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N&WICS - TA466 - Baricitinib with methotrexate - Moderate to severe rheumatoid arthritis 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. Baricitinib, with methotrexate, is recommended as an option for treating active rheumatoid arthritis in adults whose disease has responded inadequately to intensive therapy with a combination of conventional ☐Yes ☐No disease-modifying antirheumatic drugs (DMARDs), only if: 2. Baricitinib, with methotrexate, is recommended as an option for treating active rheumatoid arthritis in ☐ Yes ☐ No adults whose disease has responded inadequately to or who cannot have other DMARDs, including at least 1 biological DMARD, only if: 3. Baricitinib can be used as monotherapy for people who cannot take methotrexate because it is ☐ Yes ☐ No contraindicated or because of intolerance, when the criteria in sections 1.1 or 1.2 are met. 4. Continue treatment only if there is a moderate response measured using European League Against Rheumatism (EULAR) criteria at 6 months after starting therapy. After an initial response within 6 months, ☐ Yes ☐ No withdraw treatment if at least a moderate EULAR response is not maintained. 5. When using the DAS28, healthcare professionals should take into account any physical, psychological, sensory or learning disabilities, or communication difficulties that could affect the responses to the DAS28 ☐Yes ☐No and make any adjustments they consider appropriate. I confirm that the patient meets the criteria for treatment Name of person completing: Contact Details: Designation of person completing: Date: Trust Authorising Pharmacist Name: Date: