

**K882998 ACUTE PHASE C-REACTIVE PROTEIN LATEX AGGLU.  
TEST**Sep 7, 1988  
51 days to decisionK882998 · Product code: **DCK** · Immunology  
Source: <https://www.510kdatabase.net/k882998/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	C-reactive Protein, Antigen, Antiserum, And Control (DCK)
Date received	Jul 18, 1988
Decision date	Sep 7, 1988
Days to decision	51 days
Third-party review	No

**APPLICANT**

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Company	<b>Amtec Diagnostics International, Inc.</b>
Location	Conroe, TX, US
Contact	HELEN JOHNSON
510(k) history	2 submissions · 2 cleared · 1987-1988

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882998/>; 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 30, 2026