia); Immune system disorders - Very rare: anaphylaxis and angioedema; Nervous system disorders - Uncommon: vertigo, nervousness; Respiratory, thoracic and mediastinal disorders - Very rare: bronchospasm in predisposed patients (e.g. asthmatic patients sensitive to aspirin and other NSAIDs); Gastrointestinal disorders - Common: nausea, vomiting; Uncommon: diarrhoea, abdominal pain, constipation; Hepatobiliary disorders - Rare: hepatic transaminases increase; Skin and subcutaneous tissue disorders - Rare: erythema; Very rare: cutaneous hypersensitivity reactions, including rash; Not known: toxic epidermal necrolysis and Stevens-Johnson Syndrome. General disorders and administration site conditions disorders - Uncommon: headache, sweating/perspiration, hypothermia; Very rare: allergic reactions, exacerbated hypersensitivity reactions to paracetamol (Quincke's oedema, dyspnoea, sweating, nausea, drop in blood pressure, or even shock). Clinical/epidemiological data seem to indicate that the long-term administration of analgesics may cause nephropathy, including papillary necrosis. Caffeine: The most common undesirable effects of caffeine are nausea caused by irritation of the gastrointestinal tract, as well as insomnia and irritability due to stimulation of the central nervous system. When the recommended dosage of paracetamol-caffeine is combined with dietary intake of caffeine, the resulting dose of caffeine may increase the potential for adverse effects related to caffeine. Psychiatric disorders - Very rare: irritability, insomnia; Gastrointestinal disorders - Very rare: nausea. (11/2021) Product not subject to medical prescription. For more information, please contact the Marketing Authorization Holder. Medinfar Consumer Health - Produtos Farmacêuticos, Lda. Rua Henrique Paiva Couceiro, 27 Venda Nova, 2700-451 Amadora, Portugal



COUGH & THROAT



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Ambroxol, hydrochloride	MUCODRENOL	6 mg/ml	Syrup	200 ml bottle	Mucolytic adjuvant (expectorant)
Bromhexine, hydrochloride	TOSSEQUE	0.8 mg/ml	Syrup	200 ml bottle	Mucolytic adjuvant (expectorant)
Pine buds extract + Eucalyptol + Althea root extract	BEQUISAN	-	Oral Solution	200 ml bottle	Soothing and pleasant effect on throat, pharynx and vocal cords



ANTI- -INFLAMMATORY



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Piroxicam	REUMOXICAN	20 mg	Suppository Capsule Dispersible tablet	12 supp 20 caps / 60 caps 20 disp tabs / 60 disp tabs	Symptomatic relief of osteoarthritis rheumatoid arthritis and ankylosing spondylitis
Diethylamine Salicylate	MEDALGINAN Gel	50 mg/g	Gel	40 g tube 100 g tube	Myalgia, lumbago, strains, sprains and muscle spasms
Meloxicam	MELOXICAM GP	7.5 mg 15 mg	Tablet Tablet	20 tabs / 60 tabs 20 tabs / 60 tabs	Symptomatic relief of osteoarthritis, rheumatoid arthritis and ankylosing spondylitis



GASTRO- ENTEROLOGY

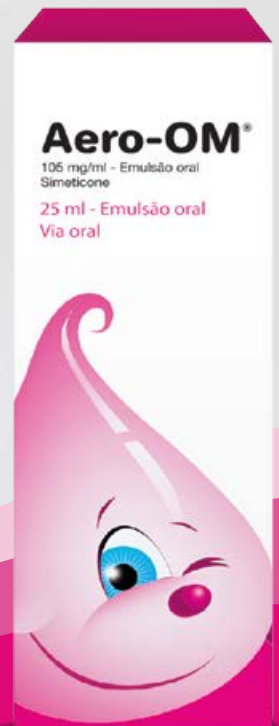


MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Omeprazole	PROTON 20 PROTON 40 OMEPAZOL GP	20 mg 40 mg	GR capsule	14 GR caps 28 GR caps 56 GR caps	GERD and heartburn
Metoclopramide	METOCLOPRAMIDA MEDINFAR	10 mg	Tablet	40 tabs	Prevention and relief of symptoms of nausea and vomiting
Domperidone	CINET	1 mg/ml	Oral suspension	200 ml bottle	Relief of symptoms of nausea and vomiting
Domperidone	CINET DOMPERIDONA GP	10 mg	Tablet	10 tabs / 20 tabs /60 tabs	Relief of symptoms of nausea and vomiting
Domperidone	CINET	10 mg	Dispersible tablet	20 disp tabs / 60 disp tabs	Relief of symptoms of nausea and vomiting
Phosphate Disodium + Monosodium Phosphate	PRECLINT	240 mg /ml + 542 mg/ml	Oral solution	2 x 45 ml bottle	Osmotic laxative (colonoscopy preparation)
Simethicone	AERO-OM	42 mg	Chewable tablet	60 tabs / 100 tabs	Functional treatment of symptoms associated with gas (abdominal distension, flatulence and meteorism)
Simethicone	AERO-OM	105 mg/ml	Oral emulsion	25 ml bottle	Symptomatic treatment of flatulent colic, difficulties in burping and regurgitation, and functional abdominal distension

