

Safety Recall

Merck Consumer Care in consultation with the U.S. Food and Drug Administration is voluntarily recalling

MiraLAX 30 Once Daily Doses Bottles

manufactured between August 2009 and June 2010

due to the potential for defective cap liners generated during the packaging process. This issue was discovered as part of the company's quality control processes. No significant adverse health consequences are expected to occur as a result of the use of these lots of MiraLAX.



Consumers with questions may contact Merck Customer support by phone at **1-800-647-2529** or visit

www.miraLAX.com.

Consumers with questions or concerns about their health should contact their doctor.

To be posted through November 15, 2010

SAFETY RECALL

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Consumers with questions may visit the Customer Service Desk or www.miraLAX.com or call **800-647-2529**.

8/15/10

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