FDA 510(k) Premarket Notification

K911539 AUROFLUID 3

CLEARED

Apr 29, 1991
28 days to decision

K911539 · Product code: **EJT** · Dental
Source: https://www.510kdatabase.net/k911539/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Gold-based Noble Metal (EJT)
Date received	Apr 1, 1991
Decision date	Apr 29, 1991
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Metalor Dental USA Corp.**
Location	Rockville Centre, NY, US
Contact	BRIAN F SCHILLER
510(k) history	67 submissions · 67 cleared · 1991-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k911539/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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