1. PRODUCT NAME
NAROPIN 2 mg/mL WITH FENTANYL 2 microgram/mL (ropivacaine hydrochloride plus fentanyl citrate)

2. QUALITATIVE AND QUANTITATIVE COMPOSITION
Each mL of solution for infusion contains ropivacaine hydrochloride 2mg and fentanyl citrate 2microgram.

For full list of excipients, see section 6.1

3. PHARMACEUTICAL FORM
NAROPIN WITH FENTANYL is a sterile, isotonic solution with nominal osmolality of 288 mOsmol/kg and pH of 4.0 – 6.0.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications
Acute Pain Management: Continuous epidural infusion e.g. postoperative or labour pain.

4.2 Dose and method of administration
Tolerability varies between patients. The lowest dosage that results in effective analgesia should be used and should be based on the status of the patient and the analgesia required. Careful observation of the patient must be maintained during the infusion.

Adults
Recommended doses for NAROPIN WITH FENTANYL for injection in the average healthy 70 kg adult for up to 72 hours.

<table>
<thead>
<tr>
<th>Epidural administration Continuous infusion</th>
<th>Volume mL/hour</th>
<th>Dosage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Naropin 2 mg/mL + Fentanyl 2 microgram/mL</td>
<td>6-14</td>
<td>12-28 mg/hr</td>
</tr>
<tr>
<td></td>
<td></td>
<td>12-28 microgram/hr</td>
</tr>
</tbody>
</table>

Children
Until further experience has been gained, NAROPIN WITH FENTANYL cannot be recommended for use in children below the age of 12 years.

Elderly or debilitated patients
Debilitated or elderly patients, including those with partial or complete heart conduction block, advanced liver disease or severe renal dysfunction should be given reduced dosage commensurate with their physical condition. Clinical studies with this group of patients have not been performed. (See section 4.4).

NAROPIN WITH FENTANYL Data Sheet
Test Dose
For epidural analgesia, a 3-5 mL test dose of a local anaesthetic, preferably containing up to 5 microgram of adrenaline (e.g. Xylocaine 2% with Adrenaline 1:200,000), should be administered. Verbal contact and repeated monitoring of the heart rate and blood pressure should be maintained for 5 minutes following the test dose after which, in the absence of signs of intravascular or intrathecal injection, the main dose may be administered.

An inadvertent intravascular injection may be recognised by a temporary increase in heart rate and an accidental intrathecal injection by signs of a spinal block. If toxic symptoms or signs occur, the infusion should be stopped immediately.

Analgesia
When calculating the dosage for postoperative analgesia, the use of intraoperative local anaesthetics and opioids should be taken into account.

When prolonged blocks are used, either by continuous infusion or repeated bolus administration, the risks of reaching a toxic plasma concentration or inducing local neural injury must be considered. Cumulative doses of up to 800 mg ropivacaine for surgery and postoperative analgesia administered over 24 hours were well tolerated, as were postoperative continuous epidural infusions of ropivacaine with 2 microgram fentanyl at rates up to 14 mL/hour for 72 hours.

4.3 Contraindications
1. **Hypersensitivity**
   NAROPIN WITH FENTANYL solutions are contraindicated in patients with known hypersensitivity to local anaesthetics of the amide type or a known intolerance to fentanyl.

2. **Bronchial asthma.**
   See also section 4.4.

3. **Head injuries and increased intracranial pressure** –
   As for any opioid analgesic, fentanyl should not be used in patients susceptible to respiratory depression, such as comatose patients who may have head injuries or a brain tumour. Fentanyl may obscure the clinical course of patients with head injury.

4. **Concomitant MAO Inhibitors**
   Severe and unpredictable potentiation by MAO inhibitors has been reported with opioid analgesics and the use of fentanyl in patients who have received MAO inhibitors within 14 days is not recommended. (See section 4.5).

5. **Myasthenia gravis**
   Fentanyl may cause muscle rigidity upon IV administration. Therefore, the need for reversal and muscle relaxants contraindicates its use in patients with a history of myasthenia gravis.

