

Adult Medicines Management Committee Meeting to be held on 12th May 2022 [virtual], 12.30-2pm Pharmacy Meeting Room, 4th Floor, Pharmacy, Oxford Road Campus: <u>Minutes</u>

Present	Area of Representation/Position	Initials
(Cha	air) Clinical Director Cardiac Anaesthesia and Intensive Care. MMC Chair (ORC, WTWA)	
	Deputy Medical Director (NMGH MMC lead (NMGH)	
	Medicines Information Pharmacist (ORC, WTWA)	
7	Medicines Optimisation Pharmacist/ Professional Secretary MMC	
	Head of Medicines Optimisation (ORC, WTWA, NMGH)	
	Medicines Optimisation Pharmacist (NMGH)	
3	High-Cost Drugs Pharmacist (WTWA)	
	High-Cost Drugs Pharmacist (ORC)	
	SMH Pharmacist	
	Microbiology Consultant	
	MHCC pharmacist.	
	Trafford Consultant	₹
	Respiratory/CF consultant	
	Critical Care- Head of Nursing.	
	High-Cost Drugs Pharmacist.	
	HIVE Pharmacist	
	Medicines Optimisation Scientist	
	SMH consultant	
	Consultant Immunologist	
n attendance		
	Pain Matron	9
Apologies for Absence		
	Lead Medicines Optimisation Pharmacist.	
in contract of	Assoc. Medical Director (MRI)	
	Diabetes Consultant, TGH	

1.Min	utes.			
116.22		Draft Minutes April 14 th 2022 Noted		
2.Mat	ters Arising			
17.22	,	See Section 10.0 for Policies approved via chairs action for NMGH transaction. Noted- Fuller discussion occurred later in the agenda.		Gara
0	3.New Drugs		200000000000000000000000000000000000000	110000000000000000000000000000000000000
ō	4.NICE		2000 00000 00000 0000 00000 00000 0000 00000 00000 0000 00000 00000	110/2001 1100 1100 1100 1100 1100 1100 1
T O 1 1	118.22	NICE TA Tracker (May 2022) Palforzia- the ongoing paediatric discussions were noted. Adults will follow lead of the paediatrics who have the larger cohort of these patients. Noted		
₹ 0)5.SSC			
1	119.22	SSC Tracker (May 2022) Noted		
	120.22	HST19 Elosulfase alfa HST19 for treating Mucopolysccharide 4A (Morquio disease) Removed from agenda as for PMMC rather than Adult MMC. Noted that previously all HSTs had come to both committees for noting routinely. However, on balance this was not thought to be needed going forward.		
0	6.GMMMG			
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	07. Amendments					
	121.22	Pristina • • • Approve	Noted that mycobacte To be adde	mycoplasma genitalia (BASSH guidelines) this application followed on from IPR last week. hat the minutes should state mycoplasma rather than eria. ed to the formulary- as per BASSH guidance.		
	08.Urgent	All	122.22	Urgent IPR requests (April 2022)		
	09.Non-urgent	CCG	123.22	IFR- CCG- Verbal update on IFR position. highlighted that there is change for CCG IFR processes.		
				There is an ongoing proposal with GM reviewing how CCG commissioned IFR therapies are processed. Currently this is at the consultation and comments stage.		
				Sequential use of Biologics IFRS: Sequential use of Biologics for CCG funded indications will no longer require an IFR through MMC as these will be reviewed by relevant MDTs.		
				Other IFR: Likely the more complex IFRs will still come to MMC e.g., Dual Biologic use for forward on to an CCG panel. noted there is likely to be a more centralised approach to IFR panels, that will reduce the inequity across GM in the current system.		
equests				At present the status quo remains, until the above is fully worked through and implemented. However HCD team are aware that there have been a few areas of confusion. Currently aware of delay for one patient that has a delayed decision, working to reduce this.		
Individual Requests			Sent out after agenda	IFR- CCG- Dual Biologic (5 th line)- Vedolizumab (4-8 weekly) added to existing Certolizumab for Crohns disease and Ankylosing Spondylitis. • **Etalked through IFR.**		
<u></u>				 noted that this IFR for dual biologics has more strength in the evidence base/rationale (i.e vedo + anti-TNF) than other Dual Biologics in previous months where there had been no evidence really at all. COS concerned that approval based on this evidence base would open the flood gates to more of these proposals. noted that the MO team had separately identified 		
				 case series where vedolizumab or dual therapies had resulted in worse outcomes for These issues should be raised within the IFR for transparency of this risk benefits at play. suggested that as Dual Biologic applications are 		
			***************************************	becoming more frequent that a set of principles be agreed including o Endorsement from both parent specialities (e.g. gastro and rheum)	,	
				 Safety data presented on the specific combination. Infection risk assessment. Further answers to questions required from the team. 		
		PBRE			s	y sisilastainaa
	10.Policies (MO) 124.22	NMGH	- Summary	of Policy adoption/adaption		
lents	1 to 7 s to 6.	all the MMPs prior to the NMGH transaction on 1st May 2022. The working group divided the work in to				
Documents		•	Policies the Policies the prescribing	at could be adopted in current form. at could not be adopted due dependence on the NCA electronic g system. (Injectable Medicines + others see paper). There is a		
				orking group that continues to work through these differences /E implementation in September.		

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-	125.22	 Policies that required amendment. (Medicines Policy, PGD Policy, NHS Foun Potassium Policy). These have been through MMC in April or via Chairs action between April and May (see below) Policies that MFT needed to adopt. (Electronic – Down Time Policy) A summary has been sent to NMGH staff. All relevant policies are available on the NMGH 'twin policy/document sites.' noted that it was agreed yesterday that from June all drug applications will come through the current MFT (ORC, WTWA) MMC meeting to include NNMGH. asked about HIVE training for NMGH due to +++ changes close to HIVE. aware of this issue and it is being raised within the working group. 	dation T	ust
-	126.22	Approved via Chairs action 26th April 2022- On agenda for noting Noted Drug Prescription and Administration Record for NMGH ePMA down time policy for NMGH Approved via Chairs action 26th April 2022- on agenda for noting.		
		Noted		
	11. Policies (MFT)			
	12.Guidelines- Ap	proved outside of meeting by MOT		
	13.Guidelines- Ha			
		Anticoagulation Documents		
	127.22	 MFT- Dalteparin Dosing guide asked for clarity from on the final conversations on the use of enoxaparin in cardiology (if the rest of the trust is using dalteparin the in the main). 		
		 Masked for clarity from the plans for warfarin charts to be rolled out more widely across the sites. Previous barrier has been warfarin charts referring to a specific LMWH. This now seems like it might be possible prior to HIVE. Masked if there was a date for NMGH to switch over to dalteparin. noted that this would not likely happen until HIVE implemented due to the 		
		reliance on the NCA electronic prescribing system. • asked about the plan for SMH. noted that there is a separate steering group still working on this. • noted that there is a plan to outline when patients admitted on other LMWH are switched to the trust LMWH in a safe and timely fashion. Approved.		
	128.22	MFT-VTE Medical Prophylaxis guidelines Approved.		
	129.22	MFT-VTE COVID Medical Prophylaxis and Treatment guidelines Approved.		
	Tabled	MFT- VTE Surgical Prophylaxis- Guideline Approved.		
	130.22	MFT-VTE Surgical Prophylaxis- Orthopaedic Guideline Approved.		
	131.22	 MFT-VTE Treatment Guideline in CKD Treatment table to be removed, and instead to point to the dalteparin dosing guide- which is colour coordinated to syringe strengths. Title needs amending to align with other VTE documents on the policy hub. Approved pending above points. 		
	132.22	MFT- Perioperative anticoagulants Approved.		
	422.00	Dishetoe Decuments	1	
	133.22	Diabetes Documents MFT- Hypersmolar Hyperglycaemic Syndrome (HHS)- Adult Guideline • Confirmed prior to the meeting that she is happy with this as the DM representative at MMC Approved.		

GM. Community Type 2 Diabetes – Management Education and Support. Cardio Renal Pathway Comments from — Derior to meeting Page 2 - amend Hb1AC to HbA1 amend HbA1 to target to 48 (not <48) for patients with type 2 DM as per NICE NG28 Guideline amend Alao consider individualised target HbA1c levels for those with problematic hypoglyceemia, certain co-morbidities and or freilly Further comments expected from — attention — the religion of the Comments during meeting Client Comments during meeting Cliently on who the authors are for feedback. Clarity as to where this SM footprint document will be approved and hosted. Suggested that this should be GMMMC, but members are aware that this has not yet been approved through this route. Clarity from — MCMFT DM pharmacist) and — Cardiology pharmacist) as to whether this replaces the existing SGLT CV reduction document on the MFT intranet. Noted- Comments to be feel back to authors. GM- Choosing who to treat with SGLT inhibitor. Similar comments to above regarding authorship and approval routes. Noted- Comments to be feel back to authors. Pain team Documents MFT Gabapentin to Pregabalin Conversion Noted- Comments to be feel back to authors. Pain team Document MFT deparent to Pregabalin equivalence and switching, Also contains more detailed dosing on renal information. Committee fet that approving this paperwork would create additional/duplicated work updating in 2-3 years litme. — Control of the meeting that the information approving the paperwork would create additional/duplicated work updating in 2-3 years litme. — Indied from an SMH perspective, that there had been a recent alert about Pregabalin in pregnancy, and that there should be signosting information to this warning. — Indied from an SMH perspective, that there had been a recent alert about Pregabalin in pregnancy, and that there is broud be signosting information to this warning. — Indied from an SMH perspective, that there had been a recent alert about Pregabalin in pregnancy, and that there is increased	•	NHS Foun	dation T
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		Noted that the document control page Foun needs updating. The medicines changes stated occurred in the previous update. There are no medicine updates in this version. Approved.			
	140.22	SMH- Transcervical Ultrasound Guided Radiofrequency Treatment of Uterine Fibroids (Sonata) Approved.			
	Sent out after agenda	COVID Adult Guidelines (Addition of Bariticinib) Noted that since this was circulated on Tuesday, that the team have made 2 x further updates. Final version will be circulated after the meeting, with final comments requested by COP on Friday 13th May.			
	14.Guidelines- OR	C/TGH/ WH/NMGH			
	15.Guidelines- ML	CO		weining was	
	16.Prescriptions				
	17.Patient Informa	tion Leaflets (PILS)			
	18.Procedures		24 (424) (444) 24 (424) 24 (424)	(100)	
	19. Out of stock m	iedicines			
	20. Other				
	141.22	Atezolizumab (Adjuvant- Introduction of Closed System Device to aid in the preparation at Ward Level.) Tisk assessment noted. Barriers to aseptic preparation noted. Closed System Device supported. Prequested that the financial implications of using Closed System vs outsourcing be quantified. Approved			
	21.Drug Safety Ale				
	142.22	Drug Safety Alert (April 2022) Noted			
Other	22.Finance				
δ	23.PGD Report				
	143.22	PGD Report (March April 2022) • Sasked that this alert be circulated after the meeting. Noted		-	
	24.AOB		11.000		
	Tabled	NMGH to be invited to MFT MMC from June 2022. Letter to be written to the following to request pharmacy, consultant and nursing representation.			
		Letters to be written.	<u> </u>	<u> </u>	