Ohio Department of Agriculture
and
Ohio Department of Health

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Mike DeWine **Governor**	**Jon Husted** **Lieutenant Governor**	**Dorothy Pelanda** **ODA Director**	**Amy Acton**, M.D. **ODH Director**

DATE: March 5, 2019

TO: Health Commissioners, Directors of Environmental Health and Interested Parties

RE: Recall Announcement (ODA/ODH) 2019-042

McDaniel Life-Line LLC Issues Voluntary Worldwide Recall of Life-Line Water

McDaniel Life-Line LLC is voluntarily recalling all lots of Life-Line Water to the consumer level. This product is being recalled because FDA analysis found the product to be contaminated with Pseudomonas aeruginosa.

Use of the contaminated product has a remote probability of necessitating medical or surgical intervention to preclude or reverse permanent damage to a body structure or function. To date, McDaniel Life-Line LLC has not received any reports of adverse events related to this recall.

The product can be taken internally or applied externally to the skin. The product is packaged in 1-gallon bottles. The affected Life-Line Water recall includes all lots.

The product was distributed in the United States and Canada to individuals via internet sales@lifelinewater.com.

McDaniel Life-Line LLC is notifying its customers, by press release, of the recalled product. Consumers that have product which is being recalled should stop using and discard.

Consumers with questions regarding this recall can contact McDaniel Life-Line by phone 806-647-1741, Monday thru Friday 8 AM-5 PM, Central Time or by e-mail lifeline@amaonline.com. Consumers should contact their physician or healthcare provider if they have experienced any problems that may be related to taking or using this product.

Adverse reactions or quality problems associated with the use of this product may be reported to FDA's MedWatch Adverse Event Reporting program either by phone, on line, by regular mail or by fax.

- Complete and submit the report Online: www.fda.gov/medwatch/report.htm
- Regular Mail or Fax: Download form www.fda.gov/MedWatch/getforms.htm or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178.

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration.