# **Newfoundland and Labrador Pharmacy Board**



### **Provincial Drug Schedules**

August 2021

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#### INTRODUCTION

#### General

In November 2014, changes were made to the Newfoundland and Labrador Drug Scheduling process in accordance with the new Pharmacy Regulations, 2014 (see Appendix A). As such, changes to the provincial drug schedules are now reviewed and approved by the Newfoundland and Labrador Pharmacy Board rather than the Minister of Health and Community Services.

This approval process will still be based on the recommendations of the National Drug Scheduling Advisory Committee, as described below. Once a recommended change is received by the Newfoundland and Labrador Pharmacy Board, it is added to the agenda for the next Board meeting for review and approval.

\*\*A recommended change is NOT effective until it has received Board approval.\*\*

#### National Drug Scheduling Advisory Committee

In May 1995, the National Association of Pharmacy Regulatory Authorities (NAPRA) approved a national scheduling model and, to ensure ongoing review and maintenance of the drugs listed in the model schedules, established the National Drug Scheduling Advisory Committee (NDSAC) in August 1995. NDSAC includes experts from across Canada as well as other relevant representatives and is mandated to advise the provincial regulatory authorities on matters relating to the placement of drugs within the national scheduling model and to continually evaluate and maintain the drug scheduling factors within the model (see Appendix B).

NDSAC meets several times throughout the year to review and analyse submissions from sponsors requesting scheduling additions or revisions, generally in response to:

- A federal government proposal to deregulate a federal prescription status drug;
- A new drug approved for sale in Canada on a non-prescription basis; or
- A request from a manufacturer, the public or other stakeholder to reassess or review the current recommended schedule status of a drug.

For a full overview of NDSAC's policies and procedures, please visit <u>napra.ca/drug-scheduling-canada-general-overview</u>.

#### SCHEDULE I – PRESCRIPTION SALE ONLY

Schedule I drugs require a prescription for sale and are provided to the public by the pharmacist following the diagnosis and professional intervention of a practitioner. They are in addition to, but subject to the same conditions as, the drugs listed in the Prescription Drug List established in accordance with section 29.1 of the Food and Drugs Act and its accompanying Regulations (Canada).

Some drugs on this schedule may appear to be non-prescription drugs since there is no "Pr" symbol directly on the product. Pharmacists must be aware of these products to prevent possible sale without a prescription.

The following table provides a current list of Schedule I drugs. **Recent changes are indicated with BOLD UNDERLINE**. Examples of the Brand Names of products containing these drugs are also provided. Please note that the list of Brand Names is not necessarily inclusive and are provided for reference only.

Chemical name as it appears in the Schedule	Examples of common Brand Names
allergy serums and extracts	•
alpha 1-proteinase inhibitor (human)	
alverine and its salts for parenteral use	
amino acid solutions for parenteral use	
aminopromazine [proquamezine] and its salts	
bacillus Calmette-Guerin vaccine	
bacitracin and its salts and derivatives for parenteral use	
calcium chloride in injectable form for parenteral nutrition	
calcium gluconate in injectable form for parenteral nutrition	
cholera vaccine except oral, inactivated, when used for prophylaxis against	
Traveler's Diarrhea due to enterotoxigenic escherichia coli [ETEC]	
chromium chloride (chromic chloride) in injectable form for parenteral nutrition	
copper chloride (cupric chloride) in injectable form for parenteral nutrition	
copper sulphate in injectable form for parenteral nutrition	
cytomegalovirus immune globulins	
dextrose injection in concentrate solutions for parenteral nutrition	
encephalitis vaccine (Japanese)	
ephedrine and its salts in preparations containing more than 8 mg per unit dose, or	
with a label recommending more than 8 mg/dose or 32 mg/day, or labelled or	
implied for use exceeding 7 days, or if indicated for other than nasal congestion	
epinephrine and its salts (other than in pre-filled syringes intended for emergency	
administration by injection in the event or anaphylactic reactions to allergens)	
erythrityl tetranitrate	
esomeprazole or its salts, except when sold for the 14-day treatment for frequent	
heartburn at a daily dose of 20 mg in package sizes of no more than 280 mg of	
esomeprazole	
ethylpapaverine and its salts	
flumazenil	
hepatitis A vaccine	
hepatitis B adult vaccine	
hepatitis B immune globulin	
herpes zoster (shingles) vaccine (live)	
homatropine and its salts for ophthalmic or parenteral use or in preparations for	
oral use containing more than 2 mg per dosage unit	
hydrocortisone or hydrocortisone acetate, when sold in a concentration that	
provides 1% or less hydrocortisone in preparations for topical use on the skin in	
adults and children 2 years of age and over in package sizes containing more than	
30g	
hydrocortisone or hydrocortisone acetate, when sold in a concentration that	
provides 1% or less hydrocortisone in preparations for topical use on the skin in	
children under 2 years of age	
immune globulin products	
isopropamide and its salts	
isosorbide and its salts	Imdur
levallorphane and its salts	

Chemical name as it appears in the Schedule	Examples of common Brand Names
lipid solutions in injectable form for parenteral nutrition	, p
Lyme Disease vaccine	
magnesium sulphate in injectable form for parenteral nutrition	
manganese and its salts in injectable form for parenteral nutrition	
metaraminol bitartrate	
mupirocin	Bactroban
naproxen and its salts, except in preparations containing the equivalent of 200 mg	Anaprox
of naproxen base* per oral dosage unit when sold in products labelled with a	Naprosyn
recommended maximum daily dose equivalent to 400 mg of naproxen base	Ναριοδήτι
The state of the s	
[*220 mg naproxen sodium approximately equivalent to 200 mg naproxen base]	
nicotinyl-tartrate	
nikethamide	
nitroglycerin (except for sublingual immediate release dosage forms)	Minitran
	Nitro-Dur
	Transderm-Nitro
	Trinipatch
omeprazole or its salts (except when sold for the 14-day treatment for frequent	- Timpaton
heartburn at a daily dose of 20 mg, in package sizes of no more than 280 mg of	
omeprazole)	
orphenadrine hydrochloride	
papaveretrine and its salts	
papaverine and its salts	
pentaerythritol tetranitrate	
potassium salts in preparations for injection	
proquamezine [aminopromazine] and its salts for internal use	
quinidine salts	
rabies immune globulin	
rabies vaccine	
rho D immune globulin	
selenium in injectable form for parenteral nutrition	
sodium acetate in injectable form for parenteral nutrition	
sodium chloride in injectable form for parenteral nutrition	
sodium iodine in injectable form for parenteral nutrition	
sodium phosphate in injectable form for parenteral nutrition	
streptokinase/streptodormase	
succinlycholine and its salts	
tetanus immune globulin	
typhoid vaccines / salmonella typhi vaccines	
vaccines (except those part of a routine immunization program in most/all	
provinces & territories: cholera vaccine (oral, inactivated) when used for	
prophylaxis against traveller's diarrhea and due to enterotoxigenic escherichia coli	
(ETEC); diphtheria toxoid, haemphilus influenzae type b, hepatitis B pediatric,	
influenza, measles, mumps, pertussis, pneumococcus, poliomyelitis, rubella,	
tetanus toxoid; and those requiring special enhanced public access due to disease	
outbreaks: meningococcus)	
varicella vaccine (chicken pox)	
varicella zoster immune globulin	
vitamin K (except Vitamin K1 and Vitamin K2 sold (a) for external use in humans;	
or (b) in an oral dosage form for use in humans if the maximum recommended daily	
dose is 0.120 mg or less)	
vitamins in injectable form for parenteral nutrition	
yellow fever vaccine	
zinc chloride in injectable form for parenteral nutrition	
zinc sulphate in injectable form for parenteral nutrition	

