

MYLOC CR

1. Product Name

Myloc CR, 23.75 mg, 47.5 mg, 95 mg and 190 mg, modified-release tablets.

2. Qualitative and Quantitative Composition

Each modified-release tablet contains 23.75 mg, 47.5 mg, 95 mg or 190 mg of metoprolol succinate.

For the full list of excipients, see section 6.1.

3. Pharmaceutical Form

Modified-release tablets of metoprolol succinate will be referred to as controlled release tablets in the following text.

Myloc CR tablets are a controlled delivery, prolonged release tablet formulation, containing membrane coated drug pellets embedded in a rapidly-disintegrating tablet.

Myloc CR 23.75 mg tablets: white to off-white, oblong, biconvex, film-coated tablets with break scores on both sides. The tablet can be divided into equal doses. Each tablet contains 23.75 mg of the active substance metoprolol succinate (corresponding to 9.71 mg of free drug base).

Myloc CR 47.5 mg tablets: white to off-white, oblong, biconvex, film-coated tablets with break scores on both sides. The tablet can be divided into equal doses. Each tablet contains 47.5 mg of the active substance metoprolol succinate (corresponding to 19.42 mg of free drug base).

Myloc CR 95 mg tablets: white to off-white, oblong, biconvex, film-coated tablets with break scores on both sides. The tablet can be divided into equal doses. Each tablet contains 95 mg of the active substance metoprolol succinate (corresponding to 38.84 mg of free drug base).

Myloc CR 190 mg tablets: white to off-white, oblong, biconvex, film-coated tablets with break scores on both sides. The tablet can be divided into equal doses. Each tablet contains 190 mg of the active substance metoprolol succinate (corresponding to 77.69 mg of free drug base).

4. Clinical Particulars

4.1 *Therapeutic indications*

- Hypertension. To reduce blood pressure and to reduce the risk of cardiovascular and coronary mortality (including sudden death), and morbidity
 - Angina pectoris
 - Symptomatic mild to severe chronic heart failure as an adjunct to other heart failure therapy to: increase survival, reduce hospitalisation, improve left ventricular function, improve New York Heart Association (NYHA) functional class and improve Quality of Life
 - Cardiac arrhythmias, especially supraventricular tachycardia, reduction of ventricular rate in atrial fibrillation and ventricular extrasystoles
 - Maintenance treatment after myocardial infarction
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- Hyperthyroidism
- Functional heart disorder with palpitations
- Migraine prophylaxis
- Paediatric hypertension.

4.2 Dose and method of administration

Dosage should always be adjusted to the patient's individual requirements.

Myloc CR is intended for once-daily treatment and can be taken with or without food. Myloc CR tablets should be swallowed with liquid. The tablets and the divided halves must not be chewed or crushed.

The following dosage recommendations may be taken as a guide:

Hypertension

The recommended dose in patients with mild to moderate hypertension is 47.5 mg Myloc CR given once daily. In patients not responding to 47.5 mg the dose can be increased to 95 to 190 mg once daily or combined with other antihypertensive agents.

Long-term antihypertensive treatment with metoprolol in daily doses of 95 to 190 mg has been shown to reduce total mortality, including sudden cardiovascular death, stroke and coronary events in hypertensive patients.

Angina pectoris

95 to 190 mg daily given as a single dose in the morning. Myloc CR can be combined with other antianginal agents if needed.

Chronic heart failure

The dose of Myloc CR should be individually adjusted in patients with chronic heart failure stabilised on other heart failure treatment. A recommended initial dose during the first two weeks is a 23.75 mg tablet once daily. It is recommended that patients with NYHA functional classes III-IV begin with half a 23.75 mg tablet once daily for the first week. It is recommended that the dose then be doubled every second week up to a maximum target dose of 190 mg Myloc CR once daily (or to the highest tolerated dose). During long-term treatment the aim should be to reach 190 mg Myloc CR once daily (or the highest tolerated dose).

At each dose level the patient should be carefully evaluated with regard to tolerability. In the case of hypotension, a decrease in concomitant medication may be necessary. Initial hypotension does not necessarily mean that the dose cannot be tolerated in chronic treatment but the patient should be kept at the lower dose until stabilised.

Cardiac arrhythmias

95 to 190 mg daily, given as a single dose once daily.

