

Notes of the meeting of the South and West Devon Formulary Interface Group

Friday 14th November 2014, 2pm – 4.30pm

The Watermark, Erme Court, Leonards Road, Ivybridge PL21 0SZ

Present:	Andrew Gunatilleke (AG), Consultant, Chair Iain Roberts (IR), Lead MO Pharmacist Emma Hewitt (EH), Joint Formulary Pharmacist Larissa Sullivan (LS), Interface Pharmacist Bill Nolan (BN), GP Petrina Trueman (PT), Joint Formulary Pharmacist Jeremy Morris (JM), Formulary Pharmacist Phil Melluish (PM), GP Paul Manson (PLM), Lead MO Pharmacist Carol Webb (CW), Joint Formularies Technician Margaret Hinchliffe (MH) Mark Stone (MS), Community Pharmacist Steve Cooke (SC), Chief Pharmacist	South Devon NHS Trust South Devon & Torbay CCG NEW Devon CCG NEW Devon CCG South Devon & Torbay CCG NEW Devon CCG Plymouth Hospitals NHS Trust South Devon & Torbay CCG NEW Devon CCG NEW Devon CCG Lay member Devon LPC Plymouth Community Healthcare
In attendance	Tony Perkins (TP), Medicines Optimisation Pharmacist (For item 4) Alan Desmond (AD), Consultant Gastroenterologist Hilary Pearce (HP), Clinical Effectiveness Pharmacist (For item 13)	NEW Devon CCG South Devon NHS Trust NEW Devon CCG
Apologies	Paul Humphriss (PH), Head of Medicines Management Elena Mercer (EM), Formulary Pharmacist David Gwynne (DG), GP	Torbay and Southern Devon Health and Care NHS Trust South Devon NHS Trust NEW Devon CCG

1. **Welcome:** apologies as noted above.

2. **Notes of last meeting:**

The notes of the meeting of 12th September 2014 were agreed

Action list outstanding from the previous minutes, not on the agenda:

- Brand names for epilepsy treatments, it was commented that this issue has been outstanding for some time.

Action: to follow up the issue of brand name advice for epilepsy treatments in the formulary. IR

- Lipids: this piece of work is still being consulted on.

Osteoporosis

The proposed entry into the formulary was presented and discussed. This piece of work had been conducted across Devon by Stuart Kyle from North Devon District Hospital and circulated to secondary care clinicians in the acute trusts for comment.

- It was agreed to change risedronate from blue to green first-line. It was also agreed to change denosumab from amber to blue second-line. There are on-going discussions about the service currently provided by Torbay Hospital which may be changing to ask GPs to provide denosumab treatment for osteoporosis.

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- The comment was made about denosumab now having a license to treat men.
Action: to check this licensing of denosumab for men and amend the wording of the guidance accordingly. CW
- The MHRA information on calcium levels and ONJ for patients receiving denosumab has been added.
- Vitamin D, Invita D3 to be added to the formulary. Colecalciferol capsules to be removed from the formulary.

It was agreed to add this clinical guidance to the formulary, aligning the information across Devon.

3. Melatonin update

The background to the proposed formulary entry was given to the meeting. There was discussion about the Drug Tariff costs of the unlicensed liquid preparations. It was agreed to add a note about the preferred option due to cost, 5mg/5ml.

It was agreed to add these notes to the formulary.

4. DuoResp Spiromax® inhaler

IR and TP declared an interest on behalf of the CCG. Teva are likely to be sponsoring some training on inhaler technique.

DuoResp Spiromax® is a bioequivalent inhaler to Symbicort® and it is proposed that this be added into the formulary. It is only licensed for adults at the moment. The device is different and may be easier to operate by some individuals. DuoResp Spiromax® cost 20% less than Symbicort®, giving a significant saving opportunity. It was acknowledged that wholesale switching would be difficult as patients, would need to be done at a review of patient's treatment.

It was agreed that DuoResp Spiromax® be added into the formulary

5. NICE CKD/Hypertension update

NICE published new guidance on CKD, CG182, this needs to be reflected in relevant sections of the formulary.

NASIDs – to include the renal adverse effects

Renin-angiotensin system – revised text and monitoring advice

AF – CG182 recommends apixaban in preference to warfarin in these patients. There is a differing of opinion between renal and haematology clinicians. It was agreed to leave this information until there has been confirmation from the South West Network.

