**Preamble**

1. According to the Pharmacy and Poisons Regulations (PPR)(Cap. 138A) a subsidiary legislation of the Pharmacy and Poisons Ordinance (PPO)(Cap. 138), pharmaceutical products (PP) must be registered with the Pharmacy and Poisons Board (PPB) before they can be sold, offered for sale or distributed or possessed for the purposes of sale, distribution or other use in Hong Kong.

2. The term of “pharmaceutical products” is defined in the section 2 of PPO. Application for registration of pharmaceutical products is made to the PPB, a statutory body to determine whether or not that product is a pharmaceutical product and require to be registered. Details of the provisions under PPO and PPR can be browsed at [www.legislation.gov.hk](http://www.legislation.gov.hk).

3. The guidance notes aim to provide general principles and advices to facilitate the trade to decide if products are pharmaceutical products or not. This document is not legally binding and provides only guidance.

**Definition of Pharmaceutical Product**

4. According to section 2 of the PPO, "pharmaceutical product" (藥劑製品) and "medicine" (藥物) mean any substance or combination of substances—
   
   (a) presented as having properties for treating or preventing disease in human beings or animals; or
   
   (b) that may be used in, or administered to, human beings or animals, either with a view to—
      
      (i) restoring, correcting or modifying physiological functions by exerting a pharmacological, immunological or metabolic action; or
      
      (ii) making a medical diagnosis.
General Principles for Classification

5. To determine whether a product is a pharmaceutical product or not, it is on a case by case basis and in the light of:
   - the definition set out in paragraph 4 above;
   - relevant Court precedents or legal advice from Department of Justice; and
   - following an assessment of all the available information about the product*

*Product information includes full details of product’s composition, presentation and purpose. Account will be taken of material being used to promote the product. For details, please refer to “Factors to be considered” at later paragraphs.

6. Below are the examples of products generally not considered as pharmaceutical products subject to the registration control of PPR:

(I) Proprietary Chinese medicines are exempted from the control of PPO and PPR which are subject to regulatory control under the Chinese Medicine Ordinance (Cap 549). Under Section 2 of the Chinese Medicine Ordinance (Cap. 549), proprietary Chinese medicines are defined as follows:

   "proprietary Chinese medicine" (中成藥) means any proprietary product
   a) composed solely of the following as active ingredients-
      i) any Chinese herbal medicines; or
      ii) any materials of herbal, animal or mineral origin customarily used by the Chinese; or
      iii) any medicines and materials referred to in subparagraphs (i) and (ii) respectively;
   b) formulated in a finished dose form; and
   c) known or claimed to be used for the diagnosis, treatment, prevention or alleviation of any disease or any symptom of a disease in human beings, or for the regulation of the functional states of the human body.

   Additional information related to the control of proprietary Chinese medicines may be found at the website of the Chinese Medicine Council of Hong Kong at http://www.cmchk.org.hk.

(II) A product which the average consumer would regard as something to be eaten, drunk or chewed as part of his/her diet for example, because of its taste, flavor, or nutritional value is unlikely to be classified as pharmaceutical product unless it contains one or more ingredients generally regarded as medicinal substance and indicative of a medical use.
(III) A product which the average consumer would regard as cosmetic, beauty and skin care, sunscreen, toothpastes, deodorants and antiperspirants, hair colourants and hair styling products in nature is unlikely to be classified as pharmaceutical products unless it contains one or more ingredients generally regarded as medicinal substance and indicative of a medical use.

(IV) A medical device is generally known as any instrument, apparatus, implement, machine, appliance, implant, in vitro reagent or calibrator, software, material or other similar or related article, intended by the manufacturer to be used, alone or in combination, for human beings for one or more of the specific purpose(s) of:

a) diagnosis, prevention, monitoring, treatment or alleviation of disease;
b) diagnosis, monitoring, treatment, alleviation of or compensation for an injury;
c) investigation, replacement, modification, or support of the anatomy or of a physiological process;
d) supporting or sustaining life;
e) control of conception (including contraception);
f) disinfection of medical devices;
g) providing information for medical purposes by means of in vitro examination of specimens derived from the human body;

and which does not achieve its primary intended action in or on the human body by pharmacological, immunological, or metabolic means, but which may be assisted in its intended function by such means.

