

27 June 2013 EMA/383378/2013

## Overview of invented names reviewed in June 2013 by the Name Review Group (NRG)

Adopted at the CHMP meeting of 24 – 27 June 2013

	NRG meeting 30 Jan 2013		NRG meeting 10 April 2013		NRG meeting 11 June 2013		NRG meeting 4 July 2013		NRG meeting 02 Oct 2013		NRG meeting 14 Nov 2013		2013	
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Proposed invented names	42	33	44	38	33	35								
Justification for retention of invented name *	-	2	2	2	-	2								

\*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.

7 Westferry Circus • Canary Wharf • London E14 4HB • United Kingdom

Telephone +44 (0)20 7418 8427 Facsimile +44 (0)20 7418 8409

E-mail press@ema.europa.eu Website www.ema.europa.eu



	NRG meeting NRG mee 30 Jan 2013 10 Ap 2013		April			NRG meeting 4 July 2013		NRG meeting 02 Oct 2013		NRG meeting 14 Nov 2013		2013		
Objections	Accepted	Rejected	Accepted	0	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Total number of objections raised	56	54	80	40	67	78								
Criterion - Safety concerns														
Similarity with other Invented name	49	39	63	34	59	75								
Conveys misleading therapeutic/pharmaceutical connotations		2		1	1									
Misleading with respect to composition		1	1											
Criterion - INN concerns														
Similarity with INN		8	1	1	1									
Inclusion of INN stem	3		1	2		1								
Criterion - Other public health concerns														
Unacceptable qualifiers	1	1												
Conveys a promotional message	1	3				1								
Appears offensive or has a bad connotation	2		1											
Similarity between name of individual active substance and fixed combinations and/or between fixed combinations			13	2	3	1								
Similarity between name of prodrug and related active substance														
Others					3									

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.