

**K243206 Air compression massager (ACM-PLUS-A1, ACM-PLUS-A2, ACM-PLUS-A3, ACM-PLUS-A4, ACM-PLUS-A5, ACM-PLUS-A6)**Feb 19, 2025  
141 days to decisionK243206 · Product code: **IRP** · Physical Medicine  
Source: <https://www.510kdatabase.net/k243206/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Massager, Powered Inflatable Tube (IRP)
Date received	Oct 1, 2024
Decision date	Feb 19, 2025
Days to decision	141 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sichuan Qianli-Beoka Medical Technology, Inc.</b>
Location	Chengdu, CN
Contact	Qiuju Cai
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shenzhen Reanny Medical Devices Mgmt Consulting Co., Ltd.</b>
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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**510k Database** - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k243206/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 14, 2026