

Division of Workers' Compensation Pharmacy and Therapeutics Committee

January 20, 2021
12:30pm to 2:30pm



State of California
Gavin Newsom
Governor

Agenda

- **Welcome and Introductions**
George Parisotto, Administrative Director, DWC
- **Approval of Minutes from the October 21, 2020 Meeting**
Dr. Raymond Meister, Executive Medical Director, DWC
- **MTUS Drug List v8 – *Dr. Raymond Meister***
- **Discussion:**
 - Anti-Emetics – perphenazine & granisetron - *Kevin Gorospe, DWC Consultant*
 - RxCUI refresher - *Kevin Gorospe, DWC Consultant*
 - MTUS – RxCUI consolidation model using MTUS v8 - *Kevin Gorospe, DWC Consultant*
 - *Drug Ingredient name changes*
 - *Therapeutic Category name changes*
 - *Opioid 4-day supplies*
 - Reports/Research of interest (general discussion)
 - COVID vaccination (general discussion)
- **Public Comments**
- **Review of Committee Recommendations**
- **Adjourn**

Welcome and Introductions

George Parisotto

Administrative Director, DWC

Approval of Minutes

Dr. Raymond Meister

Executive Medical Director, DWC

MTUS Drug List v8

Dr. Raymond Meister

Executive Medical Director, DWC

Anti-Emetics – perphenazine & granisetron

J. Kevin Gorospe, PharmD
DWC Consultant

Perphenazine

- Old drug (approved in 1957)
- Initially used as an antipsychotic for treatment of schizophrenia
- Current approved indications
 - treatment of schizophrenia
 - treatment of severe behavioral or psychological symptoms of dementia
 - treatment of severe nausea/vomiting
- Previously added to MTUS list as “Exempt” – (R) Depressive Disorders

Granisetron

- Approved in 1994 (brand name KYTRIL)
- Originally approved for prophylaxis of acute emesis caused by cancer chemotherapy
- Current indications
 - For chemotherapy-induced nausea/vomiting (CINV) and chemotherapy-induced nausea/vomiting prophylaxis.
 - For radiation-induced nausea/vomiting prophylaxis.
 - For the treatment of post-operative nausea/vomiting (PONV)
- The FDA-approved product label recommends against granisetron for postoperative nausea/vomiting (PONV) due to lack of efficacy and risk of QT prolongation.

Drug Ingredient	Reference Brand Name	Dosage Form	Strength	MEDI-CAL PRICE (Generic)	TOTAL DAILY UNITS (TABLETS, CAPSULES, ML)	GENERIC COST PER DAY
perphenazine	Not Applicable	Tablet	8 MG	0.4851	3	1.4553
perphenazine	Not Applicable	Tablet	4 MG	0.3766	6	2.2596
perphenazine	Not Applicable	Tablet	16 MG	3.1194	1	3.1194
perphenazine	Not Applicable	Tablet	2 MG	0.3129	12	3.7548
granisetron hydrochloride	KYTRIL	Tablet	1 MG	1.8861	2	3.7722

EXEMPT ANTI-EMETIC DRUGS BY GENERIC COST PER DAY

Drug Ingredient	Reference Brand Name	Dosage Form	Strength	MEDI-CAL PRICE (Generic)	TOTAL DAILY UNITS (TABLETS, CAPSULES, ML)	GENERIC COST PER DAY
meclizine hydrochloride	ANTIVERT	Tablet, Chewable	25 MG	0.0376	4	0.1504
promethazine hydrochloride	PHENERGAN	Tablet	25 MG	0.049	4	0.196
promethazine hydrochloride	PHENERGAN	Tablet	50 MG	0.123	2	0.246
metoclopramide hydrochloride	REGLAN	Tablet	10 MG	0.0414	6	0.2484
ondansetron hydrochloride	ZOFRAN	Tablet	8 MG	0.1085	3	0.3255
dimenhydrinate	DRAMAMINE	Tablet	50 MG	0.0479	8	0.3832
metoclopramide hydrochloride	REGLAN	Tablet	5 MG	0.0377	12	0.4524
promethazine hydrochloride	PHENERGAN	Tablet	12.5 MG	0.06	8	0.48
ondansetron hydrochloride	ZOFRAN	Tablet	4 MG	0.0819	6	0.4914
meclizine hydrochloride	ANTIVERT	Tablet	25 MG	0.1651	4	0.6604
ondansetron	ZOFRAN ODT	Tablet, Orally Disintegrating	8 MG	0.3004	3	0.9012
promethazine hydrochloride	PHENERGAN	Syrup	6.25 MG/5 ML	0.0171	60	1.026
meclizine hydrochloride	ANTIVERT	Tablet	12.5 MG	0.1586	8	1.2688
prochlorperazine maleate	COMPAZINE	Tablet	10 MG	0.3866	4	1.5464
ondansetron	ZOFRAN ODT	Tablet, Orally Disintegrating	4 MG	0.2668	6	1.6008
prochlorperazine maleate	COMPAZINE	Tablet	5 MG	0.279	8	2.232