Maintenance treatment after myocardial infarction

Long-term oral treatment with metoprolol in doses of 190 mg given once daily has been shown to reduce the risk of death (including sudden death) and to reduce the risk of reinfarction (also in patients with diabetes mellitus).

Functional heart disorder with palpitations

The recommended dosage is 95 mg once daily. If necessary, the dose may be increased to 190 mg.

Migraine prophylaxis

The recommended dosage is 95 to 190 mg once daily.

Hyperthyroidism

95 to 190 mg daily, given as a single dose in the morning. If necessary, the dose may be increased.

Impaired renal function

Dose adjustment is not needed in patients with impaired renal function.

Impaired hepatic function

Dose adjustment is not normally needed in patients suffering from liver cirrhosis because metoprolol has low protein binding (5 - 10%). When there are signs of serious impairment of liver function (e.g. shunt-operated patients) a reduction in dose should be considered.

Elderly

Dose adjustment is not needed.

Hypertension in children

The recommended initial dosage in hypertensive patients ≥ 6 years is 0.95 mg/kg Myloc CR, not exceeding 47.5 mg once daily given approximated by dose strength. In patients not responding to 0.95 mg/kg, the dose can be increased to a maximum daily dose of 1.90 mg/kg. Doses above 190 mg once daily have not been studied in children and adolescents.

Efficacy and safety of use in children < 6 years have not been studied.

4.3 Contraindications

- Bronchial asthma or other obstructive lung disorders
- Grade 2 and 3 A-V block and intranodal A-V block
- Patients with unstable decompensated cardiac failure (pulmonary oedema, hypoperfusion or hypotension), and patients with continuous or intermittent inotropic therapy acting through beta-receptor agonism
- Marked clinically relevant sinus bradycardia
- Sick-sinus syndrome (unless a permanent pacemaker is in place)
- Cardiogenic shock
- Severe peripheral arterial circulatory disorder.

Metoprolol should not be given to patients with suspected acute myocardial infarction as long as the heart rate is < 45 beats/minute, the P-Q interval is > 0.24 seconds or the systolic blood pressure is < 100 mmHg.

Myloc CR is contraindicated in patients who have shown hypersensitivity to any component of the product or to other beta-blockers.

4.4 Special warnings and precautions for use

Intravenous administration of calcium antagonists of the verapamil-type should not be given to patients treated with beta-blockers.

The risk of Metoprolol CR interfering with beta₂-receptors is less than with conventional tablet formulations of beta₁-selective blockers.

The risk of interfering with carbohydrate metabolism or masking hypoglycaemia is likely to be less with Metoprolol CR than with conventional tablet formulations of beta₁-selective blockers and much less than with non-selective beta-blockers.

Very rarely a pre-existing A-V conduction disorder of moderate degree may become aggravated (possibly leading to A-V block). If the patient develops increasing bradycardia, Metoprolol CR should be given in lower doses or gradually withdrawn.

Metoprolol CR may aggravate the symptoms of peripheral arterial circulatory disorders, mainly due to its blood pressure lowering effect.

Where Metoprolol CR is prescribed for a patient known to be suffering from phaeochromocytoma, an alpha-blocker should be given concomitantly.

Abrupt withdrawal of beta-blockade is hazardous especially in high risk patients, and therefore should not be done. If there is a need to discontinue treatment with Metoprolol CR, this should preferably be done gradually over at least two weeks when the dose is reduced by half in each step down to the final step when a whole 23.75 mg tablet is reduced to half a tablet. If symptoms occur, a slower withdrawal rate is recommended. Sudden withdrawal of beta-blockade may aggravate chronic heart failure and also increases the risk for myocardial infarction and sudden death.

The anaesthetist should be informed that the patient is receiving Metoprolol CR prior to surgery. It is not recommended to stop beta-blocker treatment in patients undergoing surgery. Acute initiation of high-dose metoprolol to patients undergoing non-cardiac surgery should be avoided, since it has been associated with bradycardia, hypotension and stroke including fatal outcome in patients with cardiovascular risk factors.

In patients taking beta-blockers anaphylactic shock assumes a more severe form.

4.5 Interaction with other medicines and other forms of interaction

Metoprolol is a metabolic substrate for the cytochrome P450 isoenzyme CYP2D6. Drugs that act as enzyme-inducing and enzyme-inhibiting substances may exert an influence on the plasma level of metoprolol. Plasma levels of metoprolol may be raised by co-administration of compounds metabolised by CYP2D6 e.g. antiarrhythmics, antihistamines, histamine-2-receptor antagonists, antidepressants, antipsychotics and COX-2 inhibitors. The plasma concentration of metoprolol is lowered by rifampicin and may be raised by alcohol and hydralazine.

