

**1. Name of the Medicinal Product**

**1.1 Product Name:** AMYN DT 250

**1.2 Strength:** Amoxicillin Tablets for Oral suspension USP 250mg

**1.3 Pharmaceutical Form:** Tablets

**2. Qualitative and Quantitative Composition**

**Qualitative declaration:**

Each Dispersible tablet contains:

Amoxicillin Trihydrate USP

Equivalent to Amoxicillin.....250 mg

### **3. Pharmaceutical form**

Dispersible Tablet

AMYN DT 250 are White to off white round shaped, flat face beveled edge tablets with breakline on one side and plain on other side without any visible defects.

### **4 Clinical Particulars**

#### **4.1 Therapeutic indications-**

Amoxicillin is a broad-spectrum antibiotic indicated for the treatment of commonly occurring bacterial infections such as:

Upper respiratory tract infections: e.g. sinusitis, acute pharyngitis.

Lower respiratory tract infections: e.g. acute exacerbations of chronic bronchitis, lobar and bronchopneumonia, uncomplicated community acquired pneumonia, H. influenzae infections.

Gastrointestinal tract infections: e.g. acute gastritis, peptic ulcer disease and invasive salmonellosis.

Skin and soft tissue infections: e.g. Cellulitis, erysipelas, osteomyelitis

Genito-urinary tract infections: e.g. cystitis, urethritis, pyelonephritis, bacteriuria in pregnancy, septic abortion, puerperal sepsis.

ENT Infections: Cervical adenitis, otitis media.

Dental infections: Dental abscess (as an adjunct to surgical management), suppurative odontogenic infections.

Listerial meningitis.

Prophylaxis of endocarditis: Amoxicillin may be used for the prevention of bacteremia associated with procedures such as dental extraction, in patients at risk of developing of endocarditis.

Strains of the following organisms are generally sensitive to the bacterial action of Amoxicillin in vitro:

Gram positive (Aerobes): Streptococcus faecalis, Streptococcus pneumoniae, Streptococcus pyogenes, Streptococcus viridans, penicillin-sensitive Staphylococcus aureus, Corynebacterium species, Bacillus anthracis, Listeria monocytogenes

Gram positive (Anaerobes): Clostridium species

Gram negative (Aerobes): Haemophilus influenzae, Eschericia coli, Proteus mirabilis, Salmonella species, Bordetela pertussis, Brucella species, Shigella species, Neisseria meningitidis, Pasteurella septica, Vibrio cholerae, Helicobacter pylori.

Amoxicillin is susceptible to degradation by  $\beta$ -lactamases and, therefore, the spectrum of activity for Amoxicillin does not include organisms that produce these enzymes, including resistant Staphylococci and all strains of Pseudomonas, Klebsiella and Enterobacter.

#### 4.2 Posology and Method of Administration

Posology:

Standard children's dosage (up to 10 years of age): For AMYN DT 250 and AMYN DT 250 Tablets:

CHILD up to 10 years: 250 mg every 8 hours doubled in severe infections.

Patients with renal impairment: In renal impairment, the excretion of the antibiotic will be delayed and depending on the degree of impairment, it may be necessary to reduce the total daily dosage according to the following scheme:

Children under 40Kg (**including older patients**):

Mild impairment (creatinine clearance > 30ml/min)      -No change in dosage

Moderate impairment (creatinine clearance 10-30ml/min)      -5mg/kg b.i.d. maximum

Severe impairment (creatinine clearance <10ml/min)      -15mg/Kg o.d.

Method of administration : Oral

#### 4.3 Contraindications

Amoxicillin is penicillin and should not be given to penicillin hypersensitive patients. Attention should be paid to possible cross-sensitivity with other  $\beta$ -lactam antibiotics e.g. cephalosporins.

#### **4.4 Special warnings and precautions for use**

Before initiating therapy with amoxicillin, careful enquiry should be made concerning previous hypersensitivity reactions to penicillins, cephalosporins.

Serious and occasionally fatal hypersensitivity (anaphylactoid) reactions have been reported in patients on penicillin therapy. These reactions are more likely to occur in individuals with a history of hypersensitivity to  $\beta$ -lactam antibiotics (see contra-indications). Erythematous (morbilliform) rashes have been associated with glandular fever in patients receiving Amoxicillin.

Prolonged use may also occasionally result in overgrowth of non-susceptible organisms.

Dosage should be adjusted in patients with renal impairment (see posology and method of administration).

#### **4.5 Interactions with other medicinal products and other forms of interactions**

In common with other broad-spectrum antibiotics, Amoxicillin may reduce the efficacy of oral contraceptives and patients should be warned accordingly. Concurrent administration of Allopurinol during treatment with Amoxicillin can increase the likelihood of allergic skin reactions.