Committee Discussion

Public Comments

RxCUI Refresher

J. Kevin Gorospe, PharmD

DWC Consultant

RxCUI

- Intent is to provide clarity on which drug/dosage form/strength combinations are exempt/non-exempt on MTUS Formulary
- RxCUI is a publicly available (non-proprietary) coding system
- Has multiple layers, for example
 - Ingredient – base drug without “salt”
 - Precise Ingredient – drug/salt forms
 - Clinical Drug or Pack – drug ingredient, dosage form, strength
 - Clinical Dose Group – e.g. oral drug, oral pill, transdermal, etc.
- Accessible at U.S. National Library of Medicine – RxNAV [RxNav \(nih.gov\)](https://www.nlm.nih.gov/rxnav/)



String Search... [Q] [↺]

Welcome to RxNav

- RxNorm Graph
- RxNorm Properties
- NDC
- RxTerms
- Pill Images
- Class View
- Interaction View
- Status

Views

- Classic
- Simple
- Table

Filters

- Human
- Vet
- Pres
- Single
- Group Form

Links

- Drug Label
- MedlinePlus
- Drug Portal

Legend

- MIN Pack Multi

Download

IN/MIN	Ingredient	PIN	Precise Ingredient	BN	Brand Name
SCDC	Clinical Drug Component			SBDC	Branded Drug Component
SCD/GPCK	Clinical Drug or Pack			SBD/BPCK	Branded Drug or Pack
SCDG	Clinical Dose Form Group	DFG	Dose Form Group	SBDG	Branded Dose Form Group



String prochlorperazine

prochlorperazine [RxCUI = 8704]

- RxNorm Graph
- RxNorm Properties
- NDC
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Legend

MIN Pack Multi

Download

IN/MIN	Ingredient (2)
H Rx S	prochlorperazine
M	isopropamide / prochlorperazine

PIN	Precise Ingredient (3)
Rx S	prochlorperazine edisylate
Rx S	prochlorperazine maleate
S	prochlorperazine mesylate

BN	Brand Name (2)
H Rx S	Compazine
H Rx S	Compro

SCDC	Clinical Drug Component (9)
S	prochlorperazine 1 MG/ML
H Rx S	prochlorperazine 10 MG
S	prochlorperazine 12.5 MG/ML
S	prochlorperazine 15 MG
S	prochlorperazine 2.5 MG
H Rx S	prochlorperazine 25 MG
S	prochlorperazine 3 MG
H Rx S	prochlorperazine 5 MG
H Rx S	prochlorperazine 5 MG/ML



SBDC	Branded Drug Component (4)
H Rx S	prochlorperazine 10 MG [Compazine]
H Rx S	prochlorperazine 25 MG [Compazine]
H Rx S	prochlorperazine 25 MG [Compro]
H Rx S	prochlorperazine 5 MG [Compazine]

SCD/GPCK	Clinical Drug or Pack (13)
H Rx S	2 ML prochlorperazine 5 MG/ML Injection
S	prochlorperazine 1 MG/ML Oral Solution
S	prochlorperazine 10 MG Extended Release Oral Capsule
H Rx S	prochlorperazine 10 MG Oral Tablet
S	prochlorperazine 12.5 MG/ML Injectable Solution
S	prochlorperazine 15 MG Extended Release Oral Capsule
S	prochlorperazine 2.5 MG Rectal Suppository
S	prochlorperazine 25 MG Oral Tablet
H Rx S	prochlorperazine 25 MG Rectal Suppository
S	prochlorperazine 3 MG Buccal Tablet

SBD/BPCK	Branded Drug or Pack (4)
H Rx S	Compazine 10 MG Oral Tablet
H Rx S	Compazine 25 MG Rectal Suppository
H Rx S	Compazine 5 MG Oral Tablet
H Rx S	Compro 25 MG Rectal Suppository

SCDG	Clinical Dose Form Group (6)
S	prochlorperazine Buccal Product
H Rx S	prochlorperazine Injectable Product
S	prochlorperazine Oral Liquid Product
H Rx S	prochlorperazine Oral Product
H Rx S	prochlorperazine Pill
H Rx S	prochlorperazine Rectal Product