4.4 Special warnings and precautions for use
Regional anaesthetic procedures should always be performed in a properly equipped and staffed area. Equipment and medicines necessary for monitoring and emergency resuscitation should be immediately available. Patients receiving major blocks should have an IV line inserted before the blocking procedure. The clinician responsible should be appropriately trained and familiar with diagnosis and treatment of side effects, systemic toxicity and other complications. (See section 4.9).
Patients in poor general condition due to ageing or other compromising factors such as partial or complete heart conduction block, advanced liver disease or severe renal dysfunction require special attention although regional anaesthesia is frequently the optimal anaesthetic technique in these patients. To reduce the risk of potentially serious adverse reactions, attempts should be made to optimise the patient’s condition before major blocks are performed, and the dosage should be adjusted accordingly. Patients treated with anti-arrhythmic drugs class III (e.g. amiodarone) should be under close surveillance and ECG monitoring considered, since cardiac effects may be additive.

There have been rare reports of cardiac arrest during the use of NAROPIN for epidural anaesthesia of peripheral nerve blockade, especially after unintentional accidental intravascular administration in elderly patients and in patients with concomitant heart disease. In some instances, resuscitation has been difficult. Should cardiac arrest occur, prolonged resuscitative efforts may be required to improve the possibility of a successful outcome.

NAROPIN is metabolised in the liver and should therefore be used with caution in patients with severe liver disease and repeated doses may need to be reduced due to delayed elimination. Normally there is no need to modify the dose in patients with impaired renal function when used for single dose or short term treatment. Acidosis and reduced plasma protein concentration, frequently seen in patients with chronic renal failure may increase the risk of systemic toxicity (see section 4.2).

Epidural anaesthesia may lead to hypotension and bradycardia. The risk of such effects can be reduced, e.g. by injecting a vasopressor. Hypotension should be treated promptly with a sympathomimetic, repeated as necessary.

Adequate facilities should be available for post-operative monitoring and ventilation. Resuscitative equipment, oxygen and a opioid antagonist should be readily available to manage apnoea.

NAROPIN is possibly porphyrinogenic and should only be prescribed to patients with acute porphyria when no safer alternative is available. Appropriate precautions should be taken in the case of vulnerable patients.

NAROPIN WITH FENTANYL should be used with caution in patients with severe impairment of pulmonary function because of the possibility of respiratory depression (e.g. chronic obstructive pulmonary disease, patients with decreased respiratory reserve, or any patient with potentially compromised respiration). In such patients, opioids may further decrease respiratory drive and increase airway resistance. Respiratory depression caused by opioid analgesics can be reversed by opioid antagonists. Consult individual product information (nalorphine or naloxone) before employing opioid antagonists.

Fentanyl may cause muscle rigidity, particularly involving the muscles of respiration. This effect is related to the dose and speed of injection and may be reduced by slow infusion. It is likely to arise following epidural injection. However, if this effect occurs, it may be managed by the use of assisted or controlled respiration and, if necessary, by administration of a neuromuscular blocking agent compatible with the patient’s condition. Nonepileptic myoclonic movements can occur.

Fentanyl should be administered with caution to patients with liver and kidney dysfunction because of the importance of these organs in the metabolism and excretion of drugs.

Fentanyl may produce bradycardia, which may be treated with atropine; however, it should be used with caution in patients with cardiac bradyarrhythmias.
As has been observed with all opioid analgesics, episodes suggestive of Sphincter of Oddi Spasm may occur with fentanyl.

**Concomitant use of Monoamine Oxidase Inhibitors (MAOIs)**
Fentanyl is not recommended for use in patients who require the concomitant administration of MAOIs due to the risk of serotonin toxicity (see section 4.5).

**Concomitant use of Benzodiazepines and other Central Nervous System (CNS) Depressants**
Profound sedation, respiratory depression, coma, and death may result from the concomitant use of fentanyl with benzodiazepines or other CNS depressants (e.g., non-benzodiazepine sedatives/hypnotics, anxiolytics, tranquilizers, muscle relaxants, general anaesthetics, medicines with antihistamine-sedating actions such as antipsychotics, other opioids, alcohol). Because of these risks, reserve concomitant prescribing of these drugs for use in patients for whom alternative treatment options are inadequate.