#### SCHEDULE II - SALE ONLY FROM A NO PUBLIC ACCESS AREA OF A DISPENSARY

Schedule II drugs, while less strictly regulated, do require professional intervention from the pharmacist at the point of sale and possibly referral to a practitioner. While a prescription is not required, the drugs are available only from the pharmacist and must be retained within an area of the pharmacy where there is no public access and no opportunity for patient self-selection.

Pharmacists have a professional responsibility to ensure patients are knowledgeable about the appropriate selection of a Schedule II drug depending on symptoms, compatibility with other drugs being used, efficacy, possible adverse drug reactions, expected outcomes and what to do if outcomes are not achieved.

The following table provides a current list of Schedule II drugs. **Recent changes are indicated with <u>BOLD UNDERLINE</u>**. Examples of the Brand Names of products containing these drugs are also provided. Please note that the list of Brand Names is not necessarily inclusive and are provided for reference only.

Chemical name as it appears in the Schedule	Examples of common Brand Names
acetaminophen and ibuprofen in oral, fixed-dose combinations, in package sizes	·
containing either more than 20,000 mg of acetaminophen or more than 6,000 mg	
of ibuprofen	
acetarsol	
acetylcysteine	
acetylsalicylic acid (ASA) and its salts (in oral preparations containing 80 mg or	Asaphen Chewable 80mg
less per dosage unit and intended for pediatric use OR rectal preparations	Asatab 80mg Chewable
containing 150 mg or less per dosage unit, in package sizes containing no more	<b>3</b>
than 1.92 g of acetylsalicylic acid)	
adiphene and its salts for parenteral use	
allethrins	
amylocaine and its salts (for ophthalmic or parenteral use)	
anisotropine and its salts	
anthralin	
antihemophilic factor, human	
antipyrine (except otic preparations)	
apomorphine and its salts	
arginine and its salts	
artemisia, its preparations, extracts and compounds (except in trace amounts in	
homeopathic preparations)	
belladonna alkaloids and their salts and derivatives (except in preparations for	
topical use or in trace amounts in homeopathic preparations)	
benoxinate hydrochloride (oxybuprocaine) for ophthalmic or parenteral use	
bentiromide	
benzalkonium and its salts (liquid preparations in concentrations greater than 2%)	
benzethonium chloride (liquid preparations in concentrations greater than 1%)	
benzocaine and its salts (for ophthalmic or parenteral use)	
benzyl benzoate	
boric acid and its salts (in preparations for systemic or ophthalmic use in	
concentrations greater than 2%, except in contact lens solutions intended to be	
rinsed off prior to insertion in the eye.)	
buclizine	
bufexamac	
bupivacaine and its salts (for ophthalmic or parenteral use)	
butacaine and its salts (for ophthalmic or parenteral use)	
calcium disodium edentate	
camphor (in oleaginous vehicles and in liquid forms in concentrations greater than	
11%)	
cantharides, their preparations and derivatives	
charcoal (activated) for use in poisoning treatment	
chloroprocaine and its salts (for ophthalmic or parenteral use)	
cholecystokinin	
cholera vaccine (oral, inactivated) when used for prophylaxis against Traveler's	Dukoral
Diarrhea due to enterotoxigenic escherichia coli [ETEC]	
choline bitartrate (for parenteral use)	

Chemical name as it appears in the Schedule	Examples of common Brand Names
chymopapain (for parenteral use)	
chymotrypsin (for ophthalmic and parenteral use)	
cinchocaine (dibucaine) and its salts (for opthalmic or parenteral use)	
clidinium and its salts	
clobetasone butyrate (when sold in a concentration of 0.05% clobetasone butyrate	Specto EczemaCare Medicated Cream
in cream preparations for topical use on the skin)	Openio Edzernadare Midaldatea Ordani
coal tar (in concentrations greater than 10%)	Odans Liquor Carbonis Detergens
codeine and its salts (in preparations exempted from the Regulations to the	- Gadilo Elquoi Galborilo Botorgorio
Controlled Drugs and Substances Act – i.e. not more than 8 mg or its equivalent of	
codeine phosphate per solid form or not more than 20 mg or its equivalent of	
codeine phosphate per 30 mL in a liquid preparation)	
4,	
( <u>Please Note</u> : These preparations should be stored in	
accordance with the Regulations to the Controlled Drugs	
and Substances Act)	
collagenase (as a debriding agent)	
crotamiton	Eurax
cyclandelate	
cyclazocine and its salts	
cyclomethacaine and its salts (for ophthalmic or parenteral use)	
cyclopentamine and its salts	
cyclopentolate and its salts (except in products for ophthalmic or parenteral use)	
cyproheptadine and its salts	
desoxyribonuclease [pancreatic dormase]	
dextrose (sclerosing agents)	
diclofenac diethylamine when sold as a single medicinal ingredient for topical use	Voltaren Emulgel Extra Strength products (in package
on the skin for not more than 7 days - in concentrations greater than 1.16% and	sizes greater than 112 gm)
less than or equal to 2.32% - in package sizes containing greater than 2.6g of	3 ,
diclofenac diethylamine	
dicyclomine and its salts (except for topical use and lozenges)	
dihydroquinidine and its salts (except phenylbarbiturate)	
diiodohydroxyquin (for topical use on the skin)	
dimenhydrinate and its salts (for parenteral use)	Gravol IM injection
diperodon and its salts (except for topical use)	
diphenhydramine and its salts and preparations (for parenteral use)	
diphenhydramine and its salts and preparations (for topical use in concentrations of	
greater than 2%)	
diphtheria toxoid	
dyclonine (except for topical use on mucous membranes)	
ephedrine and its salts in single entity products (in preparations containing no more	
than 8 mg per unit dose, with a label recommending no more than 8 mg/dose or 32	
mg/day and for use for not more than 7 days, and indicated for nasal congestion)	
epinephrine and its salts (in pre-filled syringes intended for emergency	Epipen / Epipen Jr.
administration by injection in the event of anaphylactic reactions to allergens.)	
esdepallethrin/piperonyl butoxide	
ethanolamine oleate	
ethoheptazine and its salts	
ethyl chloride (except in trace amounts)	
fibrin	
fibrinolysin	
fluticasone propionate, when sold for the treatment of allergic rhinitis in a nasal	
spray that delivers 50 mcg/spray for those 18 years of age and older, in package	
sizes containing more than <b>360</b> metered sprays	
gentian violet (for application to skin or mucous membranes)	
glucagon	
glycopyrrolate and its salts	
hemophilus influenzae type B vaccine	
heparin and its salts (except for topical use)	
hepatitis B pediatric vaccine	Obtaine
herpes zoster (shingles) vaccine (non-live recombinant)	Shingrix
histamine and its salts (except for topical use)	