Additional information on the control of medical devices may be found at the website of the Department of Health’s Medical Device Control Office at www.mdco.gov.hk.

(V) Whole human blood; or any human blood component, other than plasma prepared by a method involving an industrial process, or under highly manipulation.

**What is a pharmaceutical product?**

7. As mentioned above, when determining whether a particular product comes within the definition of pharmaceutical product is on a case by case basis, and product information includes full details of product’s composition, presentation, purpose and promotional material will be assessed by considering the relevant factors.
Factors relevant to deciding whether a product is a pharmaceutical product

8. In order to assess a product is regarded as a “pharmaceutical product”, it is essential to consider and make known of whether:

(a) the substance or combination of substances is present in the product;
(b) the substance or combination of substances is medicinal or not;
(c) the product is in pharmaceutical dose form (i.e. capsule, tablet, etc.), and the way it is to be used;
(d) the use(s) indicated on the label, packaging/package inserts, promotional materials is/are within the scope of uses under the definition;
(e) any essentially similar pharmaceutical products registered in Hong Kong; and
(f) the product may pose any risk to the public.

9. Since each product is considered individually, it is not possible to provide a simple list of substances which will be considered as pharmaceutical products. However, as considering whether the substance is medicinal or not, it may be helpful to refer to the substances listed under the heading “A” of the Poisons List whose uses are essentially medicinal. For other examples of substance which are not included in the Poisons List but may generally be regarded as medicinal, please refer to Appendix 1.

10. Some substances which are commonly found in the health food products lacks of scientific evidence to support their medicinal use, and are normally NOT regarded as medicinal substance. For examples of those substances, please refer to Appendix 2.

11. As considering whether the use of the product is fall within the definition of pharmaceutical products, the context in which the medicinal claims of usage made in the labeling, packaging/package inserts, promotional materials and the overall presentation will be taken into account.

12. Some words or phrases which may present the product as having properties for treating or preventing disease are medicinal claims. Although it is not possible to produce an indicative list of all kinds of medicinal claims of usage in this guidance, some examples which are considered as medicinal claims are listed in Appendix 3 for reference.
13. Claims to “maintain”, “help to maintain”, or “support” health or a healthy lifestyle are normally not considered as medicinal in themselves. Examples of those claims are listed in Appendix 4 for reference.

14. Therefore, if a product is found to contain medicinal substance(s) at a reasonable amount with medicinal claims, it would generally be classified as a “pharmaceutical product”.

15. On the contrary, a product carries claims without any evidence may contravene the Trade Description Ordinance Cap. 362, Laws of Hong Kong.

16. For some cases, the general line for deciding whether the product is or is not pharmaceutical had been reached by the PPB. For details of those cases, please refer to Appendix 5.

**Disclaimer**

The guidance notes are only intended to provide general information on the classification of products as “pharmaceutical products” and should not be considered as a substitute for legal or other professional advice. Whenever necessary, please refer to the Pharmacy and Poisons Ordinance and Regulations for details of the requirements. The Department of Health accepts no liability for any loss or damaged caused, arising directly, or indirectly, in connection with reliance on the contents of the guidance notes.

Drug Office,
Department of Health
December 2015
Appendix 1

Examples of substances other than the one listed in the Poisons List may generally be regarded as medicinal are as following:

- Alfacalcidol
- Androstenediol
- Androstenedione
- Antibiotic
- Aspirin
- 1,4 Butanediol (BD)
- Calcitriol
- Climbazole [Note: Please see Appendix 5 for the exceptions]
- Digestive enzymes, e.g. papain, pancreatin, lipase, cellulase, bromelain, amylase, etc.
- Hydroxyapatite
- Laetrile (amygdalin)
- N-N-Dimethylglycine (DMG, pangamic acid)
- Paracetamol
- Selenium for anti-dandruff use
- Sennosides
- Tars (coal tar or pine tar)
Examples of substances which are normally NOT regarded as medicinal are as following:

