

29 May 2024 EMA/217318/2024 Human Medicines Division

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

Adopted at the CHMP meeting of 27-30 May 2024

	NRG meeting 13-14 Feb 2024		NRG meeting 16 April 2024		NRG meeting 18-19 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							112	73
Justification for retention of (invented) name **	8	4	0	1							8	5

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

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<sup>\*\*</sup>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 13-14 Feb 2024		NRG meeting 16 April 2024		NRG meeting 18-19 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	71	84							142	238
Similarity with other (invented) name	57	120	58	66							115	186
Conveys misleading therapeutic connotations	1	5	0	1							1	6
Conveys misleading pharmaceutical connotations	1	1	2	2							3	3
Misleading with respect to composition	0	0	0	0							0	0
Similarity with INN	7	8	3	4							10	12
Inclusion of INN stem	0	0	1	2							1	2
Unacceptable qualifiers	0	0	1	1							1	1
Conveys a promotional message	0	6	1	3							1	9
Appears offensive or has an inappropriate connotation	0	2	2	0							2	2
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations	0	0	0	0							0	0
Similarity between name of prodrug and related active substance	0	0	0	0							0	0
Inclusion of common umbrella segment	0	0	0	0							0	0
Potential difficulties in pronunciation	0	5	1	2							1	7
Others	5	7	1	4							6	11

See Guideline on the <u>Acceptability of Names for Human Medicinal Products Processed through the Centralised Procedure (EMA/CHMP/287710/2014)</u> for detailed explanations of criteria used.