

Effective Date: 06/01/23

Buckeye Health Plan Medicaid Criteria Updates –Q2 2023

Buckeye Health Plan (BHP) routinely reviews their Prior Authorization (PA) and Medical Necessity (MN) criteria. Decisions on PA and MN criteria content are coordinated with input from pharmacy and medical practitioners, Buckeye Health Plan representatives, and review of current available medical literature and professional standards of practice. Below is the list of changes to the Medicaid criteria this quarter.

For the most current program description you may call Provider Services at 1-866-296-8731 (TTY/TTD

Policy/ Coverage Criteria Guideline	Applicable Business	Revision Summary Description
		Clinically Significant Change(s)
CP.PHAR.16 Palivizumab (Synagis)	Commercial, HIM, Medicaid	2Q 2023 annual review: for CLD added bronchodilator therapy as an additional option to confirm appropriateness of therapy in the second year of life per AAP guidance; references reviewed and updated
CP.PHAR.50 Binimetinib (Mektovi)	Commercial, HIM, Medicaid	2Q 2023 annual review: for melanoma added limited resectable melanoma and added off-label criteria for histiocytic neoplasms per NCCN category 2A recommendation; references reviewed and updated.
CP.PHAR.60 Capecitabine (Xeloda)	HIM, Medicaid	2Q 2023 annual review: collapsed off-label criteria for anal carcinoma and added to NCCN recommended (off-label) criteria set; added ampullary adenocarcinoma and extrapulmonary neuroendocrine carcinoma to NCCN recommended (off-label) list; RT4: per updated PI, updated FDA approved indications for colorectal cancer and breast cancer, added gastric/esophageal/gastroesophageal junction cancer and pancreatic cancer criteria (removed from off-label list), removed criterion "member does not have severe renal impairment (creatinine clearance < 30 mL/min)" as severe renal impairment is no longer a contraindication as updated in Appendix C, updated section V; references reviewed and updated.
CP.PHAR.64 Topotecan (Hycamtin)	Commercial, HIM, Medicaid	2Q 2023 annual review: updated off-label criteria for endometrial carcinoma to include "as second line or subsequent therapy" per NCCN compendium; references reviewed and updated
CP.PHAR.69 Sorafenib (Nexavar)	Commercial, HIM, Medicaid	2Q 2023 annual review: for generic redirection removed "if available" as generic is now available; per NCCN Compendium added additional ovarian cancer subtypes; references reviewed and updated.
CP.PHAR.71 Lenalidomide (Revlimid)	Commercial, HIM, Medicaid	2Q 2023 annual review: per NCCN Compendium updated MM criteria updated maintenance therapy following autologous hematopoietic stem cell transplantation to include option for carfilzomib or bortezomib with dexamethasone, for myelodysplastic syndrome added SF3B1 mutation status, for myelofibrosis-associated anemia, added "in combination with prednisone



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		taper", updated off-label criteria for systemic light chain amyloidosis to include combination therapy, for classic Hodgkin lymphoma changed "as third-line" to "as fourth-line" to align with
		NCCN Hodgkin Lymphoma guideline, for HIV related B-cell lymphoma, post-transplant
		lymphoproliferative disorder of B-cell lymphomas and high grade B-cell diffuse lymphoma added "in combination with Monjuvi for non-transplant candidates", added off-label criteria for
		POEMS syndrome per NCCN 2A recommendation; references reviewed and updated.
CP.PHAR.72 Dasatinib (Sprycel)	Commercial,	2Q 2023 annual review: for MLNE added NCCN supported use in the blast phase; added off-
	HIM, Medicaid	label use in melanoma; modified continued approval duration for Medicaid and HIM lines of business from 6 to 12 months; references reviewed and updated.
CP.PHAR.73 Sunitinib (Sutent)	Commercial,	2Q 2023 annual review: for RCC adjuvant therapy added clarification that clear cell histology is
	HIM, Medicaid	required per NCCN and prescribing information; for pNET added additional options for
	,	recurrent and advanced disease per NCCN; added pheochromocytoma/paraganglioma as NCCN supported off-label uses; references reviewed and updated.
CP.PHAR.74 Erlotinib (Tarceva)	Commercial,	2Q 2023 annual review: added NCCN Compendium supported off-label uses for pigmented
CF.FHAR./4 Ellottillo (Talceva)	HIM, Medicaid	
	milvi, iviedicald	villonodular synovitis/tenosynovial giant cell tumor and melanoma; modified continued
		approval duration for Medicaid and HIM lines of business from 6 to 12 months; references reviewed and updated.
CP.PHAR.76 Nilotinib (Tasigna)	Commercial,	
Cr.FHAR./0 Mionino (Tasigna)	,	2Q 2023 annual review: added NCCN Compendium supported off-label uses for pigmented
	HIM, Medicaid	villonodular synovitis/tenosynovial giant cell tumor and melanoma; modified continued
		approval duration for Medicaid and HIM lines of business from 6 to 12 months; references reviewed and updated.
CD DILAD 77 Tomografianida	IIIM Madiasid	
CP.PHAR.77 Temozolomide	HIM, Medicaid	2Q 2023 annual review: for astrocytoma, replaced term anaplastic (no longer used in
(Temodar)		compendium) with IDH mutant per NCCN, added WHO grading per NCCN, added "disease is
		refractory and progressive despite treatment with procarbazine and nitrosourea" and added
		Appendix B per FDA label; per NCCN, removed criteria for mucosal melanoma (downgraded to
		2B recommendation) and angiosarcoma as use is no longer recommended, updated off label-use
		criteria for solitary fibrous tumor to require use with bevacizumab, added off-label criteria for
		extrapulmonary poorly differentiated neuroendocrine carcinoma, large or small cell carcinoma,
		mixed neuroendocrine-non-neuroendocrine neoplasms, pediatric diffuse high-grade gliomas,
		updated off-label criteria for astrocytoma, oligodendroglioma and uterine sarcoma to align with
CD DIIAD 70 Thatida mida (Thata mid)	Commercial	NCCN; references reviewed and updated.
CP.PHAR.78 Thalidomide (Thalomid)	Commercial,	2Q 2023 annual review: for myeloproliferative neoplasms added prescribed in combination with
	HIM, Medicaid	prednisone per NCCN 2A recommendation; for aphthous stomatitis/ulcers, updated dose from
		100 to 400 mg per day in initial criteria per Clinical Pharmacology and referenced trial
		(Jacobson et al); clarified MM dosing in continued therapy criteria; revised oral oncology
		generic (if available) redirection language to align with template; references reviewed and
		updated.



