

PRODUCT MONOGRAPH

<sup>T/C</sup> **DOM-CLONAZEPAM**

(Clonazepam Tablets, USP)

0.5 mg and 2 mg

Anticonvulsant

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## <sup>T/C</sup> DOM-CLONAZEPAM

(Clonazepam Tablets, USP)

### PART I: HEALTH PROFESSIONAL INFORMATION

#### SUMMARY PRODUCT INFORMATION

| <b>Route of Administration</b> | <b>Dosage Form / Strength</b> | <b>Clinically Relevant Nonmedicinal Ingredients</b>   |
|--------------------------------|-------------------------------|---|
| Oral                           | Tablets 0.5 mg and 2 mg       | lactose<br><i>For a complete listing see Dosage Forms, Composition and Packaging section.</i> |

#### INDICATIONS AND CLINICAL USE

DOM-CLONAZEPAM (clonazepam) has been found useful when used alone or as an adjunct in the management of myoclonic and akinetic seizures and petit mal variant (Lennox-Gastaut syndrome).

DOM-CLONAZEPAM may be of some value in patients with absence spells (petit mal) who have failed to respond to succinimides.

Up to nearly one-third of the patients in some studies have shown a loss of anticonvulsant activity, often within the first three months of administration of clonazepam. In some cases dosage adjustment may re-establish efficacy.

##### **Geriatrics** (>65 years of age)

There is no clinical trial experience with clonazepam in seizure disorder patients 65 years of age and older (See DOSAGE AND ADMINISTRATION).

##### **Pediatrics** (<18 years of age)

For a brief description see WARNINGS AND PRECAUTIONS, Special Populations, Pediatrics (<5 years of age) and DOSAGE AND ADMINISTRATION, Recommended Dose and Dosage Adjustment, Children.

#### CONTRAINDICATIONS

DOM-CLONAZEPAM (clonazepam) should not be used in patients with a history of sensitivity to benzodiazepines including clonazepam, or any of its excipients. DOM-CLONAZEPAM is

also contraindicated in patients with clinical or biochemical evidence of severe hepatic insufficiency, severe respiratory insufficiency, myasthenia gravis, sleep apnea syndrome, and in patients with narrow angle glaucoma.

## **WARNINGS AND PRECAUTIONS**

### **General**

A paradoxical increase in seizure activity or the appearance of new seizure types has occurred in a very few patients during treatment with clonazepam. When used in patients in whom several different types of seizures coexist, clonazepam may increase the incidence or precipitate the onset of generalized tonic-clonic seizures (grand mal). These phenomena may require the addition of appropriate anticonvulsants or an increase in their dosage. The concomitant use of valproic acid and clonazepam may produce absence status.

The abrupt withdrawal of clonazepam, particularly in those patients on long-term, high dose therapy, may precipitate status epilepticus. Therefore, as with any other anticonvulsant, gradual withdrawal is essential when discontinuing clonazepam. While clonazepam is being gradually withdrawn, the simultaneous substitution of incremental doses of another anticonvulsant may be indicated.

Clonazepam should be used only with particular caution in patients with spinal or cerebellar ataxia, and in the event of acute intoxication with alcohol or drugs.

Patients with a history of depression and/or suicide attempts should be kept under close supervision.

Anterograde amnesia may occur using therapeutic dosages of benzodiazepines, with the risk increasing at higher doses (see ADVERSE REACTIONS).

### **Concomitant use of alcohol/CNS depressants**

The concomitant use of DOM-CLONAZEPAM with alcohol or/and CNS depressants should be avoided. Such concomitant use has the potential to increase the clinical effects of clonazepam possibly including severe sedation, clinically relevant respiratory and/or cardiovascular depression (see DRUG INTERACTIONS). Patients should be advised against the concurrent use of alcohol and other CNS depressant drugs.

### **Medical history of alcohol or drug abuse**

DOM-CLONAZEPAM should be used with extreme caution in patients with a history of alcohol or drug abuse. Benzodiazepines have produced habituation, dependence and withdrawal symptoms similar to those noted with barbiturates and alcohol. The risk of dependence increases with dose and duration, and is greater in patients with a medical history of alcohol and drug abuse (See WARNINGS AND PRECAUTIONS, Dependence and Tolerance).

### **Lactose intolerance**

Lactose is a non-medicinal ingredient in DOM-CLONAZEPAM. Therefore, patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicine.

### **Porphyria**

In patients with porphyria, DOM-CLONAZEPAM has to be used with care because it may have a porphyrogenic effect.

### **Dependence and Tolerance**

With long-term DOM-CLONAZEPAM treatment at the therapeutic doses, development of physical and psychic dependence may occur (see ADVERSE REACTIONS). The risk of dependence increases with dose and duration of treatment; it is also greater in patients with a medical history of alcohol and/or drug abuse.

Once physical dependence has developed, abrupt termination of treatment will be accompanied by withdrawal symptoms. The possibility that such effects may also occur following short-term use, especially at high doses, or if the daily dose is reduced rapidly or abruptly discontinued should be considered. Symptoms of withdrawal include tremor, sweating, agitation, sleep disturbances and anxiety, headaches, muscle pain, extreme anxiety, tension, restlessness, confusion, and irritability. In severe cases the following symptoms may occur: derealization, depersonalization, hyperacusis, numbness and tingling of the extremities, hypersensitivity to light, noise and physical contact or hallucinations. Since the risk of withdrawal symptoms is greater after abrupt discontinuation of treatment, abrupt withdrawal of the drug should be avoided and treatment - even if only of short duration - should be terminated by gradually reducing the daily dose.

### **Driving and Hazardous Activities**

Patients receiving DOM-CLONAZEPAM should be cautioned against engaging in hazardous occupations requiring complete mental alertness, such as operating machinery or driving a motor vehicle. Sedation, amnesia and impaired muscular function are effects of benzodiazepines that can adversely affect the ability to drive or operate machinery. This effect is increased if the patient has had alcohol.

Driving, operating machinery and other hazardous activities should be avoided altogether or at least during the first few days of treatment. The decision on this question rests with the patient's physician and should be based on the patient's response to treatment and the dosage involved. They also should be warned against the concomitant use of alcohol and other CNS depressant drugs.

