Edinburgh Pain Assessment Tool (EPAT $^{\odot}$) VITAL SIGNS AND PAIN ASSESSMENT CHART – STEP 1

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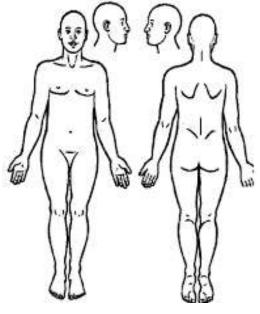
➤ Patients with severe pain

➤ Patients requiring a rapidly increasing opioid dose

➤ Patients with opioid-induced drowsiness

Edinburgh Pain Assessment Tool (EPAT ©) -Step 2

Name:	Ward:	Date/Time
		,



Severity Score	A Most Severe	В	С
Worst Pain in Last 24 hrs (0-10)			
Least Pain in Last 24 hrs (0-10)			

0 = No Pain 10 = Worst Pain Imaginable

➤ Patients with movement-related pain

➤ Pain unrelieved by initial management

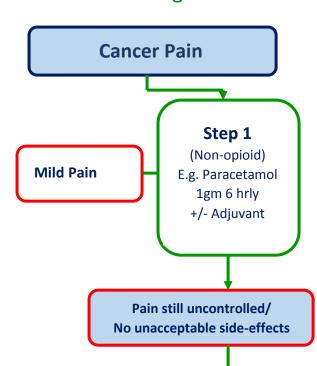
	Does your p Yes □	ain disturb yo No□	our sleep?			
Is your pain? Shooting or Stabbing □ Tingling or Pricking □ >	Pins & Ne	edles□ Ho	ot or burning \Box			
Do any areas of your skin feel numb or strange of Yes □ No □ Detail:						
Does moving or any other activity make your pa	in worse?	Ye	s □ No□			
Does your pain come on suddenly at rest? Consider neuropathic pain. Starting gabapentin/starting and		djuvants	s □ No□			
Does moving or any other activity make your pa	in worse?	Ye.	s □ No□			
Does your pain come on suddenly at rest?		Ye	s □ No□			
Is the patient experiencing movement-related or spontaneous pain? Consider bone pain Use WHO analgesic ladder – See EPAT algorithm. Give PRN analgesia before movement Consider NSAID's / Palliative Radiotherapy / Bisphosphonates						
What makes your pain better?						
Remember: non-pharmaco Consider: Position change / Relaxation / Physiot	_		acupuncture			
Is there anything worrying or concerning you	about vour	pain? Discus	ss with patient.			

Remember: anxiety/depression may co-exist with severe pain.
Consider referral to your Specialist Team for patients
who have persistent pain:

