Oregon Health Authority

2020 Mental Health Parity Analysis Report

for

InterCommunity Health Network

February 2021





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Overview of Oregon's Mental Health Parity Analysis

Mental Health Parity (MHP) regulations are intended to ensure that coverage and access to services for the treatment of mental health (MH) and substance use disorder (SUD) conditions are provided in parity with treatments provided for medical and surgical (M/S) needs. The required analysis of MH benefits is governed by federal regulations. The Mental Health Parity and Addiction Equity Act of 2008 (MHPAEA) governs how MH/SUD treatments delivered by managed care organizations and limitations on MH/SUD benefits are comparable to and applied no more stringently than the limitations applied to M/S benefits. Provisions of the MHPAEA became applicable to the Oregon Health Plan (OHP) in October 2017 when the Medicaid Parity Final Rule (42 Code of Federal Regulations [CFR] §438 Subpart K) went into effect. The rule requires parity in key areas:

- Aggregate lifetime and annual dollar limits (AL/ADLs).
- Financial requirements (FRs—such as copays).
- Quantitative treatment limitations (QTLs—such as day and visit limits).
- Non-quantitative treatment limitations (NQTLs—such as prior authorization [PA] and provider network admission requirements).

Additional MHP regulations require that criteria for medical necessity determinations for MH/SUD benefits must be made available to beneficiaries and providers upon request, as well as the reason for denial of reimbursement or payment for MH/SUD benefits. States must also implement monitoring procedures to ensure continued compliance and to identify when changes in benefit design or operations could affect compliance and require an updated analysis.

To meet the requirements, the Oregon Health Authority (OHA) conducted an initial MHP Analysis of OHP's full delivery system in 2018. OHA's 15 coordinated care organizations (CCOs) and Oregon Health Plan Fee-for-Service (OHP FFS) participated in the initial MHP Analysis, which included an inventory of all MH/SUD and M/S benefits offered to OHP members and the limitations applied to those benefits to ensure that limitations (e.g., day limits, PA requirements, or network admission standards) for MH and SUD services were comparable to and applied no more stringently than those for M/S services provided under OHP. Results of the initial analysis were reported in August 2018; and in 2019, the CCOs implemented corrective actions in areas lacking parity.

For 2020, OHA tasked Health Services Advisory Group, Inc. (HSAG), with conducting a follow-up MHP Analysis across the CCOs, in part due to each of the CCOs entering into new five-year contracts with the State, to determine if the existing benefits and any NQTLs remained compliant with the MHP regulations in 42 CFR §438 Subpart K. HSAG conducted the MHP Analysis in 2020 based on the August 2018 results, any implemented corrective actions, and any additional changes to benefits design or operations that may impact parity. This report provides information on and results of the 2020 MHP Analysis for InterCommunity Health Network (IHN).



Components of the 2020 MHP Analysis

In accordance with 42 CFR §438 Subpart K, MHP applied to all OHP benefits delivered through OHA's managed care delivery system, including those delivered through a combination of managed care and FFS delivery systems. HSAG developed a protocol and tools to carry out the analysis activity based on the initial 2018 MHP Analysis and in alignment with guidance outlined in the toolkit provided by the Centers for Medicare & Medicaid Services (CMS): *Parity Compliance Toolkit Applying Mental Health and Substance Use Disorder Parity Requirements to Medicaid and Children's Health Insurance Programs*. ¹⁻¹ The 2020 MHP Analysis also referenced Oregon's Mapping Guide ¹⁻² that assigned benefits to MH/SUD and M/S groupings based on International Classification of Diseases, Tenth Revision (ICD-10) diagnoses and mapped into four prescribed classifications as published in the March 30, 2016, Federal Register, Vol. 81, No. 61¹⁻³ as illustrated in Figure 1-1.

Figure 1-1—MHP: Four Prescribed Classifications

Inpatient Outpatient Prescription Drug Emergency Care

OHP Benefit Packages

While all OHP benefit packages were delivered in accordance with the same Medicaid essential health benefits structure, the delivery of those benefits was categorized by OHP benefit package based on enrollment. Table 1-1 identifies the four OHP benefit packages evaluated in the 2020 MHP Analysis. Since each benefit package involves the delivery of Medicaid essential health benefits covered by both CCOs and OHP FFS, HSAG conducted an analysis of each CCO's NQTLs, and then against the OHP FFS NQTLs.

Table 1-1—OHP Benefit Packages

Benefit Package	Benefit Types Covered	Evaluation	
CCOA	Physical Health, Behavioral Health, Dental Health	CCO MH/SUD and FFS MH/SUD	
ССОВ	Physical Health, Behavioral Health	compared to CCO M/S	
CCOE	Behavioral Health	CCO MH/SUD and FFS MH/SUI	
CCOG	Behavioral Health, Dental Health	compared to FFS M/S	

¹⁻¹ The CMS *Parity Compliance Toolkit Applying Mental Health and Substance Use Disorder Parity Requirements to Medicaid and Children's Health Insurance Programs* and additional CMS resources related to MHP can be accessed at: https://www.medicaid.gov/medicaid/benefits/behavioral-health-services/parity/index.html.

¹⁻² The Oregon Mapping Guide includes definitions, links, and resources important for the MHP Analysis. It also maps all Oregon Medicaid benefits to the classifications required for the MHP Analysis. It can be accessed on OHA's MHP webpage at: https://www.oregon.gov/OHA/HSD/OHP/Pages/MH-Parity.aspx.

¹⁻³ Federal Register. Volume 81, No. 61/Wednesday, March 30, 2016. Available at: https://www.govinfo.gov/content/pkg/FR-2016-03-30/pdf/FR-2016-03-30.pdf. Accessed on: Dec 4, 2020.



Non-Quantitative Treatment Limitations

Because Oregon Medicaid does not permit the use of QTLs (e.g., day and visit limits), HSAG's analysis focused on assessing NQTLs in the OHP delivery system. NQTLs are health care management limitations on the scope or duration of benefits through the use of managed care processes, such as PA or network admission standards. "Soft limits," benefit limits that allow for an individual to exceed limits or allow for limits to be "waived" based on medical necessity, are also considered NQTLs. Examples of NQTLs include:

- Medical management standards limiting or excluding benefits based on medical necessity or appropriateness criteria.
- Standards for provider admission to participate in a network and reimbursement rates.
- Restrictions based on geographic location, facility type, or provider specialty.
- Fail-first policies or step therapy protocols.
- Exclusions based on failure to complete a course of treatment prior to allowing authorization of a subsequent treatment.

