FDA 510(k) Premarket Notification

CLEARED

K902402 HIGH SPEED NON-FIBER OPTIC SWIVEL HANDPIECE

Aug 29, 1990
90 days to decision

K902402 · Product code: **EBF** · Dental
Source: https://www.510kdatabase.net/k902402/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Material, Tooth Shade, Resin (EBF)
Date received	May 31, 1990
Decision date	Aug 29, 1990
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Dentalez Group**
Location	Lancaster, PA, US
Contact	GARY FOSS
510(k) history	13 submissions · 13 cleared · 1988-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k902402/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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