Drug and Therapeutics Committee - Minutes -approved

Date / Time 14th January 2016

Venue The Board Room, Alderson House, HRI

Chair Prof A Morice, Chair, Professor of Respiratory Medicine

Notes / Action Points Mrs Wendy Hornsby, Senior Pharmacy Technician

Quorate: Yes / No Yes

Attendance Mr S Gaines, Professional Secretary, Senior Principal Pharmacist – Clinical Services

Mr D Corral, Chief Pharmacist, Clinical Director Therapy & Therapeutics

Mr P O'Brien, Deputy Chief Pharmacist

Mrs J Lyon, Head of Medicines Management, North Yorks and Humber CSU

Prof M Lind, Vice Chair, Professor of Oncology Mrs C Grantham, Medicines Management Nurse

Dr H Klonin, Consultant Paediatrician

Dr E Williamson, Consultant Microbiologist

Apologies Dr O Ogunbambi, Consultant Rheumatologist

Mr P Renwick, Vascular Surgeon

Agenda No	Item	Discussion	Decision Made	Action	Lead	Due Date	Progress /Date Closed
2016.01.01	Apologies	As above.					
2016.01.02	Declarations of Interest	None	Noted.	No further action			01/16
2016.01.03	Minutes of the previous meeting	The minutes were accepted as a true record.	Noted.	No further action			01/16
2016.01.04	Action Tracker	Opioid Conversion Chart Awaiting amended version before adding to website.	WH to chase.	WH to chase	WH	11/15	
		New Product Requests ML has not yet written to applicants but formulary has been updated.	ML to write.	ML to write	ML	01/16	
		New Product Requests SG has asked endocrinology for their preferred first line option. They have responded with empagliflozin and the formulary has been amended to say this.	Action complete.	No further action			01/16
		NICE Guidance Vortioxetine has been added to formulary as via chairs approval, prior to discussion at HFT & HERPC.	Action complete.	No further action			01/16
		NICE Guidance Hepatitis C medicines list still to be added to main body of formulary. LC (Infectious Diseases Pharmacist) is updating format.	WH liaising with ID pharmacist.	WH to update formulary	WH	01/16	
		NICE Guidance TA368 Apremilast - SG had checked - to be discussed on agenda under this month's NICE guidance.	Action complete.	No further action			01/16

NICE Guidance	Action	No further action			01/16
NG 24 Blood Transfusion – SG has asked Dr Saleh to discuss at thrombosis committee.	complete.	Tro fairner denom			0.7.10
NICE Guidance NG24 Blood Transfusion – FU not present to discuss if raised within surgery dept.	FU not present.	FU to feed back next time	FU	01/16	
MHRA DSU Vemurafenib – ML has discussed potentiation of radiation toxicity with Prof Maraveyas.	Action complete.	No further action			01/16
Dalteparin/Tinzaparin Switch KMc had provided figures to HEY pharmacy on costs to primary care.	Action complete.	No further action			01/16
Dalteparin /Tinzaparin Switch WH added to MMIG agenda for discussion.	Action complete.	No further action			01/16
Dalteparin/Tinzaparin Switch DC had added to OGC agenda for discussion, but the meeting was cut short and therefore not discussed. This will be on the February agenda for discussion.	Discuss next time.	DC to feed back next time	DC	01/16	
The committee further discussed issues surrounding the switch and agreed to discuss further once outcomes from OQC, HERPC and Safe Medication Practice Committee were known.					
Guidelines for Acute Perioperative Pain Relief in Adults SG has meet with Caroline Weetman to make amend minor amendments to the guideline.	Action complete.	No further action			01/16
Correspondence Received ML has not yet written to Mr Burnett about approval of Tafluprost, but it has been added to formulary.	ML to write.	ML to write	ML	01/16	
Correspondence Received Nutilis switch to Nutilis clear was discussed at MMIG and it was agreed to switch at HEY and review the situation in 6 months'	Action complete.	No further action			01/16

		time.					
2016.01.05	New Product Requests	Sucroferric oxyhydroxide (Velphoro) – Katherine Durrans (Highly Specialised Renal Dietician) The application was submitted with a view to phasing out lanthanum and swapping patients over to Velphoro. First line choices would still be calcium-based products and sevelamer as Sucroferric Oxyhydroxide has a higher rate of adverse effects compared to sevelamer.	Approved.	AM to write to applicants WH to update formulary	AM/WH	02/16	
		Cutimed Sorbact Gel – Karen Harrison/Angela Oswald This application was submitted by the Tissue Viability Team who have already discussed this at the wound group and recommended it for use. The dressing is interactive so will be sourced via pharmacy.	Approved.	WH to liaise with applicant over which size products from range are required.	WH	02/16	
		Naloxegol – Dr Elaine Boland Application submitted in line with TA345. The committee agreed that the HERPC constipation guideline would need to be updated, and should include the reinforced message that prophylactic laxatives should always be considered for patients prescribed opioids.	Approved for consultant use, to ensure it was used appropriately.	SG to liaise with MM over update of guidance	SG	02/16	
		Lubiprostone – Prof G Duthie Application submitted in line with TA318. The committee agreed that the constipation guideline would need to emphasize that lubiprostone was available for short term treatment (2-4 weeks) via gastroenterology or GI surgery specialists only.	Approved for specialist use as a red drug.	SG to liaise with MM over guidance	SG	02/16	
2016.01.06	NICE Guidance	NICE Guidance NG27 Transition between inpatient hospital settings and community or care home settings for adults with social care needs	Noted.	No further action			01/16
		NG28 Type 2 Diabetes in adults: management. This guideline includes reference to repaglinide in note 3, page 20 but the Trust formulary includes nataglinide only.	Noted. SG to ask ML to discuss with endocrinology.	SG to liaise with ML	SG	02/16	

