

K051342 MEDICAL INTERPOROUSJun 2, 2006
375 days to decisionK051342 · Product code: **EFT** · DentalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051342/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026