

**K014140 RENAX HEMODIALYSIS BLOOD TUBING SET**Nov 18, 2002  
336 days to decisionK014140 · Product code: **FJK** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k014140/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Tubing, Blood, With And Without Anti-regurgitation Valve (FJK)
Date received	Dec 17, 2001
Decision date	Nov 18, 2002
Days to decision	336 days
Third-party review	No
Summary / Statement	Statement
Other names	SUNDER HEMODIALYSIS BLOOD TUBING SET

**APPLICANT**

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Company	<b>Sunder Biomedical Tech. Co., Ltd.</b>
Location	Taichung City, TW
Contact	TONY HUNG
510(k) history	3 submissions · 3 cleared · 2001-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k014140/> 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 June 29, 2026