

**K854977 FIBER MESH**Feb 20, 1986  
70 days to decisionK854977 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k854977/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 12, 1985
Decision date	Feb 20, 1986
Days to decision	70 days
Third-party review	No

**APPLICANT**

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Company	<b>Puldent Corp. of America</b>
Location	Mchenry, IL, US
Contact	KENNETH J BERK
510(k) history	6 submissions · 6 cleared · 1981-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854977/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 2, 2026