Observational studies have demonstrated that concomitant use of opioid analgesics and benzodiazepines increases the risk of medicine-related mortality compared to use of opioid analgesics alone. Because of similar pharmacological properties, it is reasonable to expect similar risk with the concomitant use of other CNS depressant drugs with opioid analgesics (see section 4.5).

If the decision is made to prescribe a benzodiazepine or other CNS depressant concomitantly with an opioid analgesic, prescribe the lowest effective dosages and minimum durations of concomitant use. In patients already receiving an opioid analgesic, prescribe a lower initial dose of the benzodiazepine or other CNS depressant than indicated in the absence of an opioid, and titrate based on clinical response. If an opioid analgesic is initiated in a patient already taking a benzodiazepine or other CNS depressant, prescribe a lower initial dose of the opioid analgesic, and titrate based on clinical response. Follow patients closely for signs and symptoms of respiratory depression and sedation.

Advise both patients and caregivers about the risks of respiratory depression and sedation when fentanyl is used with benzodiazepines or other CNS depressants (including alcohol and illicit drugs). Advise patients not to drive or operate heavy machinery until the effects of concomitant use of the benzodiazepine or other CNS depressant have been determined. Screen patients for risk of substance use disorders, including opioid abuse and misuse, and warn them of the risk for overdose and death associated with the use of additional CNS depressants including alcohol and illicit drugs (see section 4.5).

**4.5 Interaction with other medicines and other forms of interaction**
Ropivacaine should be used with caution in patients receiving other local anaesthetics or agents structurally related to amide-type local anaesthetics e.g. certain anti-arrhythmics, such as lignocaine and mexiletine, since the toxic effects are additive. Specific interactions studies with ropivacaine and anti-arrhythmic drugs class III (e.g. amiodarone) have not been performed, but caution is advised (see section 4.4).

**CYTOCHROME P450**
In healthy volunteers ropivacaine clearance was reduced by 77% during co-administration of fluvoxamine, a potent competitive inhibitor of P4501A2. CYP1A2 is involved in the formation of 3-hydroxy-ropivacaine, a major metabolite. Thus strong inhibitors of CYP1A2, such as fluvoxamine and enoxacin, given concomitantly with NAROPIN can cause a metabolic interaction leading to an increased ropivacaine plasma concentration. Prolonged
administration of ropivacaine should therefore be avoided in patients treated with strong inhibitors of CYP1A2 such as fluvoxamine and enoxacin.

**BENZODIAZEPINES AND OTHER CENTRAL NERVOUS SYSTEM (CNS) DEPRESSANTS**

Benzodiazepines and other CNS depressant drugs, e.g. barbiturates, neuroleptics, other sedatives/hypnotics, anxiolytics, tranquilizers, muscle relaxant general anaesthetics, drugs with antihistamines-sedating actions such as antipsychotics, other opioids and alcohol will have additive or potentiating effects with fentanyl. Due to additive pharmacologic effect, the concomitant use of benzodiazepines or other CNS depressants including alcohol, increases the risk of respiratory depression, profound sedation, coma, and death.

Reserve concomitant prescribing of these drugs for use in patients for whom alternative treatment options are inadequate. Limit dosages and durations to the minimum required. Follow patients closely for signs of respiratory depression and sedation (see Section 4.4).

**SEROTONERGIC MEDICINES INCLUDING MONOAMINE OXIDASE INHIBITORS (MAOIs)**

Caution is advised when fentanyl is coadministered with drugs that affect the serotonergic neurotransmitter systems.

The development of a potentially life-threatening serotonin syndrome may occur with the concomitant use of serotonergic drugs such as Selective Serotonin Reuptake Inhibitors (SSRIs) and Serotonin Norepinephrine Reuptake Inhibitors (SNRIs), and with drugs which impair metabolism of serotonin (including Monoamine Oxidase Inhibitors [MAOIs]), antidepressants, migraine medicines and sibutramine. This may occur within the recommended dose.

