SESLHD POLICY COVER SHEET



NAME OF DOCUMENT	Domperidone for treatment of low breastmilk supply		
TYPE OF DOCUMENT	Policy		
DOCUMENT NUMBER	SESLHDPD/287		
DATE OF PUBLICATION	November 2021		
RISK RATING	Low		
LEVEL OF EVIDENCE	National Safety and Quality Health Standard Standard 1 – Clinical Governance Standard 2 – Partnering with Consumers Standard 4 – Medication Safety Standard 6 – Communicating for Safety		
REVIEW DATE	November 2026		
FORMER REFERENCE(S)	SESLHDPD/287		
EXECUTIVE SPONSOR or EXECUTIVE CLINICAL SPONSOR	Clinical Stream Director, Women's and Children's Health		
AUTHOR	SESLHD Lactation Group Lily Byun Senior Pharmacist RHW		
POSITION RESPONSIBLE FOR THE DOCUMENT	CMC Women's& Children's Clinical Stream Alison Brown <u>Alison.Brown3@health.nsw.gov.au</u>		
FUNCTIONAL GROUP(S)	Women's and Babies Health		
KEY TERMS	Domperidone, lactation, breast milk		
SUMMARY	This policy outlines the management of low breast milk supply and the role of domperidone.		

COMPLIANCE WITH THIS DOCUMENT IS MANDATORY This Policy is intellectual property of South Eastern Sydney Local Health District. Policy content cannot be duplicated.



SESLHDPD/287

1. POLICY STATEMENT

This policy outlines the management of low breastmilk supply and the role of domperidone.

2. AIMS

- To help prevent early cessation of breast feeding due to low milk supply
- To ensure domperidone is prescribed appropriately and in conjunction with nonpharmacological therapies

3. TARGET AUDIENCE

- Medical staff
- Midwifery and nursing staff

4. **RESPONSIBILITIES**

Midwives, nurses and doctors caring for women with low breastmilk supply should follow this policy.

5. DEFINITION

Low milk supply is the one of the most common reasons given for early weaning, therefore it is imperative the condition is diagnosed accurately and if confirmed, managed appropriately. Undersupply may be real, or perceived. Mothers may perceive their infant's need for frequent feeding and comfort as a problem with milk supply. Awareness of normal feeding patterns and growth and the developmental stages of infants can help mothers to be more reassured about their own infant's feeding behaviour.

6. **DOCUMENTATION –** Procedure/Forms

6.1 Procedure

- Ensure a low milk supply exists (perceived vs actual supply) and seek input from lactation services
- Take a full history of mother, baby and birth. An adequate milk supply is dependent on sufficient glandular tissue, intact nerve pathways and ducts, adequate hormones, hormone receptors and adequate frequent, effective milk removal and stimulation
- Ensure non-pharmacological approaches have been trialled such as:
 - Correct positioning and attachment (whilst observing an entire feed), and manage any nipple trauma
 - Increase the number of breastfeeds: wake the infant more often and/or offer the breast for comfort instead of using a dummy/pacifier
 - Massaging breasts prior to feeds and breast compressions during feeds may increase milk transfer
 - $\circ\;$ Educate the mother regarding infant hunger and satiety cues and the signs of effective milk transfer



SESLHDPD/287

 Decrease non-medically prescribed or unnecessary use of artificial infant formula

- Implement 'switch feeding': change the infant from one breast to the other several times during a feed when swallowing has ceased to keep the infant alert and to increase milk intake
- o Increase skin-to-skin contact
- Additional breast stimulation and drainage through double regular expressing after or between breastfeeds
- Good maternal nutrition, rest, relaxation and domestic support and reduce smoking, caffeine and use of alcohol
- Inform the woman that domperidone will increase milk supply ONLY in conjunction with frequent breast drainage (frequent breastfeeds/expressing at least eight feeds every 24 hours)
- Ensure mother does not have any contraindications to treatment with domperidone:
 - Significant personal or family history of cardiac arrhythmia, underlying cardiac disease or electrolyte disturbances
 - In situations when stimulation of gastric motility may be dangerous
 - Prolactin releasing tumour (prolactinoma)
 - Moderate/severe hepatic impairment
 - Lactose intolerance
- Ensure mother is not taking any other medications that may prolong the QT interval and/or inhibit the metabolism of domperidone:
 - o Ketoconazole
 - o Erythromycin
 - Methadone
 - Citalopram/escitalopram
 - Other CYP3A inhibitors which can prolong the QT interval such as fluconazole, voriconazole, clarithromycin and amiodarone
- Discuss the benefits and risks of domperidone use with mother to ensure she is making an informed decision
- Reassure mother that domperidone is safe in lactation. Very low levels are detectable in milk as the molecule is poorly lipid soluble and highly protein bound in maternal plasma.