Aero-OM[®]

**FEWER MOMENTS OF COLIC
MORE MOMENTS AERO-OM[®]**

GAS | COLIC | BLOATED TUMMY



NEWBORNS, BABIES AND CHILDREN

ESSENTIAL INFORMATION COMPATIBLE WITH THE SUMMARY OF PRODUCT CHARACTERISTICS

Name of the medicinal product: Aero-OM, 105 mg/ml, oral emulsion Qualitative and quantitative composition: One millilitre of oral emulsion contains 105 mg of simethicone as active substance. Excipient with known effect: Sucrose - 300 mg per millilitre of oral emulsion Pharmaceutical form: Oral emulsion. Translucent syrupy white liquid. Therapeutic indications: Symptomatic treatment of flatulent colic, difficulties in burping and regurgitation, and functional abdominal distension. Posology and method of administration: Oral route. Shake before using. Aero-OM oral emulsion: Children aged up to 1 year: 5 to 10 drops 3 or 4 times a day (20 to 40 mg/dose). Children aged 15 years: 10 drops 3 or 4 times a day (40 mg/dose). The duration of treatment should not exceed 10 days. Contraindications: Hypersensitivity to the active substance or to any of the excipients. Undesirable effects: In view of the absence of intestinal absorption, the risk of undesirable effects with simethicone is limited. The incidence of adverse events in published clinical trials is low (1.6%, based on close to 2500 patients exposed), and are primarily gastrointestinal alterations (overall incidence of 1%). The undesirable effects are classified according to the MedDRA convention, by system, organ, class and by frequency as follows: Very common ($\geq 1/10$) Common ($\geq 1/100$ to $<1/10$) Uncommon ($\geq 1/1000$ to $<1/100$) Rare ($\geq 1/10,000$ to $<1/1000$) Very rare ($<1/10,000$) Unknown (cannot be estimated from the available data). Gastrointestinal alterations: Uncommon: diarrhoea abdominal pain, ileus, nausea and vomiting. Immune system disorders: Rare: hypersensitivity reactions, such as rash, itching, swelling of the face and tongue, and breathing difficulty were also reported. (07/2023) This medicinal product is not subject to medical prescription. For more information, please contact the Marketing Authorization Holder. Medinfar Consumer Health - Produtos Farmacêuticos, Lda. Rua Henrique Paiva Couceiro, 27 Venda Nova, 2700-451 Amadora, Portugal.

ANTIVIRAL



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Aciclovir	CICLOVIRAL	200 mg 400 mg 800 mg	Tablet	25 tabs 25 tabs 25 tabs / 35 tabs / 60 tabs	Herpes simplex and varicella zoster infections
Aciclovir	CICLOVIRAL CREME	50 mg/g	Cream	10 g tube	Cold sores treatment (Herpes simplex infection)

ANTI-BACTERIAL



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Ciprofloxacin, hydrochloride	CIPROFLOXACINA GP	250 mg 500 mg	Tablet	8 tabs / 16 tabs 16 tabs	Bacterial infection



