

K760199 ELECTRORETINOGRAPHAug 4, 1976
28 days to decisionK760199 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k760199/> 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 June 4, 2026