Signs of serotonin syndrome include confusion, agitation, fever, sweating, ataxia, hyperreflexia, diarrhoea and clonus (spontaneous, induced and myoclonus). Immediate withdrawal of the serotonergic medicines usually brings about a rapid improvement. Treatment depends on the nature and severity of the symptoms.

Severe and unpredictable potentiation of opiate effects and/or serotonergic effects by MAOIs has been reported with fentanyl. Fentanyl should not be used within 14 days after discontinuation of treatment with MAOIs. (See section 4.3).

**NITROUS OXIDE**

Nitrous oxide has been reported to produce cardiovascular depression when given with high doses of fentanyl.

**AMIODARONE**

Profound bradycardia, sinus arrest and hypotension have occurred when patients receiving amiodarone have been given fentanyl for anaesthesia.

### 4.6 Fertility, pregnancy and lactation

**Pregnancy**

Apart from obstetric use, there are no adequate data on the use of ropivacaine in pregnancy. Animal studies do not indicate direct or indirect harmful effects with respect to pregnancy, embryonal/foetal development, parturition or postnatal development (see PRECLINICAL SAFETY DATA).
Opioid analgesics may cause respiratory depression in the newborn infant. These products should only be used during labour after weighing the needs of the mother against the risk to the foetus. The safe use of fentanyl has not been established with respect to possible adverse effects upon foetal development. Therefore it should be used in women of childbearing potential only when in the judgement of the physician the potential benefits outweigh the possible hazards.

**Lactation**

Fentanyl may enter the maternal milk. Therefore, breastfeeding is not recommended for 24 hours following the administration of the medicine.

The excretion of ropivacaine or its metabolites in human milk has not been studied. Based on the milk/plasma concentration ratio in rats, the estimated daily dose to a pup will be about 4% of the dose given to the mother. Assuming that the milk/plasma concentration ratio in humans is of the same order, the total ropivacaine dose to which the baby is exposed by breast feeding is far lower than by exposure *in utero* in pregnant women at term.

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

Depending on the dose, NAROPIN WITH FENTANYL may have a very mild effect on mental function and co-ordination even in the absence of overt CNS toxicity and may temporarily impair locomotion and alertness.

### 4.8 Undesirable effects

Adverse reactions to ropivacaine are rare in the absence of overdosage or inadvertent intravascular injection. Adverse reactions to fentanyl are similar to those observed with other opioid agonist analgesics. The adverse reactions to NAROPIN WITH FENTANYL are similar to those observed with the individual agents.

These adverse reactions are in general dose related and may result from high plasma levels caused by excessive dosage, rapid absorption or may result from a hypersensitivity, idiosyncrasy or diminished tolerance on the part of the patient. They should be distinguished from the physiological effects of the nerve block itself e.g. a decrease in blood pressure and bradycardia during epidural anaesthesia.

Pronounced acidosis, hyperkalaemia or hypoxia in the patient may increase the risk and severity of toxic reactions.

The following list of adverse events is based upon experience with the monotherapies in their usual dosage range. These events are considered to be of clinical importance, regardless of causal relationship.

**Table of Adverse Drug Reactions**

(Pooled data from all types of blocks)

<table>
<thead>
<tr>
<th>Category</th>
<th>Adverse Effects</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Very common</strong></td>
<td>Vascular Disorders: Hypotension</td>
</tr>
<tr>
<td>(&gt;1/10)</td>
<td>Gastrointestinal Disorders: Nausea</td>
</tr>
<tr>
<td><strong>Common</strong></td>
<td>Nervous System Disorders: Paraesthesia, Dizziness, Headache&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>(&gt;1/100)</td>
<td>Cardiac Disorders: Bradycardia&lt;sup&gt;a&lt;/sup&gt;, Tachycardia</td>
</tr>
<tr>
<td></td>
<td>Vascular Disorders: Hypertension</td>
</tr>
</tbody>
</table>
Gastrointestinal Disorders: Vomiting

