	Queensland Government
CONT	Government

NIKI T34™, T34™ and BodyGuard™ T **Subcutaneous Medication Infusion Chart**

(Affix	identification label here)

URN:

Family name:

Given name(s):

NOT A VALID PRESCRIPTION UNLESS **IDENTIFIERS PRESENT**

Address:

Date of birth: \square M \square F \square

NIKI T34™, T34™ and E	BodyGuard™ T Syringe Pump Pro	blem Solving
Troubleshooting alerts and	alarms	
Screen information	Result/cause	Possible actions
Program nearly complete	Alert: Program is about to end/syringe is almost empty.	Prepare to change syringe or discontinue pump use.
Low battery	Alert: Battery is almost depleted.	Prepare to change battery.
Pump paused too long	Alarm: The syringe pump has been stopped/paused for more than 2 minutes without any key presses.	Either press key to resume the infusion, press to continue pause for another two minutes or turn the power off.
Syringe empty, remove syringe	Alarm: Current infusion program has completed/syringe is empty.	Prepare to change the syringe or discontinue pump use.
End battery	Alarm: Battery will fail imminently.	Change battery.
Syringe displaced Check syringe	Alarm: One or more of the syringe detection sensors is not detecting.	Check the syringe and re-seat as necessary. Check screen messages for assistance.
Occlusion/empty syringe Check line	Alarm: Clamped set, occluded or kinked. Actuator has reached the minimum travel position.	Release the clamp, flush/replace the access device or clear the occlusion.
System error (high priority alarm) Press and hold !! key for details. If problem persists send pump for service.	Alarm: An internal system error has occurred.	If error recurs: Take pump out of use. Press key to obtain error message. Record error code and summary of fault and return to designated service centre.
Error. Startup motmov fail. If problem persists send pump for service.		
Date and time Incorrect date/time. Press ▶ key to restore	Alarm: The internal backup battery has been depleted and date/time values have been reset.	Press the key , then enter current date and time.

Technical problem/error and failure identification

- The syringe pump alarms if an internal system fault has been detected and the unit will be inoperative.
- The user may be prompted to power off and restart, which may rectify the error.
- If the problem cannot be rectified: power off and remove from patient use.
- Refer to the Technical Service Manual for full details of all technical alarms
- Follow local policy and/or contact your authorised Medical Engineering Department for advice.

The Event Log within the NIKI T34™, T34™ and BodyGuard™ T syringe pump will record the error/alarm event.

Subcutaneous Cannula Sites

- If the Occlusion alarm continues after problem solving actions have occurred, remove and replace the subcutaneous cannula as it may be kinked under the skin.
- If frequent (i.e. every 24 to 48 hours) re-siting of the subcutaneous cannula is required due to site reactions, consider:
- » Changing to a less irritant medicine¹.
- » Adding dexamethasone 1mg to the infusion as an anti-inflammatory agent1.
- » Using NIKI T34™, T34™ or BodyGuard™ T syringe pumps to separate medicines and increase infusion dilution.

Therapeutic Guidelines Limited. Palliative Care: Subcutaneous drug administration in palliative care (Appendix 10.1). [Internet]. West Melbourne (Victoria): Therapeutic Guidelines Limited; July 2016 [eTG August 2020 edition]. Available from: Palliative Care Subcutaneous Drug Administration in Palliative Care (Appendix 10.1)

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Queensland

NIKI T34[™], T34[™] and BodyGuard[™] T Subcutaneous Medication Infusion Chart

Facility:

(Affix identification label here) URN: NOT A VALID Family name:

PRESCRIPTION UNLESS **IDENTIFIERS PRESENT**

NIKI T34™

, T34™ AND BODYGUARD™

T SUBCUTANEOUS MEDICATION INFUSION CHART

Address:

Given name(s):

Date of birth: M F Sex:

First Prescriber to Print Patient Name and Check Label Correct:

This form is suitable for use in Hospitals and Residential Aged Care Facilities.

Practice Tips for NIKI T34™, T34™ and BodyGuard™ T Syringe Pump

Remember the NIKI T34[™], T34[™] and BodyGuard[™] T syringe pump is pre-programed to run over 24 hours for palliative care patients.

To standardise practice, it is recommended to:

- Use 30mL Luer lock syringes. Luer lock syringes prevent risk of disconnection. A 30mL syringe is the largest size syringe that can be contained within the NIKI T34™, T34™ and BodyGuard™ T syringe pump safety Lockbox.
- Use sodium chloride 0.9% as the diluent of choice, unless incompatible with medicine to be infused.
- Draw up medicine(s) plus sodium chloride 0.9% to a combined volume of 20mL to dilute the infusion as much as practical.
- Complete the documentation record sheet on a regular basis, as required by policy and procedures of the local service provider.
- Contact the prescriber for advice if more than three doses of breakthrough medicine are required in a day for a particular symptom.
- Change the extension set every 72 hours or whenever medicine orders are changed.

Battery Guide Niki T34™ Any 9V alkaline disposable battery. Battery life up to 72 hours.

T34™ and BodyGuard™ T

9V alkaline disposable battery with IEC code 6LR61 (code will be written on the battery casing).

