Pascual Laboratories, Inc.

Manufacturing and Product Information Catalogue

Dear Valued Customer,

Since 1946, the PascualLab Manufacturing Plant has been committed to building strong, healthy families through the production of safe, quality, and innovative healthcare products.

But more than just a production facility, we are also a professional and passionate family that is driven by the purpose of enriching quality of life in every community we touch. This guiding mission was passed on to my generation by my grandparents and PascualLab's founders, Leonora and Isosceles, and my father, Abraham, and is now deeply embedded in our manufacturing and product development processes. It is also reflected in the wide spectrum of family-friendly health and wellness products that we manufacture.

As one of the biggest production facilities in the Philippines processing both herbal and pharmaceutical products, you can be guaranteed that our efforts are directed towards ensuring that we provide safe and quality solutions for our customers. We are certified and licensed by the Food and Drug Administration of the Philippines (FDA) for Good Manufacturing Practices (GMP), and in adherence to Quality Management System standards, the PascualLab Manufacturing Plant received its ISO 9001:2015 Certification and HALAL Certificate from Islamic Da'Wah Council of the Philippines (IDCP).

It is with gratitude for the unwavering trust of our clients and consumers throughout our more than 70 years in the business that we continuously improve our facilities and methods to exceed and set industry standards both locally and abroad. We research and engineer on breakthrough yet purposefully innovative products with your family in mind.

We share your passion for health and life, and we offer you quality, innovative, and effective solutions here at PascualLab. Thank you for your partnership and support.

Mr. Jose Augusto Pascual
President and CEO of Pascual Laboratories

Dear Customers,

At the PascualLab Manufacturing Plant, our commitment goes beyond the delivery of excellent products and services. We build lasting relationships with our clients while helping their businesses progress forward and attain their goals.

More than the awards and recognitions, we find value in knowing that we have provided you with the best and most cost-efficient solutions.

More than a reliable manpower, we are a professional family that understands your needs and the greater purpose that our products and output serve, to the well-being of your consumers.

More than just accreditations, we work with government agencies and certifying bodies to help improve on the technology and industry standards available today, as we uplift local practices and the quality of products available in the market.

More than innovation, we formulate our products purposefully, with relevant health issues in mind.

You can trust us to be responsive to your product and manufacturing requirements, and deliver it in the caring and professional manner that only PascualLab can.

Engr. Jun Porte, Jr.
Vice President
Product Supply Group

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Pascual Laboratories, Inc.

From Our Family to Yours, Since 1946

The PascualLab Manufacturing Plant is a subsidiary of Pascual Laboratories, Inc. (PascualLab), a leading pharmaceutical company based in the Philippines. We are licensed by the Philippine Food and Drug Administration and certified for Good Manufacturing Practices (GMP). We are also an ISO 9001:2015 Certified facility, adhering to international Quality Management System standards and HALAL certified by the Islamic Da'Wah Council of the Philippines (IDCP).

The PascualLab Manufacturing Plant began as an initial venture of our founders Isosceles and Leonora Pascual in 1946, following the end of World War II. The married couple were both Chemistry graduates of the University of the Philippines. Their early operations were confined to a single-room laboratory at their home in Balut Island, Tondo, Manila, with focus on the formulation and manufacturing of medicines to cure tuberculosis and vitamin deficiency.

Pascual Laboratories, Inc.

PascualLab grew into a large-scale manufacturing facility in the 1950s, and became a full-blown corporation by the '60s. Our manufacturing plant was moved to its present-day location in Balagtas, Bulacan in the '70s, while our corporate headquarters transferred to Quezon City.

The '80s and '90s saw our company progressing into the new millennium as a formidable player, led by our Chairman, 2nd generation Pascual Family member, Dr. Abraham F. Pascual. Propelled by his visionary leadership, PascualLab experienced significant growth through the introduction of pioneering and innovative products, time-tested formulas, and manufacturing expertise in producing high quality, safe and family-friendly healthcare solutions.

Today, under the helm of 3rd generation Pascual Family member Jose Augusto G. Pascual, the PascualLab Manufacturing Plant is one of the largest

Pascual Laboratories, Inc.

production facilities in the Philippines, processing both pharmaceutical, herbal and consumer products in solid, semi-solid, and liquid forms. We supply the product requirements of our wholly-owned subsidiary, Pascual Consumer Healthcare Corp. (PCHC), and provide toll manufacturing services to local and off-shore pharmaceutical companies.

We are dedicated to and passionate about providing safe and quality solutions for our customers while continuously elevating plant quality controls, equipment, and production capabilities to comply with global industry standards. As quality standards are implemented at every stage, from formulation to manufacturing, our healthcare products are not just meant for cure, but feature purposeful innovation, with the goal of allowing families to live a better life, together.

From our family to yours, this is our #GalingPascual story.

Manufacturing

Our Strong Manufacturing Heritage

The PascualLab Manufacturing Plant operates across 5.2 hectares of land in Balagtas, Bulacan, and features modern machineries and technology that we utilize in the formulation and production of natural and ethical health and wellness products, in solid, semi-solid, and liquid forms. A separate facility also caters to the production of steroids and cosmetics.

With over seven decades of manufacturing experience and research, our company continues to optimize its facilities, increase plant capabilities, and upgrade our manufacturing practices, as we strive to offer top-notch products and services to our clients and consumers.

Quality is the Heart of our Business

We proudly manufacture our products following the stringent Good Manufacturing Practices (GMP) for pharmaceutical and herbal products, based on PIC/S requirements.

We have embedded the principles of GMP in every aspect of our manufacturing operations through standard operating procedures, in the design and construction of facilities and the end-to-end processes of our supply chain - from material sourcing up to finished goods release. This is done to consistently ensure that we manufacture our products according to their registered specifications and quality attributes.

Likewise, we maintain a methodical and tedious process in assuring that all products manufactured and

Quality is the Heart of our Business

released by our Manufacturing Plant are of quality, safe, and effective:

- Raw materials are sampled and inspected during every delivery and subjected to incoming materials quality control testing prior to release for production use.
- Pertinent information during manufacturing and packaging are collected and properly documented into batch processing records.
- Batch processing is performed according to approved manufacturing and packaging instructions.
- Precise manufacturing controls are maintained throughout the production process. We conduct

Quality is the Heart of our Business

routine in-process sampling and testing as basis for release of the product to succeeding stages of manufacturing.

