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**N&WICS - TA352 - Vedolizumab - Treating moderately to severely active Crohn's disease after prior therapy**

Before providing patient identifiable data on this form, please confirm that the patient (or in the case of a minor or vulnerable adult with the parent/legal guardian/carer) has given appropriate explicit consent for sensitive personal information on this form to be passed to the CCG and/or CSU for processing this funding request and validating subsequent invoices. Consent given: ☐

If there is more than one NICE-approved treatment available, a discussion between the responsible clinician and the patient has taken place about the advantages and disadvantages of the treatments available. This has taken into consideration therapeutic need and whether or not the patient is likely to adhere to treatment. The most appropriate, least expensive, will be chosen (taking into account administration costs, dosage and price per dose) unless an order of preference is stated in the TAs. ☐

<b>Patient NHS No:</b>	<b>Trust:</b>	<b>Practice Name:</b>
<b>Patient Hospital No:</b> <input type="text"/>	<b>Consultant Making Request:</b> <input type="text"/>	<b>Practice Postcode:</b>
<b>Patient's Initials and DoB:</b>		<b>Practice Code:</b>
<b>Notification Email Address:</b> <input type="text"/> (@NHS.net account ONLY)		<b>Contact name &amp; number:</b> <input type="text"/>
<b>Start date of requested treatment:</b> <input type="text"/>	<b>Provider:</b> <input type="checkbox"/> Private <input type="checkbox"/> NHS <b>Supplier:</b> <input type="checkbox"/> Homecare <input type="checkbox"/> Hospital	<b>Sub-Type:</b> <input type="text"/> N/A <input type="button" value="v"/> (if applicable)

By completing this form, you confirm that you intend to use the requested medicinal product described below as agreed in the local commissioning statement. Any requests which fall outside of this use will require an individual funding request (IFR).

For support regarding IFRs, please contact: [norfolkicd@nhs.net](mailto:norfolkicd@nhs.net)

For support regarding the criteria listed below, please contact: [norfolknontariff@nhs.net](mailto:norfolknontariff@nhs.net)

<b>Please indicate whether patient meets the following NICE criteria:</b>	<b>Please tick</b>
1. Vedolizumab is recommended as an option for treating moderately to severely active Crohn's disease only if: <ul style="list-style-type: none"> <li>• a tumour necrosis factor-alpha inhibitor has failed (that is, the disease has responded inadequately or has lost response to treatment) or</li> <li>• a tumour necrosis factor-alpha inhibitor cannot be tolerated or is contraindicated. Vedolizumab is recommended only if the company provides it with the discount agreed in the patient access scheme.</li> </ul>	<input type="checkbox"/> Yes <input type="checkbox"/> No
2. Vedolizumab should be given as a planned course of treatment until it stops working or surgery is needed, or until 12 months after the start of treatment, whichever is shorter. At 12 months, people should be reassessed to determine whether treatment should continue. Treatment should only continue if there is clear evidence of ongoing clinical benefit. For people in complete remission at 12 months, consider stopping vedolizumab, resuming treatment if there is a relapse. People who continue vedolizumab should be reassessed at least every 12 months to decide whether continued treatment is justified.	<input type="checkbox"/> Yes <input type="checkbox"/> No
3. As per NICE guidance, treatment should normally be started with the least expensive drug (considering the dose required, price per dose and any additional administration costs).  Please confirm that the choice of drug is considered the most cost-effective treatment for this individual patient, noting that alternative treatment options recommended by NICE may be lower in cost, including the use of biosimilar medications if applicable.  Where an alternative lower cost treatment is available and has not been tried by the patient, please confirm that the clinical rationale for prescribing has been included in the patients' medical record for the purpose of auditing.	<input type="checkbox"/> Yes <input type="checkbox"/> No
4. The product is being used as described by local commissioning position.	<input type="checkbox"/> Yes <input type="checkbox"/> No
<b>I confirm that the patient meets the criteria for treatment</b>	

Name of person completing:	<input type="text"/>	Contact Details:	<input type="text"/>
Designation of person completing:	<input type="text"/>	Date:	<input type="text"/>
Trust Authorising Pharmacist			
Name: <input type="text"/>			
Date: <input type="text"/>			