Renal and Urinary Disorders: Urinary retention, Oliguria

General Disorders and Administration Site Conditions: Temperature elevation, Rigor, Back pain, Chest Pain, Insomnia

Uncommon (>1/1,000)

Psychiatric Disorders: Anxiety

Nervous System Disorders: Symptoms of CNS toxicity (Convulsions, Grand mal convulsions, Seizures, Light headedness, Circumoral paraesthesia, Numbness of the tongue, Hyperacusis, Tinnitus, Visual disturbances, Dysarthria, Muscular twitching, Tremor), Hypoaesthesia

Vascular Disorders: Syncope

Respiratory, Thoracic and Mediastinal Disorders: Dyspnoea

General Disorders and Administration Site Conditions: Hypothermia

Rare (>1/10,000)

Cardiac Disorders: Cardiac arrest, Cardiac arrhythmias

General Disorders and Administration Site Conditions: Allergic reactions (anaphylactic reactions, angioneurotic oedema and urticaria)

Unknown

Cough, Constipation

a These reactions are more frequent after spinal anaesthesia.
b These symptoms usually occur because of inadvertent intravascular injection, overdose or rapid absorption (see section 4.9).

CLASS RELATED ADVERSE DRUG REACTIONS

This section includes complications related to anaesthetic technique regardless of the local anaesthetic used.

Neurological complications: Neuropathy and spinal cord dysfunctions (e.g. anterior spinal artery syndrome, arachnoiditis, cauda equina syndrome), have been associated with intrathecal and epidural anaesthesia.

Total spinal block: Total spinal block may occur if an epidural dose is inadvertently administered intrathecally, or if a too large an intrathecal dose is administered.

Neurological: Neuropathy and spinal cord dysfunction (e.g. anterior spinal artery syndrome, arachnoiditis, cauda equina syndrome) have been associated with regional anaesthesia, regardless of the local anaesthetic drug used. Convulsions have been observed following unintended intravascular injection of ropivacaine.

Other: Blurred vision, miosis, diaphoresis, post operative mental depression, paradoxical CNS excitation, delirium and spasm of the Sphincter of Oddi.
CLINICAL TRIALS USING NAROPIN 2 mg/mL WITH FENTANYL 2 microgram/mL

Clinical trials in patients undergoing epidural infusion for postoperative pain show that the following events were more common in patients receiving NAROPIN WITH FENTANYL than in the group receiving NAROPIN alone: pruritis, ileus, hypomagnesaemia, hypoglycaemia, atelectasis, urine abnormalities, laboratory test abnormalities.

The following events were less common in the NAROPIN WITH FENTANYL group than the plain NAROPIN group: hypothermia, chest pain, vasospasm, coughing.

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

The symptoms of NAROPIN WITH FENTANYL overdosage are expected to reflect those seen with overdosage of the individual drugs. Systemic toxic reactions may involve the central nervous system and the cardiovascular system.

Accidental intravascular injections of local anaesthetics may cause immediate (within seconds to a few minutes) systemic toxic reactions. In the event of overdose, systemic toxicity appears later (15-60 minutes after injection) due to the slower increase in local anaesthetic blood concentration.

SYMPTOMS

Acute Systemic Toxicity

Systemic toxic reactions primarily involve the central nervous system (CNS) and the cardiovascular system (CVS). Such reactions are caused by high blood concentration of a local anaesthetic, which may appear due to (accidental) intravascular injection, overdose or exceptionally rapid absorption from highly vascularised areas, (see section 4.4). CNS reactions are similar for all amide local anaesthetics, while cardiac reactions are more dependent on the drug, both quantitatively and qualitatively.

Central Nervous System Toxicity is a graded response with symptoms and signs of escalating severity. The first symptoms are usually light-headedness, circumoral paraesthesia, numbness of the tongue, hyperacusis, tinnitus and visual disturbances. Dysarthria, muscular twitching or tremors are more serious and precede the onset of generalised convulsions. These signs must not be mistaken for neurotic behaviour.

