

South East London Integrated Medicines Optimisation Committee Formulary recommendation

Reference	147
Intervention:	Colchicine for the secondary prevention of ischaemic heart disease in adults
	(Colchicine is a plant alkaloid with anti-inflammatory properties)
Date of Decision	October 2023
Date of Issue:	November 2023 (Original time limited approval for 12 months. Extended in
	April 2025 to April 2026 to enable further collection of data for outcomes report)
Recommendation:	
Further Information	 Cardiovascular disease (CVD) describes a range of conditions that affect the heart and blood vessels caused by the process of atherosclerosis. Ischemic heart disease falls into this category. Colchicine tablets are accepted for use in South East London for the secondary prevention of ischaemic heart disease (including after acute myocardial infarction and in chronic coronary disease) in high risk patients. Colchicine may only be considered in this setting if the following criteria are met: Use is for the secondary prevention of CVD disease AND Established treatments recommended by NICE* have been optimised (for example: ACE inhibitors, beta blockers, statins) AND Use is in high risk patients, defined as: Previous spontaneous acute myocardial infarction (diagnosed according to the universal MI criteria) with or without persistent ST-segment elevation OR Previous stroke or intervention for peripheral arterial disease (i.e., evidence of atherosclerotic disease affecting >1 vascular bed) OR Established diagnosis of diabetes mellitus OR Systemic Coronary Risk Estimation 2 (SCORE2) or Systemic Coronary Risk Estimation 2 (SCORE2-OP) algorithm 10-year risk of fatal and non-fatal myocardial infarction or stroke >10%
	 specialist. In line with clinical trials describing the use of colchicine in this setting, colchicine is recommended at a dose of 500 micrograms once daily. Colchicine is not licensed* for use in this indication (off-label use). Informed consent should be gained from the patient before treatment is initiated. Treatment with colchicine in this setting would be continued long-term if tolerated. When making a decision to initiate colchicine, consideration should also be given to the polypharmacy aspects and the risks associated with this for individual patients.
	 Patients should also be advised to stop colchicine and seek advice from their GP or CVD specialist in the following scenarios: If gastrointestinal symptoms occur (e.g. nausea, vomiting, abdominal pain, diarrhoea) despite a low or lactose free diet and / or There are symptoms suggestive of infection (e.g. fever, stomatitis, sore throat, prolonged bleeding, bruising or skin disorders) If after assessment by the GP it is deemed that the only cause is colchicine, then colchicine should be discontinued by the GP and the initiating CVD specialist informed of the discontinuation due to adverse effects. Colchicine has a narrow therapeutic index and is extremely toxic in overdose. Refer to the <u>summary of product characteristics</u> for colchicine for further information. This approval was orginally time limited to 12 months to enable experience of use. The time limit was extended in April 2025 to April 2026 at the

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	applicant's request. , A report summarising outcomes with colchicine over this period will be presented back to the committee by April 2026. This report will be coordinated across all Trusts in SEL by the original formulary applicant and will include:
	- The total number of patients initiated on colchicine (including the proportion
	from SEL) for the secondary prevention of ischaemic heart disease
	 Whether the use of colchicine is in line with this formulary recommendation and the rationale for any deviation.
	 Reporting on patient related outcomes, including:
	(i) Adverse effects reported and their themes
	(ii) Number of patients discontinuing treatment due to adverse effects
<u> </u>	* NICE guidance includes: NICE guideline <u>185</u> (Acute Coronary Syndrome), NICE Guideline <u>181</u> (CVD)
Shared Care/ Transfer of	
care required:	N/A
Cost Impact for	The application estimates that ~ 3,850 patients in SEL per annum will be eligible
agreed	for treatment with colchicine in this setting. This is a broader patient cohort than
patient group	agreed in the "Further Information" section.
	• The cost of colchicne is around £9.60 per anum per patient.
	• This would equate to £37,000 (or ~£1,850 per 100,000 population per year) for
	the broader patient cohort.
Usage Monitoring &	
Impact Assessment	 Monitor and audit usage and outcomes from use of colchicine in this setting as
	outlined in the "For information" section and report back the Committee by
	April 2026. SEL Borough Medicines Teams:
	 Monitor exception reports from GPs if inappropriate prescribing requests are
	made to primary care
Evidence reviewed	References (from evidence review)
	1. NICE NG185: Acute coronary syndromes. (November 2020)
	 NICE CG181: Cardiovascular disease: risk assessment and reduction, including lipid modification (February 2023) NICE CKS: Lipid modification – CVD prevention (January 2023)
	 Visseren, F.L. et al. (2021) "2021 ESC guidelines on Cardiovascular Disease Prevention in Clinical Practice," European Heart Journal, 42(34), pp. 3227–3337.
	 Colchicine - summary of product characteristics (SmPC) - (EMC). Available <u>here</u> [Accessed 14/02/2023]
	6. Shah B, Allen N, Harchandani B, Pillinger M, Katz S, Sedlis SP, Echagarruga C, Samuels SK, Morina P, Singh P, et al.
	Effect of colchicine on platelet-platelet and platelet-leukocyte interactions: a pilot study in healthy subjects. Inflammation. 2016; 39:182–189.
	 Mastrocola R, Penna C, Tullio F, Femminò S, Nigro D, Chiazza F, Serpe L, Collotta D, Alloatti G, Cocco M, et al. Pharmacological inhibition of NLRP3 inflammasome attenuates myocardial ischemia/reperfusion injury by activation of RISK and mitochondrial pathways.Oxid Med Cell Longev. 2016; 2016;5271251.
	8. Diaz-Arocutipa, C. et al. (2021) "Efficacy and safety of Colchicine in post–acute myocardial infarction patients: A systematic review and meta-analysis of randomized controlled trials," Frontiers in Cardiovascular Medicine, 8.
	 Hemkens, L.G. et al. (2016) "Colchicine for prevention of cardiovascular events," Cochrane Database of Systematic Reviews [Preprint].
	 Tardif, JC. et al. (2019) "Efficacy and safety of low-dose colchicine after myocardial infarction," New England Journal of Medicine, 381(26), pp. 2497–2505.
	 Nidorf, S.M. et al. (2020) "Colchicine in patients with chronic coronary disease," New England Journal of Medicine, 383(19), pp. 1838–1847.
	12. Opstal, T.S.J. et al. (2022) 'Long-term efficacy of colchicine in patients with chronic coronary disease: Insights from Iodoco2', Circulation, 145(8), pp. 626–628.
	 Samuel, M. et al. (2020) "Cost-effectiveness of low-dose colchicine after myocardial infarction in the Colchicine Cardiovascular Outcomes Trial (colcot)," European Heart Journal - Quality of Care and Clinical Outcomes, 7(5), pp. 486–495
	14. European Society of Cardiology Guidelines for the management of acute coronary syndromes, August 2023. Available <u>here</u> , last accessed 18.10.23 (not in evidence review)

NOTES:

- a) SEL IMOC recommendations and minutes are available publicly via the website.
- b) This SEL IMOC recommendation has been made on the cost effectiveness, patient outcome and safety data available at the time. The recommendation will be subject to review if new data becomes available, costs are higher than expected or new NICE guidelines or technology appraisals are issued.
- c) Not to be used for commercial or marketing purposes. Strictly for use within the NHS.

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