MHP regulations hold that no NQTL can be applied to MH/SUD benefits and services that is not comparable to or is more stringent than those applied to M/S benefits and services in each benefit classification regarding processes, strategies, evidentiary standards, or other factors. HSAG assessed policies and procedures as written and operational processes for compliance with parity requirements by classification (e.g., inpatient [IP] and outpatient [OP]) of services. The 2018 MHP Analysis compared NQTLs for services that address MH/SUD diagnoses with services that address M/S diagnoses across the OHP benefit packages. Comparability was assessed as to the reason an NQTL was used, the evidence that supported its use, and the process for its implementation. The stringency criterion assessed the rigor with which the NQTLs were applied, the evidence for the level of stringency, and penalties and exceptions associated with limitations. Comparability and stringency are defined in Figure 1-2.

NQTL ANALYSIS

STRINGENCY

The comparability of the processes, strategies, evidentiary standards, and other factors (in writing and in operation) used in applying the NQTL to MH/SUD benefits and M/S benefits.

STRINGENCY

The stringency with which the processes, strategies, evidentiary standards, and other factors (in writing and operation) are applied to MH/SUD benefits and M/S benefits.

Figure 1-2—MHP Analysis Comparability and Stringency



NQTL Categories

Similar to the Initial 2018 MHP Analysis, HSAG assessed for comparability and stringency criteria across six specific NQTL categories in the OHP delivery system. The six categories are described below.

- Category I—Utilization Management Limits Applied to Inpatient Services: Utilization management (UM) processes implemented through PA, concurrent review (CR), and retrospective review (RR) that may also be used to ensure medical necessity for MH/SUD and M/S services.
- Category II—Utilization Management Limits Applied to Outpatient Services: UM processes applied to OP MH/SUD and M/S services through PA, CR, and RR to ensure medical necessity.
- Category III—Prior Authorization for Prescription Drug Limits: PA as a means of determining whether particular medications will be dispensed. PA of prescription drugs limits the availability of specific medications.
- Category IV—Provider Admission—Closed Network: Closed networks as they impose limits to providers seeking to join a panel of approved providers.
- Category V—Provider Admission—Network Credentialing: Network enrollment/credentialing requirements imposed, including provider admission requirements such as state licensing requirements and exclusions of specific provider types, that may result in limitations.
- Category VI—Out-of-Network/Out-of-State Limits: Out-of-network (OON) and out-of-state (OOS) limits that affect how members access OON and OOS providers and address how OHA and the CCOs ensure necessary access to providers not eligible to be reimbursed or not in a CCO's network.



2. Process and Methodology

Building from the initial 2018 MHP Analysis, HSAG worked with OHA and the CCOs to conduct a follow-up MHP Analysis that evaluated changes to benefits design and operations that may impact parity. The 2020 MHP Analysis identified and addressed differences between the policies and standards governing limitations applied to MH/SUD services as compared to M/S services. Differences in how limits were applied to MH/SUD services as compared to M/S services were evaluated for continued compliance with MHP regulations to ensure evidence-based, quality MH/SUD care.

Analysis Activities for 2020

The 2020 MHP Analysis activities are illustrated in Figure 2-1 and described below.

Protocol and Tool Development/ Dissemination

Pre-Analysis Webinar

Documentation Submission

Desk Review

Conference Calls

Reporting Action Planning and Implementation

Figure 2-1—2020 MHP Analysis Activities

- 1. **Protocol and Tool Development and Dissemination:** HSAG developed and disseminated an MHP Analysis Protocol that presented details and guidance to OHA and CCOs on the analysis process and included tools in which to conduct the 2020 MHP Analysis Activity. The tools utilized for the analysis, identified below, were based on OHA's initial analysis of MHP and were developed using guidance outlined in the CMS *Parity Compliance Toolkit Applying Mental Health and Substance Use Disorder Parity Requirements to Medicaid and Children's Health Insurance Programs*.
 - MHP Evaluation Questionnaire—Questions referencing the six NQTL categories, to identify changes that may impact parity.
 - MHP Reporting Template—Documentation of changes and additions to NQTLs previously reported in 2018, organized by the six NQTL categories.
 - MHP Required Documentation Template—UM and credentialing data across MH/SUD and M/S benefits and providers.
- 2. **Pre-Analysis Webinar:** HSAG conducted a pre-analysis webinar on July 15, 2020, with OHA and the CCOs to provide an overview of MHP regulations, details of the protocol and tools, specifics of the analysis timeline, and examples of MHP scenarios for reference.
- 3. **Documentation Submission:** OHA and the CCOs were required to submit documentation that included responses to the MHP Evaluation Questionnaire and completed templates, along with supporting documentation, by August 31, 2020.



- 4. **Desk Review:** HSAG conducted a desk review of all submitted MHP Evaluation Questionnaires, the MHP Reporting Template, and required and supporting documentation (e.g., policies and procedures, benefit schedules, and delegate agreements) to analyze policies and operational practices that impact MHP and determine preliminary analysis findings.
- 5. **Conference Calls:** HSAG conducted conference calls to discuss preliminary analysis findings and areas in need of clarification. Additional information and documentation were requested at that time, as necessary to support the MHP Analysis.
- 6. **Reporting:** HSAG compiled analysis results and documented MHP determinations for each CCO and as compared to OHP FFS, identifying areas in which MHP had not been achieved and corrective actions were required to ensure future parity. OHA and each CCO had an opportunity to review report drafts prior to finalizing the reports.
- 7. **Corrective Action Planning and Implementation:** HSAG will work with OHA and the CCOs to develop and implement corrective action plans to achieve compliance with MHP requirements.