		Edinburgh	Pain Assessment Tool (E	PAT©) – Treatment Revie	ew Chart		
Name:		DOB:		nit/Hospital No:	Ward:		. <u></u>
Date & Time							
Worst Pain Now	(0-10)	(0-10)	(0-10)	(0-10)	(0-10)	(0-10)	Analgesic
Worst Pain Site (specify)							InterventionsBreakthroughAnalgesia
Pain Distress	Is your pain distressing: ☐Yes ☐No Comment:	Is your pain distressing: ☐Yes ☐No Comment: ———————————————————————————————————	Is your pain distressing: ☐Yes ☐No Comment: ———————————————————————————————————	Is your pain distressing: ☐Yes ☐No Comment: ————————————————————————————————————	Is your pain distressing: ☐Yes ☐No Comment: ———————————————————————————————————	Is your pain distressing: ☐Yes ☐No Comment: ————————————————————————————————————	 ↑Dose Regular Analgesia Regular Analgesia
Opioid toxicity Stop & Check Daily!	□Shadows-corner of your eyes □ Drowsiness □Vivid dreams □Hallucinations □Confusion □Jerking/twitching Check-opioid dose may not suit	□ Shadows-corner of your eyes □ Drowsiness □ Vivid dreams □ Hallucinations □ Confusion □ Jerking/twitching Check-opioid dose may not suit	□Shadows-corner of your eyes □ Drowsiness □Vivid dreams □Hallucinations □Confusion □Jerking/twitching Check-opioid dose may not suit	□Shadows-corner of your eyes □ Drowsiness □Vivid dreams □Hallucinations □Confusion □Jerking/twitching Check-opioid dose may not suit	□Shadows-corner of your eyes □ Drowsiness □Vivid dreams □Hallucinations □Confusion □Jerking/twitching Check-opioid dose may not suit	□Shadows-corner of your eyes □ Drowsiness □Vivid dreams □Hallucinations □Confusion □Jerking/twitching Check-opioid dose may not suit	Changed Adjuvant Drug Prescribed Adjuvant Dose
Intervention (specify)	□Analgesia □ Non-Pharmacological □ Other	□Analgesia □ Non-Pharmacological □ Other □	□Analgesia □ Non-Pharmacological □Other	□Analgesia □ Non-Pharmacological □ Other	□Analgesia □ Non-Pharmacological □ Other	□Analgesia □ Non-Pharmacological □ Other	Non- Pharmacological Interventions Position change Physiotherapy Heat
Outcome (Review <u>1 hr</u> after analgesia)	Worst pain (0-10) Is your pain distressing now: ☐Yes ☐No	Worst pain (0-10) Is your pain distressing now: ☐Yes ☐No ————————————————————————————————————	Worst pain (0-10) Is your pain distressing now: □Yes □No	Worst pain (0-10) Is your pain distressing now: □Yes □No	Worst pain (0-10) Is your pain distressing now: □Yes □No	Worst pain (0-10) Is your pain distressing now: □Yes □No ————————————————————————————————————	ColdTENSMassageAcupunctureRelaxation
Next Review (Date & Time)							

Managing Cancer Pain

► Green for go!



(Weak Opioid)

F.g. Codeine

Moderate Pain

E.g. Codeine 30-60mg 6 hrly +/- Non-opioid +/- Adjuvant

Step 2

Pain still uncontrolled/
No unacceptable side-effects

Moderate →
Severe Pain

(WHO Analgesic Ladder)

(Opioid) Oramorph 5mg 4 hrly + PRN

+/- Non-opioid

Step 3

+/- Adjuvant

Titrate Until

Pain Relief

Edinburgh Pain

Assessment Tool EPAT©

Patient's pain is usually:

• Continuous requiring Regular analgesia

If pain is intermittent:

(See 'Breakthrough Cancer Pain' algorithm)

Cancer pain requires careful assessment:

- Cause
- Severity
- Treatment Response

Analgesics:

- Steps 1, 2 & 3 WHO Analgesic Ladder
- Adjuvant Analgesics (some examples

•	•
Steroids	Dexamethasone
NSAIDs	Ibuprofen
Anticonvulsants	Gabapentin
Tricyclic antidepressants	Amitriptyline

Management:

- Start on Appropriate Step
- Consider route any absorption issues?
- Prescribe regular analgesia & PRN
- PRN dose usually <u>1/6th</u> 24hr opioid dose
- > Reassess pain & Titrate Upwards
- Until Pain Controlled
- Or Unacceptable Opioid Side Effects
- Convert to long acting analgesia when Pain control stable e.g. MST Continus
- Prescribe Regular Laxative
 - -- PRN anti-emetic
- Reduce opioid Frail ElderlyRenal/Liver Dysfunction

Step 3

!Monitor for signs of opioid toxicity

Dose Titration – see EPAT© examples

Opioid Toxicity

<u>Signs:</u> Shadows-at corner of eyes/Drowsiness Vivid Dreams/ Hallucinations/ Confusion/Jerking Management:

- Reduce opioid dose
- ➤ Hydrate IV/SC fluids if required
- Consider Adjuvant Therapies
 - Opioid Switch
 - Non-drug Measures
- Prescribe antipsychotic e.g. Haloperidol if confusion/hallucinations/agitation present

!Uncontrolled pain

If patient still have uncontrolled pain +/unacceptable side-effects, discuss with your
Specialist Team

EPAT© Dose Titration Examples

Example 1 – Continuous Pain

Mr A is prescribed Oramorph 5 mg 4 Hourly and Oramorph 5 mg PRN for breakthrough pain – he has required 3 extra doses of Oramorph over the last 24 hours.

