

**K901882 URESIL ENDOSCOPIC LASER OVERTUBE**Jul 27, 1990  
92 days to decisionK901882 · Product code: **KOQ** · Dental  
Source: <https://www.510kdatabase.net/k901882/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Adhesive, Denture, Carboxymethylcellulose Sodium (40-100&) (KOQ)
Date received	Apr 26, 1990
Decision date	Jul 27, 1990
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Uresil Corp.</b>
Location	Mchenry, IL, US
Contact	M GOLDBERG,MD
510(k) history	45 submissions · 44 cleared · 1981-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901882/>; 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 May 22, 2026