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CP.PHAR.90 Crizotinib (Xalkori)	Commercial,	2Q 2023 annual review: added off-label NCCN-supported indications of cutaneous melanoma
	HIM, Medicaid	and uterine sarcoma; references reviewed and updated.
CP.PHAR.103 Immune Globulins	Commercial,	2Q 2023 annual review: added limitation of use for HyQvia and Privigen; removed HCPCS code
	HIM, Medicaid	C9270; added HCPCS Codes J1460, J1554, J1558, J1560; removed references to Carimune NF
		due to product discontinuation; references reviewed and updated.
CP.PHAR.107 Regorafenib (Stivarga)	Commercial,	2Q 2023 annual review: for GIST per prescribing information and NCCN clarified previous
	HIM, Medicaid	treatment requiring imatinib and Sutent, added per NCCN Compendium off label uses in
		combination with everolimus and SDH mutation positive disease; for soft tissue sarcoma
		removed solitary fibrous tumor as this off-label use is no longer NCCN Compendium supported,
		for pleomorphic rhabdomyosarcoma clarified disease is advanced or metastatic, for non-
		adipocytic sarcoma clarified use is for subsequent therapy for advanced, metastatic, recurrent
		unresectable or recurrent stage IV disease; references reviewed and updated.
CP.PHAR.112.Ponatinib (Iclusig)	Commercial,	2Q 2023 annual review: for ALL added age requirement of 18 years or older, clarified HIM
	HIM, Medicaid	approval durations to be consistent with Medicaid line of business; references reviewed and
		updated.
CP.PHAR.116 Pomalidomide	Commercial,	2Q 2023 annual review: for MM, added requirement that Pomalyst must be prescribed in
(Pomalyst)	HIM, Medicaid	combination with dexamethasone per PI and NCCN; added off-label criteria for POEMS and
		added requirement for Pomalyst to be prescribed as a single agent for primary CNS lymphomas
		per NCCN 2A compendium recommendation; references reviewed and updated.
CP.PHAR.127 Encorafenib (Braftovi)	Commercial,	2Q 2023 annual review: for melanoma criteria added limited resectable melanoma, and for colon
	HIM, Medicaid	and rectal cancer added appendiceal adenocarcinoma per NCCN category 2A recommendation;
		references reviewed and updated.
CP.PHAR.145 Deferasirox (Exjade,	Commercial,	Added Parkinson disease to section III with rationale in Appendix E.
Jadenu)	HIM, Medicaid	
CP.PHAR.146 Deferoxamine	Commercial,	Added Parkinson disease to section III with rationale in Appendix D.
(Desferal)	HIM, Medicaid	
CP.PHAR.147 Deferiprone (Ferriprox)	Commercial,	Added Parkinson disease to section III with rationale in Appendix E.
	Medicaid	
CP.PHAR.172 Histrelin (Vantas,	Commercial,	Added Commercial line of business; added off-label use criteria for gender dysphoria or gender
Supprelin LA)	HIM-Medical	transition.
	Benefit,	
	Medicaid	
CP.PHAR.174 Nafarelin (Synarel)	HIM, Medicaid	Added off-label use criteria for gender dysphoria or gender transition.
CP.PHAR.176 Paclitaxel protein-	Commercial,	2Q 2023 annual review: removed criterion for prior anthracycline therapy for non-triple negative
bound (Abraxane)	HIM, Medicaid	breast cancer per NCCN; added ampullary adenocarcinoma and cervical cancer as additional
		NCCN supported indications (off-label); removed HCPCS/CPT code 96413 and 96415;
		references reviewed and updated.



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CP.PHAR.227 Pertuzumab (Perjeta)	Commercial, HIM, Medicaid	2Q 2023 annual review: for breast cancer, added option for Perjeta without taxanes and chemotherapy for members previously treated with chemotherapy and trastuzumab without pertuzumab and revised docetaxel to taxane-containing chemotherapy per NCCN 2A recommendation; for colorectal cancer, removed requirement for no previous use of a HER2 inhibitor therapy; added unresectable or metastatic HER2-positive gallbladder cancer and cholangiocarcinoma to NCCN recommended uses (off-label); references reviewed and updated.
CP.PHAR.228 Trastuzumab Biosimilars Trastuzumab- Hyaluronidase	Commercial, HIM, Medicaid	2Q 2023 annual review: added gallbladder cancer and cholangiocarcinoma as NCCN supported off-label indication; references reviewed and updated.
CP.PHAR.236 Darbepoetin alfa (Aranesp)	Commercial, HIM, Medicaid	2Q 2023 annual review: per NCCN for MDS continuation of therapy modified treatment response assessment to occur after at least 8 weeks of therapy (previously this was 12 weeks); per NCCN Compendium for MDS added approval pathway for lower risk (IPSS low/intermediate-1) disease associated with symptomatic anemia with del(5q); references reviewed and updated.
CP.PHAR.237 Epoetin alfa (Epogen, Procrit), Epoetin alfa-epbx (Retacrit)	Commercial, HIM, Medicaid	2Q 2023 annual review: per NCCN for MDS continuation of therapy modified treatment response assessment to occur after at least 8 weeks of therapy (previously this was 12 weeks); per NCCN Compendium for MDS added approval pathway for lower risk (IPSS low/intermediate-1) disease associated with symptomatic anemia with del(5q); for cancer indications and other indications sections clarified redirection requirements to include an option for Retacrit requests where no redirection is required; for zidovudine induced anemia continuation of therapy added requirement to confirm member continues to receive zidovudine therapy; references reviewed and updated.
CP.PHAR.239 Dabrafenib (Tafinlar)	Commercial, HIM, Medicaid	2Q 2023 annual review: moved the following indications: hepatobiliary cancers, CNS cancers, ovarian, fallopian and peritoneal cancers from off-label criteria and added ampullary adenocarcinoma, pancreatic adenocarcinoma, salivary gland tumor, thyroid carcinoma (papillary, follicular, Hürthle) to solid tumor criteria (per NCCN 2A recommendation), as they are classified as solid tumors; references reviewed and updated. Template changes per continued therapy section not applicable.
CP.PHAR.240 Trametinib (Mekinist)	Commercial, HIM, Medicaid	2Q 2023 annual review: moved the following indications: hepatobiliary cancers, CNS cancer, ovarian, fallopian, and peritoneal cancers, and metastatic uveal melanoma from off-label criteria and added ampullary adenocarcinoma, pancreatic adenocarcinoma, salivary gland tumor, thyroid carcinoma (papillary, follicular, Hürthle) to solid tumor criteria (per NCCN 2A recommendation), as they are classified as solid tumors; for NSCLC updated oral oncology generic redirection language to align with other indications in policy; references reviewed and updated. Template changes per continued therapy section do not apply.
CP.PHAR.241 Abatacept (Orencia)	Medicaid	2Q 2023 annual review: for pJIA, PsA, and RA, added TNFi criteria to allow bypass if member has had history of failure of two TNF blockers; references reviewed and updated.



