



Chelsea and Westminster Hospital NHS Foundation Trust Trust Medicines Group

Summary of Main Points from the Meeting held on Monday 10th July 2023

2. Minutes and Summary Notes from last meeting

This meeting was held via Teams. The Minutes and Summary notes of the Medicines Group Meeting held on 15th May 2023 were approved. 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 in due course.

3. Matters Arising

The Group noted the matters arising from the previous meeting.

4. Business to be transacted by the Medicines Group

4.1 Formulary Applications

Full Applications

- Nil

Ex-Panel Applications

- Tostran[®] Testosterone 2% Gel**

Requested by: GUM

Proposal: Add to the formulary in line with its licensed indication - Replacement therapy for male hypogonadism when testosterone deficiency has been confirmed by clinical features and biochemical tests.

Rational: Delivers half the testosterone dose as Testogel 16.2mg/g pump: Testogel 16.2mg/g gel pump delivers: 20mg in 1 pumps Tostran 2% gel pump delivers: 10mg in 1 pump This provides more flexibility with dosing where lower doses of testosterone are needed such as in cases of a suprathreshold serum testosterone levels or haematological toxicity, whilst maintaining an accurate dosing mechanism with the pump formulation.

Outcome: Approved for addition to the Trust Formulary

- Tolterodine 2mg SR Capsule**

Requested by: Urology

Proposal: Add to the formulary in line with its licensed indication - Symptomatic treatment of urge incontinence and/or increased urinary frequency and urgency as may occur in patients with Overactive Bladder Syndrome.

Rational: Tolterodine 4mg SR Capsules currently included on the formulary. Addition of 2mg capsules provides more flexibility with dosing.

Outcome: Approved for addition to the Trust Formulary

Removals

- Nil

NICE Approved drug applications

- TA877 Finerenone for treating chronic kidney disease in type 2 diabetes (23/03/2023) Proposal: Add to the formulary in line with NICE TA877

Outcome: Approved for addition to the Trust Formulary for use in line with NICE TA877

Pharmacoeconomic Board requests

- Bevacizumab - Refractory upper GI bleeding**

Approved by the Pharmacoeconomic Board on 24/05/2023

Outcome: Noted



- Infliximab for granulomatous disorder**
Approved by the Pharmacoeconomic Board on 30/05/2023
Outcome: Noted
- Infliximab for TB IRIS**
Approved by the Pharmacoeconomic Board on 02/06/2023
Outcome: Noted
- Infliximab for TB IRIS**
Approved by the Pharmacoeconomic Board on 06/06/2023
Outcome: Noted
- Tocilizumab for TB IRIS**
Approved by the Pharmacoeconomic Board on 03/07/2023
Outcome: Noted

4.2 Trust Medicines Policy

- TMP: Section 3 - Prescribing**

Update to the circumstances where a prescription may be issued to a patient at a hospital out-patient consultation in line with NWL prescribing guidance.

- Adult IV Administration Guide**

- ***Terlipressin***

Recent updates have suggested that there are less side effects with terlipressin given as an infusion compared to bolus doses. It is unlicensed but is detailed in the Medusa Monograph and other Hepatology centres are administering via infusion.

Outcome: Approved

- ***Infliximab***

Additional administration method has been added to the existing Infliximab monograph: In adult patients on maintenance therapy who have tolerated at least 3 initial 2-hour infusions (induction phase) consideration may be given to administering subsequent infusions over 1 hour.

Outcome: Approved

- ***Ferric Derisomaltose (Previously Monofer)***

The trade name removed from this brand of Iron as it is now to be referred to by the INN 'Ferric Derisomaltose'

Outcome: Approved

4.3 Medicines Optimisation

- Nil

4.4 NICE Technical Appraisals and Guidance

a) NICE Technology Appraisals

TA appraisals published since previous TMG meeting: 20

- TA886 Olaparib for adjuvant treatment of BRCA mutation-positive HER2-negative high-risk early breast cancer after chemotherapy (10/05/2023)**

Formulary status / Action

Not included on the formulary

Action: Add to the formulary following receipt of an application form from the Oncology Department



- TA887 Olaparib for previously treated BRCA mutation-positive hormone-relapsed metastatic prostate cancer (10/05/2023)**
Formulary status / Action
Not included on the formulary
Action: Add to the formulary following receipt of an application form from the Oncology Department

- TA888 Risankizumab for previously treated moderately to severely active Crohn's disease (17/05/2023)**
Formulary status / Action
Included on the formulary for another indication
Numbers likely to treat at CWH site: 5-10 patients per year
Numbers likely to treat at WMUH site: 5-10 patients per year

- TA889 Ciltacabtagene autoleucel for treating relapsed or refractory multiple myeloma (Terminated appraisal) (17/05/2023)**
Formulary status / Action
Action: Nil - Terminated Appraisal

- TA890 Difelikefalin for treating pruritus in people having haemodialysis (17/05/2023)**
Formulary status / Action
Not included on the formulary
Action: Nil - Not applicable as condition not treated at CWFT

- TA891 Ibrutinib with venetoclax for untreated chronic lymphocytic leukaemia (31/05/23)**
Formulary status / Action Included on the formulary for another indication
Numbers likely to treat at CWH site: 6 patients per year
Numbers likely to treat at WMUH site: 6 patients per year

- TA892 Mosunetuzumab for treating relapsed or refractory follicular lymphoma (31/05/23)**
Formulary status / Action
Action: Nil - Not recommended