Chemical name as it appears in the Schedule	Examples of common Brand Names
homatropine and its salts (for oral use in concentrations of 2 mg or less per dosage	
unit)	
human insulin	
human papillomavirus vaccine	
hyaluronic acid and its salts (preparations in concentration of 5% or more)	
hyaluronidase	
hydroquinone (topical preparations)	
hydroxyephedrine and its salts	
hyoscine and its salts and derivatives (except for parenteral use)	Buscopan tablets
hyoscyamine and its salts and derivatives (except for topical use)	
influenza vaccine	
insulin	
iodinated glycerol	
iodine and its salts and derivatives (except in topical preparations or in oral doses	
of 1 mg or less per day)	
iodochlorhydroxyquin (for topical use)	
ipecac & its extracts & derivatives (when used as an emetic)	
iron and its salts and derivatives (in preparations with more than 30mg elemental	Feramax products
iron per unit or 5ml liquid)	Neo-Fer / Neo-Fer Cf
	Palafer capsules / suspension / Palafer CF
30mg elemental iron equals:	
ferrous ascorbate 250mg	
ferrous fumarate 92mg	
ferrous gluconate 257mg	
ferrous sulphate 150mg	
levargorphane and its salts	
levonordefrine	
levonorgestrel (when sold in concentrations of 0.75 mg per oral dosage unit	
(except when labelled to be taken as a single dose of 1.5 mg and in package sizes	
containing no more than 1.5 mg levonorgestrel, packaged and labelled for	
emergency contraception.))	
lidocaine and its salts (for ophthalmic or parenteral use or topical use on mucous	Xylocaine Viscous
membranes except lozenges)	
lindane	
loperamide and its salts in products marketed for pediatric use – under 12 years of	
age	
magnesium sulphate (for parenteral use)	
mannitol and its salts	
measles vaccine	
meningococcus vaccine	
mepivacaine and its salts (for ophthalmic or parenteral use)	
metathoheptazine and its salts	
methantheline	
methdilazine and its salts	
methenamine and its salts (except for topical use)	
metheptazine and its salts	
methocarbamol (for parenteral use)	
methyl salicylate (in liquid dosage forms in concentrations greater than 30%	
methylene blue (for parenteral use)	
monobenzone	
monoethanolamine oleate	
mumps vaccine	
naloxone hydrochloride injection, when indicated for emergency use for opioid	
overdose	
overdose naloxone hydrochloride nasal spray, when indicated for emergency use for opioid	
overdose naloxone hydrochloride nasal spray, when indicated for emergency use for opioid overdose	
naloxone hydrochloride nasal spray, when indicated for emergency use for opioid	

<sup>&</sup>lt;sup>1</sup> Available on the Standards, Guidelines, Policies and Positions page of the NLPB website: <a href="https://nlpb.ca/pharmacy-practice/standards-guidelines-policies/">https://nlpb.ca/pharmacy-practice/standards-guidelines-policies/</a>
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Chemical name as it appears in the Schedule	Examples of common Brand Names
naphazoline and its salts (in nasal preparations for pediatric use)	
niacin [nicotinic acid] (in extended release formulations)	
nitroglycerin (sublingual immediate release dosage forms)	Nitrolingual Pumpspray
	Nitrostat 0.3mg / 0.6mg tablets
norepinephrine and its salts [levarteronol, noradrenaline]	
omeprazole or its salts, when sold for the 14-day treatment for frequent heartburn	Heartburn Control (Apotex)
at a daily dose of 20mg, in package sizes of no more than 280mg of omeprazole	Omep
oxymetazoline and its salts (in nasal preparations for pediatric use)	
oxyquinoline	
paroxypropione	
pentagastrin and its salts	
permethrin and its derivatives	<ul> <li>Kwellada-P cream rinse / lotion</li> <li>Nix cream rinse / Nix dermal cream</li> </ul>
pertussis vaccine	
phenol (preparations with concentrations greater than 20%)	
phenoxybenzamine and its salts	
phenylephrine and its salts and preparations (in nasal preparations in	
concentrations of 2.5% or less for pediatric use)	
physostigmine salicylate (for oral or topical use)	
piperazine and its salts	
pneumococcal polysaccharide vaccine	
pneumococcal 7-valent conjugate vaccine	
pneumococcal 13-valent conjugate vaccine	
poliomyelitis vaccine	
polyacrylamide	
potassium salts (in preparations containing greater than 5 mmol per dose)	
povidone-iodine (vaginal preparations, except in concentrations of 5 % or less	
pramoxine and its salts (for ophthalmic or parenteral use)	
prilocaine and its salts (for ophthalmic or parenteral use)	
procaine and its salts (for ophthalmic or parenteral use)	
promethazine and its salts (except for topical use)	
propantheline and its salts	
proparacaine and its salts (for ophthalmic or parenteral use)	
propylhexidine	
protamine and its salts	
pseudoephedrine and its salts and preparations in single entity products	
pyrantel and its salts	Combantrin products
pyrethrins	
pyrethrins/piperonyl butoxide	R&C Shampoo with Conditioner
pyrivinium and its salts	
racemethionine	
rose bengal	
rotavirus vaccine	
rubella vaccine	
rue and its preparations and extracts	
salicylic acid and its salts (in topical preparations in concentrations greater than	
40%)	
silver nitrate	
sincalide	
sodium acetate (for parenteral use)	
sodium biphosphate (for parenteral use)	
sodium chloride (single ingredient solutions for parenteral or ophthalmic use in	
concentrations greater than 0.9%) [Does not apply to contact lens products	
intended to be rinsed off prior to insertion in the eye.]	
sodium citrate (for parenteral use)	
sodium iodide (for sclerosing)	
sodium phosphate (for parenteral use)	