### **Hepatic**

The safety and efficacy of clonazepam in patients with hepatic impairment has not been studied (See DOSAGE AND ADMINISTRATION, ACTION AND CLINICAL PHARMACOLOGY).

### **Renal**

The safety and efficacy of clonazepam in patients with renal impairment has not been studied. Clonazepam and its metabolites are excreted by the kidneys; to avoid excessive accumulation, caution should be exercised in the administration of the drug to patients with impaired renal function (See DOSAGE AND ADMINISTRATION, ACTION AND CLINICAL PHARMACOLOGY).

### **Respiratory**

Respiratory depression may occur following administration of clonazepam. This effect may be aggravated by pre-existing airway obstruction or brain damage or if other medications which depress respiration have been given. As a rule, this effect can be avoided by careful adjustment of the dose to individual requirements.

Treatment with clonazepam should be instituted with caution in patients with chronic respiratory diseases (See CONTRAINDICATIONS).

Hypersecretion in the upper respiratory passages has at times been a troublesome adverse reaction during clonazepam therapy, especially in small mentally retarded children who ordinarily have difficulty handling secretions. Therefore special attention must be paid to maintaining patency of the airways.

### **Carcinogenesis**

See TOXICOLOGY.

### **Special Populations**

#### **Pregnant Women:**

In a reproductive study in rabbits, administration of clonazepam was associated with an increased incidence of cleft palate and other anomalies at two dose levels (see TOXICOLOGY: Teratogenicity).

Reports indicate an association between the use of anticonvulsant drugs and an elevated incidence of birth defects in children born to epileptic women taking such medication during pregnancy. The incidence of congenital malformations in the general population is regarded to be approximately 2%; in children of treated epileptic women this incidence may be increased two to three-fold. The increase is largely due to specific defects, e.g., congenital malformations of the heart, and cleft lip and/or palate. Nevertheless, the great majority of mothers receiving anticonvulsant medications deliver normal infants.

Data are more extensive with respect to diphenylhydantoin and phenobarbital, but these drugs are also the most commonly prescribed anticonvulsants. Some reports indicate a possible similar association with the use of other anticonvulsant drugs, including trimethadione and paramethadione. However, the possibility also exists that other factors, e.g., genetic predisposition or the epileptic condition itself may contribute to or may be mainly responsible for the higher incidence of birth defects.

The preceding considerations should be borne in mind and clonazepam should be used in women of childbearing potential only when the expected benefits to the patient warrant the possible risk to a fetus. Administration of high doses in the last trimester of pregnancy or during labour can cause irregularities in the heartbeat of the unborn child and hypothermia, hypotonia, mild respiratory depression and poor feeding in the neonate. Moreover, infants born to mothers who took benzodiazepines chronically during the later stages of pregnancy may have developed physical dependence and may be at some risk for developing withdrawal symptoms in the postnatal period.

Anticonvulsant drugs should not be discontinued in patients in whom the drug is administered to prevent major seizures, because of the strong possibility of precipitating status epilepticus with attendant hypoxia and risk to both the mother and the unborn child. With regard to drugs given for minor seizures, the risk of discontinuing medication prior to or during pregnancy should be weighed against the risk of congenital defects in the particular case and with the particular family history.

Epileptic women of childbearing age should be encouraged to seek professional counsel and should report the onset of pregnancy promptly to their physician. Where the necessity for continued use of anti-epileptic medication is in doubt, appropriate consultation might be indicated.

**Nursing Women:** Although the active ingredient of clonazepam has been found to pass into the maternal milk in small amounts only, mothers receiving DOM-CLONAZEPAM should not breast-feed their infants.

**Pediatrics (< 5 years of age):** Because of the possibility that adverse effects on physical or mental development of the child could become apparent only after years, a risk-benefit consideration of the long-term use of DOM-CLONAZEPAM is important in pediatric patients.

**Geriatrics:** There is no clinical trial experience with clonazepam in seizure disorder patients 65 years of age and older. In general elderly patients should be started on low doses of DOM-CLONAZEPAM and observed closely.

### **Monitoring and Laboratory Tests**

Periodic liver function tests and blood counts are recommended during long-term therapy with clonazepam.

## **ADVERSE REACTIONS**

### **Adverse Drug Reaction Overview**

#### **Most Frequent Adverse Reactions:**

The most frequently occurring adverse reactions of clonazepam are referable to CNS depression. Experience to date has shown that drowsiness has occurred in approximately 50% of patients and

ataxia in approximately 30%. In some cases, these may diminish with time. Behaviour problems have been noted in approximately 25% of patients.

Somnolence, slowed reaction, muscular hypotonia, muscle weakness, dizziness, ataxia occur relatively frequently and are usually transient and generally disappear spontaneously in the course of the treatment or on reduction of the dosage. They can be partially prevented by increasing the dose slowly at the start of treatment.

**Serious and Important Adverse Reactions:**

An increased risk for falls and fractures has been recorded in elderly benzodiazepine users.

Allergic reactions and a very few cases of anaphylaxis have been reported to occur with benzodiazepines.

Release of hostility and other paradoxical effects such as irritability, excitability, restlessness, agitation, aggressiveness, delusion, rages, nightmares, hallucinations, psychosis, inappropriate behaviour and other adverse behavioural effects are known to occur with the use of benzodiazepines.

If these occur, use of the drug should be discontinued.

Anterograde amnesia may occur using therapeutic doses of benzodiazepines, the risk increasing with higher doses. Effects of anterograde amnesia may be associated with inappropriate behaviour.

Chronic use (even at therapeutic doses) may lead to the development of physical dependence: discontinuation of the therapy may result in withdrawal or rebound phenomena (See WARNINGS-and PRECAUTIONS-Dependence and Tolerance).

**Post Market Adverse Drug Reactions**

Other adverse reactions listed by system, are:

**Body as Whole:** Fever, coated tongue.

**Cardiovascular System:** Palpitations, cardiac failure including cardiac arrest.

**Digestive System:** increased salivation, nausea, vomiting, anorexia, constipation, diarrhea, encopresis, dry mouth, increased appetite, abdominal pain, epigastric symptoms and hepatomegaly.

**Endocrine System:** gynecomastia, isolated cases of reversible development of premature secondary sexual characteristics in children (incomplete precocious puberty).

**Hemic and Lymphatic System:** Anemia, leukopenia (WBC below 4000/cu mm), decreased platelet count, eosinophilia and lymphadenopathy.



