

**K900884 P.R.G. PERCUTANEOUS REPLACEMENT  
GASTROSTOMY**Jul 2, 1990  
126 days to decisionK900884 · Product code: **KGC** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k900884/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Tube, Gastro-enterostomy (KGC)
Date received	Feb 26, 1990
Decision date	Jul 2, 1990
Days to decision	126 days
Third-party review	No

**APPLICANT**

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Company	<b>Applied Medical Technologies</b>
Location	Mchenry, IL, US
Contact	PAT GILPIN
510(k) history	23 submissions · 20 cleared · 1981-1998

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900884/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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