- TA893 Brexucabtagene autoleucel for treating relapsed or refractory B-cell acute lymphoblastic leukaemia in people 26 years and over (07/06/2023)**
Not applicable to CWFT - Condition not treated at CWH and WMUH site

- TA894 Axicabtagene ciloleucel for treating relapsed or refractory follicular lymphoma (07/06/2023)**
Formulary status / Action
Action: Nil - Not recommended

- TA895 Axicabtagene ciloleucel for treating relapsed or refractory diffuse large B-cell lymphoma after first-line chemoimmunotherapy (07/06/2023)**
Not applicable to CWFT - Condition not treated at CWH and WMUH site

- TA896 Bulevirtide for treating chronic Hepatitis D (07/06/2023)**
Formulary status / Action
Not included on the formulary
Action: Nil - Not applicable as CWFT not a commissioned site

- TA897 Daratumumab with bortezomib and dexamethasone for previously treated multiple myeloma (06/06/2023)**
Formulary status / Action Included on the formulary for this indication
Numbers likely to treat at CWH site: 6 patients per year
Numbers likely to treat at WMUH site: 6 patients per year

- TA898 Dabrafenib plus trametinib for treating BRAF V600 mutation-positive advanced non-small-cell lung cancer (14/06/2023)**
Formulary status / Action Included on the formulary for another indication



Numbers likely to treat at CWH site: TBC
Numbers likely to treat at WMUH site: TB

TA899 Esketamine for treating major depressive disorder in adults at imminent risk of suicide (06/06/2023) Formulary status / Action
Action: Nil - Terminated Appraisal

TA900 Tixagevimab plus cilgavimab for preventing COVID-19 (14/06/2023)
Formulary status / Action
Action: Nil - Not recommended

TA901 Cemiplimab for treating recurrent or metastatic cervical cancer (20/06/2023) Formulary status / Action
Action: Nil - Terminated Appraisal

TA902 Dapagliflozin for treating chronic heart failure with preserved or mildly reduced ejection fraction (21/06/2023)
Formulary status / Action Included on the formulary for another indication
Numbers likely to treat at CWH site: TBC
Numbers likely to treat at WMUH site: 100 patients per year

TA903 Darolutamide with androgen deprivation therapy and docetaxel for treating hormone-sensitive metastatic prostate cancer (21/06/2023)
Formulary status / Action Included on the formulary for another indication
Numbers likely to treat at CWH site: 3 patients per year
Numbers likely to treat at WMUH site: 0 patients per year

TA904 Pembrolizumab with lenvatinib for previously treated advanced or recurrent endometrial cancer (21/06/2023)
Formulary status / Action Included on the formulary for another indication
Not applicable to CWFT - Condition not treated at CWH and WMUH site

TA905 Upadacitinib for previously treated moderately to severely active Crohn's disease (21/06/2023)
Formulary status / Action
Included on the formulary for another indication
Numbers likely to treat at CWH site: 20 patients per year
Numbers likely to treat at WMUH site: 5-10 patients per year

b) NICE Highly Specialised Technology Appraisals

HST appraisal published since previous TMGG meeting: 2

HST25 Lumasiran for treating primary hyperoxaluria type 1 (19/04/2023)
Action: Nil - Not applicable to CWFT

HST26 Eladocagene exuparvovec for treating aromatic L-amino acid decarboxylase deficiency (19/04/2023)
Action: Nil - Not applicable to CWFT

4.5 IVIG requests

IVIG Issues for April 2023 - CW Site
There were 11 IVIG issues in December 2022, with 4 new requests
Outcome: Noted



IVIG Issues for April 2023 - WMUH Site

There were 8 IVIG issues in March 2023, with 5 new requests

Outcome: Noted

IVIG issues for May 2023 - CWH Site

here were 16 IVIG issues in May 2023 with 7 new requests

Outcome: Noted

IVIG issues for May 2023 - WMH Site

There were 8 IVIG issues in March 2023, with 2 new requests

Outcome: Noted

4.6 Items for noting

Quarterly Controlled Drug Summary Report - Q4 2022/23

Quarterly Controlled Drug Summary Report for Q3 2022/23

Outcome: Noted

Quarterly Controlled Drugs Accountable Officer Report - Q4 2022/23

Quarterly CD Accountable Officer Report for Q3 2022/23

Outcome: Noted

Medication Safety Bulletin: Intravenous Iron

Medication Safety Bulletin relating to Intravenous Iron

Outcome: Noted

Medication Safety Bulletin: Intravenous Iron preparations

Medication Safety Bulletin: Reporting Intravenous iron Preparations

Outcome: Noted

MHRA Drug Safety Update - April 2023

MHRA update for April 2023

Outcome: Noted

Trust response to MHRA Risk Alerts relating to antimicrobials:

Nitrofurantoin - Pulmonary and Hepatic Toxicity

Aminoglycoside - Ototoxicity

Outcome: Plan agreed and actions noted

MHRA Drug Safety Update - May 2023

MHRA update for May 2023

Outcome: Noted

MHRA Drug Safety Update - June 2023

MHRA update for June 2023

Outcome: Noted

4.7 Meeting minutes for noting

Medication Safety Group meeting minutes - May 2023

Minutes from Medical Safety Group meeting held May 2023

Outcome: Noted

Antimicrobial Stewardship Group meeting minutes - May 2023

Minutes from Antimicrobial Stewardship Group meeting held May 2023

Outcome: Noted

5. Any other business



Nil

6. Date of next meeting

Next meeting

Date 25th September 2023

Time: 8am-9am

Location: via Teams