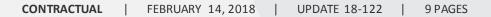
## **PROVIDER***Update*



## Medication Trend Updates and Formulary Changes – 1st Quarter 2018

This update includes information regarding the use of codeine and tramadol in children and nursing mothers, an opioid prescribing update for Medi-Cal members, the Health Net Prescription Drug Prior Authorization or Step Therapy Exception Request Form, and changes to the Health Net of California, Inc., Health Net Community Solutions, Inc. and Health Net Life Insurance Company (Health Net) commercial *Recommended Drug Lists* (*RDLs*), *Medi-Cal RDL* and *Medicare Part D Formularies* for the first quarter of 2018.

### CODEINE AND TRAMADOL SHOULD NOT BE PRESCRIBED FOR CHILDREN AND NURSING MOTHERS

On April 20, 2017, the U.S. Food and Drug Administration (FDA) issued an announcement restricting the use of codeine and tramadol (generic for Ultram<sup>®</sup>) in children and nursing mothers. The updated changes include:

- A contraindication to the codeine and tramadol labels alerting users that codeine should not be used to treat pain or cough, and tramadol should not be used to treat pain in children younger than age 12.
- A new contraindication to the tramadol label warning against its use in children younger than age 18 to treat pain after surgery to remove the tonsils and/or adenoids.
- A new warning to codeine and tramadol labels recommending against their use in adolescents between ages 12–18 who are obese and have conditions such as obstructive sleep apnea or severe lung disease.
- A warning that breastfeeding is not recommended when taking codeine or tramadol due to the risk of serious adverse reactions in breastfed infants, which can include excess sleepiness, difficulty breastfeeding, or serious breathing problems that could result in death.

This advice is also consistent with the September 2016 Policy Statement from the American Academy of Pediatrics titled *Codeine: Time To Say "No."* 

For more information, visit the following websites:

- www.fda.gov/Drugs/DrugSafety/ucm549679.htm
- http://pediatrics.aappublications.org/content/early/2016/09/15/peds.2016-2396

### **OPIOID PRESCRIBING UPDATE FOR MEDI-CAL MEMBERS**

In response to the national opioid epidemic and recommendations from the Centers for Disease Control and Prevention (CDC), and to ensure proper use of opioid medications and safety of our members, Health Net will be implementing changes in opioid utilization. Starting in April 2018, prescribers may be required to submit a prior authorization and a signed pain contract for Medi-Cal members when opioid prescriptions exceed plan quantity or frequency limits. For more information, review the following resources:

 CDC Opioid Prescribing Guidelines: www.cdc.gov/drugoverdose/prescribing/guideline.html.



### THIS UPDATE APPLIES TO **CALIFORNIA** PROVIDERS:

#### • Physicians

- Participating Physician Groups
- Hospitals
- O Ancillary Providers

#### LINES OF BUSINESS:

- HMO/POS/HSP
- PPO
- EPO
- Medicare Advantage (HMO)
- Medi-Cal
  - Kern
  - Los Angeles
    - Molina
  - Riverside
  - Sacramento
  - San Bernardino
  - San Diego
  - San Joaquin
  - Stanislaus
  - Tulare

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- CDC's *Guideline for Prescribing Opioids for Chronic Pain* factsheet: www.cdc.gov/drugoverdose/pdf/Guidelines\_Factsheet-a.pdf.
- Prescription Drug Monitoring Program [Controlled Substance Utilization Review and Evaluation System (CURES) 2.0]. All prescribers are required to enroll in and access CURES reports to establish whether or not a member is receiving controlled substances from other providers. For further information, visit https://oag.ca.gov/cures.