Unconsciousness and grand mal convulsions may follow, which may last from a few seconds to several minutes. Hypoxia and hypercarbia occur rapidly during convulsions due to the increased muscular activity, together with the interference with respiration and possible loss of functional airways. In severe cases apnoea may occur. Acidosis, hyperkalaemia, hypocalcaemia and hypoxia increase and extend the toxic effects of local anaesthetics.

Recovery follows the redistribution of the local anaesthetic agent from the central nervous system and subsequent metabolism and excretion. Recovery may be rapid unless large amounts of the agent have been injected.

Cardiovascular Toxicity may be seen in severe cases and is generally preceded by signs of toxicity in the central nervous system. In patients under heavy sedation or receiving a general anaesthetic, prodromal CNS symptoms may be absent. Hypotension, bradycardia, arrhythmia and even cardiac arrest may occur as a result of high systemic concentrations of
local anaesthetics, but in rare cases cardiac arrest has occurred without prodromal CNS effects.

Overdose due to fentanyl may result in narcosis (which may be preceded by marked skeletal muscle rigidity), cardiorespiratory depression accompanied by cyanosis, followed by a fall in body temperature, circulatory collapse, coma and possibly death.

TREATMENT
If signs of acute systemic toxicity appear, injection of the local anaesthetic should be stopped immediately and CNS symptoms (convulsion, CNS depression) must promptly be treated with appropriate airway/respiratory support and the administration of anticonvulsant drugs.

If circulatory arrest should occur, immediate cardiopulmonary resuscitation should be instituted. Optimal oxygenation and ventilation and circulatory support as well as treatment of acidosis are of vital importance.

If cardiovascular depression occurs (hypotension, bradycardia), appropriate treatment with intravenous fluids, vasopressor and or inotropic agents should be considered. Children should be given doses commensurate with their age and weight.

Should cardiac arrest occur, a successful outcome may require prolonged resuscitative efforts.

If depressed respiration is associated with muscular rigidity, an intravenous neuromuscular blocking agent might be required to facilitate assisted or controlled respiration.

The patient should be carefully observed for 24 hours; body warmth and adequate fluid intake should be maintained.

If severe or persistent hypotension occurs, the possibility of hypovolaemia should be considered and managed with appropriate parenteral fluid therapy.

A specific opioid antagonist, such as nalorphine or naloxone, should be available for use as indicated to manage respiratory depression. This does not preclude the use of more immediate countermeasures. The duration of respiratory depression following overdosage of fentanyl is usually longer than the duration of opioid antagonist action.

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

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Ropivacaine
Ropivacaine has both anaesthetic and analgesic effects. At high doses it produces surgical anaesthesia with motor block, while at lower doses it produces sensory block (analgesia) with limited and non-progressive motor block.

The duration and intensity of ropivacaine sensory block are not improved by the addition of adrenaline.
Ropivacaine, like other local anaesthetics, causes reversible blockade of impulse propagation along nerve fibres by preventing the inward movement of sodium ions through the cell membrane of the nerve fibres.

Local anaesthetics may have similar effects on other excitable membranes, e.g. in the brain and myocardium. If excessive amounts of medicine reach the systemic circulation, symptoms and signs of toxicity may appear, emanating from the central nervous and cardiovascular systems.

Cardiac effects measured in vivo in animal studies showed that ropivacaine has a lower cardiac toxicity than bupivacaine.

Pregnant ewes showed no greater sensitivity to ropivacaine than non-pregnant ewes.

Healthy volunteers exposed to intravenous infusions tolerated ropivacaine well. The clinical experience with this medicine indicates a good margin of safety.

Indirect cardiovascular effects (hypotension, bradycardia) may occur after epidural administration, depending on the extent of the concomitant sympathetic block.

**Fentanyl**

Fentanyl is an opioid analgesic. A dose of 100 microgram is approximately equivalent in analgesic activity to 10 mg of morphine or 75 mg of pethidine. The principal actions of therapeutic value are analgesia and sedation. Alterations in respiratory rate and alveolar ventilation associated with opioid analgesics may last longer than the analgesic effect. As the dose of opioid is increased, the decrease in pulmonary exchange becomes greater. Large doses may produce apnoea. Fentanyl appears to have less emetic activity than either morphine or pethidine.