**Metabolic and Nutritional Disorders:** transient elevations of transaminase and alkaline phosphatase, weight gain or loss, dehydration.

**Musculoskeletal System:** Low back pain.

**Nervous System:** Nystagmus, dysarthria, vertigo, insomnia, tiredness, lassitude, dysdiadokinesis, aphonia, withdrawal and coma. Isolated reports of akinesia, hemiparesis, tremor, headache and choreiform movements have been received. Minor changes in EEG patterns, specifically low-voltage fast activity. With certain forms of epilepsy, an increase in the frequency of seizures during long-term treatment is possible.

Impaired concentration, restlessness, confusional state, disorientation, depression, paradoxical reactions (excitability, irritability, aggression, agitation, nervousness, hostility, anxiety, sleep disturbances, nightmares and vivid dreams), loss of libido.

**Respiratory System:** Hypersecretion in the upper respiratory passages, rhinorrhea, dyspnea and respiratory depression.

**Skin and Appendages:** nonspecific erythematous, papular and maculopapular rashes, swelling of the face and eyelids, urticaria, pigmentation changes and pruritus. Hirsutism and transient hair loss have also been reported, but drug relationship has not been established.

**Special Senses:** Particularly in long-term or high-dose treatment, reversible disorders of vision (diplopia) may occur.

**Urogenital System:** Rare instances of dysuria, nocturia, urinary incontinence, urinary retention and enuresis.

## DRUG INTERACTIONS

### **Overview**

Simultaneous administration of several anticonvulsant drugs may be considered with clonazepam, however, it should be borne in mind that the use of multiple anti-convulsants may result in an increase of central depressant adverse effects. In addition, the dosage of each drug may be required to be adjusted to obtain the optimal effect.

A paradoxical increase in seizure activity or the appearance of new seizure types has occurred in a very few patients during treatment with clonazepam. When used in patients in whom several different types of seizures coexist, clonazepam may increase the incidence or precipitate the onset of generalized tonic-clonic seizures (grand mal). These phenomena may require the addition of appropriate anticonvulsants or an increase in their dosage. The concomitant use of valproic acid and clonazepam may produce absence status.

Hepatic cytochrome P-450 3A4 is implicated in the metabolism of clonazepam to pharmacologically inactive metabolites. Therefore, concomitant use of drugs that affect the activity of cytochrome P-450 3A4 may alter the pharmacokinetics of clonazepam.

### **Drug-Drug Interactions**

*Pharmacokinetic Drug-Drug Interactions (DDI):* The antiepileptic drugs phenytoin, phenobarbital, carbamazepine, and valproate may increase the clearance of clonazepam thereby decreasing the plasma concentrations of the latter during combined treatment.

Clonazepam itself does not induce the enzymes responsible for its own metabolism.

The selective serotonin reuptake inhibitors sertraline and fluoxetine do not affect the pharmacokinetics of clonazepam when administered concomitantly.

*Pharmacodynamic Drug-Drug Interactions(DDI):*

#### *CNS-acting drugs*

Epileptic patients being treated with DOM-CLONAZEPAM must under no circumstances consume alcohol since it may alter the effect of the drug, reduce the efficacy of treatment or produce unwanted effects. Enhanced effects on sedation, respiration and hemodynamics may occur when clonazepam is coadministered with any centrally acting depressants including alcohol, narcotics, narcotic analgesics, muscle-relaxants, barbiturates, non-barbiturate hypnotics, anxiolytics/sedatives, antihistamines, phenothiazines, thioxanthene and butyrophenone classes of antipsychotic agents, monoamine oxidase inhibitors, tricyclic antidepressants, and anticonvulsants (see WARNINGS & PRECAUTIONS, Concomitant use of Alcohol/CNS Depressants; OVERDOSAGE).

In combination therapy with centrally-acting medications, the dosage of each drug must be adjusted to achieve the optimum effect. Because of the potentiation of effects that might occur, patients should be advised against the simultaneous use of other CNS depressant drugs and should be cautioned not to take alcohol during the administration of clonazepam.

### **Drug-Food Interactions**

Interactions with food have not been established. Grapefruit juice decreases the activity of cytochrome P-450 3A4, which is implicated in the metabolism of clonazepam, and may contribute to increased plasma levels of the drug.

### **Drug-Herb Interactions**

Interactions with herbal products have not been established.

### **Drug-Laboratory Interactions**

Interactions with laboratory tests have not been established.

## DOSAGE AND ADMINISTRATION

### **Dosing Considerations**

Dosage of DOM-CLONAZEPAM (clonazepam) is essentially individual and depends above all on the age of the patient. Dosage must be determined in each patient according to clinical response and tolerance. The use of multiple anticonvulsants may result in an increase of depressant adverse effects. This should be borne in mind whenever DOM-CLONAZEPAM is added to an already existing anticonvulsant regimen.

### **Recommended Dose and Dosage Adjustment**

**Children:** In order to minimize drowsiness, the initial dose for infants and children (up to 10 years of age or 30 kg of body weight) should be between 0.01 and 0.03 mg/kg/day and should not exceed 0.05 mg/kg/day given in two or three divided doses. Dosage should be increased by no more than 0.25 to 0.50 mg every third day until a maintenance dose of 0.1 to 0.2 mg/kg of body weight has been reached, unless seizures are controlled or side effects preclude further increase.

**Adults:** The initial dose for adults should not exceed 1.5 mg/day divided into three doses. Dosage may be increased in increments of 0.5 to 1 mg every three days until seizures are adequately controlled or until side effects preclude any further increase. Maintenance dosage must be individualized for each patient depending upon response. A recommended maintenance dose for adults is 8 to 10 mg/day in three divided doses. Dosages in excess of 20 mg/day should be administered with caution.

Whenever possible, the daily dose should be divided into three equal doses. If doses are not equally divided, the larger dose should be given before retiring.

**Geriatrics:** There is no clinical trial experience with clonazepam in seizure disorder patients 65 years of age and older. In general elderly patients should be started on low doses of DOM-CLONAZEPAM and observed closely.

### **Special Populations**

**Renal Impairment:** The safety and efficacy of clonazepam in patients with renal impairment has not been studied, however based on pharmacokinetic considerations no dose adjustment is required in these patients (see ACTION AND CLINICAL PHARMACOLOGY: Special Populations and Conditions).

