

**K761152 S-UREA-N, IN VITRO DIAG FOR BUN**Dec 9, 1976  
10 days to decisionK761152 · Product code: **CDL** · Chemistry  
Source: <https://www.510kdatabase.net/k761152/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Berthelot Indophenol, Urea Nitrogen (CDL)
Date received	Nov 29, 1976
Decision date	Dec 9, 1976
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Advanced Biomedical Methods, Inc.</b>
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1976-1994

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

Device record: <https://www.510kdatabase.net/k761152/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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