Chemical name as it appears in the Schedule	Examples of common Brand Names
sodium picosulphate for oral purgatives, 10 mg per pack (when found in	Pico-Salex
preparations with magnesium oxide 3.5g and citric acid 12g)	Purg-Odan
sodium tetradecyl sulphate	
stramonium, its preparations, extracts and compounds	
streptokinase (as a debriding agent)	
strontium and its salts (for parenteral use)	
sutilains	
tetanus toxoid	
tetracaine and its salts (for ophthalmic and parenteral use)	
tetrahydrozoline (in nasal preparations for pediatric use)	
thrombin	
thyroglobulin	
thyrotropin	
triamcinolone acetonide in an aqueous nasal spray that delivers 55mcg per	
metered spray for adults and children 12 years of age and older, in package sizes	
containing more than 120 metered sprays	
urea in topical preparations in concentrations greater than 25%	
vaccines. which are part of a routine immunization program in most/all provinces	
and territories	
vitamins (any parenterals not included in Schedule I)	
xylometazoline and its salts (in nasal preparations for pediatric use (0.05%))	
xylose	

#### SCHEDULE III - PHARMACY SALE ONLY

Schedule III drugs may present risks to certain populations in self-selection. Although available without a prescription, these drugs are to be sold from the self-selection area of the pharmacy which is operated under the direct supervision of the pharmacist. Such an environment is accessible to the patient and clearly identified as the "professional services area" of the pharmacy. The pharmacist is available, accessible and approachable to assist the patient in making an appropriate self-care selection.

The following table provides a current list of Schedule III drugs. **Recent changes are indicated with BOLD UNDERLINE**. Examples of the Brand Names of products containing these drugs are also provided. Please note that the list of Brand Names is not necessarily inclusive and are provided for reference only.

Chemical name as it appears in the Schedule	Examples of common Brand Names
acetaminophen and ibuprofen in oral, fixed-dose combinations, in	
package sizes containing 20,000 mg or less of acetaminophen and	
6,000 mg or less of ibuprofen	
acetaminophen in sustained release formulations (in strengths of greater	Tylenol Arthritis Pain 8H (100's, 170's, 200's)
than 650 mg per unit or in package sizes of more than 50 units)	Tylenol Muscle & Body 8H (72's, 110's)
acetylsalicylic acid and its salts (in products intended for oral adult use in	Aspirin 81mg Daily Low Dose / 81mg Quick Chews
strengths of 81mg per dosage unit and 650mg or greater per dosage	Entrophen 81mg Daily Low Dose / 81mg Chewable
unit, and in rectal preparations containing more than 150 mg per dosage	Novasen 650m
unit)	Novasen osom
amylocaine and its salts (in preparations for topical use on mucous	
membranes except lozenges)	
anethole trithione	
antazoline and its salts	Refresh Eye Allergy Relief
antipyrine (for otic use)	Auralgan
bacitracin and its salts and derivatives (for ophthalmic use)	·
benzonatate	
bisacodyl and its salts (except when sold in concentrations of 5mg or	
less per oral dosage unit in package sizes containing no more than	
105mg of bisacodyl or 10mg or less per rectal dosage unit/suppository	
in package sizes containing no more than 50mg of bisacodyl)	
brompheniramine and its salts as a single entity for the treatment of	
allergies	
bupivacaine and its salts (for topical use on mucous membranes except	
lozenges)	
calcium polycarbophil	
carbinoxamine and its salts	
cerapon	
cetirizine and its salts (in concentrations of 10 mg equivalent to 8.5 mg	Reactine Children's Liquid / Fast Melt
or less of cetirizine base per dosage unit) in products marketed for	
pediatric use (under 12 years of age)	
chlophedianol and its salts	
chloroprocaine and its salts (for topical use on mucous membranes	
except lozenges)	
chlorzoxazone and its salts	Acetazone Forte
cimetidine and its salts (in concentrations of 100 mg or less per dosage	
unit)	
clemastine and its salts	
clotrimazole and its salts (in preparations for vaginal use)	Canesten vaginal products
	Clotrimaderm vaginal creams
danthron	
dehydrocholic acid and its salts	
desloratadine and its salts and preparations in products marketed for	Aerius Kids Syrup
pediatric use (under 12 years of age)	

Chemical name as it appears in the Schedule	Examples of common Brand Names
dexbrompheniramine and its salts	
dexchlorpheniramine and its salts	
dextromethorphan and its salts	<ul> <li>Balminil DM products / Night-Time</li> <li>Benylin All in One Cold &amp; Flu products / "Cough" products / DM products / Mucus &amp; Phlegm Relief products / Tickly Throat Cough</li> <li>Buckley's Complete products / Daytime &amp; Nighttime Cold and Sinus</li> <li>Coricidin Cold &amp; Flu Extra Strength</li> <li>Dimetapp DM Cough &amp; Cold</li> <li>Jack &amp; Jill Cough &amp; Cold Liquid</li> <li>NeoCitran Extra Strength Total Cold / Ultra Strength Total Flu</li> <li>Robitussin Complete / Cough &amp; Cold / Cough Control / CoughGels / Total Cough, Cold &amp; Flu</li> <li>Tylenol "Cough" products / Cold Daytime &amp; Nighttime / Plus Mucus Relief / Flu Daytime &amp; Nighttime</li> <li>Vicks Children's Nyquil Cold &amp; Cough / Dayquil Cold &amp; Flu / Dayquil "Cough" products / Nyquil Cold &amp; Flu / Nyquil "Cough" products</li> </ul>
diclofenac diethylamine when sold as a single medicinal ingredient for topical use on the skin for not more than 7 days - in concentrations greater than 1.16% and less than or equal to 2.32% - in package sizes containing no more than 2.6g of diclofenac diethylamine	Voltaren Emulgel Extra Strength (in package sizes less than 112gm)
(Please Note: Pharmacists are advised that in areas here there is evidence of abuse or a particular concern about abuse, dimenhydrinate products should not be located in a self-selection area of the pharmacy.  Please also see NLPB Guidance regarding the sale of dimehydrinate <sup>2</sup> )	<ul> <li>Gravol products</li> <li>generic and store brands</li> </ul>
dimethothiazine	
dimeticone 100 cSt solution 50% w/w for topical use in the treatment of head lice	NYDA
diphenhydramine and its salts and preparations for topical use in concentrations of 2% or less when sold in containers of greater than 300mg of diphenhydramine hydrochloride	<ul><li>Benadryl cream / spray</li><li>Calamine lotion + antihistamines</li></ul>
diphenhydramine and its salts and preparations (except for parenteral use)  diphenylpyraline doxylamine and its salts (except those sold for nausea and vomiting of pregnancy)	<ul> <li>Advil Cold &amp; Flu / Nighttime</li> <li>Aleve Nighttime</li> <li>Benadryl oral products</li> <li>Benylin All in One Cold &amp; Flu Night</li> <li>Buckley's Night Time Mixture</li> <li>Jack &amp; Jill Bedtime</li> <li>Neo Citran Total Cold Night</li> <li>Nytol products except Nytol Natural Source</li> <li>Sinutab Sinus Nighttime Extra Strength</li> <li>Sleep-Eze products except V Natural</li> <li>Tylenol Children's Cold &amp; Cough Nighttime / Complete Cold, Cough &amp; Flu Nighttime</li> <li>Unisom products except Unisom-2</li> <li>ZZZ-Quil</li> <li>Buckley's Complete Nighttime / Nighttime Cold &amp; Sinus</li> <li>NyQuil Cold &amp; Flu Nighttime / Complete Cold &amp; Flu / Cough / Sinus</li> <li>Robitussin Complete Nighttime</li> </ul>

