

**K071210 MODIFICATION TO FIDIS CONNECTIVE 10*, MODEL
MX006**Dec 19, 2007
232 days to decisionK071210 · Product code: **LKO** · Immunology
Source: <https://www.510kdatabase.net/k071210/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anti-rnp Antibody, Antigen And Control (LKO)
Date received	May 1, 2007
Decision date	Dec 19, 2007
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomedical Diagnostics (Bmd) SA
Location	Marne La Vallee Cedex 2, FR
Contact	COURIVAUD CHRISTELLE
510(k) history	10 submissions · 10 cleared · 2005-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k071210/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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