

K882170 RX ASPENJun 17, 1988
24 days to decisionK882170 · Product code: **EJH** · DentalSource: <https://www.510kdatabase.net/k882170/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Metal, Base (EJH)
Date received	May 24, 1988
Decision date	Jun 17, 1988
Days to decision	24 days
Third-party review	No

APPLICANT

Company	Jeneric/Pentron, Inc.
Location	Wallingford, CT, US
Contact	PRASAD, PH.D.
510(k) history	78 submissions · 78 cleared · 1988-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882170/>, 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 May 26, 2026