- Animal cartilage
- Amino acids e.g. alanine, arginine, citrulline, cysteine, cystine, glycine, histidine, isoleucine, leucine, lysine, phenylalanine, serine, tyrosine, etc. (except injection form),
- Apple cider vinegar
- Bee pollen
- Bioflavonoids, e.g. diosmin, hesperidin, quercetin, rutin, etc.
- Biotin
- Brewer’s yeast
- Caffeine
- Camphor (external preparations)
- Casein
- Chitosan
- Chlorophyll
- Choline
- Chondroitin
- Coenzyme Q10 (ubidecarenone)
- Collagen
- Colostrum
- Creatine
- DHEA (dehydroepiandrosterone)
- Fibers from fruits and vegetables
- Fish liver oils
- Fish oils
- Gamma aminobutyric acid (GABA)
- Glucosamine (except injection form)
- Goat's milk
- Grape seed extract (pycnogenol)
- Herbal substances, e.g. bilberry, blueberry, cranberry, echinacea, garcinia cambogia, ispaghula husk, psyllium husk (plantago ovata), rose hips, saw palmetto, etc. (except belladonna, cascara, ephedra and yohimbe)
- Hydroquinone
- Lactic acid producing organisms, e.g. bifidobacterium, lactobacillus
- Lecithin
- Lutein
- Melatonin
- Menthol (external preparations)
- Minerals, e.g. calcium, copper, iodine, iron, magnesium, zinc, etc. (except injection form)
- Nucleic acids, e.g. deoxyribonucleic acid (DNA), ribonucleic acid (RNA), etc.
- Omega-3 triglycerides, e.g. docosahexanoic acid (DHA), eicosapentaenoic acid (EPA), etc.
- Phystosterols
- Seaweeds, e.g. kelp
- Simethicone/dimethicone for topical use
- Squalene
- Urea
- Vitamins (except injection form and other preparations under Appendix 5)
- Wheat germ oil
- Whey protein
- Zeaxanthin
Appendix 3

Examples of words or phrases listed below indicated association with medicinal claims.

- “this product helps to prevent heart disease”
- “prevents osteoporosis”
- “treatment or management of obesity”
- “headlice treatment”
- “prevents/relieves allergies”
- “prevents acne/pimple”
- “this product heals cold sores”
- “cures athlete’s foot”
- “frequent use of the product can alleviate pimples”
- “anti-gingivitis mouthwash”
- “use of the product can prevent infections”
- “frequent use can fight cold and flu”
- “remedy for hay fever”
- “this product relieves occasional constipation/diarrhoea”
- “prevention of travel sickness”
- “eases heartburn and indigestion”
- “treats mouth ulcers”
- “fights periodontal diseases”
- “prevents periodontitis”
Appendix 4

Examples of words or phrases listed below are normally not considered as medicinal in themselves.

- anti-plaque
- teeth whitening/polishing
- prevents teeth decay/prevents teeth cavity
- removes teeth stains
- relieves teeth sensitivity
- fights bad odor
- cleanses acne-prone skin
- energizes skin
- helps to prevent signs of aging
- hypoallergenic
- soothes sensitive skin
- smoothes wrinkles
- skin whitening
- fades dark pigmented areas
- improves skin conditions and relieves dryness
- anti-dandruff (without coal tar, pine tar, selenium, etc.)
- vitalizes hair
- improves general health
### Case 1: Insect repellant product

<table>
<thead>
<tr>
<th>Topic</th>
<th>Substante: DEET</th>
</tr>
</thead>
<tbody>
<tr>
<td>Decision</td>
<td>DEET containing insect repellant do not satisfy the definition of pharmaceutical product under section 2 of PPO, and therefore is not subject to the registration requirement.</td>
</tr>
</tbody>
</table>

### Case 2: Anti-dandruff product

<table>
<thead>
<tr>
<th>Topic</th>
<th>Substance: Zinc pyrithione</th>
</tr>
</thead>
<tbody>
<tr>
<td>Decision</td>
<td>Products containing Zinc pyrithione and presented as anti-dandruff products without medicinal claim, will not be regarded as pharmaceutical products.</td>
</tr>
</tbody>
</table>