**Hepatic Impairment:** The safety and efficacy of clonazepam in patients with hepatic impairment has not been studied. No data are available on the influence of hepatic disease on clonazepam pharmacokinetics (see ACTION AND CLINICAL PHARMACOLOGY: Special Populations and Conditions).

## OVERDOSAGE

**Symptoms:** Benzodiazepines commonly cause drowsiness, ataxia, dysarthria and nystagmus. Overdose of clonazepam is seldom life-threatening if the drug is taken alone, but may lead to areflexia, apnea, hypotension, cardiorespiratory depression and coma. Coma, if it occurs, usually lasts a few hours but it may be more protracted and cyclical, particularly in elderly patients. Benzodiazepine respiratory depressant effects are more serious in patients with respiratory disease.

Benzodiazepines increase the effects of other central nervous system depressants, including alcohol.

### *Treatment*

In managing overdose, consider the possibility of multiple drug involvement.

Monitor the patient's vital signs and institute supportive measures as indicated by the patient's clinical state. In particular, patients may require symptomatic treatment for cardiorespiratory effects or central nervous system effects.

Further absorption should be prevented using an appropriate method e.g. treatment within 1-2 hours with activated charcoal. If activated charcoal is used, airway protection is imperative for drowsy patients. In case of mixed ingestion, gastric lavage may be considered, however not as a routine measure. Induction of vomiting is not generally recommended.

As in overdose with other benzodiazepines, dialysis is of no known value in clonazepam overdose.

If CNS depression is severe consider the use of flumazenil (ANEXATE®) a benzodiazepine receptor antagonist. The following should be kept in mind when flumazenil is used in the treatment of benzodiazepine overdose:

- Flumazenil should only be administered under closely monitored conditions. In view of the short half life (about 1 hour) and duration of action of flumazenil, and the possible need for repeat doses, the patient should be closely monitored until all possible central benzodiazepine effects (e.g., re-sedation) have subsided.
- Particular caution is necessary when using flumazenil in cases of multiple drug overdose, since the toxic effects (cardiac arrhythmias and/or convulsions) of other psychotropic drugs, especially cyclic antidepressants, may increase as the effects of benzodiazepines subside. Flumazenil is contraindicated in patients who are showing signs of serious cyclic antidepressant overdose.

**Warning:** The benzodiazepine receptor antagonist flumazenil is not indicated in patients with epilepsy who have been treated with benzodiazepines. Antagonism of the benzodiazepine effect in such patients may provoke seizures.

Refer to the prescribing information for flumazenil, for further information on the correct use of this drug.

For management of a suspected drug overdose, contact your regional Poison Control Centre immediately.

## **ACTION AND CLINICAL PHARMACOLOGY**

### **Mechanism of Action**

Clonazepam has pharmacological properties characteristic of the benzodiazepine class of drugs. Clonazepam has sedative, hypnotic and anticonvulsant properties. As an anticonvulsant it is useful in the management of minor motor seizures (myoclonic seizures) and may be of some value in selected patients with absence spells (petit mal) who have failed to respond satisfactorily to the succinimides. Clonazepam is capable of suppressing the spike and wave discharge in absence seizures (petit mal) and decreasing the frequency, amplitude, duration and spread of discharge in minor motor seizures.

### **Absorption:**

Clonazepam is rapidly and almost completely absorbed after oral administration of clonazepam tablets. Peak plasma concentrations of clonazepam are reached in 1-4 hours. The absorption half-life is around 25 minutes. The absolute bioavailability is 90%.

Plasma concentrations of clonazepam at steady state for a once-daily dosage regimen are 3-fold higher than those after a single oral dose; the predicted accumulation ratios for two times and three times daily regimens are 5 and 7, respectively. Following multiple oral doses of 2 mg three times daily steady-state pre-dose plasma concentrations of clonazepam averaged 55 ng/ml. The plasma concentration-dose relationship of clonazepam is linear. The target anticonvulsant plasma concentrations of clonazepam range from 20 to 70 ng/ml.

### **Distribution:**

Clonazepam distributes very rapidly to various organs and body tissues with preferential uptake by brain structures.

The distribution half-life is approximately 0.5-1 hour. The volume of distribution is 3 l/kg. The plasma protein binding is 82-86%.

### **Metabolism:**

Clonazepam is extensively metabolized by reduction to 7-amino-clonazepam and by N-acetylation to 7-acetamino-clonazepam. Hydroxylation at the C-3 position also occurs. Hepatic cytochrome P-450 3A4 is implicated in the nitroreduction of clonazepam to pharmacologically inactive metabolites.

The metabolites are present in urine both as free and conjugated (glucuronide and sulphate) compounds.

**Elimination:**

The mean elimination half-life is 30-40 hours. The clearance is 55 mL/min.

50-70% of the dose is excreted in the urine and 10-30% in feces as metabolites. The urinary excretion of unchanged clonazepam is usually less than 2% of the administered dose.

The elimination kinetics in children are similar to those observed in adults.

**Special Populations and Conditions**

**Neonates:** The elimination half-life and clearance values in neonates are of the same order of magnitude of those reported in adults.

**Geriatrics:** The pharmacokinetics of clonazepam in the elderly has not been established.

**Hepatic Failure:** The influence of hepatic disease on clonazepam pharmacokinetics has not been investigated.

**Renal Failure:** Renal disease does not affect the pharmacokinetics of clonazepam. Based on pharmacokinetic criteria, no dose adjustment is required in patients with renal disease.

**STORAGE AND STABILITY**

Store between 15 and 30°C. Keep in a tightly closed, light resistant container.

**DOSAGE FORMS, COMPOSITION AND PACKAGING**

**0.5 mg** Each round, orange, biconvex tablet, debossed with “Clonazepam” on one side and “pms” score “0.5” on the other side contains 0.5 mg of clonazepam and the following non medicinal ingredients: corn starch, FD&C Yellow # 6 lake, lactose, magnesium stearate, and microcrystalline cellulose.

Available in HDPE bottles of 100, 500 and 1000 tablets.

**2 mg** Each round, white, biplane tablet with bevelled edges, debossed with “Clonazepam” on one side and “2.0” under score line on the other side contains 2.0 mg of clonazepam and the following non medicinal ingredients: corn starch, lactose, magnesium stearate, and microcrystalline cellulose.

