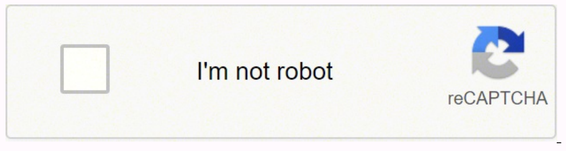
Fda philippines joint affidavit of undertaking form



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Fda philippines joint affidavit of undertaking form



ARTICLES OF INCORPORATION AND BY-LAWS - STOCK CORPORATION

Articles of Incorporation

Of

BALANCED SOLUTION ADVERTISING CORPORATION

(Name of the Corporation)

KNOW ALL MEN BY THESE PRESENTS:

The undersigned incorporators, all of legal age and are residents of the Philippines, have this day voluntarily agreed to form a stock corporation under the laws of the Republic of the Philippines;

AND WE HEREBY CERTIFY:

FIRST: That the name of said corporation shall be

BALANCED SOLUTION ADVERTISING CORPORATION

SECOND: That the primary purpose for which such corporation is incorporated:

1. To earn profit by rendering advertising services to clients.
2. To manage, operate, provide legal and professional advertising services, and do all other lawful things as may be necessary, incidental, or convenient to carry on the business as provided herein.
3. To engage in business transactions related to enhance the image of our clients, publicize their new products and programs, and promote and convince people that their services or products are worth of utilizing or not.

Provided that the corporation shall not solicit, accept or take investment/placements from the public neither shall it issue investment contracts.

THIRD: That the principal office of the corporation is located in

Republic of the Philippines)
Province of Cagayan) S. S.
City of Tuguegarao)
x----------x

AFFIDAVIT OF LOSS

I, **VICENTE FERRER,** Filipino, of legal age and a resident of San Juan, Tuao, Cagayan, after first having been duly sworn to in accordance with law deposes and states:

1. That I was the driver of that Nissan Patrol owned by the Provincial Government of Cagayan bearing Plate Number **SCR 964**;

2. It is unfortunate that said Plate Number was accidentally detached from said Nissan Patrol without my notice;

3. That I went back to the places I have traveled in search of the said Plate Number but all efforts to locate the same proved futile, thus, it is presumed lost for all intents and purposes;

4. That the said plate was never lent to another person or entity nor confiscated by any other law enforcement agency;

5. That I bind myself to surrender the said plate to the district office that issued the same in the event that said plate is recovered;

6. That I have executed this affidavit to attest to the veracity of the foregoing statements and to secure a replacement of the aforementioned Plate Number.

IN WITNESS WHEREOF, I have hereunto affixed my signature this __ day _____ 2021 in Tuguegarao City, Cagayan, Philippines.

VICENTE FERRER
affiant

SUBSCRIBED AND SWORN to before me this __ day _____ 2021 in Tuguegarao City, Cagayan, Philippines.

Notary Public

Doc. No.
Page No.
Book No.
Series of



Exempt from VRA while sealed
Access for VRA when made public

Agency Case Number: _____

To Clerk:
1. This affidavit must be kept sealed until charges related to the warrant are filed. It may be unsealed and made confidential (case) after the warrant is issued. Crim. R. 17(c) (1) & (3).
2. When you unseal the affidavit after charges are filed, you must redact it to determine if it contains confidential information under the Victim's Rights Act as provided in Admin. Bulletin 11 sec. 09 before releasing it to the public.

IN THE DISTRICT/SUPERIOR COURT FOR THE STATE OF ALASKA AT _____

In the Matter of:)
)
SEARCH WARRANT AUTHORIZING)
ELECTRONIC MONITORING)
)
Court No. [DR:] _____
_____/

Affidavit Supporting Request For Electronic Monitoring Warrant
(to measure and record a conversation in which one participant has consented to the recording)

Being first duly sworn, I, _, state that:

1. I am: a police officer; a federal agent; a state investigator;
☐ currently assigned to a _ investigative unit; ☐ _

2. Criminal charges: have ☐ have not ☐ been filed concerning this matter.
If charges have been filed, the court case number is: _____

3. I have reason to believe that statements about criminal activity will be made as follows:

The person expected to make the statements is _____
Description: Male Female
☐ Caucasian ☐ Black ☐ Hispanic ☐ Alaska Native ☐ _____
Approximate _____ age _____ Other: _____

The statements are expected to be made ☐ in person ☐ by telephone at _____
_____ who will be at or near (city & state): _____

The statements are expected to occur between the dates of _____
and _____

The conversation is expected to include statements about criminal activity involving:
☐ controlled substances (list) ☐
☐ sexual assault/abuse
☐ _____

