

**K880241 &apos;HILT-WAY&apos; RESUSCITATOR**Mar 31, 1988  
71 days to decisionK880241 · Product code: **CAE** · AnesthesiologySource: <https://www.510kdatabase.net/k880241/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Airway, Oropharyngeal, Anesthesiology (CAE)
Date received	Jan 20, 1988
Decision date	Mar 31, 1988
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>Medifield , Ltd.</b>
Location	England, GB
Contact	RICHARD GILL
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880241/>, 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 27, 2026