NHS Portsmouth CCG South Eastern Hampshire CCG Fareham and Gosport CCG Portsmouth Hospitals NHS Trust Southern Health NHS Foundation Trust Solent NHS Trust

Area Prescribing Committee Meeting, 1.00pm on Friday 16th October 2020 Via MS Teams Draft Notes

5.20.1	Attendance Alastair Bateman (Chair), Jo Williams (secretary), Mike Stewart, Nick Moore, Vanessa Lawrence, Phil Foster, Karen Atkinson, Simon Cooper, Jon Durand, Kieran Hand, Helen McHale, Debby Crockford, Luke Groves Apologies for absence	
	Kevin Vernon, Charlie Mitchell, Jason Peett	
5.20.1.1	Declarations of Interest None to declare. JW to re-circulate form to members who need to submit annual declaration.	
5.20.2	DRAFT Notes of last meeting Accepted as an accurate record.	
	Action log APC action log October 2020.docx	
5.20.3	Matters arising None Chairs approvals: Care homes re-use of medicine scheme document was supported by the committee.	
5.20.4	Formulary Management – applications for approval	
5.20.4.1	Oral semaglutide Submitted by Phil Newland-Jones & Iain Cranston The committee received a joint application from PHU and UHS for the addition of oral semaglutide to the formulary. It was acknowledged that no NICE TA is expected for this preparation, however the SMC has supported it's inclusion on to Scottish formularies. The application request that oral semaglutide is available as a second line option for those patients where subcutaneous would be considered but where it is unsuitable. There are bioavailability concerns and the requirement to follow counselling advice. There does not appear to be advice available on switching between subcutaneous and oral preparations.	
	APC decision The committee accept oral semaglutide for restricted use as per the application request as a second line option where a patient would be considered for semaglutide but where subcutaneous use is not advisable. JW to discuss formulary status with Iain Cranston and to bring back to the committee for discussion.	JW

5.20.5	Drug therapy and shared care guidance for approval	
5.20.5.1	PHU Shared Care Guidance for Denosumab (XGEVA)	
	Submitted by Beverly Miell	
	This is an update of the current Xgeva shares care guideline with minor	
	updates including addition of dose of Adcal D3 and patient	
	responsibilities.	
	APC members noted that the new template had not been used for the	
	update so it was requested that the information on the agreed consent	
	process was added, in addition to changing the primary care suggested	
	calcium/vitamin D3 options to those included on the formulary.	
	APC decision	
	The committee support the approval of the guideline following the	
	requested changes being made.	
5.20.6	Items for note/consultation	
5.20.6.1	NICE Guidance	
	NICE developments:	
	NICE updates August 2020	
	TA 641: Brentuximab vedotin in combination for untreated systemic	
	anaplastic large cell lymphoma	
	Brentuximab vedotin with cyclophosphamide, doxorubicin and	
	prednisone (CHP) is recommended, within its marketing authorisation,	
	as an option for untreated systemic anaplastic large cell lymphoma in	
	adults. It is only recommended if the company provides brentuximab	
	vedotin according to the commercial arrangement. Resource impact: This technology is commissioned by NHSE. No	
	significant resource impact is expected.	
	Action required: The formulary entry for Brentuximab vedotin will be	
	updated with a link to NICE TA641.	
	apaded with a link to those 17.011.	
	TA 642: Gilteritinib for treating relapsed or refractory acute myeloid	
	leukaemia	
	Gilteritinib monotherapy is recommended as an option for treating	
	relapsed or refractory FLT3-mutation-positive acute myeloid leukaemia	
	(AML) in adults only if the company provides gilteritinib according to the	
	commercial arrangement.	
	Gilteritinib should not be given as maintenance therapy after a	
	haematopoietic stem cell transplant.	
	Resource impact: This technology is commissioned by NHS England,	
	a local resource impact planner has been developed as the price of the technology is commercial in confidence.	
	Action required: Gilteritinib will be added to the formulary as a Red	
	agent for use as per NICE criteria.	
	agenties are por rived emonal	
	TA 643: Entrectinib for treating ROS1-positive advanced non-small-cell	
	lung cancer	
	Entrectinib is recommended, within its marketing authorisation, as an	
	option for treating ROS1-positive advanced non-small-cell lung cancer	
	(NSCLC) in adults who have not had ROS1 inhibitors. It is	
	recommended only if the company provides entrectinib according to the	
	commercial arrangement.	
	Resource impact: This technology is commissioned by NHSE. No	
	significant resource impact is expected.	
	Action required: Entrectinib will be added to the formulary as a Red agent for use as per NICE criteria.	