### Case 3: Antiseptic and disinfectant product

<table>
<thead>
<tr>
<th>Substance in the product</th>
<th>Maximum concentration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Benzalkonium salts</td>
<td>6% when used diluted or in rinse-off preparations</td>
</tr>
<tr>
<td></td>
<td>1% when used undiluted or in leave-on applications</td>
</tr>
<tr>
<td>Benzethonium salts</td>
<td>0.1%</td>
</tr>
<tr>
<td>Cetrimide</td>
<td>3%</td>
</tr>
<tr>
<td>Cetylpyridinium salts</td>
<td>2.5% when used diluted or in rinse-off preparations</td>
</tr>
<tr>
<td></td>
<td>0.3% when used undiluted or in leave-on applications</td>
</tr>
<tr>
<td>Chlorhexidine salts</td>
<td>20% when used diluted or in rinse-off preparations</td>
</tr>
<tr>
<td></td>
<td>2% when used undiluted or in leave-on applications</td>
</tr>
<tr>
<td>Chloroxyculenol</td>
<td>4.8% when used diluted or in rinse-off preparations</td>
</tr>
<tr>
<td></td>
<td>0.5% when used undiluted or in leave-on applications</td>
</tr>
<tr>
<td>Substance</td>
<td>Concentration</td>
</tr>
<tr>
<td>----------------------------</td>
<td>--------------------------------</td>
</tr>
<tr>
<td>Dichloroxylenol</td>
<td>2%</td>
</tr>
<tr>
<td>Ethyl alcohol</td>
<td>All concentrations</td>
</tr>
<tr>
<td>Isopropyl alcohol</td>
<td>All concentrations</td>
</tr>
<tr>
<td>Phenoxyisopropanol</td>
<td>2% in rinse-off applications or used diluted</td>
</tr>
<tr>
<td></td>
<td>1% in leave-on applications or used undiluted</td>
</tr>
<tr>
<td>Salicylic acid</td>
<td>2%</td>
</tr>
<tr>
<td>Thymol</td>
<td>1%</td>
</tr>
<tr>
<td>Triclocarban</td>
<td>1%</td>
</tr>
<tr>
<td>Triclosan</td>
<td>2% when used diluted</td>
</tr>
<tr>
<td></td>
<td>1% when used undiluted</td>
</tr>
</tbody>
</table>

**Decision:** Antiseptic and disinfectant products are not classified as pharmaceutical product if the following condition fulfilled:
- Containing the substance specified in above; and
- No medicinal claims; and
- Not labeled for use on broken skin.

**Case 4:**

<table>
<thead>
<tr>
<th>Topic:</th>
<th>DMAA containing products</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substance:</td>
<td>1,3-Dimethylamylamine (DMAA)</td>
</tr>
<tr>
<td>Decision:</td>
<td>The Registration Committee of the Pharmacy and Poisons Board decided to regulate 1,3-Dimethylamylamine (DMAA) as pharmaceutical product with effect from 1 April 2013, after considering the pharmacological effects of DMAA, its potential risk of causing adverse effects and the international situations in the control of DMAA.</td>
</tr>
</tbody>
</table>

**Case 5:**

<table>
<thead>
<tr>
<th>Topic :</th>
<th>Hair care or cosmetic products containing climbazole</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substance:</td>
<td>Climbazole</td>
</tr>
</tbody>
</table>
| Decision:                    | i). Hair care products containing climbazole not exceeding 2% in rinse-off products, or 0.5% in leave-on products are not considered as pharmaceutical products under the PPO, unless they are labelled for medicinal uses; and  
   ii). Cosmetic products containing climbazole as a preservative with a maximum concentration of 0.5% are not considered as pharmaceutical products under the PPO, unless they are labelled for medicinal uses. |
Case 6:

<table>
<thead>
<tr>
<th>Topic</th>
<th>Vitamin products</th>
</tr>
</thead>
<tbody>
<tr>
<td>Substance</td>
<td>Vitamins</td>
</tr>
<tr>
<td>Decision</td>
<td>In addition to the scenarios mentioned in the previous appendices, vitamin products are not considered as pharmaceutical products unless they belong to the following categories in oral dose form: a) vitamin A with not less than 10,000 I.U. daily dose; b) vitamin B3 (nicotinic acid) with more than 200 mg daily dose; c) vitamin D with more than 1,000 I.U. daily dose; and d) vitamin K except vitamins K1 or K2 with 120 mcg or less daily dose.</td>
</tr>
</tbody>
</table>