FDA 510(k) Premarket Notification

K812331 VTS-1000 VISUAL TESTING SYSTEM

CLEARED

Nov 16, 1981
90 days to decision

K812331 · Product code: **HLX** · Ophthalmic
Source: https://www.510kdatabase.net/k812331/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photostimulator, Ac-powered (HLX)
Date received	Aug 18, 1981
Decision date	Nov 16, 1981
Days to decision	90 days
Third-party review	No

APPLICANT

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