- Finished product release is only initiated once the quality control tests, batch records review, and statistical inspection on the finished product have been found satisfactory, as reflected in the Certificate of Analysis and the GMP Certificate issued by Quality.

The Quality System fostered by PascualLab is continuously improved to tide changes in industry standards, technology, and customer feedback. All are done and led by our highly-skilled and qualified personnel with utmost care and vigilance.

Product Development

We partner with multinational and local pharmaceutical companies through technology transfer process and innovate on established product technologies according to the capabilities and facilities available in the local market, while still ensuring that the quality requirements of the products are complied with. This product development approach offers our subsidiaries and clients value-adding benefits compared to importation of finished goods.

A Professional Family Committed to Our Partners

We are a fast-growing family of over 200 highly-trained workers with diverse manufacturing experience. It is our management's top priority that they are provided with an optimal and safe working environment, and a pleasant professional and family-oriented working atmosphere. With a strong frontline of pharmacists, chemists, engineers and healthcare professionals, we continue to provide quality and safe products that address the needs of our clients and consumers.

Our professional and collaborative management team strives to delight customers with the care and efficiency with which we manage and respond to their needs. For over 70 years, PascualLab has been a trusted partner in pharmaceutical manufacturing.

Products

Vitex negundo L.
Lagundi Leaf
ASCOF® Forte

If symptoms persist, consult your doctor.

FORMULATION

Each Ascof tablet contains:
Vitex negundo L. (Powdered Lagundi) Leaves ..... 300 mg
Each Ascof Forte tablet contains:
Vitex negundo L. (Powdered Lagundi) Leaves ..... 600 mg
Each Ascof Forte capsule contains:
Vitex negundo L. (Powdered Lagundi) Leaves ..... 600 mg

PHARMACOLOGIC CATEGORY

Herbal Medicine (Cough Remedy/Anti-Asthma)

INDICATION

300 mg and 600 mg Tablet:

For the relief of mild to moderate cough due to common colds, flu and mild to moderate acute bronchitis; for relief of reversible mild to moderate bronchospasm in adults and children 2 years of age and older with obstructive airway disease such as asthma and chronic bronchitis.

600 mg Capsule:

For the relief of mild to moderate cough due to common colds and flu. For the treatment of bronchospasm in acute bronchial asthma, chronic bronchitis and other bronchopulmonary disorders. For relief of reversible, mild to moderate bronchospasm (prophylactic/maintenance medication) in adults and children with obstructive airway disease.

DOSAGE AND ADMINISTRATION

Ascof 300 mg Tablet

Adults:
2 tablets 3-4 times a day.
Children (7 to 12 years old):
1 tablet 3-4 times a day
Or as prescribed by the physician.

Ascof Forte 600 mg Tablet

Adults:
1 tablet 3 -4 times a day.
Children (13 years old and above):
1 tablet 3 times a day
Children (2 to 12 years old):
As prescribed by the physician.

HOME REMEDY

Vitex negundo L.
Lagundi Leaf

ASCOF® Forte

Ascof Forte 600 mg Capsule
Adults:
1 capsule 3-4 times a day.
Children 7-12 years old: 1/2 tablet 3-4 times a day.
Or as prescribed by the physician.

AVAILABILITY

Ascof 300 mg Tablet
Box of 100's in Foil strip x 10's
Ascof Forte 600 mg Tablet
Box of 12's in aluminium foil strip x 4's
Box of 60's in aluminium foil strip x 6's
Box of 120's in aluminium foil strip x 6's
Ascof Forte 600 mg Capsule
Box of 60's in blister pack x 10's
Box of 120's in blister pack x 10's

STORAGE CONDITIONS

Store at temperatures not exceeding 30°C

SHELF-LIFE

300 mg Tablet – 48 months
600 mg Tablet – 60 months
600 mg Capsule – 24 months

LEADTIME

90 days

REGISTRATION NO.

Ascof 300 mg Tablet – HMR-18
Ascof Forte 600 mg Tablet – HMR-01
Ascof Forte 600 mg Capsule – HMR-31

If symptoms persist, consult your doctor.

HOME REMEDY

Vitex negundo L.
Lagundi Leaf

ASCOF® Forte

FORMULATION

Each 5 mL (teaspoonful) of Ascof Forte (Menthol Flavor) syrup contains
the extract from *Vitex negundo* L. (Dried Lagundi leaves) ..... 600 mg
Each 5 mL (teaspoonful) of Ascof Forte (Menthol & Sugar Free Flavor) syrup contains
the extract from *Vitex negundo* L. (Dried Lagundi leaves) ..... 600 mg

PHARMACOLOGIC CATEGORY

Herbal Medicine (Cough Remedy/Anti-Asthma)

INDICATION

For the relief of mild to moderate cough due to common colds, flu and mild to moderate acute bronchitis; for relief of reversible mild to moderate bronchospasm in adults and children 2 years of age and older with obstructive airway disease such as asthma and chronic bronchitis.

DOSAGE AND ADMINISTRATION

Ascof Forte Syrup (Menthol) (Menthol & Sugar Free) (Bottle)
Adults:
5 mL (1 teaspoon) 3 to 4 times a day. Dose may be increased as needed.
Children (13 years old and above):
5 mL (1 teaspoon) 3 times a day.
Children (2 to 12 years old):
As prescribed by the physician.
Ascof Forte Syrup (Menthol) (Menthol & Sugar Free) (Sachet)
Adults: 5 mL (1 sachet) 3 to 4 times a day. Dose may be increased as needed, per physician's advice.
Children (13 years old and above):
5 mL (1 sachet) 3 times a day. Dose may be increased as needed, per physician's advice.
Chidren (2 years to 12 years old):
As prescribed by the physician.

If symptoms persist, consult your doctor.