<sup>&</sup>lt;sup>2</sup> Available on the Drug Schedules page of the NLPB website: <a href="https://nlpb.ca/pharmacy-practice/provincial-drug-schedules/">https://nlpb.ca/pharmacy-practice/provincial-drug-schedules/</a>

Chemical name as it appears in the Schedule	Examples of common Brand Names
ephedrine and its salts in combination products (in preparations	Examples of common brain names
containing no more than 8 mg per unit dose, with a label recommending	
no more than 8 mg/dose or 32 mg/day and for use for not more than 7	
days, and indicated for nasal congestion) [Note: Pharmacists are	
advised that in areas where there is evidence of abuse or particular	
concern about abuse, ephedrine products should NOT be located in a	
self- selection area of the pharmacy]	
esomeprazole or its salts, when sold for the 14-day treatment for	Nexium 24HR
frequent heartburn at a daily dose of 20 mg, in package sizes of no more	TOMAIN E ITH
than 280 mg of esomeprazole	
famotidine and its salts, when sold in concentrations of 20 mg or less per	
oral dosage unit and indicated for the treatment of heartburn, in package	
sizes containing more than 600 mg of famotidine	
fexofenadine hydrochloride (in products marketed for paediatric use	
(under 12 years of age))	
fluconazole when sold in a concentration of 150 mg per oral dosage unit	CanesOral
and indicated for the treatment of vaginal candidiasis, in package sizes	Diflucan One
containing no more than 150 mg of fluconazole	Monicure
fluticasone propionate, when sold for the treatment of allergic rhinitis in a	
nasal spray that delivers 50 mcg/spray for those 18 years of age and	
older, in package sizes containing no more than <b>360</b> metered sprays	
fractar	
glyceroargentinate	
gramicidin and its salts and derivatives (for ophthalmic use)	Optimyxin eye/ear drops
	Polysporin Eye and Ear drops
haloprogin	
ibuprofen and its salts, containing 400mg or less per oral dosage unit	Advil single-entity 200mg products (more than 90 tablets)
(when sold in package sizes exceeding 18,000mg)	Advil single-entity 400mg products (more than 45 tablets)
	Motrin single-entity 200mg products (more than 90 tablets)
	Motrin single-entity 300mg products (more than 60 tablets)
	Motrin single-entity 400mg products (more than 45 tablets)
ibuprofen or its salts, when sold in a modified-release oral dosage form	Advil 12 Hour
that provides 600 mg or less per dosage unit	7 Advii 12 Hodi
isopropyl myristate in concentration of 50% (for use in the treatment of	Resultz
head lice)	1 TOOLILE
lactulose	
levonorgestrel (when sold in concentrations of 0.75 mg per oral dosage	Plan B
unit to be taken as a single dose of 1.5 mg, packaged and labelled for	
emergency contraception, in package sizes containing no more than 1.5	
mg of levonorgestrel)	
lidocaine and its salts (for otic use)	Polysporin Plus Pain Relief ear drops
lidocaine and prilocaine (eutectic mixture)	EMLA cream and patches
loratadine and its salts and preparations in products marketed for	Claritin Kids Syrup
pediatric use (under 12 years of age)	- Glantin Hudo Gyrup
meclizine and its salts (when sold in concentrations of 25mg or less per	
dosage unit)	
mepivacaine and its salts (for topical use on mucous membranes except	
lozenges)	
methocarbamol (except for parenteral use)	Motrin Platinum Muscle and Body
· · (· · · · · · · · · · · · · · · · ·	Robax Platinum / Robaxacet / Robaxin / Robaxin 750 / Robaxisal
	Tylenol Extra Strength Back Pain
miconazole and its salts (for vaginal use)	Monistat vaginal products
mineral tar (except shampoos with concentrations less than 5%)	
Timoral tar (except sharipees with concentrations less than 576)	I

