

HYDRORPHONE

Warnings
high alert medication
(see Potential Hazards section)

Classification
opioid analgesic

Alternate Names
DILAUDID

Indications

- symptomatic relief of severe pain
- treatment of opioid use disorder
- [Indications and/or prescribing restrictions apply](#)

Reconstitution and Stability

- store vials at room temperature and protect from light
- a yellowish discolouration may develop; this has not been associated with a loss of potency

Compatibility

- compatible with dextrose 5%, sodium chloride 0.9%, dextrose-sodium chloride combinations, and lactated ringers

Preparation and Administration

Administration Route	Approved	Preparation and Administration Instructions	Required Monitoring
Subcutaneous	YES	<i>opioid use disorder:</i> <ul style="list-style-type: none">▪ doses less than 25 mg, use 10 mg/mL vial▪ doses of 25 mg or more, use 50 mg/mL vial	Intermediate Monitoring
Continuous Subcutaneous Infusion	YES	standard solution concentrations: <u>2 mg/mL:</u> 100 mg in 50 mL; 200 mg in 100 mL <u>10 mg/mL:</u> <ul style="list-style-type: none">○ remove 10 mL from 50 mL bag, then add 500 mg to bag○ remove 20 mL from 100 mL bag, then add 1000 mg to bag <u>50 mg/mL:</u> transfer 2500 mg to an empty sterile bag (empty bag obtained from stores or pharmacy) <i>rate must be controlled with an infusion pump</i>	Intermediate Monitoring
Intramuscular	YES	<i>opioid use disorder:</i> <ul style="list-style-type: none">▪ doses less than 25 mg, use 10 mg/mL vial▪ doses of 25 mg or more, use 50 mg/mL vial	Intermediate Monitoring

Preparation and Administration table continues on next page. This monograph has 3 pages.

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continuation of Preparation and Administration table

Administration Route	Approved	Preparation and Administration Instructions	Required Monitoring
IV direct	YES	<p><u>1 mg or less</u>: dilute 2 mg (1 mL from 2 mg/mL vial) with 9 mL NS or SWFI to make a 0.2 mg/mL concentration; discard excess volume in syringe before administration of calculated dose.</p> <p><u>doses greater than 1 mg</u>: use either 2 mg/mL or 10 mg/mL vial; dilute dose to 5 mL with NS or SWFI</p> <p>administer <u>slowly</u> over at least 2 to 3 minutes</p> <ul style="list-style-type: none"> ○ <i>opioid naïve patients</i>, doses of 0.5 mg or greater should be administered IV intermittent ○ <i>highly opioid tolerant patients</i>, 50 mg/mL vial may be used on a case specific basis ○ <i>opioid use disorder</i>: <ul style="list-style-type: none"> ▪ IV doses must be given IV direct ▪ doses less than 25 mg, use 10 mg/mL vial ▪ doses of 25 mg or more, use 50 mg/mL vial ▪ administer undiluted, flush slowly over 2 to 3 minutes: <ul style="list-style-type: none"> with 10 mL NS flush (for peripheral IV) with 20 mL NS flush (for central line e.g. PICC/CVC) 	Intermediate Monitoring
IV intermittent	YES	dilute in 50 mL; infuse over 15 to 30 minutes	Intermediate Monitoring
Continuous IV infusion	YES	<p>standard solution concentrations:</p> <p><u>0.5 mg/mL</u>: 50 mg in 100 mL</p> <p><u>2 mg/mL</u>: 100 mg in 50 mL; 500 mg in 250 mL</p> <p><u>10 mg/mL</u>:</p> <ul style="list-style-type: none"> ○ remove 10 mL from 50 mL bag, then add 500 mg to bag ○ remove 20 mL from 100 mL bag, then add 1000 mg to bag <p><u>50 mg/mL</u>: transfer 2500 mg or 5000 mg to an empty sterile bag (empty bag obtained from stores or pharmacy)</p> <p><i>rate must be controlled with an IV infusion pump</i></p>	Intermediate Monitoring
Patient Controlled Analgesia (PCA)	YES	<p>syringe standard solution concentration:</p> <p><u>0.6 mg/mL</u> in NS; <i>rate must be controlled with a PCA pump</i></p> <p>BC Women's Hospital: <u>0.2 mg/mL</u> in NS; <i>via CADD pump</i></p>	Intermediate Monitoring or per PCA site protocol
Epidural	YES	<p>bolus by physician only or per epidural site protocol</p> <p>continuous epidural infusion - initiated by physician (use preservative-free solution)</p> <p>standard solution concentration:</p> <p><u>10 mcg/mL</u>: in NS 250 mL</p> <p><i>rate must be controlled with an epidural pump</i></p>	Intermediate Monitoring or per epidural site protocol
Intrathecal	YES	<p>initiated by physician (use preservative-free solution)</p> <p>standard solution concentrations:</p> <p><u>0.1 mg/mL</u>: 10 mg in NS 100 mL</p> <p><u>0.2 mg/mL</u>: 20 mg in NS 100 mL</p>	Intermediate Monitoring or per intrathecal site protocol

This monograph has 3 pages

HYDRORomphone

Dosage

Dosing is individualized based on patient response and tolerance.

Usual dosing for opioid naïve patient:

- **IV direct:** 0.1 to 0.4 mg every 6 to 10 minutes until analgesia achieved; usual maximum 2 mg/hour
- **subcutaneous and IM:** 0.5 to 1 mg every 3 to 4 hours
- **IV intermittent:** 0.5 to 1 mg every 3 to 4 hours
- **continuous IV infusion or continuous subcutaneous infusion:** 0.1 to 0.6 mg/hour titrated to response; there is no maximum dose (palliative patients may require greater than 20 mg/hour)
- **epidural:** 0.5 to 2 mg; may follow with epidural infusion of 0.1 to 0.2 mg/hour as per site specific protocol
- **intrathecal:** dose as per site protocol.

Opioid tolerant patient:

- *opioid use disorder:*
 - Single doses up to 200 mg or higher IV Direct/IM/SUBCUT and daily total doses of up to 500 mg or higher have been used.
 - Dose is determined based on previous use, and no maximum dose has been established.

Switching from oral to parenteral routes:

- subcutaneous or IM dose is one-half the oral immediate release dose given at the same interval
- continuous IV or subcutaneous infusion dose is one-half the oral dose per 24 hours

Potential Hazards of Parenteral Administration

- respiratory/circulatory depression and orthostatic hypotension (may be increased with rapid IV injection)
- pain and/or burning at the subcutaneous injection site
- mild erythema may occur at the IM injection site
- antidote: naloxone
- **confirm resuscitative equipment is readily available**
- high alert medication:
 - HYDRORomphone vials or ampoules containing more than 2 mg
 - when given by these routes: epidural infusions, intrathecal, patient controlled analgesia

Important Implications

Contraindications/Cautions

- caution in patients with convulsive disorders, cranial injuries, respiratory insufficiency, cardiac arrhythmias, reduced blood volume,
- geriatric or debilitated patients, patients with hepatic dysfunction or severe renal dysfunction and those on other CNS depressants should receive reduced doses and carefully titrated.
- Pregnancy/Lactation: refer to Lexicomp or Micromedex

Side effects

- respiratory depression, circulatory depression and orthostatic hypotension
- restlessness, nightmares, tremors, euphoria, dizziness, confusion
- nausea, vomiting, constipation, pruritis

Monitoring

- severity of pain, time and frequency of breakthrough pain

Other

- HYDRORomphone is at least 5 times more potent than morphine.