FDA 510(k) Premarket Notification

K901882 URESIL ENDOSCOPIC LASER OVERTUBE

CLEARED

Jul 27, 1990
92 days to decision

K901882 · Product code: **KOQ** · Dental
Source: https://www.510kdatabase.net/k901882/

SUBMISSION DETAILS

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

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