

25 April 2014 EMA/218433/2014

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

Adopted at the CHMP meeting of 22 - 25 April 2014

	NRG meeting 29 Jan 2014		NRG meeting 26 Mar 2014		NRG meeting 15 May 2014						NRG meeting 26 Nov 2014			
	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Proposed invented names	35	37	47	41										
Justification for retention of invented name *	1	6	0	5										

\*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.

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 29 Jan 2014		NRG meeting 26 Mar 2014		NRG meeting 15 May 2014		NRG meeting 02 July 2014		NRG meeting 01 Oct 2013		NRG meeting 26 Nov 2014		2014	
Objections	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected	Accepted	Rejected
Total number of objections raised	76	45	82	53										
Criterion - Safety concerns														
Similarity with other Invented name	62	36	72	46										
Conveys misleading therapeutic/pharmaceutical connotations		1		1										
Misleading with respect to composition														
Criterion - INN concerns														
Similarity with INN	4	1	4	2										
Inclusion of INN stem	3		4	3										
Criterion - Other public health concerns														
Unacceptable qualifiers	2													
Conveys a promotional message	2	4												
Appears offensive or has a bad connotation	1	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										

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.