DFG	Dose Form Group (6)
H Rx S	Buccal Product
HV Rx S	Injectable Product
HV Rx S	Oral Liquid Product
HV Rx S	Oral Product
HV Rx S	Pill
HV Rx S	Rectal Product

SBDC	Branded Dose Form Group (4)
H Rx S	Compazine Oral Product
H Rx S	Compazine Pill
H Rx S	Compazine Rectal Product
H Rx S	Compro Rectal Product

PIN

Precise Ingredient (3)

Rx S prochlorperazine edisylate

Rx S prochlorperazine maleate

S prochlorperazine mesylate

SCD/GPCK**Clinical Drug or Pack (13)**

H Rx S 2 ML prochlorperazine 5 MG/ML Injection

S prochlorperazine 1 MG/ML Oral Solution

S prochlorperazine 10 MG Extended Release Oral Capsule

H Rx S prochlorperazine 10 MG Oral Tablet

S prochlorperazine 12.5 MG/ML Injectable Solution

S prochlorperazine 15 MG Extended Release Oral Capsule

S prochlorperazine 2.5 MG Rectal Suppository

S prochlorperazine 25 MG Oral Tablet

H Rx S prochlorperazine 25 MG Rectal Suppository

S prochlorperazine 3 MG Buccal Tablet

H Rx S prochlorperazine 5 MG Oral Tablet

S prochlorperazine 5 MG Rectal Suppository

SCDG**Clinical Dose Form Group (6)****S** prochlorperazine Buccal Product**H Rx S** prochlorperazine Injectable Product**S** prochlorperazine Oral Liquid Product**H Rx S** prochlorperazine Oral Product**H Rx S** prochlorperazine Pill**H Rx S** prochlorperazine Rectal Product

MTUS – RxCUI consolidation model using MTUS v8

J. Kevin Gorospe, PharmD
DWC Consultant

RxCUI Consolidation

- P&T request to consolidate Non-Exempt under a single RxCUI
 - Would reduce the size of the list (fewer entries)
 - Non-Exempt drugs require authorization
- Consolidation at Dose Form level whenever possible
 - “Ingredient” and “Precise Ingredient” level too broad and would include dosage forms not suitable for the MTUS list (e.g. drugs for infusion)

Most Non-Exempt drugs could be consolidated under a single “Dose Form” RxCUI, e.g., “Oral Product” for acyclovir

SCD/GPCK		Clinical Drug or Pack (7)
H Rx S	acyclovir 200 MG Oral Capsule	
S	acyclovir 200 MG Oral Tablet	
H Rx S	acyclovir 40 MG/ML Oral Suspension	
H Rx S	acyclovir 400 MG Oral Tablet	
H Rx S	acyclovir 50 MG Buccal Tablet	
S	acyclovir 80 MG/ML Oral Suspension	
H Rx S	acyclovir 800 MG Oral Tablet	

SCDG	Clinical Dose Form Group (1)	DFG
H Rx S	acyclovir Oral Product	HvRx S Oral Product

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In some instances, Dose Form too broad and included multiple drug ingredients that appear on the MTUS list, for example betamethasone topical or drugs with individual allowances (e.g. 4-day Special Fill for specific drugs). In these cases, drugs were not consolidated.

SCD/GPCK		Clinical Drug or Pack (16)
		betamethasone 0.0005 MG/MG Topical Ointment
S		betamethasone 0.0005 MG/MG Topical Gel
H Rx S		betamethasone 0.0005 MG/MG Topical Ointment
H Rx S		betamethasone 0.001 MG/MG Topical Ointment
S		betamethasone 0.005 MG/MG Topical Ointment
S		betamethasone 0.25 MG/ML Topical Cream
H Rx S		betamethasone 0.5 MG/ML Topical Cream
H Rx S		betamethasone 0.5 MG/ML Topical Lotion
H Rx S		betamethasone 0.5 MG/ML Topical Spray
H Rx S		betamethasone 1 MG/ML Topical Cream
H Rx S		betamethasone 1 MG/ML Topical Lotion
H Rx S		betamethasone valerate 1.2 MG/ML Topical Foam

SCDG	Clinical Dose Form Group (1)	DFG
H Rx S	betamethasone Topical Product	HvRx S Topical Product