Patients receiving concomitant treatment with sympathetic ganglion blocking agents, other beta-blockers (i.e. eye drops) or monoamine oxidase inhibitors should be kept under close surveillance.

If concomitant treatment with clonidine is to be discontinued, the beta-blocker medication should be withdrawn several days before clonidine.

Increased negative inotropic and chronotropic effects may occur when metoprolol is given together with calcium antagonists of the verapamil and diltiazem type. In patients treated with beta-blockers, intravenous administration of calcium antagonists of the verapamil type should not be given.

Beta-blockers may enhance the negative inotropic and negative dromotropic effect of antiarrhythmic agents (of the quinidine type and amiodarone).

Digitalis glycosides, in association with beta-blockers, may increase atrioventricular conduction time and may induce bradycardia.

In patients receiving beta-blocker therapy, inhalation anaesthetics enhance the cardiodepressant effect.

Concomitant treatment with indomethacin or other prostaglandin synthetase inhibiting agents may decrease the antihypertensive effect of beta-blockers.

Under certain conditions, when adrenaline is administered to patients treated with beta-blockers, cardioselective beta-blockers interfere much less with blood pressure control than non-selective beta-blockers.

The dosages of oral antidiabetics may have to be readjusted in patients receiving beta-blockers.

4.6 Fertility, pregnancy and lactation

Pregnancy

As with most medicines, Metoprolol CR should not be given during pregnancy and lactation unless its use is considered essential. In general, beta-blockers reduce placental perfusion, which has been associated with growth retardation, intrauterine death, abortion and early labour. It is therefore suggested that appropriate maternofetal monitoring be performed in pregnant women treated with metoprolol. As with all antihypertensive agents, beta-blockers may cause side effects (e.g. bradycardia) in the foetus and in the newborn and breast-fed infant.

Breast-feeding

The amount of metoprolol ingested via breast-milk seems to be negligible as regards beta-blocking effect in the infant if the mother is treated with metoprolol doses within the normal therapeutic range.

4.7 Effects on ability to drive and use machines

Patients should know how they react to Metoprolol CR before they drive or use machines because occasionally dizziness or fatigue may occur.

4.8 Undesirable effects

Metoprolol CR is well tolerated and adverse reactions have generally been mild and reversible. The following events have been reported as adverse events in clinical trials or reported from routine use, mostly with conventional metoprolol (metoprolol tartrate). In many cases a relationship to treatment with metoprolol has not been established. The following definitions of frequencies are used: Very common ($\geq 10\%$), common (1 to 9.9%), uncommon (0.1 to 0.9%), rare (0.01 to 0.09%) and very rare ($<0.01\%$).

Cardiovascular system

Common: Bradycardia, postural disorders (very rarely with syncope), cold hands and feet, palpitations.

Uncommon: Deterioration of heart failure symptoms, cardiogenic shock in patients with acute myocardial infarction*, first degree heart block, oedema, precordial pain.

Rare: Disturbances of cardiac conduction, cardiac arrhythmias.

Very rare: Gangrene in patients with pre-existing severe peripheral circulatory disorders.

* Excess frequency of 0.4% compared with placebo in a study of 46,000 patients with acute myocardial infarction where the frequency of cardiogenic shock was 2.3% in the metoprolol group and 1.9% in the placebo group in the subset of patients with low shock risk index. The shock risk index was based on the absolute risk of shock in each individual patient derived from age, sex, time delay, Killip class, blood pressure, heart rate, ECG abnormality, and prior history of hypertension. The patient group with low shock risk index corresponds to the patient in which metoprolol is recommended for use in acute myocardial infarction.

Central Nervous System

Very common: Fatigue.

Common: Dizziness, headache.

Uncommon: Paraesthesiae, muscle cramps.

Gastrointestinal

Common: Nausea, abdominal pain, diarrhoea, constipation.

Uncommon: Vomiting.

Rare: Dry mouth.

Haematologic

Very rare: Thrombocytopenia.

Hepatic

Rare: Liver function test abnormalities.

Very rare: Hepatitis.

Metabolism

Uncommon: Weight gain.

Musculoskeletal

Very rare: Arthralgia.