HOME REMEDY

Vitex negundo L.
Lagundi Leaf
ASCOF® Forte

AVAILABILITY
Ascof Forte Syrup (Menthol) (Menthol & Sugar Free) (Bottle)
Amber Glass bottle x 60 mL and 120 mL
Ascof Forte Syrup (Menthol) (Menthol & Sugar Free) (Sachet)
Box of 60's in PET Foil sachet x 5 mL

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C

SHELF-LIFE
Ascof Forte Syrup (Menthol) (Bottle) – 36 months
Ascof Forte Syrup (Menthol & Sugar Free) (Bottle) – 24 months
Ascof Forte Syrup (Menthol) (Menthol & Sugar Free) (Sachet) – 24 months

LEADTIME
90 days

REGISTRATION NO.
Ascof Forte Syrup (Menthol) (Bottle) – HMR-19
Ascof Forte Syrup (Menthol & Sugar Free) (Bottle) – HMR-32
Ascof Forte Syrup (Menthol) (Sachet) – HMR-85
Ascof Forte Syrup (Menthol & Sugar Free) (Sachet) – HMR-83

If symptoms persist, consult your doctor.

HOME REMEDY

Vitex negundo L.
Lagundi Leaf
ASCOF®

FORMULATION

Each 5 mL (teaspoonful) of Ascof (Strawberry Flavor) syrup contains
the extract from *Vitex negundo* L. (Dried Lagundi leaves) ..... 300 mg
Each 5 mL (teaspoonful) of Ascof (Ponkan Flavor) syrup contains
the extract from *Vitex negundo* L. (Dried Lagundi leaves) ..... 300 mg

PHARMACOLOGIC CATEGORY

Herbal Medicine (Cough Remedy)

INDICATION

For the relief of mild to moderate cough due to common colds, flu and mild to moderate acute bronchitis; for relief of reversible mild to moderate bronchospasm in adults and children 2 years of age and older with obstructive airway disease such as asthma and chronic bronchitis.

DOSAGE AND ADMINISTRATION

Ascof Syrup (Strawberry and Ponkan) (Bottle and Sachet)

Adults:
10 mL (2 teaspoonfuls) 3 to 4 times a day. Dose may be increased as needed, per physician's advice.
Children:
15 mg/kg/dose (or 0.25 mL/kg/dose) administered 3 times a day.
2-4 years old (10 to 15.5 kg): ½ to 1 teaspoonful 3 times a day.
4-6 years old (15.5 to 20 kg): 1 teaspoonful 3 times a day.
6-12 years old (20-40 kg): 1 ½ - 2 teaspoonfuls 3 times a day.
13 years old and above (over 40 kg): 2 teaspoonfuls 3 times a day.
Or as prescribed by the physician.

If symptoms persist, consult your doctor.

HOME REMEDY

Vitex negundo L.
Lagundi Leaf

ASCOF®

AVAILABILITY
Ascof Forte Syrup (Strawberry and Ponkan) (Bottle)
Amber bottle with measuring cup x 60 mL and 120 mL
Ascof Forte Syrup (Strawberry and Ponkan) (Sachet)
Box of 60's in PET Foil sachet x 10 mL

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C

SHELF-LIFE
Ascof Syrup (Strawberry) (Bottle) – 24 months
Ascof Syrup (Ponkan) (Bottle) – 36 months
Ascof Syrup (Strawberry and Ponkan) (Sachet) – 24 months

LEADTIME
90 days

REGISTRATION NO.
Ascof Syrup (Strawberryl) (Bottle) – HMR-29
Ascof Syrup (Ponkan) (Bottle) – HMR-14
Ascof Syrup (Strawberry) (Sachet) – HMR-82
Ascof Syrup (Ponkan) (Sachet) – HMR-84

If symptoms persist, consult your doctor.

HOME REMEDY

POVIDONE-IODINE

ALFADINE

75 mg/mL (7.5% w/v)
Cleansing Solution
ANTISEPTIC

If symptoms persist, consult your doctor.

FORMULATION
Each mL contains
Povidone-Iodine ..... 75 mg
(equivalent to available iodine ..... 7.5 mg

CLASSIFICATION
Antiseptic

INDICATION(S)
As a broad-spectrum bacteriocide, also effective against molds, viruses and protozoans. May be used for the treatment and/or prevention of infections in wounds (accidental, surgical), cuts, burns and ulcers; as a surgical scrub

DOSAGE AND ADMINISTRATION
Pour a small amount of Povidone-Iodine (ALFADINE) Skin Cleanser on the palm (equivalent to 1/3 of the bottle cap) and dilute with a palm-full of water. Work into a lather on the skin like liquid soap for 60 seconds. Rinse thoroughly with water.

For body odor, use regularly until odor is removed.

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C.

AVAILABILITY
60mL, 120mL

SHELF LIFE
24 mos.

LEADTIME
90 days

REGISTRATION NO.
DRHR-1374

HOME REMEDY

POVIDONE-IODINE

ALFAFRESH

7.5% Cleansing Solution
ANTISEPTIC
FEMININE WASH

If symptoms persist, consult your doctor.

FORMULATION
Each mL contains
Povidone-Iodine ..... 75 mg
(equivalent to available iodine ..... 7.5 mg)

CLASSIFICATION
Antiseptic

INDICATION(S)
As a broad-spectrum bacteriocide, also effective against yeast, protozoans and fungi. May be used for the treatment of vaginitis secondary to *Candida*, *Trichomonas* or mixed-type infections.

DOSAGE AND ADMINISTRATION
Pour a small amount of Povidone-Iodine (ALFAFRESH) Gyne Solution on the palm (equivalent to 1/3 of bottle cap) and dilute with a palm-full of water. Wash on the external genitals for 60 seconds or less. If to be used as a regular wash for protection, use only twice a week. Rinse thoroughly with water.

For prompt relief of external genital itching, dilute similarly as for protection against infection.

'For red days, use in the morning and evening. Rinse thoroughly with water after use.'

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C.

AVAILABILITY
25mL, 60mL, 120mL

SHELF LIFE
24 mos.