### NEW PRESCRIPTION DRUG PRIOR AUTHORIZATION OR STEP THERAPY EXCEPTION REQUEST FORM

Effective January 1, 2018, providers must use the updated Prescription Drug Prior Authorization or Step Therapy Exception Request Form for members with prescription medication benefits enrolled in commercial and Medi-Cal plans. The form must be completed and submitted for all prior authorization prescription requests. A copy of the updated form is attached to this update for reference, and is also available on the Health Net provider website at provider.healthnet.com under *Pharmacy Information* and in the Provider Library under *Forms*. Health Net processes medication authorization requests within 72 hours for nonurgent requests and 24 hours for exigent requests for commercial members. Medication authorization requests for Medi-Cal members continue to be processed within 24 hours.

Beginning January 1, 2018, providers must submit prior authorization prescription requests on the updated form, or the request will be rejected. Also, effective January 1, 2018, the previous version of the Prescription Drug Prior Authorization Request Form (also referred to as the SB866 form) is no longer accepted.

### CHANGES TO THE RECOMMENDED DRUG LIST AND MEDICARE PART D FORMULARIES

The Health Net Pharmacy and Therapeutics (P&T) Committee, which comprises practicing physicians, pharmacists and other health care professionals, reviews medications on the Health Net *RDLs* and *Formularies* for commercial and Medi-Cal members, and the *Medicare Part D Formularies* for Medicare members each quarter to determine medications to remain on or be moved to a different tier. A list of some recent changes is provided beginning on page 4. The list contains brand-name prescription medications, status, alternatives, and comments for the first quarter of 2018.

Complete lists of the *RDL*s, *Formularies* and *Medicare Part D Formularies* are available on the Health Net provider website at provider.healthnet.com by selecting *Pharmacy Information* or *Provider Library*. Other pharmacy-related provider updates, prior authorization criteria and pharmacy forms are also available online under *Pharmacy Information*.

### PHARMACY HELP LINE

For additional information regarding changes to the commercial Health Net *RDL*, Health Net Medi-Cal *RDLs* or *Medicare Part D Formularies*, contact the appropriate pharmacy telephone numbers listed below:

- **Pharmacy Services** (commercial): 1-800-548-5524, option #3; fax 1-800-314-6223
- Pharmacy Service Center (Medi-Cal and Medicare): 1-800-867-6564; fax 1-800-977-8226
- Health Net Clinical Pharmacy Line (clinical programs): 1-800-782-2221

### ADDITIONAL INFORMATION

Providers are encouraged to access Health Net's provider portal online, as listed in the table below, for real-time information, including eligibility verification, claims status, prior authorization status, plan summaries, and more.

If you have questions regarding the information contained in this update, contact the applicable Health Net Provider Services Center within 60 days at:

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ENHANCEDCARE PPO (IFP)	1-844-463-8188	provider.healthnetcalifornia.com	
ENHANCEDCARE PPO (SBG)	1-844-463-8188	provider.healthnet.com	provider_services@healthnet.com
HEALTH NET EMPLOYER GROUP HMO, POS, HSP, PPO, & EPO	1-800-641-7761	provider.healthnet.com	

Line of Business	Telephone Number	Provider Portal	Email Address
INDIVIDUAL FAMILY PLAN	1-888-926-2164	provider.healthnetcalifornia.com	
MEDICARE (INDIVIDUAL)	1-800-929-9224	provider.healthnetcalifornia.com	provider_services@healthnet.com
MEDICARE (EMPLOYER GROUP)	1-800-929-9224	provider.healthnet.com	
MEDI-CAL	1-800-675-6110	provider.healthnet.com	N/A