Available in HDPE bottles of 100, 500 and 1000 tablets.

## PART II: SCIENTIFIC INFORMATION

### PHARMACEUTICAL INFORMATION

#### Drug Substance

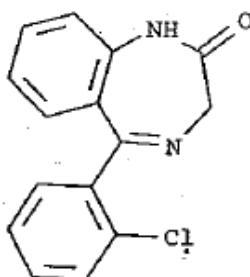
Proper Name: clonazepam

Chemical name: 5-(2-chlorophenyl)-1,3-dihydro-7-nitro-2H-1,4-benzodiazepin-2-one.

Molecular formula:  $C_{15}H_{10}ClN_3O_3$

Molecular mass: 419.46 g/mol

Structural formula:



#### Description

Light yellow powder, having a faint odor. Melts at about 239°. Insoluble in water; sparingly soluble in acetone and in chloroform; slightly soluble in alcohol and in ether.

## CLINICAL TRIALS

Single oral doses of clonazepam to healthy volunteers gives maximum blood levels of drug, in 1 to 3 hours. The half-life of the parent compound ranges from approximately 18 to 50 hours. The major route of excretion of clonazepam is the urine.

### Comparative Bioavailability Studies

A comparative bioavailability study was performed using 18 normal human male volunteers. The rate and extent of absorption after a single oral dose of the test product DOM-CLONAZEPAM 2 mg manufactured by Dominion Pharmacal vs. the reference product Rivotril® 2 mg manufactured by Hoffman Laroche was measured and compared. The results can be summarized as follows:

### SUMMARY OF COMPARATIVE BIOAVAILABILITY DATA

|   |  |  |  |  |
|---|--|--|--|--|
| <b>Clonazepam<br/>(1 x 2 mg)<br/>From measured data</b><br><br><b>Geometric Mean<br/>Arithmetic Mean (CV %)</b> |  |  |  |  |
|---|--|--|--|--|

| Parameter                     | Test*                    | Reference†               | % Ratio of Geometric Means | Confidence Interval (90%) |
|-------------------------------|--------------------------|--------------------------|----------------------------|---------------------------|
| AUC <sub>T</sub><br>(ng.h/mL) | 392.02<br>398.22 (17.48) | 364.95<br>372.26 (19.86) | 107.42%                    | 99.78 – 115.63            |
| AUC<br>(ng.h/mL)              | 459.71<br>466.40 (16.84) | 428.70<br>434.63 (16.70) | 107.23%                    | 100.45-114.47             |
| C <sub>MAX</sub><br>(ng/mL)   | 10.56<br>10.71 (15.90)   | 10.03<br>10.21 (19.19)   | 105.33%                    | 95.57-116.08              |
| T <sub>MAX</sub> §<br>(h)     | 2.57 (1.81)              | 2.56 (2.31)              |                            |                           |
| T <sub>1/2</sub> ¶<br>(h)     | 35.93 (7.32)             | 34.06 (7.06)             |                            |                           |

\* DOM-CLONAZEPAM 2 mg tablets, Dominion Pharmacal Inc, Canada

† Rivotril® 2 mg tablets, Hoffman-La Roche Ltd, Canada

§ Expressed as the arithmetic mean (CV%) only.

¶ Expressed as the arithmetic mean (CV%) only.



A comparative bioavailability study of DOM-CLONAZEPAM 0.5 mg tablets was performed versus Rivotril® 0.5 mg tablets, Hoffman-La Roche Ltd., Canada, 2 mg (4 x 0.5 mg) oral administration to 26 male volunteers in the fasting state. Pharmacokinetic and bioavailability data were measured and the results are summarized in the following table:

**SUMMARY TABLE OF THE COMPARATIVE BIOAVAILABILITY DATA**

|  |
|--|
| <p><b>Clonazepam<br/>(4 x 0.5 mg)<br/>From measured data</b></p> <p><b>Geometric Mean<br/>Arithmetic Mean (CV %)</b></p> |
|--|

| Parameter                         | Test*                   | Reference†              | % Ratio of Geometric Means | Confidence Interval (90%) |
|-----------------------------------|-------------------------|-------------------------|----------------------------|---------------------------|
| AUC <sub>0-72h</sub><br>(ng.h/mL) | 272.35<br>276.82 (17.1) | 261.83<br>265.88 (18.0) | 104.0                      | (98.5 - 109.8)            |
| AUC <sub>T</sub><br>(ng.h/mL)     | 328.96<br>337.02 (20.2) | 302.50<br>308.80 (20.4) | 108.8                      | (101.7 - 116.3)           |
| AUC <sub>∞</sub><br>(ng.h/mL)     | 410.96<br>421.20 (21.0) | 375.80<br>382.72 (19.1) | 109.4                      | 101.2 - 118.1)            |
| C <sub>MAX</sub><br>(units)       | 8.64<br>8.74 (16.6)     | 7.85<br>7.97 (18.4)     | 110.1                      | (104.5 - 115.9)           |
| T <sub>MAX</sub> §<br>(h)         | 1.94 (0.73)             | 2.60 (1.42)             |                            |                           |
| T <sub>½</sub>   <br>(h)          | 47.77 (11.7)            | 42.39 (16.8)            |                            |                           |

\* DOM-CLONAZEPAM 0.5 mg tablets, Dominion Pharmacal Inc, Canada

† Rivotril® 0.5 mg tablets, Hoffman-La Roche Ltd, Canada

§ Expressed as the arithmetic mean (CV%) only.

|| Expressed as the arithmetic mean (CV%) only.

## DETAILED PHARMACOLOGY

The pharmacological profile of clonazepam is the same as that of other anxiolytic sedative benzodiazepines. Its basic anticonvulsive properties are also similar to those of other diazepamines.

### **Relative Potency of Clonazepam and Other Anticonvulsants (Experimental Tests)**

The following table gives an indication of the relative potency of clonazepam and other anticonvulsants in various experimental tests in animals.

