Click here to access the guidelines/NICE algorithm N&WICS - TA210 - Modified-release dipyridamole - Vascular 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 Sub-Type: Supplier: ☐ Homecare ☐ Hospital requested N/A 🗸 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. Clopidogrel is recommended as an option to prevent occlusive vascular events: • for people who have had an ischaemic stroke or who have peripheral arterial disease or multivascular ☐ Yes ☐ No disease or • for people who have had a myocardial infarction only if aspirin is contraindicated or not tolerated. 2. Modified-release dipyridamole in combination with aspirin is recommended as an option to prevent occlusive vascular events: □Yes □No • for people who have had a transient ischaemic attack or • for people who have had an ischaemic stroke only if clopidogrel is contraindicated or not tolerated.

| * for people who have had an ischaemic stroke or who have peripheral arterial disease or multivascular disease or
| * for people who have had a myocardial infarction only if aspirin is contraindicated or not tolerated.
| 2. Modified-release dipyridamole in combination with aspirin is recommended as an option to prevent occlusive vascular events:
| * for people who have had a transient ischaemic attack or
| * for people who have had an ischaemic stroke only if clopidogrel is contraindicated or not tolerated.
| 3. Modified-release dipyridamole alone is recommended as an option to prevent occlusive vascular events:
| * for people who have had an ischaemic stroke only if aspirin and clopidogrel are contraindicated or not tolerated or
| * for people who have had a transient ischaemic attack only if aspirin is contraindicated or not tolerated.
| 4. Treatment with clopidogrel to prevent occlusive vascular events should be started with the least costly licensed preparation.
| * Geople currently receiving clopidogrel or modified-release dipyridamole either with or without aspirin outside the criteria in 1.1, 1.2 and 1.3 should have the option to continue treatment until they and their clinicians consider it appropriate to stop.
| * Geople currently receiving clopidogrel or modified potential properties of the option to continue treatment until they and their clinicians consider it appropriate to stop.
| * Geople currently received per dose and any additional administration costs).
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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		□Yes □No
auditing.		
7. 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	1	
Name:		
Date:		