FDA 510(k) Premarket Notification

K832980 VXTRA OCUTOME KIT

CLEARED

Oct 19, 1983
47 days to decision

K832980 · 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