MHP Analysis Methodology

HSAG reviewers conducted a desk review of submitted MHP Analysis tools and supporting documentation to further clarify reported changes and additions to previously reported NQTLs from the initial MHP Analysis conducted in 2018. More specifically, HSAG evaluated responses to the MHP Evaluation Questionnaire to identify changes to benefits design and operations within OHA and each CCO that may impact MHP, cross-referenced questionnaire responses with changes and additions reported in the MHP Reporting Template, and reviewed supporting documentation submitted by OHA and the CCOs. Supporting documentation included, but was not limited to, UM policies, workflow diagrams, program descriptions, prescription drug formularies, and network admission/credentialing policies. HSAG conducted the 2020 MHP Analysis based on this information to determine compliance with parity guidelines, including ensuring that policies followed standard industry practice, allowed for little to no exception or variation, incorporated established State definitions and guidelines, included staff members qualified to make the decisions and complete the tasks assigned and appropriate oversight.

Information obtained via scheduled conference calls was also evaluated in relation to changes and additions reported. Differences in how limits were applied to MH/SUD services as compared to M/S services, in relation to comparability and stringency standards displayed in Table 2-1, were evaluated across the six NQTL categories for continued compliance with MHP regulations. Each CCO's NQTLs were additionally evaluated against OHP FFS MH/SUD and M/S NQTLs based on the structure of OHP benefit packages referenced in Section 1 of this report.



Table 2-1—Comparability and Stringency Standards

Comparability and Stringency Standard		Question Description
Benefits in Which NQTLs Apply	1.	To which benefits is an NQTL assigned? Purpose: To describe the NQTL assigned to MH/SUD and M/S benefits (e.g., PA, scope of services, and time frames).
Comparability of Strategy	2.	Why is the NQTL assigned to these benefits? Purpose: To describe for what reasons or purpose the NQTL is assigned (e.g., ensure medical necessity, prevent overutilization, and comply with State and federal requirements).
Comparability of Evidentiary Standard	3.	What evidence supports the rationale for the assignment? Purpose: To describe the evidence to support the rationale (e.g., benchmarks, standards that form the basis of the rationale, and State and federal requirements).
Comparability of Processes	4.	What are the NQTL procedures? Purpose: To describe the NQTL process and evidence needed to support NQTL determinations (e.g., documentation requirements, timelines, and steps for the CCO and members/providers).
Stringency of Strategy	5.	How frequently or strictly is the NQTL applied? Purpose: To describe the frequency of application, frequency of medical necessity and appropriateness reviews, level of discretion in how the NQTL is applied, triggers for review and re-review, etc.
Stringency of Evidentiary Standard	6.	What standard supports the frequency or rigor with which the NQTL is applied? Purpose: To describe standards that the CCO uses to determine the frequency or rigor of NQTL procedures.

Analysis Results for 2020

Results of the analysis are incorporated in Section 3 of this report. The results identify overall compliance with MHP regulations across the six NQTL categories in relation to comparability and stringency. Limitations or other operational processes found to impact parity are reported as findings. Required actions are also presented to support future compliance with MHP requirements as applicable.



3. MHP Analysis Results

HSAG derived 2020 MHP Analysis results from the evaluation and observation of information obtained from IHN. More specifically, the information and observations used for the evaluation included the following tools, documentation, and conversations:

- Responses to the 2020 MHP Evaluation Questionnaire.
- Reported data in the 2020 MHP Reporting Templates pertaining to NQTL categories.
- Information obtained from IHN's submitted 2020 MHP data using the Required Documentation Template and supporting documentation as provided.
- Observations from conversations during the conference call conducted with the CCO.

Results of the MHP Analysis are detailed below. Limitations or other operational processes found to impact parity are reported as findings, along with corresponding required actions. Appendices A and B include IHN's completed MHP questionnaire and finalized MHP reporting details by each NQTL category, respectively.

Overall Assessment

IHN was responsible for delivering MH/SUD and M/S Medicaid benefits to members in all four benefit packages (CCOA, CCOB, CCOE, and CCOG), whereas OHP FFS was fully managing M/S benefits for CCOE and CCOG benefit packages. IHN was providing MH/SUD benefits prescribed by OHA did not have UM delegation agreements with any other entity. HSAG evaluated IHN's application of NQTLs to MH/SUD and M/S benefits in terms of comparability and stringency across the six NQTL categories.

Most of IHN's policies and documentation identified standardized processes that applied to both MH/SUD and M/S benefits, including a UM and service authorization policy, a PA list, and an internal monitoring and oversight policy, among many others. The CCO did not have separate policies for the management of benefits based on benefit package (i.e., CCOA, CCOB, CCOE, and CCOG).

For limits applied to IP and OP health benefits, IHN and its delegates used UM processes to manage MH/SUD and M/S benefits. The purpose of the CCO's UM processes was to ensure coverage, medical necessity, appropriate treatment in the least restrictive environment that maintains the safety of the individual, compliance with federal and State requirements, and the prevention of unnecessary overutilization. IHN reported that the evidence used to apply UM to MH/SUD and M/S included Oregon Administrative Rules (OARs), Health Evidence Review Commission (HERC) Prioritized List (PL) and guidelines, and Milliman Care Guidelines (MCG). Both the CCO and OHP FFS allowed 90-day RR for MH/SUD and M/S when providers failed to obtain authorization, with some consideration to requests beyond the time frame. The application of authorization limits and the frequency and rigor in which they were applied to authorization requests was comparable across both MH/SUD and M/S benefits and to OHP FFS's application across both benefit types. Regarding interrater reliability (IRR), IHN conducted



regular reviews of MH/SUD and M/S authorizations using an 85 percent testing standard, which ensured a greater level of monitoring for consistency of authorization reviews than OHP FFS's annual M/S authorization reviews using an 80 percent testing standard. HSAG determined this discrepancy to be a quality issue for M/S benefit authorizations rather than a parity concern.

HSAG's analysis of IHN processes and operations did not reveal any MHP concerns for the authorization of prescription drugs across the benefit packages. The application of PA for MH/SUD prescription drugs was comparable to PA for M/S prescription drugs. Prescription drug authorization requirements, guidelines, procedures, and 24-hour responses were determined to be no more stringently applied to MH/SUD benefit requests than to M/S requests.

