FDA 510(k) Premarket Notification

K792702 VASPAR SEALANT

CLEARED

Jan 24, 1980
28 days to decision

K792702 · Product code: **KEQ** · Pathology
Source: https://www.510kdatabase.net/k792702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Mounting, Water Soluble (KEQ)
Date received	Dec 27, 1979
Decision date	Jan 24, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Meridian Diagnostics, Inc.**
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999