cicloviral[®] cream

YOUR SHIELD AGAINST HERPES



ESSENTIAL INFORMATION COMPATIBLE WITH THE SUMMARY OF PRODUCT CHARACTERISTICS

Name of the medicinal product: CICLOVIRAL 50 mg/g Cream Qualitative and quantitative composition: Each gram of cream contains 50 mg of acyclovir (5% of acyclovir, w/w). Excipients: Each gram of cream contains 67.5 mg cetostearyl alcohol; Each gram of cream contains 400 mg propylene glycol. Pharmaceutical form: Cream Therapeutic indications: CICLOVIRAL cream is indicated in the treatment of labial herpes resulting from infection by the Herpes simplex virus type 1. Posology and method of administration: Method of administration: cutaneous use. Adults and children aged 12 or older: CICLOVIRAL cream should be applied five times daily, at approximately 4-hour intervals, omitting the nighttime period. CICLOVIRAL cream should be applied to lesions or imminent lesions, as early as possible after the infection has started. In recurrent episodes, it is particularly important to start treatment in the early stages (prodrome or erythema), or when lesions appear. Studies in patients with labial herpes have demonstrated that treatment is still effective when started after the lesions have appeared (papular and vesicular stages). Treatment should be continued for 4 days. If healing has not occurred after 4 days, treatment may be continued for up to 10 days. If lesions are still present after 10 days, users should be advised to consult a doctor. Children aged 2 to 12 years: The CICLOVIRAL cream doses used in adults are recommended, under medical surveillance, for the following reasons: the stratum corneum is comparable in children and adults; no safety concerns are raised regarding the potential limit of systemic absorption of acyclovir from CICLOVIRAL cream; and there is no evidence to suggest the existence of any differences between age groups regarding the natural history of recurrent episodes of labial herpes. Users should wash their hands before and after applying the cream. Users should also avoid touching or rubbing the lesions, or even touching the lesions with a towel, to avoid aggravating or transferring the infection. Contraindications: Hypersensitivity to acyclovir or any of the excipients. Undesirable effects A transient burning sensation may occur sporadically, following the application of CICLOVIRAL cream. Some cases of mild drying and flaking of the skin have been reported following cutaneous application of acyclovir. Erythema, pruritus and contact dermatitis may occur in rare cases (12/2011) This medicinal product is not subject to medical prescription. For more information, please contact the Marketing Authorization Holder. Medinfar Consumer Health - Produtos Farmacêuticos, Lda. Rua Henrique Paiva Couceiro, 27 Venda Nova, 2700-451 Amadora, Portugal



CARDIOLOGY



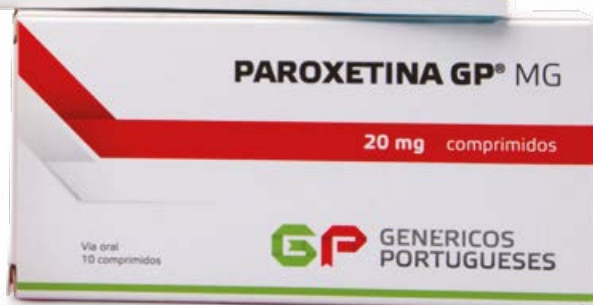
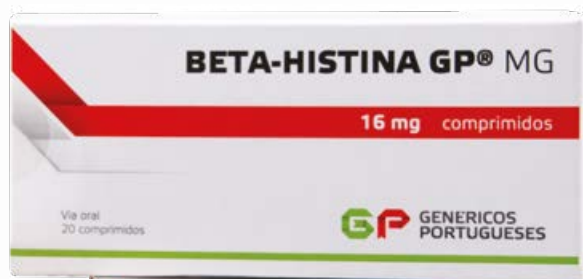
MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Acetylsalicylic Acid	ASP	100 mg	Tablet	20 tabs / 30 tabs / 60 tabs	Anti-thrombotic
Amiodarone, hydrochloride	AMIODARONA GP	200 mg	Tablet	10 tabs / 60 tabs	Hypertension and angina
Amlodipine, besilate	AMLODIPINA GP	5 mg 10 mg	Tablet	10 tabs / 60 tabs 60 tabs	Tachycardia and arrhythmia associated with several conditions
Captopril	CAPTOPRIL GP	25 mg 50 mg	Tablet	20 tabs / 60 tabs	Hypertension and heart failure
Carvedilol	CARVEDILOL GP	6.25 mg 25 mg	Tablet	14 tabs / 56 tabs	Hypertension, coronary heart disease and heart failure
Indapamide	INDAPAMIDA GP	2.5 mg	Film coated tablet	15 tabs / 60 tabs	Hypertension



CARDIOLOGY



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Lisinopril	LISINOPRIL GP	5 mg 20 mg	Tablet	10 tabs / 60 tabs 60 tabs	Hypertension and heart failure
Losartan, potassium	LOSARTAN GP	50 mg 100 mg	Film coated tablet	14 tabs / 56 tabs	Hypertension and heart failure
Losartan, potassium + Hydrochlorothiazide	LOSARTAN / HCTZ GP	50 mg + 12.5 mg	Film coated tablet	14 tabs / 56 tabs	Hypertension
Ramipril	RAMIPRIL GP	1.25 mg 2.5 mg 5 mg 10mg	Capsule	14 tabs / 56 tabs 56 caps 56 caps 56 caps	Hypertension, heart and renal failure
Ticlopidine, hydrochloride	TICLOPIDINA GP	250 mg	Film coated tablet	10 tabs / 60 tabs	Anti-thrombotic



