

**K030333 LATEX EXAMINATION GLOVE,POWEDERED TYPE II,  
LOW MODULUS,WITH PROTEIN LABLING CLAIM**Mar 21, 2003  
49 days to decisionK030333 · Product code: LYY · General Hospital  
Source: <https://www.510kdatabase.net/k030333/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Jan 31, 2003
Decision date	Mar 21, 2003
Days to decision	49 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Perusahaan Getah Asas Sdn. Bhd.</b>
Location	42000 Port Kelang, S.D.E., MY
Contact	JAMES F LOGAN
510(k) history	30 submissions · 30 cleared · 1989-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030333/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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