FDA 510(k) Premarket Notification

K760199 ELECTRORETINOGRAPH

CLEARED

Aug 4, 1976
28 days to decision

K760199 · Product code: **HLZ** · Ophthalmic
Source: https://www.510kdatabase.net/k760199/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Corneal (HLZ)
Date received	Jul 7, 1976
Decision date	Aug 4, 1976
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Lkc Technologies, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1976-2015