

## New Zealand Data Sheet

Clopine® can cause agranulocytosis. Its use should be limited to patients:

- with schizophrenia who are non-responsive to or intolerant of classical antipsychotic agents, or with schizophrenia or schizoaffective disorder who are at risk of recurrent suicidal behaviour (*see section 4.1*),
- who have initially normal leukocyte findings (white blood cell count (WBC)  $\geq 3500/\text{mm}^3$  ( $3.5 \times 10^9/\text{L}$ ), and absolute neutrophil counts (ANC)  $\geq 2000/\text{mm}^3$  ( $2.0 \times 10^9/\text{L}$ )),
- and in whom regular white blood cell counts and absolute neutrophil counts can be performed as follows: weekly during the first 18 weeks of therapy, and at least every 4 weeks thereafter throughout treatment. Monitoring must continue throughout treatment and for 4 weeks after complete discontinuation of Clopine.

Prescribing physicians should comply fully with the required safety measures. At each consultation, a patient receiving Clopine should be reminded to contact the treating physician immediately if any kind of infection begins to develop. Particular attention should be paid to flu-like complaints such as fever or a sore throat and to other evidence of infection, which may be indicative of neutropenia.

Clopine must be dispensed in accordance with appropriate local dispensing guidelines. The following conditions apply to the sale, supply and use of Clopine in New Zealand under the consent notice from Medsafe. Douglas draws prescribers, nurses and patients attention to the following criteria:

Clozapine may only be prescribed by:

- Registered medical practitioners as defined in the Health Practitioners Competence Assurance Act 2003 who are certified by the Medical Council of New Zealand as competent in the scope of practice of psychiatry;
- Medical practitioners employed as registrars in the branch of psychiatry who are under the supervision of the persons referred to above;
- Medical officers who:
  - o are in the employment of a District Health Board; and
  - o are under the supervision of persons who are registered medical practitioners as defined in the Health Practitioners Competence Assurance Act 2003 who are certified by the Medical Council of New Zealand as competent in the scope of practice of psychiatry.
- Registered medical practitioners as defined in the Health Practitioners Competence Assurance Act 2003 who are registered with the Medical Council of New Zealand within the vocational scope of practice of general practice. The general practitioner must be continuing the prescribing of clozapine for a specific patient whose illness is well controlled in collaboration, or following consultation, with a Community Mental Health Team.

Persons prescribing clozapine must comply with appropriate local treatment guidelines. The medicine must be dispensed in accordance with appropriate local dispensing guidelines.

Brand swapping between clozapine products is discouraged and should occur on the advice of a clinician.

Prescribers and dispensers should verify that the patient has not previously developed an adverse reaction to clozapine that contraindicates further use of any clozapine containing product.

Clozapine is associated with an increased risk of myocarditis which has, in rare cases, been fatal. The increased risk of myocarditis is greatest in the first two months of treatment. Fatal cases of cardiomyopathy have also been reported rarely. Myocarditis and cardiomyopathy should be suspected in patients who experience persistent tachycardia at rest especially in the first two months of treatment and/or palpitations, arrhythmias, chest pain and other signs and symptoms of heart failure (e.g. unexplained fatigue, dyspnoea, tachypnoea) or symptoms that mimic myocardial infarction. If myocarditis or cardiomyopathy is suspected, Clopine treatment should promptly be stopped and the patient immediately referred to a cardiologist.

## 1. PRODUCT NAME

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Clopine® 25 mg tablet

Clopine® 50 mg tablet

Clopine® 100 mg tablet

Clopine® 200 mg tablet

Clopine® 50 mg/mL oral suspension

## 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

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Each Clopine 25 mg tablet contains 25 mg of clozapine

Each Clopine 50 mg tablet contains 50 mg of clozapine

Each Clopine 100 mg tablet contains 100 mg of clozapine

Each Clopine 200 mg tablet contains 200 mg of clozapine

Each 1 mL of Clopine 50 mg/mL oral suspension contains 50 mg of clozapine

For a full list of excipients, *see section 6.1.*

## 3. PHARMACEUTICAL FORM

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The tablet can be divided into equal doses using the break line.

Clopine 25 mg tablets are round yellow flat bevel-edged tablets engraved "25" over a pressure sensitive break line on one face, the other face plain.

Clopine 50 mg tablets are round yellow flat bevel-edged tablets engraved "50" over a pressure sensitive break line on one face, the other face plain.

Clopine 100 mg tablets are round yellow flat bevel-edged tablets engraved "100" over a pressure sensitive break line on one face, the other face plain.

Clopine 200 mg tablets are oval shaped, yellow tablets engraved "200" on one face, and a pressure sensitive break line on the other face.

Clopine 50 mg/mL oral suspension is a free-flowing yellow suspension

## 4. CLINICAL PARTICULARS

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### 4.1. Therapeutic indications

Treatment-resistant schizophrenia

Clopine is indicated in patients with treatment-resistant schizophrenia, i.e. patients with schizophrenia who are non-responsive to or intolerant of classic antipsychotics.

Non-responsiveness is defined as a lack of satisfactory clinical improvement despite the use of adequate doses of at least two marketed antipsychotics prescribed for adequate durations.

Intolerance is defined as the impossibility of achieving adequate clinical benefit with classic antipsychotics because of severe and untreatable neurological adverse reactions

Clopine tablets and Suspension are only indicated in patients aged 16 years and above.

### 4.2. Dose and method of administration

Brand swapping between clozapine products is discouraged and should only occur on the advice of a clinician.

#### Dose

The dosage must be adjusted individually. For each patient, the lowest effective dose should be used. Cautious titration and a divided dosage schedule are necessary to minimise the risks of hypotension, seizure, and sedation.

Initiation of clozapine treatment must be restricted to those patients with a white blood cell (WBC) count  $\geq 3500/\text{mm}^3$  ( $3.5 \times 10^9/\text{L}$ ) and an absolute neutrophil count (ANC)  $\geq 2000/\text{mm}^3$  ( $2.0 \times 10^9/\text{L}$ ) and within standardised normal limits.

Dose adjustment is indicated in patients who are also receiving medicinal products that have pharmacokinetic interactions with clozapine, such as benzodiazepines or selective serotonin reuptake inhibitors (*see section 4.5*).

In patients with a history of seizures or suffering from renal or cardiovascular disorders (note: severe renal or cardiovascular disorders are contraindications) the initial dose should be 12.5 mg given once on the first day, and dosage increase should be slow and in small increments.

