NHS England South West

Subject: Specialised Services Circular (SSC 2641)
Sent on behalf of: Chair of the SW Specialised Service Circular Group

Dear Colleagues,

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

SSC Number	SSC Title	Trusts approved to prescribe in accordance with the SSC, providing appropriate internal governance arrangements are in place
2641	**NICE Technology Appraisal Final Draft Guidance: Selinexor with dexamethasone for treating relapsed or refractory multiple myeloma after 4 or more treatments**	All South West Acute Trusts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

NHS England South West

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12 April 2024

All Chief Executives

All Medical Directors

All Chief Pharmacists

Dear Colleagues,

Re: NICE Technology Appraisal Final Draft Guidance: Selinexor with dexamethasone for treating relapsed or refractory multiple myeloma after 4 or more treatments

I am writing to advise you regarding the funding position on the recently published NICE Technology Appraisal Final Draft Guidance (FDG) for selinexor with dexamethasone for treating relapsed or refractory multiple myeloma after 4 or more treatments.

The FDG can be found at: https://www.nice.org.uk/guidance/awaiting-development/gid-ta11223.

NICE in their FDG published on 9 April 2024 has stated that:

Selinexor plus dexamethasone is recommended, within its marketing authorisation, for treating multiple myeloma in adults when:

- they have had 4 or more treatments, and
- the condition is refractory to at least 2 proteasome inhibitors, 2 immunomodulatory agents and an anti-CD38 monoclonal antibody (penta-refractory), and
- the condition has progressed on the last treatment

Selinexor will be available via the Cancer Drugs Fund (CDF) from 9 April 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. 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.

NHS England will then routinely commission selinexor in patients with relapsed or refractory multiple myeloma, incorporating these treatment criteria, including those contained within this letter from 90 days after the day of publication of the final guidance.

In addition:

- Trusts must ensure that they are purchasing selinexor at the agreed proposed 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 any patients registered on the company led compassionate use programme who meet the clinical criteria for the drug are registered via Blueteq in order for the CDF to pick up the costs of their ongoing treatment. Patients who do not meet the clinical criteria should continue to receive drug via the manufacturer.
- Trusts must ensure that, until 90 days after publication of the final guidance from NICE, only invoices for the drug procurement costs of selinexor 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 selinexor 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 (SELIN2) and meet the clinical criteria on the registration form during the interim funding period.
- **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 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.

NHS England South West

Yours sincerely,

Luke Culverwell
Deputy Director of Specialised Commissioning

Tracey Williams
Principal Pharmacist