



ACI NSW Agency
for Clinical
Innovation



Ketamine Infusion Prescription and Observation Chart - adult Education Slide Presentation

A presentation prepared by the
Pain Interest Group Nursing Issues
in association with the Agency of Clinical Innovation
Pain Management Network

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MARCH 31, 2017

Ketamine infusion Prescription and Observation Chart - adult

The ketamine infusion prescription and observation chart for adult patients has been developed by a team of experts in the field of acute pain including clinical nurse consultants, anaesthetists and pharmacy representatives.

This chart has been designed for the administration of subcutaneous or intravenous ketamine infusions for patients who may have acute or chronic pain. This chart may not be appropriate for ketamine infusion administration in palliative care.

Standardisation of this chart promotes best practice in pain assessment and management of adverse effects in those patients who are receiving a ketamine infusion for pain management.

Aim of this presentation:

This presentation aims to explain

- **how to use the chart** to record the prescription, administration and discarding of ketamine as a continuous infusion.
- **how to complete** the clinical observations.
- **guidelines on the management of patients** who are receiving ketamine as a continuous infusion including the management of adverse effects.

Ketamine Infusion Prescription and Observation Chart - adult

NSW Health

KETAMINE INFUSION (ADULT)

Ketamine Infusion Management Guidelines
(For detailed information regarding PCA prescribing and management refer to local hospital PCA policy)

- Observations on this form to be recorded either 2 hourly or 4 hourly as indicated on the prescription section of this form or more frequently if patient's clinical condition warrants.
 - If PCA (Patient controlled analgesia) in use, document pain scores on the PCA chart only. Record observations according to PCA management guidelines.
- Infusion pump settings to be checked at the commencement of each shift, on patient transfer and when the syringe or bag is changed.
- The cannula site (subcutaneous or intravenous) must be checked each shift for signs of redness, swelling or tenderness.

Managing Adverse Effects

- Managing dysphoric effects such as hallucinations, unpleasant dreams or visual disturbances: contact the relevant pain service or equivalent medical officer. A medical officer may consider a dose reduction of the ketamine infusion or the addition of a benzodiazepine. (e.g. midazolam)

REFER TO YOUR LOCAL CLINICAL EMERGENCY RESPONSE SYSTEM (CERS) PROTOCOL FOR INSTRUCTIONS ON HOW TO MAKE A CALL TO ESCALATE CARE FOR YOUR PATIENT

APPROPRIATE CLINICAL CARE FOR PATIENTS WITH YELLOW ZONE OBSERVATIONS:

- ENSURE THAT THE ACUTE PAIN SERVICE OR EQUIVALENT MEDICAL OFFICER IS CONTACTED

YELLOW ZONE RESPONSE

IF YOUR PATIENT HAS ANY YELLOW ZONE OBSERVATIONS YOU **MUST** FOLLOW THE YELLOW ZONE RESPONSE INSTRUCTIONS ON THE NSW STANDARD OBSERVATION CHARTS AND INITIATE APPROPRIATE CLINICAL CARE AS STATED ABOVE

Front page:

- Ketamine Infusion
- Management Guidelines
- Yellow Zone Response instructions

NO WRITING Page 1 of 8

Attach ADR Sticker

ALLERGIES & ADVERSE DRUG REACTIONS (ADR)

Not known Unknown (Use appropriate box or complete details below)

Drug (or other)	Reaction/Type/Date	Initials

Sign: _____ Print: _____ Date: _____

Ketamine Infusion (Adult)
Prescription is valid for a maximum of 7 days unless ceased earlier.
Refer to local hospital policy for standardised Ketamine solutions
Observations for this patient to be recorded: 2 hourly OR 4 hourly.

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate:mg per hour =mL per hour Infusion range (if applicable): FROM:mg per hour =mL per hour TO:mg per hour =mL per hour

Date: _____ Prescriber's signature: _____ Print your name: _____ Contact: _____ Pharmacy: _____

Revised prescription is valid until this chart is completed unless ceased earlier

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate:mg per hour =mL per hour Infusion range (if applicable): FROM:mg per hour =mL per hour TO:mg per hour =mL per hour

Date: _____ Prescriber's signature: _____ Print your name: _____ Contact: _____ Pharmacy: _____

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Infusion start rate:mg per hour =mL per hour Infusion range (if applicable): FROM:mg per hour =mL per hour TO:mg per hour =mL per hour

Date: _____ Prescriber's signature: _____ Print your name: _____ Contact: _____ Pharmacy: _____

Revised prescription is valid until this chart is completed unless ceased earlier

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate:mg per hour =mL per hour Infusion range (if applicable): FROM:mg per hour =mL per hour TO:mg per hour =mL per hour

Date: _____ Prescriber's signature: _____ Print your name: _____ Contact: _____ Pharmacy: _____

CEASE KETAMINE ACCORDING TO INSTRUCTIONS IN THE MEDICAL RECORD
Refer to entry in the medical record written on _____ Date: _____ Time: _____

