FDA 510(k) Premarket Notification

K820337 IMPACT, MODEL 303

CLEARED

Mar 2, 1982
22 days to decision

K820337 · Product code: **GCX** · General Hospital
Source: https://www.510kdatabase.net/k820337/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Suction, Operating-room, Wall Vacuum Powered (GCX)
Date received	Feb 8, 1982
Decision date	Mar 2, 1982
Days to decision	22 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/k820337/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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