FDA 510(k) Premarket Notification

K882170 RX ASPEN

CLEARED

Jun 17, 1988
24 days to decision

K882170 · Product code: **EJH** · Dental
Source: https://www.510kdatabase.net/k882170/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Metal, Base (EJH)
Date received	May 24, 1988
Decision date	Jun 17, 1988
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Jeneric/Pentron, Inc.**
Location	Wallingford, CT, US
Contact	PRASAD, PH.D.
510(k) history	78 submissions · 78 cleared · 1988-2002