		Status		Health	Net Formulary Altern	ative(s)	
Medication	Commercial 3-Tier Plan (4-Tier Plan)	Medicare Part D Value <sup>1</sup>	Medi-Cal	Commercial (Tier 1 or 2)	Medicare Part D Value <sup>1</sup> (Tier 1, 2, 3, or 6)	Medi-Cal	Comments
ORAL MEDICATION	IS						
ldhifa <sup>®</sup> (enasidenib) tablet	Tier 2* (Specialty Tier*)	Tier 5*	F*, **				Treatment of adult patients with relapsed or refractory acute myeloid leukemia (AML) with an IDH2 mutation as detected by an FDA-approved test
Nerlynx <sup>®</sup> (neratinib) tablet	Tier 2* (Specialty Tier*)	Tier 5*	F*, **			Herceptin <sup>®</sup> (under medical benefit)*	Extended adjuvant treatment of adult patients with early stage human epidermal growth factor receptor 2 (HER2)- overexpressed/amplified breast cancer, to follow adjuvant trastuzumab-based therapy
Symproic <sup>®</sup> (naldemedine) tablet	Tier 3* (Tier 4*)	NF	NF**	lactulose, polyethylene glycol 3350 powder (Miralax <sup>®</sup> )	lactulose, polyethylene glycol 3350 packet (Miralax)	docusate sodium (Colace <sup>®</sup> ), lactulose, polyethylene glycol 3350 (Miralax), bisacodyl (Dulcolax <sup>®</sup> )	Treatment of opioid-induced constipation (OIC) in adult patients with chronic non-cancer pain

### HEALTH NET RECOMMENDED DRUG LIST (RDL), MEDI-CAL FORMULARY AND MEDICARE PART D FORMULARY CHANGES

		Status		Health	Net Formulary Altern		
Medication	Commercial 3-Tier Plan (4-Tier Plan)	Medicare Part D Value <sup>1</sup>	Medi-Cal	Commercial (Tier 1 or 2)	Medicare Part D Value <sup>1</sup> (Tier 1, 2, 3, or 6)	Medi-Cal	Comments
ORAL MEDICATION	IS, CONTINUED						
Vosevi® (sofosbuvir/ velpatasvir/ voxilaprevir) tablet	Tier 3* (Specialty Tier*)	NF	NF	Mavyret <sup>™</sup> Mavyret is the preferred product. For 3- tier plan: Mavyret is Tier 3 with prior authorization (PA). For 4-tier plan, it is specialty tier with PA		Mavyret*	<ul> <li>Treatment of adult patients with chronic hepatitis C virus (HCV) infection without cirrhosis or with compensated cirrhosis (Child-Pugh A) who have:</li> <li>Genotype 1, 2, 3, 4, 5, or 6 infection and have previously been treated with an HCV regimen containing an NS5A inhibitor. (In clinical trials, prior NS5A inhibitor experience included daclatasvir, elbasvir, ledipasvir, or welpatasvir.)</li> <li>Genotype 1a or 3 infection and have previously been treated with an HCV regimen containing sofosbuvir without an NS5A inhibitor. [In clinical trials, prior treatment experience included sofosbuvir without an NS5A inhibitor. [In clinical trials, prior treatment experience included sofosbuvir without an NS5A inhibitor. [In clinical trials, prior treatment experience included sofosbuvir with or without any of the following: peginterferon alfa/ribavirin, ribavirin, HCV NS3/4A protease inhibitor (boceprevir, simeprevir or telaprevir).]</li> <li>Additional benefit of Vosevi over sofosbuvir/velpatasvir was not shown in adults with genotype 1b, 2, 4, 5, or 6 infection previously treated with sofosbuvir without an NS5A inhibitor</li> </ul>

