FDA 510(k) Premarket Notification

K810265 VISIO-BAND

CLEARED

Feb 23, 1981
21 days to decision

K810265 · Product code: **KLE** · Dental
Source: https://www.510kdatabase.net/k810265/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Feb 2, 1981
Decision date	Feb 23, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Espe-Premier Sales Co.**
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1981-1981