



Certificate of Drug Master File (DMF)

Service Period
June 1, 2020 – May 31, 2021

This certifies that:

Ilhaplast, S.A.
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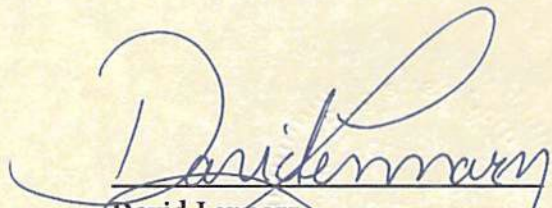
is a Drug Master File holder with the U.S. Food and Drug Administration pursuant to part 314 of Title 21, US Code of Federal Regulations, such filing having been verified as currently effective on the date hereof by Registrar Corp.

Drug Master File Number: **30228**
Status: **Active**
Type: **III**
Subject: **PHARMACEUTICAL PRIMARY PACKAGING**

Registrar Corp will confirm that such filing remains effective upon request and presentation of this certificate until May 31, 2021, unless terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. Filing of a Drug Master File does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of filing of Drug Master File is misleading. The U.S. Food and Drug Administration does not issue a certificate of Drug Master Files. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

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David Lennarz
Executive Director
Registrar Corp
Dated: April 16, 2020