



DMF 042506

DMF ACKNOWLEDGEMENT

SHANGHAI JIA TIAN PHARMACEUTICAL PACKAGING CO., LTD.
ATTENTION: LEON HUANG, QUALITY. DIRECTOR
NO: 3058 TINGFENG ROAD ZHUJING TOWN, JINSHAN DISTRICT
SHANGHAI, ZIP CODE: 201500, CHINA

Dear Leon Huang,

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

<u>DMF NUMBER ASSIGNED:</u>	042506
<u>DATE OF SUBMISSION:</u>	AUGUST 16, 2025
<u>DMF TYPE:</u>	III
<u>SUBJECT (TITLE):</u>	FLUORINE COATED ALUMINUM CANS FOR AEROSOLS
<u>HOLDER:</u>	SHANGHAI JIA TIAN PHARMACEUTICAL PACKAGING CO., LTD.
<u>SUBMITTED BY:</u>	SHANGHAI JIA TIAN PHARMACEUTICAL PACKAGING CO., LTD.
<u>AGENT:</u>	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.

You are expected to:

- Adhere to the statement of commitment you have provided.
- Provide the following submissions to the DMF:
 - Letters of Authorization (LOAs) granting permission to a third party (authorized party) or self to reference the DMF and for FDA to review the DMF. Listing an authorized party in the Annual Report 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 (with DMF number) is also not sufficient to authorize that party to reference the DMF.
 - Any change, addition or deletion of information
 - Annual Reports to the DMF containing:
 - 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.

U.S. Food and Drug Administration
Silver Spring, MD 20993
www.fda.gov