

25 June 2025 EMA/148455/2025 Human Medicines Division

Overview of (invented) names reviewed in June 2025 by the Name Review Group (NRG)

Adopted at the CHMP meeting of 24 July 2025

	NRG meeting 11-12 February		NRG meeting 8-9 April		NRG meeting 11-12 June		NRG meeting 23-24 September		NRG meeting 18-19 November		2025 total	
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Proposed (invented) names	47	38	56	35	31	33					134	106
Proposed INN+MAH/TM names	1	0	1	0	2	0					4	0
Re-use/re-confirmation applications	2	0	0	0	3	1					5	1
Justification for retention of (invented) name *	3	3	4	0	1	5					8	8

^{*}In case of objections to the proposed (invented) name(s), the applicant may justify the retention of the proposed invented name using the relevant justification form available on the EMA website.



	NRG meeting 11-12 February		NRG meeting 8-9 April		NRG meeting 11-12 June		NRG meeting 23-24 September		NRG meeting 18-19 November		2025 total	
	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed
Total number of objections raised	78	163	65	139	72	139					215	441
Similarity with other (invented) name	49	143	48	113	46	113					95	256
Conveys misleading therapeutic connotations	4	2	0	0	1	1					5	3
Conveys misleading pharmaceutical connotations	0	0	0	0	0	0					0	0
Similarity with INN	5	8	9	11	15	12					29	31
Inclusion of INN stem	1	1	0	0	0	0					1	1
Unacceptable qualifiers	1	1	0	0	0	0					1	1
Conveys a promotional message	8	4	1	8	4	3					13	15
Appears offensive or has an inappropriate connotation	0	0	4	1	3	0					7	1
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations	0	0	0	0	0	0					0	o
Similarity between name of prodrug and related active substance	0	0	0	0	0	0					0	0
Inclusion of common umbrella segment	0	0	0	0	0	0					0	0
Potential difficulties in pronunciation	6	1	3	1	2	3					11	5
Others	0	0	0	0	2	2					2	2

See <u>Guideline on the acceptability of names for human medicinal products processed through the centralised procedure (EMA/CHMP/287710/2014 - Rev. 7)</u> for detailed explanations of criteria used.