TA 644: Entrectinib for treating NTRK fusion-positive solid tumours

Entrectinib is recommended for use within the Cancer Drugs Fund as an option for treating neurotrophic tyrosine receptor kinase (NTRK) fusion-positive solid tumours in adults and children 12 years and older if:

- the disease is locally advanced or metastatic or surgery could cause severe health problems and
- · they have not had an NTRK inhibitor before and
- they have no satisfactory treatment options.

It is recommended only if the conditions in the managed access agreement for entrectinib are followed.

Resource impact: This technology is commissioned by NHS England, from the Cancer Drugs Fund.

Action required: Entrectinib will be added to the formulary as a Red agent for use as per NICE criteria.

NG180: Perioperative care in adults

This guideline covers care for adults (aged 18 and over) having elective or emergency surgery, including dental surgery. It covers all phases of perioperative care, from the time people are booked for surgery until they are discharged afterward. The guideline includes recommendations on preparing for surgery, keeping people safe during surgery and pain relief during recovery.

NG 181: Rehabilitation for adults with complex psychosis

This guideline covers mental health rehabilitation for adults with complex psychosis. It aims to ensure people can have rehabilitation when they need it and promotes a positive approach to long-term recovery. It includes recommendations on organising rehabilitation services, assessment and care planning, delivering programmes and interventions, and meeting people's physical healthcare needs.

NG 178: COVID 19 rapid guideline: renal transplantation

Updated guidance: in **August 2020**, NICE added recommendations for regional networks on responding to changes in local prevalence of COVID-19. NICE aligned recommendations for donors and recipients with their COVID-19 guideline on arranging planned care in hospitals and diagnostic services.

NG 174: COVID-19 rapid guideline: children and young people who are immunocompromised

Updated guidance: in **August 2020**, NICE updated the recommendation on safeguarding to remove a link to government guidance that has been withdrawn.

NG 172: COVID-19 radip guideline: gastrointestinal and liver conditions treated with drugs affecting the immune response

Updated guidance: in **August 2020**, NICE updated recommendations on modifications to care in line with their COVID-19 rapid guideline on arranging planned care in hospitals and diagnostic services.

NG 160: COVID-19 rapid guideline: dialysis service delivery

Updated guidance: in **August 2020** NICE changed the recommendation on how long people need to stay in the cohort of patients known to have COVID-19.

NG 125: Surgical site infections: prevention and treatment

Updated guidance: in August 2020, NICE added links to the NICE guideline on perioperative care in adults for additional recommendations on intravenous fluids, cardiac monitoring and blood glucose control in adults.

CG 134: <u>Anaphylaxis: assessment and referral after emergency</u> treatment

Updated guidance: in **August 2020** NICE added advice on prescribing adrenaline injectors before discharge after emergency treatment.

NICE updates September 2020

TA 645: Avelumab with axitinib for untreated advanced renal cell carcinoma

Avelumab with axitinib is recommended for use within the Cancer Drugs Fund as an option for untreated advanced renal cell carcinoma in adults. It is recommended only if the conditions in the managed access agreement for avelumab with axitinib are followed.

Resource impact: This is a Cancer Drugs Fund Technology. **Action required**: The formulary entries for avelumab and axitinib will be updated with a link to NICE TA 645.

TA 649: Polatuzumab vedotin with rituximab and bendamustine for treating relapsed or refractory diffuse large B-cell lymphoma

Polatuzumab vedotin with rituximab and bendamustine is recommended, within its marketing authorisation, as an option for treating relapsed or refractory diffuse large B-cell lymphoma in adults who cannot have a haematopoietic stem cell transplant. It is recommended only if the company provides polatuzumab vedotin according to the commercial arrangement.

Resource impact: This technology is commissioned by NHS England. A local resource impact calculator is provided as the price is commercial in confidence.

Action required: Polatuzumab will be added to the area prescribing formulary with a link to NICE TA 649. The formulary entries for rituximab and bendamustine will be updated to include a link to NICE TA 649.

TA 650: Pembrolizumab with axitinib for untreated advanced renal cell carcinoma

Pembrolizumab with axitinib is not recommended, within its marketing authorisation, for untreated advanced renal cell carcinoma in adults.