CENTRAL NERVOUS SYSTEM



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Beta-histine, hydrochloride	BETA-HISTINA GP	16 mg	Tablet	20 tabs / 60 tabs	Meniere's disease
Fluoxetine, hydrochloride	FLUOXETINA GP	20 mg	Capsule	14 tabs / 56 caps	Major depressive episodes, obsessive-compulsive disorder, bulimia nervosa
Paroxetine, hydrochloride	PAROXETINA GP	20 mg	Tablet	10 tabs / 60 tabs	Depression, anxiety disorders

GLUCOCORTICOID



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Deflazacort	DEFLAZACORTE GP	6 mg 30 mg	Tablet	20 / 60 tabs 20 tabs	Anti-inflammatory and immunosuppressant

WARTS TREATMENT



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Salicylic Acid	SALICYLIC ACID	167 mg/g	Solution	10 ml bottle	Warts and callosities treatment

CALCIUM FOLINATE



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Calcium Folate	MEDIFOLIN	15 mg	Capsule	20 caps	Antidote of folic acid antagonist, megaloblastic anaemia

ANTI-HISTAMINIC



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Cetirizine, dihydrochloride	CETIRIZINA GP	10 mg	Film coated tablet	20 tabs	Relief of nasal and ocular symptoms of seasonal and perennial allergic rhinitis, and of chronic idiopathic urticarial

ANTIDISLIPIDEMIC



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Simvastatin	SINVASTINA GP	10 mg 20 mg 40 mg	Film coated tablet	60 tabs 60 tabs 60 tabs	Hypercholesterolemia and cardiovascular prevention

BENIGN PROSTATIC HYPERTROPHY



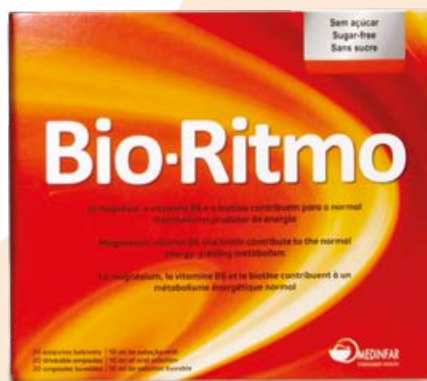
MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Finasteride	FINASTERIDA GP	5 mg	Film coated tablet	60 tabs	Treatment and control of benign prostatic hyperplasia



VITAMINS & MINERALS



MAIN INGREDIENTS	BRAND	DOSAGE FORM	PACK	INDICATION/USE
Magnesium, Manganese, Copper, Vitamin C, Vitamin B1, Vitamin B2, Vitamin B6	MAGNORANGE ACTIVE	Tablet	30 tabs	Normal energy yielding metabolism, reduction of tiredness and fatigue and normal functioning of the nervous system
Magnesium	MAGNORANGE SENIOR	Oral solution	20 x 10 ml ampoules	Normal energy yielding metabolism, reduction of tiredness and fatigue and maintenance of normal bones
Magnesium, Vitamin C, Ginkgo biloba extract, Vitamin B5, Vitamin B6, Vitamin B1	MAGNORANGE FOCUS	Tablet	60 tabs	Maintenance of memory and good cognitive function, normal physiological function, normal energy yielding metabolism
Echinacea Extract, Vitamin C, Vitamin B6, Zinc, Vitamin D	MAGNORANGE IMUNITAS	Tablet	60 tabs	Normal function of the immune system, normal energy yielding metabolism



VITAMINS & MINERALS



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Magnesium	MAGNORAL	1028.4 mg /10 ml	Oral solution	20 x 10 ml ampoules	Magnesium deficiency and prevention
Arginine Aspartate + Vitamin B6 + Biotin + Magnesium	BIO RITMO	-	Oral solution	20 x 10 ml ampoules	Normal energy yielding metabolism
Zinc	ZITREX	-	Capsules	60 caps	Protection of cells from oxidative stress and maintenance of normal skin, hair and nails