#### Switching from a previous antipsychotic therapy to Clopine

It is generally recommended that Clopine should not be used in combination with other antipsychotics. When Clopine therapy is to be initiated in a patient undergoing oral antipsychotic therapy, it is recommended that the dosage of other antipsychotics be

reduced or discontinued by gradually tapering it downwards. Based on the clinical circumstances, the prescribing physician should judge whether or not to discontinue the other antipsychotic therapy before initiating treatment with Clopine.

#### Treatment-resistant schizophrenia

Starting therapy Clopine should be started with 12.5 mg (half a 25 mg tablet) once or twice on the first day, followed by one or two 25 mg tablets on the second day. If well tolerated, the daily dose may then be increased slowly in increments of 25 mg to 50 mg in order to achieve a dose level of up to 300 mg/day within 2 to 3 weeks. Thereafter, if required, the daily dose may be further increased in increments of 50 mg to 100 mg at half-weekly or, preferably, weekly intervals.

#### Therapeutic dose range

In most patients, antipsychotic efficacy can be expected with 300 to 450 mg/day given in divided doses. Some patients may be treated with lower doses, and some patients may require doses up to 600 mg/day. The total daily dose may be divided unevenly, with the larger portion being taken at bedtime.

#### Maximum dose

To obtain a full therapeutic benefit, a few patients may require larger doses, in which case judicious increments (not exceeding 100 mg) are permissible up to 900 mg/day. The possibility of increased adverse reactions (in particular seizures) occurring at doses over 450 mg/day must be borne in mind.

#### Maintenance dose

After achieving maximum therapeutic benefit, many patients can be maintained effectively on lower doses. Careful downward titration is therefore recommended. Treatment should be maintained for at least 6 months. If the daily dose does not exceed 200 mg, once daily administration in the evening may be appropriate.

#### Ending therapy

In the event of planned termination of Clopine therapy, a gradual reduction in dose over a 1- to 2-week period is recommended. If abrupt discontinuation is necessary (e.g. because of leucopenia), the patient should be carefully observed for the recurrence of psychotic symptoms and symptoms related to cholinergic rebound (*see section 4.4*).

#### Re-starting therapy

In patients in whom the interval since the last dose of Clopine exceeds 2 days, treatment should be reinitiated with 12.5 mg (half a 25 mg tablet) given once or twice on the first day. If this dose is well tolerated, it may be feasible to titrate the dose to the therapeutic level more quickly than is recommended for initial treatment. However, in any patient who has previously experienced respiratory or cardiac arrest with initial *dosing* (*see section 4.4*), but was then able to be successfully titrated to a therapeutic dose, re-titration should be done with extreme caution.

## ***Special populations***

### **Cardiovascular disorders**

In patients suffering from cardiovascular disorders (note: severe cardiovascular disorders are contraindications) the initial dose should be 12.5 mg given once on the first day, and dosage increase should be slow and in small increments.

### **Renal impairment**

In patients with mild to moderate renal impairment the initial dose should be 12.5 mg given once on the first day, and dosage increase should be slow and in small increments.

### **Hepatic impairment**

Patients with hepatic impairment should receive Clopine with caution along with regular monitoring of liver function tests (*see section 4.4*)

### **Patients 60 years of age and older**

It is recommended that treatment in patients 60 years and older is initiated at a particularly low dose (12.5 mg given once on the first day) with subsequent dose increments restricted to 25 mg/day.

### ***Paediatric population***

No paediatric studies have been performed.

### **Children and adolescents**

The safety and efficacy of clozapine in children and adolescents up to 16-years of age have not been established.

### **Method of Administration**

Clopine is administered orally.

## **4.3. Contraindications**

- Known hypersensitivity to clozapine or to any of the excipients of Clopine (*see section 6.1*).
- Patients unable to undergo regular blood tests.
- History of toxic or idiosyncratic granulocytopenia/agranulocytosis (with the exception of granulocytopenia/agranulocytosis from previous chemotherapy).
- Impaired bone marrow function.
- Uncontrolled epilepsy.
- Alcoholic and other toxic psychoses, drug intoxication, comatose conditions.
- Circulatory collapse and/or CNS depression of any cause.
- Severe renal or cardiac disorders (e.g. myocarditis).
- Active liver disease associated with nausea, anorexia or jaundice; progressive liver disease, hepatic failure.
- Paralytic ileus.

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

##### Agranulocytosis

Prescribers and dispensers should verify that the patient has not previously developed an adverse reaction to clozapine that contraindicates further use of any clozapine containing product.

Because of the association of clozapine with **agranulocytosis**, the following precautionary measures are mandatory:

Medicines known to have a substantial potential to depress bone marrow function should not be used concurrently with Clopine. In addition, the concomitant use of long-acting depot antipsychotics should be avoided because of the impossibility of removing these medications, which may be potentially myelosuppressive, from the body rapidly in situations where this may be required, e.g. granulocytopenia.

Patients with a history of primary bone marrow disorders may be treated only if the benefit outweighs the risk. They should be carefully reviewed by a haematologist prior to starting Clopine.

Patients who have low white blood cell (WBC) counts because of benign ethnic neutropenia should be given special consideration and may be started on Clopine after agreement of a haematologist.

##### White Blood Cell (WBC) counts and Absolute Neutrophil Count (ANC) monitoring

White blood cell (WBC) and differential blood counts must be performed within 10 days prior to starting Clopine treatment to ensure that only patients with normal leukocyte and absolute neutrophil counts ( $WBC \geq 3500/\text{mm}^3$  ( $\geq 3.5 \times 10^9/\text{L}$ ) and  $ANC \geq 2000/\text{mm}^3$  ( $\geq 2.0 \times 10^9/\text{L}$ )) will receive Clopine. After the start of Clopine treatment, regular WBC count and ANC must be performed and monitored weekly for 18 weeks, and thereafter at least every four weeks throughout treatment, and for 4 weeks after complete discontinuation of Clopine.

Prescribing physicians should comply fully with the required safety measures. At each consultation, the patient should be reminded to contact the treating physician immediately if any kind of infection begins to develop. Particular attention should be paid to flu-like complaints such as fever, or a sore throat and to other evidence of infection, which may be indicative of neutropenia. A differential blood count must be performed immediately if any symptoms or signs of an infection occur.

