FDA 510(k) Premarket Notification

K051342 MEDICAL INTERPOROUS

CLEARED

Jun 2, 2006
375 days to decision

K051342 · Product code: **EFT** · Dental
Source: https://www.510kdatabase.net/k051342/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cleanser, Denture, Over The Counter (EFT)
Date received	May 23, 2005
Decision date	Jun 2, 2006
Days to decision	375 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bonyf AG**
Location	Silver Creek, NY, US
Contact	DONNA MARIE HARTNETT
510(k) history	1 submissions · 1 cleared · 2006-2006