

**K952325 ET TUBE PLACEMENT DETECTOR (WITH COLIBRI INDICATOR)**

Jul 31, 1995  
74 days to decision

K952325 · Product code: **CCK** · Anesthesiology  
Source: <https://www.510kdatabase.net/k952325/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	May 18, 1995
Decision date	Jul 31, 1995
Days to decision	74 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Icor AB</b>
Location	Bromma, SE
Contact	ANDRAS GEDEON
510(k) history	16 submissions · 16 cleared · 1986-1996

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

Device record: <https://www.510kdatabase.net/k952325/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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