



Chelsea and Westminster Hospital NHS Foundation Trust Trust Medicines Group

Summary of Main Points from the Meeting held on Monday 27th July 2020

2. Minutes and Summary Notes from last meeting

On account of the Trust response to the Covid-19 Pandemic, this meeting was held via Zoom.

The summary notes will be disseminated and published on the Trust intranet. A quarterly summary report will be drafted and forwarded to the Trust Patient Safety Group meeting for inclusion on the agenda for September 2020.

3. Matters Arising

The Group noted the matters arising from the previous meeting.

4. Business to be transacted by the Medicines Group

a) Formulary Applications

Full Applications

Melatonin 1mg & 5mg Prolonged-Release Tablets (Slenyto®)

Requested by Paediatrics for the treatment of insomnia in children and adolescents aged 2-18 with Autism Spectrum Disorder (ASD) and / or Smith-Magenis syndrome, where sleep hygiene measures have been insufficient.

This is the only licenced preparation for use in Paediatrics. Many patients with disorders such as autism struggle with the liquid preparation and would benefit from the mini pill option which can be hidden in food to increase compliance.

Paediatric expect 20 patients to be treated at each hospital site who would benefit from switching from liquid melatonin to Slenyto®.

Melatonin liquid (Colonis®) (£0.87/mg) vs Melatonin (Slenyto®) (£0.45/mg)

Outcome: Additional information to be sought to support the decision

Medroxyprogesterone Acetate (MPA) 104mg in 0.65ml suspension for Injection (Sayana-Press®) Requested by GUM for its licensed indication for long-term female contraception.

Sayana-Press[®] is adminstered by subcutaneous injection (one injection per 13 weeks) thereby facilitating self-administration at home, enhancing patient experience and reducing hospital clinic time.

Having this available as an option will hgely help with the management fo appointments during the Covid Pandemic period.

Sayana-Press[®] is not currently included on the NWLIF and therefore it would be expected that a application is made to the panel for its inclusion, to increase flexibility for women should they subsequently wish to obtain contraceptives via their GP.

Cost comparison:

- Sayana-Press® (Medroxyprogesterone Acetate 105mg/0.65ml Injection (SC every 13 weeks) £6.90 per Injection
- Depo-Provera® (Medroxyprogesterone Acetate 150mg/ml Injection (IM every 12 weeks) £6.10 per Injection

Outcome: Approved for addition to the formulary

Ex-panel





Daratumumab (Darzalex®) 1,800mg Solution for Injection

This a new SC product presented as a 1800 mg of daratumumab solution for subcutaneous injection. The dosing schedule will vary according to the indication, but this remains the same as per the IV dosing schedule for each indication.

For use in line with the following NICE Technology Appraisals for daratumumab which include a recommended dose schedule using 16mg/kg body weight by intravenous (IV) infusion.

- Daratumumab with bortezomib and dexamethasone for previously treated multiple myeloma. Technology appraisal guidance (TA573)
- Daratumumab monotherapy for treating relapsed and refractory multiple myeloma. Technology appraisal guidance (TA510)

Both treatments are funded from the Cancer Drugs Fund (CDF). NHS England and NHS Improvement can confirm that the new SC Daratumumab product will be funded within the NHS provided it remains a cost neutral alternative to the IV formulation.

New patients may be initiated on the SC version as an alternative to the IV formulation following discussion between the individual patients and their clinician. Patients currently receiving IV daratumumab can be switched to SC daratumumab following the advice in the manufacturers SPC. Approved for addition to the formulary by Chair's action on 06/07/2020 - For noting

Outcome: Noted

Human Papillomavirus 9-Valent Vaccine (Gardasil 9®) Deferred to next meeting

Removals

Nil

NICE Approved drug applications

• TA628 - Lorlatinib for previously treated ALK-positive advanced non-small-cell lung cancer Approved by NICE in May 2020

Outcome: Approved for addition to the formulary

TA631 - Fremanezumab for preventing migraine

Approved by NICE in June 2020

Outcome: Approved for addition to the formulary

Pharmacoeconomic Board requests

• Nil

4.2 Trust Medicines Policy

- TMP: Section 21: Patient Group Directions
 - Full review and update undertaken
 - Addition of:
 - Approval process for PGDs used by services externally contracted by the Trust
 - Supply of medicines under PGD by posting or where the patient is absent

Outcome: Approved

Updates to Trust Adult IV Administration Guide

- Update to the Fosfomycin IV monograph to include reconstitution/dilution instructions as now diluting in IV fluid bag rather than hanging a pre-diluted glass bottle.
- New Monograph: Ocrelizumab (Ocrevus[®])

Outcome: Approved





4.3 Medicines Optimisation

• Trust Medicines Optimisation Annual Report 2019/20

Medicines optimisation encompasses a range of activities intended to improve the way that medicines are selected, procured, prescribed, dispensed and administered.