##### Low WBC count and/or ANC

If during the first 18 weeks of Clopine therapy, the WBC count falls to between  $3500/\text{mm}^3$  ( $3.5 \times 10^9/\text{L}$ ) and  $3000/\text{mm}^3$  ( $3.0 \times 10^9/\text{L}$ ) and/or the ANC falls to between  $2000/\text{mm}^3$  ( $2.0 \times 10^9/\text{L}$ ) and  $1500/\text{mm}^3$  ( $1.5 \times 10^9/\text{L}$ ), haematological evaluations must be performed at least twice weekly.

After 18 weeks of Clopine therapy, haematological evaluations should be performed at least twice weekly if the WBC count falls to between  $3000/\text{mm}^3$  ( $3.0 \times 10^9/\text{L}$ ) and  $2500/\text{mm}^3$  ( $2.5 \times 10^9/\text{L}$ ) and/or the ANC falls to between  $1500/\text{mm}^3$  ( $1.5 \times 10^9/\text{L}$ ) and  $1000/\text{mm}^3$  ( $1.0 \times 10^9/\text{L}$ ). In addition, if, during Clopine therapy, the WBC count is found

to have dropped by a substantial amount from baseline, a repeat WBC count and a differential blood count should be performed.

A substantial drop is defined as a single drop of  $3000 \text{ mm}^3$  ( $3.0 \times 10^9/\text{L}$ ) or more in the WBC count or a cumulative drop of  $3000 \text{ mm}^3$  ( $3.0 \times 10^9/\text{L}$ ) or more within three weeks.

Immediate discontinuation of Clopine is mandatory if the WBC count is less than  $3000/\text{mm}^3$  ( $3.0 \times 10^9/\text{L}$ ) or the ANC is less than  $1500/\text{mm}^3$  ( $1.5 \times 10^9/\text{L}$ ) during the first 18 weeks of therapy, or if the WBC count is less than  $2500/\text{mm}^3$  ( $2.5 \times 10^9/\text{L}$ ) or the ANC is less than  $1000/\text{mm}^3$  ( $1.0 \times 10^9/\text{L}$ ) after the first 18 weeks of therapy. WBC counts and differential blood counts should then be performed daily and patients should be carefully monitored for flu-like symptoms or other symptoms suggestive of infection. Following discontinuation of Clopine, haematological evaluation is required until haematological recovery has occurred.

If Clopine has been withdrawn and WBC count falls further to below  $2000/\text{mm}^3$  ( $2.0 \times 10^9/\text{L}$ ) and/or the ANC falls below  $1000/\text{mm}^3$  ( $1.0 \times 10^9/\text{L}$ ), the management of this condition must be guided by an experienced haematologist. If possible, the patient should be referred to a specialised haematological unit, where protective isolation and the administration of GM-CSF (granulocyte-macrophage colony stimulating factor) or G-CSF (granulocyte colony stimulating factor) may be indicated. It is recommended that the colony stimulating factor therapy to be discontinued when the neutrophil count has returned to a level above  $1000/\text{mm}^3$  ( $1.0 \times 10^9/\text{L}$ ).

Patients in whom Clopine has been discontinued as a result of white blood cell deficiencies (see above) must not be re-exposed to Clopine.

It is recommended that the haematological values be confirmed by performing two blood counts on two consecutive days; however, Clopine should be discontinued after the first blood count.

**Table 1: Blood monitoring during the first 18 weeks of Clopine therapy**

Blood cell count		Action required
WBC/ $\text{mm}^3$ (/L)	ANC/ $\text{mm}^3$ (/L)	
$\geq 3500$ ( $> 3.5 \times 10^9$ )	$\geq 2000$ ( $> 2.0 \times 10^9$ )	Continue Clopine treatment.
3000-3500 ( $3.0 - 3.5 \times 10^9$ )	1500-2000 ( $1.5 - 2.0 \times 10^9$ )	Continue Clopine treatment, sample blood twice weekly until counts stabilise or increase.
$< 3000$ ( $< 3.0 \times 10^9$ )	$< 1500$ ( $< 1.5 \times 10^9$ )	Immediately stop Clopine treatment, sample blood daily until haematological abnormality is resolved, and monitor for infection. Do not re-expose the patient.

**Table 2: Blood monitoring after 18 weeks of Clopine therapy**

Blood cell count		Action required
WBC/mm <sup>3</sup> (/L)	ANC/mm <sup>3</sup> (/L)	
≥3000 (>3.0 x 10 <sup>9</sup> )	≥1500 (> 1.5 x 10 <sup>9</sup> )	Continue Clopine treatment.
2500-3000 (2.5 - 3.0 x 10 <sup>9</sup> )	1000-1500 (1.0 - 1.5 x 10 <sup>9</sup> )	Continue Clopine treatment, sample blood twice weekly until counts stabilise or increase.
<2500 (<2.5 x 10 <sup>9</sup> )	<1000 (<1.0 x 10 <sup>9</sup> )	Immediately stop Clopine treatment, sample blood daily until haematological abnormality is resolved, and monitor for infection. Do not re-expose the patient.

In the event of interruption of therapy for non-haematological reasons

Patients who have been on Clopine for more than 18 weeks and have had their treatment interrupted for more than 3 days but less than 4 weeks should have their WBC count and ANC monitored weekly for an additional 6 weeks. If no haematological abnormality occurs, monitoring at intervals not exceeding 4 weeks may be resumed. If Clopine treatment has been interrupted for 4 weeks or longer, weekly monitoring is required for the next 18 weeks of treatment.

Other precautions

Eosinophilia

In the event of **eosinophilia**, discontinuation of Clopine is recommended if the eosinophil count rises above 3000/mm<sup>3</sup> (3.0 x 10<sup>9</sup>/L). Therapy should be re-started only after the eosinophil count has fallen below 1000/mm<sup>3</sup> (1.0 x 10<sup>9</sup>/L).

Thrombocytopenia

In the event of **thrombocytopenia**, discontinuation of Clopine is recommended if the platelet count falls below 50 000/mm<sup>3</sup> (50.0 x 10<sup>9</sup>/L).

Cardiovascular disorders

In patients suffering from cardiovascular disorders (note: severe cardiovascular disorders are contraindications) the initial dose should be 12.5 mg given once on the first day, and dosage increase should be slow and in small increments (*see section 4.2*).

Hypotension

**Orthostatic hypotension**, with or without syncope, can occur during clozapine treatment. Rarely (about one case per 3,000 clozapine-treated patients), collapse can be profound and may be accompanied by cardiac and/or respiratory arrest. Such events are



more likely to occur during initial titration in association with rapid dose escalation; on very rare occasions they occurred even after the first dose. Therefore, patients commencing Clopine treatment require close medical supervision.