Dosing

Domperidone 10mg (one tablet) three times daily. A response to treatment should be evident within 7 days, with maximal effects likely to be achieved after 2 to 4 weeks. There is little evidence to support prolonged treatment. Treatment should not be continued for more than 4 weeks.

Once an adequate breast milk supply is achieved, women may benefit from titrating the dose downwards over 1 to 2 weeks before ceasing, avoiding an abrupt withdrawal of treatment.



SESLHDPD/287

Provide patient with SESLHD Increasing your Supply of Breastmilk leaflet. (Appendix A)

Domperidone use in low breast milk is an off-label indication therefore complete the SESLHD-Exceptional Use of Medicine Consent Form. (Appendix B)

Prescribing

- Inpatient: Prescribe domperidone on the eMEDS
- Outpatient: Provide patient with a private prescription.

Side-effects

- Common dry mouth, headache
- Uncommon urticarial rash, insomnia
- Rare loss of balance, palpitations, swelling of feet, restlessness

6.2 Forms

- eMEDS
- Electronic Medical Records including (eMaternity)

7. REFERENCES

Powers NG, Montgomery A, Academy of Breastfeeding Medicine Protocol Committee: ABM Clinical Protocol #9: Use of galactogogues in initiating or augmenting the rate of maternal milk secretion (first revision January 2011). Breastfeed Med 2011; 6(1):41-49

Broddribb, W Academy of Breastfeeding Medicine Protocol Committee: ABM Clinical Protocol #9: Use of Galactogogues in Initiating or Augmenting Maternal Milk Production,Second Revision 2018 Breastfeed Med Volume 13, Number 5, 2018

Grzeskowlak LE, Amlr L, Pharmacological management of low milk supply with domperidone: separating fact from fiction. Med J Aust 2014; 201(5):257-58

Grzeskowlak LE, Smithers LG, Amlr LH and Grivell RM, Domperidone for increasing breast milk volume in mothers expressing breast milk for their preterm infants: a systematic review and meta-analysis. BJOG 2018;125;1371 1378.

Thanigasalam K, Taylor T, Domperidone and breast milk. Obstet Gynae Magazine 2014; 16(4):40-41

Motilium Product Infomation (2020) Retrieved 2/5/21 from eMIMS (via CIAP) Drugs and Lactation Database (LactMed) [Internet]. Bethesda (MD): National Library of Medicine (US); 2006–. Domperidone. 2021 Apr 19. PMID: 30000430.



SESLHDPD/287

Hale TW. Medications and Mothers' Milk. 2012. Hale Publishing, L.P. 15th Edition. Texas, USA

Da Silva OP, Knoppert DC, Angelini MM, Forret PA. Effect of domperidone on milk production in mothers of premature newborns: a randomized, double-blind, placebo-controlled trial. CMAJ.2001;164:17-21

Petraglia F, De Leo V, Sardelli S, Pieroni ML, D'Antona N, Genazzani AR. Domperidone in defective and insufficient lactation. Eur J Obstet Gynecol Reprod Biol.1985;19:281- 287

