PTHC and Guildford and Waverley CCG Palliative Care GP Update

Dr Nick Dando, Consultant in Palliative Medicine
Debbie Hindson, Nurse Consultant
Dr Jacqui Phillips, Consultant in Palliative Medicine



Aims

 Update on service development at PTHC with a particular emphasis on new community-focussed models of care

 Link in with developing models of care for patients with non-malignant life-limiting illness in G+W

 Symptom control update including cancer pain management and symptom control at the end of life



Non malignant service development 2012 - 2017

Chronic respiratory disease

Neurodegenerative disease

 Individualised 3 month blocks of care under the Day Hospice

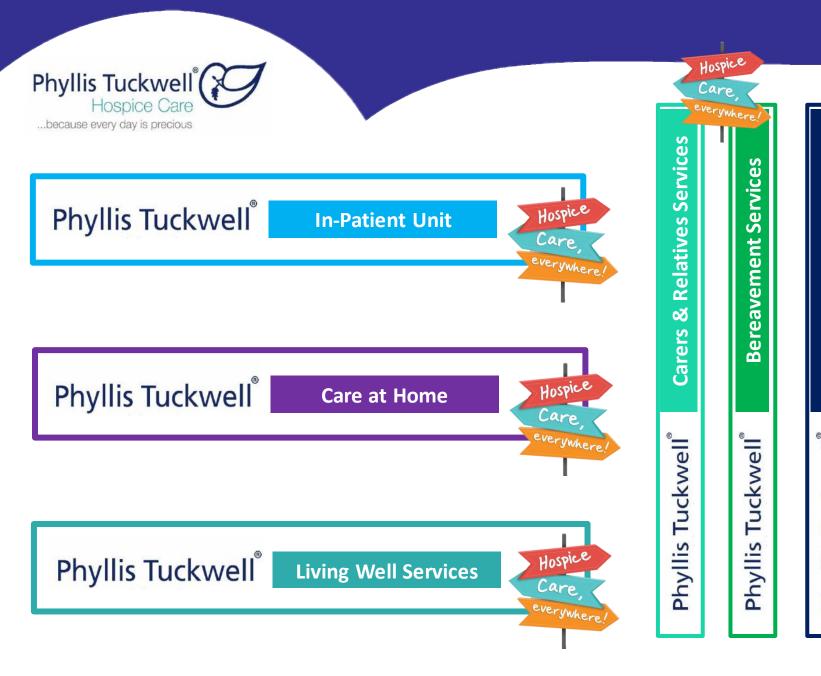
 Appointment of Debbie Hindson (Nurse Consultant) and Dr Jacqui Phillips (Non-malignant medical lead)



The Beacon Centre April 2015







Education & Training

Phyllis Tuckwell

Living Well

Focusing on the 'Rest of Life' we are here to help patients, and those closest to them, to:

- maximise their physical health
- help to cope with changes
- improve wellbeing
- be as independent as possible to get more out of life.



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Symptom control update

Dr Nick Dando



Aims

- Opioid update
- Transdermal opioids
- General prescribing at the end of life
- Prescribing in renal impairment

Management of terminal agitation

Management of secretions



Oral to transdermal Fentanyl conversion

PANG/PCF/BNF

PO Morphine 100mg (50mg BD)

Divide 100

Transdermal fentanyl 1mg in 24 hours

Multiple 1000

Transdermal fentanyl 1000 micrograms in 24 hours

Divide 24

= 41.6 micrograms per hour

Fentanyl patch 37microgram/hr with PRN morphine IR 15mg PO

 This conversion is for patients with stable pain and careful monitoring of pain and side effects should be undertaken when switching to a patch.

Transdermal Fentanyl	PO Morphine	
Micrograms/hr	mg/24 hours	PRN (mg)
12	30 - 59	5
25	60 – 89	10
37	90 - 119	15
50	120 - 149	20
75	180 - 239	30
100	240 - 299	40

Adapted from: Quick prescribing guide: Use of transdermal fentanyl patches (PCF 5) and Opioid Equivalence for Transdermal Patches Conversion Guide (PANG 2016).

