Pipemidic acid capsules

**PHARMACOLOGICAL ACTION:**
Deblaston contains the antimicrobial substance Pipemidic acid which is related to Nalidixic acid.

Pipemidic acid is absorbed from the intestine, reaches effective levels in the tissues, is excreted in the urine in high concentrations in the active form and is metabolized only to a small extent.

In-vitro Deblaston is effective against Gram negative organisms, except Serratia marcescens, also effective against Pseudomonas aeruginosa.

It is not effective against Gram positive organisms especially Strep. faecalis.

**INDICATIONS:**
Urethrits, cystitis, pyelonephritis and acute prostatitis due to susceptible organisms.

**CONTRA-INDICATIONS:**
Deblaston should not be administered to children or adolescents during the growth phase and is contra-indicated in pregnancy and lactation.

Deblaston is contra-indicated in severe renal failure with anuria or oliguria (serum creatinine above 3 mg%, creatinine clearance below 10 mL/min.).

Contra-indicated in patients with hepatic disease or patients in whom hepatic dysfunction develops on treatment.

**DOSAGE AND DIRECTIONS FOR USE:**
Two capsules should be taken twice daily - morning and evening with meals - for at least 5 days. Treatment should not exceed 10 days in duration. At the end of this time, it is advisable to change to another anti-bacterial agent so as to counter any possible development of resistance on the part of the micro-organisms.

**SIDE-EFFECTS AND SPECIAL PRECAUTIONS:**
Gastro-intestinal symptoms such as anorexia, gastric discomfort, abdominal distention and pain, nausea, vomiting, thirst, diarrhoea, constipation, fatigue, dizziness, headache and stomatitis may occur. There have been reports of skin reactions (pruritis, erythema and other eruptions). Anaphylactic shock can occur.

During treatment excessive exposure to sunlight or ultraviolet radiation should be avoided.

In the event of an allergic reaction, treatment should be discontinued at once and your doctor should be consulted.

As adequate information is not yet available regarding the use of Deblaston in patients with existing epilepsy or epileptiform conditions, it should not be given to such patients without careful consideration of the risks and the benefits.

The development of cross-resistance between Pipemidic acid and Nalidixic acid is in principle to be expected. However, mutants resistant to Nalidixic acid may be sensitive to Pipemidic acid.

Elevation of BUN, creatine levels, SGOT and SGPT as well as a decrease in leucocyte counts can occur. False high values of urine protein are sometimes recorded by certain measuring methods. To avoid this it is recommended that the boiling method or the test paper method be used.

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Information for Health Care Professionals

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Taj Pharmaceuticals Limited is a pharmaceutical company founded and based in India. The company manufacturers pharmaceutical formulations and API for India and other countries of the world. The company was established in 1995 as an enterprise and in 2004 became a public limited company. As per Mumbai Pharmaxil and Chemixil association the company manufacturers and exports to countries like Albania, Argentina, Austria, Chile and Iraq. In 1995 pharmaceutical wing only has a schedule M certification for pharmaceuticals products manufacturing in India. Taj Pharmaceuticals established its manufacturing unit in Gujarat because of government policies in 1999 with WHO / GMP licence. The company in 2003 revived all the old manufacturing units and approached the FDA Gujarat for 4000 new pharmaceuticals drug permissions for the first time in India.

According to the Indian Trade Mark the company owns about 450 brands and 4600 generic manufacturing permissions in India. According to the export data analysis the company was the largest exporter of generic medicines to the Europe and Middle East countries.

www.tajpharma.com

The company medicines are present in France, Georgia, Egypt and CIF countries.

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