HSAG also did not identify any MHP concerns for the provider admission and OON/OOS NQTL categories based on an analysis of IHN's processes, strategies, and evidentiary standards. The CCO consistently applied credentialing and recredentialing processes to both MH/SUD and M/S providers, and no denials were reported for MH/SUD providers seeking credentialing during the reporting period. Similarly, the same PA processes and evidentiary standards described in NQTL categories I, II, and III were applied to OON and OOS coverage of MH/SUD and M/S benefits across the benefit packages, which also resulted in parity determination across both benefit types in the respective NQTL category. OHP FFS did not use memorandums of understanding (MOUs) to secure OON providers to deliver Medicaid benefits to members but instead enrolled providers. The inconsistency was not identified as a parity concern as the comparison of two different processes was determined to be inapplicable for the MHP analysis.

Overall, IHN's application of NQTLs for MH/SUD benefits across the NQTL categories and all benefits packages was determined to be comparable to and no more stringently applied than to M/S benefits. IHN achieved full compliance with MHP requirements, receiving no findings.

Table 3-1 presents HSAG's overall assessment of IHN's compliance based on the analysis of the comparability of NQTL strategies and the stringency applied by IHN when implementing NQTLs.

NQTL Category	Comparability	Stringency
Category I—UM Limits Applied to Inpatient Services	Compliant	Compliant
Category II—UM Limits Applied to Outpatient Services	Compliant	Compliant
Category III—Prior Authorization for Prescription Drug Limits	Compliant	Compliant
Category IV—Provider Admission—Closed Network	Compliant	Compliant
Category V—Provider Admission—Network Credentialing	Compliant	Compliant
Category VI—Out-of-Network/Out-of-State Limits	Compliant	Compliant

Table 3-1—Overall MHP Analysis Results—Comparability and Stringency



Findings and Required Actions

Based on the strategy and evidence provided by IHN, including reported changes in operations and practices, PA and credentialing data, and discussions during prescheduled conference calls, HSAG analyzed the parity of MH/SUD benefits as compared to M/S benefits. Overall, IHN's application of NQTLs for MH/SUD benefits across the NQTL categories and all benefits packages was determined to be comparable to and no more stringently applied than to M/S benefits. IHN achieved full compliance with MHP requirements, receiving no findings.

Data Analysis Results

IHN submitted UM data in the MHP Required Documentation Template, identifying PA counts and denial data for IP, OP, and prescription drug benefits. The reporting also included data on provider admission counts and terminations/denials. The completed templates included data from the period of January 1, 2020, through June 30, 2020. An analysis of the data reported is presented in the text below pertaining to the following categories:

- Utilization Management for Inpatient/Outpatient Services (NQTL Categories I and II).
- Utilization Management for Prescription Drugs (NQTL Category III).
- Enrollment/Credentialing Decisions (NQTL Categories IV and V).

Utilization Management for Inpatient/Outpatient Services

IHN provided requested UM data for IP and OP services pertaining to authorization request counts and outcomes of requests. Table 3-2 presents IHN's counts for IP and OP PAs by benefit type, identifying the number of PA requests denied, appealed, and overturned.

Prior Authorization Counts by Benefit Type # of PA % of PA # of PA % of PA # of PA % of PA **Denials Denials Denials Denials Benefit** # of PA Requests Requests Requests **Denied Denied Appealed Appealed** Overturned Overturned Type MH/SUD 1,142 63 5.52% 3 4.76% 0.00% M/S 57 5,269 593 11.25% 9.61% 26 4.38% 656 10.23% 60 **Total** 6,411 9.15% 26 3.96%

Table 3-2—Prior Authorization Counts for Inpatient and Outpatient Services

Observations

HSAG's analysis of IHN's PA data for IP and OP benefits did not reveal any concerns related to MHP due to a low rate of denials for MH/SUD PA requests. The following data points were observed:



- Of the total 6,411 IP and OP PA requests reported, 10.23 percent were denied.
- Of the 63 MH/SUD PA requests denied, representing 5.52 percent of the 1,142 MH/SUD PA requests, only three denials resulted in an appeal.
- The 63 reported MH/SUD denials represented 9.60 percent of total denials.

Utilization Management for Prescription Drugs

IHN provided requested data pertaining to prescription drug authorization request counts and outcomes. Table 3-3 presents IHN's PA counts for formulary and non-formulary prescription drug PA requests, identifying the number of requests overturned.

Prior Authorization Counts (Formulary and Non-Formulary) # of PA % of PA # of PA # of PA % of PA % of PA # of PA Requests Requests **Denials Denials** Denials **Denials Denied Appealed Overturned** Requests **Denied Appealed Overturned** 1,330 33.63% 50 3.76% 28 2.11% 3,955

Table 3-3—Prior Authorization Counts for Prescription Drugs

Observations

HSAG's analysis of IHN's counts for prescription drug PA requests did not reveal any concerns related to parity. The following data points were observed:

- Of the total 3,955 prescription drug PA requests reported, 33.63 percent were denied.
- Less than 4 percent (3.76 percent) of the 1,330 prescription drug PA request denials were appealed, with 28 PA denials resulting in an overturned decision.
- The majority of denied prescription drug PA requests were denied for a "not covered" categorical reason.

Enrollment/Credentialing

IHN provided requested data pertaining to provider enrollment requests and outcomes. Table 3-4 presents IHN's enrollment/credentialing counts by provider type, identifying the number of terminations and denials, which includes applications not accepted.



Table 3-4—Enrollment/Credentialing Counts by Provider Type

Enrollment/Credentialing Counts by Provider Type						
Provider Type	Avg. # Enrolled Providers	# Providers Terminated	% Terminated	# of Cred. Requests	# of Cred. Requests Denied/Not Accepted	% of Cred. Requests Denied/Not Accepted
MH/SUD	1,746	13	0.74%	51	13	25.49%
M/S	10,157	113	1.11%	255	55	21.57%
Total	11,903	126	1.06%	306	68	22.22%

Observations

HSAG's analysis of IHN's provider credentialing data did not reveal any parity concerns. The following data points were observed:

- Of the 11,903 reported average number of providers enrolled during the reporting period, 14.67 percent were MH/SUD providers.
- The total denial rate for all provider types was 22.22 percent
- Of the 51 MH/SUD providers seeking credentialing during the reporting period, 25.49 percent were denied credentialing, which was fairly comparable to the 21.57 percent M/S denial rate.
- While there were only 13 MH/SUD provider credentialing denials, 46.15 percent of those denials were due to an "out of area" denial reason.

