

Phone: +1-757-224-0177 * Fax: +1-757-224-0179 * Email: info@registrarcorp.com * Website: www.registrarcorp.com

May 20, 2020

Vago International Manufacturing SA de CV Emilio Carranza 239 Centro de Monterrey Monterrey, NL 64000 Mexico

Re: 2020 Electronic Drug Listing Certificates

Good Day:

We are pleased to provide the attached Certificates of Electronic Drug Listing issued by Registrar Corp. Your drug listings are now active with the U.S. FDA. Your firm's drug 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 fax so that we may correct any problems for you.

Registrar Corp also will send you a color copy of your Certificates of Electronic Drug Listing by email. You may wish to use this electronic version to forward copies of your company's Certificate of Electronic Drug Listing to your customers and suppliers so they are aware that your company has complied with the U.S. registration and listing requirements for this drug product. Please note, however, that pursuant to 21 CFR § 207.39, "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." Any representation that creates an impression of official approval because of registration or possession of a registration number or NDC number is misleading and constitutes misbranding." This means the certificate does not denote endorsement or approval by the U.S. FDA, and it should not be used to suggest such an inference.

If the NDC number is shown on a drug label, it shall be placed as outlined in 21CFR207.35(b)(3). 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. Registrar Corp will be pleased to update your listing or help with additional drug listings.

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 fax, 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.

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

Sincerely,

David Lennarz Vice President

Registrar Corp is a private registration agent not affiliated with the U.S. Food and Drug Administration.



CERTIFICATE OF ELECTRONIC DRUG LISTING

This certifies that:

Vago International Manufacturing SA de CV Emilio Carranza 239 Centro de Monterrey Monterrey, NL 64000 Mexico

has listed the referenced drug 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: Ivgel Non-Sterile Solution (Hydroxyethl), 1000 ml

Labeler: Vago International Manufacturing SA de CV

Manufacturer: Vago International Manufacturing SA de CV

NDC/DLS Number: 74967-001-01 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.

Registrar Corp

144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179 info@registrarcorp.com • www.registrarcorp.com

David Lennarz Executive Director

Registrar Corp

Dated: May 20, 2020



CERTIFICATE OF ELECTRONIC DRUG LISTING

This certifies that:

Vago International Manufacturing SA de CV Emilio Carranza 239 Centro de Monterrey Monterrey, NL 64000 Mexico

has listed the referenced drug 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: IVGEL Hand Sanitizer, 60mL

Labeler: Vago International Manufacturing SA de CV
Manufacturer: Vago International Manufacturing SA de CV

NDC/DLS Number: 74967-000-02 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.

Registrar Corp

144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179 info@registrarcorp.com • www.registrarcorp.com David Lennarz

Executive Director Registrar Corp

Dated: 1 20, 2020