

**K100616 KNOCKOUT SR MEDIUM, KNOCKOUT SR XENOFREE  
MEDIUM MODEL 10828, 12618**

May 20, 2010  
77 days to decision

K100616 · Product code: **NDS** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k100616/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Culture, Ex Vivo, Tissue And Cell (NDS)
Date received	Mar 4, 2010
Decision date	May 20, 2010
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Life Technologies Corporation</b>
Location	Grand Island, NY, US
Contact	KELLI TANZELLA
510(k) history	7 submissions · 7 cleared · 2010-2025

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

Device record: <https://www.510kdatabase.net/k100616/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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