



BUCCOLAM

Procedural steps taken and scientific information after the authorisation

Application number	Scope	Opinion/ Notification ¹ issued on	Commission Decision Issued ² / amended on	Product Information affected ³	Summary
IA/0039	B.II.b.2.a - Change to importer, batch release arrangements and quality control testing of the FP - Replacement/addition of a site where batch control/testing takes place	03/08/2018	n/a		
N/0038	Minor change in labelling or package leaflet not connected with the SPC (Art. 61.3 Notification)	25/07/2018		Labelling	

¹ Notifications are issued for type I variations and Article 61(3) notifications (unless part of a group including a type II variation or extension application or a worksharing application). Opinions are issued for all other procedures.

² A Commission decision (CD) is issued for procedures that affect the terms of the marketing authorisation (e.g. summary of product characteristics, annex II, labelling, package leaflet). The CD is issued within two months of the opinion for variations falling under the scope of Article 23.1a(a) of Regulation (EU) No. 712/2012, or within one year for other procedures.

³ SmPC (Summary of Product Characteristics), Annex II, Labelling, PL (Package Leaflet).



IB/0037	B.II.b.5.f - Change to in-process tests or limits applied during the manufacture of the finished product - Addition or replacement of an in-process test as a result of a safety or quality issue	24/05/2018	n/a		
IAIN/0036	B.II.b.2.c.1 - Change to importer, batch release arrangements and quality control testing of the FP - Replacement or addition of a manufacturer responsible for importation and/or batch release - Not including batch control/testing	02/02/2018		Annex II and PL	
PSUSA/10118 /201609	Periodic Safety Update EU Single assessment - midazolam (oromucosal solution, treatment of prolonged, acute, convulsive seizures)	06/04/2017	n/a		PRAC Recommendation - maintenance
R/0032	Renewal of the marketing authorisation.	01/04/2016	26/05/2016	SmPC, Annex II, Labelling and PL	
IA/0033	A.5.b - Administrative change - Change in the name and/or address of a manufacturer/importer of the finished product, including quality control sites (excluding manufacturer for batch release)	02/05/2016	n/a		
PSUSA/10118 /201509	Periodic Safety Update EU Single assessment - midazolam (oromucosal solution, treatment of prolonged, acute, convulsive seizures)	17/03/2016	n/a		PRAC Recommendation - maintenance
IG/0603/G	This was an application for a group of variations. A.1 - Administrative change - Change in the name and/or address of the MAH	03/12/2015	26/05/2016	SmPC, Annex II, Labelling and PL	

	A.5.a - Administrative change - Change in the name and/or address of a manufacturer/importer responsible for batch release				
IG/0621	C.I.8.a - Introduction of or changes to a summary of Pharmacovigilance system - Changes in QPPV (including contact details) and/or changes in the PSMF location	16/10/2015	n/a		
IA/0028	B.II.b.3.a - Change in the manufacturing process of the finished or intermediate product - Minor change in the manufacturing process	09/10/2015	n/a		
IA/0027	B.II.b.2.a - Change to importer, batch release arrangements and quality control testing of the FP - Replacement/addition of a site where batch control/testing takes place	16/07/2015	n/a		
PSUSA/10118 /201409	Periodic Safety Update EU Single assessment - midazolam (oromucosal solution, treatment of prolonged, acute, convulsive seizures)	12/03/2015	n/a		PRAC Recommendation - maintenance
IB/0025/G	This was an application for a group of variations. A.5.b - Administrative change - Change in the name and/or address of a manufacturer/importer of the finished product, including quality control sites (excluding manufacturer for batch release) A.7 - Administrative change - Deletion of manufacturing sites	29/10/2014	n/a		

IA/0024	B.II.b.3.a - Change in the manufacturing process of the finished or intermediate product - Minor change in the manufacturing process	26/09/2014	n/a		
IAIN/0023	C.I.8.a - Introduction of or changes to a summary of Pharmacovigilance system - Changes in QPPV (including contact details) and/or changes in the PSMF location	02/07/2014	n/a		
PSUV/0012	Periodic Safety Update	10/04/2014	n/a		PRAC Recommendation - maintenance
IB/0020	B.II.b.1.e - Replacement or addition of a manufacturing site for the FP - Site where any manufacturing operation(s) take place, except batch-release, batch control, primary and secondary packaging, for non-sterile medicinal products	28/03/2014	n/a		
IB/0021/G	This was an application for a group of variations. B.II.e.2.c - Change in the specification parameters and/or limits of the immediate packaging of the finished product - Deletion of a non-significant specification parameter (e.g. deletion of an obsolete parameter) B.II.e.2.z - Change in the specification parameters and/or limits of the immediate packaging of the finished product - Other variation	26/03/2014	n/a		
IB/0019	Update of section 6.6 of the SmPC to include information on how to administer Buccolam as already introduced in the Package Leaflet. In addition several editorial corrections including an update of the entire	20/03/2014	05/03/2015	SmPC, Annex II, Labelling and PL	