	Status			Health	Net Formulary Altern	ative(s)	
Medication	Commercial 3-Tier Plan (4-Tier Plan)	Medicare Part D Value <sup>1</sup>	Medi-Cal	Commercial (Tier 1 or 2)	Medicare Part D Value <sup>1</sup> (Tier 1, 2, 3, or 6)	Medi-Cal	Comments
INHALATION PREP	ARATIONS			· · · · · · · · · · · · · · · · · · ·			
Seebri <sup>™</sup> Neohaler <sup>®</sup> (glycopyrrolate) inhalation powder	Tier 3 QL (Tier 4 QL)	NF	NF (QL)	Advair Diskus, <sup>®</sup> Atrovent <sup>™</sup> HFA, Breo <sup>®</sup> Ellipta, <sup>®</sup> Incruse Ellipta, <sup>®</sup> Serevent <sup>®</sup> Diskus, Symbicort, <sup>®</sup> Spiriva, <sup>®</sup> Tudorza <sup>™</sup> Pressair <sup>™</sup>	Advair Diskus, Breo Ellipta, Foradil <sup>®</sup> Aerolizer, <sup>®</sup> Incruse Ellipta, Serevent Diskus, Spiriva, Tudorza Pressair	Advair Diskus, Atrovent HFA, Breo Ellipta, Combivent Respimat,® Incruse Ellipta, Serevent Diskus, Symbicort, Spiriva, Tudorza Pressair	Long-term maintenance treatment of airflow obstruction in patients with chronic obstructive pulmonary disease (COPD), including chronic bronchitis and/or emphysema Quantity limit is 1 inhaler per month
INJECTABLE PREPA	RATIONS			·			
Kevzara <sup>®</sup> (sarilumab) prefilled syringe	Medical benefit (Specialty Tier*)	Tier 5*	NF**	Disease- modifying antirheumatic drugs (DMARDs): methotrexate, sulfasalazine, leflunomide, hydroxychloroqu ine Preferred biologic DMARDs at Specialty Tier*: Humira, <sup>®</sup> Enbrel, <sup>®</sup> Xeljanz, <sup>®</sup> XR	DMARDs: methotrexate, sulfasalazine, leflunomide, hydroxychloroquine	DMARDs: methotrexate**, sulfasalazine, leflunomide**, hydroxychloroqu ine Biologic DMARDS: Humira*, **, Simponi Aria <sup>®</sup> *, **, Remicade <sup>®</sup> *, **	Treatment of adult patients with moderately to severely active rheumatoid arthritis (RA) who have had an inadequate response or intolerance to one or more DMARDs For commercial line of business, coverage of Kevzara requires a trial of Humira and Enbrel, followed by Xeljanz or Xeljanz XR

	Status			Health	Net Formulary Altern			
Medication	Commercial 3-Tier Plan (4-Tier Plan)	Medicare Part D Value <sup>1</sup>	Medi-Cal	Commercial (Tier 1 or 2)	Medicare Part D Value <sup>1</sup> (Tier 1, 2, 3, or 6)	Medi-Cal	Comments	
INJECTABLE PREPA	INJECTABLE PREPARATIONS, CONTINUED							
Tremfya <sup>®</sup> (guselkumab) prefilled syringe	Medical benefit (Specialty Tier*)	Tier 5*	NF**	methotrexate, cyclosporine Preferred biologic DMARDs at Specialty Tier*: Humira, Cosentyx®	methotrexate, cyclosporine (Part B vs. Part D determination)	methotrexate**, cyclosporine** Biologic DMARDS: Humira*, **, Stelara <sup>®</sup> *, **, Remicade*, **	Treatment of adult patients with moderate-to-severe plaque psoriasis who are candidates for systemic therapy or phototherapy For commercial line of business, coverage of Tremfya requires a trial of Humira and Cosentyx	

<sup>1</sup>Medicare Part D Value Formulary = Health Net Seniority Plus Amber I (HMO SNP), Health Net Seniority Plus Amber II (HMO SNP), Health Net Healthy Heart (HMO) \*Prior authorization (PA) is required to verify member eligibility and that the member satisfies clinical protocols to ensure appropriate use of the medication. \*\*CCS = California Children's Services: Refer to w ww.dhcs.ca.gov for the local telephone number to determine member's coverage eligibility.

