

ORPHENADRINE CITRATE- orphenadrine citrate tablet, extended release

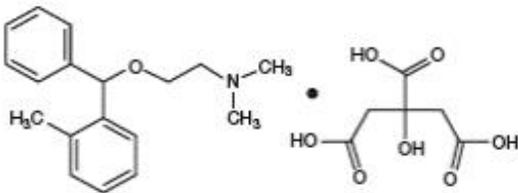
A-S Medication Solutions

Orphenadrine Citrate Extended-Release Tablets

Rx Only

DESCRIPTION

Orphenadrine citrate is the citrate salt of orphenadrine. It occurs as a white, crystalline powder having a bitter taste. It is practically odorless; sparingly soluble in water, slightly soluble in alcohol. The chemical name of orphenadrine citrate is (\pm)-*N,N*-Dimethyl-2-[(*o*-methyl- α -phenylbenzyl)oxy]ethylamine citrate (1:1) having molecular formula $C_{18}H_{23}NO \cdot C_6H_8O_7$ and molecular weight of 461.51. It has the following structural formula:



Each tablet for oral administration contains 100 mg orphenadrine citrate. Each Orphenadrine citrate extended- release tablet contains the following inactive ingredients: hydroxypropyl methylcellulose, lactose monohydrate and magnesium stearate.

CLINICAL PHARMACOLOGY

The mode of therapeutic action has not been clearly identified, but may be related to its analgesic properties. Orphenadrine citrate does not directly relax tense muscles in man. Orphenadrine citrate also possesses anti-cholinergic actions.

INDICATIONS AND USAGE

Orphenadrine citrate extended-release tablets are indicated as an adjunct to rest, physical therapy, and other measures for the relief of discomfort associated with acute painful musculoskeletal conditions.

CONTRAINDICATIONS

Orphenadrine citrate extended-release tablets are contraindicated in patients with glaucoma, pyloric or duodenal obstruction, stenosing peptic ulcers, prostatic hypertrophy or obstruction of the bladder neck, cardio-spasm (mega-esophagus) and myasthenia gravis. Orphenadrine citrate tablets are contraindicated in patients who have demonstrated a previous hypersensitivity to the drug.

WARNINGS

Some patients may experience transient episodes of light-headedness, dizziness or syncope. Orphenadrine may impair the ability of the patient to engage in potentially hazardous activities such as operating machinery or driving a motor vehicle; ambulatory patients should therefore be cautioned accordingly.

PRECAUTIONS

Confusion, anxiety and tremors have been reported in few patients receiving propoxyphene and orphenadrine concomitantly. As these symptoms may be simply due to an additive effect, reduction of dosage and/or discontinuation of one or both agents is recommended in such cases. Orphenadrine citrate should be used with caution in patients with tachycardia, cardiac decompensation, coronary insufficiency, cardiac arrhythmias.

Safety of continuous long-term therapy with orphenadrine has not been established. Therefore, if orphenadrine is prescribed for prolonged use, periodic monitoring of blood, urine and liver function values is recommended.

Pregnancy

Pregnancy Category C

Animal reproduction studies have not been conducted with orphenadrine. It is also not known whether orphenadrine can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Orphenadrine should be given to a pregnant woman only if clearly needed.

Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

ADVERSE REACTIONS

Adverse reactions of orphenadrine are mainly due to the mild anticholinergic action of orphenadrine, and are usually associated with higher dosage. Dryness of the mouth is usually the first adverse effect to appear. When the daily dose is increased, possible adverse effects include: tachycardia, palpitation, urinary hesitancy or retention, blurred vision, dilatation of pupils, increased ocular tension, weakness, nausea, vomiting, headache, dizziness, constipation, drowsiness, hypersensitivity reactions, pruritus, hallucinations, agitation, tremor, gastric irritation, and rarely urticaria and other dermatoses. Infrequently, an elderly patient may experience some degree of mental confusion.

These adverse reactions can usually be eliminated by reduction in dosage. Very rare cases of aplastic anemia associated with the use of orphenadrine tablets have been reported. No causal relationship has been established.

DRUG ABUSE AND DEPENDENCE

Orphenadrine has been chronically abused for its euphoric effects. The mood elevating effects may occur at therapeutic doses of orphenadrine.

OVERDOSAGE

Orphenadrine is toxic when overdosed and typically induces anticholinergic effects. In a review of orphenadrine toxicity, the minimum lethal dose was found to be 2 to 3 grams for adults; however, the range of toxicity is variable and unpredictable.

Treatment for orphenadrine overdose is evacuation of stomach contents (when necessary), charcoal at repeated doses, intensive monitoring, and appropriate supportive treatment of any emergent anticholinergic effects.

DOSAGE AND ADMINISTRATION

Adults

Two tablets per day; one in the morning and one in the evening.

HOW SUPPLIED

Product: 50090-0169

NDC: 50090-0169-0 20 TABLET, EXTENDED RELEASE in a BOTTLE

NDC: 50090-0169-1 30 TABLET, EXTENDED RELEASE in a BOTTLE

NDC: 50090-0169-2 14 TABLET, EXTENDED RELEASE in a BOTTLE

NDC: 50090-0169-5 60 TABLET, EXTENDED RELEASE in a BOTTLE

Orphenadrine Citrate



ORPHENADRINE CITRATE

orphenadrine citrate tablet, extended release

Product Information

| | | | | |
|--|---|---|-------------------------------|---------------------------|
| Product Type | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:50090-0169(NDC:43386-480) | |
| Route of Administration | ORAL | | | |
| Active Ingredient/Active Moiety | | | | |
| Ingredient Name | | Basis of Strength | Strength | |
| ORPHENADRINE CITRATE (UNII: X0A40N8I4S) (ORPHENADRINE - UNII:AL805O9OG9) | | ORPHENADRINE CITRATE | 100 mg | |
| Inactive Ingredients | | | | |
| Ingredient Name | | | Strength | |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) | | | | |
| LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) | | | | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | | | | |
| Product Characteristics | | | | |
| Color | WHITE | Score | no score | |
| Shape | ROUND | Size | 10mm | |
| Flavor | | Imprint Code | NL4 | |
| Contains | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:50090-0169-0 | 20 in 1 BOTTLE; Type 0: Not a Combination Product | 06/21/2016 | |
| 2 | NDC:50090-0169-1 | 30 in 1 BOTTLE; Type 0: Not a Combination Product | 11/28/2014 | |
| 3 | NDC:50090-0169-2 | 14 in 1 BOTTLE; Type 0: Not a Combination Product | 11/28/2014 | |
| 4 | NDC:50090-0169-5 | 60 in 1 BOTTLE; Type 0: Not a Combination Product | 11/28/2014 | |
| Marketing Information | | | | |
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| ANDA | ANDA040284 | 06/19/1998 | | |

Labeler - A-S Medication Solutions (830016429)

Establishment

| Name | Address | ID/FEI | Business Operations |
|--------------------------|---------|-----------|--|
| A-S Medication Solutions | | 830016429 | RELABEL(50090-0169) , REPACK(50090-0169) |