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Additional changes

- In addition to RxCUI consolidation other changes were made that deviate from the latest MTUS v8
 - Drug ingredient names corrected to match FDA listings, e.g.
 - bromfenac changed to bromfenac sodium
 - emedastine ophth changed to emedastine difumarate ophth
 - Consolidate camphor/menthol/methyl salicylate into single entry as reference brand BEN GAY ULTRA contains all three; leave menthol as standalone for listed reference brand BIOFREEZE
 - Brand drug names corrected to match ingredient, dosage form, and strength of specific products, e.g.
 - prednisolone sodium phosphate ophth (ingredient corrected to add “phosphate”) changed reference brand to INFLAMASE FORTE; the listed OMNIPRED is actually prednisolone acetate

More Changes

- Therapeutic categories were corrected/changed for consistency
 - isocarboxazid, phenelzine, selegiline, tranylcypromine from “Antidepressants” to “Antidepressants (MAOIs)”
 - nefazodone from “Antidepressants (SSRI)” to “Antidepressants (Serotonin Modulators)”
- 4-day Special Fill and Peri-Op quantities for specific opioids were entered per prior P&T recommendation
- P&T recommendations on Exempt status were incorporated
- Drugs no longer on the market were removed
- Various name and therapeutic classification change recommendations will be communicated to ACOEM

Previous P&T Recommendations

Drug Ingredient	Reference Brand Name	Exempt/Non-Exempt*	Special Fill	Peri-Op	Drug Class	Dosage Form	Strength	RxCUI	Comments
bromfenac sodium ophth	BROMSITE	Non-Exempt	Not Applicable	Not Applicable	Ophthalmic Agents (NSAID)	Solution, ophthalmic	0.075%	1790141	Move bromfenac sodium from exempt to non-exempt status
bromfenac sodium ophth	BROMDAY, XIBROM	Non-Exempt	Not Applicable	Not Applicable	Ophthalmic Agents (NSAID)	Solution, ophthalmic	0.09%	578018	Move bromfenac sodium from exempt to non-exempt status
bromfenac sodium ophth	PROLENSA	Non-Exempt	Not Applicable	Not Applicable	Ophthalmic Agents (NSAID)	Solution, ophthalmic	0.07%	1375917	Move bromfenac sodium from exempt to non-exempt status
diclofenac potassium	CATAFLAM	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Tablet	25 MG	857702	Move all oral systemic diclofenac from exempt to non-exempt status
diclofenac potassium	CATAFLAM	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Tablet	50 MG	855942	Move all oral systemic diclofenac from exempt to non-exempt status
diclofenac potassium	ZIPSOR	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Capsule	25 MG	858342	Move all oral systemic diclofenac from exempt to non-exempt status
diclofenac potassium	CAMBIA	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Powder for Solution	50 MG	859063	Move all oral systemic diclofenac from exempt to non-exempt status
indomethacin	TIVORBEX	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Capsule	20 MG	1490727	Recommend to the Administrative Director that 20mg and 40 mg indomethacin be changed from exempt to non-exempt
indomethacin	TIVORBEX	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Capsule	40 MG	1491529	Recommend to the Administrative Director that 20mg and 40 mg indomethacin be changed from exempt to non-exempt
meloxicam	VIVLODEX	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Capsule	5 MG	1722349	Recommend to Administrative Director that capsule form of meloxicam (5mg and 10mg strengths) be changed from exempt to non-exempt.
meloxicam	VIVLODEX	Non-Exempt	Not Applicable	Not Applicable	Analgesics - Anti-Inflammatory (NSAID)	Capsule	10 MG	1722357	Recommend to Administrative Director that capsule form of meloxicam (5mg and 10mg strengths) be changed from exempt to non-exempt.
naloxone hcl	EVZIO	Non-Exempt	Not Applicable	Not Applicable	Antidotes and Specific Antagonists	Injection IM/SC	2 MG/ 0.4ML	1855730	Recommend that the Administrative Director keep Narcan (nasal) exempt and change designation of Evzio (auto-injector) from exempt to non-exempt.

Committee Discussion

Public Comments

Reports/Research of Interest

General Discussion

Committee Discussion

Public Comments

COVID Vaccination

- ACOEM COVID-19 (Coronavirus) Guideline was last updated on December 14, 2020

General Discussion



State of California
Gavin Newsom
Governor

California Dept. of Public Health

- Information available
 - Main Website
 - <https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/Immunization/ncov2019.aspx>
 - Community Vaccine Advisory Committee
 - <https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/COVID-19/Community-Vaccine-Advisory-Committee.aspx>
 - Drafting Guidelines Workgroup
 - <https://www.cdph.ca.gov/Programs/CID/DCDC/Pages/COVID-19/Drafting-Guidelines-Workgroup.aspx>

Committee Discussion

Public Comments

Review of Recommendations

Adjournment