#### Myocarditis/Cardiomyopathy

Tachycardia that persists at rest, accompanied by arrhythmias, shortness of breath or signs and symptoms of heart failure, may rarely occur during the first month of treatment and very rarely thereafter. The occurrence of these signs and symptoms necessitates an urgent diagnostic evaluation for **myocarditis**, especially during the titration period. Therefore, the possibility of myocarditis should be considered in patients receiving Clopine who present with unexplained fatigue, dyspnoea, tachypnoea, fever, chest pain, tachycardia, palpitations, other signs and symptoms of heart failure, ECG changes (such as ST-T wave abnormalities) or arrhythmias. It is not known whether eosinophilia is a reliable predictor of myocarditis. The incidence of myocarditis, reported globally is rare (<0.1%) during the first month of therapy and very rare (<0.01%) thereafter. Some cases of myocarditis have been reported to be fatal (incidence approximately 0.2 cases/100,000 patient years). Most reported cases of myocarditis have occurred in the first month of therapy, therefore there should be a high index of suspicion in the first 6-8 weeks of therapy.

If the diagnosis of myocarditis is confirmed, Clopine should be discontinued. There have been postmarketing reports of myocarditis including fatal cases. Later in treatment, the same signs and symptoms may very rarely occur and may be linked to cardiomyopathy. Further investigation should be performed and if the diagnosis is confirmed, the treatment should be stopped unless the benefit clearly outweighs the risk to the patient. If patients are diagnosed with cardiomyopathy while on Clopine treatment, there is potential to develop mitral valve incompetence. Mitral valve incompetence has been reported in cases of cardiomyopathy related to clozapine treatment. These cases of mitral valve incompetence reported either mild or moderate mitral regurgitation on two-dimensional echocardiography (2DEcho) (*see section 4.8*).

Monitoring of standing and supine blood pressure is necessary during the first weeks of treatment in patients with Parkinson's disease.

Patients with clozapine-induced myocarditis or cardiomyopathy should not be re-exposed to Clopine.

#### Myocardial infarction

In addition, there have been postmarketing reports of **myocardial infarction** including fatal cases. Causality assessment was difficult in the majority of these cases because of serious pre-existing cardiac disease and plausible alternative causes.

#### Seizures

Clopine may lower seizure threshold. In patients with a history of **seizures**, or suffering from renal or cardiovascular disorders (note: severe renal or cardiovascular disorders are contraindications) the initial dose should be 12.5 mg given once on the first day, and dosage increase should be slow and in small increments (*see section 4.2*)

### Anticholinergic effects

Clozapine exerts **anticholinergic activity**, which may produce adverse effects throughout the body. Careful supervision is indicated in the presence of **prostatic enlargement** and **narrow-angle glaucoma**. Probably on account of its anticholinergic properties, Clozapine has been associated with varying degrees of **impairment of intestinal peristalsis**, ranging from **constipation** to **intestinal obstruction, faecal impaction, paralytic ileus, megacolon** and **intestinal infarction/ischaemia** (*see section 4.8*). On rare occasions, these cases have proved fatal. Since complications have been associated with delayed diagnosis, patients should be questioned about their bowel habits.

### Fever

During Clozapine therapy, patients may experience transient temperature elevations above 38°C, with the peak incidence within the first 3 weeks of treatment. This fever is generally benign. Occasionally, it may be associated with an increase or decrease in the WBC count. Patients with fever should be carefully evaluated to rule out the possibility of an underlying infection or the development of agranulocytosis. In the presence of high fever, the possibility of neuroleptic malignant syndrome (NMS) must be considered.

### Falls

Clozapine may cause seizures, somnolence, postural hypotension, motor and sensory instability, which may lead to falls and, consequently, fractures or other injuries. For patients with diseases, conditions, or medications that could exacerbate these effects, complete fall risk assessments when initiating antipsychotic treatment and recurrently for patients on long-term antipsychotic therapy.

### Neuroleptic Malignant Syndrome (NMS)

If the diagnosis of **NMS** is confirmed, Clozapine should be discontinued immediately and appropriate medical measures should be administered.

### Metabolic Changes

Atypical antipsychotic drugs, including clozapine, have been associated with **metabolic changes** that may increase cardiovascular/cerebrovascular risk. These metabolic changes may include hyperglycaemia, dyslipidemia, and body weight gain. While atypical antipsychotic drugs may produce some metabolic changes, each drug in the class has its own specific risk profile.

### Hyperglycaemia and Diabetes Mellitus

Hyperglycaemia, in some cases extreme and associated with ketoacidosis or hyperosmolar coma or death, has been reported in patients with atypical antipsychotics including clozapine. Assessment of the relationship between atypical antipsychotic use and glucose abnormalities is complicated by the possibility of an increased background risk of diabetes mellitus in patients with schizophrenia and the increasing incidence of diabetes mellitus in the general population. Given these confounders, the relationship between atypical antipsychotic use and hyperglycaemia-related adverse events is not completely understood. However, epidemiological studies suggest an increased risk of treatment-emergent hyperglycaemia related adverse events in patients treated with the atypical antipsychotics studied. Precise risk estimates for hyperglycaemia-related adverse events in patients treated with atypical antipsychotics are not available. The

available data are insufficient to provide reliable estimates of differences in hyperglycaemia related adverse event risk among the marketed atypical antipsychotics.

Patients with an established diagnosis of diabetes mellitus who are started on atypical antipsychotics should be monitored regularly for worsening of glucose control. Patients with risk factors for diabetes mellitus (e.g. obesity, family history of diabetes) who are starting treatment with atypical antipsychotics should undergo fasting blood glucose testing at the baseline and periodically during treatment. Any patient treated with atypical antipsychotics should be monitored for symptoms of hyperglycaemia including polydipsia, polyuria, polyphagia, and weakness. Patients who develop symptoms of hyperglycaemia during treatment with atypical antipsychotics should undergo fasting blood glucose testing. In some cases, hyperglycaemia has resolved when the atypical antipsychotic was discontinued; however, some patients required continuation of anti-diabetic treatment despite discontinuation of the suspect drug. In patients with significant treatment-emergent hyperglycaemia, discontinuation of Clopine should be considered.

There is a risk of altering the metabolic balance resulting in slight impairment of glucose homeostasis and a possibility of unmasking a pre-diabetic condition or aggravating pre-existing diabetes.

#### Dyslipidemia

Undesirable alterations in **lipids** have been observed in patients treated with atypical antipsychotics, including clozapine. Clinical monitoring, including baseline and periodic, follow up lipid evaluations in patients using clozapine, is recommended.