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Diabetes Pharmacist (ML) to discuss this with the endocrinology team.					
 NG29 Intravenous fluid therapy in children and young people in hospital HK advised the committee that the HEY guideline had just been updated but that she would be reviewing this again. 	HK will discuss with paediatric Pharmacist (AK) outside of meeting.	No further action			01/16
NG30 Oral health promotion: general dental practice Fluoride toothpaste & mouthwash were already formulary.	Noted.	No further action			01/16
 NG31 Care of dying adults in the last days of life SG had already asked MM to include Just In Case box prescribing on the next MMIG agenda for discussion. 	Noted.	No further action for D&T			01/16
NG32 Older people: independence and mental wellbeing	Noted.	No further action			01/16
 TA369 Ciclosporin for treating dry eye disease that has not improved despite treatment with artificial tears 	Noted, Ikervis 0.2% on formulary.	No further action			01/16
TA370 Bortezomib for previously untreated mantle cell lymphoma	Noted, on formulary.	No further action			01/16
TA371 Trastuzumab emtansine (Kadcyla) for treating HER2-positive, unresectable locally advanced or metastatic breast cancer after treatment with trastuzumab and a taxane – not recommended by NICE, however ML commented that he currently had a patient in full remission who had received this treatment.	Was on CDF list. Current patients can continue with treatment.	No further action			01/16
 TA 372 Apremilast for treating active psoriatic arthritis – not recommended by NICE. Will not be commissioned by the CCGs, so only available to new patients via IFR. 	Was on Trust formulary.	WH to amend formulary to say for existing patients only	WH	02/16	

		 TA373 Abatacept, adalimumab, etanercept and tocilizumab for treating juvenile idiopathic arthritis 	Noted, on formulary.	No further action			01/16
		 TA374 Erlotinib and gefitinib for treating non-small-cell lung cancer that has progressed after prior chemotherapy ML noted that all HEY patients are EGFR-TK mutation tested. 	Noted, erlotinib is on formulary.	No further action			01/16
		Osimertinib AZD9291 EAMS Public Assessment Report The MHRA EAMS documents had been circulated for awareness.	ML agreed to submit an application if a patient presents who would benefit.	No further action			01/16
2016.01.07	MHRA Drug Safety update December 2015	 Thalidomide – decreased dose in patients >75yrs Mycophenolate not to be used in pregnancy Bisphosphonates risk of osteonecrosis 	Noted, all formulary	No further action required			01/16
2016.01.11	Correspondence Received	CCG Representative at D&T Committee AM has received a reply from Dr Dan Roper of the Hull CCG explaining that the job description for the new Hull GP prescribing lead has been altered to include their attendance at HERPC, only not D&TC. ER CCG was currently advertising for a GP prescribing lead, so the post was vacant. In spite of this, the committee still felt that GP attendance at D&T was valuable and AM agreed to respond to Dr Roper. JLy agreed that she would also discuss with the CCG's, although it is possible with the CSU restructuring this could be JLy's last meeting. The chair thanked JLy for her	AM to write to Dr Roper. JLy to discuss with CCG's.		AM	02/16	
2016.01.01	Chairs Approvals	 Valuable contribution to D&T committee. Thiotepa – CNS Lymphoma – Dr James Bailey Deferiprone – Superficial Siderosis – Dr A Raman Tacrolimus SR (Envarsus Brand) – Prophylaxis of Organ Rejection – Dr M Edey Genvoya (Elvitagrevir/Cobicistat/Emtricitabine/Tenofovir 	Noted.	No further action			01/16

		Alafenamide Fumarate) – HIV – Dr Thaker Natamycin Eye Drops – Fusarium Keratitis – Mr Stewart Recommended on Royal College Guidelines, so Mr Stewart will submit an full new drug application soon.					
2016.01.13	Issues to escalate to OQC	Potential dalteparin to tinzaparin switch, as above.	Concerns to be discussed at OQC.	DC to discuss at OQC, as D&TC representative	DC	02/16	
2016.01.14	Any other Business	SG informed the committee of a new product Idarucizumab Praxbind ®, which is manufactured by Boehringer as a reversal agent for dabigatran. Boehringer have offered two free vials for treatment of the first patient. Thrombosis committee have discussed the product and expect that Haematology will submit a new product request to D&TC, as well as reviewing the reversal of anticoagulation guideline. SG informed the committee that a meeting had taken place between pharmacy, primary care and dermatology to discuss apremilast and emollients. The committee agreed that if dermatology wished to remove/add several emollients to the	Await new drug application from Haematology. SG to feedback to MM.	No further action	SG	02/16	01/16
		formulary it would be acceptable to submit a paper detailing the proposed changes, rather than a large number of new product requests for similar items.					
2016.01.01	Date and Time of Next Meeting	Thursday 11 th February 2016, 8.15am – 9.30am. The Board Room, Alderson House, HRI					