DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
Silver Spring, MD 20993

DMF 036700

DMF ACKNOWLEDGEMENT

JAIN ALUFOILS PVT. LTD.
Attention: MR. MAHAVEER PRASAD JAIN, DIRECTOR
SURVEY NO.110/2/4, HIMALAYA INDUSTRIAL STATE
OPP. SILVASSA MUNCIPAL COUNCIL, AMILI
SILVASSA-3960230, U.T. OF D.N.H. & D.D., INDIA

Dear Mr. Mahaveer Prasad Jain,

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

DMF NUMBER ASSIGNED:	036700
DATE OF SUBMISSION:	DECEMBER 30, 2021
DMF TYPE:	III
SUBJECT (TITLE):	PRINTED/COATED/PLAIN/LAMINATED ALUMINIUM BLISTER FOIL, PRINTED/COATED/PLAIN/LAMINATED POLYSTER FILM, PRINTED/COATED/PLAIN/LAMINUATED BOPP FILM, PRINTED/COATED/PLAIN/LAMINATED POLY FILM, PAPER & OTHER FLEXIBLE PACKAGING FILM PRINTED & LAMINATED
HOLDER:	JAIN ALUFOILS PVT. LTD.
SUBMITTED BY:	JAIN ALUFOILS PVT. LTD.
AGENT:	NONE

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

Your DMF will be reviewed only in connection to 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.

Submissions in Paper (in two copies) or physical media should be sent 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

For information on various DMF submissions, example of letter templates and DMF Guidance for Industry, check the DMF website at https://www.fda.gov/drugs/forms-submission-requirements/drug-master-files-dmfs.

The holder of the DMF is responsible for compliance with 21 CFR314.420 as interpreted in "The Guideline for Drug Master Files" at https://www.fda.gov/drugs/drug-master-files-dmfs/guideline-drug-master-files-dmf.