HEALTH MINISTRY.

OFFICE OF SANITARY REGULATION AND DEVELOPMENT

GENERAL DIRECTION OF SANITARY CONTROL

OF GOODS AND SERVICES.

REQUIREMENTS FOR APPLICATION TO REGISTRATION OR REVISION FOR IMPORTED GOOD SUPPLIES.

1.- Number of importer's health department license, and the warehouse's activity.

2.- Formula of the product in letterheaded paper of the manufacturer in the country of origin.

3.- Registration number in case of revision.

4.- Results of the physic/chemical and microbiological analyses of the finished product, executed by a credited laboratory in the country of origin.

5.- Certificate of free sale issued by the sanitary authority in the country of origin. In case of products of US origin, that certificate can be substituted by an analysis of the product, made by a laboratory with FDA certificate.

6.- Full description of packaging (primary and secondary package if the case) including original labels of the country of origin and the project in Spanish covering the specifications of the Mexican regulations.

7.- Representation letter of the manufacturer in the country of origin issued to the importer, granting the representtion confirmed by the Mexican Consul at the country of origin.

8.- Notarized letter of power of attorney with photograph in the name and issued to the person entitled to do the registration procedures.