

| <b>somapacitan (Sogroya®)</b>   |  |   |                                      |   |
|---|--|---|--------------------------------------|---|
| <b>SMC Drug ID</b>  | <b>Conditions</b>  | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2629   | <p>for the replacement of endogenous growth hormone (GH) in children aged 3 years and above, and adolescents with growth failure due to growth hormone deficiency (paediatric GHD), and in adults with growth hormone deficiency (adult GHD).</p> <p>SMC restriction: for children aged 3 years and above and adolescents with growth failure due to growth hormone deficiency (paediatric GHD).</p> | Not routinely available as the ADTC is waiting for further advice from local clinical experts | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |  |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8726/somapacitan-sogroya-abbreviated-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8726/somapacitan-sogroya-abbreviated-final-oct-2024-for-website.pdf</a> |  |   |                                      |   |

| <b>linzagolix (Yselty®)</b>   |   |  |                                      |   |
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| SMC2631   | <p>the treatment of moderate to severe symptoms of uterine fibroids in adult women of reproductive age.</p> <p>SMC restriction: for use in patients when conventional first-line treatments (such as tranexamic acid, hormonal contraceptives and intrauterine devices) have failed or are considered unsuitable.</p> | Not routinely available as there is a local preference for alternative medicines | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |  |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8732/linzagolix-yselty-final-april-2024-issued-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8732/linzagolix-yselty-final-april-2024-issued-oct-2024-for-website.pdf</a> |   |  |                                      |   |
| <b>pembrolizumab (Keytruda®)</b>  |   |  |                                      |   |
| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>  | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2688   | in combination with platinum-containing chemotherapy as neoadjuvant treatment, and then continued as monotherapy as adjuvant treatment, for the treatment of resectable non-small cell lung carcinoma at high risk of recurrence in adults.   | Not routinely available as not recommended for use in NHS Scotland               | 18/11/2024                           | 16/10/2024  |
| <b>Other Decision Specified :</b> Economic analysis.  |   |  |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8733/pembrolizumab-keytruda-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8733/pembrolizumab-keytruda-final-oct-2024-for-website.pdf</a>                           |   |  |                                      |   |

| <b>axicabtagene cilocleucel (Yescarta®)</b>   |   |   |                                      |   |
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| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2695   | for the treatment of adult patients with diffuse large B-cell lymphoma (DLBCL) and high-grade B-cell lymphoma (HGBL) that relapses within 12 months from completion of, or is refractory to, first-line chemoimmunotherapy. | Not routinely available as the ADTC is waiting for further advice from local clinical experts | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8728/axicabtagene-yescarta-resubmission-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8728/axicabtagene-yescarta-resubmission-final-oct-2024-for-website.pdf</a> |   |   |                                      |   |
| <b>tenecteplase (Metalyse®)</b>   |   |   |                                      |   |
| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2697   | in adults for the thrombolytic treatment of acute ischaemic stroke within 4.5 hours from last known well and after exclusion of intracranial haemorrhage.   | Available in line with local or regional guidance   | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8727/tenecteplase-metalyse-abbreviated-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8727/tenecteplase-metalyse-abbreviated-final-oct-2024-for-website.pdf</a>   |   |   |                                      |   |

| <b>quizartinib (Vanflyta®)</b>  |   |   |                                      |   |
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| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2699   | in combination with standard cytarabine and anthracycline induction and standard cytarabine consolidation chemotherapy, followed by quizartinib single-agent maintenance therapy for adult patients with newly diagnosed acute myeloid leukaemia (AML) that is FLT3-ITD positive.                     | Not routinely available as the ADTC is waiting for further advice from local clinical experts | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8725/quizartinib-vanflyta-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8725/quizartinib-vanflyta-final-oct-2024-for-website.pdf</a>                                   |   |   |                                      |   |
| <b>bismuth subcitrate potassium/ metronidazole/ tetracycline (Pylera®)</b>  |   |   |                                      |   |
| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2701   | In combination with omeprazole, for the eradication of Helicobacter pylori and prevention of relapse of peptic ulcers in patients with active or a history of H. pylori associated ulcers. SMC restriction: restricted to use in accordance with clinical guidelines for the eradication of H. pylori | Available in line with local or regional guidance   | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8729/bismuth-subcitrate-pylera-abbreviated-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8729/bismuth-subcitrate-pylera-abbreviated-final-oct-2024-for-website.pdf</a> |   |   |                                      |   |

| <b>lebrikizumab (Ebglyss®)</b>  |   |   |                                      |   |
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| SMC2707   | <p>for the treatment of moderate-to-severe atopic dermatitis in adults and adolescents 12 years and older with a body weight of at least 40 kg who are candidates for systemic therapy.</p> <p>SMC restriction: patients who have had an inadequate response to an existing systemic immunosuppressant such as ciclosporin, or in whom such treatment is considered unsuitable and where a biologic would otherwise be offered.</p> | Not routinely available as the ADTC is waiting for further advice from local clinical experts | 11/11/2024                           | 03/01/2025  |
| <b>Other Decision Specified :</b>   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8731/lebrikizumab-ebglyss-final-oct-2024-updated-161024-for-website.pdf">https://scottishmedicines.org.uk/media/8731/lebrikizumab-ebglyss-final-oct-2024-updated-161024-for-website.pdf</a> |   |   |                                      |   |
| <b>enzalutamide (Xtandi®)</b>   |   |   |                                      |   |
| <b>SMC Drug ID</b>  | <b>Conditions</b>   | <b>Decision</b>   | <b>Date published on SMC Website</b> | <b>Date of decision / Expected date of decision</b> |
| SMC2742   | as monotherapy or in combination with androgen deprivation therapy for the treatment of adult men with high-risk biochemical recurrent (BCR) non-metastatic hormone-sensitive prostate cancer (nmHSPC) who are unsuitable for salvage radiotherapy.   | Not routinely available as not recommended for use in NHS Scotland                            | 11/11/2024                           | 16/10/2024  |
| <b>Other Decision Specified :</b> Non-Submission.   |   |   |                                      |   |
| <b>Web Link :</b> <a href="https://scottishmedicines.org.uk/media/8730/enzalutamide-xtandi-non-sub-final-oct-2024-for-website.pdf">https://scottishmedicines.org.uk/media/8730/enzalutamide-xtandi-non-sub-final-oct-2024-for-website.pdf</a>                 |   |   |                                      |   |