This report summarises the activities of groups responsible for the management of medicines at Chelsea and Westminster Hospital NHS Foundation Trust, describes developments throughout the 2019-20 year and reports on results of external assessments and peer reviews.

Action: Group members to review and forward any comments or suggestions for change to DR by 7th August 2020

4.4 NICE Technical Appraisals and Guidance

- a) NICE Technical Appraisals
- 1 Appraisal published in May 2020 since last meeting
- 8 Appraisals published in June 2020
- 2 Appraisals published in July 2020

TA626 - Avatrombopag for treating thrombocytopenia in people with chronic liver disease needing a planned invasive procedure

Recommendations

Formulary status / Action

Add to the formulary following receipt of a signed application form from the Hepatology Team.

TA630 - Larotrectinib for treating NTRK fusion-positive solid tumours

Formulary status / Action

Not applicable - CWFT not commissioned

Action: To be queried with NHS England as it would be expected that CWFT is a commissioned site.

TA631 - Fremanezumab for preventing migraine

Formulary status / Action

Added to the formulary following receipt of a signed application form from the Neurology Team - See Section 4.1

TA632 - Trastuzumab emtansine for adjuvant treatment of HER2-positive early breast cancer Formulary status / Action

Currently included on the formulary.

Numbers likely to treat at CWH site: 0 patients per year - Condition not treated at CW site

Numbers likely to treat at WMUH site: 4-6 patients per year

TA633 - Ustekinumab for treating moderately to severely active ulcerative colitis

Formulary status / Action

Currently included on the formulary.

Numbers likely to treat at CWH site: 15 patients per year Numbers likely to treat at WMUH site: 6 patients per year

TA634 - Daratumumab with lenalidomide and dexamethasone for untreated multiple myeloma

Formulary status / Action

Nil - Terminated appraisal

TA635 - Eculizumab for treating refractory myasthenia gravis (Terminated appraisal)

Formulary status / Action

Nil - Terminated appraisal





TA636 - Eculizumab for treating refractory myasthenia gravis (Terminated appraisal) Formulary status / Action Nil - Terminated appraisal

TA637 - Ranibizumab for treating diabetic retinopathy (Terminated appraisal) Formulary status / Action
Nil - Terminated appraisal

TA638 - Atezolizumab with carboplatin and etoposide for untreated extensive-stage small-cell lung cancer

Formulary status / Action

Currently included on the formulary.

Numbers likely to treat at CWH site: 5-6 patients per year

Numbers likely to treat at WMUH site: 0 patients per year - Condition not treated at WMUH site

TA639 - Atezolizumab with nab-paclitaxel for untreated PD-L1-positive, locally advanced or metastatic, triple-negative breast cancer

Formulary status / Action

Currently atezolizumab included on the formulary for other indication.

Numbers likely to treat at CWH site: 0 patients per year - Condition not treated at CW site

Numbers likely to treat at WMUH site: 3-5 patients per year

Add nab-paclitaxel to the formulary following receipt of a signed application form from the WM Oncology Team.

a) NICE Highly Specialised Technologies published since last meeting

0 Highly Specialised Technologies published

4.5 IVIG requests

IVIG Issues for May 2020 - WMUH Site

There were 12 IVIG issues in May 2020, with 8 new requests:

Outcome: Noted

IVIG Issues for June 2020 - CWH Site

There were 8 IVIG issues in June 2020, with 4 new requests:

Outcome: Noted

IVIG Issues for June 2020 - WMUH Site

Outcome: Noted

4.6 Items for noting

• Medication Safety Bulletin - National Patient Safety Review of Medication-related incidents Medication safety Bulletin relating to National Patient Safety Review of Medication - Related Incidents published July 2020

Outcome: Noted

Patient Group Directions Log - July 2020

Patient Group Directions Log for July 2020

Outcome: Noted

• NWL moving to single CCG





Letter from NWL providing an update on NWL moving to single CCG by April 2021

Outcome: Noted

MHRA Drug Safety Update - June 2020

MHRA update for June 2020

Outcome: Noted

4.7 Meeting minutes for noting

Nil

4.8 Additional papers to go to Trust Patient Safety Group

Trust Medicines Optimisation Annual Report 2019/20

5. Any other business

Nil

6. Date of next meeting Date: 28th September Time: 8am-9am Location: Via Zoom

Closing date: 11th September 2020