Registrar Corp
144 Research Drive, Hampton, Virginia 23666 USA
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March 28, 2025

Alcoholera De Zapopan S.A. de C.V.
Vicente Guerrero No. 295
Agua Blanca Industrial
Zapopan, JA 45235
Mexico

Re: 2025 Electronic API Product Listing Certificates

Good Day:

We are pleased to provide the attached Certificates of Electronic API Product Listing issued by Registrar Corp. Your API listings are now active with the U.S. FDA. Your firm's APIs identified by the Certificates has been listed with the U.S. FDA in compliance with Title 21 of the United States Code of Federal Regulations (CFR). If there are any errors in the listing information on the Certificates, please notify us in writing, preferably by email.

Registrar Corp also will send you a copy of your Certificates of Electronic API Listing by mail. You may wish to use this electronic version to forward copies of your company's Certificates of Electronic API Listing to your customers and suppliers, so they are aware that your company has complied with the U.S. registration and listing requirements for the referenced drug products. Please note, however, that pursuant to 21 CFR § 207.77(a), "Registration of an establishment or listing of a drug does not denote approval of the establishment, the drug, or other drugs of the establishment, nor does it mean that a product may be legally marketed. Any representation that creates an impression of official approval or that a drug is approved or is legally marketable because of registration or listing is misleading and constitutes misbranding." Accordingly, these Certificates do not denote endorsement or approval by the U.S. FDA, and it should not be used to suggest such an inference.

Per 21 CFR § 201.2, "The National Drug Code (NDC) number is requested but not required to appear on all drug labels and in all drug labeling, including the label of any prescription drug container furnished to a consumer." Please also note, that if you revise your labeling, an updated listing must be submitted. If your company wishes to introduce another drug product into U.S. commerce, a separate listing will be required for that new drug.

U.S. FDA requires reviewing drug listing filings every June and December per 21 CFR § 207.57(b). Listings that have no changes must be certified between October 1st and December 31st each year. U.S. FDA does inactivate listings that are not kept up to date or certified. Update drug listing information in MyFDA (myfda.com) when there are any changes to the manufacturer's information or to the product.

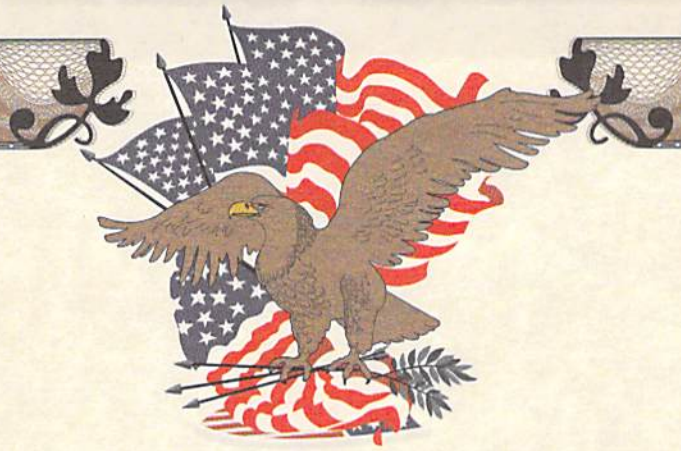
There is an additional requirement for listed drug products. Manufacturers are required to report the amounts of listed drug product manufactured to U.S. FDA. Per Guidance for Industry, Reporting Amount of Listed Drugs and Biological Products Under Section 510(j)(3) of the FD&C Act, FDA recommended deadline is March 31st. Registrar Corp also assists with this submission process for the listings we manage for your company.

As your U.S. Agent and Registrant Contact, Registrar Corp will continue to serve as a communications link between the U.S. FDA and your company for your electronic submissions. If we receive any correspondence from the U.S. FDA for your company regarding your electronic submissions, we will notify you by email, or phone. In addition, we will be pleased to complete electronic submissions for any drug products not already listed with the U.S. FDA which you may introduce to market, submit updates to already filed listings or assist with reporting amounts to FDA.

