Indiana State Department of Health
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Mitchell E. Daniels, Jr.
Governor

Gregory N. Larkin, M.D., F.A.A.F.P.
State Health Commissioner

DATE: June 8, 2011

TO: All Local Health Departments
Attn: Chief Food Inspection Officer

FROM: A. Scott Gilliam, MBA, CP-FS
Director, Food Protection Program

SUBJECT: US Nutrition Recall

SUGGESTED ACTION: **Unclassified Recall; Daily Multiple for Women 50 + Tablets, because certain lots contain undeclared Fish (fish gelatin); Recommend notification of affected stores via phone, fax or e-mail.**

From the information provided by FDA, the products being recalled may have been distributed in the State of Indiana. The products listed below were distributed throughout the United States via retail stores and through the Kaiser Permanente Health Organization. Detail information is not available at this time. In addition, if any recalled products are found, please notify this office at 317-233-7360.

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Recall -- Firm Press Release

FDA posts press releases and other notices of recalls and market withdrawals from the firms involved as a service to consumers, the media, and other interested parties. FDA does not endorse either the product or the company.

US Nutrition Issues Allergy Alert On Undeclared Fish In Daily Multible For Women 50 + Tablets

Contact:
Consumer:
888-534-6370

Media:
Brian Ellis
804-675-8140

FOR IMMEDIATE RELEASE - June 7, 2011 - **US Nutrition, of Ronkonkoma, New York** is recalling **Daily Multiple for Women 50 + Tablets**, because certain lots contain undeclared **Fish (fish gelatin)**. People who have an allergy or severe sensitivity to **Fish** run the risk of a serious or life-threatening allergic reaction if they consume these products.

This recall was initiated after it was discovered that this product contains **Fish (fish gelatin)** but was distributed in packaging that did not list the presence of Fish.

The tablet product was sold under the names **Daily Multiple for Women 50 +, One Daily for Women 50 Plus, One Daily Women's 50 +**, and **Women's 50+ Daily Formula**, under the brands shown in the table below.

The products listed below were distributed **throughout the United States** via retail stores and through the Kaiser Permanente Health Organization.

Product No.	Description	UPC Codes	Label	Expiration Dates	Lot Numbers
17850	CVS Daily Multiple for Women 50+, 50 tablets	0 50428 12299 0	CVS	12/13 12/13 01/14 01/14 02/14 03/14	318871-02 318871-03 330303-01 330303-02 337049-01 337050-01
17850	Giant Eagle One Daily Women's 50+, 50 tablets	0 30034 06682 6	Giant Eagle	01/14	330419-05
17851	CVS Daily Multiple for Women 50+, 100 tablets	0 50428 12867 1	CVS	12/13 01/14 01/14 01/14 02/14 03/14 03/14 03/14 03/14	318870-01 330303-03 330419-01 330419-02 337049-02 337050-02 338119-01 338119-03 338130-01
17874	Kaiser Permanente Women's 50+ Daily Formula, 120 tablets	0 01798 43712 2	Kaiser Permanente	12/13	318871-04
54381	Equate One Daily for Women 50 Plus, 50 tablets	6 81131 02390 0	Equate (Walmart)	12/13 12/13 12/13 12/13	318871-01 330301-01 330302-01 330302-02
54381	Equate One Daily for Women 50 Plus, 50 tablets	6 81131 02390 0	Equate (Walmart)	02/14 02/14 02/14	337047-01 334048-01 330419-03

As of this date, there have been no adverse reaction complaints reported relating to this recall.

Lots other than those listed above are not affected by this recall.

Consumers who have any of these products in their possession should return the product to the place of purchase or call 1-(888)-534-6370, Monday – Friday 9 am – 7 pm EDT for further instructions.

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RSS Feed for FDA Recalls Information[1] [what's this?[2]]

Photo: Product Labels[3]