FDA 510(k) Premarket Notification

K222547 Electric nasal aspirator

CLEARED

Oct 20, 2022
58 days to decision

K222547 · Product code: **BTA** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k222547/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Portable, Aspiration (manual Or Powered) (BTA)
Date received	Aug 23, 2022
Decision date	Oct 20, 2022
Days to decision	58 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Xinlianfeng Technology Co.,Ltd**
Location	Shenzhen, CN
Contact	Ma Qiang
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	**Shenzhen Reanny Medical Devices Management Consulting Co., Ltd.**
Contact	Reanny Wang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov