

16 July 2024 EMA/302580/2024 Human Medicines Division

Overview of (invented) names reviewed in June 2024 by the Name Review Group (NRG)

Adopted at the CHMP meeting of 22-25 July 2024

	NRG meeting 13-14 Feb 2024		NRG meeting 16 April 2024		NRG meeting 18 June 2024		NRG meeting 24-25 September 2024		NRG meeting 19-20 November 2024		2024 total	
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Proposed (invented) names*	69	35	43	38	44	31					156	104
Justification for retention of (invented) name **	4	4	0	1	2	4					6	9

*Includes invented names, INN+MAH/TM and re-use applications.

**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.

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	NRG meeting 13-14 Feb 2024		NRG meeting 16 April 2024		NRG meeting 18 June 2024		NRG meeting 24-25 September 2024		NRG meeting 19-20 November 2024		2024 total	
	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed	Endorsed	Not endorsed
Total number of objections raised	71	154	79	153	60	112					210	419
Similarity with other (invented) name	57	120	65	117	47	132					169	369
Conveys misleading therapeutic connotations	1	5	0	1	1	0					2	6
Conveys misleading pharmaceutical connotations	1	1	2	2	0	1					3	4
Similarity with INN	7	8	3	4	4	8					14	20
Inclusion of INN stem	0	0	1	2	0	0					1	2
Unacceptable qualifiers	0	0	1	1	0	1					1	2
Conveys a promotional message	0	6	1	3	0	1					1	10
Appears offensive or has an inappropriate connotation	0	2	2	0	1	0					3	2
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations	0	0	0	0	0	0					0	0
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	0	5	1	2	5	4					6	11
Others	5	7	1	4	3	2					9	13

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.