Opioid Pain Management Clinic for Chronic Non-Cancer Pain





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Disclaimer

I have previously spoken about 'The Opioid Pain Management Clinic at Guy's and St. Thomas' Hospital' at a Grunenthal-sponsored Knowledge Exchange (March 2017) and GP Educational Meeting (July 2017).

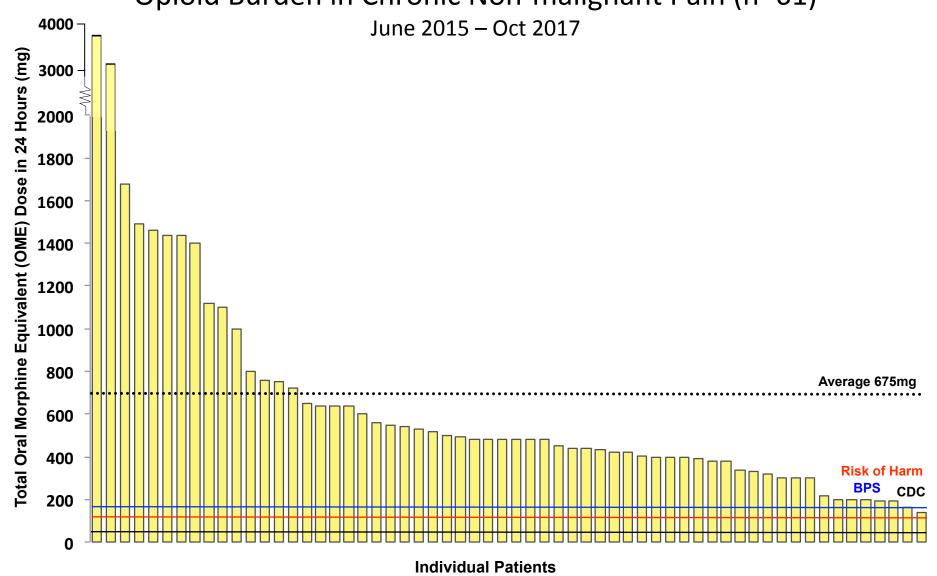
Outline

- Serious adverse effects of high dose chronic opioids
- GSTT 'Opioid Pain Management Clinic'
- 'Opioids Aware' & CDC Guidelines
- Considerations in managing acute pain on a background of chronic opioids

Serious Adverse Effects of Chronic High Dose Opioids

- Fatality due to respiratory depression and accidental drug overdose
- Addiction/ opioid dependence syndrome
- Endocrine dysfunction (sex hormones opioid-induced androgen deficiency)
- Secondary effects of hypogonadism:
 - Bone metabolism osteoporosis and increased fracture risk
 - Anxiety
 - Depression
 - Glucose intolerance
 - Increased cardiac risk
- Suppression of immune function (innate and acquired)
- Link with breast cancer in animal models
- Structural and functional changes within the brain
- Opioid-induced hyperalgesia (OIH)

GSTT Opioid Pain Management Clinic: Opioid Burden in Chronic Non-malignant Pain (n=61)



Rationalise, Optimise & Educate



GSTT Opioid Pain Management Clinic

- Monthly
- Multidisciplinary: Pain Specialist, Nurse, Clinical Psychologist
- Referral criteria >120mg OME (previously >300mg OME)
- Nurse-led second triage
- New & F/Us 45 mins
- Questionnaires: Patient-related outcomes
- Education: BPS, 'Opioids Aware' & DfT 'Drug Driving'
- Treatment agreement (and consent forms)
- Named GP prescriber
- Exit strategy
- Urine & blood testing



Urine & Blood Testing

Urine Immunoassay



- Purpose of testing
- Simultaneous, qualitative detection of 10 drugs and their metabolites
- 98% Accuracy

SYMBOL	TARGET DRUG	CONC.
THC	MARIJUANA	50 ng/ml
COC	COCAINE	300 ng/ml
PCP	PHENCYCLIDINE	25 ng/ml
MOR	OPIATES	2000 ng/ml
MET	METHAMPHETAMINE	1000 ng/ml
MTD	METHADONE	300 ng/ml
AMP	AMPHETAMINE	1000 ng/ml
BAR	BARBITURATES	300 ng/ml
BZO	BENZODIAZEPINES	300 ng/ml
TCA	TRICYCLIC ANTIDEPRESSANTS	30 ng/ml