Chemical name as it appears in the Schedule	Examples of common Brand Names
naproxen and its salts, in preparations containing the equivalent of 200 mg of naproxen base* per oral dosage unit (when sold in products labelled with a recommended maximum daily dose equivalent to 400 mg of naproxen base, and in package sizes exceeding the equivalent of 6,000 mg of naproxen base.)  [*220 mg naproxen sodium approximately equivalent to 200 mg	Aleve (100 tablets or more)
naproxen base] nystatin and its salts and derivatives (in topical preparations for use on	Nyaderm cream
the skin) orphenadrine citrate	generic brands
oxethazine	
oxybuprocaine and its salts (for topical use on mucous membranes except lozenges)	
phenyltoloxamine and its salts	
polymyxin B and its salts and derivatives (for ophthalmic use)	Optimyxin eye/ear drops     Polysporin Eye and Ear drops
pramoxine and its salts (for topical use on mucous membranes except lozenges)	
prilocaine and its salts (for topical use on mucous membranes except lozenges)	
procaine and its salts (for topical use on mucous membranes except lozenges)	
promethazine and its salts (for topical use)	
proparacaine and its salts (for topical use on mucous membranes except lozenges)	
(Please Note: Pharmacists are advised that in areas where there is evidence of abuse or particular concern about abuse, pseudoephedrine products should NOT be located in a self-selection area of the pharmacy)  ranitidine and its salts, when sold in concentrations of 150mg or less per	<ul> <li>Advil Cold &amp; Sinus products / Children's Advil Cold / Cold &amp; Flu</li> <li>Aerius Dual Action 12 Hour</li> <li>Allegra D</li> <li>Benadryl Total</li> <li>Benylin All-in-One Cold &amp; Flu / Cough &amp; Chest Congestion / "D" products</li> <li>Buckley's Cough &amp; Cold / Cough, Cold &amp; Flu / Cold &amp; Sinus</li> <li>Claritin Allergy + Sinus</li> <li>NeoCitran Flu with Mucous Relief / Flu Night</li> <li>Reactine Complete</li> <li>Robitussin Cough &amp; Cold / Total Cough, Cold &amp; Flu</li> <li>Sinutab Sinus / Sinus &amp; Allergy</li> <li>Sudafed Head Cold &amp; Sinus / Sinus / Sinus Advance</li> <li>Tylenol Cold &amp; Sinus / Complete Cold, Cough &amp; Flu</li> <li>Tylenol Children's Cold / Cold &amp; Cough / Cold &amp; Stuffy Nose / Complete Cold, Cough &amp; Fever</li> </ul>
oral dosage unit & indicated for the treatment of heartburn, in package sizes containing more than 4,500mg of ranitidine	
sodium biphosphate (cathartics) sodium cromoglycate (in solutions in concentrations of 2% or less for	Cromolyn eye drops
ophthalmic or nasal use)	Opticrom     Rhinaris-CS Anti-Allergic Nasal Mist
sodium phosphate (cathartics)	
tetracaine and its salts (for topical use on mucous membranes except lozenges)	
tioconazole and its salts (in preparations for vaginal use) triethanolamine oleate	
triethanolamine salicylate (in concentrations greater than 20%)	
tripelennamine and its salts	
triprolidine	

Chemical name as it appears in the Schedule	Examples of common Brand Names
tyrothricine	
triamcinolone acetonide in an aqueous nasal spray that delivers 55mcg	Nasacort Allergy 24HR
per metered spray for adults and children 12 years of age and older, in	
package sizes containing no more than 120 metered sprays	
vegetable tar (except shampoos in concentrations of 5% or less)	

#### **UNSCHEDULED DRUG PRODUCTS**

Many drugs have been reviewed by the National Drug Scheduling Advisory Committee (NDSAC) but have not been assigned any place in the above schedules. As well, many drugs that are included in the schedules are included based on specific parameters such as strength and dosage form. That same drug in another strength or dosage form may be considered "Unscheduled".

Unscheduled drugs can be sold without professional supervision. Adequate information is available for the patient to make a safe and effective choice and labelling is deemed sufficient to ensure the appropriate use of the drug. These drugs may be sold from any retail environment.

The following table lists all drugs that are currently specifically recommended for unscheduled status by NDSAC. It is not intended to reflect all drugs not otherwise captured by the Provincial Drug Schedules. **Recent changes are indicated with BOLD UNDERLINE**. Examples of the Brand Names of products containing these drugs are also provided. Please note that the list of Brand Names is not necessarily inclusive and are provided for reference only.

Chemical name	Examples of common Brand Names
acetaminophen (in immediate release tablets, capsules, suppositories or	Atasol Forte
liquid)	Tempra products
	Tylenol products
acetaminophen in sustained release formulations (up to and including 650	Tylenol Arthritis Pain 8H
mg per unit, in package sizes containing no more than 50 units)	Tylenol Muscle & Body 8H
acetylsalicylic acid and its salts (in products for oral use in strengths of	Alka-Seltzer products
325mg and 500mg per dosage unit)	Anacin products
	Aspirin Regular / Extra Strength
	Novasen 325mg
bacitracin and its salts (for topical use)	Bacitin ointment
	Band Aid adhesive bandages plus antibiotic
	Ozonol Antibiotics Plus
	Polysporin Antibiotic ointment / Triple Antibiotic ointment /
	Complete
benzoyl peroxide (preparations of 5% or less as a single ingredient)	Advantage Acne Control Kit
	Benzagel products
	Clean & Clear Persa-Gel 5
	Clearasil Daily Clear Acne Treatment Cream BP Plus
	Continuous Control Acne Cleanser
	Deep Pore Acne Vanishing Treatment
	Emergency Acne Vanishing Facial Cleanser
	Spectro AcneCare Wash
bisacodyl and its salts (when sold in concentrations of 5mg or less per oral	
dosage unit in package sizes containing no more than 105mg of	
bisacodyl or 10mg or less per rectal dosage unit/suppository, in package	
sizes containing no more than 50mg of bisacodyl)	
brompheniramine and its salts in combination products for the relief of	Dimetapp liquid
cough and cold symptoms	Robitussin Children's Cold
butenafine (1% cream)	
cetirizine and its salts (in concentrations of 10 mg equivalent to 8.5 mg or	Reactine 10mg tablets (single-entity)
less of cetirizine base per dosage unit) in products marketed for adult use	the state of the state (state)
(12 years of age and older)	
charcoal (activated) except for use in poisoning treatment	
chloral hydrate (for topical use)	
chlorpheniramine and its salts and preparations	Benylin Cold & Sinus Night
	Chlor-Tripolon
	Tylenol Extra Strength Sinus Nighttime
cinnamedrine	

