FDA 510(k) Premarket Notification

K801690 VALIANT

CLEARED

Aug 13, 1980
21 days to decision

K801690 · Product code: **EJJ** · Dental
Source: https://www.510kdatabase.net/k801690/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Amalgam (EJJ)
Date received	Jul 23, 1980
Decision date	Aug 13, 1980
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**L.D. Caulk Co.**
Location	Mchenry, IL, US
Website	https://www.dentsplysirona.com
510(k) history	20 submissions · 19 cleared · 1976-1980

L.D. Caulk Co. is a Dental device manufacturer based in McHenry, US. The company specializes in restorative and preventive Dental products and materials. L.D. Caulk Co. has received FDA 510(k) clearances from total submissions. The company's regulatory activity spans from 1976 to 1980, with all submissions focused on Dental devices. This represents a historical record; the company has not received clearances in more than five years and is no longer active. Historical cleared devices include composite restoratives, dental amalgamators, calcium hydroxide products, and retra...