State of California
Secretary of State

This Certificate is not valid for use anywhere within the United States of America, its territories or possessions.

APOSTILLE
(Convention de La Haye du 5 octobre 1961)

1. Country: Pays / País:	United States of America
This public document Le présent acte public / El presente documento público	
2. has been signed by a été signé par ha sido firmado por	*J. Cranston*
3. acting in the capacity of agissant en qualité de quien actúa en calidad de	*Notary Public, State of California*
4. bears the seal / stamp of est revêtu du sceau / timbre de y está revestido del sello / timbre de	*J. Cranston , Notary Public, State of California*

Certified
Attesté / Certificado

5. at à / en	Sacramento, California	**6. the** le / el día	*7th day of June 2018*
7. by par / por	Secretary of State, State of California		
8. N° sous n° bajo el número	*72986*		
9. Seal / stamp: Sceau / timbre: Sello / timbre:		**10. Signature:** Signature: Firma:	

This Apostille only certifies the authenticity of the signature and the capacity of the person who has signed the public document, and, where appropriate, the identity of the seal or stamp which the public document bears.
This Apostille does not certify the content of the document for which it was issued.
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Cette Apostille atteste uniquement la véracité de la signature, la qualité en laquelle le signataire de l'acte a agi et, le cas échéant, l'identité du sceau ou timbre dont cet acte public est revêtu.
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Esta Apostilla certifica únicamente la autenticidad de la firma, la calidad en que el signatario del documento haya actuado y, en su caso, la identidad del sello o timbre del que el documento público esté revestido.
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Este certificado no constituye una Apostilla en virtud del Convenio de La Haya de 5 de octubre de 1961 cuando se presenta en un país que no es parte del Convenio. En estos casos, el certificado debe ser presentado a la sección consular de la misión que representa a ese país.

Allergan

5 June 2018

To Whom It May Concern:

I, Meredith Sewell, certify that after sufficient due diligence and to the best of my knowledge, the attached document is a true and correct copy of the Certificate of GMP Compliance of a Manufacturer, certificate number: 18363/M11938, issued on 2018-01-23.

Respectfully,

Meredith Sewell
Executive Director
Regulatory Submissions

A notary public or other officer completing this certificate verifies only the identity of the individual who signed the document to which this certificate is attached, and not the truthfulness, accuracy, or validity of that document.

State of California)
County of Orange)

On June 5, 2018 before me, J. Cranston, Notary Public, personally appeared Meredith Sewell, who proved to me on the basis of satisfactory evidence to be the person(s) whose name(s) is/are subscribed to the within instrument and acknowledged to me that he/she/they executed the same in his/her/their authorized capacity(ies), and that by his/her/their signature(s) on the instrument the person(s), or the entity upon behalf of which the person(s) acted, executed the instrument.

I certify under PENALTY OF PERJURY under the laws of the State of California that the foregoing paragraph is true and correct.

WITNESS my hand and official seal.

Signature J Cranston

J. CRANSTON
Commission # 2134121
Notary Public - California
Orange County
My Comm. Expires Dec 16, 2019

2525 Dupont Drive • Irvine, CA • 92612
Tel: 714-246-4500

HPRA
An tÚdarás Rialála Táirgí Sláinte
Health Products Regulatory Authority

Health Products Regulatory Authority

CERTIFICATE NUMBER: *18363/M11938*

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER [1, 2]

Part 1

Issued following an inspection in accordance with :
Art. 111(5) of Directive 2001/83/EC as amended

The competent authority of Ireland confirms the following:
The manufacturer: ***Allergan Sales LLC***
Site address: ***8301 Mars Drive, Waco, Texas, 76712, United States***

Has been inspected in connection with marketing authorisation(s) listing manufacturers located outside of the European Economic Area in accordance with Art. 111(4) of Directive 2001/83/EC transposed in the following national legislation:
Medicinal Products (Control of Manufacture) Regulations 2007 to 2013.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on ***2017-10-13*** , it is considered that it complies with :

- The principles and guidelines of Good Manufacturing Practice laid down in Directive 2003/94/EC [3]

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection. However, this period of validity may be reduced or extended using regulatory risk management principles by an entry in the Restrictions or Clarifying remarks field. This certificate is valid only when presented with all pages and both Parts 1 and 2. The authenticity of this certificate may be verified in EudraGMDP. If it does not appear, please contact the issuing authority.

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[1] *The certificate referred to in paragraph 111(5) of Directive 2001/83/EC and 80(5) of Directive 2001/82/EC, shall also be required for imports coming from third countries into a Member State.*

[2] *Guidance on the interpretation of this template can be found in the Help menu of EudraGMDP database.*

[3] *These requirements fulfil the GMP recommendations of WHO.*

An tÚdarás Rialála Táirgí Sláinte, Teach Kevin O'Malley, Ionad Phort an Iarla, Ardán Phort an Iarla, Baile Átha Cliath 2, Éire
Health Products Regulatory Authority, Kevin O'Malley House, Earlsfort Centre, Earlsfort Terrace, Dublin 2, Ireland
T: +353 1 676 4971 • F: +353 1 676 7836 • info@hpra.ie • www.hpra.ie

Part 2

Human Medicinal Products

1 MANUFACTURING OPERATIONS	
1.1	**Sterile products**
	1.1.1 Aseptically prepared (processing operations for the following dosage forms) 1.1.1.3 Semi-solids 1.1.1.4 Small volume liquids
1.2	**Non-sterile products**
	1.2.1 Non-sterile products (processing operations for the following dosage forms) 1.2.1.11 Semi-solids
1.5	**Packaging**
	1.5.1 Primary Packing 1.5.1.11 Semi-solids
	1.5.2 Secondary packing
1.6	**Quality control testing**
	1.6.1 Microbiological: sterility *1.6.2 Microbiological: non-sterility* *1.6.3 Chemical/Physical*

2018-01-23

Name and signature of the authorised person of the Competent Authority of Ireland

CERTIFIED HPRA

Dearbhla Cullen

Ms. Dearbhla Cullen
Health Products Regulatory Authority
Tel: *+35 31676 4971*
Fax: *+35 31676 7836*