Page 1 of 2 AO 11.30 (03/20) (3)
CR 39(b) (3)(F) & (c) Criminal Rules 37 & 11
AFFIDAVIT IN SUPPORTING REQUEST FOR ELECTRONIC MONITORING WARRANT

The Philippines is a growing market for cosmetics, with the GlobalData analysis company that previews the industry that is worth the PHP 228,8 billion (4,2 billion dollars) within the 2023. Contributing to this potential for recovery is the growing population of the country and the reduction of unemployment rates. The citizens of the nation's archipelago appreciate the interactions of beauty and social, leading the demand for cosmetic products in the Philippines. Incredibly, there is an opportunity to have here for enterprising people and companies. After all, cosmetics are an umbrella term. Products such as perfumes, hair dyes, lipsticks, soaps, lotions and other skin care products are all in high demand and will continue to be so in the short and long term. If you are interested in importing, selling or distributing cosmetic products in the Philippines, it is first and foremost necessary to secure the necessary permits from the Food and Drug Administration (FDA). Through the registration process you will show potential customers who are interested in their health and safety. In this article you will get the lowdown on how to register cosmetic products in the country.regulation of cosmetic products in the cosmetic standards of Philippinesmain in PhilippinESSTE Philippine Cosmetics Regulatory Environment is based mainly on the Cosmetics Directive ASEAN (ACD), an initiative to promote trade between the members of the Association of Southeast Asian Nations (ASEAN) is harmonising the requirements of the technical sector while maintaining the standards of safety and quality. To this directive, cosmetic products are "the substance or preparation intended to be placed in contact with the outer parts of the human body... or with the teeth and mucous membranes of the oral cavity with a view exclusively or principally to their cleansing, perfuming or changing their appearance, and body and/ or protecting them or keeping them in good condition. Several other regulations have also been issued by which are linked to the ASEAN rationalisation initiative. Among them is the Administrative Order (AO) 2016-0003 (Â¢ â¯¬ ÂGuidelines on requirements and procedures for unified food and drug licensing), which aims to improve approval processes for cosmetic products and strengthen the institution " Post-market surveillance activities by unifying and harmonizing requirements FDA regulatory body licensing.Â¨ in the Philippine Center for Cosmetics Regulation and Research (CCRR) is the main arm of the FDA responsible for the regulation of the cosmetic product. This department further branches out into two divisions – (1) the Licensing and Registration Division and (2) Research on Product Research and the Standard Development Division.STEPS in the Registration of Cosmetic Products in the PhilippinesSiclense to operate (LTO) is required a company to obtain a Before that can legally import, distribute, sell wholesale and / or the manufacture of cosmetics. Company at Dayanan Consulting, we help our clients facilitate their application process with the FDA from start to finish, from LTO labeling. They only need to provide us with the necessary information and we help prepare the necessary documents. But if you choose to try to do it yourself, here are the requirements to protect it to one: Notarized petition form from the owner, the incorporator or the authorized representativeJoint Affidavit of the company signed by the owner and an authorized individual (an allied health scientist professional) claiming to understand the FDA regulation for LTO; Note that the authorized individual must also submit their professional license or board certification of the master file registrationsite of your business “This is required by manufacturers of drugs, devices and cosmetics) Proof of employment or addresses Plan and Surroundings Plan and Surroundings Business Location MapRISK PlanFeafter Management Application Preparing these requirements, you can now proceed to file them with the FDA. Note Note If you are just renewing your LTO, the presentation of the first, second and last items would be sufficient. You will therefore have to await the results of the FDA assessment. For manufacturers, pre-opening inspection is necessary as part of the FDA's post-marketing surveillance activities. The FDA can inspect all covered plants at any time, so it is necessary to be prepared and conform to all standards. Notification before the product can be released on the market, the FDA must be notified before using the procedures outlined in the Cosmetic Notification v. 2.0 User manual for candidates. You must provide the following information in your cosmetic product notification: brand name/ product name/ product variants "The full name of the product must be provided in the following sequence: brand name, line name (if applicable) product name and shade number if a single shade is notified (e.g. ABC Brand XYZ Eyshadow Shade 1). If there are different shades, the shade name/ number for each shade must be declared. the category of cosmetic product for which notification is being sought. Details of the product- the validity of the notification can be one, two or three years at the option of the applicant. This refers to the function or use of the product and not the instructions for use, e.g. To hydrate the face, hand, etc. Product presentations- Select if you are applying for (1) a single product, (2) a range of similar variants in composition for the same use but differs in colors, flavours, etc. (3) palette (s) in a range of a product type, or (4) products combined in a single kit. Local company responsible for placing the cosmetic product on the market -indicate the local company the placing of cosmetic products on the market. This company must also have a valid LTO number. If the company responsible for placing the cosmetic product on the market is a distributor, then the applicant also declare the details of the direct supplier or manufacturer of the cosmetic product, ie country of manufacture, address, and altri.194; 160informazioni on the factory Â ¢ Â Ô refers to the manufacturer's information and / or provider of the cosmetic product notificato.Persona representing the local company Â ¢ Â Â refers to the company representative who can be contacted in the event FDA questions about the cosmetic product or whether it is additional documents needed to certify lâ applicazione.194; 160; List of product ingredients Â ¢ Â Â all the ingredients of the product must be specified using the nomenclature dellâ latest edition of the standard references (International Cosmetic Ingredient Dictionary, British Pharmacopoeia, United States Pharmacopoeia, Chemical Synthesis Services). Botanical and botanical extract must be identified for its genus and its species. The genus can be abbreviated. The functions and the percentages of the ingredients must be declared if it is dâ substances