Additional Requirement Results

HSAG requested information from IHN on the required availability of medical necessity determinations regarding MH/SUD benefits to members, potential members, and contracting providers upon request, and how reasons for denial of reimbursement or payment for MH/SUD benefits were made available to members. IHN policies outlined the CCO's notice of adverse benefit determination process and how notices are to include denial reasons for members. A review of IHN's website showed that the CCO had resources available on its website for members that included information on MH/SUD benefits available, a prescription drug formulary, and a prior authorization list. As a result, HSAG determined that IHN was compliant with the additional administrative MHP requirements.



4. Improvement Plan Process

To the extent MHP findings or concerns were found, OHP and all CCOs are required to complete and submit an improvement plan addressing corrective actions/interventions to resolve all MHP findings. The improvement plan template is provided in Appendix C. For each of the findings documented in Section 3 of this report, IHN must identify the following:

- Interventions planned by the organization to achieve MHP compliance.
- Individual(s) responsible for ensuring that the planned interventions are completed.
- Proposed timeline for completing each planned intervention with the understanding that most corrective actions/interventions can be completed within three months and no longer than six months. Corrective actions/interventions requiring additional time will need to include specific information to determine the appropriateness of the extended time frame.

The improvement plan is due to HSAG no later than 30 days following the organization's receipt of the final 2020 MHP Analysis report. The improvement plan should be uploaded electronically to OHA's deliverables reporting email address: cco.mcoopeliverableReports@dhsoha.state.or.us. HSAG will review the improvement plan using the following criteria to evaluate the sufficiency of each corrective action/intervention identified in the improvement plan to bring performance into compliance:

- Completeness of the improvement plan document in addressing each finding and identifying a responsible individual, a timeline/completion date, and specific corrective actions/interventions that the organization will take.
- Degree to which the planned corrective actions/interventions are anticipated to bring the organization into compliance with MHP requirements.
- Appropriateness of the timeline for the corrective actions/interventions given the nature of the finding.

Once reviewed, HSAG will communicate to the organization whether the improvement plan is approved. If any corrective actions/interventions are determined to not meet the requirements related to correlating findings, HSAG will identify the discrepancies and require resubmission of the improvement plan until it is approved by HSAG. Quarterly reviews of improvement plan progress will be conducted with each CCO via desk reviews and conference calls as necessary to ensure that all planned activities and interventions are completed.

HSAG will be available for technical assistance related to corrective actions/interventions. The CCO may contact either of the following HSAG representatives for assistance:

Melissa Isavoran, Associate Executive Director misavoran@hsag.com 503.839.9070 Barb McConnell, Executive Director bmcconnell@hsag.com 303.717.2105



Appendix A. MHP Evaluation Questionnaire

IHN submitted its completed MHP Evaluation Questionnaire, which identified changes or additions to benefits design and operations that may impact MHP corresponding with the six NQTL categories. The questionnaire served as a guide for OHA and the CCOs in that responses were used to identify and further document such changes and additions in the finalized MHP NQTL Reporting Tables located in Appendix B of this report.

General Questions for CCOs				
Ques	tion	Yes/No		
1.	Did the CCO add, change, or eliminate delegated administrative functions to a new or for an existing subcontractor (e.g., UM, provider admission, etc.)? Documentation Required: Provide contractual requirements (e.g., scope of work) for delegated administrative functions.	☐ Yes ⊠ No		
2.	Did the CCO add or exclude any specific classifications of drugs from its formulary?	☐ Yes ⊠ No		
Utiliz	ation Management (IP, OP, and Rx) Changes in CCO—MH Parity Analysis Sections I, II, and III			
Ques	tion	Yes/No		
1.	Did the CCO change payment arrangements with some/all providers (e.g., FFS to sub-capitation, per diem to DRG, reduction in payment levels to specific provider types or for specific benefits)?	☐ Yes ⊠ No		
2.	Did the CCO add or remove numerical limits (e.g., number of units) to MH/SUD or M/S benefits? Urine drug screens require prior approval after 12 units, per year.	⊠ Yes □ No		
3.	Did the CCO add or remove non-numerical benefit limits (e.g., scope or duration of benefits, medical necessity criteria, etc.) to MH/SUD or M/S benefits?	☐ Yes ⊠ No		
4.	Did the CCO change timelines for authorization requests for MH/SUD or M/S benefits?	☐ Yes ⊠ No		



6. Did the CCO change qualifications for reviewers that can authorize or deny requests? 7. Did the CCO develop or add medical necessity/level of care criteria for MH/SUD or M/S benefits? 8. Did the CCO change the method for monitoring consistency of MNC application for MH/SUD or M/S benefits (e.g., standards for consistency of MNC, reliability adherence criteria)? 9. Did the CCO change/modify penalties for failure to request/receive authorization for MH/SUD or M/S benefits (e.g., payment reductions, exceptions or waivers of penalties)? Retro authorization policy went into effect May 2018. 10. Did the CCO change frequency, time frames, or conditions of utilization review for MH/SUD or M/S benefits (e.g., RR or CR time frames or conditions)? 11. What is the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns experienced during the last full calendar year separately for MH/SUD and M/S for each classification (IP, OP, and Rx)? MH drugs are a carve out and covered by FFS OHP. See report submission and documentation. Documentation Required: Provide lists that identify the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e., IP, OP, and Rx). For Rx, include a list identifying the number of drugs subject to PA.	No Yes No Yes No Yes No Yes No				
 7. Did the CCO develop or add medical necessity/level of care criteria for MH/SUD or M/S benefits? S. Did the CCO change the method for monitoring consistency of MNC application for MH/SUD or M/S benefits (e.g., standards for consistency of MNC, reliability adherence criteria)? 9. Did the CCO change/modify penalties for failure to request/receive authorization for MH/SUD or M/S benefits (e.g., payment reductions, exceptions or waivers of penalties)? Retro authorization policy went into effect May 2018. 10. Did the CCO change frequency, time frames, or conditions of utilization review for MH/SUD or M/S benefits (e.g., RR or CR time frames or conditions)? 11. What is the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns experienced during the last full calendar year separately for MH/SUD and M/S for each classification (IP, OP, and Rx)? MH drugs are a carve out and covered by FFS OHP. See report submission and documentation. Documentation Required: Provide lists that identify the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e., hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e., 	Yes No Yes No Yes No Yes				
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carve out and covered by FFS OHP. See report submission and documentation. Documentation Required: Provide lists that identify the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e.,	Yes				
See report submission and documentation. *Documentation Required: Provide lists that identify the number of coverage requests, denials, appeals, appeal overturns, hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e.,	No				
hearings, and hearing overturns for the last full calendar year separately for MH/SUD and M/S for each classification (i.e.,					
Provider Network Admission Changes in CCO—MH Parity Analysis Sections IV and V					
Question	es/No				
1. Did the CCO change its network status from open (accepting new provider applications) to closed (not accepting new					
provider applications for certain provider types) or from closed to open?	Yes				
2. Did the CCO add, remove, or change provider admission requirements (e.g., special training, education, experience),					
including as a result of State licensing changes, for any MH/SUD or M/S providers?	No				