AMÉLIORE VOS PERFORMANCES



1 à 3 ampoules/ jour
au cours des principaux repas

SANS SUCRE

SANS ALCOOL



ASPARTATE
D'ARGININE

VITAMINE B6

BIOTINE

20 AMPOULES EN PLASTIQUE PVC
INCASSABLES
UN CONDITIONNEMENT INNOVANT

Bio-Ritmo®



Bio-ritmo est un complément alimentaire. Les compléments alimentaires ne doivent pas être utilisés comme substituts d'une alimentation variée. La vitamine B6, la biotine et le magnésium contribuent à un métabolisme énergétique normal ainsi qu'au fonctionnement normal du système nerveux et aussi à des fonctions psychologiques normales. La vitamine B6 et le magnésium contribuent à réduire la fatigue. La vitamine B6 contribue au fonctionnement normal du système immunitaire. La biotine contribue au maintien d'une peau et des cheveux normaux. Le magnésium contribue au maintien d'une assaturation et d'une dentition normale. Sujets jeunes de plus de 11 ans et les adultes: 1 à 3 ampoules par jour, au cours des principaux repas. Ne pas dépasser les doses journalières recommandées. Les colorants (E 102 et E 124) peuvent réagir de façon croisée avec l'acide acétylsalicylique. Bio-ritmo ne doit pas être pris en cas d'allergie à l'un des ingrédients. Bio-ritmo doit être pris avec précaution chez les patients présentant une insuffisance rénale sévère. Si vous êtes enceinte ou que vous allaitez, demandez l'avis de votre médecin. En cas de doute, consulter la personne responsable de sa mise sur le marché. MEDINFAR CONSUMER HEALTH - PRODUTOS FARMACÊUTICOS, LDA., Rua Henrique Paiva Couceiro, N° 27 Venda Nova, 2700-451 Amadora.



HAIR LOSS



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Minoxidil	TRICOVIVAX	20 mg/ml 50 mg/ml	Cutaneous solution	60 ml / 100 ml bottle (1, 2 and 3 bottles packs)	Androgenic and areata alopecia

PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
OTÉZIA SPH HAIR LOTION	Anti-hair loss lotion with an active peptide complex, significantly reduces hair loss and improves the scalp	100 ml spray	Biotin, Niacinamide, Hydrolyzed Keratin, Synthetic Peptides
TRIZATEK	Food supplement containing vitamins and minerals that contribute to healthy hair and nails	60 tablets / 180 tablets	L-Cystine, Keratine, Zinc,Iron, Selenium, Niacin, Pantothenic Acid, Vitamin B6, Biotin
TRIKARE C+ SHAMPOO ANTI-HAIR LOSS	Anti-hair loss shampoo formulated for hair loss of various origins, including androgenetic alopecia and thin or fragile hair; suitable for daily use and as an adjunct to anti-hair loss treatments	100 ml / 200 ml tube	Caffeine, Serenoa Repens Extract, Hydrolyzed Keratin



