

K832980 VXTRA OCUTOME KITOct 19, 1983
47 days to decisionK832980 · Product code: **HQC** · Ophthalmic
Source: <https://www.510kdatabase.net/k832980/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Sep 2, 1983
Decision date	Oct 19, 1983
Days to decision	47 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k832980/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026