

**K032938 BRIGHTWAY BRAND LATEX EXAMINATION  
GLOVE-(POWDERED, STERILE), CONTAINING 150 UGM OR  
LESS OF WATER EXTRACTABLE PROTEIN PER**Nov 20, 2003  
59 days to decisionK032938 · Product code: LYY · General Hospital  
Source: <https://www.510kdatabase.net/k032938/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Latex Patient Examination Glove (LYY)
Date received	Sep 22, 2003
Decision date	Nov 20, 2003
Days to decision	59 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Brightway Holdings Sdn. Bhd.</b>
Location	Klang, Selangor Darul Ehsan, MY
Contact	G. BASKARAN
510(k) history	24 submissions · 24 cleared · 1999-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032938/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 1, 2026