

K823105 RESI-WALKERNov 5, 1982
16 days to decisionK823105 · Product code: **ITJ** · Physical MedicineSource: <https://www.510kdatabase.net/k823105/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Walker, Mechanical (ITJ)
Date received	Oct 20, 1982
Decision date	Nov 5, 1982
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Orthion Corp.
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1981-1983

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Device record: <https://www.510kdatabase.net/k823105/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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