FDA 510(k) Premarket Notification

K881992 SYRETTE

CLEARED

Jul 26, 1988
75 days to decision

K881992 · Product code: **EKJ** · Dental
Source: https://www.510kdatabase.net/k881992/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Burnisher, Operative (EKJ)
Date received	May 12, 1988
Decision date	Jul 26, 1988
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Oratec Corp.**
Location	Herndon, VA, US
Contact	KELNER, PHD
510(k) history	1 submissions · 1 cleared · 1988-1988