

**K812902 ACUPRESS VIBRATOR**Nov 10, 1981  
25 days to decisionK812902 · Product code: **IRO** · Physical MedicineSource: <https://www.510kdatabase.net/k812902/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vibrator, Therapeutic (IRO)
Date received	Oct 16, 1981
Decision date	Nov 10, 1981
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Eicona Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k812902/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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