3.	Were any of the CCO's providers denied credentialing due to network closure (if applicable) or based on credentialing requirements? See report submission and documentation. Documentation Required: Provide a list of the number and percentage of providers denied credentialing (relative to those seeking credentialing, including the number of applications not accepted) or terminated from credentialing and provide the credentialing determination.	⊠ Yes □ No
4.	Did the CCO add or remove any MH/SUD or M/S provider types that are eligible for credentialing/reimbursement for services?	☐ Yes ⊠ No
Out-c	of-Network/Out-of-State Limit Changes in CCO—MH Parity Analysis Section VI	
Ques		Yes/No
Ques		Yes/No ☐ Yes ☐ No



Appendix B. Finalized MHP NQTL Reporting Tables

IHN submitted a completed MHP Reporting Template, which identified changes or additions to NQTLs that may impact MHP. HSAG synthesized the changes and additions to NQTLs with those reported in the CCO's 2018 MHP Analysis. Below are the finalized MHP NQTLs reported and assessed for the 2020 MHP Analysis by each of the six NQTL categories across MH/SUD and M/S benefits. Each NQTL was addressed based on comparability and stringency standards.

Category I—Utilization Management Limits Applied to Inpatient Services

NQTL: UM limits including PA, CR, RR, and IRR

Benefit Package: CCOA, CCOB, CCOE, and CCOG for adults and children

Classification: IP and emergency care

Overview: MH/SUD and M/S IP benefits require notification for emergency admissions. PA is not required for emergency care, but is applied to most other IP benefits including residential treatment. PA and CR are applied to IP benefits to confirm coverage, assure services are medically necessary and delivered in the least restrictive environment, and reduce overutilization of these high-cost services. These rationalizations were identified as indicators 1, 2 and 4 as listed in comparability and stringency Standard #2 below, which cross-reference to indicators used by OHP FFS. HSAG analyzed NQTLs applied to IP benefits based on information provided related to all six comparability and stringency standards as listed below. The benefit packages were analyzed as follows:

- **Benefit packages A and B:** MH/SUD benefits in columns 1 (CCO MH/SUD) and 2 (FFS MH/SUD) compared using indicators 1–4 to M/S benefits in column 3 (CCO M/S). These benefit packages include MH/SUD IP benefits managed by the CCO and OHA through its subcontractors, Comagine Health and Keystone Peer Review Organization (KEPRO), as compared to M/S IP benefits in column 3 managed by the CCO.
- **Benefit packages E and G:** MH/SUD benefits in columns 1 (CCO MH/SUD) and 2 (FFS MH/SUD) compared using indicators 1, 2, and 4 to M/S benefits in column 4 (FFS M/S). These benefit packages include MH/SUD IP benefits managed by the CCO and OHA through Comagine Health and KEPRO, as compared to M/S IP benefits in column 4 managed by OHA.