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CP.PHAR.244 Anakinra (Kineret)	Medicaid	2Q 2023 annual review: for RA, added TNFi criteria to allow bypass if member has had history of failure of two TNF blockers; updated appendix D with general information for CAPS;
		references reviewed and updated.
CP.PHAR.246 Canakinumab (Ilaris)	Commercial,	2Q 2023 annual review: no significant changes; updated off-label dosing for Appendix B;
(2)	HIM, Medicaid	references reviewed and updated.
CP.PHAR.247 Certolizumab (Cimzia)	Medicaid	2Q 2023 annual review: for PsA and RA, added TNFi criteria to allow bypass if member has had
(history of failure of two TNF blockers; references reviewed and updated.
CP.PHAR.253 Golimumab (Simponi,	Medicaid	2Q 2023 annual review: for AS, pJIA, PsA, and RA, added TNFi criteria to allow bypass if
Simponi Aria)		member has had history of failure of two TNF blockers; reference reviewed and updated.
CP.PHAR.259 Natalizumab (Tysabri)	Commercial,	2Q 2023 annual review: for CD, added TNFi criteria to allow bypass if member has had history
	HIM, Medicaid	of failure of two TNF blockers; for MS, to be inclusive of members continuing therapy from a
		different benefit, revised Medicaid/HIM continued approval duration to reference the duration of
		total treatment received rather than the number of re-authorizations; references reviewed and
		updated.
CP.PHAR.260 Rituximab (Rituxan,	Commercial,	2Q 2023 annual review: criteria added for off-label use in NS; for RA, added TNFi criteria to
Riabni, Ruxience, Truxima, Rituxan	HIM, Medicaid	allow bypass if member has had history of failure of two TNF blockers; removed nephrotic
Hycela)		syndrome in other diagnoses/indications section in initial and continued therapy; continued
		therapy approval duration for DM updated to 1 month; references reviewed and updated.
CP.PHAR.261 Secukinumab	Medicaid	2Q 2023 annual review: updated off-label dosing in Appendix B; for AS and PsA, added TNFi
(Cosentyx)		criteria to allow bypass if member has had history of failure of two TNF blockers; references
		reviewed and updated.
CP.PHAR.263 Tocilizumab (Actemra)	Medicaid	2Q 2023 annual review: RT4: revised criteria for COVID-19 emergency authorized use to FDA-
		approved indication; removed Appendix K since Actemra does not have EUA and is approved
		for COVID-19; updated off-label dosing in Appendix B; references reviewed and updated.
CP.PHAR.264 Ustekinumab (Stelara)	Medicaid	2Q 2023 annual review: updated off-label dosing in Appendix B; for CD, PsO, PsA, and UC,
		added TNFi criteria to allow bypass if member has had history of failure of two TNF blockers;
		references reviewed and updated.
CP.PHAR.265 Vedolizumab (Entyvio)	Medicaid	2Q 2023 annual review: for UC and CD, added TNFi criteria to allow bypass if member has had
		history of failure of two TNF blockers; updated off-label dosing for Appendix B; added high
		risk factors for postoperative occurrence to Appendix E to align with other CD policies;
	~	references reviewed and updated.
CP.PHAR.294 Osimertinib (Tagrisso)		2Q 2023 annual review: for NSCLC adjuvant treatment updated allowable stages from stage IB-
	HIM, Medicaid	IIIA to stage IB–IIIB per NCCN off-label support; references reviewed and updated.
CP.PHAR.306 Ofatumumab (Arzerra,	Commercial,	2Q 2023 annual review: for Arzerra, removed B-cell lymphoma criteria, SLL criteria, and off-
Kesimpta)	HIM, Medicaid	label CLL uses per updated NCCN guidelines and limited commercial availability; for
		Kesimpta, applied template changes to continued therapy section, and for MS, to be inclusive of
		members continuing therapy from a different benefit, revised Medicaid/HIM continued approval