Guidelines for Strong Opioids at Phyllis Tuckwell Hospice. 2017



Buprenorphine

- Buprenorphine is a partial mu agonist which only becomes significant at high doses of Buprenorphine (3-5mg/24hrs).
- It is also a kappa receptor antagonist, which as a consequence can increase mu receptor expression on membrane surfaces, and an agonist for the opioid receptor like 1 (ORL-1) receptor.
- The drug is effective in cancer pain, neuropathic pain and other pain presentations and is associated with less analgesic tolerance.
- These clinical benefits have resulted in increased prescription of buprenorphine in some cancer centres, with more patients being discharged into the community with the medication

Oral to transdermal buprenorphine conversion

Buprenorphine conversion (morphine:buprenorphine 100:1)

Morphine sulfate 25 mg PO BD = 50mg PO in 24 hours

Divide by 100

Buprenorphine 0.5mg in 24 hours

Multiple 1000

Buprenorphine 500 micrograms in 24 hours

Divide by 24

Buprenorphine 20.8 micrograms/hour = BuTrans Patch 20 micrograms/hour

Breakthrough dose = 50 divided by 6 = 8.3mg PO morphine PRN

Buprenorphine Equivalents

Transdermal Buprenorphine	PO Morphine	
	mg/24 hours	PRN (mg)
5microgram/hr	12	2
10microgram/hr	24	5
20microgram/hr	48	10
35 microgram/hr	84	15
52.5 microgram/hr	126	20
70 microgram/hr	168	30

Adapted from Quick Practice Guide: Use of transdermal buprenorphine. PCF 5. Guidelines for Strong Opioids at Phyllis Tuckwell Hospice. 2017



General Prescribing Guidelines in EOLC

Paín	Morphine sulfate	2.5-5mg	SC	PRN
Anxiety	Mídazolam	2.5-5mg	sc	PRN
Nausea	Haloperidol	0.5- 1.5mg	SC	PRNTDS
Chest secretions	Hyoscine butylbromide	20mg	SC	PRN

If three or more PRN doses needed in 24 hours, start a syringe driver with equivalent drug doses

Example

Water for	Morphine sulfate 15mg	vía sub cut syringe
Injection	Haloperidol 3mg	dríver over 24 hours
, rgooccore	Hyoscine butylbromide 60mg	011 0 0 0 1 0 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 1 0 0 0 0 1 0 0 0 0 1 0 0 0 0 1 0 0 0 0 0 1 0



Alfentanil

- Synthetic lipophillic opioid
- Alfentanil may be considered for patients with renal impairment or intolerance to Diamorphine/ Morphine/Oxycodone
- Morphine 30mg PO = Alfentanil 1mg
- Please note Alfentanil has a short duration of action (30min). It may therefore require more frequent PRN dosing, with careful monitoring for side effects.



Prescribing in renal impairment

Paín	Oxycodone	1.25 mg	sc	4 hourly PRN
Anxiety	Mídazolam	2.5mg	sc	PRN
Nausea	Haloperidol	0.5- 1.5mg	sc	PRNTDS
Chest secretions	Hyoscine butylbromide	20mg	SC	PRN

If a background syringe driver is needed for pain then use alfentanil

Example

Waterfor	Alfentaníl 500 mícrograms	sub cut over 24 hours
Injection		



Terminal Agitation

- Patients may develop delirium and confusion at the end of life
- Spectrum of behaviour
- If acutely delirious, consider higher doses of haloperidol rather than midazolam as benzodiazepines may exacerbate confusion

Example

 Haloperidol 1.5 - 2.5mg SC PRN for delirium (max 12mg in 24 hours)



Chest secretions

- Consider the causes
- Assess the impact on the patient
- Reassure and support family
- Consider use of antisecretory medication

Respiratory or oro-pharyngeal secretions present in a dying patient Reposition patient and provide reassurance. Consider stopping any clinically assisted hydration/ nutrition Give verbal information, and offer written information to carers at the Consider use of either Glycopyrronium 200mcg sc stat OR Hyoscine butylbromide 20mg sc stat Consider use of syringe driver either Glycopyrronium 600-1200mcg/24hrs OR Hvoscine butvlbromide 60-120mg/ 24hrs Think: Consider alternative causes of secretions if no response to above. Eg. pulmonary oedema, infection or gastric Think: Is there distress or breathlessness? Is an opioid or benzodiazepine necessary? Think:

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Is there a role for

suction?

Guidelines for the management of noisy respiratory secretions at the end-of-life. PTHC 2017.



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