Brands that meet this requirement are:

- WINC 9V premium alkaline battery
- Battery World ultra alkaline battery
- Battery life approx: 24 hours for T34™ and 50 hours for BodyGuard™ T

Commencing an Infusion

- Prime the extension set using the PURGE function on the NIKI T34™, T34™ and BodyGuard™ T syringe pump before connecting it to the patient.
- Purging can use up to 2mL of the infusion volume, hence the duration of the infusion can be two to two and a half hours shorter for that day.
- Once the infusion has commenced, remember to press and hold the information button to lock the NIKI T34[™], T34[™] and BodyGuard[™] T syringe pump keyboard.

Terminal Care Prescribing

- For information concerning terminal phase medicine doses and opioid conversions download the NPS MedicineWise PalliMEDS app.
- When using a NIKI T34[™], T34[™] and BodyGuard[™] T syringe pump for terminal phase symptom management, it is best practice to also prescribe anticipatory medicines for pain, agitation, nausea and respiratory tract secretions.
- When commencing patients on a NIKI T34™, T34™ or BodyGuard™ T syringe pump any existing opioid therapy should be reassessed. In particular, consider whether opioid patches should be continued.
- Opioid breakthrough dosing is usually 10% of the total (24 hour) background opioid dose.

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Queensland	(Affix identification label here)				
Government	URN:				
NIKI T34™, T34™ and BodyGuard™ T	Family name:	NOT A VALID			
Subcutaneous Medication Infusion Chart	Given name(s):	PRESCRIPTION UNLESS IDENTIFIERS PRESENT			
A44-ch ABB Odislass NKDA	Address:				
Attach ADR Sticker Unknown	Date of birth:	Sex: M F			
(see Medication Chart for details)	First Prescriber to Print Patient Name and Check Label Correct:				
Sign Date					
Opioid patch to remain? Yes No Not appl	icable				

Prescr	ription	Should be reviewed daily and is	valio	d for up to seven (7) days		
Date	Time (24hr)	Medication		Dose to be delivered in	Preso	Pharmaceutica	
		(print generic name)		24 hours	Print your name	Signature	review (sign and date)
			Ф				
			chloride				
			sodium				
			s %6				
			0				
			with				
			Dilute				

Nursing Calculation and Administration Record									
Date									
Time									
Medication/Ampoule conc.	Volume								
1	mL								
1	mL								
1	mL								
1	mL								
1	mL								
1	mL								
1	mL								
Sodium chloride 0.9%	mL								
Total volume									
Prepared by									
Checked by									
Rate (mL/hr)	mL/hr								
Total volume at commencement of infusion (after priming)	mL								
Date site changed									
Volume discarded at end of infusion	mL								
Discarded by/ Witnessed by									
Pharmacist									

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Given name(s):

(Affix identification label here)

				Date of bi	rth:		Sex:	M F	l
Site Check and D	evice (Check F	Record	Monitoring of	f the infusion r	nust be docur	mented ev	ery four (4) hours	
AIN SCORE Ask the patient to	Date	Time (24hr)	Pain score	Site check	Device check	Volume left (mL)	Sign	Comments	
rate the pain by a numerical scale		04:00	(0–10)		(√) □Y □N	` ′			
0–10 (0 = No pain,		08:00							
10 = Worst pain					UY UN				_
imaginable) Document the score		12:00			Y N				_
-		16:00			Y N				
Check infusion site		20:00			Y N				
Record the results		24:00			Y N				_
(e.g. B1 = Mild		04:00			Y N				
swelling)		08:00			UY UN				
Redness		12:00			∐Y ∐N				
Swelling		16:00			□Y □N				
Tenderness/ hardness		20:00			□Y □N				
Leakage		24:00			Y N				
Urticaria		04:00			□Y □N				
Haematoma		08:00			□Y □N				
None		12:00			\square Y \square N				
Mild Moderate		16:00			\square Y \square N				
Severe		20:00			□Y □N				
Very severe		24:00			\square Y \square N				
EVICE CHECK		04:00			\square Y \square N				
check the device,		08:00			□Y □N				
olution and line using ne check list below.		12:00			□Y □N				
Required volume		16:00			\square Y \square N				
(mL) has been		20:00			\square Y \square N				
delivered		24:00			$\square_{Y} \square_{N}$				
Connections are secure		04:00			□Y □N				_
Solution is clear		08:00			□Y □N				
Lines are kink free		12:00			□Y □N				
Battery life adequate (refer to relevant		16:00			□Y □N				
product information)		20:00			Y N				
Volume remaining is sufficient to maintain		24:00			Y N				
continuous infusion		04:00			□Y □N				_
as prescribed		08:00							
Record results:					Y N				
Y (Yes) if all checks positive		12:00			Y N				
N (No) if any check		16:00			□Y □N				_
negative		20:00			Y N				
If N, document problem and action in		24:00			Y N				_
progress notes		04:00			Y N				
		08:00			Y N				
		12:00			Y N				
		16:00			□Y □N				
		20:00			□Y □N				
		24:00			\square Y \square N				

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