Date	Revision No.	Author and Approval			
February 2015	0	Endorsed be SESLHD Clinical Quality Council			
August 2015	0	Drafted by: Mariella De Rosa Senior Pharmacist RHW Claudelle Miles Clinical Midwifery Consultant (CMC), Lactation RHW			
August 2015	0	Endorsed by Executive Sponsor to proceed to Draft for Comment			
November 2015	0	Endorsed by SESLHD D&QUMC			
October 2021	1	Minor review by Katy Hunt CMC Lactation RHW, Lily Byun Senior Pharmacist RHW, Dr Debra Kennedy Director Mothersafe, Alison Brown CMC WCCS & the SESLHD Lactation Specialist Group.			
		Domperidone dose reduced following consultation with Mothersafe.			
		Appendices added.			
		Approved by Manager, Women's and Children's Health Stream (Executive Sponsor position vacant).			
		To be tabled at Quality Use of Medicines Committee.			
November 2021	1.1	Endorsed by SESLHD Quality Use of Medicines Committee.			

8. REVISION & APPROVAL HISTORY

Appendix A – Increasing your Supply of Breastmilk

South Eastern Sydney

Increasing your Supply of Breastmilk

July 2021

Many mothers worry about producing enough breastmilk for their babies and many stop breastfeeding because they feel like they don't have enough milk. If you are concerned that your breastmilk supply is low, it is important to seek advice from a breastfeeding specialist, like your Midwife, Lactation Consultant, Australian Breastfeeding Association counsellor or GP.

You will know your baby is getting enough milk if:

- They have at least eight to twelve breastfeeds within 24 hours.
- They have six to eight pale coloured, wet cloth nappies or five to six pale, odourless heavily wet, disposable nappies over a 24 hour period after the first few days.
- They are contented after most feeds.
- They have good skin and muscle tone.
- They show signs of growth or weight gain, after losing 5-10% of birth weight in the first week and they are then back to birth weight by 2-3 weeks.

Bowel movements vary greatly in breastfed babies but should be at least two soft yellow stools every 24 hours for the first 6 weeks. Infrequent bowel patterns in older babies (after 6 weeks) is not a sign of constipation. Breastmilk is so good there is nothing to waste.

Things to try if you are worried about your breastmilk supply:

- Check that your baby is positioned and attached correctly. Any nipple damage or distortion can mean baby is not attached properly and will thus be receiving less milk.
- Increase breast stimulation by increasing how often you feed or the number of times you express, including night time.
- Ask for a breastfeeding specialist to observe a whole feed. They may be able to make some suggestions on how you can feed your baby more effectively.
- Feed from one breast then offer the second breast. Offer both breasts a second time.
- Squeeze your breast for ten seconds while baby is feeding.
- Offer a 'top-up' breastfeed if your baby is unsettled.
- Offer another breastfeed for comfort, rather than using a dummy.
- Encourage skin-to-skin contact.
- Avoid giving your baby other fluids or food unless it is necessary for their health.
- Try to rest, drink adequate fluids and have a well-balanced diet.

Trim No. T15/29360

- Limit caffeine (tea, coffee, cola and chocolate), nicotine and alcohol. Too much can decrease your milk supply.
- Accept practical help at home.

Date: November 2021

Page 5 of 7







SESLHDPD/287

- Surround yourself with supportive people.
- Use of medication to increase supply would only be suggested if other means have been unsuccessful. Medication will have the best chance of working if you also continue increased breast stimulation and removal of milk.

Domperidone (motilium) to increase breastmilk supply

Domperidone is normally used to treat nausea and vomiting, but it can also increase production of the milk producing hormone prolactin. It may take a week before you notice an increase in your breast milk supply.

It is important to continue frequent breastfeeds that is a minimum of 8 every 24 hours, and/or expressing to help your breasts make more milk whilst taking domperidone.

Dosage

Take 1 tablet (10 mg), three times a day, e.g. 6 am, 2 pm, 10 pm. You should see a response within 7 days but the full effect may take 2-4 weeks.

Once a good milk supply is achieved, begin decreasing the dose over 1-2 weeks before stopping the medicine all together. There is little evidence to support treatment with domperidone for more than one month but seek advice from a Lactation Consultant or your breastfeeding specialist.

Possible effects on mother

Tell your doctor if you have any underlying medical conditions or if you are on other medications. A small number of mothers may complain of a dry mouth, skin rash, headache, thirst or drowsiness. If side effects are severe stop the medication and seek medical advice.

