FDA 510(k) Premarket Notification

K830379 PHAKO KIT

CLEARED

Apr 5, 1983
60 days to decision

K830379 · 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