FDA 510(k) Premarket Notification

K813399 ERG-JET ELECTRODE

CLEARED

Jan 22, 1982
49 days to decision

K813399 · Product code: **HLZ** · Ophthalmic
Source: https://www.510kdatabase.net/k813399/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Corneal (HLZ)
Date received	Dec 4, 1981
Decision date	Jan 22, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Nicolet Biomedical Instruments**
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k813399/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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