Possible effects on baby

There is no record of harmful side effects for babies. However, a small amount of the domperidone will pass through to the breastmilk.

Other Options

Sometimes herbal/naturopathic preparations may be suggested. There is little researched information available on dosage, effectiveness and safety for either mother or baby.

Resources

- Your Midwife, Child and Family Health Nurse, or Lactation Consultant
- Australian Breastfeeding Association <u>www.breastfeeding.asn.au</u> Helpline: 1800 686 268.
- Mother Safe: Medications in Pregnancy & Lactation Service) Ph: 02 9382 6539 or 1800 647 848 if outside the Sydney Metropolitan area or visit <u>www.mothersafe.org.au</u>
- Mothersafe 9382 6539
- If you need an interpreter, call Translating and Interpreting Service (TIS) on 131 450

Endorsed November 2021. Reviewed by consumers in development stage July 2021. Should you wish to discuss any aspect of this information please send an email RHWfeedback@health.nsw.gov.au

Royal Hospital for Women Barker Street, Randwick, NSW 2031 Telephone : 02 9382 6111 www.sesIhd.health.nsw.gov.au/rhw/



SESLHD POLICY



Domperidone for treatment of low breastmilk supply

SESLHDPD/287

Appendix B – Consent for Exceptional Medicine Use

	stillit. Health	FAMILY NAME		MRN				
	NEW AND ADDRESS OF ADD	GIVEN NAME						
	Facility:	D.O.B/ / M	.0.					
		ADORESS						
	CONSENT FOR EXCEPTIONAL							
	CONSENT FOR EXCEPTIONAL USE OF MEDICINE	LOCATION / WARD						
		COMPLETE ALL DETAILS OF	RAFFIXF	PATIENT LAB	EL HERE			
020025	use of a registered medication in an individual patie	ceptional use of medicine includes medication used under the Special Access Scheme (SAS) and some off-label of a registered medication in an individual patient. See SESIH Area Drug Committee Decision Algorithm for aluation of Medicines for Individual Patient Use to confirm justification for exceptional use.						
SEIO	Advice to patient/s carers: This drug is not registered in Australia for use in the cor the Therapeutic Goods Administration of the Australian for exceptional use if an individual patient has a serious unsuccessful or is inappropriate. There may be unknow treating doctor before the commencement of treatment. discuss this with you. In addition, should this treatment significant information has become available.	Department of Health and Ageing, a underlying disease or condition an n side effects. You should discuss t If you have not done this, please a be ongoing, you should ask the trea	and it ma d standa he know sk the tre	ay only be con and therapy have n side effects eating doctor	sidered s been with the now to			
	Written informed consent is required prior to treatment	with this drug.		-				
)	Drug name and form:							
	The condition requiring treatment:							
SNI.	Alternative therapies that may be considered:							
SINDING MARGIN - NO WRITING	Potential risks associated with this treatment:							
NIC I								
AR								
BINDING MARGIN	Expected benefits of treatment:							
BIN								
_	Details of additional written material provided:							
	Details of additional written material provided.							
	STATEMENT OF CONSENT BY PATIENT I have read the above information and statement of accept this liability. I acknowledge that the nature, r explained to my satisfaction. Before signing this do relating to any possible harm I might suffer as a res	eason for use, and possible risks cument I have been given the op	s of the t portunit	treatment ha ty to ask que	ve been stions			
	Signature of patient/carer:			Date:	II			
	By patient, if over 16 years. Otherwise please state relationsh if adult patient unable to give consent, by guardian/spouse/de if patient between 14-16 years, patient plus parent to sign. If u	facto/caregiver/Guardianship Board.	_					
	I, Prof/Dr patient the nature, purpose and risks of the drug tre	(Dr name prin atment to be employed.	nted), ha	we fully exp	ained to the			
	Signature of authorised prescriber: Admitting Medical Officer (Only for Category A SAS may be AMO's Reg	(strar)		Date:	II			
	Both signatures witnessed by:							
	Signature of witness:			Date:	II			
011216	Original to stay in patient t	ile Copy to pharmacy with drug	order					
60168								
8	NC	WRITING			Page 1 of			