Formulation	Immediate release vs sustained; liquid to tablets	
Dosing interval	Scheduled rather than PRN to avoid toxicity-withdrawal; increase dosing interval to address frequency	
Rate of wean	Slow; 10% of total daily dose every 1-2 weeks; let patient choose which dose is decreased	
Rebound pain	Adjuvants - Gabapentinoids are opioid-sparing (caution with polypharmacy); or non-opioid strategies/ psychological support	
Withdrawal	Education; taper more slowly; Lofexidine/ Clonidine	
Opioid rotation	Weaning strategy – reduce OME by 25-30% to account for incomplete cross-tolerance	
Frequency of pharmacy dispensing	weekly, alternate day or daily; named GP prescriber	
End point of wean	Ideally <50mg OME; <120mg for PMP or SCS (motivation to wean)	
Support	Increase frequency of visits; Clinical Pyschology/ Nurse	

Opioids Aware

www.fpm.ac.uk/faculty-of-pain-medicine/opioids-aware

- In Jan 2016, FPM/RCoA launched 'Opioids Aware'
- Evidence-based online prescribing resource to support healthcare professionals & patients
- Broad support developed in collaboration with medical royal colleges, RPS, BPS, Public Health England, NHS England, NICE, CQC
- Links to other sources



Opioids Aware

www.fpm.ac.uk/faculty-of-pain-medicine/opioids-aware

- Doses >120mg OME risk of harm escalates with no added benefit
- Avoid prescribing 'risk multipliers' (BDZ, Pregabalin)
- Identify patients at risk of:
 - Opioid dependence
 - Aberrant drug-related behaviour
- Opioid trials
 - Goal orientated
 - Dose-limited
 - Time-limited

Best Professional Practice

Opioids and the law, writing opioid prescriptions, patient safety, reporting harms, record keeping, prescribing

The Condition, The Patient, The Context

Assessment and challenges of long-term pain, the role of medicines, a stepped approach to opioid prescribing

Clinical Use of Opioids

Opioids for different types of pain, their effectiveness and harms, dependence and addiction

A Structured Approach to Opioid Prescribing

Patient assessment, the opioid trial, long-term prescribing, stopping opioids, equivalents, the addicted patient

Information for Patients

Types of pain, thinking about starting opioid medication and frequently asked questions about taking opioids

Opioids Aware

www.fpm.ac.uk/faculty-of-pain-medicine/opioids-aware

- 1. Opioids are very good analgesics for acute pain and for pain at the end of life but there is little evidence that they are helpful for long term pain.
- 2. A small proportion of people may obtain good pain relief with opioids in the long-term if the dose can be kept low and especially if their use is intermittent (however it is difficult to identify these people at the point of opioid initiation).
- 3. The risk of harm increases substantially at doses above an oral morphine equivalent of 120mg/day, but there is no increased benefit.
- 4. If a patient is using opioids but is still in pain, the opioids are not effective and should be discontinued, even if no other treatment is available.
- Chronic pain is very complex and if patients have refractory and disabling symptoms, particularly if they are on high opioid doses, a very detailed assessment of the many emotional influences on their pain experience is essential.

CDC Guidelines (March 2016)

- For Primary Care Clinicians
- 12 Recommendations
- Caution when prescribing opioids at any dose
- Prescribe lowest effective dose to reduce risks of opioid use disorder and overdose
- Reassess individual risk/benefit when prescribing doses of >50mg; avoid prescribing at >90mg
- Overdose risk doubles at 50-99mg OME/day; increases up to x9 at 100mg+ OME/day

CDC RECOMMENDATIONS

DETERMINING WHEN TO INITIATE OR CONTINUE OPIOIDS FOR CHRONIC PAIN

OPIOIDS ARE NOT FIRST-LINE THERAPY Nonpharmacologic therapy and no are preferred for chronic pain. Cli therapy only if expected benefits f anticipated to outweigh risks to th

should be combined with nonpha pharmacologic therapy, as approp-

ESTABLISH GOALS FOR PAIN Before starting opioid therapy for ch establish treatment goals with all cafor pain and function, and should or will be discontinued if benefits do n should continue opioid therapy only improvement in pain and function ti

DISCUSS RISKS AND BENEFIT Before starting and periodically du realistic benefits of opioid therapy

OPIOID SELECTION, DOSAG

USE IMMEDIATE-RELEASE OF When starting opioid therapy for di prescribe immediate-release opio) long-acting (ER/LA) opioids.

USE THE LOWEST EFFECTIVE When opioids are started, clinicians effective desage. Clinicians should opioids at any dosage, should care individual benefits and risks when a50 morphine milligram equivalent increasing dosage to >90 MME/day titrate desage to x90 MME/day.

PRESCRIBE SHORT DURATION Long-term opioid use often begins When opioids are used for acute c the lowest effective dose of immed prescribe no greater quantity than of pain severe enough to require of often be sufficient; more than seve

LEARN MORE

EVALUATE BENEFITS AND HARMS FREQUENTLY

Clinicians should evaluate benefits and harms with patients within 1 to 4 weeks of starting opioid therapy for chronic pain or of dose escalation. Clinicians should evaluate benefits and herms of continued therapy with patients every 3 months or more frequently. If benefits do not outweigh harms of continued opioid therapy, clinicians should optimize other therapies and work with patients to taper opioids to lower dosages or to taper and discontinue opioids.