#### Weight Gain

Weight gain has been observed with atypical antipsychotic use, including clozapine. Clinical monitoring of **weight** is recommended.

#### Risk of Thromboembolism

Since Clopine may cause sedation and weight gain, thereby increasing the risk of **thromboembolism**, immobilisation of patients should be avoided.

#### Cerebrovascular adverse events

An increased risk of **cerebrovascular adverse events** has been seen in the dementia population with some atypical antipsychotics. The mechanism for this increased risk is not known. An increased risk cannot be excluded for other antipsychotics or other patient populations. Clopine should be used with caution in patients with risk factors for stroke.

#### Prolongation of QT Interval

As with other antipsychotics, caution is advised in patients with known cardiovascular disease or family history of QT prolongation.

As with other antipsychotics, caution should be exercised when Clopine is prescribed with medicines known to increase the QTc interval.

## Parkinson's Disease

Physicians should weigh the risks versus the benefits when prescribing clozapine to patients with **Parkinson's Disease** or dementia with Lewy Bodies (DLB) since both groups may be at increased risk of Neuroleptic Malignant Syndrome as well as having an increased sensitivity to antipsychotics. The manifestation of this increased sensitivity can include confusion, obtundation, postural instability with frequent falls, in addition to extrapyramidal symptoms.

## Suicide

The possibility of a **suicide** attempt is inherent in schizophrenia and close supervision of high-risk patients should accompany therapy.

## ***Special populations***

### Liver disease

Patients with stable pre-existing liver disorders may receive Clopine, but must undergo regular liver function tests. Such tests should be performed immediately in patients who develop symptoms of possible liver dysfunction such as nausea, vomiting and/or anorexia during Clopine treatment. If the elevation of the values is clinically relevant or if symptoms of jaundice occur, treatment with Clopine must be discontinued. It may be resumed (*see section 4.3 - Restarting therapy*) only when the results of liver function tests are normal. In such cases, liver function should be closely monitored after re-introduction of Clopine.

### Renal impairment

In patients suffering from mild to moderate renal impairment, an initial dose of 12.5 mg/day (half a 25 mg tablet) is recommended (*see section 4.3*).

### Patients aged 60 years and older

It is recommended that treatment is initiated at a particularly low dose (12.5 mg given once on the first day) and subsequent dose increments be restricted to 25 mg/day.

Clinical studies with clozapine did not include sufficient numbers of subjects aged 60 years and over to determine whether or not they respond differently from younger subjects.

Orthostatic hypotension can occur with Clopine treatment and there have been rare reports of tachycardia, which may be sustained, in patients taking Clopine. Patients aged 60 years and over, particularly those with compromised cardiovascular function, may be more susceptible to these effects.

Patients aged 60 years and over may also be particularly susceptible to the anticholinergic effects of clozapine, such as urinary retention and constipation.

### Patients aged 60 years and older with Dementia-related Psychosis

In patients aged 60 years and over with dementia-related psychosis, the efficacy and safety of clozapine have not been studied. Observational studies suggest that patients aged 60 years and over with dementia-related psychosis treated with antipsychotic drugs are at an increased risk of death. In the published literature, risk factors that may

predispose this patient population to increased risk of death when treated with antipsychotics include sedation, the presence of cardiac conditions (e.g. cardiac arrhythmias) or pulmonary conditions (e.g. pneumonia, with or without aspiration). Clopine is not approved for the treatment of dementia-related behavioural disturbances.

#### Rebound, withdrawal effects

If abrupt discontinuation of Clopine is necessary (e.g. because of leucopenia), the patient should be carefully observed for the recurrence of psychotic symptoms and symptoms related to cholinergic rebound such as profuse sweating, headache, nausea, vomiting and diarrhoea.

### **4.5. Interaction with other medicines and other forms of interaction**

#### **Pharmacodynamic-related Interactions**

Anticipated pharmacodynamic interactions resulting in concomitant use not being recommended Medicinal products known to have a substantial potential to depress bone marrow function should not be used concurrently with Clopine (*see section 4.4*).

As with other antipsychotics, caution should be exercised when Clopine is prescribed with medicines known to increase the QTc interval or cause electrolyte imbalance.

#### Observed pharmacodynamic interactions to be considered

Particular caution is recommended when Clopine therapy is initiated in patients who are receiving (or have recently received) a benzodiazepine or any other psychotropic agent, as these patients may have an increased risk of circulatory collapse, which, on rare occasions, can be profound and may lead to cardiac and/or respiratory arrest.

Concomitant use of lithium or other CNS-active agents may increase the risk of development of neuroleptic malignant syndrome (NMS).

Rare but serious reports of seizures, including onset of seizures in non-epileptic patients, and isolated cases of delirium where clozapine was co-administered with valproic acid have been reported. These effects are possibly due to a pharmacodynamic interaction, the mechanism of which has not been determined.

#### Anticipated pharmacodynamic interactions to be considered

Clozapine may enhance the central effects of alcohol, MAO inhibitors and CNS depressants such as narcotics, antihistamines, and benzodiazepines.

Because of the possibility of additive effects, caution is essential when substances possessing anticholinergic, hypotensive, or respiratory depressant effects are given concomitantly.

Owing to its anti-alpha-adrenergic properties, clozapine may reduce the blood pressure, increasing the effect of noradrenaline, or other predominantly alpha-adrenergic agents and reverse the pressor effect of adrenaline.

## **Pharmacokinetic-related Interactions**

Clozapine is a substrate for many CYP 450 isoenzymes, in particular 1A2 and 3A4. The risk of metabolic interactions caused by an effect on an individual isoform is therefore minimised. Nevertheless, caution is called for in patients receiving concomitant treatment with other substances that are either inhibitors or inducers of these enzymes.

No clinically relevant interactions have been observed thus far with tricyclic antidepressants, phenothiazines or type 1C antiarrhythmics, which are known to bind to cytochrome P450 2D6.

### Observed pharmacokinetic interactions to be considered

Concomitant administration of substances known to induce cytochrome P450 enzymes may decrease the plasma levels of clozapine.

- Substances known to induce the activity of 3A4 and with reported interactions with clozapine include, for instance, carbamazepine, phenytoin and rifampicin.

Concomitant administration of substances known to inhibit the activity of cytochrome P450 isozymes may increase the plasma levels of clozapine.

