

**K182950 Acare Suction Unit**Aug 23, 2019  
304 days to decisionK182950 · Product code: **JCX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k182950/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Ward Use, Portable, Ac-powered (JCX)
Date received	Oct 23, 2018
Decision date	Aug 23, 2019
Days to decision	304 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acare Technology Co., Ltd.</b>
Location	Shanghai, CN
Contact	Kevin Tsai
510(k) history	2 submissions · 2 cleared · 2011-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>Ruscert Technology Co., Ltd.</b>
Contact	Ming-Yie Jan

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182950/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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