He describes his cancer-related pain as 'almost constant' and states the Oramorph is providing only partial pain relief – he has no apparent side-effects on assessment.

<u>Titration</u> – Oramorph 5 mg 4 hourly – 5 mg x 6 = 30 mg + PRN doses 5 mg x 3 = 15 mg

Total <u>24 Hr</u> Dose Requirement = Oramorph <u>45 mg</u>

New Titration Dose: Oramorph 45 mg ÷ 6 = 7.5 mg 4 Hourly & PRN dose 7.5 mg

Example 2 – Breakthrough Cancer Pain (End-of-dose-failure)

Mrs B is receiving Oramorph 4 Hourly and Oramorph 30 mg PRN for breakthrough pain – she is finding her pain is generally well controlled for approximately 3 hours then it returns. She is using 2 PRN doses for breakthrough pain daily with reluctance, as she feels drowsy if a PRN dose is required closely (within an hour) of her regular 4 Hourly dose.

Titration – Oramorph 30 mg 4 hourly – 30 mg x 6 = 180 mg + PRN doses 30 mg x 2 = 60 mg

Total 24 Hr Dose Requirement = Oramorph 240 mg

New Titration Dose: Oramorph 240 mg ÷ 6 = 40 mg 4Hourly & PRN dose 40 mg

!Remember: If end-of-dose-failure is problematic and PRN doses are not being utilised – gently titrate

Oramorph dose (25-30%) and consider switching to long acting preparations

Example 3 - Breakthrough Cancer Pain (Incident Pain)

Mr C is prescribed Oramorph 80 mg 4 Hourly and Oramorph 80 mg PRN for breakthrough pain – he related that when lying down his pain is more controlled and he requires 2 doses of PRN Oramorph over the day. When he attempts to mobilise however his pain becomes very severe and he requires a further 2 or 3 PRN doses daily prior to mobilising with only partial effect.

<u>Titration</u> – Oramorph 80 mg 4 hourly – 80 mg x 6 = 480 mg + PRN doses 80 mg x 2 = 160 mg*

Total 24Hr Dose Requirement = Oramorph 640 mg

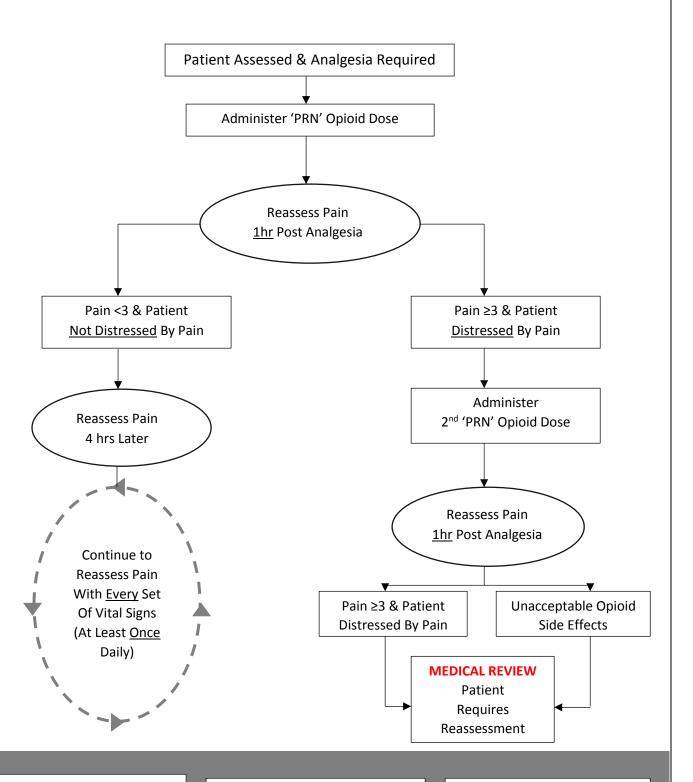
New Titration Dose: Oramorph 640 mg ÷ 6 (rounded to) 100 mg 4 Hourly & PRN dose 100 mg