Histamine assays and skin wheal testing in man indicate that clinically significant histamine release rarely occurs with fentanyl. Recent assays in man show no clinically significant histamine release at doses up to 50 microgram/kg. Fentanyl preserves cardiac stability and blunts stress-related hormonal changes at higher doses.

Fentanyl produces minimal cortical depression and may act by filling receptor sites located in the thalamus, midbrain and spinal cord. A specific morphine antagonist (e.g. nalorphine or naloxone) produces reversal of respiratory, cardiovascular, miotic and motor incoordination effects and also produces reversal of analgesia, euphoria and sedation. Rigidity of the diaphragm and intercostal muscles can be eliminated by suxamethonium. Cholinergic effects such as bradycardia are reversed by atropine.

As with longer-acting opioid analgesics, the duration of the respiratory depressant effect of fentanyl may be longer than the analgesic effect. The following observations have been reported concerning altered respiratory response to CO2 stimulation following administration of fentanyl to man:

1. Diminished sensitivity to CO2 stimulation may persist longer than depression of respiratory rate. Fentanyl frequently slows the respiratory rate.
2. Altered sensitivity to CO2 stimulation has been demonstrated for up to four hours following a single intravenous dose of 600 microgram fentanyl to healthy volunteers.
3. Duration and degree of respiratory depression is dose related.
4. The peak respiratory effect of a single intravenous dose of fentanyl is noted 5 to 15 minutes following injection. (See also section 4.4 concerning respiratory depression).
5.2 Pharmacokinetic properties

Ropivacaine

Ropivacaine has a chiral centre and is the pure S-(-)-enantiomer. Ropivacaine has a pKₐ of 8.1 and a distribution ratio of 141 (25°C n-octanol/phosphate buffer pH 7.4).

The plasma concentration of ropivacaine depends upon the dose, the route of administration and the vascularity of the injection site. Ropivacaine follows linear pharmacokinetics and the maximum plasma concentration is proportional to the dose.

Ropivacaine shows complete and biphasic absorption from the epidural space with half-lives of the two phases of the order of 14 minutes and 4 hours. The slow absorption is the rate-limiting factor in the elimination of ropivacaine, which explains why the apparent elimination half-life is longer after epidural than after intravenous administration.

The pharmacokinetic profile of ropivacaine following experimental IV administration is summarised below:

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Plasma clearance</td>
<td>440 mL/min</td>
</tr>
<tr>
<td>Unbound plasma clearance</td>
<td>8 L/min</td>
</tr>
<tr>
<td>Renal clearance</td>
<td>1 mL/min</td>
</tr>
<tr>
<td>Volume of distribution at steady-state</td>
<td>47 L</td>
</tr>
<tr>
<td>Unbound volume of distribution at steady-state</td>
<td>819 L</td>
</tr>
<tr>
<td>Terminal half-life</td>
<td>1.8 h</td>
</tr>
<tr>
<td>Unbound fraction</td>
<td>0.06</td>
</tr>
<tr>
<td>Hepatic extraction ratio</td>
<td>0.4</td>
</tr>
<tr>
<td>Major metabolite</td>
<td>3-OH-ropivacaine</td>
</tr>
</tbody>
</table>

Ropivacaine is mainly bound to α₁-acid glycoprotein in plasma with an unbound fraction of about 6%. An increase in total plasma concentrations during continuous epidural infusion has been observed, related to a postoperative increase of α₁-acid glycoprotein.

Variations in unbound, i.e. pharmacologically active, concentration have been much less than in total plasma concentration.

Ropivacaine readily crosses the placenta and equilibrium in regard to unbound concentration will be rapidly reached. The degree of plasma protein binding in the foetus is less than in the mother, which results in lower total plasma concentrations in the foetus.