LEADTIME
90 days

REGISTRATION NO.
DRHR-1375

HOME REMEDY

POVIDONE-IODINE

ALFADINE

100 mg/mL (10% w/v)
Topical Solution
ANTISEPTIC

If symptoms persist, consult your doctor.

FORMULATION
Each mL contains
Povidone-Iodine ..... 100 mg
(equivalent to available iodine ..... 10 mg)

CLASSIFICATION
Antiseptic

INDICATION(S)
Disinfectant and antiseptic mainly for the treatment of contaminated wounds and pre-operative preparation of the skin and the mucous membranes

DOSAGE AND ADMINISTRATION
Wash clean with mild soap and dry the affected skin by dabbing with sterile cotton/tissue. Soak sterile cotton balls with Povidone-Iodine (ALFADINE) solution. Apply generously on affected areas in a circular manner starting from center. Bandage with non-stick gauze for larger or deep wounds. Superficial wounds are best left exposed. Re-apply 3 to 4 times a day.

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C.

AVAILABILITY
10mL, 20mL, 60mL, 120mL

SHELF LIFE
24 mos.

LEADTIME
90 days

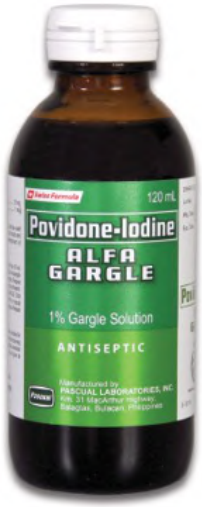
REGISTRATION NO.
DRHR-1373

HOME REMEDY

POVIDONE-IODINE

ALFAGARGLE

1% Gargle Solution
ORAL ANTISEPTIC



If symptoms persist, consult your doctor.

FORMULATION
Each mL contains
Povidone-Iodine ..... 10 mg
(equivalent to available iodine ..... 1 mg)

CLASSIFICATION
Antiseptic

INDICATION(S)
As a broad-spectrum bacteriocide, may be used for the treatment and/or prevention of mouth and throat sores. May reduce the development of bad breath.

DOSAGE AND ADMINISTRATION
For sore throat and mouth infections: Pour 20 mL (4 teaspoons) of Povidone-Iodine (ALFAGARGLE) into a glass and gargle for 30 seconds. Repeat 2-4 times daily or as required. As mouthwash: Use 20 mL (4 teaspoons) of ALFAGARGLE full strength or diluted with an equal amount of water. Gargle for 30 seconds. Repeat as desired.

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C.

AVAILABILITY
60mL, 120mL, 240mL

SHELF LIFE
36 months

LEADTIME
90 days

REGISTRATION NO.
DRHR-1372

HOME REMEDY

LYSOZYME hydrochloride
DEQUALINIUM chloride

QUADEZYME

FORMULATION

Each tablet contains:

Lysozyme hydrochloride	20 mg
Dequalinium chloride, BP	250 mcg

INDICATION(S)

For the relief of productive and non-productive cough and nasal congestion associated with common colds and allergic conditions.

DOSAGE AND ADMINISTRATION

Adults and children over 6 years old:
One lozenge to be sucked slowly every 2 hours as needed

Children over 6 years:
Maximum 3 tablets daily.

Adults:
Maximum 6 tablets daily.

or as prescribed by the physician.

STORAGE CONDITIONS

Store at temperatures not exceeding 30°C.

AVAILABILITY

Box of 20's in foil strip x 4's
Box of 500's in foil strip x 10's

SHELF LIFE

36 months

LEADTIME

90 days

REGISTRATION NO.

DRHR-566

If symptoms persist, consult your doctor.

HOME REMEDY

Blumea balsamifera L. (Sambong Leaf)

FORMULATION
Each tablet contains:
Blumea balsamifera (L.) Sambong leaves ..... 500 mg

CLASSIFICATION
Anti-urolithiasis

DOSAGE AND ADMINISTRATION
Diuretic: 1 tablet 3 times a day.

Anti-urolithiasis: 2 tablets 3 times a day.
Or as prescribed by the physician.

STORAGE CONDITIONS
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box x 60's

SHELF LIFE
24 mos.

LEADTIME
90 days

If symptoms persist, consult your doctor.

HOME REMEDY

ASCORBIC ACID
POTEN-CEE®

FORMULATION

Each Oral Drops contains:
Ascorbic acid (Vitamin C) 100mg/mL
Each Syrup contains:
Ascorbic acid (Vitamin C) ..... 100 mg/5 mL
Each Sugar Coated Tablet contains:
Ascorbic acid (Vitamin C) ..... 500 mg
Each Film-Coated Tablet contains:
Ascorbic acid (Vitamin C) ..... 500 mg
Each Forte tablet contains:
Ascorbic acid (Vitamin C) 1000mg

PHARMACOLOGIC CATEGORY

Vitamin

INDICATION(S)

For the prevention and treatment of Vitamin C deficiency.

DOSAGE AND ADMINISTRATION

Oral Drops:
Below 3 months 0.1 to 0.3ml (3-9 drops)
9-21 months 0.3 to 0.7 mL (9-12 drops)
To be taken once a day or as prescribed by the physician.

Syrup:
1 to 2 years 2.5 ml (1/2 tsp.)
3 to 6 years 5ml (1 tsp)
7 to 12 years 10 ml (2 tsp)
To be taken once a day or as prescibed by the physician.

