FDA 510(k) Premarket Notification

K833283 RX ORY PENTRON C & B

CLEARED

Nov 28, 1983
67 days to decision

K833283 · Product code: **EIS** · Dental
Source: https://www.510kdatabase.net/k833283/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Remover, Crown (EIS)
Date received	Sep 22, 1983
Decision date	Nov 28, 1983
Days to decision	67 days
Third-party review	No

APPLICANT

Company	**Genesis Industries, Inc.**
Location	Walker, MI, US
510(k) history	17 submissions · 17 cleared · 1983-2000