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		duration to reference the duration of total treatment received rather than the number of re-
		authorizations; references reviewed and updated.
CP.PHAR.319 Ipilimumab (Yervoy)	Commercial,	2Q 2023 annual review: updated FDA indication for RCC to mirror PI; revised NSCLC criteria
	HIM, Medicaid	to include additional requirements related to mutation status, added off-label use for MSI-
		H/dMMR ampullary adenocarcinoma, bone cancer, brain metastases, and Kaposi sarcoma per
		NCCN compendium; references reviewed and updated.
CP.PHAR.339 Durvalumab (Imfinzi)	Commercial,	2Q 2023 annual review: for NSCLC per NCCN Compendium added recurrent or advanced
	HIM, Medicaid	disease and additional actionable molecular biomarkers that could be negative for use in
		combination with Imjudo and platinum therapy, added off-label continuation maintenance
		therapy; added off-label use for cervical cancer; clarified maximum 12 month continued
		approval duration applies only to stage II-III NSCLC; references reviewed and updated.
CP.PHAR.342 Brigatinib (Alunbrig)	Commercial,	2Q 2023 annual review: added off-label NCCN-supported indications of Erdheim-Chester
	HIM, Medicaid	disease and uterine sarcoma; references reviewed and updated. Template verbiage pertaining to
	,	continued therapy does not apply.
CP.PHAR.349 Ceritinib (Zykadia)	Commercial,	2Q 2023 annual review: added off-label NCCN-supported indications of Erdheim-Chester
	HIM, Medicaid	disease and uterine sarcoma; removed capsule formulation as this was discontinued; references
	Timit, madada	reviewed and updated. Template verbiage pertaining to continued therapy section does not
		apply.
CP.PHAR.364 Guselkumab (Tremfya)	Medicaid	2Q 2023 annual review: for PsA, added TNFi criteria to allow bypass if member has had history
		of failure of two TNF blockers; updated dosing in Appendix B to reflect dosing for redirected
		indications; references reviewed and updated
CP.PHAR.369 Alectinib (Alecensa)	Commercial,	2Q 2023 annual review: added off-label NCCN-supported indications of diffuse large B-cell
()	HIM, Medicaid	lymphoma, Erdheim-Chester disease, and uterine sarcoma; references reviewed and updated.
	1111111, 11110 0110 0110	Template verbiage pertaining to continued therapy section does not apply.
CP.PHAR.380 Cobimetinib (Cotellic)	Commercial,	2Q 2023 annual review: for Melanoma criteria, added stage III melanoma as adjuvant therapy,
	HIM, Medicaid	limited resectable melanoma, and requirements for trial of Tafinlar/Mekinist, updated off-label
	1111111, 11110 0110 0110	criteria for CNS cancers to include WHO grade 2, or 3 adult oligodendroglioma and WHO grade
		2, 3, or 4 adult IDH-mutant astrocytoma, per NCCN-supported 2A recommendation; removed
		anaplastic glioma and grade 2 glioma as terminology is no longer used in NCCN compendium;
		references reviewed and updated.
CP.PHAR.385 Corticosteroids for	Commercial,	Added Dextenza to policy; revised dosing frequency for Ozurdex from q4 months to q3 months
ophthalmic injection (Dextenza,	HIM, Medicaid	per literature review, market analysis, and specialist feedback; updated HCPCS code for Xipere.
Iluvien, Ozurdex, Retisert, Xipere,		r, manus analysis, and specialist reducin, aparted fields code for rapele.
Yutiq)		
CP.PHAR.386 Tildrakizumab-asmn	Medicaid	2Q 2023 annual review: added HCPCS code; for PsO, added TNFi criteria to allow bypass if
(Ilumya)		member has had history of failure of two TNF blockers; references reviewed and updated.
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CP.PHAR.406 Lorlatinib (Lorbrena)	Commercial,	2Q 2023 annual review: added off-label NCCN-supported indications of diffuse large B-cell
	HIM, Medicaid	lymphoma, Erdheim-Chester disease, IMT, and uterine sarcoma; references reviewed and
		updated.
CP.PHAR.417 Brexanolone (Zulresso)	Commercial,	2Q 2023 annual review: shortened the trial durations of antidepressant agent from 8 weeks to 4
	HIM, Medicaid	weeks; references reviewed and updated. Template verbiage pertaining to continued therapy
		does not apply.
CP.PHAR.418 Dexrazoxane (Totect)	Commercial,	2Q 2023 annual review: updated FDA approved indication to mirror PI; clarified that use is
	HIM, Medicaid	limited to the pediatric population for Ph-negative ALL and Hodgkin lymphoma; added off-label
		use for soft tissue sarcoma to criteria under doxorubicin-induced cardiomyopathy per NCCN 2A
		recommendation; reference reviewed and updated.
CP.PHAR.419 Elapegademase-lvlr	Commercial,	2Q 2023 annual review: added hematologist specialty option to criteria; references reviewed and
(Revcovi)	HIM, Medicaid	updated.
CP.PHAR.426 Risankizumab-rzaa	Medicaid	2Q 2023 annual review: updated off-label dosing in Appendix B; for PsA and CD, added TNFi
(Skyrizi)	1,10d10d1d	criteria to allow bypass if member has had history of failure of two TNF blockers; references
(SKJ1121)		reviewed and updated.
CP.PHAR.443 Upadacitinib (Rinvoq)	Medicaid	2Q 2023 annual review: for RA, PsA, AS, and UC, added TNFi criteria to allow bypass if
CI.III/IIC.443 Opadacitimo (Kinivoq)	Wicdicaid	member has had history of failure of two TNF blockers; updated off-label dosing for Appendix
		B; references reviewed and updated.
CP.PHAR.447 Mercaptopurine	Commercial,	2Q 2023 annual review: added by-passing of redirection if state regulations do not allow step
	HIM, Medicaid	therapy in certain oncology settings; clarified HIM approval durations align with Medicaid;
(Purixan)	HIM, Medicald	
		references reviewed and updated. Template verbiage pertaining to continued therapy does not
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CP.PHAR.462 Ozanimod (Zeposia)	Commercial,	2Q 2023 annual review: for UC, added TNFi criteria to allow bypass if member has had history
	HIM, Medicaid	of failure of two TNF blockers; for MS, to be inclusive of members continuing therapy from a
		different benefit, revised continued approval duration to reference the duration of total treatment
		received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.481 Idecabtagene Vicleucel	Commercial,	2Q 2023 annual review: added additional option to currently required measurable disease
(Abecma)	HIM, Medicaid	requirement to allow for progressive disease as defined by IMWG; clarified requirement for
		diagnosis of <i>relapsed or refractory</i> multiple myeloma; references reviewed and updated.
CP.PHAR.503 Sutimlimab-jome	Commercial,	2Q 2023 annual review: RT4: removed requirement for history of at least one documented blood
(Enjaymo)	HIM, Medicaid	transfusion within 6 months (initial criteria), revised required increase in hemoglobin level from
		2 to 1.5 g/dL (continued criteria), and modified evidence of positive response from being both of
		the following to just one of the following per revised FDA indication and new data from the
		CADENZA study; corrected hemoglobin-related continued criteria from > to ≥ per pivotal trial
		design; removed inactive HCPCS codes; references reviewed and updated.



