Registrar Corp
144 Research Drive, Hampton, Virginia 23666 USA
Phone: +1-757-224-0177 * Fax: +1-757-224-0179 * Email: info@registrarcorp.com * Website: www.registrarcorp.com

October 23, 2014
Organics Corporation of America dba Ambix Laboratories
55 West End Road
Totowa, NJ 07512
United States

Re: Drug Establishment Registration Renewal and Listing Certificates

Good Day,

We enclose the Certificate of Registration issued by Registrar Corp verifying that your firm's drug establishment registration has been electronically submitted to FDA for fiscal year 2015 (FY2015). We also enclose Certificates of Electronic Drug Listing for your products which are currently valid with FDA.

We advise that FDA has changed its requirement for annual registration. The Food and Drug Administration Safety and Innovation Act of 2012 ("FDASIA"), which was signed into law on July 9, 2012, requires that all registrants renew their Drug Establishment Registrations between October 1 and December 31 of each year. Your next renewal period will be October 1 to December 31, 2015. You will be sent documents to verify the accuracy of your information at that time.

Registrar Corp will send you color copies of your certificates by email. You may wish to use the electronic version to forward copies of your company's certificates to your customers and suppliers so they are aware that your company has complied with FDA's registration requirements. Please note 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 that the enclosed certificate does not denote endorsement or approval by the U.S. FDA, and it should not be used to suggest such an inference.

As your 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 questions or need additional help with FDA compliance.

Sincerely,

David Lennarz
Vice President

FY2015

CERTIFICATE OF REGISTRATION

This certifies that:

Organics Corporation of America dba Ambix Laboratories
55 West End Road
Totowa, NJ 07512
United States

is registered with the U.S. Food and Drug Administration for the statutory filing period applicable to U.S. FY 2015 pursuant to part 207 of Title 21, U.S. Code of Federal Regulations.

DUNS Number:	**061061164**
Labeler Code:	**10038**
Registrant Contact:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

Filing was performed during the October 1 - December 31, 2014 statutory period, and renewal is not required until the next statutory period of October 1- December 31, 2015. Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate, until the end of the year stated 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. 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 registration or possession of registration number or NDC number is misleading and constitutes misbranding. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

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Russell K. Statman
Executive Director
Registrar Corp
Dated: October 23, 2014

2015

CERTIFICATE OF ELECTRONIC DRUG LISTING

This certifies that:

Organics Corporation of America dba Ambix Laboratories
55 West End Road
Totowa, NJ 07512
United States

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:	**Ambix First Aid Antiseptic & Pain Relieveing Cream, 25g**
Labeler Code:	**10038**
Product Code:	**250**
Package Code:	**25**
NDC Number:	**10038-250-25**
Registrant Contact:	**Registrar Corp** 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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Product Trade Name:	**Ambix First Aid Antiseptic & Pain Relieveing Cream, 42.5g**
Labeler Code:	**10038**
Product Code:	**250**
Package Code:	**42**
NDC Number:	**10038-250-42**
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