



DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration  
Silver Spring MD 20993

DMF 030678

## DMF ACKNOWLEDGEMENT

SILILABEL S.A.  
ATTENTION: ENG. NICOLAS GERZENSTEIN  
PTEFRONDIZI 2501, PILAR  
PROVINCIA DE BUENOS AIRES, ARGENTINA

Dear Eng. Nicolas Gerzenstein,

The Food and Drug Administration acknowledges receipt of the following Drug Master File (DMF) submission:

**DMF NUMBER ASSIGNED:** 030678  
**DATE OF SUBMISSION:** AUGUST 14, 2016  
**DMF TYPE:** III  
**SUBJECT (TITLE):** BLISTER MATERIALS  
**HOLDER:** SILILABEL S.A.  
**SUBMITTED BY:** SILILABEL S.A.

All subsequent correspondence to this DMF should be identified with the information as provided above.

Your DMF will be reviewed only in connection with a New Drug Application, Abbreviated New Drug Application, Investigational New Drug Application, Biological License Application, New Animal Drug Application, Abbreviated New Animal Drug Application, Investigational New Animal Drug Application, or DMF it is intended to support when a Letter of Authorization (LOA) is submitted to the DMF and a copy of the LOA is submitted in the application e.g., NDA, that references the DMF.

You are responsible for compliance with 21 CFR 314.420. See "The Guideline for Drug Master Files" <http://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionRequirements/DrugMasterFilesDMFs/ucm073164.htm>

You are required to submit all changes in information regarding the DMF (21 CFR 314.420(c)). In particular the FDA must be notified of any changes in the holder name or ownership and/or the agent and/or the name of the contact person.

The types of information to be submitted may be found at the DMF Web Site. See "**Submission of Amendments, Annual Reports, and Letters of Authorization.**"

You are expected to:

- Adhere to the statement of commitment you have provided.
- Provide the following submissions to the DMF:

Reference ID: 3973021

- a. Letters of Authorization (LOAs) granting permission to a third party (authorized party) to reference the DMF and for FDA to review the DMF. Listing an authorized party in the Annual Report (see below) is not sufficient to authorize that party to reference the DMF. Submission of a copy of the LOA to the authorized party without submitting the original LOA to the DMF is also not sufficient to authorize that party to reference the DMF.
- b. Annual Reports to the DMF containing:
  - i. Date(s) of the amendment(s) reporting changes since the last Annual Report or the original DMF filing date, whichever is most recent or a statement that no amendments have been submitted since the last Annual Report or the original DMF filing date, whichever is most recent.
  - ii. A complete list of all parties authorized to make reference to the DMF, identifying by name, reference number, volume, date, and page number the information that each person is authorized to incorporate and the date of the LOA or a statement that there are no Authorized Parties.
  - iii. A list of all parties whose authorization has been withdrawn, if applicable.

Submissions containing multiple types of information e.g. administrative changes, an annual report, or changes in technical information should specify the different types of information in the header in the cover letter.

Electronic submissions that are 10GB or smaller in size must be submitted through the Electronic Submission Gateway (ESG). Submissions that are over 10GB may be submitted on physical media (such as compact disc)<sup>1</sup> to the following address:

Food and Drug Administration  
Center for Drug Evaluation and Research  
Central Document Room  
Drug Master File Staff  
5901-B Ammendale Road  
Beltsville MD 20705-1266

If you have any questions, please email [dmfquestion@cder.fda.gov](mailto:dmfquestion@cder.fda.gov)

Sincerely,  
*{See appended electronic signature page}*  
Vathsala Selvam  
Drug Master File  
Division of Life Cycle API/ONDP/OPQ  
Center for Drug Evaluation and Research  
Food and Drug Administration

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<sup>1</sup> See FDA eCTD Web Page for further information.  
<http://www.fda.gov/Drugs/DevelopmentApprovalProcess/FormsSubmissionRequirements/ElectronicSubmissions/ucm153574.htm>.

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**This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.**  
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/s/  
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CLAUDE THEOPHIN  
08/16/2016