ASSESSING RISK AND ADDRESSING HARMS

USE STRATEGIES TO MITIGATE RISK

Before starting and periodically during continuation of opioid. therapy, clinicians, should evaluate risk factors for opioid-related harms. Clinicians should incorporate into the management plan strategies to mitigate risk, including considering offering nalesone when factors that increase risk for opioid overdose. such as history of overdose, history of substance use disorder, higher opioid dosages (>50 MME/day), or concurrent benzodiazepine use, are present.

REVIEW POMP DATA

Clinicians should review the patient's history of controlled substance prescriptions using state prescription drug monitoring program (PDMP) data to determine whether the patient is receiving opioid dosages or dangerous combinations that put him or her at high risk for overdose. Clinicians should review PDMP data when starting opioid therapy for chronic pain and periodically during opioid therapy for chronic pain, ranging from every prescription to every 3 months.

USE URINE DRUG TESTING

10 When prescribing opicids for chronic pain, clinicians should use urine drug testing before starting opioid therapy and consider urine drug testing at least annually to assess for prescribed medications as well as other controlled prescription. drugs and illicit drugs.

AVOID CONCURRENT OPIOID AND BENZODIAZEPINE PRESCRIBING

> Clinicians should avoid prescribing opioid pain medication and benzodiazepines concurrently whenever possible.

OFFER TREATMENT FOR OPIGID USE DISORDER

Clinicians should offer or arrange evidence-based treatment Justially medication-assisted treatment with buprenorphine or methadone in combination with behavioral therapies) for patients with opioid use disorder.

Walcoone: a drug that can reverse the effects of opioid overfose

Benzodiazepine: sometimes called "benzo," is a sedative often used to breat arriety, insomnia, and other conditions

POMP: a prescription drug monitoring program is a statewide electronic database that tracks all controlled substance prescriptions



Americans, aged 12 or older, either abused or were dependent on prescription opioids in 2014

Medication-assisted treatment: treatment for opioid use disorder including medications such as buprenorphine or methadone

CDC Guidelines (March 2016)

6. PRESCRIBE SHORT DURATIONS FOR ACUTE PAIN

- Long-term opioid use often begins with treatment of acute pain
- When opioids are used for acute pain, prescribe the lowest effective dose of immediate-release opioids
- Prescribe no greater quantity than needed for the expected duration of pain severe enough to require opioids
- 3 Days or less will often be sufficient;
 >7 days will rarely be needed.

CDC RECOMMENDATIONS

DETERMINING WHEN TO INITIATE OR CONTINUE OPIOIDS FOR CHRONIC PAIN

- 1 Prioritis ARR NOT FIRST-LINE THERAPY heapthreadlegible therapy and nemptical sharmscollagic therapy are preferred for chronic pain. Clinicians should consider opicid therapy only if expended benefits for both pain and function are earlicipeded to obvelegif risks to the obtent. If opicide are used, view should be combined with nonpharmscripp; therapy and nonopicid pharmscripp; therapy, as appropriate.
- ESTABLISH GOALS FOR PAIN AND FUNCTION

 Before starting color therapy for chronic pain, clinicians should establish treatment goals with all patients, including realistic goals for pain and function, and should consider how opcid therapy will be discontinued if benefits do not outweigh risks. Clinicians should continue opid of therapy only if there is clinicially miseningful improvement in pain and function that commonly is skiss to patient safely.

Nonpharmacologic therapies and nonopioid medications include:

- Nonopioid medications such as acetaminophen, ibuprofen, or certain medications that are also used for decression or seizures
- Physical treatments (eg, exercise therapy, weight loss)
- Behavioral treatment (eg. CBT)
- Interventional treatments (eg. injections)
- 3 Before starting and periodically during opioid therapy, clinicians should discuss with patients known risks and resistic benefits of opioid therapy and patient and clinician responsibilities for managing therapy.

OPIDID SELECTION, DOSAGE, DURATION, FOLLOW-UP, AND DISCONTINUATION

- USE IMMEDIATE-RELEASE OPIOIDS WHEN STARTING When starting opioid therapy for chronic pain, clinicians should prescribe immediate-release opioids instead of extended-release/ long-eating (ERVLA) opioids.
- USE THE LOWEST EFFECTIVE OOSE. When opinide are stands, chickers shauld prescribe the lowest effective despective. The chickers shauld prescribe, the lowest effective despective. Chickers should use equation when prescribing opinids at any dosage, should carefully respects evidence of individual centerflower considering interesting dosage to a50 morphite milligram equivalents IMMEDitay, and should avoid increasing dosage to 300 MEDITAY or carefully justify a decision to three dosage to 300 MEDITAY.
- PRESCRIBE SHORT DURATIONS FOR ACUTE PAIN Long-term opioid use often begins with treatment of acute pain. When opioids are used for acute pain, clinicians should prescribe the lowest effective close of limed dire-recess opioids and should prescribe no greater quantity than needed for the expected duration of pain severe enough to require opioids. Three days or lists will often be sufficient, more than seven days will rend be needed.

