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Opioid Dose Equivalence

In order to calculate an oral Morphine Equivalent Daily Dose (oMEDD), multiply the current daily opioid dose by the conversion factor in column 3.

Calculation example for a patient taking oxycodone 40mg/day:

- oMEDD of oxycodone 40 mg/day = 40 x 1.5 = morphine 60 mg/day = oMEDD 60 mg
- Given that calculated oMEDD is ≥ 40 mg, recommend take home naloxone (THN) and provide education

CURRENT OPIOID	CONVERSI	ON FACTOR	PROPRIETARY NAMES	
ORAL (SWALLOWED) PREPARATIONS				
Note: Modified release formulations are marked MR				
Morphine	mg/day	1	Anamorph, Kapanol (MR), MS Contin (MR), MS Mono (MR), Ordine, Sevredol	
Oxycodone	mg/day	1.5	Endone, OxyContin (MR), OxyNorm, Targin (MR)	
Hydromorphone	mg/day	5	Dilaudid, Jurnista (MR)	
Codeine	mg/day	0.13	Aspalgin, Codalgin, Panadeine, Panadeine Forte, Mersyndol, Nurofen Plus, others	
Dextropropoxyphene	mg/day	0.1	Di-Gesic, Doloxene	
Tramadol	mg/day	0.2	Durotram-XR (MR), Tramal, Tramadol SR (MR), Zydol, Zydol SR (MR), others	
Tapentadol	mg/day	0.3	Palexia-SR (MR), Palexia-IR	
SUBLINGUAL PREPARATIONS				
Buprenorphine	mg/day	40	Suboxone, Subutex, Temgesic	
	R	ECTAL PREF	PARATION	
Note: Absorption from rectal administration is highly variable				
Oxycodone	mg/day	1.5	Proladone	
TRANSDERMAL PREPARATIONS				
Buprenorphine	mcg/hr	2	Norspan	
Fentanyl	mcg/hr	3	Denpax, Durogesic, Dutran, Fenpatch, Fentanyl Sandoz	
		NTERAL PRE		
Morphine	mg/day	3	DBL morphine sulphate injection, DBL morphine tartrate injection	
Oxycodone	mg/day	3	OxyNorm FI	
Hydromorphone	mg/day	15	Dilaudid FI, Dilaudid-HP FI	
Codeine	mg/day	0.25	Codeine phosphate injection USP	
Pethidine	mg/day	0.4	Pethidine injection BP	
Fentanyl	mcg/day	0.2	DBL fentanyl injection, Sublimaze	
Sufentanil	mcg/day	2		
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CURRENT oMEDD	Take Home Naloxone (THN) to mitigate harm for outpatients		
Adjust the following recommendations based on individual and contextual risks for opioid harm			
oMEDD 1 - 40mg	Consider THN and patient-education		
oMEDD > 40mg	Recommend THN and patient-education		
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Practical considerations

- 1. This opioid dose equivalence table is intended for comparison of different opioid and opioid formulations in individual patients or in patient cohorts.
- 2. Caution is required if opioid dose equivalence tables are used to guide opioid switching, as the administration of a calculated 'equivalent' dose of the replacement opioid may lead to overdosage.
- 3. It should be noted that there is considerable variability in pharmacokinetics and pharmacodynamics of the different opioids, within and between individual patients. In addition, interactions with non-opioid drugs can strongly influence opioid pharmacokinetics.
- 4. Modified-release formulations can be sub-classified as delayed- or extended- release. Extended release of a drug can be achieved using sustained- or controlled-release delivery systems. When the opioid regimen includes modified- and immediate-release preparations, both should be included in calculation of the oMEDD.
- 5. Methadone, fentanyl lozenges and neuraxial opioids are not included in this table due to their complex and variable pharmacokinetics.
- 6. The conversion factors listed are derived from pooled data in the peer-reviewed literature and pharmaceutical company product information.
- 7. Opioid Safety and TakeHome Naloxone (THN)
- 7.1. The oMEDD dose at which THN should be recommended remains a matter of conjecture and is more nuanced and complex than a simple number.
- 7.2. The literature suggests that there is no safe dose of opioid and that THN should be considered with all/every opioid prescription based on risk assessment. The 2022 Queensland study showed that widely available THN and brief education about recognizing an overdose and how to use THN saved an estimated 3 lives per day from inadvertent over doses.²⁷
- 7.3. Risk factors such as frailty, age <18 or >65 years, weight <50 kg, comorbid medical conditions (respiratory conditions, liver or kidney disease, hypothyroid, body mass), concomitant medications, palliative care, disordered sleep, mental health disorder, and a child <18 in household, need to be considered in determining individual prescription and patient risk.
- 7.4. Although an oMEDD >40 mg and an oMEDD >100 mg are respectively associated with a four-fold and eleven-fold increased risk of a cardiorespiratory event.²⁰ The risk factors listed above and co-prescription of other sedatives, such as benzodiazepines, gabapentanoids and alcohol use elevates the risk and makes it nonlinear.
- 7.5. FPM ANZCA provides the following guidance on take home naloxone (THN):
 - a. Prescribers <u>consider THN</u> and patient-education with **ALL out-patient** opioid prescriptions, adjusted for individual risk
 - Prescribers <u>recommend THN</u> and patient-education with an oMEDD > 40 mg, adjusted for individual risk