- Substances known to inhibit the activity of the major isozymes involved in the metabolism of clozapine and with reported interactions include, for instance, cimetidine, erythromycin (3A4), fluvoxamine (1A2), perazine (1A2), ciprofloxacin (1A2) and oral contraceptives (1A2, 3A4, 2C19).
- The plasma concentration of clozapine is increased by caffeine (1A2) intake and decreased by nearly 50% following a 5-day caffeine-free period.
- Elevated clozapine plasma concentrations also have been reported in patients receiving the substances in combination with selective serotonin re-uptake inhibitors (SSRIs) such as paroxetine (1A2), sertraline, fluoxetine or citalopram.

### Anticipated pharmacokinetic interactions to be considered

Concomitant administration of substances known to induce cytochrome P450 enzymes may decrease the plasma levels of clozapine.

- Known inducers of 1A2 include, for instance, omeprazole and tobacco smoke. In cases of sudden cessation of tobacco smoking, the plasma clozapine concentration may be increased, thus leading to an increase in adverse effects.

Concomitant administration of substances known to inhibit the activity of cytochrome P450 isozymes may increase the plasma levels of clozapine.

- Potent inhibitors of CYP3A, such asazole antimycotics and protease inhibitors, could potentially also increase clozapine plasma concentrations; no interactions have been reported to date, however.

## **4.6. Fertility, pregnancy and lactation**

### Women of childbearing potential and contraceptive measures

Some female patients treated with antipsychotics other than Clopine may become amenorrhic. A return to normal menstruation may occur as a result of switching from

other antipsychotics to Clopine. Adequate contraceptive measures must, therefore, be ensured in women of childbearing potential.

#### Pregnancy

Reproduction studies in animals have revealed no evidence of impaired fertility or harm to the foetus due to clozapine. However, the safe use of clozapine in pregnant women has not been established. Therefore, Clopine should be used during pregnancy only if the expected benefit clearly outweighs any potential risk.

#### *Non-teratogenic effects*

Neonates exposed to antipsychotic drugs, during the third trimester of pregnancy are at risk for extrapyramidal and/or withdrawal symptoms following delivery. There have been reports of agitation, hypertonia, hypotonia, tremor, somnolence, respiratory distress and feeding disorder in these neonates. These complications have varied in severity; while in some cases symptoms have been self-limited, in other cases neonates have required intensive care unit support and prolonged hospitalisation.

Antipsychotic drugs, including clozapine, should be used during pregnancy only if the potential benefit justifies the potential risk to the foetus.

#### Breast-feeding

Animal studies suggest that clozapine is excreted in breast milk and has an effect in the suckling offspring; therefore, mothers receiving Clopine should not breast-feed. Potentially clozapine may cause sedation, cardiovascular instability, restlessness or irritability or seizures in the nursing infant.

### **4.7. Effects on ability to drive and use machines**

Owing to the ability of Clopine to cause sedation and lower the seizure threshold, activities such as driving or operating machinery should be avoided, especially during the initial weeks of treatment.

### **4.8. Undesirable effects**

The adverse effects of clozapine are most often predictable based on its pharmacological properties with the exception of agranulocytosis (*see section 4.4*).

The most serious adverse reactions experienced with clozapine are agranulocytosis, seizure, cardiovascular effects and fever (*see section 4.4*). The most common side effects are drowsiness/sedation, dizziness, tachycardia, constipation, and hypersalivation.

Data from the clinical trials experience showed that a varying proportion of clozapine-treated patients (from 7.1 to 15.6%) discontinued treatment due to an adverse event, including only those that could be reasonably attributed to clozapine. The more common events considered to be causes of discontinuation were leukopenia; somnolence; dizziness (excluding vertigo); and psychotic disorder.

**Table 3: Treatment-emergent adverse experience frequency estimate from spontaneous and clinical trial reports**

Frequencies are defined as very common ( $\geq 1/10$ ); common ( $\geq 1/100$  to  $< 1/10$ ); uncommon ( $\geq 1/1,000$  to  $< 1/100$ ); rare ( $\geq 1/10,000$  to  $< 1/1,000$ ); very rare ( $< 1/10,000$ ), not known (cannot be estimated from the available data).

<b>Blood and lymphatic system disorders</b>	
Common:	Leukopenia/decreased WBC/neutropenia, eosinophilia, leukocytosis
Uncommon:	Agranulocytosis
Rare:	Anaemia
Very rare:	Thrombocytopenia, thrombocythaemia
<b>Metabolism and nutrition disorders</b>	
Common:	Weight gain
Rare:	Impaired glucose tolerance, new onset diabetes, diabetes aggravated
Very rare:	Ketoacidosis, hyperosmolar coma, severe hyperglycaemia, hypercholesterolemia, hypertriglyceridemia
<b>Psychiatric disorders</b>	
Common:	Dysarthria
Uncommon:	Dysphemia
Rare:	Agitation, restlessness
<b>Nervous system disorders</b>	
Very common:	Drowsiness/sedation, dizziness
Common:	Blurred vision, headache, tremor, rigidity, akathisia, extrapyramidal symptoms, seizures/convulsions/myoclonic jerks,
Uncommon:	Neuroleptic malignant syndrome
Rare:	Confusion, delirium
Very rare:	Tardive dyskinesia, obsessive-compulsive symptoms
<b>Eye disorders</b>	
Common:	Blurred vision
<b>Cardiac disorders *</b>	
Very common:	Tachycardia
Common:	ECG changes



Rare:	Circulatory collapse, arrhythmias, myocarditis**, pericarditis
Very rare:	Cardiomyopathy
<b>Vascular disorders</b>	
Common:	Hypertension, postural hypotension, syncope
Rare:	Thromboembolism
<b>Respiratory, thoracic and mediastinal disorders</b>	
Rare:	Aspiration of ingested food, pneumonia and lower respiratory tract infection which may be fatal
Very rare:	Respiratory depression/arrest
<b>Gastrointestinal disorders</b>	
Very common:	Constipation, hypersalivation
Common:	Nausea, vomiting, dry mouth
Rare:	Dysphagia
Very rare:	Parotid gland enlargement, intestinal obstruction/paralytic ileus/faecal impaction, abdominal discomfort, heartburn
<b>Hepatobiliary disorders</b>	
Common:	Elevated liver enzymes
Rare:	Hepatitis, cholestatic jaundice, pancreatitis
Very rare:	Fulminant hepatic necrosis
<b>Skin and subcutaneous tissue disorders</b>	
Very rare:	Skin reactions
<b>Renal and urinary disorders</b>	
Common:	Urinary incontinence urinary retention
Very rare:	Interstitial nephritis, nocturnal enuresis
<b>Reproductive system and breast disorders</b>	
Very rare:	Priapism
<b>General disorders and administration site conditions</b>	
Common:	Fatigue, benign hyperthermia, disturbances in sweating/temperature regulation
Uncommon	Neuroleptic malignant syndrome
Very rare:	Sudden unexplained death
<b>Investigations</b>	

Rare:	Increased CPK
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\*Very rare events of ventricular tachycardia and QT prolongation which may be associated with Torsades De Pointes have been observed although there is no conclusive causal relationship to the use of this medicine.