Prolongation of prothrombin time has been reported rarely in patients receiving Amoxicillin. Appropriate monitoring should be undertaken when anticoagulants are prescribed concurrently.

It is recommended that when testing for the presence of glucose in urine during Amoxicillin treatment, enzymatic glucose oxidase methods should be used.

Due to the high urinary concentrations of Amoxicillin, false positive readings are common with chemical methods.

Probenecid decreases the renal tubular secretion of Amoxicillin. Concurrent use with Amoxicillin may result in increased and prolonged blood levels of Amoxicillin.

#### **4.6 Pregnancy and Lactation**

##### **Pregnancy:**

Animal studies with Amoxicillin have shown no teratogenic effects. However, treatment with Amoxicillin may be considered appropriate when the potential benefits outweigh the potential risks associated with treatment.

##### **Lactation:**

Amoxicillin may be given during lactation. With the exception of the risk of sensitization associated with the excretion of trace quantities of Amoxicillin in breast milk, there are no known detrimental effects for the breast-fed infant.

#### **4.7 Effects on ability to drive and use machine**

Adverse effects on the ability to drive or operate machinery have not been observed.

#### **4.8 Undesirable effects**

Side effects, as with other penicillins, are uncommon and mainly of a mild and transitory nature.

Hypersensitivity reactions: If any hypersensitivity occurs, the treatment should be discontinued.

Skin rash, pruritis and urticaria have been reported occasionally. Rarely, skin reaction such as erythema multiforme and Steven-Johnson syndrome, toxic epidermal necrolysis and bullous and exfoliative dermatitis have been reported. As with other antibiotics, severe allergic reactions including angioneurotic oedema, anaphylaxis, serum sickness and hypersensitivity vasculitis have been reported rarely.

Gastrointestinal reactions: Effects include nausea, vomiting and diarrhea. Intestinal candidiasis and antibiotic associated colitis (including pseudo-membranous colitis and hemorrhagic colitis) have been reported rarely. Intestinal nephritis can occur rarely.

Hepatic effects: A moderate rise in AST and/or ALT has been occasionally noted but the significance of this is unclear. As with other  $\beta$ -lactam antibiotics, hepatitis and cholestatic jaundice have been reported rarely.

Hematological effects: As with other  $\beta$ -lactam antibiotics, reversible leucopenia (including severe neutropenia or agranulocytosis), reversible thrombocytopenia and hemolytic anemia have been reported rarely.

Prolongation of bleeding time and prothrombin time has also been reported rarely.

CNS effects: CNS effects have been reported rarely. They include hyperkinesias, dizziness and convulsions. Convulsions may occur in patients with impaired renal function or in those receiving high doses.

Miscellaneous: Superficial tooth discoloration has been reported rarely and mostly with the dispersible tablets. It can usually be removed by brushing.

#### **4.9 Overdose**

Problems of over dosage with amoxicillin are unlikely to occur. If encountered, gastrointestinal effects such as nausea, vomiting and diarrhea may be evident and should be treated symptomatically with attention to the water/electrolyte balance. During the administration of high doses of amoxicillin, adequate fluid intake and urinary output must be maintained. Amoxicillin can be removed from the circulation by hemodialysis.

#### **5.0 Pharmacological properties**

##### **5.1 Pharmacodynamic properties**

Amoxicillin is a semi-synthetic aminopenicillin of the  $\beta$ -lactam group of antibiotics. It has a broad spectrum of antibacterial activity against many Gram-positive and Gram-negative micro-organisms, acting through the inhibition of biosynthesis of cell wall mucopeptide. It is rapidly bactericidal and possesses the safety profile of penicillin.

##### **5.2 Pharmacokinetic properties**

Amoxicillin is well absorbed. Oral administration, usually at convenient t.d.s. dosage, produces high serum levels, independent of the time at which food is taken. Amoxicillin is not highly protein bound; approximately 18% of total plasma drug content is bound to protein. Amoxicillin diffuses readily into

most body tissues and fluids, with the exception of the brain and spinal fluid. Inflammation generally increases the permeability of the meninges to penicillins and this may apply to amoxicillin. The elimination half-life is approximately 1 hour. The major route of elimination for amoxicillin is via the kidney. Approximately 60-70% of Amoxicillin is excreted unchanged in urine during the first 6 hours after administration of a standard dose. Amoxicillin is partly excreted in the urine as the inactive penicilloic acid in quantities equivalent to 10-25% of the initial dose.

### **5.3 Pre-clinical safety data**

An oral dose of 4000mg amoxicillin/kg was essentially without effect on the barbiturate sleep, electroshock convulsion and pain response to tail pinch in mice, and on the body temperature, blood pressure, heart rate, electrocardiogram pattern and urine volume in unanesthetized rats. Similarly, an oral dose of 1000mg amoxicillin/kg produced no effect on the blood pressure, heart rate and intestinal motility in unanesthetized dogs.

