

K780890 CAPP TERMINAL DEVICEJun 13, 1978
13 days to decisionK780890 · Product code: **IQX** · Physical MedicineSource: <https://www.510kdatabase.net/k780890/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hook, External Limb Component, Mechanical (IQX)
Date received	May 31, 1978
Decision date	Jun 13, 1978
Days to decision	13 days
Third-party review	No

APPLICANT

Company	Hosmer Dorrance Corp.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1977-1982

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Device record: <https://www.510kdatabase.net/k780890/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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