



EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

19 September 2019
EMA/CHMP/35552/2019
Committee for Medicinal Products for Human Use (CHMP)

Colchicine tablet 0.5 mg and 1 mg product-specific bioequivalence guidance

Draft Agreed by Pharmacokinetics Working Party (PKWP)	January 2019
Adopted by CHMP for release for consultation	28 February 2019
Start of public consultation	8 March 2019
End of consultation (deadline for comments)	30 June 2019
Draft Agreed by Pharmacokinetics Working Party (PKWP)	September 2019
Adopted by CHMP	19 September 2019
Date of coming into effect	1 April 2020

Keywords	<i>Bioequivalence, generics, colchicine</i>
-----------------	--

Official address Domenico Scarlattilaan 6 • 1083 HS Amsterdam • The Netherlands

Address for visits and deliveries Refer to www.ema.europa.eu/how-to-find-us

Send us a question Go to www.ema.europa.eu/contact **Telephone** +31 (0)88 781 6000

An agency of the European Union



Colchicine tablet 0.5 mg and 1 mg product-specific bioequivalence guidance

Disclaimer:

This guidance should not be understood as being legally enforceable and is without prejudice to the need to ensure that the data submitted in support of a marketing authorisation application complies with the appropriate scientific, regulatory and legal requirements.

Requirements for bioequivalence demonstration (PKWP)*

BCS Classification**	BCS Class: <input type="checkbox"/> I <input checked="" type="checkbox"/> III <input type="checkbox"/> Neither of the two Background: Colchicine is considered a high solubility compound with limited absorption.
Bioequivalence study design <i>in case a BCS biowaiver is not feasible or applied</i>	single dose
	cross-over
	healthy volunteers
	<input checked="" type="checkbox"/> fasting <input type="checkbox"/> fed <input type="checkbox"/> both <input type="checkbox"/> either fasting or fed
	Strength: 1 mg Background: Highest strength recommended. However, it is also possible to use the lower strength for a drug with linear pharmacokinetics and high solubility.

	Number of studies: One
Analyte	<input checked="" type="checkbox"/> parent <input type="checkbox"/> metabolite <input type="checkbox"/> both
	<input checked="" type="checkbox"/> plasma/serum <input type="checkbox"/> blood <input type="checkbox"/> urine
	Enantioselective analytical method: <input type="checkbox"/> yes <input checked="" type="checkbox"/> no
Bioequivalence assessment	Main pharmacokinetic variables: C_{max} , AUC_{0-t} Background/justification:
	90% confidence interval: 80.00– 125.00% for C_{max} and 90.00-111.11% for AUC_{0-t} . Background: Colchicine is a narrow therapeutic index drug.

* As intra-subject variability of the reference product has not been reviewed to elaborate this product-specific bioequivalence guideline, it is not possible to recommend at this stage the use of a replicate design to demonstrate high intra-subject variability and widen the acceptance range of C_{max} . If high intra-individual variability ($CV_{intra} > 30\%$) is expected, the applicants might follow respective guideline recommendations.

** Applying for a BCS-based biowaiver is restricted to highly soluble drug substances with known human absorption and considered not to have a narrow therapeutic index (NTI). As colchicine is considered a NTI drug, a BCS biowaiver is not possible.