



Your medicines and health products,
our concern

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Your letter from	Your reference	Our reference	Annex	Date
		AFMPS/DGPRE/R&D/LSA 978.119		03 MARS 2017

Onderwerp
Titre de l'objet
Subject

Goedkeuring van een klinische proef op 13/02/2017
Approbation d'un essai clinique le 13/02/2017
Authorisation of a clinical trial dated 13/02/2017

Trial Title: Dossier OGM : B/BE/16/BVW1: A Phase 1, Multicenter, Open-label Trial to Evaluate the Safety of Talimogene Laherpaprepvec Injected into Liver Tumors

EudraCT: 2014-005386-67

Chère Madame, Cher Monsieur,

Conformément à l'article 12 de la Loi du 7 mai 2004 relative aux expérimentations sur la personne humaine, j'ai décidé d'autoriser l'essai clinique ci-dessus mentionné.

Salutations sincères,

Pour la Ministre des Affaires sociales et de la Santé publique

Geachte Mevrouw, Geachte Heer,

In overeenstemming met artikel 12 van de wet van 7 mei 2004 inzake experimenten op de menselijke persoon, heb ik besloten de hierboven vermelde proef goed te keuren.

Met de meeste hoogachting,

Voor de Minister van Sociale Zaken en Volksgezondheid

Dr. Greet Musch

Unofficial translation

In accordance with article 12 of the Law of 7 May 2004 concerning experiments on the human person, I have decided to authorise the above mentioned clinical trial.

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Votre lettre du	Vos références	Nos références	Annexe(s)	Date
		AFMPS/DGP/RE/R&D/LSA 963195		13 FEV. 2017

Dossier OGM : B/BE/16/BVW1 (2014-005386-67): A Phase 1, Multicenter, Open-label Trial to Evaluate the Safety of Talimogene Laherparepvec Injected into Liver Tumors

Geachte Mevr. Houben,

Hierbij informeren wij u dat de vergunning krachtens het Koninklijk besluit van 21 februari 2005 tot reglementering van de doelbewuste introductie in het leefmilieu evenals van het in de handel brengen van genetisch gemodificeerde organismen of van producten die er bevatten u wordt toegekend op basis van het gunstig advies van de Adviesraad voor Bioveiligheid daterend van 13 december 2016 en dit volgens de voorwaarden hernomen in de conclusie van bovenvermeld advies, dat wil zeggen:

- The notifier and the investigators must strictly apply the trial protocol, and all the safety instructions as described in the dossier.
- Any protocol amendment has to be previously approved by the Competent Authority.
- The notifier is responsible to verify that each study center has qualified personnel experienced in handling infectious material and that the investigator has the required authorisations to perform the clinical trial activities inside the hospital (laboratory, pharmacy, hospital room, consultation room, ...) according to the Regional Decrees transposing Directive 2009/41/EC on the contained use of genetically modified micro-organisms.
- For the transport of the IMP the notifier should conform to the transportation rules regarding transport of GMO's.
- The Biosafety Advisory Council should be informed within 2 weeks when the first patient starts the treatment and the last patient receives the last treatment.

- At the latest six months after the last visit of the last patient included in the trial, the notifier must send to the competent authority for the attention of the Biosafety Advisory Council a report with details concerning the biosafety aspects of the project. This report shall contain at least:
 - o The total number of patients included in the trial and the number of patients included in Belgium;
 - o A summary of all adverse events marked by the investigators as probably or definitely related to the study medication;
 - o A report on the accidental releases, if any, of the recombinant HSV-1.

Met hoogachting,

Maggie De Block
Ministre des Affaires Sociales
et de la Santé publique



Marie Christine Marghem
Ministre de l'Energie, de
l'Environnement et du
Développement durable

