

**K812289 CELESTIN DILATOR**Sep 1, 1981  
19 days to decisionK812289 · Product code: **KNQ** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k812289/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal (KNQ)
Date received	Aug 13, 1981
Decision date	Sep 1, 1981
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Inmed Corp.</b>
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1981-1986

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Device record: <https://www.510kdatabase.net/k812289/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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