\*\*The prompt discontinuation of Clopine therapy is warranted upon suspicion of myocarditis (see section 4.4)

***AEs from spontaneous reports and literature (frequency unknown)***

The following post-marketing adverse effects were derived from experience with clozapine via spontaneous case reports and literature cases and have been categorised according to MedDRA system organ class (Table 4). Because these have been reported voluntarily from a population of uncertain size and are subject to confounding factors, these post-marketing AEs have been categorised with a frequency of “unknown” since it is not possible to reliably estimate their frequency. Adverse effects are listed according to system organ classes (SOC) in MedDRA.

Within each system organ class, AEs are presented in order of decreasing seriousness.

**Table 4: Adverse drug reactions from spontaneous reports and literature (frequency not known)**

<b>Infections and infestations</b>
Sepsis
<b>Immune system disorders</b>
Angioedema, leukocytoclastic vasculitis
<b>Endocrine disorders</b>
Pseudophaeochromocytoma
<b>Nervous system disorders</b>
Cholinergic syndrome, EEG changes, pleurothotonus
<b>Metabolism and nutrition disorders</b>
Obesity
<b>Cardiac disorders</b>
Myocardial infarction sometimes fatal, myocarditis sometimes fatal, chest pain/angina pectoris, palpitations, atrial fibrillation, mitral valve incompetence associated with clozapine-related cardiomyopathy
<b>Vascular disorders</b>
Hypotension
<b>Respiratory, thoracic and mediastinal disorders</b>
Pleural effusion, sleep apnoea syndrome, nasal congestion
<b>Gastrointestinal disorders</b>

Megacolon sometimes fatal, intestinal infarction/ischaemia sometimes fatal, diarrhoea, abdominal discomfort/heartburn/dyspepsia, colitis
<b>Hepatobiliary disorders</b>
Hepatic steatosis, hepatic necrosis, hepatotoxicity, hepatic fibrosis, hepatic cirrhosis, liver disorders including those hepatic events leading to life-threatening consequences such as liver injury (hepatic, cholestatic and mixed), liver failure which may be fatal and liver transplant
<b>Skin and subcutaneous tissue disorders</b>
Pigmentation disorder
<b>Musculoskeletal &amp; connective tissue disorders</b>
Rhabdomyolysis, muscle weakness, muscle spasms, muscle pain, systemic lupus erythematosus
<b>Renal and urinary disorders</b>
Renal failure, nocturnal enuresis
<b>Reproductive system and breast disorders</b>
Retrograde ejaculation
<b>General disorders and administration site conditions</b>
Polyserositis
<b>Injury, poisoning and procedural complications</b>
Falls (associated with clozapine-induced seizures, somnolence, postural hypotension, motor and sensory instability)

#### Reporting of suspected adverse reactions

Reporting suspected adverse reactions after authorisation of the medicine is important. It allows continued monitoring of the benefit/risk balance of the medicine. Healthcare professionals are asked to report any suspected reactions

<https://nzphvc.otago.ac.nz/reporting/>

#### **4.9. Overdose**

In cases of acute intentional or accidental clozapine overdosage, for which information on the outcome is available, to date, the mortality is about 12%. Most of the fatalities were associated with cardiac failure or pneumonia caused by aspiration and occurred at doses above 2000 mg.

There have been reports of patients recovering from an overdose in excess of 10 000 mg. However, in a few adult individuals, primarily those not previously exposed to clozapine, the ingestion of doses as low as 400 mg led to life-threatening comatose conditions and, in one case, to death. In young children, the intake of 50 mg to 200 mg resulted in strong sedation or coma without being lethal.

### Signs and symptoms

Drowsiness, lethargy, areflexia, coma, confusion, hallucinations, agitation, delirium, extrapyramidal symptoms, hyperreflexia, convulsions; hypersalivation, mydriasis, blurred vision, thermolability; hypotension, collapse, tachycardia, cardiac arrhythmias; aspiration pneumonia, dyspnoea, respiratory depression or failure.

### Treatment

There are no specific antidotes for clozapine.

Gastric lavage and/or the administration of activated charcoal within the first 6 hours after Clopine ingestion. (Peritoneal dialysis and haemodialysis are unlikely to be effective.) Symptomatic treatment under continuous cardiac monitoring, surveillance of respiration, monitoring of electrolytes and acid-base balance. The use of adrenaline should be avoided in the treatment of hypotension because of the possibility of a 'reverse adrenaline effect. Close medical supervision is necessary for at least 5 days because of the possibility of delayed reactions.

For advice on the management of overdose please contact the National Poisons Centre on 0800 POISON (0800 764766).

## **5. PHARMACOLOGICAL PROPERTIES**

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### **5.1. Pharmacodynamic properties**

Pharmacotherapeutic group: Antipsychotic agent (ATC code NO5A H02)

#### Mechanism of action

Clozapine has been shown to be an antipsychotic agent that is different from classic antipsychotics. In pharmacological experiments, the compound does not induce catalepsy or inhibit apomorphine- or amphetamine-induced stereotyped behaviour. It has only weak dopamine receptor-blocking activity at D<sub>1</sub>, D<sub>2</sub>, D<sub>3</sub> and D<sub>5</sub> receptors, but shows high potency for the D<sub>4</sub> receptor, in addition to potent anti-alpha-adrenergic, anticholinergic, antihistaminic, and arousal reaction-inhibiting effects. It has also been shown to possess antiserotonergic properties.

#### Pharmacodynamic effects

Clinically clozapine produces rapid and marked sedation and exerts antipsychotic effects in patients with schizophrenia resistant to other antipsychotic agents. In such cases, Clozapine has proven effective in relieving both positive and negative schizophrenic symptoms in short- and long-term trials

Clozapine is unique in that it produces virtually no major extrapyramidal reactions such as acute dystonia and tardive dyskinesia. Furthermore, parkinsonian-like side effects and akathisia are rare. In contrast to classical antipsychotics, clozapine produces little or no prolactin elevation, thus avoiding adverse effects such as gynaecomastia, amenorrhoea, galactorrhoea, and impotence.

