

**Subject: Specialised Services Circular (SSC)**  
**Sent on behalf of: Chair of the SW Specialised Service Circular Group**

Dear Colleagues,

Please find attached the following Specialised Services Circular(s):

<b>SSC Number</b>	<b>SSC Title</b>	<b>Trusts approved to prescribe in accordance with the SSC, providing appropriate internal governance arrangements are in place</b>
2730	<b>NICE Technology Appraisal Final Draft Guidance: Elranatamab for treating relapsed or refractory multiple myeloma after 3 therapies</b>	All South West Acute Trusts

**Is an implementation plan required from all SW trusts (regardless of commissioned status) for this SSC? No**

**For all other South West region trusts this is for information only.**

Trusts should ensure that use is registered on the Blueteq system (if appropriate).

Treatment will only be funded where the drugs minimum dataset is fully and accurately populated.

Please direct any queries to: [england.speccomm-southwest@nhs.net](mailto:england.speccomm-southwest@nhs.net)

**All Chief Executives**

**All Medical Directors**

**All Chief Pharmacists**

Specialised Commissioning  
South West  
NHS England  
100 Temple Street  
Bristol  
BS1 6AG  
Email: [england.speccomm-southwest@nhs.net](mailto:england.speccomm-southwest@nhs.net)

04 November 2024

Dear Colleagues,

**Re: NICE Technology Appraisal Final Draft Guidance: Elranatamab for treating relapsed or refractory multiple myeloma after 3 therapies.**

I am writing to advise you regarding the funding position on the recently published NICE Technology Appraisal Final Draft Guidance (FDG) for elranatamab for treating relapsed or refractory multiple myeloma after 3 therapies. This provider letter replaces SSC2670 published on 21<sup>st</sup> June 2024.

The FDG can be found at: <https://www.nice.org.uk/guidance/indevelopment/gid-ta10918>

NICE in their FDG published on 29<sup>th</sup> October 2024 has stated that:

Elranatamab is recommended with managed access as an option for treating relapsed and refractory multiple myeloma in adults, only after 3 or more lines of treatment (including an immunomodulatory drug, a proteasome inhibitor and an anti-CD38 antibody) when the myeloma has progressed on their last treatment.

Elranatamab has been available via the Cancer Drugs Fund (CDF) since 21<sup>st</sup> June 2024 in line with these recommendations and according to a set of treatment criteria which translates the NICE recommendation into a clinical guide as to use in practice. The treatment criteria have been updated to reflect the updated NICE FDG. These treatment criteria can be found on the national CDF list at <https://www.england.nhs.uk/cancer/cdf/cancer-drugs-fund-list/> or on the application form(s) on the Blueteq site.

Elranatamab will be available via the Cancer Drugs Fund (CDF) under the terms of the managed access agreement (MAA) and commercial access agreement (CAA), until NICE re-reviews these indications.

In addition:

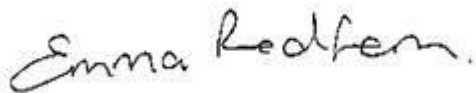
- Trusts must ensure that they are purchasing elranatamab at the agreed patient access scheme (PAS) discounted price. This discounted price will be applied

automatically at point of invoice and applies to all indications. Trusts should refer to the CAP portal for further information on the PAS price. The CAP portal is available at <https://nhsengland.sharefile.eu/Authentication/Login>

- Trusts must ensure that only invoices for the drug procurement costs of elranatamab in this indication are directed to the CDF and that they are also submitting complete and accurate information via the CDF minimum dataset (MDS).
- In line with the terms and conditions included in the NHS Standard Contract and as per the agreement that Cancer Services are commissioned with Trusts, Schedule 6a Reporting Requirements for drugs will apply. Payment of Trust invoices will be contingent on the completion of the MDS record and this information being made available in a timely way.
- Trusts must ensure they are registering elranatamab use on SACT. The SACT dataset is a mandated dataset as part of the Health and Social Care Information Standards. This is listed as a Schedule 6 national information requirement within the NHS Standard Contract.
- Patients must be registered via Blueteq (ELR1) and meet the clinical criteria on the registration form.
- **Payment of Trust invoices will be contingent on Blueteq registration, the full SACT and CDF MDS record applicable to the drug being completed and this information being made available in a timely way.**
- Trusts must ensure that local governance aspects (e.g. technical issues, education & training, patient information) have been identified and addressed for all staff groups (as appropriate) in order to permit the safe delivery of this therapy.

I would be grateful if you could cascade this information to relevant clinical teams within your organisation to support the consistent adoption of the policy nationally.

With best wishes,



Emma Redfern  
Medical Director NHS England South West  
Region



Tracey Williams  
Principal Pharmacist