Click here to access the guidelines/NICE algorithm N&WICS - TA187 - Infliximab - Active fistulising Crohn's disease 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. Patient **Practice** Trust: NHS No: Name: **Patient** Consultant **Practice** Hospital Making Postcode: No. Request: Patient's **Practice** Initials and Code: DoB: Notification Contact Email (@NHS.net account ONLY) name & Address: number: Provider: Start date ☐ Private ☐ NHS of Supplier: Sub-Type: ☐ Homecare ☐ Hospital requested treatment: (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 For support regarding the criteria listed below, please contact: norfolknontariff@nhs.net Please indicate whether patient meets the following NICE criteria: Please tick 1. Infliximab, within its licensed indication, is recommended as a treatment option for people with active fistulising Crohn's disease: • whose disease has not responded to conventional therapy (including antibiotics, drainage and immunosuppressive treatments), or who are intolerant of or have contraindications to conventional therapy. ☐Yes ☐No • Infliximab should be given as a planned course of treatment until treatment failure (including the need for surgery) or until 12 months after the start of treatment, whichever is shorter. • People should then have their disease reassessed (see Q4) to determine whether ongoing treatment is still clinically appropriate. 2. Treatment with Infliximab should only be continued if there is clear evidence of ongoing active disease as determined by clinical symptoms, biological markers and investigation, including endoscopy if necessary. • Specialists should discuss the risks and benefits of continued treatment with patients and consider a trial withdrawal from treatment for all patients who are in stable clinical remission. □Yes □No • People who continue treatment with infliximab or adalimumab should have their disease reassessed at least every 12 months to determine whether ongoing treatment is still clinically appropriate. • People whose disease relapses after treatment is stopped should have the option to start treatment again. 3. When using the CDAI and Harvey-Bradshaw Index, healthcare professionals should take into account any physical, sensory or learning disabilities, or communication difficulties that could affect the scores and ☐Yes ☐No make any adjustments they consider appropriate. 4. Treatment with infliximab should only be started and reviewed by clinicians with experience of TNF ☐ Yes ☐ No inhibitors and of managing Crohn's disease. 5. 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.		□Yes □No
6. The product is being used as described by local commissioning position.		□Yes □No
I confirm that the patient meets the criteria for treatment		
Name of person completing:	Contact Details:	
Designation of person completing:	Date:	
Trust Authorising Pharmacist	<u> </u>	