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

ASCORBIC ACID POTEN-CEE®

Sugar Coated Tablet and Film Coated Tablet:
Take 1 tablet daily. For increased resistance against colds and flu: Take 2-3 tablets daily.
Forte:
1 tab 1x a day for forte or as recommended by your physician

STORAGE
Store at temperatures not exceeding 30°C

AVAILABILITY
Oral Drops:
Amber glass bottle x 30 mL box of 1's

Syrup:
Amber glass bottle 60ml box of 1's
Amber glass bottle 120ml box of 1's
Amber glass bottle 250ml box of 1's

Sugar Coated Tablet:
Amber blister pack x 10's Box of 20's
Amber blister pack x 10's Box of 100's
Amber blister pack x 10's Box of 500's

Film Coated Tablet:
Alu/orange PVC blister Pack x 10's Box of 100's

Forte Tablet:

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

ASCORBIC ACID
POTEN-CEE®

Aluminium Foil Strip x 6's Box of 30's
Aluminium Foil Strip x 4's Box of 100's

SHELF LIFE
Drops and Syrup Tablets: 24 months
Sugar Coated Tablets: 36 months
Film Coated: 48 months
Forte Tablets: 24 months

LEADTIME
90 days

REGISTRATION NO.
Oral Drops - DR-X5295
Syrup - DR-X3754
Sugar Coated Tablets - DR-9468
Film Coated Tablets - DRHR-1014
Forte Tablets - DR-XY32613

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

Phenylpropanolamine Hydrochloride +
Paracetamol +
Chlorphenamine Maleate +
Salicylamide

FORMULATION
Each tablet contains:

Phenylpropanolamine Hydrochloride	25mg
Paracetamol	325 mg
Chlorphenamine Maleate	2mg
Salicylamide	50 mcg

CLASSIFICATION
Cough and Cold Remedy

DOSAGE AND MODE OF ADMINISTRATION
Adult: 1 tablet 3-4 times a day.
Children: 1/2 tablet 3-4 times a day.
Children below 6 years: according to physician.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Plastic Bottle x 1000's

SHELF LIFE
60 months

LEADTIME
90 days

REGISTRATION NO.
DR-XY35895

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

Thiamine Hydrochloride (Vit. B_1) + Pyridoxine Hydrochloride (Vit. B_6) + Cyanocobalamin (Vit. B_{12}) (VITAMIN B_1+B_6+B_{12})

FORMULATION
Each tablet contains:

Thiamine HCl (Vitamin B1)	100 mg
Pyridoxine HCl (Vitamin B6)	5 mg
Cyanocobalamin (Vitamin B12)	50 mcg

PHARMACOLOGIC CATEGORY
Vitamin B Complex

INDICATION(S)
This product contains B-complex vitamins. These nutrients are required for normal nerve function and are used as adjunct in the management of neuromuscular pain.

DOSAGE AND ADMINISTRATION
1-2 tablets daily.
Or as prescribed by the physician.

STORAGE
Store at temperatures not exceeding 30°C.
Keep all medicines out of reach of children.

AVAILABILITY
Box of 100's in strip foil x 10's
Box of 500's in strip foil x 10's

SHELF LIFE
36 months

LEADTIME
90 days

REGISTRATION NO.
DR-XY27819

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

Dextromethorphan HBr +
Guaifenesin +
Phenylephrine HCl +
Chlorpheniramine Maleate

Dexeryl

FORMULATION
Each tablet contains:
Dextromethorphan hydrobromide, USP ..... 15 mg
Guaifenesin, USP ..... 75 mg
Phenylephrine hydrochloride, USP ..... 5 mg
Chlorphenamine maleate, USP ..... 1 mg

PHARMACOLOGIC CATEGORY
Antitussive/Antihistamine/Nasal Decongestant

INDICATION(S)
For the relief of productive and non-productive cough and nasal congestion associated with common colds and allergic conditions.

DOSAGE AND ADMINISTRATION
Adult: 1 tablet 3-4 times a day.
Children 6 years and older: In proportion or as directed by the physician.
Not indicated for use of children under 6 years of age.

STORAGE
Store at temperatures not exceeding 30°C.
Protect from light and moisture.

AVAILABILITY
Box by 100's in orange colorless PVC/ Alu x 10's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
DR-XY35276

If symptoms persist, consult your doctor.

OVER-THE-COUNTER

OraCare™

INGREDIENTS
REGULAR: Purified Water, Stabilized Chlorine Dioxide, Peppermint Flavor
COOL: Purified Water, Stabilized Chlorine Dioxide, Peppermint, Winsense

CLASSIFICATION
Cosmetic

DIRECTIONS FOR USE
After brushing, rinse with 15mL OraCare Mouthrinse for 60 seconds. No need to dilute with water.

STORAGE
Keep away from heat

AVAILABILITY
REGULAR: 80mL, 250mL, 500mL
COOL: 80mL, 250mL, 500mL

SHELF LIFE
24 months

LEADTIME
90 days

NOTIFICATION NO.
NN-1000002870600 - Regular
NN-1000002821930 - Cool

OVER-THE-COUNTER

℞

ALVERINE CITRATE

Profenil

FORMULATION
Each tablet contains:
Alverine citrate, BP (bis-gamma-phenylpropylethylamine) ..... 60 mg

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOGIC CATEGORY
Antispasmodic

INDICATION
Alverine citrate is an antispasmodic that acts directly on intestinal and uterine smooth muscle. It is used for the relief of smooth muscle spasm in the treatment of gastrointestinal disorders such as irritable bowel syndrome. It is also used in the treatment of dysmenorrhea. Spastic conditions of the gastrointestinal tract, gastric hyperacidity, spasm associated with peptic ulcer, spastic colitis, cholecystitis, biliary dyskinesia and the spasm attendant to diarrhea; spastic conditions of the genitourinary tract attributable to inflammation and caculi; certain primary dysmenorrheas; as an aid in cystoscopic, esophagoscopic and gastroscopic examinations.

DOSAGE AND ADMINISTRATION
Oral: 1 or 2 tablets after meals 1 to 3 times daily or as indicated. When treating spasm associated with peptic ulcer, administer tablets 1/2 hour before meals. In dysmenorrhea, 1 or 2 tablets 4 times daily starting at onset of discomfort.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box by 100's in Alu-PVC blister pack x 10's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
DR-XY36817

PRESCRIPTION

℞

AMLODIPINE BESILATE

FORMULATION
Amlodipine besilate 5mg Tablet
Each tablet contains:
Amlodipine (as besilate) 5mg
Amlodipine besilate 10mg Tablet
Each tablet contains:Amlodipine (as besilate) 10mg

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOGIC CATEGORY
Antihypertensive

INDICATION(S)
For the treatment of hypertension and prophylaxis of angina

DOSAGE AND ADMINISTRATION
Hypertension
Usual Initial Dose: 5 mg once daily, increased if necessary to 10 mg once daily. Similar doses are given in the treatment of stable angina and Prinzmetal's angina.
Or as prescribed by the physician.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Amlodipine besilate 5 mg tablet: PVC film opaque white blister pack of 10's, box of 100 Tablets
Amlodipine besilate 10 mg tablet: PVC film opaque white blister pack of 10's, box of 100 Tablets

SHELF LIFE
36 mos.

