

K990223 MODIFICATION OF CLEARPLAN EASY FERTILITY MONITORFeb 24, 1999
30 days to decisionK990223 · Product code: **NGE** · Chemistry
Source: <https://www.510kdatabase.net/k990223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Test, Luteinizing Hormone (lh), Over The Counter (NGE)
Date received	Jan 25, 1999
Decision date	Feb 24, 1999
Days to decision	30 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Unipath , Ltd.
Location	Bedford Mk41 Oqg, GB
Contact	Louise Roberts
510(k) history	73 submissions · 73 cleared · 1986-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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