Clinical Commissioning Policy: Infliximab for Refractory or Progressive Neurosarcoidosis (Adults and Post-pubescent Children) [200501P]
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Commissioning Position

Summary

Infliximab will be available as a treatment option through routine commissioning for refractory or progressive neurosarcoidosis within the criteria set out in this document.

The policy is restricted to certain age groups as there is insufficient evidence to confirm safety and it is not recommended to be used in those age groups not included in the policy.

Executive Summary

Equality Statement

Promoting equality and addressing health inequalities are at the heart of NHS England’s values. Throughout the development of the policies and processes cited in this document, we have:

• Given due regard to the need to eliminate discrimination, harassment and victimisation, to advance equality of opportunity, and to foster good relations between people who share a relevant protected characteristic (as cited under the Equality Act 2010) and those who do not share it; and
• Given regard to the need to reduce inequalities between patients in access to, and outcomes from healthcare services and to ensure services are provided in an integrated way where this might reduce health inequalities

Plain Language Summary

About Neurosarcoidosis

Sarcoidosis can present in a variety of ways, ranging from a mild, acute self-limiting disease to chronic disease involving several organs and causing severe symptoms and functional impairment. It is characterised by the presence of lumps caused by clusters of inflammatory cells (non-caseating granulomas) in the organs. When the brain and central nervous system is affected, it is called neurosarcoidosis.

Neurosarcoidosis is uncommon but potentially serious manifestation of the disease, which occurs in approximately 5% of people with sarcoidosis. Approximately 100 people are diagnosed with neurosarcoidosis in the UK each year (Sarcoidosis UK, 2018).

While the cranial nerves are most frequently affected, neurosarcoidosis can involve other nervous system tissues including the meninges (the membranes enclosing the brain and spinal cord); brain parenchyma (the functional brain tissue), especially the hypothalamic region; spinal cord; peripheral nerve and muscle. There is a specific subgroup of neurosarcoidosis involving the brain parenchyma which can cause tissue destruction. This is the group of patients who are likely to benefit from infliximab, as it can prevent tissue destruction.

Neurosarcoidosis has no known cure. Spontaneous remission has been observed, but long-term therapy often is required. Treatment alleviates symptoms that are severe or progressive.
Immunosuppression is the principal method of controlling the disease, and corticosteroids are the cornerstone of therapy.

About Current Treatments

Most patients with sarcoidosis do not require treatment and often make a full recovery. Around a third have more serious disease involving different organs and require therapies such as steroids and drugs that suppress the immune system. The majority of patients with neurosarcoidosis require treatment of some type, which may include steroids and oral immunosuppressants. In most cases this is all that is required to put patients into remission. However, in the invasive parenchymal form of neurosarcoidosis, these treatments have been shown to be ineffective.

About the New Treatment

Infliximab belongs to a group of medicines called ‘biological drugs. It is a type of drug which works by reducing the effect of a chemical called tumour necrosis factor-alpha (TNF-α). TNF-α is released in response to a disease or infection as part of the body’s immune response. Infliximab is used to treat a number of medical conditions including Crohn’s disease, ulcerative colitis and certain types of arthritis such as rheumatoid arthritis and ankylosing spondylitis.

What We Have Decided

NHS England has carefully reviewed the evidence to treat refractory or progressive neurosarcoidosis with infliximab. We have concluded that there is enough evidence to make the treatment available at this time.

Committee Discussion

The Clinical Panel considered that the studies are of low quality and are not well structured but do show the quantum of benefit may be quite high. The treatment was supported provided the clinical body can identify those patients with rapidly progressive disease and apply strictly the stop criteria.

The Clinical Priorities Advisory Group considered the policy proposition and supporting documentation. See the committee papers (link) for full details of the evidence.