Chemical name	Examples of common Brand Names
clotrimazole and its salts (in preparations for topical use)	Canestan 1% Topical Cream
coal tar (in concentrations up to and including 10%)	
desloratidine and its salts and preparations (in products marketed for adult use – 12 years and older)	Aerius tablets
diclofenac diethylamine when sold as a single medicinal ingredient for topical use on the skin in concentrations of not more than 1.16% for not more than 7 days	Voltaren Emulgel regular strength products
diphenhydramine and its salts and preparations for topical use in concentrations of 2% or less when sold in containers of 300 mg or less of diphenhydramine hydrochloride	Benadryl Itch Relief Stick
docosanol 10% for topical use	Abreva ointment
docusate and its salts	<ul> <li>Colace products</li> <li>Senokot S</li> <li>Soflax products</li> </ul>
famotidine and its salts, when sold in concentrations of 20mg or less of famotidine per oral dosage unit and indicated for the treatment of heartburn, in package sizes containing no more than 600mg of famotidine	Pepcid AC 10mg (60 tablets or less)     Pepcid AC Maximum Strength 20mg (30 tablets or less)
fexofenadine HCl (in products marketed for adult use – 12 years and older)	Allegra 12 hour / 24 hour
gramicidin and its salts (for topical use)	Polysporin Cream / Complete / Plus Pain Relief / For Kids / Triple Antibiotic Ointment
guaifenesin	<ul> <li>Balminil Expectorant</li> <li>Benylin "E" / Mucous &amp; Phlegm</li> <li>Robitussin Mucus &amp; Phlegm</li> </ul>
ibuprofen and its salts, containing 400mg or less per oral dosage unit (when sold in package sizes of up to 18,000mg)	<ul> <li>Advil single-entity 200mg products (90 tablets or less)</li> <li>Advil single-entity 400mg products (45 tablets or less)</li> <li>Motrin single-entity 200mg products (90 tablets or less)</li> <li>Motrin single-entity 300mg products (60 tablets or less)</li> <li>Motrin single-entity 400mg products (45 tablets or less)</li> </ul>
ketoconazole and its salts (as a shampoo)	Nizoral 2% shampoo
lidocaine and its salts (for topical use on the skin, including lozenges)	<ul> <li>After Sun Soothing Spray</li> <li>Afterburn</li> <li>Bactine First Aid Spray</li> <li>Band-Aid Brand Antiseptic Wash</li> <li>Ozonol Antibiotics Plus</li> <li>Polysporin Complete / For Kids / Plus Pain Relief</li> <li>Solarcaine First Aid Lidocaine Spray / Medicated First Aid Lotion / Medicated Lidocaine Gel</li> </ul>
loperamide and its salts in products marketed for adult use – 12 years and older)	Imodium products
loratadine and its salts and preparations in products marketed for adult use (12 years of age and older)	Claritin adult products (single entity)
miconazole and its salts (for topical use)	<ul><li>Micatin Cream / Spray</li><li>Monistat Derm Cream</li></ul>
minoxidil, when sold in preparations for topical use in adults in concentrations of 5% or less, for human use only	
naphazoline and its salts (in nasal preparations for adult use and in ophthalmic products)	<ul> <li>Albalon</li> <li>Clear Eyes / Clear Eyes Allergy / Extra Strength Redness Relief</li> <li>Naphcon A</li> <li>Opti-Tears Allergy</li> <li>Refresh Redness Relief</li> <li>Soothe Allergy / Redness</li> <li>Visine for Allergy with Antihistamine</li> </ul>

Chemical name	Examples of common Brand Names
naproxen and its salts, in preparations containing the equivalent of 200 mg of naproxen base* per oral dosage unit (when sold in products labelled with a recommended maximum daily dose equivalent to 400 mg of naproxen base, and in package sizes of the equivalent of 6,000 mg of naproxen base or less.)	• Aleve (24's)
[*220 mg naproxen sodium approximately equivalent to 200 mg naproxen base]	
oxiconazole (1% for topical use)	
oxymetazoline (in nasal preparations for adult use and in ophthalmic products)	<ul> <li>Claritin Allergy Decongestant nasal spray</li> <li>Dristan Long Lasting nasal sprays</li> <li>Drixoral nasal sprays</li> </ul>
pheniramine	<ul> <li>Dristan Nasal Mist</li> <li>Naphcon-A</li> <li>NeoCitran Cold &amp; Sinus Night / Cold &amp; Sore Throat Night</li> <li>Opti-Tears Allergy</li> <li>Soothe Allergy</li> <li>Visine for Allergy with Antihistamine</li> </ul>
phenylephrine and its salts and preparations (for oral use, in nasal preparations for adults and in ophthalmic preparations in concentrations of 2.5% or less)	<ul> <li>Benylin Cold &amp; Sinus / Cold &amp; Sinus Plus caplets</li> <li>Dayquil Sinus</li> <li>Dimetapp liquid</li> <li>Dristan Tablets / Nasal Mist</li> <li>NeoCitran Cold &amp; Congestion / Cold &amp; Sinus Night / Cold &amp; Sore Throat Night</li> <li>Robitussin's Children's Cold</li> </ul>
polyethylene glycol (topical administration)	
polyethylene glycol 3350 as a single ingredient oral product indicated as a laxative to treat occasional constipation	<ul> <li>Clearlax</li> <li>Lax-A-Day</li> <li>Pegalax</li> <li>Purelax</li> <li>Restoralax</li> </ul>
polymyxin and its salts and derivatives (topical)	<ul> <li>Band aid adhesive bandages plus antibiotic</li> <li>Ozonol Antibiotics Plus</li> <li>All Polysporin topical products</li> </ul>
pramoxine and its salts for topical application on the skin and including lozenges	<ul> <li>Aveeno Anti-Itch Lotion</li> <li>Gold Bond Anti-Itch lotion / Medicated Anti-Itch cream</li> <li>Polysporin Itch Relief / Poly-To-Go</li> </ul>
propylene glycol (topical application)	
pyrilamine	Menstrual Midol Complete / PMS Midol Complete
ranitidine and its salts, when sold in concentrations of 150mg or less per oral dosage unit and indicated for the treatment of heartburn, in package sizes containing no more than 4500mg of ranitidine	<ul><li>Zantac 75</li><li>Zantac 150</li></ul>
sodium tartrate tetrahydrozoline (for ophthalmic use and in nasal preparations for adults)	Clear Eyes Triple Action Relief     Visine Allergy / Multi-Symptom / Original / Red Eye
tioconazole and its salts (in preparations for topical use)	
triethanolamine salicylate (in concentrations up to and including 20%) [trolamine]	<ul><li>Aspercreme</li><li>Myoflex products</li></ul>
xylometazoline and its salts (in nasal preparations for adults(0.1%))	<ul><li>Balminil Decongest spray</li><li>Otrivin Cold &amp; Allergy products</li></ul>

## APPENDIX A PROVINCIAL REGULATIONS

### **DRUG SCHEDULES**

- 13. (1) The board may adopt the drug schedules established under the National Association of Pharmacy Regulatory Authorities National Drug Schedules System.
- (2) Notwithstanding subsection (1), a pharmacist in charge may, where he or she considers it necessary, exercise a higher degree of control over a particular drug than what is contemplated in the schedules.

