

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

Food and Drug Administration  
5600 Fishers Lane, HFS-681  
Rockville, MD 20857



**Date:** December 10, 2012

The U.S. Food and Drug Administration (FDA) is hereby providing you with a confirmation copy of the information FDA received regarding registration of your facility with the FDA as required by 21 CFR Part 1, Subpart H, and the Public Health Security and Bioterrorism Preparedness and Response Act of 2002. The FDA Registration of Food Facilities database shows:

<b>Food Facility Name:</b>	ND PHARMA & BIOTECH
<b>Food Facility Registration Number:</b>	11014242610
<b>PIN:</b>	[REDACTED]

<b>Street Address Line 1:</b>	[REDACTED]		
<b>Street Address Line 2:</b>	[REDACTED]		
<b>City:</b>	[REDACTED]	<b>State/Province:</b>	[REDACTED]
<b>ZIP/Postal Code:</b>	[REDACTED]		
<b>Country/Area:</b>	SPAIN		

You may want to review the accuracy of the registration in the attached paper document. **If the information is correct, no action is necessary.**

If the information is incorrect, you must update your registration within 15 days of receipt of this letter.

You can update your registration electronically via the FDA Industry Systems web site at <https://www.access.fda.gov/>.

If you have not used the FDA Industry Systems previously, you first will need to create an account.

There are tutorials at the above web address that will explain how to do so. If you did not create this registration online, you will need to link the registration to your account so you can view, update or cancel it.

Alternatively, you may submit an update by mail or fax to:

U.S. Food and Drug Administration  
5600 Fishers Lane, HFS-681  
Rockville, MD USA 20857  
Fax: 301-436-2804

You should include a copy of this letter with your mailed or fax response along with your request.

Please use Form 3537 if submitting a paper update to your registration, and be sure to check in Section 1 of the form the items applicable to the update.

You may contact the FDA Industry Systems Help Desk via telephone at: 1-800-216-7331 (Domestic) or 301-575-0156 if you have questions.

Thank You.

FDA Unified Registration and Listing and Prior Notice System Helpdesk