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CP.PHAR.529 Relugolix (Orgovyx),	Commercial,	2Q 2023 annual review: for prostate cancer, removed specific diagnosis characteristic
relugolix-estradiol-norethindrone	HIM, Medicaid	requirements to align with current approach for the other GnRH agents (all are recommended for
(Myfembree)		use in the same place in therapy per NCCN); references reviewed and updated.
CP.PHAR.530 Tepotinib (Tepmetko)	Commercial,	2Q 2023 annual review: For NSCLC that is <i>MET</i> exon 14 skipping-positive, added exclusion for
	HIM, Medicaid	previous progression with a MET exon 14 skipping mutation-targeted regimen per NCCN
		Compendium and New Century Health criteria; added monotherapy criterion per NCCN and
		New Century Health criteria; references reviewed and updated.
CP.PHAR.531 Umbralisib (Ukoniq)	Commercial,	2Q 2023 annual review: removed initial approval criteria for marginal zone lymphoma and
	HIM, Medicaid	follicular lymphoma as use is not supported by the FDA and NCCN; removed table from
		Appendix B; references reviewed and updated.
CP.PHAR.533 Ciltacabtagene	Commercial,	2Q 2023 annual review: added additional option to currently required measurable disease
Autoleucel (Carvykti)	HIM, Medicaid	requirement to allow for progressive disease as defined by IMWG; clarified requirement for
, ,		diagnosis of relapsed or refractory multiple myeloma; removed J9999 HCPCS code; references
		reviewed and updated.
CP.PHAR.538 Tivozanib (Fotivda)	Commercial,	2Q 2023 annual review: clarified requirement that RCC is of clear cell histology per NCCN and
	HIM, Medicaid	pivotal clinical trial inclusion criteria, updated <i>Appendix B</i> to remove references to regimens for
	,	non-clear cell histology; references reviewed and updated.
CP.PHAR.577 Tralokinumab-ldrm	Commercial,	2Q 2023 annual review: modified list of agents for which concurrent use is not allowed to
(Adbry)	HIM, Medicaid	include non-asthma biologic immunomodulators; clarified that topical corticosteroids
	,	requirement is for corticosteroids of different molecular identities and expanded examples of
		medium to very high potency topical corticosteroids in Appendix B; removed low potency
		topical corticosteroids from Appendix B; references reviewed and updated.
CP.PHAR.582 Lutetium Lu 177	Commercial,	2Q 2023 annual review: added clarification to approval duration is for up to a total of 6 doses;
vipivotide tetraxetan (Pluvicto)	HIM, Medicaid	revised continued therapy approval duration from 12 to 6 months; for continued therapy added
vipivonae tenanetan (1 iavieto)	Timvi, ivicalcula	requirement that member has not received ≥ 6 doses (infusions) of Pluvicto; added piflufolastat
		F-18 as an additional radioactive diagnostic agent for identification of PSMA-positive disease;
		updated <i>Appendix D</i> examples of androgen deprivation therapy per NCCN; removed inactive
		HCPCS code A9699; references reviewed and updated.
CP.PHAR.583 Pacritinib (Vonjo)	Commercial,	2Q 2023 annual review: for MF added criteria for lower-risk disease per NCCN 2A
C1.111ARC.363 1 actitimo (vonjo)	HIM, Medicaid	recommendation and added criteria for higher-risk disease with platelets $\geq 50 \times 10^9$ /L per NCCN
	1111vi, ivicuicalu	1 recommendation; for continued therapy section updated FDA maximum dosing to mirror PI;
		provided details on risk stratification in Appendix D; references reviewed and updated.
		New
CP.PHAR.619 Nedosiran (DCR-	Commercial,	Policy created premeptively
PHXC)	HIM, Medicaid	1 oney created premeptivery
CP.PHAR.620 Pirtobrutinib (Jaypirca)	Commercial,	Policy created
C1.111AK.020 1 Intoolutiiio (saypiica)	HIM, Medicaid	1 oney created
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CP.PHAR.621 Ublituximab-xiiy	Commercial,	Policy created: adapted from existing criteria for non-preferred MS agents in line with prior
(Briumvi)	HIM, Medicaid	SDC recommendations/P&T approved clinical guidance.
CP.PHAR.622 Lenacapavir (Sunlenca)	Commercial,	Policy created
	HIM, Medicaid	
CP.PHAR.623 Elacestrant (Orserdu)	Commercial,	Policy created
	HIM, Medicaid	
CP.PHAR.624 Ferric Pyrophosphate	Commercial,	Policy created per February SDC.
Citrate (Triferic)	HIM, Medicaid	
	1	No Significant Change(s)
CP.PHAR.43 Sapropterin (Kuvan)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.65 Imatinib (Gleevec)	Commercial,	2Q 2023 annual review: no significant changes; clarified for melanoma imatinib should be used
	HIM, Medicaid	following BRAF-targeted therapy; references reviewed and updated.
CP.PHAR.68 Gefitinib (Iressa)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.75 Bexarotene (Targretin	Commercial,	2Q 2023 annual review: no significant changes; removed off-label criteria related to mycosis
Capsules, Gel)	HIM, Medicaid	fungoides/Sezary syndrome as those are subtypes of CTCL, an already covered FDA approved
		indication; references reviewed and updated.
CP.PHAR.88 Belimumab (Benlysta)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.92 Tetrabenazine	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Xenazine)	HIM, Medicaid	
CP.PHAR.105 Bosutinib (Bosulif)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.108 Omecetaxine (Synribo)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.120 Sipuleucel-T	Commercial,	2Q 2023 annual review: no significant changes; updated <i>Appendix D</i> examples of androgen
(Provenge)	HIM, Medicaid	deprivation therapy per NCCN; clarified <i>estimated</i> life expectancy of > 6 months requirement
		consistent with NCCN; references reviewed and updated.
CP.PHAR.135 Baricitinib (Olumiant)	Medicaid	2Q 2023 annual review: no significant changes; updated off-label dosing in Appendix B;
		references reviewed and updated.
CP.PHAR.152 Laronidase	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Aldurazyme)	HIM, Medicaid	
CP.PHAR.153 Eliglustat (Cerdelga)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	



