FDA 510(k) Premarket Notification

K820331 IMPACT, MODEL 302

CLEARED

Feb 23, 1982
15 days to decision

K820331 · Product code: **JCX** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k820331/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Ward Use, Portable, Ac-powered (JCX)
Date received	Feb 8, 1982
Decision date	Feb 23, 1982
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Impact Instrumentation, Inc.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1981-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820331/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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