Ropivacaine is extensively metabolised, predominantly by aromatic hydroxylation. In total 86% of the dose is excreted in the urine after intravenous administration of which only about 1% relates to unchanged medicine. The major metabolite is 3-hydroxyropivacaine, about 37% of which is excreted in the urine, mainly conjugated. Urinary excretion of 4-hydroxyropivacaine, the N-dealkylated metabolite and the 4-hydroxy-dealkylated accounts for 1-3%. Conjugated and unconjugated 3-hydroxyropivacaine shows only detectable concentrations in plasma. 3-hydroxy- and 4-hydroxyropivacaine have a local anaesthetic activity although less than that of ropivacaine.

During epidural infusion, both PPX and 3-hydroxy-ropivacaine are the major metabolites excreted in the urine. Total PPX concentration in the plasma was about half of that of total ropivacaine, however, mean unbound concentrations of PPX was about 7 to 9 time higher than that of unbound ropivacaine following continuous epidural infusion up to 72 hours. The threshold for CNS-toxic unbound plasma concentrations of PPX in rats is about twelve times higher than that of unbound ropivacaine.
Impaired renal function has little or no influence on ropivacaine pharmacokinetics. The renal clearance of PPX is significantly correlated with creatinine clearance. A lack of correlation between total exposure, expressed as AUC, with creatinine clearance indicates that the total clearance of PPX includes a non-renal elimination in addition to renal excretion. Some patients with impaired renal function may show an increased exposure to PPX resulting from a low non-renal clearance. Due to the reduced CNS toxicity of PPX as compared to ropivacaine the clinical consequences are considered negligible in short-term treatment.

There is no evidence of *in vivo* racemisation of ropivacaine.

**Fentanyl**

The pharmacokinetics of fentanyl can be described by a three-compartment model, with a distribution time of 1.7 minutes, redistribution of 13 minutes and a terminal elimination half-life of 219 minutes. The volume of distribution for fentanyl is 4 L/kg.

Fentanyl plasma protein binding capacity increases with increasing ionisation of the drug. Alterations in pH may affect its distribution between plasma and the central nervous system. It accumulates in skeletal muscle and fat, and is released slowly into the blood.

Fentanyl is primarily transformed in the liver and demonstrates a high first pass clearance with approximately 75% of an intravenous dose excreted in urine, primarily as metabolites with less than 10% representing the unchanged drug. Approximately 9% of the dose is recovered in the faeces, primarily as metabolites.

6. **PHARMACEUTICAL PARTICULARS**

6.1 **List of excipients**

Sodium chloride
Water for injection
Hydrochloric acid and sodium hydroxide for pH adjustment.

6.2 **Incompatibilities**

Not applicable

6.3 **Shelf life**

24 months.

For storage after opening, see section 6.6.

6.4 **Special precautions for storage**

Store below 30°C. Do not freeze.

6.5 **Nature and contents of container**

100 mL & 200 mL Polybag® infusion bags in Sterile AstraZeneca Theatre Pack™ (5's)

6.6 **Special precautions for disposal and other handling**

*NAROPIN with FENTANYL* injection solutions do not contain a preservative. Each Polybag is intended for single use only, not exceeding 24 hours. Any solution remaining from an opened Polybag should be discarded.
The intact container must not be re-autoclaved. A blistered container should be chosen when a sterile outside is required.

7. MEDICINE SCHEDULE
Controlled Drug B3.

8. SPONSOR
Pharmacy Retailing (NZ) Limited trading as Healthcare Logistics
58 Richard Pearse Drive
Airport Oaks
Auckland
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Telephone: (09) 918 5100
Email: aspen@aspenpharma.co.nz

9. DATE OF FIRST APPROVAL
3 February 2000

10. DATE OF REVISION OF THE TEXT
27 April 2018

SUMMARY TABLE OF CHANGES

<table>
<thead>
<tr>
<th>Section changed</th>
<th>Summary of new information</th>
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<tbody>
<tr>
<td>4.4</td>
<td>Update of warning regarding hypotension or bradycardia</td>
</tr>
<tr>
<td>4.5</td>
<td>Update on serotonin syndrome information</td>
</tr>
<tr>
<td>4.8</td>
<td>Additional information on undesirable effects</td>
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