CP.PHAR.154 Imiglucerase	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Cerezyme)	HIM, Medicaid	
CP.PHAR.155 Cysteamine oral	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Cystagon, Procysbi)	HIM, Medicaid	
CP.PHAR.156 Idursulfase (Elaprase)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.158 Agalsidase beta	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Fabrazyme)	HIM, Medicaid	
CP.PHAR.159 Sebelipase alfa	Commercial,	2Q 2023 annual review: no significant changes; added definition of "suboptimal clinical
(Kanuma)	HIM, Medicaid	response" for determining the need for further dose increases; references reviewed and updated.
CP.PHAR.160 Alglucosidase	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Lumizyme)	HIM, Medicaid	
CP.PHAR.161 Galsulfase (Naglazyme)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
, , , , , , , , , , , , , , , , , , ,	HIM, Medicaid	
CP.PHAR.162 Elosulfase alfa	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Vimizim)	HIM, Medicaid	
CP.PHAR.164 Miglustat (Zavesca)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.229 Ado-trastuzumab	Commercial,	2Q 2023 annual review: no significant changes; clarified for NSCLC that disease is recurrent,
(Kadcyla)	HIM, Medicaid	advanced, or metastatic per NCCN; references reviewed and updated.
CP.PHAR.230 AbobotulinumtoxinA	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Dysport)	HIM, Medicaid	
CP.PHAR.238 Methoxy polyethylene	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
glycol-epoetin beta (Mircera)	HIM, Medicaid	
CP.PHAR.242 Adalimumab (Humira)	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
Humira Biosimilars		
CP.PHAR.243 Alemtuzumab	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Lemtrada)	HIM, Medicaid	
CP.PHAR.245 Apremilast (Otezla)	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
CP.PHAR.248 Dalfampridine	Commercial,	2Q 2023 annual review: no significant changes; added generic redirection to continued therapy
(Ampyra)	HIM, Medicaid	section; references reviewed and updated.
CP.PHAR.249 Dimethyl fumarate	Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy
(Tecfidera), diroximel fumarate,		from a different benefit, revised continued approval duration to reference the duration of total
monomethyl fumarate		treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.250 Etanercept (Enbrel)	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.



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CP.PHAR.251 Fingolimod (Gilenya, Tascenso ODT)	Medicaid	2Q 2023 annual review: no significant changes; added redirection to generic for Gilenya 0.5 mg requests per SDC; RT4: for Tascenso ODT, added new 0.5 mg dosage strength and updated indication/criteria to remove prior upper age limit and weight requirement per revised FDA labeling; to be inclusive of members continuing therapy from a different benefit, revised continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.252 Glatiramer (Copaxone, Glatopa)	Commercial, Medicaid	2Q 2023 annual review: no significant changes; added generic redirection to continued therapy section; to be inclusive of members continuing therapy from a different benefit, revised Medicaid continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.254 Infliximab (Avsola, Inflectra, Remicade, Renflexis)	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
CP.PHAR.255 Interferon beta-1a (Avonex, Rebif)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy from a different benefit, revised Medicaid/HIM continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.256 Interferon beta-1b (Betaseron, Extavia)	Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy from a different benefit, revised continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.257 Ixekizumab (Taltz)	Medicaid	2Q 2023 annual review: no significant changes; updated off-label dosing for Appendix B; references reviewed and updated.
CP.PHAR.258 Mitoxantrone (Novantrone)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; clarified lymphoma criteria per NCCN; references reviewed and updated.
CP.PHAR.262 Teriflunomide (Aubagio)	Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy from a different benefit, revised continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.266 Rilonacept (Arcalyst)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; updated appendix B dosing for recurrent pericarditis to align with 2020 JACC Management of Acute and Recurrent Pericarditis guideline; references reviewed and updated.
CP.PHAR.267 Tofacitinib (Xeljanz Xeljanz XR)	Medicaid	2Q 2023 annual review: no significant changes; updated off-label dosing in Appendix B; references reviewed and updated.
CP.PHAR.271 Peginterferon beta-1a (Plegridy)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy from a different benefit, revised Medicaid/HIM continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.272 Sonidegib (Odomzo)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated. Template verbiage per continued therapy section does not apply.



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CP.PHAR.273 Vismodegib (Erivedge)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated. Template
	HIM, Medicaid	verbiage does not apply to continued therapy section.
CP.PHAR.298 Afatinib (Gilotrif)	Commercial,	2Q 2023 annual review: no significant changes; added Tagrisso as an example of EGFR tyrosine
	HIM, Medicaid	kinase inhibitor therapy within criteria per NCCN NSCLC treatment algorithm; references
		reviewed and updated. Template verbiage not applicable to continued therapy section.
CP.PHAR.316 Cabazitaxel (Jevtana)	HIM, Medicaid	2Q 2023 annual review: no significant changes; RT4 – added 45 mg/4.5 mL and 60 mg/6 mL
		concentrations; updated <i>Appendix D</i> examples of androgen deprivation therapy per NCCN;
		removed 60 mg/3 mL dose form as product was discontinued; references reviewed and updated.
		Template verbiage does not apply to continued therapy section.
CP.PHAR.335 Ocrelizumab (Ocrevus)	Commercial,	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy
	HIM, Medicaid	from a different benefit, revised Medicaid/HIM continued approval duration to reference the
		duration of total treatment received rather than the number of re-authorizations; references
		reviewed and updated.
CP.PHAR.337 Telotristat ethyl	Commercial,	2Q 2023 annual review: no significant changes; added redirection to generic telotristat for brand
(Xermelo)	HIM, Medicaid	Xermelo requests; updated Appendix C to include contraindication per PI; references reviewed
		and updated.
CP.PHAR.340 Valbenazine (Ingrezza)	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
CP.PHAR.341 Deutetrabenazine	Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Austedo)		
CP.PHAR.343 Edaravone (Radicava)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.344 Midostaurin (Rydapt)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated. Template
	HIM, Medicaid	verbiage pertaining to continued therapy section does not apply.
CP.PHAR.346 Sarilumab (Kevzara)	Medicaid	2Q 2023 annual review: no significant changes; updated off-label dosing for Appendix B;
		references reviewed and updated.
CP.PHAR.374 Vestronidase alfa-vjbk	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Mepsevii)	HIM, Medicaid	
CP.PHAR.375 Brodalumab (Siliq)	Medicaid	2Q 2023 annual review: no significant changes; updated Appendix B to include all relevant
		formulations of MTX; references reviewed and updated.
CP.PHAR.376 Apalutamide (Erleada)	Medicaid	2Q 2023 annual review: no significant changes; updated <i>Appendix D</i> examples of androgen
		deprivation therapy per NCCN; references reviewed and updated.
CP.PHAR.378 Ibalizumab-uiyk	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Trogarzo)	HIM, Medicaid	
CP.PHAR.416 Caplacizumab-yhdp	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Cablivi)	HIM, Medicaid	



CP.PHAR.421 Onasemnogene	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated. Template
Abeoarvovec (Zolgensma)	HIM, Medicaid	verbiage pertaining to continued therapy does not apply.
CP.PHAR.422 Cladribine (Mavenclad)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.427 Siponimod (Mayzent)	Commercial,	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy
	HIM, Medicaid	from a different benefit, revised continued approval duration to reference the duration of total
		treatment received rather than the number of re-authorizations; references reviewed and updated.
CP.PHAR.468 Aducanumab-avwa	HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Aduhelm)		
CP.PHAR.469 Belantamab Mafodotin-	Commercial,	2Q 2023 annual review: no significant changes, removed inactive HCPCS code C9069;
blmf (Blenrep)	HIM, Medicaid	references reviewed and updated.
CP.PHAR.471 Fosdenopterin (Nulibry)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
	HIM, Medicaid	
CP.PHAR.474 Remestemcel-L	Commercial,	2Q 2023 annual review: no significant changes as drug is not yet FDA-approved; references
(Ryoncil)	HIM, Medicaid	reviewed and updated.
	,	
CP.PHAR.475 Sacituzumab	Commercial,	2Q 2023 annual review: no significant changes; updated commercial LOB approval language to
Govitecan-hziy (Trodelvy)	HIM, Medicaid	standard language with addition of "whichever is longer"; references reviewed and updated.
	,	RT4: added new indication for treatment of HR-positive, HER2-negative breast cancer who have
		received endocrine-based therapy and at least two additional systemic therapies in the metastatic
		setting.
CP.PHAR.478 Selpercatinib (Retevmo)	Commercial,	2Q 2023 annual review: no significant changes; for thyroid cancer, removed requirement that
,	HIM, Medicaid	disease is not amenable to radioactive iodine therapy for DTC as this is redundant with
	,	immediately preceding criterion; references reviewed and updated.
CP.PHAR.479	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
Decitabine/Cedazuridine (Inqovi)	HIM, Medicaid	
CP.PHAR.482 Isatuximab-irfc	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Sarclisa)	HIM, Medicaid	
CP.PHAR.483 Lisocabtagene	Commercial,	2Q 2023 annual review: no significant changes; modified AIDS-related DLBCL to HIV-related
Maraleucel (Breyanzi)	HIM, Medicaid	per NCCN Compendium; references reviewed and updated.
CP.PHAR.486 Bimatoprost Implant	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(Durysta)	HIM, Medicaid	
CP.PHAR.504 Voclosporin (Lupkynis)	Commercial,	2Q 2023 annual review: no significant changes; references reviewed and updated.
(30pu)1115)	HIM, Medicaid	
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CP.PHAR.514 Pralsetinib (Gavreto)	Commercial, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.521 Avalglucosidase Alfangpt (Nexviazyme)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.526 Fibrinogen Concentrate [Human] (Fibryga, RiaSTAP)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.527 Narsoplimab (OMS721)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes as drug is not yet FDA-approved; references reviewed and updated.		
CP.PHAR.528 Odevixibat (Bylvay)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.534 Insulin Delivery Systems (V-Go, Omnipod, InPen)	Commercial, HIM-Medical Benefit, Medicaid	2Q 2023 annual review: no significant changes; for V-Go, revised minimum age requirement from 21 years to 18 years per user guide; references reviewed and updated.		
CP.PHAR.535 Melphalan Flufenamide (Pepaxto)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.536 Ophthalmic Riboflavin (Photrexa, Photrexa Viscous)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
CP.PHAR.537 Ponesimod (Ponvory)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; to be inclusive of members continuing therapy from a different benefit, revised continued approval duration to reference the duration of total treatment received rather than the number of re-authorizations; references reviewed and updated.		
CP.PHAR.575 Tebentafusp-tebn (Kimmtrak)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; removed inactive HCPCS codes C9095, J9999; references reviewed and updated.		
CP.PHAR.581 Faricimab (Vabysmo)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes, removed inactive HCPCS codes J3590 and C9097; references reviewed and updated.		
CP.PHAR.584 Sodium Phenylbutyrate/Taurursodiol (Relyvrio)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated.		
Strategy Development Committee (SDC) Criteria changes based on SDC decisions				
CP.PHAR.157 Taliglucerase alfa (Elelyso)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; references reviewed and updated. Per February SDC and prior clinical guidance, added redirections to Cerdelga and Cerezyme; added Appendix B; added HIM line of business and retired HIM.PA.162.		
CP.PHAR.163 Velaglucerase alfa (VPRIV)	Commercial, HIM, Medicaid	2Q 2023 annual review: no significant changes; added weight requirement and max dose limits to align with previously Corporate P&T-approved approach for max dose limits when switching from imiglucerase; references reviewed and updated. Per February SDC and prior clinical		



