United States Food and Drug Administration

Southwest Import District

Notice of FDA Action

Entry Number: AM4-0179590-2

Notice Number: 3
July 7, 2008

Importer:
Elias Berry Mondragon
219 S. Pine Street
Santa Maria, CA 93456

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Port of Entry: 2506, Otay Mesa, CA
Carrier: BERRY MONDRAGON, ELIAS (MX-38;
Date Received: June 30, 2008
Arrival Date: June 30, 2008

Filer of Record: Guillermo Lizarraga Int'l Customs Broker, San Diego, CA 92154
Consignee: Elias Berry Mondragon, Santa Maria, CA 93456

HOLD DESIGNATED

Summary of Current Status of Individual Lines

	Line ACS/FDA	Product Description	Quantity	Current Status
*	001/001	JALAPENO PEPPER	192 CT	Released 07-07-2008
	001/002	JALAPENO PEPPER	192 CT	Product Collected by FDA 07-01-2008
	001/003	JALAPENO PEPPER	192 CT	May proceed 07-01-2008
	001/004	JALAPENO PEPPER	192 CT	May proceed 07-01-2008
	001/005	JALAPENO PEPPER	192 CT	May proceed 07-01-2008

* = Status change since the previous notice. Read carefully the sections which follow for important information regarding these lines.

@ = Consignee ID

FDA will not request redelivery for examination or sampling, if the products not released by FDA are moved, following USCS conditional release to a location within the local metropolitan area or to a location approved by the FDA office at the number below.

All products in this entry not listed above may proceed without FDA examination. This notice does not constitute assurance the products involved comply with provisions of the Food, Drug, and Cosmetic Act or other related acts, and does not preclude action should the products later be found violative.

LINES RELEASED

Line ACS/FDA	Product Description
001/001	JALAPENO PEPPER

Aleta T. Flores, Compliance Officer (Region/District)
U.S. Food and Drug Administration
9777 Via De La Amistad, Rm. 128
San Diego, CA 92154

(619) 661-3250 ext. 102
(619) 661-3195 (FAX)
ALETA.FLORES@FDA.HHS.GOV

These products are released. This notice does not constitute assurance that the product released complies with all provisions of the Food, Drug, and Cosmetic Act, or other related Acts, and does not preclude action should the product later be found violative.

Notice Prepared For: The District Director, U.S. Food and Drug Administration
Notice Prepared By: ATF