Potentially serious adverse reactions caused by clozapine therapy are granulocytopenia and agranulocytosis occurring at an estimated incidence of 3% and 0.7% respectively (see Special warnings and precautions for use).

## Clinical efficacy and safety

### ***Clinical studies in treatment-resistant schizophrenia***

#### *Clozapine Study 30*

In a double-blind clinical trial performed in 319 treatment-resistant patients, clinically relevant improvement was observed within 6 weeks in about 30% of the clozapine-treated patients. Two open-label trials in which patients were treated for 12 months, showed clinically relevant improvement in 37% of patients within the first 6 weeks of treatment and in an additional 39%-44% of patients by the end of 12 months. The improvement was defined as a reduction of more than 20% from baseline in Brief Psychiatric Rating Scale Score. In addition, improvement in some aspects of cognitive dysfunction has been described.

## **5.2. Pharmacokinetic properties**

### Absorption

The absorption of orally administered clozapine is 90% to 95%; neither the rate nor the extent of absorption is influenced by food. Clozapine is subject to moderate first-pass metabolism, resulting in an absolute bioavailability of 50% to 60%.

### Distribution

In steady-state conditions, when given twice daily, peak blood levels occur on an average at 2.1 hours (range: 0.4 to 4.2 hours), and the volume of distribution is 1.6 L/kg. Clozapine is approximately 95% bound to plasma proteins.

### Biotransformation/metabolism

Clozapine is almost completely metabolised before excretion by CYP1A2 and 3A4, and to some extent by CYP2C19 and 2D6. Of the main metabolites, only the desmethyl metabolite was found to be active. Its pharmacological actions resemble those of clozapine, but are considerably weaker and of short duration.

### Elimination

Its elimination is biphasic, with a mean terminal half-life of 12 hours (range: 6 to 26 hours). After single doses of 75 mg, the mean terminal half-life was 7.9 hours; it increased to 14.2 hours when steady-state conditions were reached by administering daily doses of 75 mg for at least 7 days.

Only trace amounts of unchanged drug are detected in the urine and faeces, approximately 50% of the administered dose being excreted as metabolites in the urine and 30% in the faeces.

### Linearity/non-linearity

Dosage increases from 37.5 mg to 75 mg and 150 mg given twice daily were found to result during steady state in linearly dose-proportional increases in the area under the plasma concentration/time curve (AUC) and in the peak and minimum plasma concentrations.

### 5.3. Preclinical safety data

Preclinical data reveal no special hazard for humans based on conventional studies of safety pharmacology, repeated dose toxicity, genotoxicity and carcinogenic potential (for reproductive toxicity, see Fertility, pregnancy and lactation).

#### Mutagenicity

Clozapine and/or its metabolites were devoid of genotoxic potential when investigated for induction of gene mutations, chromosome aberrations and primary DNA-damage in a spectrum of in vitro mutagenicity tests. No clastogenic activity was observed in vivo (bone marrow micronucleus test in mice).

#### Carcinogenicity

In Sprague-Dawley (CD) rats treated in the diet for 24 months, maximum tolerated doses of 35 mg/kg per day revealed no carcinogenic potential of clozapine. Likewise, no evidence of tumorigenic effects was obtained in two 78-week feeding studies in Charles River (CD) mice. In the first study, oral dose levels of up to 64 mg/kg were administered to males, and of up to 75 mg/kg to females respectively. In the second study, the drug intake achieved for both sexes was 61 mg/kg per day.

#### Reproductive toxicity

No embryotoxic or teratogenic potential of clozapine was observed in rats or rabbits. In male rats treated for 70 days prior to mating, fertility was unaffected.

In female rats, fertility as well as pre- and postnatal development of the offspring was not adversely affected by oral clozapine treatment prior to mating. When rats were treated during the later part of pregnancy and during lactation, survival rates of the youngs from lactating dams, treated at dose levels up to 40 mg/kg body weight, were lowered and the youngs were hyperactive. However, there was no lasting effect on pup development after weaning.

## 6. PHARMACEUTICAL PARTICULARS

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### 6.1. List of excipients

**Clopine tablets** contain the following excipients:

Lactose monohydrate, magnesium stearate, microcrystalline cellulose, povidone, purified water, sodium starch glycolate

**Clopine 50 mg/mL oral suspension** contains the following excipients:

Glycerol, hydrochloric acid, monobasic sodium phosphate dihydrate, povidone, purified water, sodium hydroxide, sodium methyl hydroxybenzoate, sodium propyl hydroxybenzoate, sorbitol, xanthan gum

### 6.2. Incompatibilities

Not applicable

### 6.3. Shelf life

**Clopine tablets:** 60 months from date of manufacture

#### **Clopine 50 mg/mL oral suspension:**

**Unopened:** 36 months from date of manufacture

**Opened:** 3 months after opening

### 6.4. Special precautions for storage

**Clopine tablets:** Store at or below 30°C, Protect from light.

**Clopine 50 mg/mL oral suspension:** Store at or below 25°C.

### 6.5. Nature and contents of container

#### **Clopine tablets:**

Transparent PVC/PVDC/ Aluminium Foil Blister Strips in a cardboard carton containing 50 or 100 tablets.

HDPE bottles with polypropylene child-resistant, a tamper-evident cap containing 50 or 100 tablets.

Not all pack sizes may be marketed

#### **Clopine 50 mg/mL oral suspension:**

Bottle, glass, 125 mL amber bottle, polypropylene child-resistant, tamper-evident cap packaged with a graduated oral dispenser

### 6.6. Special precautions for disposal and other handling

Any unused medicine or waste material should be disposed of in accordance with local requirements.

## 7. MEDICINE SCHEDULE

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Prescription Medicine.

## 8. SPONSOR

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Douglas Pharmaceuticals Ltd

P O Box 45 027

Auckland 0651

New Zealand

Phone: (09) 835 0660

## 9. DATE OF FIRST APPROVAL

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**Clopine tablets:** 03 June 2010

**Clopine 50 mg/mL oral suspension:** 30 August 2012

## 10. DATE OF REVISION OF THE TEXT

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6 July 2017

Summary table of changes

<u>Section Changed</u>	<u>Summary of new information</u>
All	Revised to reflect SPC format
4.4, 4.8	Updated safety information regarding gastrointestinal hypomotility