The Condition

Sarcoidosis is a systemic granulomatous disease of unknown cause characterised by multi-organ involvement. Almost 90% of patients have lung involvement but many of these will also show involvement of other organs, typically the skin and eyes (uveitis). Overall mortality in sarcoidosis is 1-5%, usually due to pulmonary, cardiac or neurological involvement or their complications (Sarcoidosis UK, 2018).

Neurological sarcoidosis is uncommon and occurs in approximately 5-10% of people with sarcoidosis. About 100 people are diagnosed with neurosarcoidosis affecting the brain and spinal cord in the UK each year (NICE Evidence Review, 2018). It can affect the central nervous system (CNS, the brain and spinal cord) and peripheral nervous system (including the 12 cranial nerves supplying the head and neck). When the nerves are affected there may be weakness, numbness of the face, problems with hearing, weakness of the tongue, difficulty swallowing, or double vision. Other patients have more serious disease affecting the CNS such as meningoencephalitis (inflammation of the lining of the brain and spinal cord) or vasculitis (inflammation of the blood vessels in the brain).

Current Treatments

Usually, oral corticosteroids are the first line therapy for patients with progressive disease or significant symptoms, with a maintenance dose given for 6-24 months. Other immunosuppressive or anti-inflammatory treatments are considered when corticosteroids are
failing to control disease, the side effects are intolerable, or corticosteroids are contraindicated (typically when patients also have diabetes mellitus and osteoporosis). Methotrexate or azathioprine are the most common second line treatments (Warrell et al. 2014).

In neurosarcoidosis, around 80% respond to 1st line and 2nd line oral treatment but around 20% need further treatment (NICE Evidence Review, 2018).

**New Treatment**

Infliximab has been used in the treatment of neurosarcoidosis that does not respond to first- or second-line treatments. Infliximab works by reducing the effect of a chemical called tumour necrosis factor-alpha (TNF-α), which is released in response to disease or infection as part of the body’s immune response. It is currently licensed as a treatment for other immune-mediated diseases such as rheumatoid arthritis and Crohn’s disease (Sarcoidosis UK, 2018). However, infliximab does not have a license or marketing authorisation for sarcoidosis.

A review of the current literature of infliximab in the treatment of neurosarcoidosis has been carried out and informs the development of this clinical commissioning policy.

**Epidemiology and Needs Assessment**

The cause of sarcoidosis is unknown. It is characterised by nodules of inflammation (non-caseating granulomas) and scarring in affected organs.

Around 1 in every 10,000 people in the UK has sarcoidosis. The incidence is highest in people aged 30 to 50 years and appears to be higher in Afro-Caribbean people and marginally higher in women (Sarcoidosis UK, 2018). Sarcoidosis in children is extremely rare. Many infectious and neoplastic diseases can simulate sarcoidosis.

The presentation of sarcoidosis varies considerably from mild, acute self-limiting disease to chronic disease involving several organs and causing severe symptoms and functional impairment. The most commonly affected organ is the lung, which is affected in more than 90% of people with sarcoidosis.

The natural course of sarcoidosis is difficult to predict and there are significant differences in the severity of disease and the organs involved. The prognosis is generally good, and sarcoidosis resolves in most people within two to five years.

It is estimated that 5-10% of cases have neurological involvement (Sarcoidosis UK, 2018). The NICE Evidence Review (2018) suggests about 100 people are diagnosed with neurosarcoidosis affecting the brain and spinal cord in the UK each year. Patients with neurosarcoidosis have a particularly poor prognosis and present with severe acute events, e.g. optic neuritis and blindness, acute hydrocephalus and coma or progressive lower limb weakness (Sarcoidosis UK, 2018). Around 80% respond to 1st and 2nd line oral treatment, but around 20% need further treatment.

Disease related mortality in sarcoidosis is reported to be about 5%, with the most common cause of death being from lung, cardiac and neurological disease that is refractory to therapy (Warrell et al. 2014).

**Evidence Summary**

NHS England has concluded that there is sufficient evidence to support a policy for the routine commissioning of this treatment for the indication.