*If all PRN doses are included in titration where pain is mainly Incident, it is highly likely that patients will experience unacceptable opioid side-effects

→ Initially aim for comfort at rest

→ Then titrate PRN dose & consider other interventions to optimise Breakthrough Pain control

Cancer Pain – Administering 'PRN' Opioids



Opioids - Management

Prescribe:

- Regular & PRN Analgesia
- Regular Laxative
- PRN anti-emetic

Monitor: Signs Opioid Toxicity
Always Reassess Pain!

PRN Opioids - Dose

PRN Opioid Usually <u>1/6th</u>
Total <u>24hr</u> Opioid Dose

Caution: Frail Elderly

Renal Dysfunction

Hepatic Impairment (Dose Reduction May Be Required)

Assessment - Timing

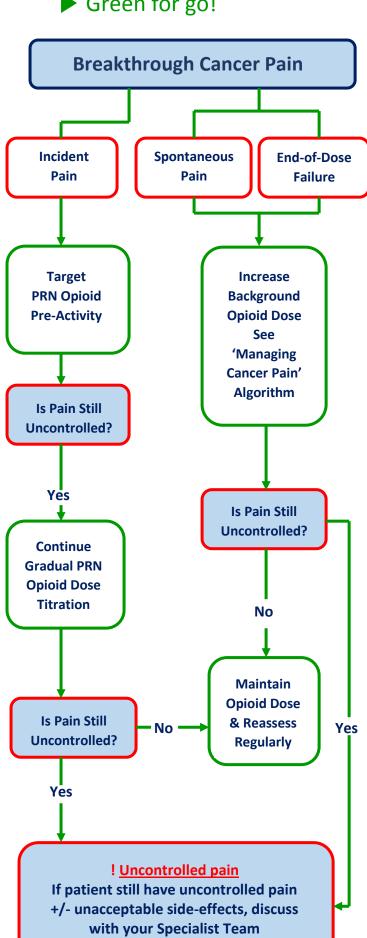
<u>1hr</u> – Oral Opioid Dose 30mins – Parenteral Opioid

More Immediate Review:

- Distressed Patients
- Using Fast Onset Opioids
 e.g. OTFC (Actiq)

Managing Breakthrough Cancer Pain

Green for go!



Edinburgh Pain

Assessment Tool EPAT©

Breakthrough Cancer Pain (BTCP):

A transitory exacerbation of pain occurring on a background of adequately controlled baseline pain (Portenoy et al 2004).

Patient's pain is usually of:

- Sudden Onset
- Severe Intensity
- Short Duration (average 30 mins)

Breakthrough Pain Types:

- Incident: Related to activity e.g. Movement/ Micturition/ Coughing
- Spontaneous: Unexpected/At any time
- End-of-dose-failure: **Insufficient Background Dose**

Management:

> Incident Pain:

<u>Target</u> PRN opioid dose <u>Pre-Activity</u> e.g. Movement-Related Pain

- Oramorph 30 mins before mobilising
- Spontaneous Pain: Prescribe Immediate Release Opioid Increase background Opioid Dose Consider Adjuvant/(dose titration)
- > End-of-dose-failure: Increase Background Opioid Dose

All Breakthrough Cancer Pain:

- Assess most appropriate Immediate Release Opioid (&Dose) for Patient
- Is Specialist Intervention Required?
- Consider Non-opioids & Adjuvants
- Consider Non-drug Measures
 - Heat/ TENS/ Activity Modification
 - ! Stop titrating opioid dose if patient experiencing unacceptable side-effects

Analgesics:

- Steps 1, 2 & 3 WHO Analgesic Ladder & Adjuvant analgesics
- PRN Opioid Dose for BTCP usually <u>1/6th</u> 24hr total opioid dose
- Seek Specialist Advice
 - Pain is of very short duration
 - Side effects prevent ↑ PRN opioid

! Remember Breakthrough Cancer Pain is often associated with opioid toxicity

Edinburgh Pain

Assessment Tool EPAT©

Patients' pain may be:

- Continuous 'throbbing'/'sharp'/'boring'
- Spontaneous unpredictable
- Movement-related initiated on walking/ lying down/ sitting or standing
- Localised or Radiates on movement

! Always assess for any underlying contributing non-malignant bone pain

Cancer Induced Bone Pain

Managing

Green for go!