#### **Convulsant Test Oral ED<sub>50</sub> Values (mg/kg) in Mice and Humans**

| <b>Drug</b>      | <b>Max. Human Therapeutic Dose (mg/kg)</b> | <b>Metrazol Seizures</b> | <b>Thiosemicarbazide Seizures</b> | <b>30% Strychnine Threshold</b> | <b>Maximum Electroshock</b> |
|------------------|--|--------------------------|-----------------------------------|---------------------------------|-----------------------------|
| Clonazepam       | 0.40                                       | 0.08 - 0.16              | 0.73                              | 2.1                             | 8.4                         |
| Diazepam         | 0.43                                       | 0.8 - 1.4                | 3.4                               | 6.2                             | 9.0                         |
| Chlordiazepoxide | 1.43                                       | -                        | 27.0                              | 22.2                            | 17.2                        |
| Phenobarbital    | 8.5  | 8.0 - 27.0               | 63                                | 37.2                            | 7.3                         |
| Trimethadione    | 25.7                                       | 300                      | 770                               | -                               | 490                         |
| DPH              | 7.7  | -                        | 7800                              | 7300                            | 8.7                         |

Clonazepam is effective in reducing photomyoclonic responses in baboons in doses under 0.5 mg/kg i.m. However, seizures evoked by local application of benzylpenicillin or strychnine do not respond well to systemic administration of clonazepam. Other CNS effects noted in several species at varying doses include taming, disinhibitory, sedative, ataxic, and hypnotic effects.

Blood pressure in dogs is lowered and vascular responses to serotonin and noradrenaline are inhibited by clonazepam in doses between 1 and 4 mg/kg i.v. There is a slight myocardial depressant action at these doses. Other pharmacologic effects occur only at higher doses in which gross CNS depressant effects are observed.

Metabolic pathways are similar in several species and the chief metabolites, 7-amino and 7-acetyl amino derivatives, have been isolated in urine of rats, dogs and humans. Hydroxylation also occurs as a prominent metabolic pathway. Metabolites are excreted primarily in urine, approximately 50% of an oral dose is excreted within seven days. Excretion of the drug plus metabolites increases as the dose increases.

## TOXICOLOGY

### **Acute Toxicity:**

The following LD<sub>50</sub> values have been calculated for clonazepam:

| Species       | Dose (mg/kg) and Route |      |          |
|---------------|------------------------|------|----------|
|               | Oral                   | i.p. | i.v.     |
| Mouse         | >4000                  | >800 | 2.85±0.1 |
| Rat (adult)   | >4000                  | -    | -        |
| Rat (neonate) | 550±120                | -    | -        |
| Rabbit        | >2000                  | -    | -        |

Signs of toxicity include decreased motor activity, ataxia, piloerection and tremors.

**Chronic Toxicity:** Rats were fed clonazepam in the diet for 18 months in concentrations corresponding to 5, 20 and 50 mg/kg/day. No gross drug-related toxicity was evident. Slight and transient elevations in liver function tests appeared in high dose animals corresponding to increases in liver weights, but these findings were not accompanied by histologic evidence of liver damage.

A study in dogs was conducted in which animals received clonazepam in doses of 3, 10 and 30 mg/kg/day for 12 months. Weight gain was reduced in mid- and high-dose animals compared to controls. The following significant changes in laboratory values were noted: a decrease in hemoglobin and hematocrit values in mid- and high-dose animals, a decreased albumin/globulin ratio due to decreased albumin and increased globulins in high-dose animals, increased alkaline phosphatase and bilirubin values in high-dose animals. There was a significant increase in liver weight in high-dose animals.

**Carcinogenicity:** No 2-year carcinogenicity studies have been conducted with clonazepam. However, in an 18-month chronic study in rats no treatment-related histopathological changes were seen up to the highest tested dose of 300 mg/kg/day.

**Mutagenicity:** Genotoxicity tests using bacterial systems with *in vitro* or host mediated metabolic activation did not indicate a genotoxic liability for clonazepam.

**Impairment of Fertility:** Studies assessing fertility and general reproductive performance in rats showed a reduced pregnancy rate and impaired pup survival at doses of 10 and 100 mg/kg/day.

**Teratogenicity:** No adverse maternal or embryo-fetal effects were observed in either mice or rats following administration of oral clonazepam during organogenesis, at doses of up to 20 or 40 mg/kg/day, respectively.

In several rabbit studies following doses of clonazepam of up to 20 mg/kg/day, a low, non-

dose-related incidence of a similar pattern of malformations (cleft palate, open eyelids, fused sternebrae and limb defects) was observed.

## REFERENCES

### Preclinical

1. Blum JE, et al. Pharmakologie und toxikologie des antiepileptikums clonazepam. *Arzneimittel-Forschung* 1973;23:377-89.
2. Guerrero-Figueroa R, et al. Effects of two benzodiazepine derivatives on cortical and subcortical epileptogenic tissues in the cat and monkey. *Curr Therap Res* 1969;11:27-50.
3. Schallek W, et al. Recent developments in the pharmacology of the benzodiazepines, advances in pharmacology and chemotherapy. Academic Press, Inc., New York and London 1972;10:132-7.
4. Stark LG, et al. The anticonvulsant effects of phenobarbital, diphenylhydantoin and two benzodiazepines in the baboon *Papio Papio*. *J Pharmacol Exp Therap* 1970;173:125-32.

### Clinical

5. Barnett AM. Treatment of epilepsy with clonazepam (Ro 5-4023). *S.A. Medical Journal* 1973;47:1683-6.
6. Bergamini L et al. Elektroenzephalographische und klinische bewertung des neuen benzodiazepin Ro 5-4023. *EEG-EMG* 1970;1:182-8.
7. Browne TR. Clonazepam: a review of a new anticonvulsant drug. *Arch Neurol* 1976;33:326-32.
8. Carson MJ, et al. Treatment of minor motor seizures with clonazepam. *Develop Med Child Neurology* 1975;17:306-10.
9. Dreifus FE, et al. Serum clonazepam concentrations in children with absence seizures. *Neurology* 1975;25:255-8.
10. Fazio C, et al. Treatment of epileptic seizures with clonazepam. *Arch Neurol* 1975;32:304-7.
11. Gastaut H. The exceptional antiepileptic properties of new benzodiazepine, presented at 4<sup>th</sup> Intern Congr Neurol, New York, September 20-27, 1969.
12. Gastaut H. Propriétés anti-épileptiques exceptionnelles d'une benzodiazépine nouvelle le Ro 5-4023. *Vie Med* 1970;51:5175-88.
13. Hanson RA, et al. A new anticonvulsant in the management of minor motor seizures. *Develop Med Child Neurol* 1972;14:3-14.

