FDA 510(k) Premarket Notification

K821230 SOFTIC 49

CLEARED

May 28, 1982
31 days to decision

K821230 · Product code: **EBI** · Dental
Source: https://www.510kdatabase.net/k821230/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Apr 27, 1982
Decision date	May 28, 1982
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**The Healthcare Group Laboratories, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1980-1985