This evidence summary considers the best available evidence for infliximab (alone or in combination with other medicines) for treating neurosarcoidosis that is refractory to standard
treatments (for example, corticosteroids and immunosuppressants) or is progressive despite current treatments.

Sarcoidosis is characterised by the presence of clusters of inflammatory cells in affected organs (non-caseating granulomas). Neurosarcoidosis is a rare type of sarcoidosis that affects the central nervous system (CNS, brain and spinal cord) causing meningoencephalitis and vasculitis.

The evidence review focused on 2 multicentre retrospective case reviews undertaken in the USA (Gelfand et al. 2017, n=66) and France (Jamilloux et al. 2017, n=132, 63 with neurosarcoidosis). In summary, the results of the studies suggest that infliximab can improve outcomes for some people with refractory or progressive neurosarcoidosis, with many people reported as experiencing a complete or partial response. However, some people had residual symptoms and functional impairment after treatment or experienced no response to treatment. Also, adverse effects of infliximab were sometimes serious and led to hospitalisation or discontinuation of treatment. Relapse of neurosarcoidosis appears to be common on stopping infliximab. See below for more details.

When assessing images obtained using magnetic resonance imaging (MRI), Gelfand et al. found that, in people using infliximab for 1.5 years on average, neurosarcoidosis resolved completely in 52% (29/56), partially improved in 30% (17/56), stayed the same in 14% (8/56) and got worse in 4% (2/56). When people were assessed by their specialist after using infliximab, Gelfand et al. found that clinical signs and symptoms of neurosarcoidosis resolved completely in 29% (19/66), partially improved in 48% (32/66), stayed the same in 18% (12/66) and got worse in 3% (2/66). A favourable response with complete or partial recovery was seen in 80% of people (45/56) who had evaluations for both clinical response and MRI findings. This suggests that 8 out of 10 people experienced some improvement in both findings on MRI and clinical signs and symptoms.

In the study by Gelfand et al., neurosarcoidosis recurred in 56% of people (9/16) who had experienced remission after using infliximab for, on average, 1.5 years. Relapse occurred, on average, about 6 months after treatment was stopped. These results suggest that around half of people with neurosarcoidosis experience relapse when they have been treated with infliximab for about 1.5 years.

Gelfand et al. found that combination therapy of infliximab with another immunosuppressant was associated with higher odds of a favourable treatment response (adjusted OR 6.9, 95% CI 1.2 to 41.3, p=0.03).

In the study by Jamilloux et al., a tool was used to assess the severity of sarcoidosis in various organs in the body. The average severity score for the CNS changed from 3.78 to 2.62 after treatment with an anti-TNF (usually infliximab) in 63 people with neurosarcoidosis (p=0.001). This shows that people’s score improved by 1.16 on a 6-point scale, which suggests that, on average, their neurosarcoidosis improved. However, although individual people may feel quite a large benefit, others may experience no benefit, and it is unclear if a 1 point improvement is large enough to be important to the overall population with neurosarcoidosis.

Overall, the studies suggest that, when infliximab is used for neurosarcoidosis, its adverse effects are similar to those that are seen when it is used for the licensed indications, as listed in the summary of product characteristics; for example, infections are common and may lead to hospitalisation or discontinuation of treatment.

The results of the studies should be interpreted with caution because the studies are small, uncontrolled, and did not use standardised treatment and monitoring protocols. Weaknesses in
the studies’ design and conduct mean they are subject to bias and confounding, are difficult to interpret and cannot support firm conclusions.

The optimal treatment regimen for infliximab for treating neurosarcoïdosis is unclear because treatment regimens varied within and between the studies. The dosage of infliximab typically used was 3 to 7 mg/kg at weeks 0, 2, and 6, then every 6 to 8 weeks. The median duration of treatment in the key studies was between about 1.5 years and 1.7 years. Most people in the studies took corticosteroids and/or immunosuppressants in combination with infliximab, and it is unclear how this affects outcomes. It is also unclear what potential factors predict a good response to Infliximab treatment.