14. Hollister LE, et al. Dose-ranging studies of clonazepam in man. *Psychopharmacology Communications* 1975;1:89-92.
15. Hooshmand H. Intractable seizures; treatment with a new benzodiazepine anticonvulsant. *Arch Neurol* 1972;27:205-8.
16. Lison MP, et al. Estudo Clinico - Electrencefalografico longitudinal em pacientes epilépticos tratados. Com Ro. 5-4023, *Arquiv Neurophysiquiat* 1970;28:25-36.
17. Mekkelsen B, et al. A clinical study of benzodiazepine Ro 5-4023 (clonazepam) in the treatment of epilepsy. *Acta Neurol Scand* 1973;49:91-6.
18. Negrin P, et al. Antiepileptic properties of Ro 5-4023 by mouth. Report of 40 cases. *Electroenceph Clin Neurophysiol* 1971;31:528-34.
19. Rose SW, et al. Serum clonazepam concentrations in children with absence seizures. *Neurology* 1974;24:386-90.
20. Sjö O, et al. Pharmacokinetics and side effects of clonazepam and its 7-Amino-Metabolite in man. *Eur J Clin Pharmacol* 1975;8:249-54.
21. Turner M, et al. Clinical EEG evaluation of a new benzodiazepine derivative (Ro 5-4023) by oral administration in epileptic patients using the double-blind technique. *Electroenceph Clin Neurophysiol* 1971;31:628-30.
22. Product Monograph: Rivotril, Hoffman Laroche, Date of revision: July 8, 2009, Control Number 127819

**PART III: CONSUMER INFORMATION****<sup>T/C</sup>DOM-CLONAZEPAM  
(Clonazepam Tablets, USP)**

**This leaflet is part III of a three-part "Product Monograph" published when DOM-CLONAZEPAM was approved for sale in Canada and is designed specifically for Consumers.**

**Please read this information before you start to take your medicine. Keep this leaflet until you have finished all your tablets, as you may need to read it again. If you are helping someone else to take DOM-CLONAZEPAM, read this leaflet before you give the first tablet.**

**This leaflet is a summary and will not tell you everything about DOM-CLONAZEPAM. Contact your doctor or pharmacist if you have any questions about the drug.**

**ABOUT THIS MEDICATION****What the medication is used for:**

DOM-CLONAZEPAM is used for the treatment of certain types of seizures.

**What it does:**

DOM-CLONAZEPAM contains the active ingredient clonazepam, which belongs to a group of medicines known as benzodiazepines. DOM-CLONAZEPAM has anticonvulsant properties which help to manage seizures.

**When it should not be used:**

- If you are allergic to the group of medicines known as benzodiazepines (examples: diazepam, chlordiazepoxide, bromazepam, or flurazepam)
- If you are allergic to any of the ingredients it contains (see 'What the nonmedicinal ingredients are')
- If you suffer from lung disease.
- If you have a liver condition.
- If you have glaucoma.
- If you have myasthenia gravis.

**What the medicinal ingredient is:**

Clonazepam.

**What the nonmedicinal ingredients are:**

Corn starch, FD&C Yellow # 6 Lake (0.5 mg), magnesium stearate, and microcrystalline cellulose

**What dosage forms it comes in:**

Tablets: 0.5 mg and 2 mg

during the first few days of treatment. This effect of DOM-CLONAZEPAM may be made worse if you take alcoholic drinks. If you increase your dose or change the timings of when you take your medication this may also modify your reactions.

- You must not consume alcohol or other drugs that affect your central nervous system while taking DOM-CLONAZEPAM as this may provoke epileptic seizures.
- Always contact your doctor before stopping or reducing your dosage of DOM-CLONAZEPAM, as suddenly stopping treatment or a large decrease in dose can cause reappearance of seizures and withdrawal symptoms.
- Benzodiazepines such as DOM-CLONAZEPAM have produced dependence (addiction) and withdrawal symptoms can occur when treatment is stopped suddenly. The risk of dependence (addiction) increases with higher doses and longer duration of treatment. Symptoms of withdrawal may include shaking, sweating, sleep disturbances, agitation/restlessness, headache, muscle pain, anxiety, confusion, and irritability. In severe cases of withdrawal, symptoms may include numbness and tingling of the extremities, hallucinations (see or hear things that are not there), increased sensitivity to light, noise and physical contact.
- An increased risk of falls and fractures have been reported in elderly people who take benzodiazepines such as DOM-CLONAZEPAM.
- Memory loss may occur when DOM-CLONAZEPAM is used at therapeutic doses.
- Do not take this medicine if you are pregnant, or might become pregnant, unless advised by your doctor. Contact your doctor if you think you may be pregnant, or are intending to become pregnant.
- DOM-CLONAZEPAM passes into breast milk. Therefore, if you are breast feeding, this medicine should be avoided. Your doctor will discuss this with you.

**BEFORE you use DOM-CLONAZEPAM talk to your doctor or pharmacist if you:**

- Have a lung, liver or kidney condition.
- Have glaucoma.
- Are taking or plan on taking ANY other drugs (including herbal preparations, drugs you purchase without prescriptions, and those not prescribed by your doctor).
- Regularly drink alcohol or use recreational drugs.
- Suffer from a form of incoordination of the muscles called spinal or cerebellar ataxia.
- Have a history of depression and/or suicide attempts.
- Have rare hereditary problems of galactose intolerance.

**WARNINGS AND PRECAUTIONS**

- DOM-CLONAZEPAM may affect your ability to be alert. Driving, operating machinery and other hazardous activities should therefore be avoided altogether or at least

**INTERACTIONS WITH THIS MEDICATION**

**Tell your doctor if you are taking any other medicines including any that you have bought from a pharmacy,**

**supermarket or health food store without a prescription.**

- Some medicines may interfere with DOM-CLONAZEPAM. These medicines include:
- medicines to control seizures
- narcotics and narcotic pain relievers
- muscle relaxants
- sleeping medication
- medicines to treat your mood, such as monoamine oxidase inhibitors, tricyclic antidepressants, phenothiazines
- phenytoin, phenobarbital, carbamazepine, and valproate.

