

K830379 PHAKO KITApr 5, 1983
60 days to decisionK830379 · Product code: **HQC** · Ophthalmic
Source: <https://www.510kdatabase.net/k830379/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Feb 4, 1983
Decision date	Apr 5, 1983
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Vxtra Corp.
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1980-1983

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Device record: <https://www.510kdatabase.net/k830379/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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