

K213000 Air Compression Massager (Model LF-FT003)Nov 15, 2021
56 days to decisionK213000 · Product code: **IRP** · Physical Medicine
Source: <https://www.510kdatabase.net/k213000/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Sep 20, 2021
Decision date	Nov 15, 2021
Days to decision	56 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Wenzhou Lingfeng Electronic Technology Co., Ltd.
Location	Ruian, CN
Contact	Xiaoqun Ye
510(k) history	1 submissions · 1 cleared · 2021-2021

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](https://accessdata.fda.gov)

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