



BRGY.
PERMIT



1st
Quarter

2nd
Quarter

3rd
Quarter

4th
Quarter



REPUBLIC OF THE PHILIPPINES
CITY OF TAGUIG
BUSINESS PERMIT



Name of Corporation/Entity

VISTA C HEALTH PRODUCTS CORPORATION

Trade Name

VISTA C HEALTH PRODUCTS CORPORATION

Location of Business

2ND FLR. EL-43 / 45 SUNSHINE MALL,, WESTERN
BICUTAN CITY OF TAGUIG

Owner of Business

VISTA C HEALTH PRODUCTS CORPORATION

Ownership Type

CORPORATION

Status

RENEWAL

Account No.

006-14000355

LCN No.

11-014843

Area (Sq. m)

60.00

Date Issued

January 30, 2020

Valid Until

December 31, 2020

O.R. Number

A-4768763 - January 30, 2020

2020-00109

APPROVED BY:

Fanella Joy B. Panga - Cruz

Atty. Fanella Joy B. Panga - Cruz
Chief, BPLO

TAX DESCRIPTION	TAX BASE	AMOUNT DUE
TAX ON RETAILERS/TRADING / BUY & SELL	1,124,040.00	12,013.18 1-4 2020
MAYORS PERMIT		5,100.00 2020
ENVIRONMENTAL IMPACT FEE		534.38 1-4 2020
SANITARY INSPECTION FEE		255.00 2020
BUILDING INSPECTION FEE		204.00 2020
ELECTRICAL INSPECTION FEE		255.00 2020
PLUMBING INSPECTION FEE		127.50 2020
MECHANICAL INSPECTION FEE		255.00 2020
MEDICAL/HEALTH FEE		12.75 2020
FIRE PERMIT FEE		200.00 2020
BUSINESS PLATE (STICKER)		150.00 2020
FORM FEE		150.00 2020
SIGNBOARD FEE		178.50 2020
FIRE CODE RA 9514		901.50 2020
CEDULA CORPORATION		949.62 2020
TOTAL:		21,286.43

This permit may be revoked for violation of applicable laws or regulations and for non-payment of taxes and fees.
For illegal use of this permit, please notify BPLO at
(02) 555-7800 / 555-7818

ERASURE AND/OR ALTERATIONS WILL INVALIDATE THIS PERMIT



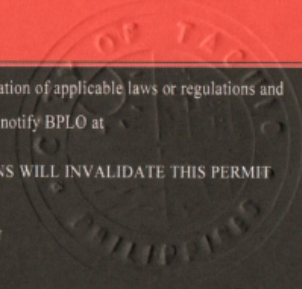
www.taguig.gov.ph



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REPUBLIC OF THE PHILIPPINES
SECURITIES AND EXCHANGE COMMISSION
SEC Building, EDSA, Greenhills
City of Mandaluyong, Metro Manila

COMPANY REG. NO. CS201221897

**CERTIFICATE OF FILING
OF
AMENDED ARTICLES OF INCORPORATION**

KNOW ALL PERSONS BY THESE PRESENTS:

This is to certify that the amended articles of incorporation of the

VISTA C HEALTH PRODUCTS CORPORATION
[Amending Article III thereof.]

copy annexed, adopted on December 03, 2013 by majority vote of the Board of Directors and by the vote of the stockholders owning or representing more than two-thirds of the outstanding capital stock, and certified under oath by the Corporate Secretary and a majority of the Board of Directors of the corporation was approved by the Commission on this date pursuant to the provision of Section 16 of the Corporation Code of the Philippines, Batas Pambansa Blg. 68, approved on May 1, 1980 and copies thereof are filed with the Commission.

Unless this corporation obtains or already has obtained the appropriate Secondary License from this Commission, this Certificate does not authorize it to undertake business activities requiring a Secondary License from this Commission such as, but not limited to acting as: broker or dealer in securities, government securities eligible dealer (GSED), investment adviser of an investment company, close-end or open-end investment company, investment house, transfer agent, commodity/financial futures exchange/broker/merchant, financing company and time shares/club shares/membership certificates issuers or selling agents thereof. Neither does this Certificate constitute as permit to undertake activities for which other government agencies require a license or permit.

