FDA 510(k) Premarket Notification

# K222669 Cryopush Cold Compression Device

**CLEARED**

Dec 5, 2022
90 days to decision

K222669 · Product code: **IRP** · Physical Medicine
Source: https://www.510kdatabase.net/k222669/

## SUBMISSION DETAILS

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|---|---|
| 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

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|---|---|
| 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

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|---|---|
| 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