

**K821230 SOFTIC 49**May 28, 1982  
31 days to decisionK821230 · Product code: **EBI** · DentalSource: <https://www.510kdatabase.net/k821230/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Resin, Denture, Relining, Repairing, Rebasing (EBI)
Date received	Apr 27, 1982
Decision date	May 28, 1982
Days to decision	31 days
Third-party review	No

**APPLICANT**

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Company	<b>The Healthcare Group Laboratories, Inc.</b>
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1980-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k821230/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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