



CONSUMER PRODUCTS SERVICES DIVISION

GOLDEN EAGLE ENTERPRISE GROUP LTD

Technical Report: (5217)345-0508
Date Received: December 11, 2017

January 30, 2018

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JOHN NG
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RM1604 NO.168
SAI YEUNG CHOI STREET
MONG KOK
KOWLOON
HONG KONG

Sample Description:	A WATER DROP ESSENTIAL OIL - MR. ROBUST		
Vendor:	GOLDEN EAGLE ENTERPRISE GROUP LTD	Sample Size:	0
Manufacturer:	N/A	Style No(s):	N/A
Buyer:	N/A	SKN/SKU No.:	N/A
Labeled Age Grade:	N/A	PO No.:	N/A
Appropriate Age Grade:	N/A	Ref #:	N/A
Client Specified Age Grade:	N/A	Country of Origin:	AUSTRALIA
Grade:			
Tested Age Grade:	N/A	Assortment No.:	N/A
UPC Code:	489703986023		

EXECUTIVE SUMMARY:

A set of reference samples was submitted for the US FDA cosmetic ingredient list review.

See findings in RESULTS & DISCUSSION Sections **for details**.

BUREAU VERITAS HONG KONG LIMITED

Alex Hui
Senior Manager
SBU of Analytical HK

AH/vc



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(I) RESULTS:

PART (I): COSMETIC INGREDIENT LIST REVIEW

SECTION	ASSESSMENT AREA	COMMENT
A	Hazardous Substances	Meet
B	Color Additives and Warning Statement	Meet
C	International Nomenclature of Cosmetic Ingredients (INCI) Name Format	See Discussion
D	Arrangement of Ingredients	Not Requested



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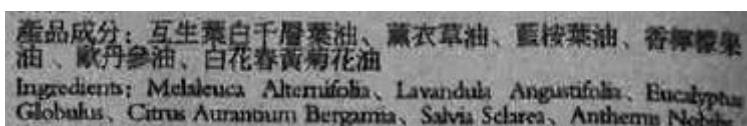
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(II) DISCUSSION :

A reference sample was submitted for cosmetic ingredients review in compliance with Code of Federal Regulations (CFR), Title 21, Section 701, 740, 73, 74 and 82. The following ingredient list was submitted for review:

INGREDIENTS :



(A) HAZARDOUS SUBSTANCES – MEET (NONE PRESENT)

With the exception of color additives, the Food and Drug Administration (FDA) does not have a list of “acceptable” ingredients for use in cosmetics. There is a list of prohibited ingredients and other hazardous substances which cannot be used in cosmetics or have limited use. These ingredients are listed in 21 CFR 250.250; AND 21 CFR, Part 700, Sections 11, 13, 14, 16, 18 and 23.

The ingredients listed are not found in the list of prohibited ingredients and other hazardous substance. According to the FDA's Cosmetic Handbook “The FD&C Act does not require that a product be tested for safety. However, the FDA strongly urges cosmetic manufactures to conduct toxicological and other appropriate tests to substantiate the safety of the cosmetics.”

(B) COLOR ADDITIVES AND WARNING STATEMENT –MEET

The INCI name of the colorant can be used only when applied to batches of the colorant that have been certified by the US FDA according to 21CFR part 80. A FDA signed certificate is issued for each batch of colorant.



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(II) DISCUSSION :

(C) INTERNATIONAL NOMENCLATURE OF COSMETIC INGREDIENTS (INCI) NAME FORMAT – SEE TABLE (I) FOR CORRECTION

The United States Code of Federal Regulations (CFR), Title 21, Part 701, Section 3 recognizes the Cosmetic, Toiletry and Fragrance Association (CTFA), International Cosmetic Ingredient Dictionary and Handbook as a source of names for cosmetic ingredient labeling (INCI name).

Any ingredient's inclusion in this book means that it is available for sale for use in cosmetic products. It does not mean that the ingredients are in compliance with the United States or any other country's laws and regulations.

Not all of the ingredients are listed in the Cosmetic, Toiletry and Fragrance Association (CTFA), International Cosmetic Ingredient Dictionary and Handbook, Eighth Edition. Moreover, common name or technical name are listed in the ingredients list and the exact INCI name are not used. They are listed in the following table:

TABLE (I) : CONVERSION OF INCI NAME

Technical / Common Name (Existing name in the ingredient list)	INCI Name (Official name for labeling)
Melaleuca alternifolia	Melaleuca alternifolia (Tea Tree) Leaf Oil
Lavandula Angustifolia	Lavandula Angustifolia (Lavender) Oil
Eucalyptus Globulus	Eucalyptus Globulus Leaf Oil
Citrus Aurantium Bergamia	Citrus Aurantium Bergamia (Bergamot) Fruit Oil
Salvia Sclarea	Salvia Sclarea (Clary) Oil
Anthemis Nobilis	Anthemis Nobilis Flower Oil

The above INCI names and others in the ingredient list shall be used in cosmetic ingredient labeling.



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(II) DISCUSSION :

(D) ARRANGEMENT OF INGREDIENTS – ORDER : NOT EVALUATED

The Food and Drug Administration (FDA), Code of Federal Regulation (CFR), Title 21, Section 701.3 requires that the label on each package of a cosmetic shall bear a declaration of the name of each ingredient in descending order of predominance and must appear on the outer packaging of the cosmetic in letters of not less than 1/16 inch or 1/32 inch for labelling area < 12 in², be conspicuously placed and easy to read.

As the percentage of ingredient was not provided so the descending order of predominance was not checked.