Subject: ALFOODACT 039-2012 Burch Equipment LLC, North Carolina, Expands Recall to Include Additional Cantaloupe Shipping Dates and to Include Honeydew Melons

Date Issued: August 16, 2012

1. REFERENCES:

- a. DLAR 4155.26/AR 40-660/NAVSUPINST 10110.8c/AFI 48-116/MCO 10110.38c, DOD Hazardous Food & Nonprescription Drug Recall System.
- b. Allied Communications Publication 121, US SUPP-1 (f).

2. BACKGROUND:

Burch Equipment LLC, North Carolina, is expanding its recall to include all of this growing season's cantaloupes and honeydew melons that may remain on the market because they may possibly be contaminated with Listeria monocytogenes. There have been no illnesses reported to date.

Listeria monocytogenes is an organism which can cause serious and sometimes fatal infections in young children, frail or elderly people, and others with weakened immune systems. Although healthy individuals may suffer only short-term symptoms such as high fever, severe headache, stiffness, nausea, abdominal pain and diarrhea, infection can cause miscarriages and stillbirths among pregnant women. The incubation period (the length of time between consuming a product and becoming ill) for Listeria monocytogenes can be 1 to 3 weeks, but may be in the range of 3 to 70 days.

Honeydew melons involved in this recall expansion do not bear any identifying stickers and were packed in cartons labeled melons. (Photo; http://www.fda.gov/Safety/Recalls/ucm315366.htm)

Consumers who may have purchased these honeydew melons should contact the store where they purchased their melons, for information about whether those melons are part of this recall.

The cantaloupes and honeydew melons involved in this expanded recall were sold to distributors between June 23rd and July 27th, in the following states: FL, GA, IL, KY, MA, MD, ME, MI, NC, NH, NJ, NY, OH, PA, SC, and VA, VT and WV. The melons may have further been distributed to retail stores, restaurants and food service facilities in other states."

Burch Equipment LLC is requesting any consumer that may have one of these cantaloupes or honeydews to discard the product.

There have been no illnesses reported to date. FDA and the North Carolina Department of Agriculture and Consumer Services are working with Burch Equipment LLC following a random sample of a cantaloupe testing positive for Listeria monocytogenes.

This recall expansion is based on FDA's finding of Listeria monocytogenes on a honeydew melon grown and packed by Burch..

3. PRODUCTION DATES/IDENTIFYING CODES:

The whole cantaloupes are identified by a red label reading Burch Farms referencing PLU # 4319. All cantaloupes involved in the recall were grown by Burch Farms, however some of the cantaloupes may have been identified with a "Cottle Strawberry, Inc." sticker referencing PLU #4319 (note: Cottle Strawberry, Inc. did not grow or process the cantaloupe involved in this recall). Cantaloupes from Burch Farms were shipped in both corrugated boxes (9 cantaloupe per case) and in bulk bins.

(Photo; http://www.fda.gov/Safety/Recalls/ucm315366.htm)

4. MANUFACTURER/DISTRIBUTOR:

Burch Equipment LLC 910-267-5781 email burch@intrstar.net.

- **5. DISTRIBUTION:** [Multiple States] FL, GA, IL, KY, MA, MD, ME, MI, NC, NH, NJ, NY, OH, PA, SC, and VA, VT and WV
- **6. REASON FOR ACTION:** Due to possibly being contaminated with Listeria monocytogenes

7. INSTRUCTIONS AND ADDITIONAL INFORMATION FOR MESSAGE RECIPIENTS:

a. Immediately inventory stocks to identify the above items and secure in a "Medical Hold" status to provide assurance of no further issue/sale/use.

POSITIVE FINDINGS should be reported to Accountable Officers/Vendor Representatives of that facility. Accountable Officers/Agency representatives/Buyers/Contracting Officers should

facility. Accountable Officer/Agency representatives/Buyers/Contracting Officers should seek/refund/credit/replacement through the normal distribution channel with which the product was received (i.e. Distribution Centers, Prime Vendors, or Manufacturers).

- b. Ships at sea are authorized to destroy or dispose of recalled products at their discretion. Documentation for the number of pounds and cases, and any additional pertinent information must be signed by the Accountable Officer and is required for the purpose of recouping to the government the cost of the product involved. In order to get credit please use a SF 364 and forward to your supporting NAVSUP Fleet Logistics Center (NAVSUP FLC) and copy furnished to NAVSUP 51. Your supporting NAVSUP FLC should forward to the account manager at DLA Troop Support. The form should include the number of the recall authorizing the survey action. Home-ported ships/galleys will utilize DD form 1149 to transfer w/ reimbursement to the PV. The PV will submit credit invoice to the account manager at DLA Troop Support.
- c. DLA Troop Support Subsistence Prime Vendors must report POSITIVE and NEGATIVE RESPONSES directly to the their DLA Troop Support Contracting Officer with a courtesy copy to the Consumer Safety Officer (dscpconssafofc@dla.mil)..
- d. DeCA, AAFES, MWR, VA, MCCS, or other non-DLA Troop Support agencies SHOULD NOT respond to the DLA Troop Support Consumer Safety Officer. These agencies should report POSITIVE and NEGATIVE responses in accordance with their agency recall policies.

- e. When corresponding with DLA Troop Support concerning this message please include this message's subject in your subject line.
- **8.** The Point of Contact for this ALFOODACT message is CW4 Tony Hemphill, Consumer Safety Officer at DLA-FTW. VOICE, DSN: 444-2922, Commercial (215) 737-2922, or by FAX, DSN: 444-7526, or Commercial, (215) 737-7526, email dscpconssafofc@dla.mil.
- **9.** Individuals or groups that would like to receive recall messages electronically can forward their email address to dscpconssafofc@dla.mil, with "add to list" in the subject line. To be removed from the list place "remove from list" in the subject line.
- 10. Previous recalls and frequently asked questions are available at the following web site:

http://www.troopsupport.dla.mil/subs/fso/alfood/alfood.asp . The navigation tool to the left allows you to also view DLA Troop Support Alerts and Archived Vendor Recalls.

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