

Ultr[®]-C

DESCRIPTION:

Ultr[®]-C tablets for oral administration.

Each tablet contains:

Methanamine	81.6 mg
Sodium Phosphate Monobasic	40.8 mg
Phenyl Salicylate	36.2 mg
Methylene Blue	10.8 mg
Hyoscyamine Sulfate	0.12 mg

HYOSCYAMINE SULFATE. [620-61-1] (3S)-endo- α -(Hydroxymethyl)-benzeneacetic acid 8-methyl-8-azabicyclo[3.2.1]oct-3-yl ester sulfate(2:1)(salt); 1 α H-5 α H-tropan-3- α -ol(-)-tropate(ester) sulfate(2:1)(salt); 3 α -tropanyl S(-)-tropate; l-tropic acid ester with tropine; l-tropine tropate. C₂₀H₂₈N₂O₇S. Hyoscyamine Sulfate is an alkaloid of belladonna. Exists as a white crystalline powder. Its solutions are alkaline to litmus. Affected by light, it is slightly soluble in water; freely soluble in alcohol; sparingly soluble in ether.

METHANAMINE. [100-97-0] 1,3,5,7-Tetraazatricyclo [3.3.1.1^{-1,3}] decane; hexamethylenetetramine; HMT; HMTA; hexamine; 1,3,5,7-tetraazaadamantane hexamethylenimine; Uritone; Urotropin. C₆H₁₂N₄; mol wt 140.19; C 51.40%, H 8.63%, N 39.96%. Methanamine (hexamethylenetetramine) exists as colorless, lustrous crystals or white crystalline powder. Its solutions are alkaline to litmus. Freely soluble in water, soluble in alcohol and in chloroform.

METHYLENE BLUE. [61-73-4] 3,7-Bis(dimethylamino) phenothiazin-5-ium chloride; C.I. Basic Blue 9; methylthioninium chloride; tetramethylnitrosonium chloride; 3,7-bis(dimethylamino) phenazothiazinone chloride. C₁₆H₁₈ClN₃S; mol wt 319.85; C 60.08%, H 5.67%, Cl 11.08%, N 13.14%, S 10.03%. Methylene Blue (Methylthionine chloride) exists as dark green crystals. It is soluble in water and in chloroform; sparingly soluble in alcohol.

PHENYL SALICYLATE. [118-55-8] 2-Hydroxybenzoic acid phenyl ester; Salol. C₁₅H₁₂O₂; mol wt 214.22; C 72.89%, H 4.71%, O 22.41%. Made by the action of phosphorus oxy-chloride on a mixture of phenol and salicylic acid. Phenyl Salicylate exists as white crystals with a melting point of 41°-43° C. It is very slightly soluble in water and freely soluble in alcohol.

SODIUM PHOSPHATE MONOBASIC. [7558-90-7] Phosphoric acid sodium salt (1:1); Sodium biphosphate; sodium diphosphoric phosphate; acid sodium phosphate; monosodium orthophosphate; primary sodium phosphate; H₂NaO₇P; mol wt 119.98; H 1.68%, Na 19.16%, O 53.34%, P 25.82%. Monohydrate, white, odorless slightly deliquescent crystals or granules. At 100° C loses all its water; when ignited it converts to metaphosphate. It is freely soluble in water and practically insoluble in alcohol. The aqueous solution is acid, pH of 0.1 molar aqueous solution at 25° C; 4.5.

Ultr[®]-C tablets contain inactive ingredients: Croscopolone, Dicalcium Phosphate, FD&C Blue #1 Aluminum Lake, Magnesium Stearate, Microcrystalline Cellulose, Polyethylene Glycol, Polyvinyl Alcohol, Stearic Acid, Talc, and Titanium Dioxide.

CLINICAL PHARMACOLOGY:

HYOSCYAMINE SULFATE is a parasympatholytic which relaxes smooth muscles and thus produces an antispasmodic effect. It is well absorbed from the gastrointestinal tract and is rapidly distributed throughout the body tissues. Most is excreted in the urine within 12 hours, 13% to 50% being unchanged. Its biotransformation is hepatic. Its protein binding is moderate.

METHANAMINE degrades in an acidic urine environment releasing formaldehyde which provides bactericidal or bacteriostatic action. It is well absorbed from the gastrointestinal tract. 70%-90% reaches the urine unchanged at which point it is hydrolyzed if the urine is acidic. Within 24 hours it is almost completely (90%) excreted; of this at a pH of 5, approximately 20% is formaldehyde. Protein binding - some formaldehyde is bound to substances in the urine and surrounding tissues. Methanamine is freely distributed to body tissue and fluids but is not clinically significant as it does not hydrolyze at pH greater than 6.8.

METHYLENE BLUE possesses weak antiseptic properties. It is well absorbed by the gastrointestinal tract and rapidly reduced to leukomethylene blue which is stabilized in some combination form in the urine. 75% is excreted unchanged.

PHENYL SALICYLATE releases salicylate, a mild analgesic for pain.

SODIUM PHOSPHATE MONOBASIC an acidifier, helps to maintain an acid pH in the urine necessary for the degradation of methanamine.

INDICATIONS AND USAGE:

Ultr[®]-C tablets are indicated for the treatment of symptoms of irritative voiding. Indicated for the relief of local symptoms, such as inflammation, hypermotility, and pain, which accompany lower urinary tract infections. Indicated for the relief of urinary tract symptoms caused by diagnostic procedures.

CONTRAINDICATIONS:

Hypersensitivity to any of the ingredients is possible. Risk benefits should be carefully considered when the following medical problems exist: cardiac disease (especially cardiac arrhythmias, congestive heart failure, coronary heart disease, and mitral stenosis); gastrointestinal tract obstructive disease; glaucoma; myasthenia gravis, acute urinary retention may be precipitated in obstructive uropathy (such as bladder neck obstruction due to prostatic hypertrophy).

WARNINGS:

Do not exceed recommended dosage. If rapid pulse, dizziness or blurring of vision occurs discontinue use immediately.

PRECAUTIONS:

Cross sensitivity and/or related problems - Patients intolerant of belladonna alkaloids or salicylates may be intolerant of this medication also. Delay in gastric emptying could complicate the management of gastric ulcers.

Pregnancy/Reproduction (FDA Pregnancy Category C) - Hyoscyamine and methanamine cross the placenta. Studies concerning the effect of hyoscyamine and methanamine on

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pregnancy and reproduction has not been done in animals or humans. Thus it is known whether **Ultr[®]-C** tablets cause fetal harm when administered to a pregnant woman or can affect reproductive capacity. **Ultr[®]-C** tablets should be given to a pregnant woman only if clearly needed.

Breast feeding - Problems in humans have not been documented; however, methanamine and traces of hyoscyamine are excreted in breast milk. Accordingly, **Ultr[®]-C** tablets should be given to a nursing mother with caution and only if clearly needed.

Prolonged use - There have been no studies to establish the safety of prolonged use in humans. No known long-term animal studies have been performed to evaluate carcinogenic potential.

Pediatric use - Infants and young children are especially susceptible to the toxic effect of the belladonna alkaloids.

Geriatric use - Use with caution in elderly patients as they may respond to usual doses of hyoscyamine with excitement, agitation, drowsiness or confusion.

ADVERSE REACTIONS:

Cardiovascular: rapid heartbeat, flushing
Central Nervous System: blurred vision, dizziness, drowsiness
Genitourinary: difficult micturition, acute urinary retention
Gastrointestinal: dry mouth, nausea and vomiting
Respiratory: shortness of breath or trouble breathing
Serious allergic reactions to this drug are rare. Seek immediate medical attention if you notice symptoms of a serious allergic reaction, including itching, rash, severe dizziness, swelling or trouble breathing.

This medication can cause urine and sometimes stools to turn blue to blue-green. This effect is harmless and will subside after medication is stopped.

Call your doctor or physician for medical advice about side effects. To report SUSPECTED ADVERSE REACTIONS, contact Mission Pharmaceutical at 1-800-845-7827 or FDA at 1-800-FDA-1088, www.fda.gov/medwatch.

Drug interactions - As a result of hyoscyamine's effects on gastrointestinal motility and gastric emptying, absorption of other oral medications may be decreased during concurrent use with this combination medication.

Urinary alkalinizers and thiazide diuretics: May cause the urine to become alkaline reducing the effectiveness of methanamine by inhibiting its conversion to formaldehyde.

Antimuscarinics: Concurrent use may intensify antimuscarinic effects of hyoscyamine because of secondary antimuscarinic activities of these medications.

Antacids/antidiarrheals: Concurrent use may reduce absorption of hyoscyamine resulting in decreased therapeutic effectiveness. Concurrent use with antacids may cause urine to become alkaline reducing effectiveness of methanamine by inhibiting its conversion to formaldehyde. Doses of these medications should be spaced 1 hour apart from doses of hyoscyamine.

Antimayesthetics: Concurrent use with hyoscyamine may further reduce intestinal motility, therefore caution is recommended.

Ketoconazole and hyoscyamine may cause increased gastrointestinal pH. Concurrent administration with hyoscyamine may result in marked reduction in the absorption of ketoconazole. Patients should be advised to take this combination at least 2 hours after ketoconazole.

Monoamine oxidase (MAO) inhibitors: Concurrent use with hyoscyamine may intensify antimuscarinic side effects.

Opioid (narcotic) analgesics may result in increased risk of severe constipation.

Sulfonamides: These drugs may precipitate with formaldehyde in the urine increasing the danger of crystalluria.

Patients should be advised that the urine may become blue to blue-green and the feces may be discolored as a result of the excretion of the methylene blue.

DRUG ABUSE AND DEPENDENCE:

A dependence on the use of **Ultr[®]-C** tablets has not been reported and due to the nature of its ingredients, abuse of **Ultr[®]-C** tablets is not expected.

OVERDOSAGE:

Emesis or gastric lavage. Slow intravenous administration of physostigmine in doses of 1 to 4 mg (0.5 to 1 mg in children), repeated as needed in one to two hours to reverse severe antimuscarinic symptoms.

Administration of small doses of diazepam to control excitement and seizures. Artificial respiration with oxygen if needed for respiratory depression. Adequate hydration. Symptomatic treatment as necessary.

If overdose is suspected, contact your local Poison Control Center or emergency room immediately. US residents can contact the US National Poison Hotline at 1-800-222-1222.

DOSAGE AND ADMINISTRATION:

Adults - one tablet orally 4 times per day followed by liberal fluid intake.

Older Children - dosage must be individualized by physician. Not recommended for use in children 6 years of age or younger.

HOW SUPPLIED:

Ultr[®]-C tablets are blue tablets and imprinted with "SAT" on one side and "902" on the opposite side. NDC 0178-0513-01, bottle of 100 Tablets.

STORAGE:

Dispense in a tight, light-resistant container as defined in the USP/NF with a child resistant closure. Store at controlled room temperature 20°-25°C (68°-77°F).

Keep in a cool, dry place. Keep container tightly closed.

Keep this and all drugs out of reach of children.

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