Psychiatric

Uncommon: Depression, concentration impaired, somnolence or insomnia, nightmares.

Rare: Nervousness, anxiety, impotence / sexual dysfunction.

Very rare: Amnesia / memory impairment, confusion, hallucinations.

Respiratory

Common: Dyspnoea on exertion.

Uncommon: Bronchospasm.

Rare: Rhinitis.

Sense organs

Rare: Disturbances of vision, dry and/or irritated eyes, conjunctivitis.

Very rare: Tinnitus, taste disturbances.

Skin

Uncommon: Rash (in the form of urticaria, psoriasiform and dystrophic skin lesions), increased sweating.

Rare: Loss of hair.

Very rare: Photosensitivity reactions, aggravated psoriasis.

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

4.9 Overdose

Symptoms

Symptoms of overdosage may include hypotension, cardiac insufficiency, bradycardia and bradyarrhythmia, cardiac conduction disturbances and bronchospasm.

Management

Care should be provided at a facility that can provide appropriate supporting measures, monitoring and supervision.

If justified, gastric lavage and/or activated charcoal can be administered.

Atropine, adreno-stimulating medicines or pacemaker to treat bradycardia and conduction disorders.

Hypotension, acute cardiac failure, and shock to be treated with suitable volume expansion, injection of glucagon (if necessary, followed by an intravenous infusion of glucagon), intravenous administration of adreno-stimulating medicines such as dobutamine with α_1 -receptor agonistic medicines added in presence of vasodilation. Intravenous use of calcium ions can also be considered.

Bronchospasm can usually be reversed by bronchodilators.

For further advice on management of overdose please contact the National Poisons Information Centre (0800 POISON or 0800 764 766).

5. Pharmacological Properties

5.1 *Pharmacodynamic properties*

Pharmacotherapeutic group: Beta blocking agents, selective, ATC code: C07AB02.

Metoprolol is a beta₁-selective beta-blocker, i.e. it blocks beta₁-receptors at doses much lower than those needed to block beta₂-receptors.

Metoprolol has an insignificant membrane-stabilising effect and does not display partial agonist activity.

Metoprolol reduces or inhibits the agonistic effect on the heart of catecholamines (which are released during physical and mental stress). This means that the usual increase in heart rate, cardiac output, cardiac contractility and blood pressure, produced by the acute increase in catecholamines, is reduced by metoprolol. During high endogenous adrenaline levels metoprolol interferes much less with blood pressure control than non-selective beta-blockers.

Metoprolol CR gives an even plasma concentration time and effect profile (beta₁-blockade) over 24 hours in contrast to conventional tablet formulations of beta₁-selective blockers.

Due to the lack of pronounced peaks in plasma concentration, the clinical beta₁-selectivity is improved with the Metoprolol CR formulation when compared to the conventional tablet formulations of beta₁-selective blockers. Furthermore the potential risk for peak plasma concentration related side-effects, such as bradycardia and leg fatigue is reduced.

When mandatory, Metoprolol CR, in combination with a beta₂-agonist, may be given to patients with symptoms of obstructive pulmonary disease. When given together with a beta₂-agonist, Metoprolol CR in therapeutic doses interferes less than non-selective beta-blockers with the beta₂-mediated bronchodilation caused by the beta₂-agonist.

Metoprolol CR interferes less with insulin release and carbohydrate metabolism than do non-selective beta-blockers.

Metoprolol CR interferes much less with the cardiovascular response to hypoglycaemia than do non-selective beta-blockers.

Short term studies have shown that Metoprolol CR may cause a slight increase in triglycerides and a decrease in free fatty acids in the blood. In some cases, a small decrease in the high density lipoproteins (HDL) fraction has been observed, although to a lesser extent than that following non-selective beta-blockers. However, a significant reduction in total serum cholesterol levels has been demonstrated after metoprolol treatment in one study conducted over several years.

Quality of life is maintained, uncompromised or improved during treatment with Metoprolol CR. An improvement in quality of life has been observed after metoprolol treatment in patients after myocardial infarction. Furthermore, Metoprolol CR has been shown to improve Quality of Life in patients with chronic heart failure.

