

K222669 Cryopush Cold Compression DeviceDec 5, 2022
90 days to decisionK222669 · Product code: **IRP** · Physical Medicine
Source: <https://www.510kdatabase.net/k222669/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Sep 6, 2022
Decision date	Dec 5, 2022
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Chengdu Cryo-Push Medical Technology Co.,Ltd
Location	Chengdu, CN
Contact	Zhang Peiyong
510(k) history	4 submissions · 4 cleared · 2022-2026

REGULATORY CONSULTANT

Consulting firm	Shenzhen Joyantech Consulting Co., Ltd.
Contact	Liz Li

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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