TA 651: Naldemedine for treating opioid-induced constipation

Naldemedine is recommended, within its marketing authorisation, as an option for treating opioid-induced constipation in adults who have had laxative treatment.

Resource impact: This technology is generally commissioned by clinical commissioning groups. NHS England commissions services for patients with constipation only when referral to a specialist centre is required. Services for opioid-induced constipation are provided in both primary and secondary care. No significant resource impact is anticipated.

Action required: Naldemedine will be added to the area prescribing formulary with a link to NICE TA 651. JW to link with PHU Gastroenterology to discuss place in therapy and appropriate formulary status.

JW

Brief product information: Naldemedine is a peripherally acting opioid receptor antagonist. Therefore it decreases the constipating effects of opioids without altering their central analgesic effects. The recommended dose is 200 micrograms once a day. BNF list price for Naldemedine 200microgram tablets is £41.72 for 28 tablets.

NG 182: Insect bites and stings: antimicrobial prescribing

This guideline sets out an antimicrobial prescribing strategy for insect and spider bites and stings in adults, young people and children aged 72 hours and over, including those that occurred while travelling outside the UK. It aims to limit antibiotic use and reduce antibiotic resistance.

NG 12: Suspected cancer: recognition and referral

This guideline covers identifying children, young people and adults with symptoms that could be caused by cancer. It outlines appropriate investigations in primary care, and selection of people to refer for a specialist opinion. It aims to help people understand what to expect if they have symptoms that may suggest cancer.

In **September 2020**, NICE clarified when to offer faecal testing for colorectal cancer to adults without rectal bleeding.

NG 159: COVID-19 rapid guideline: critical care in adults

The purpose of this guideline is to maximise the safety of patients who need critical care during the COVID-19 pandemic, while protecting staff from infection. It will also enable services to make the best use of NHS resources.

In September 2020, NICE added guidance on treatment with corticosteroids for people with severe or critical COVID-19, in line with their prescribing briefing on dexamethasone and hydrocortisone.

NG 160: COVID-19 rapid guideline: dialysis service delivery

The purpose of this guideline is to maximise the safety of patients on dialysis, while protecting staff from infection. It will also enable dialysis services to make the best use of NHS resources and match the capacity of dialysis services to patient needs if these become limited because of the COVID-19 pandemic.

In September 2020, NICE clarified their guidance for organisations on planned procedures and emergency pathways for creating dialysis access sites for patients with advanced or end-stage kidney disease.

5.20.6.2 **EAMS**

Nil received

5.20.6.3 **Portsmouth Hospitals FMG update**

The draft nots of the meeting were noted by the committee.

For noting:

- Olatuton LAR has become the CMU contract product for Octreotide. This will be added to the formulary as a Red agent.
- Tenectaplase has been accepted as an option for recanalization before planned thrombectomy in patients who would otherwise be ineligible for treatment due to staffing shortages preventing the transfer of patients whilst receiving an alteplase infusion. This indication is unlicensed.
- Tapentadol liquid has been accepted as an option for the treatment of moderate to severe pain in paediatric patients requiring a liquid preparation. Tapentadol will replace tramadol liquid, which is currently on the formulary for this indication (tramadol liquid is unlicensed).

