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Overview of invented names reviewed in September 2011 by the Name Review Group (NRG)

Adopted at the CHMP meeting of 17-20 October 2011

	NRG meeting 25 Jan 2011		22 March		NRG meeting 24 May 2011		NRG meeting 28 June 2011		NRG meeting 22 Sept 2011		NRG meeting 17 Nov 2011		2011	
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejecto	Accepted	Rejecte	Accepted	Rejected
Proposed invented names	34	68	56	50	59	41	56	55	49	41			250	255
Justification for retention of invented name *	0	2	4	6	2	4	4	0	1	2			11	14

^{*}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 EMEA website.

	NRG meeting 25 Jan 2011		NRG meeting 22 March 2011		NRG meeting 24 May 2011		NRG meeting 28 June 2011		NRG meeting 22 Sept 2011		NRG meeting 17 Nov 2011		2011	
Objections	Accepted	Rejected	Accepted	Rejecte	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Total number of objections raised	155	90	102	91	94	63	95	41	75	67			521	352
Criterion - Safety concerns														
Similarity with other Invented name	125	73	82	74	48	53	87	26	68	51			342	227
Conveys misleading therapeutic/pharmaceutical connotations	2	3	0	1	1	0	1	1	1	3			5	8
Misleading with respect to composition	3	1	5	5	6	0	0	0	0	2			14	8
Criterion - INN concerns														
Similarity with INN	8	6	5	2	4	1	3	0	2	1			22	10
Inclusion of INN stem	5	3	3	2	3	0	1	3	0	8			12	16

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Criterion - Other public health concerns														
Unacceptable qualifiers	1	0	3	2	2	0	0	1	0	0			6	3
Conveys a promotional message	1	0	2	5	0	8	3	1	3	2			9	16
Appears offensive or has a bad connotation	1	0	2	0	1	0	0	0	0	1			4	1
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations	0	0	0	0	2	1	0	0	1	0			3	1
Similarity between name of prodrug and related active substance	0	0	0	0	0	0	0	0	0	0			0	0

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.