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		guidance, added redirections to Cerdelga and Cerezyme; added Appendix B; added HIM line of business and retired HIM.PA.163.
CP.PHAR.165 Ferumoxytol	Commercial,	Per February SDC, added Commercial line of business; updated initial criteria to require failure
(Feraheme)	HIM, Medicaid	of the following: for IDA and CKD Ferrlecit and Venofer; for IDA without CKD two of
		Ferrlecit, Infed, or Venofer.
CP.PHAR.200 Mepolizumab (Nucala)	Commercial,	Per February SDC, for CRSwNP modified requirement from three intranasal steroids to require
1 /	HIM, Medicaid	only two.
CP.PHAR.231 IncobotulinumtoxinA	Commercial,	2Q 2023 annual review: Per February SDC and prior clinical guidance, added redirection
(Xeomin)	HIM, Medicaid	requirement to co-prefer Botox and Dysport for all indications except chronic sialorrhea;
		references reviewed and updated.
CP.PHAR.232 OnabotulinumtoxinA	Commercial,	2Q 2023 annual review: for chronic anal fissure, revised maximum dosing allowance up to 25
(Botox)	HIM, Medicaid	units for initial therapy and 100 units for continued therapy per treatment session; added chronic
		sialorrhea off-label indication; references reviewed and updated. Per February SDC: removed
		Dysport and/or Xeomin redirection requirement for upper and lower limp spasticity, cervical
		dystonia, blepharospasm, overactive bladder, chronic migraine, and axillary hyperhidrosis; for
		Overactive Bladder, updated criteria for adults to require use of two anticholinergic agents or
		one oral beta-3 agonist medication (previously both were required); changed all approval
		durations to 12 months for Medicaid, HIM, and Commercial.
CP.PHAR.233 RimabotulinumtoxinB	Commercial,	2Q 2023 annual review: Per February SDC and prior clinical guidance, for cervical dystonia
(Myobloc)	HIM, Medicaid	replaced Xeomin redirection with Botox to co-prefer with Dysport; references reviewed and
		updated.
CP.PHAR.234 Ferric Carboxymaltose	Commercial,	Per February SDC, added Commercial line of business; updated initial criteria to require failure
(Injectafer)	HIM, Medicaid	of the following with associated age considerations: for IDA and CKD Ferrlecit and Venofer; for
		IDA without CKD two of Ferrlecit, Infed, or Venofer; additionally, added redirection to
		Feraheme in a step-wise fashion if member has intolerance or contraindication to all preferred
CD DILL D 200 CD CT OL 1		injectable agents.
CP.PHAR.296 Pegfilgrastim (Neulasta	Commercial,	Per February SDC and prior clinical guidance, added Udenyca as step through requirement to
and biosimilars)	HIM, Medicaid	co-prefer with Ziextenzo.
CP.PHAR.336 Dupilumab (Dupixent)	Commercial,	Per February SDC, for CRSwNP modified requirement from three intranasal steroids to require
CD DILA D 201 I	HIM, Medicaid	only two.
CP.PHAR.391 Lanreotide (Somatuline	Commercial,	Per February SDC and prior clinical guidance added redirection to Sandostatin LAR depot.
Depot)	HIM, Medicaid	20 2022 amount assistant Dan Enhancem CDC and total initial anitonic to accoming fails.
CP.PHAR.480 Ferric Derisomaltose	Commercial,	2Q 2023 annual review: Per February SDC, updated initial criteria to require failure of the
(Monoferric	HIM, Medicaid	following: for IDA and CKD Ferrlecit and Venofer; for IDA without CKD two of Ferrlecit,
		Infed, or Venofer; additionally, added redirection to Feraheme in a step-wise fashion if member
		has intolerance or contraindication to all preferred injectable agents; references reviewed and
		updated.



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Retired				
CP.PHAR.166 Ferric Gluconate	One of the preferred products per Feb 2023 SDC and medical PA requirement will be removed			
(Ferrlecit)				
CP.PHAR.167 Iron Sucrose (Venofer)	One of the preferred products per Feb 2023 SDC and medical PA requirement will be removed			
CP.PHAR.520 Casirivimab and Imdevimab (REGEN-COV)	Nearing the end of the public health emergency period, and these antibodies are no longer recommended for use due to low/minimal effectiveness against current variants of concern			
CP.PHAR.532 Bamlanivimab + Etesevimab (LY-CoV555 + LY-CoV016)	Nearing the end of the public health emergency period, and these antibodies are no longer recommended for use due to low/minimal effectiveness against current variants of concern			
CP.PHAR.541 Sotrovimab (VIR-7831)	Nearing the end of the public health emergency period, and these antibodies are no longer recommended for use due to low/minimal effectiveness against current variants of concern			
CP.PHAR.571 Tixagevimab and	Nearing the end of the public health emergency period, and these antibodies are no longer			
Cilgavimab (Evusheld)	recommended for use due to low/minimal effectiveness against current variants of concern			

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