**Implementation**

Patients must be managed at, or in collaboration with, a centre commissioned to provide specialised services that has expertise in the assessment and management of neurosarcoïdosis. Patients who satisfy the eligibility criteria will be eligible for treatment with infliximab. A decision to start using infliximab for neurosarcoïdosis should be made by a multidisciplinary team (MDT) at a specialist centre, using the following criteria:

**Criteria**

Patients must have a confirmed diagnosis of neurosarcoïdosis:

- Histological proof of systemic sarcoïdosis in patients who present with neurological impairment compatible with a diagnosis of neurosarcoïdosis, confirmed via MRI. OR
- Patients presenting with isolated neurosarcoïdosis, where there is no evidence for sarcoïdosis in the systemic organs and a biopsy of neurological tissue yields evidence of granulomatous inflammation and no evidence of any alternative diagnosis (such as infection, tumour or other inflammatory disease). OR
- The clinical presentation and diagnostic evaluation suggest neurosarcoïdosis, as defined by the clinical manifestations of MRI/Cerebrospinal Fluid (CSF) and/or Electromyography (EMG)/Nerve Conduction Studies (NCS) findings typical of granulomatous inflammation of the nervous system after rigorous exclusion of other causes.

All patients must have had an infectious or neoplastic disease cause excluded.

**Treatment Starting Criteria**

Infliximab will be added to a neurosarcoïdosis treatment regimen including high dose steroids and oral immunosuppressants in adult and post-pubescent patients with progressive and refractory disease who fulfil the following criteria:

- Patients with an abnormal MRI compatible with pachymeningitis or leptomenigitis (i.e. Imaging evidence of enhancement and inflammation of the meninges and adjacent brain and/spinal cord, cranial nerves and nerve roots within the spinal canal) with an examination of the cerebrospinal fluid which has ruled out a superimposed infection or neoplastic disease who:
  - Have severe, aggressive disease with risk of rapid, permanent and profound neurological impairment early in their disease;
  - AND
  - Have failed to respond to high dose steroids and other oral immunosuppressants;
  - OR
  - Are unable to be treated with high dose steroids and oral immunosuppressant agents due to severe intolerance or toxicity.
Exclusion Criteria
- History of tuberculosis (TB)
- Any evidence of concurrent TB, fungal, viral or parasitic infection; HIV, hepatitis, active herpes simplex virus (HSV), vesicular stomatitis virus (VZV), cytomegalovirus (CMV) or Epstein-Barr virus (EBV)
- Any evidence of current infection that may deteriorate with treatment
- History of demyelinating disease
- History of malignancy including malignant myeloma
- History of moderate or severe heart failure (this is NYHA class III/IV).

Dosing
The infliximab biologic with the lowest acquisition costs should be used. This is likely to be an infliximab biosimilar.

The recommended infliximab treatment dose regimen for adults and post-pubescent children with neurosarcoidosis is induction at 0, 2 and 6 weeks at a dose of 5mg/kg. Thereafter, it is given every 4-8 weeks at a dose of 5mg/kg. It is given in hospital by intravenous infusion and can be given as a day case.

All patients will undergo an MRI every three months or earlier if required. All patients will have monitoring of serum infliximab levels and antibody formation prior to every infusion and will undergo regular monitoring for any adverse events at each appointment.

Stopping Criteria
A decision to stop using infliximab for neurosarcoidosis should be made by a multidisciplinary team (MDT) at a specialist centre, using the following criteria:
- For patients who respond and achieve drug-induced disease remission achieving full resolution of enhancement on MRI, infusions of infliximab should then be reduced in frequency to zero over a period of 12-18 months.
- If there is no improvement seen on MRI within six months.
- If there are any adverse events where harm exceeds the benefit.

