Sample Opioid Equianalgesic Doses		
Morphine Receptor Agonist	PO/PR (mg)	SQ/IV (mg)
Hydrocodone	10	n/a
Hydromorphone	5	1
Morphine	15	5
Oxycodone	10	n/a

McPherson, ML. Demystifing Opioid Conversion Calculations: A Guide to Effective Dosing. American Society of Hospital Pharmacists. 2009.

Example A: Switching from morphine 30mg PO q6h to hydromorphone PO		
1) Calculate total daily dose of morphine:		
30mg (q6h) * 4 = 120 mg		
2) Using above chart, calculate equivalent total daily dose of hydromorphone :		
15/5 = 120/x; x = 40 mg		
3) For safety, reduce calculated equivalent daily dose of hydromorphone by 50% to accomodate individual variability and/or imprecision of calculation		
40 mg * 0.50 = 20mg		
4) Use supplemental short-acting breakthrough pain medication (e.g., morphine immediate-release 5-10 mg q3h as needed)		
ANSWER: When switching from morphine 30mg PO q6h to hydromorphone PO, start with hydromorphone 20 mg total PO daily, divided as hydromorphone 4-6 mg PO q6h.		
Example B: Switching from morphine 30mg PO q6h to transdermal fentanyl patch q3d 1) Calculate 3 day a nount of morphine prescribed		
30mg (q6h) * 4 * 3 = 360 mg		
2) Estimating transdermal fentany as approximately100-fold more potent than oral morphine, calculate 3 day fentanyl dose requirement 360 mg / 100 = 3.6 mg = 3600 micrograms		
3) Divide total fentanyl dose delivered over 3 days by 72 hours to yield hourly fentanyl dose		
3600 microgra s / 72 hours = 50 micrograms / hour		
3) For safety, reduce calculated equivalent hourly dose of transdermal fentanyl by 50% to accomodate individual variability and/or imprecision of calculation		
50 micrograms / hour * 0.50 = 25 micrograms / hour		
4) Use supplemental short-acting breakthrough pain medication (e.g., morphine immediate-release 5-10 mg q3h as needed)		
ANSWER: When switching from morphine 30mg PO q6h to transdermal fentanyl q3d, start with transdermal fentanyl 25 micrograms / hour, applied q3D.		

CAUTION: As experience with opioid rotation and morphine equivalence has grown, so has appreciation that the above equivalence calculations may not work equally well in each patient (References 1,2 below). Therefore, some advocate an even more cautious approach to opioid transition (Reference 3 below):

1) Reduce the current dose by 10-30% while beginning the new opioid at a dose that would normally be used in an opioid-naïve patient or at the lowest available dose for the formulation.

2) Reduce the original total daily opioid dose by 10–25% per week while increasing the dose of the new daily opioid dose by 10–20% based on clinical need and safety.

3) Allow 3-4 weeks for this transition. Provide sufficient immediate-release opioid during this transition to prevent withdrawal and/or increased pain.

References on opioid equivalence calculations including "MEDD" (Morphine Equivalent Daily Dose)

1) Webster LR, Fine PG. Review and critique of opioid rotation practices and associated risks of toxicity. Pain Medicine 2012; 13: 562-570

2) Fudin F, Cleary JP, Schatman ME. The MEDD myth: the impact of pseudoscience on pain research and prescribing-guideline development. J Pain Research 2016; 9: 153-156

3) Webster LR, Fine PG. Overdose deaths demand a new paradigm for opioid rotation. Pain Medicine 2012; 13:571-574