	<p>PI in accordance with QRD template version 9 have also been introduced.</p> <p>C.I.z - Changes (Safety/Efficacy) of Human and Veterinary Medicinal Products - Other variation</p>				
IA/0022	A.7 - Administrative change - Deletion of manufacturing sites	27/02/2014	05/03/2015	Annex II and PL	
IAIN/0018	B.II.b.1.b - Replacement or addition of a manufacturing site for the FP - Primary packaging site	17/02/2014	n/a		
IAIN/0016/G	<p>This was an application for a group of variations.</p> <p>C.I.9.a - Changes to an existing pharmacovigilance system as described in the DDPS - Change in the QPPV and/or QPPV contact details and/or back-up procedure</p> <p>C.I.9.b - Changes to an existing pharmacovigilance system as described in the DDPS - Change(s) in the safety database and/or major contractual arrangements for the fulfilment of PhV obligations, and/or change of the site undergoing PhV activities</p> <p>C.I.9.c - Changes to an existing pharmacovigilance system as described in the DDPS - Other change(s) to the DDPS that does not impact on the operation of the PhV system</p>	17/02/2014	n/a		
IA/0017	B.II.b.3.a - Change in the manufacturing process of the finished or intermediate product - Minor change in the manufacturing process	12/02/2014	n/a		

IB/0015	B.II.e.1.z - Change in immediate packaging of the finished product - Other variation	11/02/2014	n/a		
IB/0014	B.II.b.4.z - Change in the batch size (including batch size ranges) of the finished product - Other variation	31/01/2014	n/a		
N/0011	Minor change in labelling or package leaflet not connected with the SPC (Art. 61.3 Notification)	17/01/2014	05/03/2015	Labelling and PL	The MAH applied for the introduction of a wrap-around tube label with improved visual images and a multi-strength Package Leaflet with the introduction of colour to differentiate between the strengths for optimum clarity. The labelling and PL have been updated accordingly.
IA/0013	B.II.b.4.a - Change in the batch size (including batch size ranges) of the finished product - Up to 10-fold compared to the originally approved batch size	19/12/2013	n/a		
IB/0008	B.II.b.5.b - Change to in-process tests or limits applied during the manufacture of the finished product - Addition of a new test(s) and limits	07/11/2013	n/a		
IAIN/0010	B.II.b.1.a - Replacement or addition of a manufacturing site for the FP - Secondary packaging site	06/11/2013	n/a		
IAIN/0009	B.II.b.1.a - Replacement or addition of a manufacturing site for the FP - Secondary packaging site	24/10/2013	n/a		
IA/0007/G	This was an application for a group of variations. B.III.1.a.2 - Submission of a new or updated Ph. Eur. Certificate of Suitability to the relevant Ph. Eur.	14/08/2013	n/a		

	<p>Monograph - Updated certificate from an already approved manufacturer</p> <p>B.III.1.a.2 - Submission of a new or updated Ph. Eur. Certificate of Suitability to the relevant Ph. Eur. Monograph - Updated certificate from an already approved manufacturer</p>				
IA/0006	<p>B.III.1.a.2 - Submission of a new or updated Ph. Eur. Certificate of Suitability to the relevant Ph. Eur. Monograph - Updated certificate from an already approved manufacturer</p>	17/06/2013	n/a		
IAIN/0005/G	<p>This was an application for a group of variations.</p> <p>C.I.9.a - Changes to an existing pharmacovigilance system as described in the DDPS - Change in the QPPV</p> <p>C.I.9.c - Changes to an existing pharmacovigilance system as described in the DDPS - Change of the back-up procedure of the QPPV</p> <p>C.I.9.e - Changes to an existing pharmacovigilance system as described in the DDPS - Changes in the major contractual arrangements with other persons or organisations involved in the fulfilment of pharmacovigilance obligations and described in the DD</p> <p>C.I.9.h - Changes to an existing pharmacovigilance system as described in the DDPS - Other change(s) to the DDPS that does not impact on the operation of the pharmacovigilance system</p>	02/05/2013	n/a		
IAIN/0004/G	<p>This was an application for a group of variations.</p>	13/12/2012	18/12/2013	SmPC, Annex II	

	<p>A.6 - Administrative change - Change in ATC Code/ATC Vet Code</p> <p>B.II.b.2.b.1 - Change to batch release arrangements and quality control testing of the FP - Not including batch control/testing</p>			and PL	
IAIN/0003/G	<p>This was an application for a group of variations.</p> <p>C.I.9.e - Changes to an existing pharmacovigilance system as described in the DDPS - Changes in the major contractual arrangements with other persons or organisations involved in the fulfilment of pharmacovigilance obligations and described in the DD</p> <p>C.I.9.h - Changes to an existing pharmacovigilance system as described in the DDPS - Other change(s) to the DDPS that does not impact on the operation of the pharmacovigilance system</p>	22/08/2012	n/a		
IB/0002	<p>B.II.b.5.a - Change to in-process tests or limits applied during the manufacture of the finished product - Tightening of in-process limits</p>	16/08/2012	n/a		
IB/0001	<p>B.II.e.2.b - Change in the specification parameters and/or limits of the immediate packaging of the finished product - Addition of a new specification parameter to the specification with its corresponding test method</p>	16/12/2011	n/a		