Immediate-release opioids: faster acting medication with a shorter duration of pain-relieving action

Extended release opioids: slower acting medication with a longer duration of pain-relieving action

Morphine milligram equivalents (MME/May the amount of morphine an opicial dose is equal to when prescribed, often used as a gauge of the abuse and evendose potential of the abuse and evendose potential of the an opticular time.

LEARN MORE | www.cdc.gov/drugoverdose/prescribing/guideline.html

Challenges in Managing Acute Pain in Opioid-Tolerant Patients

Biological

 Chronic stimulation of opioid receptors leads to central sensitisation (OIH/ tolerance)

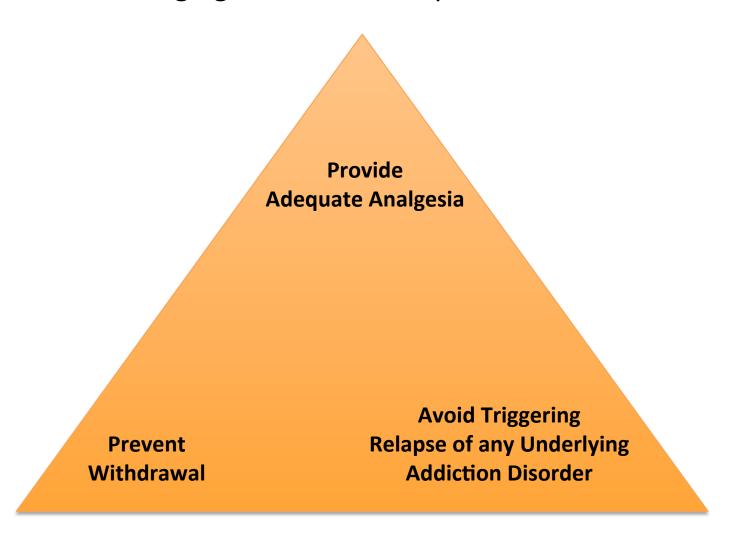
Clinician-related Barriers

- Stigmatisation of all opioid-tolerant individuals as 'addicts' or 'drug seekers'
- Possibility of reports of pain being fabricated to acquire opioids
- Diversion of prescribed opioids
- Fear of causing a drug overdose
- Lack of knowledge about opioid equivalent doses

Patient Concerns

- Fear of withdrawal/ drug cravings (OST)
- Restricted access to analgesia/ fear of pain
- Clinician distrust/ fear of discrimination/ not being taken seriously

Goals in Managing Acute Pain in Opioid-Tolerant Patients



Evaluation of Acute Pain in Opioid-Tolerant Patients

Is this an exacerbation of baseline pain?

- Development of tolerance
- Drug interaction causing decreased effectiveness of an existing regime
- Exacerbation of underlying disease

Is this a new pain?

- OIH (change in quality, pain becomes more diffuse)
- Pain in an area of known disease may reflect a new complication e.g. cord compression
- Pain unrelated to the primary disease process e.g. appendicitis

Managing Acute Pain on a Background of Chronic Opioids: Principles in Prescribing and Treating

- 1. Determine baseline daily opioid requirement prior to the onset of the acute event (OME dose in 24 hours)
- 2. Prescribe adequate doses of opioids to treat this baseline pain continue patient's previous long-acting oral or transdermal opioid
- 3. Add short-acting opioids to cover acute pain (e.g. PCA), monitor side-effects
- 4. Change from parenteral to oral formulation
- 5. If opioid titration is ineffective, consider 'opioid rotation'
- Establish whether other drugs are being misused
- 7. Optimise non-opioid analgesia:
 - Multi-modal & opioid-sparing e.g. Gabapentinoids, NSAIDs
 - Adjuvants: Ketamine, Clonidine
 - Interventional pain injections/ nerve blocks/ neuraxial analgesia
- 8. Multidisciplinary discharge planning/ community support to provide follow-up to manage the process of analgesia reduction

Summary

- Complex and challenging patients: Significant functional impairment, psychopathology and high risk for aberrant drug-related behaviour
- Clinical need to effectively manage in a multidisciplinary setting
- Significant improvement in patient outcomes with weaning
- Tailored approach: Rationalise, optimise and educate
- The problem is not opioids per se but the way in which patients are treated with opioids i.e. without appropriate indication/ precautions and with excessive doses

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