Page 2 of 8 NO WRITING

Page 1

Page 2



Agency clinical vation

Ketamine Infusion Management Guidelines:

Ketamine Infusion Management Guidelines

Managing adverse effects

Yellow Zone Response instructions



Facility:

KETAMINE INFUSION (ADULT)

FAMILY NAME		MRN
GIVEN NAME		<input type="checkbox"/> MALE <input type="checkbox"/> FEMALE
D.O.B. ____/____/____	M.O.	
ADDRESS		
LOCATION / WARD		
COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE		

Ketamine Infusion Management Guidelines

(For detailed information regarding PCA prescribing and management refer to local hospital PCA policy)

- Observations on this form to be recorded either 2 hourly or 4 hourly as indicated on the prescription section of this form or more frequently if patient's clinical condition warrants.
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Managing Adverse Effects

- Managing dysphoric effects such as hallucinations, unpleasant dreams or visual disturbances: contact the relevant pain service or equivalent medical officer. A medical officer may consider a dose reduction of the ketamine infusion or the addition of a benzodiazepine. (e.g. midazolam)

REFER TO YOUR LOCAL CLINICAL EMERGENCY RESPONSE SYSTEM (CERS) PROTOCOL FOR INSTRUCTIONS ON HOW TO MAKE A CALL TO ESCALATE CARE FOR YOUR PATIENT

APPROPRIATE CLINICAL CARE FOR PATIENTS WITH YELLOW ZONE OBSERVATIONS:

- ENSURE THAT THE ACUTE PAIN SERVICE OR EQUIVALENT MEDICAL OFFICER IS CONTACTED

YELLOW ZONE RESPONSE

IF YOUR PATIENT HAS ANY YELLOW ZONE OBSERVATIONS YOU **MUST** FOLLOW THE YELLOW ZONE RESPONSE INSTRUCTIONS ON THE NSW STANDARD OBSERVATION CHARTS AND INITIATE APPROPRIATE CLINICAL CARE AS STATED ABOVE

KETAMINE INFUSION (ADULT)



ALLERGIES & ADVERSE DRUG REACTIONS (ADR)
 All known Unknown (tick appropriate box or complete details below)

Drug (or other)	Reaction/Type/Date	Initials

Sign.....Print.....Date.....

FAMILY NAME _____ MRN _____
 MALE FEMALE
 GIVEN NAME _____
 D.O.B. ____/____/____ M.O. _____
 ADDRESS _____
 LOCATION / WARD _____

COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE
 First Prescriber to Print Patient Name and Check Label Correct: _____ Pain specialist referral
 Referring doctor name: _____
 Signature: _____
 Date: _____

Ketamine Prescription:

Patient allergy status and patient label

Ketamine Infusion (Adult)

Prescription is valid for a maximum of 7 days unless ceased earlier.
 Refer to local hospital policy for standardised Ketamine solutions
 Observations for this patient to be recorded: 2 hourly OR 4 hourly.

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate _____mg per hour = _____mL per hour
 Infusion range (if applicable) FROM: _____mg per hour = _____mL per hour TO: _____mg per hour = _____mL per hour

Date _____ Prescriber's signature _____ Print your name _____ Contact _____ Pharmacy _____

Revised prescription is valid until this chart is completed unless ceased earlier

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate _____mg per hour = _____mL per hour
 Infusion range (if applicable) FROM: _____mg per hour = _____mL per hour TO: _____mg per hour = _____mL per hour

Date _____ Prescriber's signature _____ Print your name _____ Contact _____ Pharmacy _____

Revised prescription is valid until this chart is completed unless ceased earlier

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate _____mg per hour = _____mL per hour
 Infusion range (if applicable) FROM: _____mg per hour = _____mL per hour TO: _____mg per hour = _____mL per hour

Date _____ Prescriber's signature _____ Print your name _____ Contact _____ Pharmacy _____

Revised prescription is valid until this chart is completed unless ceased earlier

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
			Sodium chloride 0.9%		(mg per mL)

Infusion start rate _____mg per hour = _____mL per hour
 Infusion range (if applicable) FROM: _____mg per hour = _____mL per hour TO: _____mg per hour = _____mL per hour

Date _____ Prescriber's signature _____ Print your name _____ Contact _____ Pharmacy _____

CEASE KETAMINE ACCORDING TO INSTRUCTIONS IN THE MEDICAL RECORD

Refer to entry in the medical record written on _____ Date:_____ Time:_____

- Ketamine prescription:**
- Route
 - Amount
 - Diluent
 - Total volume
 - Concentration
 - Infusion start rate
 - Infusion range (if used)
 - Date, prescriber, pharmacy

Three revised prescription options for alterations to ketamine prescription

CEASE instructions



Prescription: Patient identification

Prescriber to complete patient allergy and ADR section in full

Handwritten patient details or affix patient label
(First prescriber to check patient label is correct)

Attach ADR Sticker

ALLERGIES & ADVERSE DRUG REACTIONS (ADR)		
<input type="checkbox"/> Nil known <input type="checkbox"/> Unknown (tick appropriate box or complete details below)		
Drug (or other)	Reaction/Type/Date	Initials
Sign. TSmith	Print. SMITH	Date. 5/4/17