Clinical efficacy and safety

Effect in hypertension

Metoprolol CR lowers elevated blood pressure both in the standing and supine position. A short duration (a few hours) and clinically insignificant increase in peripheral resistance may be observed after the institution of metoprolol treatment. During long-term treatment total peripheral resistance may be reduced, due to reversal of hypertrophy in arterial resistance vessels. Long-term antihypertensive therapy with metoprolol has also been shown to reduce left ventricular hypertrophy and to improve left ventricular diastolic function and left ventricular filling.

In 144 paediatric patients (6 to 16 years of age) with essential hypertension, Metoprolol CR has been shown in a 4-week study to reduced placebo-corrected systolic blood pressure for the 0.95 and 1.90 mg/kg doses (4 to 6 mmHg). For diastolic blood pressure, there was a placebo-corrected reduction for the 1.90 mg/kg dose (5 mmHg) and a dose-dependent reduction for the dose range 0.19, 0.95 and 1.90 mg/kg. No apparent differences in blood pressure reduction were observed based on age, Tanner stage, or race.

In men with mild to moderate hypertension metoprolol has been shown to reduce the risk of death from cardiovascular disease, mainly due to a reduced risk for sudden cardiovascular death, to reduce the risk for fatal and non-fatal infarction and for stroke.

Effect on angina pectoris

In patients with angina pectoris metoprolol has been shown to reduce the frequency, duration and severity of both angina attacks and silent ischemic episodes and to increase the physical working capacity.

Effect in chronic heart failure

In patients with symptoms of heart failure (New York Heart Association (NYHA) II-IV) and decreased ejection fraction (≤ 0.40) Metoprolol CR when added to standard therapy has been shown to improve survival and to reduce the number of hospitalisations due to worsening heart failure. In addition, Metoprolol CR therapy has increased ejection fraction, reduced left ventricular end systolic and end diastolic volumes, improved NYHA functional class and improved Quality of Life.

In a clinical study treatment with metoprolol CR added to standard treatment with ACE inhibitors and diuretics in patients with decreased LVEF and symptoms of mild to severe chronic heart failure reduced:

- All cause mortality by 34% ($p=0.0062$ (adjusted); $p=0.00009$ (nominal))
- Combined endpoint of all cause mortality, and all cause hospitalisation (time to first event) by 19% ($p=0.00012$)
- Combined endpoint of all cause mortality, and hospitalisation due to worsening heart failure (time to first event) by 31% ($p \leq 0.00001$)
- Combined endpoint of death and heart transplantation (time to first event) by 32% ($p=0.0002$)
- Cardiovascular death by 38% ($p=0.00003$)
- Sudden death by 41% ($p=0.0002$)
- Death from worsening heart failure by 49% ($p=0.0023$)
- The pooled incidence of cardiac death and non-fatal acute MI by 39% ($p \leq 0.00001$)
- Combined endpoint of all cause mortality, hospitalisation due to worsening heart failure, and emergency room (ER) visit due to worsening heart failure (time to first event) by 32% ($p \leq 0.00001$)
- The number of hospitalisations due to worsening heart failure by 30% and the number of hospitalisations due to cardiovascular (CV) causes by 15% ($p=0.0003$).

Effect on cardiac rhythm

Metoprolol CR is suitable for regulating the heart rate as it inhibits the cardiac effects of increased sympathetic activity leading to decreased automaticity in the pacemaker cells and reduction of

supraventricular conduction velocity. Thus metoprolol controls heart rate in supraventricular tachycardia or atrial fibrillation, and in the presence of ventricular extrasystoles.

Effect on myocardial infarction

Metoprolol reduces mortality in patients with suspected or confirmed myocardial infarction mainly due to a reduction in the risk of sudden death. This effect is presumed to partly be due to the prevention of ventricular fibrillation. The anti-fibrillatory effect is believed to be due to a dual mechanism: a vagal effect within the blood-brain barrier beneficially influencing electrical stability of the heart, and a sympathetic direct cardiac anti-ischemic effect beneficially influencing contractility, heart rate and blood pressure. For both early and late intervention, the reduction in mortality is also present in high risk patients with previous cardiovascular disease; and in patients with diabetes mellitus.

Metoprolol has also been shown to reduce the risk for non-fatal myocardial infarction.

These anti-ischaemic effects of metoprolol are also reflected in a reduction in chest pain during the acute infarction phase. Metoprolol has also been shown to reduce the incidence of recurrent myocardial infarction.

Effect on hyperthyroidism

Metoprolol reduces the clinical manifestations in hyperthyroidism and can therefore be given as a supplementary medication.

