

K833800 BEECHER 20 GA. PROBEJan 4, 1984
65 days to decisionK833800 · Product code: **HQC** · Ophthalmic
Source: <https://www.510kdatabase.net/k833800/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Oct 31, 1983
Decision date	Jan 4, 1984
Days to decision	65 days
Third-party review	No

APPLICANT

Company	Beecher Medical Diagnostic Instrumentation
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1984-1987

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Device record: <https://www.510kdatabase.net/k833800/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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