## APPENDIX B SCHEDULING CRITERIA

#### FACTORS FOR THE INCLUSION OF DRUGS IN SCHEDULE I

- 1. Indications for use of the drug are identifiable only by the practitioner. Diagnosis of the indication requires intervention by the practitioner before the drug is used.
- 2. Use of the drug requires adjunctive therapy or evaluation. Adjunctive therapy could include other drugs, non-pharmacologic measures, or specialized drug delivery devices. Evaluation could include indicated laboratory or clinical assessments.
- 3. Use of the drug may produce dependency. The drug may cause addiction or become habit forming. Control of access and duration of therapy by a health care professional is required.
- 4. Serious adverse reactions to the drug are known to occur or have a recognized potential to occur at normal therapeutic dosage levels. Adverse experiences require special monitoring or intervention by a health care professional.
- 5. There exists a narrow margin of safety between the therapeutic and toxic dosages of the drug, either in the general population, or in identified subpopulations, or in patients with multiple medical problems. Safe use requires the involvement and intervention of a health care professional.
- 6. Serious interactions of the drug are known to occur. Such interactions (drug-drug, drug-food, drug-disease) require special monitoring or intervention by a health care professional.
- 7. Use of the drug has contributed to, or is likely to contribute to, the development of resistant strains of microorganisms. Appropriate use, and/or the decision to continue treatment, requires evaluation by the practitioner.
- 8. The mechanism of action of the drug is known but the consequences of widespread use are not adequately established. Unexpected effects of the drug must be evaluated and reported by a health care professional.
- 9. The therapeutic effects of a newly released drug are based on new or unknown mechanisms of action, but the consequences of widespread use are not adequately established. Close monitoring of the patient is required by a health care professional for unanticipated effects.

### FACTORS FOR THE INCLUSION OF DRUGS IN SCHEDULE II

- The initial need for a drug is normally identified by the practitioner, in addition chronic, recurrent, or subsequent therapy must be monitored by the pharmacist. A prescription should not be required to obtain a drug if the patient can understand directions for continued use through the intervention of the pharmacist. Therefore, the patient should have access to the drug for subsequent treatment and use following the first diagnosis and prescription by the practitioner. This collaborative approach enhances patient care.
- The drug must be readily available under exceptional circumstances when a prescription is not practical. Such a drug might
  be required for a serious medical situation and the patient should have access to it to prevent a possible health emergency.
  An example of such an exceptional circumstance is availability of injectable epinephrine for anaphylactic reactions.
- 3. The drug is intended for administration in a health care setting or under direction of a health care professional, or is in an injectable dosage form and is not otherwise included in Schedule I. Examples include preoperative or diagnostic agents and products used for immunization or desensitization.
- 4. Evidence of abuse of the drug has been reported, due to its inherent pharmacological action which has the potential for abuse. Monitoring by a health care professional is necessary.
- 5. The selection of the drug requires intervention by the pharmacist to confirm that an appropriate self-assessment has been made by the patient. Dosage form, for example, may be an important consideration.
- 6. Use of the drug may delay recognition or mask the symptoms of serious disease. Intervention by the pharmacist is necessary to ensure appropriate referral to the practitioner.
- 7. The drug may cause important adverse reactions, including allergies, or interacts with other drugs, foods, or disease states that cannot be adequately addressed through product labelling. Intervention by the pharmacist is necessary to assess patient risk to prevent such problems for an individual patient through interpretation and clarification of labelling.
- 8. Use of the drug requires reinforcement or an expansion of the directions for use, through pharmacist patient dialogue. Such reinforcement and expansion may include the explanation of the use of a drug delivery system.

- 9. The drug is a new ingredient for self-medication and monitoring by the pharmacist is necessary to facilitate observation and reporting of any unexpected event.
- 10. The maximum labelled dosage directions exceed the generally accepted or usual limits for Schedule III status.

#### FACTORS FOR THE INCLUSION OF DRUGS IN SCHEDULE III

- 1. The initial need for a drug is normally identified by the patient, physician, or pharmacist, but chronic, recurrent, or subsequent therapy can be monitored by the pharmacist.
- The maximum recommended duration of use of the drug is limited and specified on the product label. The pharmacist is available to explain that the consequences of not following the period of use may be serious and that persistence of symptoms may suggest an underlying ailment.
- The maximum recommended duration of use of the drug is not specified on the label, but continued use may delay recognition or mask the symptoms of serious disease. The pharmacist is available to help in interpretation of symptoms, to assist in selection of alternative therapy, or to provide appropriate referral.
- 4. The drug is used to treat a persistent, chronic or recurring condition and the availability of the pharmacist to provide advice can promote appropriate use. The pharmacist should be available to direct the patient to a practitioner for assessment if the treatment period has been inappropriate or the therapy has been ineffective.
- 5. The drug is used for self-treatment of self-limiting ailments; however, where product selection has been identified as likely to cause patient confusion and the availability of the pharmacist to provide advice can promote appropriate use. Many product selections may be confusing for the patient. These choices are further complicated by the different forms of available therapy or dosage forms.
- 6. The drug demonstrates adverse effects, including allergies, or interacts with other drugs, foods, or disease states that can be identified in product labelling, but appropriate product selection and explanation of risk may require the advice of the pharmacist. For example, individuals taking a traditional monoamine oxidase inhibitor are aware that certain drugs should be avoided (e.g., cold products) but might require assistance in selecting a safe product to use.
- 7. The drug is a new ingredient for self-selected self-medication and the availability of the pharmacist to provide advice can promote appropriate use. The pharmacist is available to answer questions about this new ingredient.
- 8. The drug has inherent pharmacologic action which has the potential for non-medical use which may result in adverse patient outcomes.
- 9. The maximum labelled dosage directions exceed the generally accepted or usual limits for unscheduled status.

## APPENDIX C VACCINES AND IMMUNE GLOBULINS QUICK REFERENCE

SCHEDULE I	SCHEDULE II
<ul> <li>Bacillus Calmette-Guerin</li> <li>Cholera (when not used for prophylaxis against Traveler's Diarrhea due to enterotoxigenic escherichia coli [ETEC])</li> <li>Hepatitis A</li> <li>Hepatitis B Adult</li> <li>Herpes Zoster (shingles)</li> <li>Japanese Encephalitis</li> <li>Rabies</li> <li>Typhoid</li> <li>Varicella (chicken pox)</li> <li>Yellow Fever</li> <li>All Immune Globulins</li> </ul>	<ul> <li>Cholera (oral, inactivated) (when used for prophylaxis against Traveler's Diarrhea due to enterotoxigenic escherichia coli [ETEC])</li> <li>Diptheria toxoid</li> <li>Haemophilus influenzae Type B</li> <li>Hepatitis B Pediatric</li> <li>Human Papillomavirus</li> <li>Influenza</li> <li>Measles</li> <li>Meningococcus</li> <li>Mumps</li> <li>Pertussis</li> <li>Pneumococcal conjugate (7-valent)</li> <li>Pneumococcal conjugate (13-valent)</li> <li>Pneumococcal polysaccharide</li> <li>Poliomyelitis</li> <li>Rotavirus</li> <li>Rubella</li> </ul>
	> Tetanus toxoid