These medicines may be affected by DOM-CLONAZEPAM or may affect how well DOM-CLONAZEPAM works. Your doctor or pharmacist can tell you what to do if you are taking any of these medicines.

**If you have not told your doctor about any of the above, tell him/her before you start taking DOM-CLONAZEPAM.**

You must not consume alcohol while taking DOM-CLONAZEPAM as its effects may worsen side effects that some patients experience with DOM-CLONAZEPAM.

Grapefruit juice may increase blood levels of DOM-CLONAZEPAM, therefore you should avoid drinking grapefruit juice while you are taking DOM-CLONAZEPAM.

In case of drug overdose, contact a health care practitioner, hospital emergency department or regional Poison Control Centre immediately, even if there are no symptoms.

**Missed Dose:**

Take the missed dose of DOM-CLONAZEPAM as soon as you remember it. However, if it is almost time for the next dose, skip the missed dose and continue your regular dosing schedule. Do not take 2 doses at the same time.

**SIDE EFFECTS AND WHAT TO DO ABOUT THEM**

Like all medications DOM-CLONAZEPAM can cause some side effects.

For most patients these side effects are likely to be minor and temporary as your body adjusts to the medicine. However, some may be serious. Consult your doctor or pharmacist as soon as you can if you do not feel well while taking DOM-CLONAZEPAM.

The most common side effects are:

- Feeling drowsy or tired, especially at the start of treatment.
- Some muscle weakness and dizziness.
- Increased salivation.

Less common possible side effects are:

- Increase secretion from the lungs may occur. Children should therefore be watched carefully as this might cause difficulties in breathing and/or severe choking and coughing.
- In rare cases changes in your blood and liver may occur and your doctor will monitor for these.
- Increased falls and fractures may occur in elderly patients.

Withdrawal-related side effects:

- With long-term DOM-CLONAZEPAM treatment development of physical and psychological dependence may occur. If treatment is stopped suddenly symptoms of withdrawal may occur, including: shaking, sweating, agitation/restlessness, sleep disturbances, anxiety (possibly extreme), headaches, muscle pain, tension, restlessness, confusion and irritability. In severe cases of withdrawal, symptoms may include numbness and tingling of the extremities, hallucinations (see or hear things that are not there), increased sensitivity to light, noise and physical contact.

**PROPER USE OF THIS MEDICATION**

**Usual dose:**

Always take the tablets exactly as your doctor tells you to. Your doctor will prescribe a suitable dose for you. The dose your doctor prescribes will depend on the nature of your illness, your reaction to the medicine, your age and body weight. The table below shows the different doses that your doctor may prescribe according to your age. Your doctor will start you on an initial low dose and gradually increase it until the desired effect is achieved.

|                                   | <b>Initial Dose</b>                  | <b>Maintenance Dose</b>            |
|-----------------------------------|--------------------------------------|------------------------------------|
| Adults                            | 1.5 mg/day or less in divided doses  | 8-10 mg/day in divided doses       |
| Children (up to 10 years or 30kg) | 0.01-0.03 mg/kg/day in divided doses | 0.1-0.2 mg/kg/day in divided doses |

The total daily dose should be taken as advised by your doctor.

Do not change the prescribed dose yourself. If you think the effect of your medicine is too weak or too strong, talk to your doctor.

Your doctor will advise you when to stop taking the medicine. Your doctor will slowly decrease the dosage as sudden discontinuation of treatment can cause the appearance of withdrawal symptoms.

**Overdose:**



**SERIOUS SIDE EFFECTS, HOW OFTEN THEY HAPPEN AND WHAT TO DO ABOUT THEM**

| Symptom / effect |   | Talk with your doctor or pharmacist |              | Stop taking drug and call your doctor or pharmacist |
|------------------|---|-------------------------------------|--------------|---|
|                  |   | Only if severe                      | In all cases |   |
| Rare             | Unusual behavioural problems (aggression, rage), sudden anxiety or excitement; restlessness, agitation, irritability; hallucinations (see or hear things that are not there) or delusions; severe sleep disturbances, nightmares, inappropriate behaviour   |                                     | √            |   |
|                  | Allergic reactions (red skin, hives, itching, swelling of the lips, face, tongue, throat, trouble breathing, wheezing, shortness of breath, skin rashes, blisters of the skin, sores or pain in the mouth or eyes)  |                                     |              | √<br><b>Immediately.</b>                            |
|                  | Depression. Symptoms may include: difficulty sleeping, changes in weight, feelings of worthlessness, guilt, regret, helplessness or hopelessness, withdrawal from social situations, family gatherings and activities with friends, reduced libido (sex drive), and thoughts of death or suicide. |                                     | √            |   |

*This is not a complete list of side effects. For any unexpected effects while taking DOM-CLONAZEPAM, contact your doctor or pharmacist.*

**REPORTING SUSPECTED SIDE EFFECTS**

You can report any suspected adverse reactions associated with the use of health products to the Canada Vigilance Program by one of the following 3 ways:

- Report online at [www.healthcanada.gc.ca/medeffect](http://www.healthcanada.gc.ca/medeffect)
- Call toll-free at 1-866-234-2345
- Complete a Canada Vigilance Reporting Form and:
  - Fax toll-free to 1-866-678-6789, or
  - Mail to: Canada Vigilance Program  
Health Canada  
Postal Locator 0701D  
Ottawa, ON K1A 0K9

Postage paid labels, Canada Vigilance Reporting Form and the adverse reaction reporting guidelines are available on the MedEffect™ Canada Web site at [www.healthcanada.gc.ca/medeffect](http://www.healthcanada.gc.ca/medeffect).

*NOTE: Should you require information related to the management of side effects, contact your health professional. The Canada Vigilance Program does not provide medical advice.*

**MORE INFORMATION**

**Reminder: This medicine has been prescribed only for you. Do not give it to anybody else. If you have any further questions, please ask your doctor or pharmacist.**

This document plus the full product monograph, prepared for health professionals, can be obtained by contacting the sponsor, DOMINION PHARMACAL at, 1-888-550-6060.

This leaflet was prepared by  
**DOMINION PHARMACAL**  
Montreal Quebec  
H4P 2T4

Last revised: May 31, 2011

**HOW TO STORE IT**

- Keep DOM-CLONAZEPAM in a cool dry place stored at room temperature (15-30°C).
- Keep this medicine out of the reach of children