Cancer Induced Bone Pain



Movement **Related Pain**

Start **Appropriate** Step On 'Managing Cancer Pain' **Algorithm**

See 'Managing Cancer **Breakthrough** Pain' **Algorithm**

Pain still uncontrolled/ No unacceptable side-effects

Increase **Opioid Dose**

Move Up

Step 1→ 2

Step 2→ 3

On Step 3?

Titrate Until

Pain Controlled Maintain Opioid Dose & Reassess Regularly

Immediate Management:

Analgesia

Specialist Options:

- Radiotherapy*
- **Bisphosphonates**
- Chemotherapy
- **Hormonal Therapy**
- **Surgery**
- **Anaesthetic Intervention**
- * Analgesic response may take 6 weeks Monitor for treatment related pain flares
- ! Remember pain may be an early warning of complications from bone metastases:
- **Pathological Fractures**
- Hypercalcaemia
- **Spinal Cord Compression**

Analgesics:

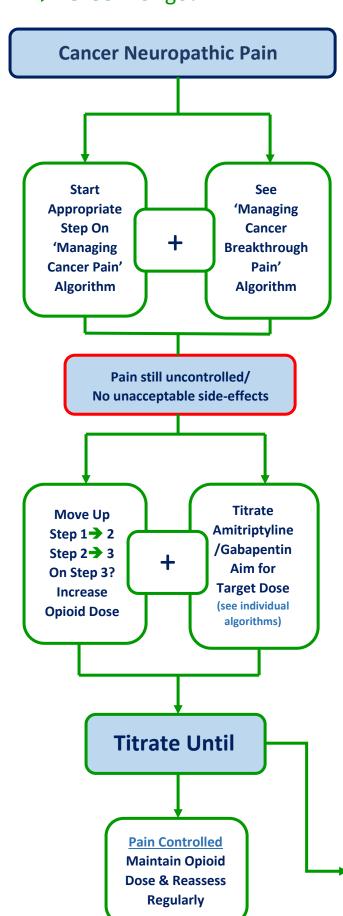
- WHO Analgesic Ladder +/- NSAIDS*
- Target different pain elements
 - **Continuous Pain**
 - **Spontaneous Pain**
 - **Movement-related Pain**
- Aim for comfort at rest
 - ↑Opioid doses are usually required for **Movement-Related & Spontaneous Pain**
- Balance pain relief against side-effects
- Consider Opioid Switch if patient is experiencing unacceptable side-effects
- Consider Non-drug Measures -**Heat/TENS**
 - ! Remember Cancer Induced Bone Pain is often associated with opioid toxicity

! Uncontrolled pain

If patient still have uncontrolled pain +/- unacceptable side-effects, discuss with your Specialist Team

Managing Cancer Neuropathic Pain

Green for go!



Edinburgh Pain

Assessment Tool EPAT©

Neuropathic Pain:

Pain initiated or caused by a primary lesion, or dysfunction in the nervous system
(IASP 1994)

Patients' pain may be:

- Continuous: 'burning'/'aching'/'heavy'
- Intermittent: 'stabbing'/'shooting'/
 'jumping'/'bursting'/'electric shocks'
 - at rest or on movement
- Triggered: initiated or worsened by light tough/ tight clothes/ bed clothes

Patients may experience:

Management:

Neuropathic pain may respond to Tricyclic Antidepressant and Anticonvulsant therapies +/- conventional analgesia.