CCO MH/SUD	FFS MH/SUD	ссо м/ѕ	FFS M/S
1. To which benefit is the	TL assigned?		
 (1, 2, 3, 4) PA and CR arrequired for planned nonemergency admissions to acute IP (in and out-of-network (OON)), PRTS, subacute and 10 days after IP SUD detoxification admission. (1, 2, 3, 4, 6) Emergency admissions require notification within one business day of admission and subsequent CR. In practice do not penalize unless pattern of noncompliance. (1, 4) Extra-contractual and experimental/investigation unproven benefit request (i.e., exceptions) are submitted through a PA-I process. 	and extra-contractual benefits are conducted by OHA consistent with the information in column 2). • (2, 4) A level-of-care review is required for SCIP, SAIP and subacute care that is conducted by an OHA designee. • (1, 4) PA for SCIP, SAIP and subacute admission is obtained through a peer-to-peer review between a Comagine psychiatrist and	 (1, 2, 3, 4) PA and CR are required for planned nonemergency admissions to IP hospital, (in and OON) and IP hospice/palliative care (excludes routine maternity, which are 7% of admissions). (1, 2, 3, 4, 6) Emergency admissions require notification within one business day of admission and subsequent CR. (1, 2, 3, 4) Skilled nursing facility benefits (after 7 days and for the first 20 days) require PA. (1, 4) Extra-contractual and experimental/investigational/unproven benefit requests (i.e., exceptions) are submitted through a PA-like process. 	 (1, 2, 4) PA and CR are required for in-state and OOS planned surgical procedures (including transplants) and associated imaging, rehabilitation and professional surgical services delivered in an inpatient setting and listed in OAR 410-130-0200, Table 130-0200-1; rehabilitation, and long term acute care (LTAC).(Notification is required for all IP admissions.) (1, 2, 4) PA, CR and RR for Behavior Rehabilitation Services (BRS) are performed by OHA, DHS or OYA designee. (1, 2, 4) PA and CR of skilled nursing facility (SNF) services. (1, 4) Requests for extracontractual and experimental/investigational /unproven benefits (i.e., exceptions) are submitted through a PA-like process.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	process, and CR, is conducted by Comagine for PRTS. • (1, 2, 4) PA, CR for AFH, SRTF, SRTH, YAP, RTF, and RTH are performed by Comagine.		
2. Why is the NQTL assigned to	these benefits?		
 (1) To ensure coverage, medical necessity and prevent unnecessary overutilization (e.g., in violation of relevant OARs and associated Health Evidence Review Commission (HERC) guidelines1). (2) Ensure appropriate treatment in the least restrictive environment that maintains the safety of the individual. (3) Maximize use of INN providers to promote costeffectiveness when appropriate. (4) To comply with federal and State requirements. (6) To preserve scarce resources. 	 (1) UM is assigned to ensure medical necessity of services and prevent overutilization (e.g., matching the level of need to the least restrictive setting using the LOCUS – Level-of-Care utilization system and LSI – Level of Service Inventory or PCSP – Person Centered Service Plan and IBL – Individually-Based Limitations) of these high cost services. (2) Ensure appropriate treatment in the least restrictive environment that maintains the safety of the individual (e.g., matching the level of need to the least restrictive setting using the LOCUS – Level-of-Care Utilization System and LSI – 	 (1) To ensure coverage, medical necessity and prevent unnecessary overutilization (e.g., in violation of relevant OARs and associated Health Evidence Review Commission (HERC) guidelines). (2) Ensure appropriate treatment in the least restrictive environment that maintains the safety of the individual. (3) Maximize use of INN providers to promote costeffectiveness when appropriate. (4) To comply with federal and State requirements. (6) To preserve scarce resources. 	 (1) PA and CR are assigned to ensure medical necessity of services and prevent overutilization (e.g., requests for care that are not medically necessary or in violation of relevant OARs, the Health Evidence Review Commission (HERC) PL and guidelines). (2) Ensure appropriate treatment in the least restrictive environment that maintains the safety of the individual. (4) To comply with federal and State requirements.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
3. What evidence supports the r	Level of Service Inventory or PCSP – Person Centered Service Plan and IBL – Individually-Based Limitations). • (4) To comply with federal and State requirements. rationale for the assignment?		
 (1, 2 and 4) ASAM, HERC PL, and guidelines and MCG. (1) UM and claims reports are reviewed for trends in overutilization as compared to prior years' utilization on a quarterly basis (1) Annual cost and utilization reports that confirm IP as a cost driver based on percentage of spend. (1) Because of the frequent absence of physical indicators of medical necessity for these services (e.g., lab tests) with higher reliance on self-report measures that increases the potential for abuse and overutilization. 	(1, 2, and 4) Health Evidence Review Commission (HERC) Prioritized List (PL) and guidelines. The HERC include 13 appointed members which include five physicians, a dentist, a public health nurse, a pharmacist and an insurance industry representative, a provider of complementary and alternative medicine, a behavioral health representative and two consumer representatives. The Commission is charged with maintaining a prioritized list of services, developing or identifying evidence-based health care guidelines and	 (1, 2 and 4) MCG, HERC PL and guidelines. (1) UM and claims reports are reviewed for trends in overutilization on a quarterly basis (1) Annual cost and utilization reports that confirm IP as a cost driver based on percentage of spend. (1) Because of the frequent absence of physical indicators of medical necessity for these services (e.g., lab tests) with higher reliance on self-report measures that increases the potential for abuse and overutilization. (2) Medical errors in the 	 (1, 2 and 4) The HERC PL and guidelines. There are more guidelines for M/S than for MH/SUD because 1) there are more technological procedures (e.g., surgery, devices, procedures and diagnostic tests); and 2) the literature is more robust. (1) InterQual. (1) PA staff reports. If the UM team identifies any services for which utilization appears to be increasing (e.g., number of requests) or it appears that the State is paying for medically unnecessary care, the UM team consults with the health analytics team to

effectiveness research. HERC

provides outcome evidence

conducting comparative

cause of death in the US.

Makary, M. & Daniel, M.

hospital is the third leading

(2) Oregon Performance Plan

services be provided in least

(OPP) requires that BH

analyze and evaluate

adjustments to PA or CR.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
restrictive setting possible. The OPP is a DOJ-negotiated Olmsted settlement. Also see Roberts, E., Cumming, J & Nelson, K., A Review of Economic Evaluations of Community Mental Health Care, Sage Journals, Oct. 1, 2005, 1-13. Accessed May 25, 2018. http://journals.sagepub.com/doi/10.1177/1077558705279307 • (2) Inherent restrictiveness of residential settings and dangers associated with seclusion and restraint. Also see Cusack, K.J., Frueh, C., Hiers, T., et. al., Trauma within the Psychiatric Setting: A Preliminary Empirical Report, Human Services Press, Inc., 2003. 453-460. • (3) Network providers' credentials have been verified and they have contracted to accept the network rate. • (4) Applicable federal and State requirements.	and clinical guidelines for certain diagnoses that may be translated into UM requirements. There are fewer guidelines for MH/SUD than for M/S. This is because 1) there are fewer technological procedures for MH/SUD (e.g., cognitive behavioral therapy and psychodynamic therapy are billed using the same codes, no surgeries, few devices); 2) the MH/SUD literature is not as robust (e.g., fewer randomized trials, more subjective diagnoses (or the ICD-10-CM diagnoses represent a spectrum) and less standardization in interventions). • (1) InterQual.	 Medical Error - The Third Leading Cause of Death in the US, BMJ, 2016;353:i2139. (3) Network providers' credentials have been verified and they have contracted to accept the network rate. (4) Applicable federal and State requirements. (6) Michael Morris, Emergency Department Boarding of Psychiatric Patients in Oregon: Report Briefing, OHA Public Health Division, OHA 0730 (12/16), February 1, 2017 pp 1-16. 	(1) Health analytics reports. The health analytics team and policy analysts refer services that have been identified to have increasing utilization to the UM team for evaluation.



