FDA 510(k) Premarket Notification

K782080 VENEER SYSTEM, MOSTIQUE LAMINATE

CLEARED

Jan 26, 1979
44 days to decision

K782080 · Product code: **KLE** · Dental
Source: https://www.510kdatabase.net/k782080/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Dec 13, 1978
Decision date	Jan 26, 1979
Days to decision	44 days
Third-party review	No

APPLICANT

Company	**Caulk Co., Div. Dentsply**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1981