PRESCRIPTION

Rx

Enalapril Maleate

PRESCRIPTION

FORMULATION
Enalapril Maleate 5 mg Tablet
Each tablet contains:
Enalapril Maleate ..... 5 mg
Enalapril Maleate 10 mg Tablet
Each tablet contains:
Enalapril Maleate ..... 10 mg
Enalapril Maleate 20 mg Tablet
Each tablet contains:
Enalapril Maleate ..... 20 mg

REGULATORY CLASSIFICATION
Prescription Drug

CLASSIFICATION
Angiotensin converting enzyme (ace) inhibitor

INDICATION(S)
Used in the treatment of hypertension and heart failure. It may also be given prophylactically to patients with asymptomatic left ventricular dysfunction to delay the onset of symptomatic heart failure and to those with left ventricular dysfunction to reduce the incidence of coronary ischaemic events, including myocardial infarction

DOSAGE AND ADMINISTRATION
Treatment of hypertension
Initial Dose: 5 mg daily
The first dose should preferably be given at bedtime. For patients with renal impairment or those who are receiving a diuretic: 2.5 mg daily
If possible the diuretic should be withdrawn 2 or 3 days before enalapril is started and resumed later if necessary.

ELDERLY PATIENTS
Initial Dose: 2.5 mg
Maintenance Dose: 10 to 20 mg once daily up to 40 mg daily in severe hypertension.
It may be given in 2 divided doses if control is inadequate with a single dose.

MANAGEMENT OF HEART FAILURE

℞

Enalapril Maleate

Initial Dose: 2.5 mg daily
Maintenance Dose: 20 mg daily as a single dose or in 2 divided doses.
Treatment should be initiated with a low dose under close medical supervision.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Enalapril Maleate 5 mg Tablet:
Box by 30's and 100's in aluminum foil strip of 10's
Enalapril Maleate 10 mg Tablet:
Box by 30's and 100's in aluminum foil strip of 10's
Enalapril Maleate 20 mg Tablet:
Box by 100's in aluminum foil strip of 10's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
5mg - DR-XY29746
10mg - DR-XY29779
20mg - DRP-7847

PRESCRIPTION

℞

Erythromycin
(as ethylsuccinate)

FORMULATION
Erythromycin ethylsuccinate 200mg suspension
Each 5mL (teaspoonful) reconstituted suspension contains:
Erythromycin (as ethylsuccinate)..... 200mg

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOGIC CATEGORY
Antibacterial

INDICATION(S)
For the treatment of upper and lower respiratory tract, skin and soft tissue inflections of mild to moderate severity caused by susceptible strains of Gram-positive and Gram-negative microoganisms

DOSAGE AND ADMINISTRATION
Adult Dose: 1-2 g daily in 2 to 4 divided doses.
For severe infections: Increase dose up to 4 g daily in divided doses.
Daily doses higher than 1 g should be given in more than 2 divided doses.

Children:
Usual Dose: 30 to 50 mg/kg daily in divided doses. Double dose in severe infections.

Dose based on age:
Infants up to 2 years: 500 mg daily in divided doses.
2 years to 8 years: 1 g daily in divided doses.
Or as prescribed by the physician.

PRESCRIPTION

℞

Erythromycin
(as ethylsuccinate)

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Amber glass bottle with white pilfer-proof plastic cap x 60 mL

SHELF LIFE
48 months

LEADTIME
90 days

REGISTRATION NO.
DRP-5991

PRESCRIPTION

℞

Erythromycin (as stearate)

FORMULATION
Tablet
Each film coated tablet contains:
Erythromycin (as stearate) ..... 250 mg
Erythromycin (as stearate) ..... 500 mg
Granules for Suspension
Each 5 mL (teaspoonful) of the reconstituted suspension contains:
Erythromycin (as ethylsuccinate) ..... 200 mg

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOCIC CATEGORY
Antibacterial

INIDCATION(S)
For the treatment of upper and lower respiratory tract, skin and soft tissue inflections of mild to moderate severity caused by susceptible strains of Gram-positive and Gram-negative microorganisms.

DOSAGE AND ADMINISTRATION
Adult Dose: 1-2 g daily in 2 to 4 divided doses.
For severe infections: Increase dose up to 4 g daily in divided doses.
Daily doses higher than 1 g should be given in more than 2 divided doses.

Children:
Usual Dose: 30 to 50 mg/kg daily in divided doses. Double dose in severe infections.

Dose based on age:
Infants up to 2 years: 500 mg daily in divided doses.
2 years to 8 years: 1 g daily in divided doses.

PRESCRIPTION

℞

Erythromycin
(as stearate)

For the prevention of streptococcal infections in patients with evidence of rheumatic fever or heart disease, who are unable to take penicillin or sulfonamides. a dose of 250 mg twice daily may be given. For the management of acne, maintenance doses as low as 250 mg daily have been used in adults but resistant strains of propionibacteria are widespread; the British National Formulary recommends a dose of 500 mg twice daily.

Or as prescribed by the physician.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
PVC-Clear Blister pack x 10's with metallized wrapper (box of 100's)

SHELF LIFE
250 and 500 mg tablets - 60 months

LEADTIME
90 days

REGISTRATION NO.
250 mg tablets - DR-XY16271
500 mg tablets - DRP-5990

PRESCRIPTION

℞

Lysozyme Hydrochloride

Flemizyme

FORMULATION
Flemizyme 30 mg tablet
Each tablet contains:
Lysozyme hydrochloride ..... 30 mg

Flemizyme 60 mg tablet
Each tablet contains:
Lysozyme hydrochloride ..... 60 mg

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOGIC CATEGORY
Adjunct to antibiotics

INDICATION
FLEMIZYME (30 mg Tablet) in inflammation of the oto-rhino-laryngeal area as what occurs in sinusitis, allergic rhinitis, pharyngitis, laryngitis, tracheitis, otitis, gingivitis, pyorrhea, etc. FLEMIZYME (60 mg Tablet) as potentiator and synergist to antibiotics during antibiotic therapy. FLEMIZYME in high doses as antiviral, in hepatitis, herpes zoster, herpes simplex, chicken pox and warts.

