

Meeting to commence at 9.30 am on Tuesday 14th September 2021 via Zoom (Meeting ID: 885 2181 3733 Passcode: 5LDwp4)

AGENDA

| To protect commercial confidentiality the meeting will not open to the public until item 9 of the agenda | | | |
|---|--|------------------------------------|--|
| 1. | Welcome and introduction | Enclosure | |
| 2. | Apologies | | |
| 3. | Declarations of interest | | |
| 4. | Minutes of the previous meeting | 1/AWMSG/0921 | |
| 5. | Appraisal 1: Full (Wales Patient Access Scheme) Nitisinone (Orfadin [®]) for the treatment of adult patients with alkaptonuria | 2 /AWMSG/0921 Appendices | |
| 6. | Appraisal 2: Full Submission (Patient Access Scheme) Everolimus (Votubia [®]) as adjunctive treatment of patients aged 2 years and older whose refractory partial-onset seizures, with or without secondary generalisation, are associated with tuberous sclerosis complex (TSC) | 3 /AWMSG/0921 Appendices | |
| 7. | Appraisal 3: Full Submission (Wales Patient Access Scheme) Amikacin liposomal (Arikayce [®]) for the treatment of non-tuberculous mycobacterial (NTM) lung infections caused by Mycobacterium avium Complex (MAC) in adults with limited treatment options who do not have cystic fibrosis | 4 /AWMSG/0921 Appendices | |
| 8. | Appraisal 4: Limited Submission (Wales Patient Access Scheme) Dolutegravir (Tivicay [®]) for the treatment of HIV-1 infected adults, adolescents and children aged at least 4 weeks weighing at least 3 kg, in combination with other antiretrovirals | 5 /AWMSG/0921 Appendices | |
| | The meeting will open to the public | | |
| 9. | Appraisal 5: Paediatric licence extension submission Etravirine (Intelence [®]) in combination with a boosted protease inhibitor and other antiretroviral medicinal products, for the treatment of human immunodeficiency virus type 1 (HIV-1) infection in antiretroviral treatment-experienced paediatric patients from 2 to <6 years of age | 6 /AWMSG/0921 Appendices | |
| The outcome of the appraisals will be uploaded to the AWMSG website after the meeting. The applicant company will have ten working days to accept the decision or lodge a request for a review which should be sent to the AWMSG Chair via AWTTC. AWMSG's recommendation will be forwarded to Welsh Government for ratification following | | | |

Chair via AWTTC. AWMSG's recommendation will be forwarded to Welsh Government for ratification following acceptance of the outcome by the applicant company or after the ten-day deadline has expired – whichever is sooner. Welsh Government takes account of the AWMSG recommendation when making the final decision on the availability of the medicine in NHS Wales.



Welcome: Powys Teaching UHB

| 10. | Chairman's report | | |
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| 11. | NPI Quarterly report | 7/AWMSG/0921 | |
| 12. | AWMSG Five-Year Medicines Strategy 2018-2023 update report | 8/AWMSG/0921 | |
| 13. | Feedback from the AWPAG meeting held 30 th June 2021 | 9 /AWMSG/0921 | |
| 14. | Wales Advice on SGLT-2i Inhibitors in Type 2 Diabetes and Cardiovascular Disease | 10/AWMSG/0921 | |
| 15. | Tramadol Educational Resources (2021 review) | 11/AWMSG/0921 | |
| 16. | National Pharmacogenetics Service for Wales | 12/AWMSG/0921 | |
| 17. | Any other business | | |
| Date and time of next meeting: Wednesday 13 th October 2021 | | | |