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5.20.6.4	Solent medicines management update	
	Update provided by Luke Groves.	
	EPMA system has been approved and agreed.	
	Solent have been working with Southampton area PCNs to	
	recruit in to joint posts	
	Solent will be producing their own rapid tranquilization	
	guidelines (previously planned to co-badge with SHFT).	
	However it was noted that the formulation of aripiprazole	
	included in the guidance is not currently listed on the area	
	prescribing formulary. Request to add as a RED agent for use in	
	SHFT and Solent – this was supported by committee members.	
5.20.6.5	Southern Health medicines management update	
0.20.0.0	Verbal update provided by Vanessa Lawrence.	
	Priadel switch work is currently on hold whilst negotiations are	
	on going with the company and NHSE regarding continuing	
	access to the market. No new patients are being initiated on	
	Priadel during this time.	
5.20.6.6	DPC update	
	The summary of the DPC meeting was not received.	
	Melatonin shared care guideline	
	The committee discussed the DPC approved shared care for Melatonin.	
	Although it was felt to be a step forward for the management of	
	melatonin use in primary care there were some questions raised	
	regarding the inclusion of the Colonis brand of melatonin and around	
	the GP/specialists responsibilities.	
	JW to link with DPC to raise these questions with the author.	JW
5.20.6.7	Wound Formulary update	011
	It was noted that there is a request for potassium permanganate to be	
	changed from Green to Amber on formularies (as per HMSG). The	
	committee were keen to discuss the merits of potassium permanganate	
	and questioned why changing the formulary status of the agent would	
	help to improve the safety of prescribing. JW will link with the HMSG to	JW
	discuss the rationale for making such a change.	
	JW also has chased a response to the previous request to remove the	
	products listed within the steroid management of wounds guidance that	
	are not on the formulary. JW to update the group with the outcome.	JW
5.20.6.8	Hampshire Medicines Safety Group	
	Verbal update was provided by HMSG members.	
5.20.6.9	Drug Safety Update and Patient Safety Alerts	
	The August and September Drug Safety Updates were noted by the	
	committee.	
5.20.6.10	Regional Medicines Optimisation Committees	
	No new guidance published	
5.20.6.11	NHSE Specialised Commissioning	
	Nil received	
5.20.6.12	Priorities committee	
E 00 0 40	Nil received	
5.20.6.13	Coroner's report – Epipen	
	The report was discussed by the committee and actions considered.	
	Suggestion included to review prescribing support messaging and	
	ensure that there is a clear message that two pens are required for all	
	patients, and that brands are not interchangeable.	
	CCG processes and the governance of messages including how	
l	decisions are made for switching between different brands are made	

	was an area for concern within the report. PF was asked to review the	PF
	governance around these processes.	
5.20.7	Any other business:	
	APC secretary cover.	
	Helen McHale was welcomed to the group who will be providing cover	
	for the professional secretarial duties for the APC during JW leave.	
	Antipsychotic reducing advice – submitted by Phil Foster	
	PF presented a new document that has been produced by the Fareham	
	and Gosport and South Eastern Hampshire Medicines Optimisation	
	Care homes Team to support the reduction and discontinuation of	
	antipsychotics being used for behavioural problems in care home	
	residents. The document has been produced with and has the support	
	of the SHFT OPMH consultants.	
	The APC committee were supportive of the document. SHFT will take it	
	through their internal processes for approval as the plan is that any	
	patient being discharged from the OPMH service with have this information sent to their GP.	
	Solent were also keen to have a similar document; they may look to co-	
	badge this document although one consultant has suggested that their	
	preference is to reduce risperidone 0.5mg in two steps.	
	January Market Strategy and Str	
	Continence Formulary update	
	PF provided an update to the group around the work that he has been	
	leading on including the establishment of a short life working group to	
	produce a HIOW continence formulary. The use of industry sponsored	
	nurses within provider Trusts continues to be a concern for the cost effective use of products and also the directing of prescriptions.	
	enective use of products and also the directing of prescriptions.	
	NPSA Emergency steroid card	
	The alert was discussed and it was noted as a topic within the HMSG.	
	AB and NM will review the availability of information within prescribing	
	systems and consider whether additional prescribing support	
	messaging is needed.	
	DC to provide comments for community pharmacy. Action is ongoing within PHU.	
	Action is origoning within F110.	
	Solent HF team request for sacubitril valsartan support to	
	prescribe – presented by Luke Groves.	
	The Solent Heart failure team are continuing to have challenges with	
	the prescribing of sacubitril valsartan (shared care product). Currently	
	there is just one non-medical prescriber within their team and they are	
	therefore asking for support from GPs to prescribe the medication whilst	
	it is being up-titrated. A revised document has been produced. Requested changes for the	
	document to get APC support:	
	Remove remark that prescribing responsibility has not been handed	
	back to GP (as whoever signs the prescription does have prescribing	
	responsibility)	
	Ensure that this is for the up-titration of doses only and not initiating the	
	product.	
	Changes at PUII	
	Changes at PHU KH informed the committee that he will be taking on a new national	
	Antimicrobial Stewardship role and will therefore be leaving PHU, this	
	will be his last APC meeting. The committee congratulated him and	
	wished him well in his future role.	<u> </u>

5.20.8	Dates of future r	neetings:		
	2020	2021	2022	_
	18 th December	19 th February	18 th February	
		23 rd April		
		18 th June		
		20 th August		
		15 th October		
		17 th December		