Effect on heart disorders with palpitations

In patients with functional heart disorder with palpitations as a major symptom (which may be due to increased sympathetic activity) Metoprolol CR is effective in reducing palpitations and improving the patient's general condition.

Effect on migraine

Metoprolol CR is suitable for prophylactic treatment of migraine.

5.2 Pharmacokinetic properties

Absorption and distribution

Metoprolol CR is completely absorbed after oral administration. The systemic bioavailability of metoprolol from a single oral dose is approximately 50%, owing to an extensive first-pass effect. The bioavailability is reduced by about 20-30% for the controlled release preparation compared to the conventional tablets, but this has been demonstrated to be of no significance for clinical efficacy, since the area under the effect curve (AUEC) for heart rate is the same as with conventional tablets. The plasma protein binding of metoprolol is low, approximately 5-10% and has a volume of distribution of 5.6 L/kg.

The controlled release tablet consists of several hundred beads of metoprolol succinate. Each bead is coated with a retarding polymeric membrane which controls the rate of metoprolol release.

The tablet disintegrates rapidly after intake whereby the beads are dispersed in the gastrointestinal tract and release metoprolol continuously for about 20 hours. The elimination half-life of metoprolol averages 3.5 hours. Thus an even metoprolol plasma concentration is achieved over a dosage interval of 24 hours. The release rate is independent of physiological factors such as pH, food and peristalsis.

The rate of elimination is not affected by the formulation but, because of the prolonged absorption phase, administration of the CR tablet results in much less variation in plasma levels and pharmacological effects during a dosing interval compared with the plain tablet.

Metabolism and elimination

Metoprolol undergoes oxidative metabolism in the liver primarily by the CYP2D6 isoenzyme. The three main metabolites have been identified, though none of them have a clinically significant β -blocking effect.

As a rule over 95% of an oral dose can be recovered in the urine. About 5% of the given dose is excreted in the urine in unchanged form, this figure rising up to 30% in isolated cases. The elimination half-life of metoprolol in plasma averages 3.5 hours (extremes: 1 and 9 hours). The total clearance rate is approximately 1 litre/minute.

The elderly show no significant changes in the pharmacokinetics of metoprolol as compared to young persons. The systemic bioavailability and elimination of metoprolol is unchanged in patients with reduced renal function, however the excretion of metabolites is reduced. Significant accumulation of metabolites was observed in patients with a glomerular filtration rate (GFR) of less than 5 mL/minute. This accumulation of metabolites does not increase the beta-blockade.

The pharmacokinetics of metoprolol is little affected by decreased liver function due to its low protein binding. However, in patients with severe liver cirrhosis and a portacaval shunt, the bioavailability may increase and the total clearance may be reduced. Patients with a portacaval anastomosis had a total clearance of approximately 0.3 litres/minute and area under the plasma concentration-time curve (AUC) values of up to 6 times higher than in healthy subjects.

The pharmacokinetic profile of metoprolol in paediatric hypertensive patients aged 6 to 17 years is similar to the pharmacokinetics described previously in adults. Metoprolol apparent oral clearance (CL/F) increased linearly with body weight.

6. Pharmaceutical Particulars

6.1 *List of excipients*

- Sugar spheres (sucrose and maize starch)
- Macrogol 6000
- Polyacrylate dispersion 30%
- talc
- povidone
- microcrystalline cellulose
- magnesium stearate
- colloidal anhydrous silica
- hypromellose
- titanium dioxide.

6.2 *Incompatibilities*

Not applicable.

6.3 *Shelf life*

2 years.

6.4 *Special precautions for storage*

Store at or below 25°C.

6.5 *Nature and contents of container*

Myloc CR 23.75 mg, 47.5 mg, 90 mg and 190 mg tablets – blister packs of 30, 90 and 100 tablets.

Not all pack types and sizes may be marketed.

6.6 *Special precautions for disposal*

Not applicable.

7. Medicines Schedule

Prescription Medicine

8. Sponsor Details

Viatris Ltd
PO Box 11-183
Ellerslie
AUCKLAND
www.viatris.co.nz
Telephone 0800 168 169

9. Date of First Approval

16 September 2010

10. Date of Revision of the Text

23 January 2023

Summary table of changes

Section	Summary of new information
6.1	Removal of claims of gluten and lactose free.
6.5	Addition of 90 pack size
8	Sponsor details updated