Bisphosphonates – it was agreed to add information from the SPCs to the monograph for the individual drugs

Vitamin D – inclusion of guidance in the formulary for prescription of vitamin D in CKD patients

Aspirin – to add information to the drug monograph and other appropriate places regarding increased bleed risk in CKD patients

Hypertension – the guidance in the formulary has been reviewed and appropriate information added to the text. New to the S&W formulary would be information on hypertension in pregnancy, diabetic patients and patients with CKD

It was agreed to add this clinical guidance to the formulary, aligning the information across Devon.

6. NICE CG184 Dyspepsia

This guidance updates that currently in the S&W Formulary. Some amendments were suggested by AD, these were agreed:

- To remove the diagnosis and treatment information on Barrett's oesophagus
- To expand the information severe conditions to use the wording in the NICE Guideline 1.6.10
- There were discussions about the terms full-dose and low-dose PPI.
- It was agreed to change the wording on continuing treatment to 'the lowest dose which controls symptoms'.

7. Dermatology Ciclosporin Western Locality shared care

Derriford wish to change the brand of ciclosporin used for their patients to Capimune® from Neoral®. The amended shared care guidance was agreed.

Action: to email the dermatologists in South Devon Hospital to ask if they are considering changing.

JM**8. TA318 Lubiprostone**

It is proposed that lubiprostone be categorised as a specialist initiated drug, this was agreed. It was agreed that it was appropriate for treatment to be initiated by a specialist clinician, to remove the words 'with experience of treating chronic idiopathic constipation'. There was discussion about treatment if lubiprostone is not effective after two weeks, it was agreed to amend to wording to indicate that treatment should be stopped, rather than reconsidered. It was suggested that the use of lubiprostone should be monitored in 12 months' time.

9. TA315 Canagliflozin

It is proposed that canagliflozin be categorised as a specialist initiated drug, this was agreed. Notes are to be added to indicate its place in treatment and that it is to be used in line with NICE, although licensed for wider use.

10. Western Locality Continence Formulary

Some amendments to the Continence Formulary have been suggested, these were agreed.

There was some discussion about a Continence Formulary to cover the South Devon area.

11. Expanded notes for stopping dementia treatments

The Formulary had been approached to increase the information given in this area. It was agreed that this information was very helpful and should be included in the formulary

12. Oxycodone brand change

The secondary care central buying contract has changed to purchasing Lynlor® standard release and Oxylan® for modified release. There was some discussion regarding this and changes that could be made in the formulary to be able to realise savings within primary care. It was agreed to recommend Longtec® modified release and Shortec® standard release. It was asked that a statement be put into the formulary that within secondary care generic prescribing for these items should be used.

13. Recent drug decisions including NICE

- CPC decision to commission second-generation antipsychotic depot injections for schizophrenia. A proposed amended formulary entry was presented where aripiprazole, olanzapine and paliperidone are included in the formulary as red, secondary care only drugs until a service agreement is produced to enable these to be administered in primary care.

It was noted that pipotiazine is to be discontinued in March, a note to be added that no new patients should be commenced on this treatment.

It was also noted that, although risperidone is to be no longer commissioned it needs to stay in the formulary for current patients. This was agreed.

It was asked that the notes on olanzapine be expanded to include the requirement for those prescribing and administering to have completed the appropriate training package. To also include that after administration observation of the patient is required for a period of 3 hours.

14. MHRA Drug Safety Updates

September:

- Denosumab, these notes have been included in the osteoporosis guidance
- Nitrofurantoin contraindicated in patients with eGFR less than 45ml/min, this information to be included in the formulary

October: This was noted.

15. Unlicensed medication

Some expanded notes on unlicensed medication are proposed to be added into the formulary. This was agreed.

It was also asked that a link to the comprehensive Derriford guidance be added and to any guidance that South Devon Hospital might have.

As this is the last Formulary meeting that Emma Hewitt will be present at, having finished her secondment, she was thanked for all the work she has completed for the formulary.

Next meeting: Friday 9th January 2015 2pm – 4:30pm The Watermark, Ivybridge PL21 0SZ

South and West Devon Formulary Group – Action log

Date	Action	Responsible	Completed
Dec 13	To brief the committee on brand names for epilepsy treatments when available	IR	
Nov 14	To follow this issue up.	IR	
Nov 14	Check the licensing of denosumab for men and amend the wording of the guidance accordingly.	CW	Treatment of men would remain within secondary care
Nov 14	To email the dermatologists in South Devon Hospital to ask if they are considering changing from Neoral® to Capimune®	JM	