DOSAGE AND ADMINISTRATION
Anti-inflammatory, Mucolytic
FLEMIZYME 30 mg Tablet
Adult: 1 tablet 3 times a day
Antiviral
FLEMIZYME 60 mg Tablet
1 tablet every 250-300 mg of antibiotic administered

As FLEMIZYME's local and systemic tolerance is excellent, doses can be increased in chronic and in severe cases.

If symptoms persist, consult your doctor.

PRESCRIPTION

℞

Lysozyme Hydrochloride

Flemizyme

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box by 100's in blister pack x 10's

SHELF LIFE
36 months

LEADTIME
90 days

REGISTRATION NO.
30 mg - DR-XY35166
60 mg - DR-XY33707

If symptoms persist, consult your doctor.

PRESCRIPTION

℞

Fluocinolone acetonide
Polymyxin B sulfate
Neomycin sulfate

Fluocine

FORMULATION
Each mL of FLUOCINE OTIC SOLUTION contains:
Fluocinolone acetonide, USP 0.025%
(equivalent to 250mcg/mL)
Polymyxin B sulfate, USP ..... 10,000 units
Neomycin sulfate, USP..... 3.5mg
(in aqueous Propylene glycol solution)

REGULATORY CLASSIFICATION
Prescription Drug

PHARMACOLOGIC CATEGORY
Anti-inflammatory/Topical Corticosteroid

INDICATION(S)
For the treatment of upper and lower respiratory tract, skin and soft tissue inflections of mild to moderate severity caused by susceptible strains of Gram-positive and Gram-negative microoganisms

DOSAGE AND ADMINISTRATION
FLUOCINE OTIC SOLUTION is intended for topical use only. The ear should be carefully cleansed and dried; FLUOCINE OTIC SOLUTION may then be applied to the affected ear with the dropper or be used to saturate a wick placed in the external auditory canal. The suggested dosage is 3 to 4 drops, 2 to 4 times daily, or more frequently as required. It is advisable to warm the drops to body temperature before they are placed in the ear in order to avoid caloric stimulation of the vestibular apparatus

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box by 100's in Alu-PVC blister pack x 10's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
DR-XY45500

If symptoms persist, consult your doctor.

PRESCRIPTION

Losartan Potassium

FORMULATION

Losartan potassium 50 mg film-coated tablet
Each film-coated tablet contains:
Losartan potassium, USP ..... 50 mg

Losartan potassium 100 mg film-coated tablet
Each film-coated tablet contains:
Losartan potassium, USP ..... 100 mg

PHARMACOLOGIC CATEGORY

Angiotensin II Receptor Blocker

INDICATION

It is used in the treatment of hypertension, particularly in patients who develop cough with Angiotensin Converting Enzyme (ACE) inhibitors, to reduce the risk of stroke in patients with left ventricular hypertrophy and in the treatment of diabetic nephropathy. It has also been used in heart failure and in myocardial infarction.

DOSAGE AND ADMINISTRATION

Hypertension
Usual dose: 50 mg once daily. The dose may be increased if necessary to 100 mg daily as a single dose or in two divided doses.
For patients with intravascular fluid depletion: Initial dose of 25 mg once daily.
Elderly over 75 years old: Initial dose of 25 mg once daily.
Similar reductions may be appropriate in patients with hepatic or renal impairment.
Children aged 6 years or over: Initial dose of 700 mcg/kg once daily with a maximum of 50 mg.
Diabetic Nephropathy
Initial dose: 50 mg once daily
Increase to 100 mg once daily depending on the blood pressure.
Or as prescribed by the physician.

PRESCRIPTION

Losartan Potassium

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box of 60's in opaque, white PVC/Aluminum Blister pack x 10's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
50 mg - DRP-5095 and DRP-5095-01
100mg - DRP-5096 and DRP-5096-01

PRESCRIPTION

Meloxicam

FORMULATION
Meloxicam 7.5 mg Tablet
Each tablet contains:
Meloxicam, USP ..... 7.5 mg
Meloxicam 15 mg Tablet
Each tablet contains:
Meloxicam, USP ..... 15 mg

PHARMACOLOGIC CATEGORY
Non-steroidal Anti-inflammatory Drug (NSAID)

INDICATION
Used in the management of rheumatoid arthritis, for the short-term symptomatic treatment and acute exacerbations of osteoarthritis, and for the symptomatic treatment of ankylosing spondylitis.

DOSAGE AND ADMINISTRATION
Use the lowest effective dose for the shortest duration consistent with individual patient treatment goals.
For acute exacerbations of osteoarthritis: 7.5 mg daily with food, increase if necessary to 15 mg once daily.
For rheumatoid arthritis and ankylosing spondylitis: 15 mg once daily with food; may be reduced to 7.5 mg daily.
Elderly patients: 7.5 mg daily
For patients on dialysis: 7.5 mg daily
Or as prescribed by the physician.

If symptoms persist, consult your doctor.

PRESCRIPTION

Meloxicam

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box x 100's in Alu/PET Strip Foil x 6's

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
DRP-5955 and DRP-5955-01

If symptoms persist, consult your doctor.

PRESCRIPTION

Metformin Hydrochloride

FORMULATION
Metformin Hydrochloride 500 mg Film-Coated Tablet
Each Film-Coated tablet contains:
Metformin Hydrochloride ..... 500 mg
Metformin Hydrochloride 850 mg Film-Coated Tablet
Each Film-Coated tablet contains:
Metformin Hydrochloride ..... 850 mg

CLASSIFICATION
Oral Hypoglycemic (Biguanide)

INDICATION(S)
For the treatment of non-insulin dependent (Type II) diabetes mellitus not responding to dietary modification.