- F indicates formulary
- NF indicates nonformulary
- NS indicates nonstructural protein
- QL indicates quantity limit
- XL indicates extended release

### PRESCRIPTION DRUG PRIOR AUTHORIZATION OR STEP THERAPY EXCEPTION REQUEST FORM

Plan/Medical Group Name: Plan/Medical Group Fax#: ()				Plan/Medical Group Phone#: () Non-Urgent				
<b>Instructions:</b> Please fill out all applicable sections on both pages completely and legibly. Attach any additional documentation that is important for the review, e.g. chart notes or lab data, to support the prior authorization or step-therapy exception_request. <b>Information contained in this form is Protected Health Information under HIPAA.</b>								
Patient Information								
First Name:	L	_ast Name:			MI:	P	hone Nur	nber:
Address:			City:				State:	Zip Code:
	] Male ] Female	Circle unit of Height (in/cm		_Weight (lb/kg):	/	Allerg	jies:	
Patient's Authorized Representat	ive (if applica	-		Authorized Repre	esentative	e Pho	ne Numb	er:
		Ins	surance	Information				
Primary Insurance Name:				Patient ID Numbe	er:			
Secondary Insurance Name:				Patient ID Numbe	er:			
		Pre	escriber	Information				
First Name:		Last Name:		Specialty:				
Address:			City:	State: Zip Code:			Zip Code:	
Requestor (if different than presc	riber):			Office Contact Pe	erson:			
NPI Number (individual):				Phone Number:				
DEA Number (if required):				Fax Number (in HIPAA compliant area):				
Email Address:								
	M	edication / Me	dical and	d Dispensing Info	rmation			
Medication Name:								
☐ New Therapy ☐ Renewal If Renewal: Date Therapy Initiate	-	rapy Exception	Request	Duration of Therap	oy (specif	ic dat	tes):	
How did the patient receive the m Paid under Insurance Name Other (explain):				Prior Auth N	Number (i	if kno	wn):	
Dose/Strength:	Freque	ncy:		Length of Therap	y/#Refills	S:	Qua	ntity:
Administration:		on 🗌 IV		] Other:				
Administration Location:	☐ Patio	ent's Home ne Care Agency patient Hospital		Long Term Ca				

### PRESCRIPTION DRUG PRIOR AUTHORIZATION OR STEP THERAPY EXCEPTION REQUEST FORM

Patient Name:		ID#:	
<b>Instructions:</b> Please fill out all applicable sections on be important for the review, e.g. chart notes or lab data, to			
1. Has the patient tried any other medications for thi	s condition?	S (if yes, complete below)	□ NO
<b>Medication/Therapy</b> (Specify Drug Name and Dosage)	Duration of Therapy (Specify Dates)	Response/Reaso	n for Failure/Allergy
2. List Diagnoses:		ICD-10:	
3. <u>Required clinical information</u> - Please provide all r exception request review.	relevant clinical informati	on to support a prior authoriz	ation or step therapy
Please provide symptoms, lab results with dates and/or ji contraindications for the health plan/insurer preferred dru evaluate response. Please provide any additional clinica information related to exigent circumstances, or required Attachments	ug. Lab results with dates r al information or comments	nust be provided if needed to es pertinent to this request for cove	stablish diagnosis, or
<b>Attestation:</b> I attest the information provided is true and a Medical Group or its designees may perform a routine at information reported on this form.	-	-	
Prescriber Signature or Electronic I.D. Verification	ion:	Date:	
<b>Confidentiality Notice</b> : The documents accompanying this are not the intended recipient, you are hereby notified th these documents is strictly prohibited. If you have receive and arrange for the return or destruction of these documents are documents are documents of the section of the section.	at any disclosure, copying, yed this information in error,	distribution, or action taken in r	eliance on the contents of
Plan/Insurer Use Only: Date/Time Request Receiv	ved by Plan/Insurer:	Date/Time of D	Decision
Fax Number <u>()</u>			
Approved Denied Comments/Information Reg	puested:		

## **PROVIDER***Update*

CONTRACTUAL | FEBRUARY 14, 2018 | UPDATE 18-122sum | 3 PAGES

## *Summary Update*: Medication Trend Updates and Formulary Changes – 1st Quarter 2018

This update includes information regarding the use of codeine and tramadol in children and nursing mothers, an opioid prescribing update for Medi-Cal members, the Health Net Prescription Drug Prior Authorization or Step Therapy Exception Request Form, and changes to the Health Net of California, Inc., Health Net Community Solutions, Inc. and Health Net Life Insurance Company (Health Net) commercial *Recommended Drug Lists* (*RDLs*), *Medi-Cal RDL* and *Medicare Part D Formularies* for the first quarter of 2018.