For more detail, see <u>PS01(PM) Opioid analgesics in chronic non-cancer pain</u>



Selected references

- Anderson R et al. Accuracy in equianalgesic dosing: conversion dilemmas. J Pain Sym Manage. 2001; 21:397-406
- 2. Behar E, Santos G-M, Wheeler E, Rowe C, Coffin PO. Brief overdose education is sufficient for naloxone distribution to opioid users. Drug and Alcohol Dependence. 2015;148:209-12.
- 3. Bruera E, Pereira J, Watanabe S et al. Opioid rotation in patients with cancer pain. A retrospective comparison of dose ratios between methadone, hydromorphone, and morphine. Cancer 1996;78(4):852-57
- Dale O, Moksnes K, Kaasa S. European Palliative Care Research Collaborative pain guidelines: Opioid switching to improve analgesia or reduce side effects. A systematic review. Palliative Medicine 2011;25:494-503
- 5. Faculty of Pain Medicine. Principles regarding the use of opioid analgesics in patients with chronic non-cancer pain. 2015
- 6. Fine PG, Portenoy RK. Establishing "Best Practices" for Opioid Rotation: Conclusions of an Expert Panel. J Pain Sym Manage 2009;38:418-425
- 7. Franklin GM. Opioids for chronic noncancer pain. Neurology. 2014;83(14):1277-84.
- 8. Galvez R, Schafer M, Hans G, Falke D, Steigerwald I. Tapentadol prolonged release versus strong opioids for severe, chronic low back pain: results of an open-label, phase 3b study. Advances in Therapy 2013; 30(3): 229-259.
- 9. Glare PA, Walsh TD. Dose-ranging study of oxycodone for chronic pain in advanced cancer. J Clin Oncol. 1993;11(5):973-8
- 10. Hagen NA, Babul N. Comparative clinical efficacy and safety of a novel controlled-release oxycodone formulation and controlled-release hydromorphone in the treatment of cancer pain. Cancer. 1997;79(7):1428-37
- 11. Houde R, Wallenstein S, Beaver W. Evaluation of analgesics in patients with cancer pain. Clin Pharm. 1966:1:59–97
- 12. Kalso E, Vainio A. Morphine and oxycodone hydrochloride in the management of cancer pain. Clin Pharmacol Ther. 1990;47(5):639-46
- 13. Kalso E, Vainio A et al. Morphine and oxycodone in the management of cancer pain: plasma levels determined by chemical and radioreceptor assays. Pharmacol Toxicol. 1990;67(4):322-28
- 14. Knotkova H, Fine PG, Portenoy RK. Opioid Rotation: The science and the limitations of the equianalgesic dose table. J Pain Sym Manage 2009;38:426-439
- 15. Lawal, O., et al. Assessment of a systematic framework to determine the equianalgesic conversion ratio between opioids: Determining the conversion ratio between tapentadol and morphine. Pain Medicine (United States) 2018; 19 (4): 869.
- 16. Mahler DL, Forrest WH. Relative analgesic potencies of morphine and hydromorphone in postoperative pain. Anesthesiology 1975;42(5):602–607



- 17. Mercadante S, Caraceni A. Conversion ratios for opioid switching in the treatment of cancer pain: a systematic review. Palliative Medicine 2011;25(5):504-515
- 18. Mercadante S, Porzio G, Aielli F, Adile C, Verna L, Ficorella C, Giarratano A, Casuccio A. Opioid switching from and to tapentadol extended release in cancer patients: conversion ratio with other opioids. Current Medical Research and Opinion 2013; 29:6, 661-666,
- 19. Pereira J, Lawlor P, Vigano A et al. Equianalgesic dose ratios of opioids: a critical review and proposals for long-term dosing. J Pain Sym Manage 2001;22:672-687
- 20. Salom C, Maravilla J, Thomas N, Juckel J, Daly C, Peacock A, et al. Evaluation of the Pharmaceutical Benefits Scheme-subsidised Take Home Naloxone Pilot 2022. https://espace.library.uq.edu.au/view/UQ:7b4d303
- 21. Sittl R, Likar R, Nautrup BP. Equipotent doses of transdermal fentanyl and transdermal buprenorphine in patients with cancer and noncancer pain: results of a retrospective cohort study. Clin Ther. 2005;27(2):225-37
- 22. Skaer TL. Dosing considerations with transdermal formulations of fentanyl and buprenorphine for the treatment of cancer pain. J Pain Research 2014;7:495-503
- 23. The Royal Australasian College of Physicians. Prescription Opioid Policy: Improving management of chronic non-malignant pain and prevention of problems associated with prescription opioid use. Sydney 2008