Criteria for Repeat Treatments
Re-treatment of such patients is permitted if the clinical assessment confirms that there is a clinical evidence of a flare, in addition to evidence of enhancement on an MRI.
Patient Pathway

- Diagnosis of neurosarcoidosis (as above)
- Subacute or progressive central neurological disorder e.g. encephalopathy, focal signs, seizures, cord or cauda equina lesion.

**Figure 1:** Patient pathway. MRI – magnetic resonance imaging. OD – once daily.

### Governance Arrangements

Any provider organisation treating patients with this intervention will be required to provide assurance that the internal governance arrangements have been completed before the medicine is prescribed. These arrangements may be through the Trust’s Drugs and Therapeutics Committee (or similar) and NHS England may ask for assurance of this process. Each provider organisation treating children with a medicine approved under the policy ‘Commissioning Medicines for Children in Specialised Services’ ([https://www.england.nhs.uk/wp-content/uploads/2017/03/commissioning-medicines-children-](https://www.england.nhs.uk/wp-content/uploads/2017/03/commissioning-medicines-children-))
specialised-services.pdf) will be required to assure itself that the internal governance arrangements have been completed before the medicine is prescribed. These arrangements may be through the Trust’s Drugs and Therapeutics Committee (or similar) and NHS England can ask for documented evidence that these processes are in place.

Provider organisations must register all patients using approved online prior approval software and ensure monitoring arrangements are in place to demonstrate compliance against the criteria as outlined.

**Mechanism for Funding**

Reimbursement for use of infliximab for patients with refractory or progressive neurosarcoidosis meeting the criteria in this policy is provided via specialised commissioning teams.

The cost will depend on the infliximab product used, the price of which is commercial in confidence. Only the infliximab product with the lowest acquisition cost will be reimbursed under this policy (likely to be a biosimilar).

**Audit Requirements**

Access to infliximab would be provided through specialist neurosarcoidosis networks with access to nationally recognised centres in this field.

Blueteq will be used to track and audit use of infliximab by clinicians, in order to ensure it is administered according to the Criteria for Commissioning.

**Policy Review Date**

This document will be reviewed when information is received which indicates that the policy requires revision. If a review is needed due to a new evidence base, then a new Preliminary Policy Proposal needs to be submitted by contacting england.CET@nhs.net

Our policies provide access on the basis that the prices of therapies will be at or below the prices and commercial terms submitted for consideration at the time evaluated. NHS England reserves the right to review policies where the supplier of an intervention is no longer willing to supply the treatment to the NHS at or below this price and to review policies where the supplier is unable or unwilling to match price reductions in alternative therapies.
Definitions

<table>
<thead>
<tr>
<th>Term</th>
<th>Description</th>
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<tbody>
<tr>
<td>Sarcoidosis</td>
<td>A systemic granulomatous disease of unknown cause characterised by multi-organ involvement.</td>
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<tr>
<td>Neurosarcoidosis</td>
<td>Sarcoidosis affecting the central nervous system including the brain and spinal cord.</td>
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<tr>
<td>Leptomeningitis</td>
<td>Inflammation of the tissues surrounding the brain or spinal cord.</td>
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<tr>
<td>Pachymeningitis</td>
<td>A rare illness which can be shown by MRI imaging to be a thickening of the tissue surrounding the brain and spinal cord when associated with an infectious, malignant or rheumatic systemic disease.</td>
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<tr>
<td>Refractory neurosarcoidosis</td>
<td>Neurosarcoidosis that has failed to respond to standard treatments (for example corticosteroids and immunosuppressants).</td>
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<tr>
<td>Progressive neurosarcoidosis</td>
<td>Sarcoidosis affecting the brain and spinal cord which is progressively worsening, despite first and second line treatments.</td>
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<tr>
<td>Infliximab</td>
<td>A drug which works by reducing the effect of a chemical called tumor necrosis factor-alpha (TNF-α). TNF-α is released in response to a disease or infection as part of the body’s immune response.</td>
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References