At concentration of 0.5mg/ml, amoxicillin had no effect on the spontaneous motilities of isolated rat uterus and rabbit ileum, and on the contraction of guinea pig ileum induced by acetylcholine, histamine and barium chloride, and of rat stomach contraction from serotonin.

Amoxicillin had no local irritating and anesthetic effect on the rabbit eyes at concentration of 4%.

QT-interval-prolonging potential of Amoxicillin was studied in a conscious dog model. Three doses of test compounds or vehicle were administered orally to male beagle dogs (n=4), and telemetry signals were recorded for 24 h after administration.

Administration of Amoxicillin did not produce any significant change in the QTc interval. Amoxicillin at a dose level of 70, 200, and 500mg/kg had no significant effect on any parameters measured at any dose. The maximum group-mean-difference in QTcF interval from the time-matched vehicle values at 70, 200, and 500mg/kg was 0%, 2%, and 4%, respectively.

The effects of enteral administration of doxycycline, **amoxicillin**, cephalexin, and enrofloxacin at therapeutic dosages for a typical duration were determined on hemostatic variables in healthy Beagle dogs.

Doxycycline (10 mg/kg, PO, q 12 h), **amoxicillin (30 mg/kg, PO, q 12 h)**, cephalexin (30 mg/kg, PO, q 12 h), and enrofloxacin (20 mg/kg, PO, q 24 h) were administered in random order to 10 healthy dogs at standard therapeutic dosages for 7 days, with a 7-day washout period between subsequent antimicrobials. In addition, 4 Beagles served as control dogs. Variables were evaluated before and after antimicrobial administration; they included platelet count, Hct, 1-stage prothrombin time (PT), activated partial thromboplastin time (PTT), fibrinogen concentration, and platelet function. Platelet function was assessed *via* buccal mucosal bleeding time, aggregation, and a platelet-function analyzer.

Administration of all antimicrobials caused a slight prolongation of 1-stage PT and activated PTT and slight decrease in fibrinogen concentration. Cephalexin caused a significant increase in 1-stage PT and activated PTT, **amoxicillin caused a significant increase in activated PTT**, and enrofloxacin caused a significant decrease in fibrinogen concentration. Platelet count or function did not differ significantly after administration of any antimicrobial.

**It was concluded that oral administration of commonly used antimicrobials in healthy dogs resulted in minor secondary hemostatic abnormalities, with no change in platelet count or function.**

Drug-related immunologic destruction of granulocytes usually develops after the second week of amoxicillin therapy but may be delayed and occur weeks or months into a course of therapy. It is characterized by a sudden fall in the peripheral neutrophil count; fever may be present. Absolute neutropenia can be severe and may place the patient at increased risk of infection. Drug-induced neutropenia may be due to antibodies to the neutrophil. The neutropenia seen with prolonged high-dose therapy with penicillins and cephalosporins is of uncertain etiology, but it may not have an immunologic basis as rechallenge is not associated with an accelerated recurrence of the neutropenia and the neutrophil count may fall more slowly.



An antibody-induced immune thrombocytopenia has been described with the penicillins and cephalosporins. These are reversed quickly when the particular drug is discontinued.

Although oral administration of amoxicillin to rabbits and guinea pigs for 30 days did not form antibodies, parenteral administration caused the formation of antibodies and hemagglutinins and passive cutaneous anaphylaxis. The cross reactivity of amoxicillin was observed in erythrocyte coagulation and passive cutaneous anaphylaxis reactions.

**6.0 Pharmaceutical Particulars**

**6.1 List of Excipients**

Microcrystalline Cellulose PH 102, Aspartame, Crospovidone, Ess. Fl. DC.  
Trusil Lemon, Flavour peppermint DC 117 quest, Magnesium Stearate,  
Purified Talc, Colloidal anhydrous silica.

**6.2 Incompatibilities**

None

**6.3 Shelf life-**

24 Months (2 years)

Proposed shelf life (after first opening container): NA

Proposed shelf life (after reconstitution or dilution): NA

**6.4 Special precautions for storage**

Store below 30°C in a dry place.

**6.5 Nature and contents of container**

Each Alu/Alu Strip contains 15 tablets & 10 tablets, 3 strips are packed in a  
Carton & 2 strips are packed in a carton.

**6.6 Instructions for use and handling**

None

**7. MARKETING AUTHORISATION HOLDER**

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**8. NUMBER(S) IN THE NATIONAL REGISTER OF FINISHED  
PHARMACEUTICAL PRODUCTS**

4914/6325/NMR/2018

**9. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Jun 19, 2023

**10. DATE OF REVISION OF THE TEXT**

Jun 19, 2023