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This advice is also consistent with the September 2016 Policy Statement from the American Academy of Pediatrics titled, *Codeine: Time To Say "No."* 

For more information, visit the following websites:

- www.fda.gov/Drugs/DrugSafety/ucm549679.htm
- http://pediatrics.aappublications.org/content/early/2016/09/15/peds.2016-2396

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# Health Net<sup>®</sup>

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Health Net of California, Inc., Health Net Community Solutions, Inc. and Health Net Life Insurance Company are subsidiaries of Health Net, Inc. and Centene Corporation. Health Net is a registered service marks feature marks feature marks feature and the property of their respective companies. All rights reserved. Confidentiality Note for Fax Transmission: This facsimile may contain confidential information. The information is intended only for the use of the individual or entity named above. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or use of the information contained in this transmission is strictly PROHIBITED. If you have received this transmission in error, please notify the sender immediately by telephone or by return fax and destroy this transmission, along with any attachments. OTH018269EH00 (2/18)

signed pain contract for Medi-Cal members when opioid prescriptions exceed plan quantity or frequency limits. For more information, review the following resources:

- CDC Opioid Prescribing Guidelines: www.cdc.gov/drugoverdose/prescribing/guideline.html.
- CDC's *Guideline for Prescribing Opioids for Chronic Pain* factsheet: www.cdc.gov/drugoverdose/pdf/Guidelines\_Factsheet-a.pdf.
- Prescription Drug Monitoring Program [Controlled Substance Utilization Review and Evaluation System (CURES) 2.0]. All prescribers are required to enroll in and access CURES reports to establish whether or not a member is receiving controlled substances from other providers. For further information, visit https://oag.ca.gov/cures.

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The form is available on the Health Net provider website at provider.healthnet.com under *Pharmacy Information* and in the Provider Library under *Forms*.

### CHANGES TO THE RECOMMENDED DRUG LISTS AND MEDICARE PART D FORMULARIES

A list of recent changes to the Health Net *RDLs* and *Formularies* is available in the complete provider update 18-122, *Medication Trend Updates and Formulary Changes – 1st Quarter 2018*. The list contains brand-name prescription medications, status, alternatives, and comments. Complete listings of the *RDLs* and *Medicare Part D Formularies* are available on the Health Net provider portal, as listed in the table below, by selecting *Pharmacy Information*.

### PHARMACY HELP LINE

For additional information regarding changes to the commercial Health Net *RDL*, Health Net Medi-Cal *RDLs* or *Medicare Part D Formularies*, contact the appropriate pharmacy telephone numbers listed below:

- Pharmacy Services (commercial): 1-800-548-5524, option #3; fax 1-800-314-6223
- Pharmacy Service Center (Medi-Cal and Medicare): 1-800-867-6564; fax 1-800-977-8226
- Health Net Clinical Pharmacy Line (clinical programs): 1-800-782-2221

### ADDITIONAL INFORMATION

To obtain a comprehensive description of the above topics, the complete update, 18-122, *Medication Trend Updates and Formulary Changes – 1st Quarter 2018,* is available on the Health Net provider portal, as listed in the table below, in the Provider Library under *Updates and Letters > 2018;* search for provider update 18-122. Providers who do not have access to the Internet may request a print copy of update 18-122 by contacting the Health Net Provider Communications Department by fax at 1-800-937-6086 or by email at provider.communications@healthnet.com.

For the most current version of the Health Net *RDLs*, visit the Health Net provider portal, as listed in the table below, under *Pharmacy Information > Drug Lists*.

Providers are encouraged to access Health Net's provider portal online, as listed in the table below, for real-time information, including eligibility verification, claims status, prior authorization status, plan summaries, and more.

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