**PLAYING
HIDE AND SEEK
WITH YOUR HAIR?**

**WIN EVERY
ROUND!**



ESSENTIAL INFORMATION COMPATIBLE WITH THE SUMMARY PRODUCT CHARACTERISTICS

TRICOVIVAX, 20 mg/ml and 50 mg/ml, cutaneous solution. Qualitative and Quantitative Composition: Tricovivax 20 mg/ml: Each millilitre of cutaneous solution contains 20 mg of minoxidil. Each millilitre of cutaneous solution contains 259 mg of propylene glycol. Tricovivax 50 mg/ml: Each millilitre of cutaneous solution contains 50 mg of minoxidil. Each millilitre of topical solution contains 350 mg of propylene glycol. Complete list of excipients: Propilenoglicol, Etanol a 96%. Pharmaceutical form: Cutaneous Solution. Therapeutic indications: **TRICOVIVAX** is indicated in the treatment of androgenic alopecia and alopecia areata. Posology and method of administration: Tricovivax is exclusively intended for external use and should only be applied to the scalp. The hair and scalp should be thoroughly dry before topical application of Tricovivax. Do not apply to areas of the body other than the scalp. Tricovivax 20 mg/ml: Apply 1 ml of the 20 mg/ml solution, every 12 hours, pressing the spray pump 6 times. Apply the solution to the center of the affected area and spread it with your fingertips to ensure that the medicinal product is uniformly distributed. For a more localized application, use the applicator provided, which should be fitted onto the top part of the spray head. This dose (1 ml) should be administered regardless of the size of the affected area. The total daily dose should not exceed 2 ml. Tricovivax 50 mg/ml: The recommended dose is 1 ml of the 50 mg/ml solution, every 12 hours. Apply the solution to the center of the affected area and spread it with your fingertips to ensure that the medicinal product is uniformly distributed. This dose (1 ml) should be administered regardless of the size of the affected area. The total daily dose should not exceed 2 ml. Method of application: According to the affected area and the application device provided. Duration of treatment: Studies conducted to the present date indicate that 4 or more months of treatment with Tricovivax may be required for noticeable hair growth. Interruption of treatment may induce a return to the initial condition within 3 to 4 months. Contraindications: Hypersensitivity to the active substance or to any of the excipients. Undesirable effects: Tricovivax 20 mg/ml: Dermatological and sensitivity reactions: The most frequently observed undesirable effects associated with use of minoxidil cutaneous solution are local skin reactions, such as scalp pruritus (1.5% of patients), dry scalp and flaking. Other effects reported were local irritation or burning sensation, including irritative dermatitis. Reported cases of local irritation and allergic contact dermatitis were partly due to the alcohol or propylene glycol present in the formulation. Additionally, wearing of wigs or other types of hairpieces may contribute to local irritation. These effects were generally moderate and only led to treatment discontinuation in rare cases. Allergic reactions including angioedema: frequency not known. Other non-specific allergic and sensitivity reactions have also been reported, including urticaria, allergic rhinitis, facial oedema, eczema, greasy skin, papular rash, folliculitis, local erythema or redness, increased hair loss, alopecia, and hypertrichosis. Cardiovascular disorders: Although not directly attributable to treatment with minoxidil topical solution, cardiovascular effects have been reported, including oedema, precordial pain (usually transient or intermittent), palpitations, increased or decreased blood pressure and/or heart rate, and ECG changes. Nervous system disorders: Although also not attributable to minoxidil, adverse events involving the nervous system have been reported occasionally, including headache (sometimes with exacerbation of pre-existing migraine), dizziness, weakness, taste changes, syncope, "empty head" sensation and vertigo. Cases of anxiety, depression and fatigue have been reported rarely. Other undesirable effects: although also not directly related to topical treatment with minoxidil, have been reported rarely or occasionally, including musculoskeletal disorders (fractures, back pain, retrosternal pain of muscular origin, tendinitis), genitourinary disorders (urinary tract infections, kidney stones, urethritis, prostatitis, epididymitis, impotence), respiratory disorders (upper respiratory tract infections, transient dyspnoea, sinusitis), digestive disorders (diarrhoea, nausea, vomiting, weight increase), blood/immune system disorders (lymphadenopathy, thrombocytopenia), endocrine disorders (breast enlargement), and other disorders (ear infection, conjunctivitis, loss of visual acuity). Tricovivax 50 mg/ml: Tricovivax 50 mg/ml: Common ($\geq 1/100$, $<1/10$); Uncommon ($\geq 1/1\ 000$, $<1/100$); ($\geq 1/10\ 000$, $<1/1000$); Unknown frequency (cannot be estimated from the available data). Skin and subcutaneous tissue disorders: Common: local skin reactions, such as scalp pruritus, dry scalp and flaking. Unknown frequency: local irritation or burning sensation, including irritative dermatitis, eczema, greasy skin, papular rash, folliculitis, local erythema or redness, increased hair loss, alopecia, and hypertrichosis. Reported cases of local irritation and allergic contact dermatitis were partly due to the ethanol or propylene glycol present in the formulation. Additionally, wearing of wigs or other types of hairpieces may contribute to local irritation. Immune system disorders: Unknown frequency: urticaria, allergic rhinitis, facial oedema, allergic reactions including angioedema. The adverse effects listed are not directly attributed to treatment with minoxidil cutaneous solution. Cardiac disorders: Unknown frequency: oedema, precordial pain (usually transient or intermittent), palpitations, increased or decreased blood pressure and/or heart rate, and ECG changes. Nervous system disorders: Uncommon: headache (sometimes with exacerbation of pre-existing migraine), dizziness, weakness, taste changes, syncope, "empty head" sensation and vertigo. Rare: Anxiety, depression, and fatigue. Musculoskeletal and connective tissue disorders: Rare or uncommon: fractures, back pain, retrosternal pain of muscular origin, tendinitis. Genital organs and breast disorders: Rare or uncommon: Prostatitis, epididymitis, and impotence. Renal and urinary disorders: Rare or uncommon: urinary tract infections, kidney stones and urethritis. Respiratory, thoracic and mediastinal disorders: Rare or uncommon: upper respiratory tract infections, transient dyspnoea and sinusitis. Gastrointestinal disorders: Rare or uncommon: diarrhoea, nausea, vomiting, weight increase. Blood and lymphatic system disorders: Rare or uncommon: lymphadenopathy and thrombocytopenia. Endocrine disorders: Rare or uncommon: breast enlargement. Ear and labyrinth disorders: Rare or uncommon: ear infections. Eye disorders: Rare or uncommon: loss of visual acuity and conjunctivitis. These medicinal products are not subject to medical prescription. For more information, please contact the Marketing Authorization Holder. Laboratório Medinfar – Produtos Farmacêuticos, S.A., Rua Henrique de Paiva Couceiro, 29, Venda Nova 2700-451 Amadora. NIF 500 384 045. (07/2019)