IN WITNESS WHEREOF, I have set my hand and caused the seal of this Commission to be affixed to this Certificate at Mandaluyong City, Metro Manila, Philippines, this 17th day of April, Twenty Fifteen.




FERDINAND B. SALES
Director

Company Registration and Monitoring Department



Republic of the Philippines
 Department of Health
FOOD AND DRUG ADMINISTRATION
 Civic Drive, Filinvest Corporate City, Alabang, Muntinlupa City



CERTIFICATE OF PRODUCT REGISTRATION

Pursuant to the provisions of Republic Act (R.A.) No. 3720 as amended, known as the Foods, Drugs, Devices and Cosmetics Act, and consistent with R.A. No. 6675, known as the Generics Act of 1988, and R.A. No. 9711, otherwise known as the Food and Drug Administration Act of 2009, the product described hereunder has been found to conform with the requirements and standards for marketing authorization of pharmaceutical products per existing regulations in force as of date hereof.

Registration Number : DR-XY45949

Generic Name : Ascorbic Acid (as Sodium Ascorbate)
Brand Name : Vista C Supreme
Dosage Strength & Form : 444.50mg Capsule (equivalent to 500mg Sodium Ascorbate)
Pharmacologic Category : Vitamin
Classification : Over-the-Counter (OTC) Drug
Approved Shelf-life : 24 months
Storage Condition : Store at temperatures not exceeding 30°C.
Packaging : White HDPE Bottle with tamper-evident PE screw cap x 100 capsules (Box of 100's)

Manufacturer : Doctors Pharmaceuticals, Inc.
 #8 Veterans Center, West Bicutan, Taguig City

Trader : Vista C Health Products Corporation
 2nd Floor, Sunshine Plaza Mall F.T.I. Complex,
 Taguig City

The marketing authorization shall be valid until 10 July 2020 subject to the conditions listed on the reverse side. No change in the formulation, labelling and commercial presentation of this product shall be made at any time during the effectivity of this registration without prior written approval of this Office.

This marketing authorization is subject to suspension, cancellation or recall should any violation of R.A. No. 3720, R.A. No. 6675 and R.A. No. 9711 and/or regulations issued thereunder involving the product be committed.

Witness My Hand and Seal of this Office, this 10 July 2017.

By Authority of the Director General
 Per FDA Order No. 2016-005

BENJAMIN G. CO, MD, FPPS, FPSECP
 Director IV
 Center for Drug Regulation and Research

REG. STATUS : Initial
 AMOUNT : Pbp9,090; Pbp510; Pbp510; Pbp1,010
 OR NUMBER : 0723691; 0821891; 0825199; 0848632
 DATE : 29 Mar 2016; 9 Feb 2017; 16 Feb 2017; 3 May 2017
 CODE : 151.114
 BAR CODE : 
 DOC TRACK : 2 0 1 6 0 3 1 7 0 9 5 0 2 3





LICENSE TO OPERATE

as
Drug Trader
is hereby granted to

VISTA C HEALTH PRODUCTS CORPORATION

2nd Floor, Sunshine Plaza Mall F.T.I. Complex, Taguig, Metro Manila

Owner: VISTA C HEALTH PRODUCTS CORPORATION
License Number: LTO-3000001233767
Application Type: Renewal
Validity of License: 16 April 2023



This LTO shall be renewed within **90 days** before its expiration, upon compliance with relevant laws, rules and regulations and the payment of fees. This LTO may be suspended, cancelled or revoked by this Office for cause if found violating RA 9711 and related issuances.

Furthermore, with this LTO, the FDA allows the establishment to apply for a market authorization [i.e. registration (CPR) or notification] for health products prior to manufacture, importation, sale or offer for sale, distribution, transfer, advertisement and/or promotion as the case may be.

BY AUTHORITY OF THE DIRECTOR GENERAL

JESUSA JOYCE N. CIRUNAY, RPh
Director IV
Center for Drug Regulation and Research

DISPLAY IN PUBLIC VIEW

Additional information required under FDA Circular 2016-006 are reflected at the second page of this LTO