DOSAGE AND ADMINISTRATION
'For Metformin Hydrochloride 500 mg Tablet:
Management of Type II diabetes mellitus in adults:
Initial: 500 mg twice daily (given with the morning and evening meals); increase daily dosage by 500 mg at weekly intervals (daily dose given in divided doses), up to a maximum of 3 g daily. Doses of up to 2 g/day may be given in 2 divided doses. If a dose of 2.5 g daily is required, it may be better tolerated 3 times daily (with meals). Allow 1-2 weeks between dose titrations. Generally, clinically significant responses are not seen at doses < 1.5 g daily. However, a lower recommended starting dose and gradual increase in dosage is recommended to minimize gastrointestinal symptoms.
For Metformin Hydrochloride 850 mg Tablet:
Management of Type II diabetes mellitus in adults:
Initial: 850 mg once or twice daily with or after meals, gradually increased if necessary to around 2 to 3 g daily.
Or as prescribed by the physician

If symptoms persist, consult your doctor.

PRESCRIPTION

Metformin Hydrochloride

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Metformin Hydrochloride 500 mg Film-Coated Tablet
Box by 100's in foil strip of 10's
Metformin Hydrochloride 850 mg Film-Coated Tablet
Box by 100's in Alu/PVC Blister Pack x 10's

SHELF LIFE
500mg - 24 months
850 mg – 36 months

LEADTIME
90 days

REGISTRATION NO.
500mg – DR-XY28742
850mg – DRP-7836

If symptoms persist, consult your doctor.

PRESCRIPTION

Metoprolol Tartrate

FORMULATION

Metoprolol tartrate 50 mg tablet
Each tablet contains:
Metoprolol tartrate, USP ..... 50 mg

Metoprolol tartrate 100 mg tablet
Each tablet contains:
Metoprolol tartrate, USP ..... 100 mg

CLASSIFICATION

Beta-adrenoceptor Blocker

INDICATION

Used in the treatment of hypotension, angina pectoris, cardiac arrhythmias and myocardial infarction. It is also used as an adjunct in the treatment of hyperthyroidism and as prophylaxis for migraine.

DOSAGE AND ADMINISTRATION

Hypertension: Initial dose of 100 mg daily by mouth, increase weekly according to the response of the patient to 400 mg daily. It may be taken as a single dose or twice daily. A usual maintenance dose is 100 mg to 200 mg daily.

Angina Pectoris: Usual dose of 50 mg to 100 mg two or three times daily by mouth.

Cardiac Arrhythmias: Usual dose is 50 mg two or three times daily by mouth, increase if necessary up to 300 mg daily in divided doses.

If symptoms persist, consult your doctor.

PRESCRIPTION

Metoprolol Tartrate

Myocardial Infarction: In patients who received the full intravenous dose, start oral treatment at 50 mg every 6 hours for 2 days. Subsequent maintenance dosage is 100 mg given twice daily by mouth. In patients who did not receive metoprolol by intravenous injection as part of early management of myocardial infarction, 100 mg twice daily by mouth may be started once condition of patient stabilizes.

Hyperthyroidism: As an adjunct, dose of 50 mg four times daily by mouth.

Migraine: For prophylaxis, doses of 100 mg to 200 mg daily in divided doses.

Or as prescribed by the physician.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Box by 100's in Aluminum foil strip x 10's

SHELF LIFE
36 months

LEADTIME
90 days

REGISTRATION NO.
50mg – DRP-5097 and DRP-5097-01
100mg – DRP-5093 and DP-5093-01

If symptoms persist, consult your doctor.

PRESCRIPTION

Simvastatin

FORMULATION
Simvastatin 10 mg tablet
Each tablet contains:
Simvastatin, USP ..... 10 mg
Simvastatin 20 mg tablet
Each tablet contains:
Simvastatin, USP ..... 20 mg
Simvastatin 40 mg tablet
Each tablet contains:
Simvastatin, USP ..... 40 mg

CLASSIFICATION
HMG-CoA Reductase Inhibitor

INDICATION(S)
It is used for the treatment of hypercholesterolemias particularly in type IIa and IIb hypolipoproteinemias.

DOSAGE AND ADMINISTRATION
Adults:
Initial dose:
20 mg once daily in the evening.
Patients who require only moderate reduction of LDL-cholesterol: Dosage may be started at 10 mg.
Patients requiring a reduction of LDL-cholesterol of >45%: 40 mg once daily.

Maintainance Dose:
Recommended dosing range of 5 - 80 mg daily as a single dose in the evening.
Doses should be individualized according to the baseline LDL-cholesterol levels.
Adjustments should be made at intervals of 4 weeks or more.

If symptoms persist, consult your doctor.

PRESCRIPTION

Simvastatin

Patients with homozygous familial hypercholesterolemia:
Adults: 40 mg in the evening or 80 mg daily in 3 divided doses of 20 mg, 20 mg and evening dose of 40 mg.
Elderly: Maximum reductions in LDL-cholesterol may be achieved with daily dose of </= 20 mg.

Patients who are concomitantly receiving cyclosporine, fibrates or niacin: Dose should not exceed 10 mg daily.

Renal Impairment:
Mild to moderate renal insufficiency:
No dosage modification is necessary

Severe renal impairment (Creatinine clearance <10 mL/minute): Initial dose of 5 mg daily with close monitoring.
Or as prescribed by the physician.

STORAGE
Store at temperatures not exceeding 30°C.

AVAILABILITY
Simvastatin 10 mg Tablet: PVC Clear Blister pack x 10's (Box of 30's)
Simvastatin 20 mg Tablet: PVC Clear Blister pack x 10's (Box of 30's)
Simvastatin 40 mg Tablet: PVC Clear Blister pack x 10's (Box of 30's)

SHELF LIFE
24 months

LEADTIME
90 days

REGISTRATION NO.
10mg – DRP-7837
20mg – DRP-7841
40mg – DRP-5935 and DRP-5935-01

If symptoms persist, consult your doctor.

PRESCRIPTION

Contacts

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Pascual