HAIR CARE



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
PROVIVAX AHA SHAMPOO	Daily use shampoo for the hygiene and care of devitalised, thin, and brittle hair, ensures effective scalp cleansing combined with repairing, nourishing, and volumizing action	200 ml tube	Panthenol, Hydrolyzed Keratin, Glycolic Acid
PROVIVAX N SHAMPOO	Daily use shampoo for dry, delicate, and weakened hair, helps to soften and protect and enhances hair luminosity	200 ml tube 400 ml bottle	Hydrolyzed Keratin, Biotin, Hamamelis Virginiana Leaf Extract



ANTI- -SEBORRHEIC



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
KETOPAN SOAP	Soap for the hygiene of oily-prone sensitive skin and control and prevention of seborrheic conditions in their multiple forms	100 g soap	Ictasol, Ceramide, Piroctone Olamine
KETOPAN FOAM	Foam for the hygiene of oily-prone sensitive skin and control and prevention of seborrheic states in their multiple forms	200 ml bottle	Piroctone Olamine, Zinc PCA
TRIKARE K SHAMPOO	Anti-seborrheic shampoo recommended for the cleansing and control of scalp irritation and scaling (seborrhoea)	200 ml tube	Glycolic Acid, Panthenol, Piroctone Olamine, Zinc PCA, Climbazole
TRIKARE DS SHAMPOO	Anti-seborrheic and anti-dandruff shampoo for the hygiene of seborrheic and irritated scalp, helps to reduce dry and oily scaling through antimicrobial, purifying, hydrating, and calming action	200 ml tube	Panthenol, Zinc Lactate, Piroctone Olamine



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Minocycline	MINOTREX	100 mg	Capsule	16 caps / 32 caps	Skin infections (acne) and other infections

PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
NIACIDE CREAM	Coadjuvant in the treatment of acne, developed for dry and sensitive skin undergoing retinoid or systemic antibiotic therapy, ensures hydration and repair, and helps to reduce skin irritation frequency	50 g tube	Niacinamide, Glycolic Acid, Salicylic Acid
NIACIDE ACNE GEL	Coadjuvant for acne treatment in oily skin, promoting the repair and correction of oil-prone areas	50 g tube	Niacinamide, Glycolic Acid, Salicylic Acid
NIACIDE DERMO-NETTOYANT	Daily cleansing of oily and acne-prone skin	200 ml tube	Salicylic Acid, Glycolic Acid, Zinc
NIACIDE ECLAIRTIN SOIN ANTI-TÂCHES	Anti-dark spot cream specially developed to reduce post-inflammatory hyperpigmentation	30 ml	Alpha-Arbutin,Kojic Dipalmitate



CICATRIZATION



MAIN INGREDIENTS	BRAND	STRENGTH	DOSAGE FORM	PACK	INDICATION/USE
Zinc Oxide	HALIBUT	150 mg/g	Ointment	30 g / 50 g/ 100 g tube	Superficial wound healing, nappy rash treatment and first degree burns

PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
HALIBUT DERMA PLUS CREAM	Repairs the skin exposed to aggressions due to excessive exposure to sunlight, fragile skin and skin discomforts	30 g tube	Glycerin, Panthenol, Zinc Oxide, Chlorhexidine Digluconate



DRY & PSORIATIC SKIN



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
XERATOP X REPAIR EMOLLIENT LOTION	Repairing lotion for dry and xerotic skin, developed to restore the skin's physiological balance and hydration by replenishing the lipid film, while providing moisturising, nourishing, and keratolytic action	400 ml bottle	Glycerin , Ammonium Lactate, Panthenol, Ceramides
XERATOP X REPAIR SHOWER GEL	Shower gel recommended for the hygiene of xerotic skin, promotes hydrolipid film restoration, comfort, and balance through physiological, moisturising, nutritive, and keratolytic agents	400 ml bottle	Paraffinum Liquidum, Ammonium Lactate, Panthenol, Cholesterol, Lactic Acid, Ceramides
PSORIL MEDINFAR LOTION	Body lotion with moisturising, nutritive and keratolytic ingredients, formulated for dry, scaly, psoriatic skin; helps to hydrate, reduce scaling, soothe redness, and relieve pruritus	200 ml tube	Sodium Lactate, Niacinamide, Ceramides, Aloe Barbadensis Leaf Juice, Panthenol

HALIBUT
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Alongside your family ...
from day one.