subject to restrictions on use, as specified in the Annexes of the ASEAN Cosmetic Directive. To apply, follow the following steps: Getting a user account CCRR requiring info@fda.gov.ph one. To proceed, you must provide the following information: e-mail address or e-mail address of the company DestinationContact No.Company NameCompany AddressProduct Classification: CosmeticsLog in at the e-portal FDA). Click on Ã ¢ Â Â New Houses226 Â Â and carry out the notification form dell'ASEAN.Scarica the notifica.126 form; ~~ ~ Â Â Â Â and assign this task to the Cashier FDA.Pagare the fee or through any bank branch or through the methods of payment BancNet on-line.Aspetta the result of domanda.Download and prints the order confirmation once your application has been received. approvate.Le organizations who submitted the correct application documents will then be a cosmetic product notification certificate that is valid for 1-3 years. Those who have insufficient documentation will be informed of the deficiencies in which or a further explanation of the details will be requested. Finally, anyone who has declared misleading information or failed to meet FDA standards will deserve a letter of disapproval. Cosmetic notifications may be reconsidered for a new validity date, in which the new validity date will be based on the date of submission of the revalidation application. Note that the same application process for cosmetic notification and-and notification applies for reassessment. In the e-portal, the previously recognised case must be selected to continue with the activity for the reassignment application. There shall be no change from the information provided during the previous application to use reconsideration. Therefore, any changes to the information will constitute a new notification application. If the notification is requested to be cancelled, the applicant may choose to cancel the application. This option is available in the re-validation application activity. Production requirements: GMP, ingredients and Labellingfor Manufacturers, your personal production processes and practices, premises, equipment, hygiene and hygiene services, etc. They shall comply with the good manufacturing practice (GMP) established by the ASEAN Guidelines. Please note that a GMP certificate of conformity will only be issued on demonstration of satisfactory compliance and will be effective until the validity of your current LTO. Subsequently, it will be released whenever your LTO is renewed. In accordance with procedures and processes, it is also necessary to ensure that ingredients used in your cosmetic products adhere to ASEAN cosmetic regulations. The Annexes to the ASEAN Cosmetic Directive provide for prohibitions and regulations on ingredients of the cosmetic product as listed below: Table 1. Annexes to the ASEANS Cosmetics Directive and the prohibitions of ingredients for cosmetic products in the Philippines Asean cosmetic directives which are prohibited from forming part of Composition of cosmetic products in Annex III Substances which are prohibited to be contained in cosmetic products, except those subject to restrictions and conditions set out in Annex IVColours which may be used in cosmetic products in Annex VII Preservatives which may be contained in cosmetic products When it comes to the labelling of cosmetic products produced in the Philippines In addition, the following products must be seen on the product Ã¢ÂÂ; packaging:Weight or volume of the product contentBrand, name of the product and its functionality Batch number Manufacturing or expiration date of the product in terms of (month/year) Country where the cosmetic product is manufactured Instructions on how the product is used Prev Special auctions (if any) List of all product ingredients Name and address of the company or person who placed the product on the market For decorative cosmetic products that are marketed in different shades of colour, all colouring agents may be listed, provided that the description includes the words, _;128; Sanctions associated with cosmetic products in the Philipetic Misbranding of cosmetics, deliberately or accidentally, will ensure that violators can get at least one year (1) or at most ten years. Fines ranging from Php 15,000 to Php 500,000 can also be imposed. Some may also have to address both issues depending on a judge;226;ÔÊ;Â decisions. On the other hand, for offenders who are manufacturers, importers or distributors of any cosmetic product, the minimum detention period would be five years while the amount of the fine ranges from Php 500,000 to Php 5 million. million.

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