Please contact us if you have any additional questions or need additional help with FDA compliance.

Sincerely,

David Lennarz
President



2025

CERTIFICATE OF API PRODUCT LISTING

This certifies that:

Alcoholera De Zapopan S.A. de C.V.
Vicente Guerrero No. 295
Agua Blanca Industrial
Zapopan, JA 45235
Mexico

has listed the referenced API product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Registrar Corp:

Product Trade Name:	**Undenatured Ethyl Alcohol, 20 L**
Labeler:	**Alcoholera De Zapopan S.A. de C.V.**
NDC/DLS Number:	**81730-001-00**
U.S. Agent/	**Registrar Corp**
Registrant Contact:	144 Research Drive, Hampton Virginia, 23666 USA
	Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

Registrar Corp will confirm that such listing remains effective upon request and presentation of this certificate until the end of the year shown above, unless terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Registrar Corp, waives, and will hold harmless and indemnify Registrar Corp from any and all claims in connection with this product, its labeling, FDA drug listing, commerce or use. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of drug listing or the assignment of an NDC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue certificates of drug listing, nor does the U.S. Food and Drug Administration recognize certificates of drug listing. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

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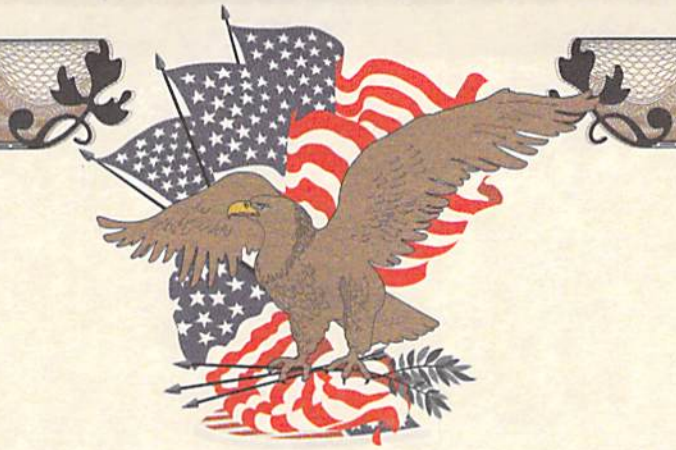
David Lennarz
Executive Director
Registrar Corp
Dated: March 28, 2025



2025

CERTIFICATE OF API PRODUCT LISTING

This certifies that:

Alcoholera De Zapopan S.A. de C.V.
Vicente Guerrero No. 295
Agua Blanca Industrial
Zapopan, JA 45235
Mexico

has listed the referenced API product with the U.S. Food and Drug Administration pursuant to part 207 of Title 21, US Code of Federal Regulations, such listing having been verified as currently effective on the date hereof by Registrar Corp:

Product Trade Name:	**Denatured Anhydrous Ethyl Alcohol, 20 L**
Labeler:	**Alcoholera De Zapopan S.A. de C.V.**
NDC/DLS Number:	**81730-002-00**
U.S. Agent/	**Registrar Corp**
Registrant Contact:	144 Research Drive, Hampton Virginia, 23666 USA
	Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

Registrar Corp will confirm that such listing remains effective upon request and presentation of this certificate until the end of the year shown above, unless terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. Holder assumes all risk, releases Registrar Corp, waives, and will hold harmless and indemnify Registrar Corp from any and all claims in connection with this product, its labeling, FDA drug listing, commerce or use. Registration of a drug establishment or drug wholesaler, or assignment of a registration number, or assignment of a NDC number does not in any way denote approval of the firm or its products by the U.S. Food and Drug Administration. Any representation that creates an impression of official approval because of drug listing or the assignment of an NDC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue certificates of drug listing, nor does the U.S. Food and Drug Administration recognize certificates of drug listing. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

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