CLEAN

PROTECTION

REPAIR
IN CASE OF
DIAPER RASH

Complete care on diaper change



BABY CARE

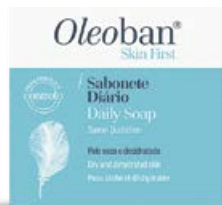


PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
HALIBUT MUDA FRALDAS LINIMENT	Gentle no-rinse cleanser for nappy changes	200 ml bottle	Helianthus Annuus Seed Oil, Glycerin, Cera Alba, Linum Usitatissimum Seed Oil
HALIBUT MUDA FRALDAS PROTECTING CREAM	Frequent use cream for nappy rash prevention	50 g/ 100 g / 150g tube	Helianthus Annuus Seed Oil, Triticum Vulgare Bran Extract, Talc, Zinc Oxide
HALIBUT MUDA FRALDAS REGENERATING OINTMENT	Regenerating ointment with 45% zinc oxide, indicated for protection and recovery of the baby's bottom skin	50g tube	Zinc Oxide
HALIBUT MUDA FRALDAS REPAIRING OINTMENT	Ointment for skin repair in cases of irritation, rash or redness in the nappy area, with miconazole to help controlling the proliferation of microorganisms	50g tube	Zinc Oxide and Miconazole

BABY CARE



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
OLEOBAN BABY CREAM	Moisturising and emollient cream for daily care of babies and newborns sensitive's skin, helps to restore hydration, reduce skin irritation and provide comfort	80g tube/ 200g tube / 450 ml bottle	Glycerine, Paraffinum Liquidum
OLEOBAN BABY BATH	Moisturising and emollient bath solution for the hygiene of babies and newborns sensitive's skin, with sunflower oil to restore the hydrolipidic film and reduces the frequency of skin irritations	300 ml / 500 ml bottle	Helianthus Annuus Seed Oil
OLEOBAN BABY BATH FOAM	2-in-1 bath foam for body and hair, ideal for gentle cleansing of baby's skin and scalp from birth, with emollient agents compatible with sensitive skin	500 ml bottle	Hydrolyzed soy protein, Glycerin
OLEOBAN BABY SOAP	Solid soap indicated for the hygiene and hydration of babies' and newborns' sensitive skin, with sunflower oil	100 g soap	Helianthus Annuus Seed Oil



BATH & HYDRATION



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
OLEOBAN DAILY CREAM	Moisturising and emollient cream that strengthens the skin barrier and helps reducing the frequency of skin dehydration	80g tube/200 g tube/ 450 g bottle/1kg bottle	Glycerin, Paraffinum Liquidum
OLEOBAN DAILY SHOWER	Moisturising and emollient shower gel for daily hygiene, formulated with oil-based cleansing agents	300ml / 500 ml bottle	Ricinus Communis Seed Oil, Panthenol, Helianthus Annuus Seed Oil
OLEOBAN SOAP	Moisturising and emollient soap for daily hygiene of dry skin, suitable for frequent and regular use	100 g soap	Helianthus Annuus Seed Oil
OLEOBAN CLEANSING CREAM	Cleansing cream for dry and sensitive skin, formulated with mild agents that respect the skin's physiological pH, combining oat extract and vegetable glycerin with calming, emollient and moisturising properties	450ml / 950 ml bottle	Oat extract, Vegetable glycerin
OLEOBAN SHOWER GEL	Shower gel with panthenol, featuring a non-oily texture suitable for daily hygiene of dry and sensitive skin	450ml / 950 ml bottle	Panthenol, Vegetable Glycerin
OLEOBAN RICH CREAM	Intensive moisturising cream for very dry skin areas such as heels, elbows, knees and feet, improves skin elasticity and helps to reduce stretch marks	170 ml jar	Butyrospermum Parkii Butter, Squalene, Hydrolyzed Linseed Extract

BATH & HYDRATION



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
OLEOBAN PLUS DERMA SHOWER OIL	Moisturising and emollient shower oil for dry and scaly skin, with antimicrobial action; adjuvant in the treatment of skin conditions such as eczema	300 ml bottle	Helianthus Annuus Seed Oil, Ictasol
OLEOBAN ANTI-ICH CREAM	Hydration of dry, sensitive, scaly and itchy skin, with polidocanol for soothing action	100 g tube	Polidocanol, Shea Butter, Argania Spinosa Oil

ANIMAL HEALTH



DERMA PRODUCTS



PRODUCT	INDICATION	PACK	MAIN INGREDIENTS
ALERDERMA ESPUMA	Shampoo adjuvant in the treatment of pruritus and cutaneous inhalant allergies in animals with allergic or atopic dermatitis	200 ml bottle	Phytosphingosine Salicyloyl
VETDERMA SHAMPOO	Shampoo with 10% urea adjuvant in the treatment of dry, peeling and itching skin	200 ml bottle	Urea



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