FAMILY NAME		MRN
GIVEN NAME		<input type="checkbox"/> MALE <input type="checkbox"/> FEMALE
D.O.B. ____ / ____ / ____	M.O.	
ADDRESS		
LOCATION / WARD		
COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE		
First Prescriber to Print Patient Name and Check Label Correct:	Pain specialist referral	
.....	Referring doctor name:	
.....	Signature:	
.....	Date:	

PRIVATE PATIENTS:
For Medicare billing purposes: A pain specialist referral from the referring doctor (name), signature and date

Ketamine Infusion (Adult)

Prescription is valid for a maximum of 7 days unless ceased earlier.
Refer to local hospital policy for standardised Ketamine solutions

Observations for this patient to be recorded: 2 hourly OR 4 hourly.

Prescription is valid for a maximum of 7 days unless ceased earlier

The frequency of observations
(2 hourly or 4 hourly)
to be determined by the
medical officer

Ketamine prescription:

**PLEASE REFER TO YOUR LOCAL HOSPITAL POLICY OR
PROCEDURE FOR PREFERRED KETAMINE DELIVERY OPTIONS**

Route	Drug (print 'ketamine')	Amount (mg)	Diluent	Total Volume (mL)	Concentration
IV	Ketamine	200 mg	Sodium chloride 0.9%	50mL	4 (mg per mL)
Infusion start rate		Infusion range (if applicable)			
4mg per hour = 1mL per hour		FROM: 4mg per hour = 1mL per hour TO: 8mg per hour = 2mL per hour			
Date	Prescriber's signature	Print your name		Contact	Pharmacy
04/4/17	<i>TSmith</i>	SMITH		7511	

Three further prescription boxes are provided for when alterations are indicated for the ketamine dose, concentration or infusion rate.

Administration and discard of ketamine:

Record of ketamine administration and ketamine discarded

Record of ketamine administration					Record of ketamine discarded				
	Date	Time	Signature 1	Signature 2	Date	Time	Total ketamine discarded (mL or mg)	Signature 1	Signature 2
1	4/4/17	10:00	<i>THall</i>	<i>SRose</i>	05/4/17	10:00	8mL	BLoh	JLucas
2									

Any ketamine remaining from a syringe or bag to be recorded on the discard section

There are 13 more rows provided for more than one syringe or bag of ketamine that is commenced

FAMILY NAME	MRN
GIVEN NAME	<input type="checkbox"/> MALE <input type="checkbox"/> FEMALE
D.O.B. ____/____/____	M.O.
ADDRESS	
LOCATION	

A patient label must be affixed or details handwritten on each page that has recorded observations

Facility:

KETAMINE INFUSION (ADULT)

Altered Calling Criteria

ALL OBSERVATIONS MUST BE GRAPHED

COMPLETE ALL DETAILS OR AFFIX PATIENT LABEL HERE

DATE		04/04/2017				
TIME		10 00	12 00	14 00	16 00	18 00
PAIN SCORE Assess pain both at rest and with relevant movement. Document "R" for rest and "M" for movement						
If PCA in use, document pain scores on PCA chart only	Severe pain					
	Moderate pain			M		
					M	
			M	R		
	Mild pain	M	R		R	M
No pain		R			R	

Pain assessment:
R = Rest, M = movement
(If a PCA chart is in use – record pain score on the PCA chart)

Yellow Zone Response:
Have pain scores previously been in the White Zone (0 to 6)?
Have additional prescribed analgesics been administered?
Has the patient been reviewed by the Acute Pain Service or equivalent Medical officer?
The assessing nurse may need to consult with the NURSE IN CHARGE to decide whether a CLINICAL REVIEW (or other CERS – Clinical Emergency Response) call should be made

Dysphoric adverse effects present	Yes					
	No	●●●●●				
INFUSION RATE mg or mL per hour (circle one)		6 mg	6 mg	8 mg	8 mg	8 mg
Cumulative dose mg or mL (circle one)			12 mg	24 mg	40 mg	56 mg
Two initials for change of ketamine program		/	/	SH KL	/	/
Ketamine program checked (initial) once per shift, on patient transfer and on change of bag or syringe		SH		SH		NH
Subcutaneous or IV cannula site secure, no signs of inflammation (check 8 hourly)		SH				NH

- Dysphoric adverse effects assessment
- Infusion rate
- Cumulative dose
- Ketamine pump program checked
- Subcut or IV site checked
- Comments
- Initial

COMMENTS

The standardisation of this chart promotes best practice in prescribing, pain assessment and management of adverse effects in those patients receiving a ketamine infusion.

Comments or questions can be directed to your implementation officer or the project leaders
Emily Edmonds or Jenni Johnson.

(contact details introduction slide)

The feedback register can be located on the ACI website:

<http://www.aci.health.nsw.gov.au/networks/pain-management/acute-pain-forms>