Amitriptyline vs Gabapentin

Is sleep pattern interrupted by pain?
- Consider Amitriptyline as 1st line

Analgesics:

- Steps 1, 2 & 3 WHO Analgesic Ladder & Adjuvant Analgesics (see above examples)
- Always prescribe regular analgesia
- <u>Balance</u> pain relief against side-effects
 If Patient experiencing opioid side-effects
 - titrate Adjuvant Analgesic only (1st line)
 - consider Opioid Switch (2nd line)
- <u>Consider Non-drug Measures If rubbing</u> area helps pain try TENS
- ! Remember patients with neuropathic pain are often susceptible to opioid toxicity

! Uncontrolled pain

If patient still have uncontrolled pain +/- unacceptable side-effects, discuss with your Specialist Team

Edinburgh Pain Assessment Tool EPAT© **Starting Amitriptyline*** Green for Go! Red for No! ¹For younger patients and inpatients: Start 25 mg nocte Start Amitriptyline 10-25 mg nocte Increase to 50 mg nocte at day 3 (check 1 &2) • Increase to 75 mg nocte at week 2 ! Remember to balance against side-effects ²For frail/elderly/infirm and outpatients: Increase dose as Are side-effects • Start 10 mg nocte acceptable? per 1 & 2 Increase to 25 mg at day 3 • Increase to 50 mg at week 2 • Increase to 75 mg at week 3-4 (as tolerated) ! Seek advice if Amitriptyline is not effective or side-effects prevent dose increases Yes No No ! Drug Interaction: Avoid using amitriptyline in patients taking SSRI's **Amitriptyline Side Effects:** Sleepiness/ dizziness/Delirium/ Dry Mouth/ **Blurred Vision/ Constipation in the elderly** Is dose Has target controlling dose been No If side-effects are intolerable: patient's reached? • Exclude other causes for these symptoms pain? - they may not be due to Amitriptyline Reduce to the last tolerated dose* and/or Yes Yes stop the amitriptyline • Consider discussing with your Specialist Team *Consider a more gradual dose increase Is pain still uncontrolled? e.g. 10 mg increments Maintenance: Maintain Consider Maintain dose when analgesic benefit Dose & Alternative adjuvant achieved Reassess e.g. Gabapentin **Continue to monitor patient tolerance** regularly (see 'Starting Avoid abrupt withdrawal after prolonged use Gabapentin' algorithm) ! Uncontrolled Pain *If in doubt check BNF for drug If patient still has uncontrolled interactions/contraindications pain, discuss with Is pain still your Specialist Team uncontrolled?

Edinburgh Pain Assessment Tool EPAT© **Starting Gabapentin*** Green for Go! Red for No! Dose Increases: [300 mg every 24 hrs] Day 1: 300 mg nocte Day 2: 300 mg bd Start Amitriptyline 10-25 mg nocte Day 3: 300 mg tid (check 1 &2) Target dose: Gabapentin may need to be titrated to 600 mg tid before being effective ! Remember to balance against side-effects Increase dose by 300 Are side-effects ¹Is patient frail/elderly/infirm? mg every 24 hrs acceptable? Start 100 mg nocte • Increase by 100 mg every 24-48hrs if tolerated (see side-effects box) Maintain dose when analgesia reached ²Is there renal dysfunction? Use reduced doses & discuss with ward Yes No No pharmacist ! Seek advice if Gabapentin is not effective or side-effects prevent dose increases **Gabapentin Side Effects:** Sleepiness/ Dizziness/Ataxia/Tremor Has target If side-effects are intolerable: Is dose dose been • Exclude other causes for these symptoms controlling No reached? - they may not be due to Gabapentin patient's (600 mg tid) Reduce to the last tolerated dose* and/or pain? stop the Gabapentin If side-effects settle: Yes Yes • Consider a more gradual dose increase e.g. 100 mg or 300 mg every 48-72 hrs ! Remember if the dose is kept the same mild side-effects may settle over a few days Is pain still uncontrolled? Maintenance: Maintain Maintain dose when analgesic benefit dose & achieved reassess Yes **Continue to monitor patient tolerance** every 24 hrs Avoid abrupt withdrawal after prolonged use ! Uncontrolled Pain If patient still has uncontrolled *If in doubt check BNF for drug pain, discuss with interactions/contraindications your Specialist Team