

18 December 2014 EMA/790636/2014 Human Medicines Evaluation

Overview of (invented) names reviewed in November 2014 by the Name Review Group (NRG)

Adopted at the CHMP meeting of 15 - 18 December 2014

	NRG m	eeting	NRG m	eeting	NRG m	eeting	NRG me	eting	NRG m	eeting	NRG m	eeting	2014	
	29 Jan	2014	26 Ma	r 2014	15 Ma	y 2014	02 July	2014	01 Oct	2014	26 Nov	2014		
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Proposed invented names	35	37	47	41	54	32	61	54	48	37	77	52	322	253
Justification for retention of invented name *	1	6	0	5	1	9	2	4	3	0	3	4	10	28

^{*}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		NRG meeting		NRG meeting		NRG meeting		NRG meeting		NRG meeting		2014	
	29 Jan	2014	26 Mar	2014	15 Ma	y 2014	02 Jul	y 2014	01 Oct 2	2013	26 No	ov 2014		
Objections	Accepted	Rejected	Accepted	Rejected										
Total number of objections raised	76	45	82	53	47	38	114	65	65	28	101	39	485	268
Criterion - Safety concerns														
Similarity with other Invented name	62	36	72	46	42	28	96	53	52	22	90	38	414	223
Conveys misleading therapeutic/pharmaceutical connotations		1		1		1	1					1	1	4
Misleading with respect to composition						2			1		1		2	2
Criterion - INN concerns														
Similarity with INN	4	1	4	2	2	1	9	3	7	4	2		28	11
Inclusion of INN stem	3		4	3		1	3				3		13	4
Criterion - Other public health concerns														
Unacceptable qualifiers	2					1					1		3	1
Conveys a promotional message	2	4				1	5	4	1				8	9
Appears offensive or has a bad connotation	1	1			1				1				3	1
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations														
Similarity between name of prodrug and related active substance														
Others	2	2	2	1	2	3		5	3	2	4		13	13

See Guideline on the Acceptability of Names for Human Medicinal Products Processed through the Centralised Procedure (CPMP/328/98 Rev. 5) for detailed explanations of criteria used.