CCO MH/SUD	FFS MH/SUD	ссо м/ѕ	FFS M/S
• (6) Michael Morris, Emergency Department Boarding of Psychiatric Patients in Oregon: Report Briefing, OHA Public Health Division, OHA 0730 (12/16), February 1, 2017 pp 1-16.			
4. What are the NQTL procedure			
• Standard requests are approved within 14 days. Expedited requests within 72 hours.	 Timelines for gender reassignment surgery authorizations: (OHA) Standard requests are to be processed within 14 days. Timelines for child residential authorizations: (OHA) OHA provides the initial authorization (level-of-care review) within three days of receiving complete requests for SCIP, SAIP or subacute. (Comagine) Authorization requests for PRTS are submitted prior to admission or within 14 days of an emergency admission. An emergency admission is acceptable only under unusual and extreme 	Timelines for authorizations: Standard requests are approved within 14 days. Expedited requests within 72 hours.	 Timelines for authorizations: All in-state and out-of-state (OOS) emergency admissions, LTAC, and IP rehabilitation require notification. Notification is preferred within 24 hours of admission, but there is no timeline requirement. Notification allows the State to conduct case management and discharge planning, but does not limit the scope or duration of the benefit. PA is required before admission. OARs require emergency requests be processed within one business day, urgent requests within three business days and standard requests within 14 days.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	circumstances, subject to RR by Comagine. Timelines for adult residential and YAP authorizations: (Comagine Health)		
	• Emergency requests are processed within one business day, urgent within two business days, and standard requests within 10 business days.		
 Documentation requirements: May approve based on review of documentation received from individual completing MH assessment. Provider must provide the diagnostic and CPT codes and a rationale that demonstrates medical necessity for the procedure. 	Documentation requirements (OHA): • PA documentation requirements for non-residential MH/SUD benefits include a form that consists of a cover page. Diagnostic and CPT code information and a rationale for medical necessity must be provided, plus any additional supporting documentation. • The documentation requirement for level-of-care assessment for SCIP, SAIP	Provider must provide the diagnostic and CPT codes and a rationale that demonstrates medical necessity for the procedure.	PA documentation requirements include a form that consists of a cover page. Diagnostic and CPT code information and a rationale for medical necessity must be provided, plus any additional supporting documentation.
	and subacute is a psychiatric evaluation. Other information may be reviewed when available.		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
CCO MH/SUD	Documentation requirements for PRTF CONS and CR for PRTF, SCIP and SAIP (Comagine): • PRTS CONS requires documentation that supports the justification for child residential services, including: - A cover sheet detailing relevant provider and recipient - Medicaid numbers; - Requested dates of service; - HCPCS or CPT Procedure code requested; and	CCO M/S	FFS M/S
	requested; and - Amount of service or units requested; - A behavioral health assessment and service plan meeting the requirements described in OAR 309-019-0135 through 0140; or - Any additional supporting clinical information supporting medical justification for the services requested;		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	- For substance use disorder services (SUD), the Division uses the American Society of Addiction Medicine (ASAM) Patient Placement Criteria second edition-revised (PPC-2R) to determine the appropriate level of SUD treatment of care.		
	There are no specific documentation requirements for CR of PRTS, SCIP or SAIP.		
	Documentation requirements (Comagine Health):		
	Documentation may include assessment, service plan, plan-of-care, Level-of-Care Utilization System (LOCUS), Level of Service Inventory (LSI), PCSP, IBL, or other relevant documentation.		
Method of document submission:	Method of document submission (OHA):	Method of document submission:	Method of document submission:
Information may be submitted via fax, phone, electronic submission	For non-residential MH/SUD services, paper (fax) or online PA requests are submitted prior to the delivery of	Information may be submitted via fax, phone, electronic submission	Paper (fax) or online PA requests are submitted prior to the delivery of services for which PA is required.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
through the provider portal or by secure email.	services for which PA is required. • For SCIP, SAIP and subacute level-of-care review, the OHA designee may accept information via fax, mail or secure email and has also picked up information. Supplemental information may be obtained by phone. Method of document	through the provider portal or by secure email.	
	 submission (Comagine): Packets are submitted to Comagine by mail, fax, email or web portal for review for child residential services. Telephonic clarification may be obtained. Psychiatrist to psychiatrist review is telephonic. Method of document submission (Comagine Health): 		
	Providers submit authorization requests for adult MH residential to Comagine Health by mail, fax, email or via portal, but documentation must still be faxed if the request is through		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	portal. Telephonic clarification may be obtained.		
 Qualifications of reviewers: Licensed (e.g., MD, LCW, PhD) clinical reviewers can approve authorization requests relative to MNC (e.g., MCG, ASAM, HERC PL and associated guideline notes, OARs, Plan benefit coverage, federal rules). Only physicians can deny authorization requests. 	 Qualifications of reviewers (OHA): OHA M/S staff conduct PA and CR (if applicable) for gender reassignment surgery. The OHA designee is a licensed, master's-prepared therapist that reviews psychiatric evaluations to approve or deny the level-of-care requested. Psychiatric consultation is available if needed. Qualifications of reviewers (Comagine): Two reviewers with QMHP designation make residential authorization decisions. Two psychiatrists make CONS determinations. Qualifications of reviewers (Comagine Health): Comagine Health QMHPs must meet minimum qualifications (see below) and demonstrate the ability to conduct and review an assessment, including 	 Qualifications of reviewers: Nurses may authorize services relative to MNC (e.g., MCG, HERC PL and associated guideline notes, OARs, Plan benefit coverage, federal rules). Only Medical Directors may deny based on lack of medical necessity. 	• Nurses may authorize and deny authorization requests relative to OAR, HERC PL guidelines and associated notes, and other industry guidelines (e.g., AIM for radiology).



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
CCO MIH/SUD	identifying precipitating events, gathering histories of mental and physical health, substance use, past mental health services and criminal justice contacts, assessing family, cultural, social and work relationships, and conducting/reviewing a mental status examination, complete a DSM diagnosis, and write and supervise the implementation of a PCSP. • A QMHP must meet one of the follow conditions: — Bachelor's degree in nursing and licensed by the State of Oregon; — Bachelor's degree in occupational therapy and licensed by the State of Oregon; — Graduate degree in psychology; — Graduate degree in	CCO IM/S	FFS IM/S
	social work; – Graduate degree in		
	recreational, art, or music therapy;		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
Criteria: MCG, ASAM, HERC PL and associated guideline notes, OARs, Plan benefit coverage, federal rules. Recently moved to an automated system based on MCG, so frequency and style of review has changed.	 Graduate degree in a behavioral science field; or A qualified Mental Health Intern, as defined in 309-019-0105(61). Criteria (OHA): Authorizations for nonresidential MH/SUD services are based on the HERC PL and guidelines; Oregon Statute, OAR, and federal regulations; InterQual guidelines; and evidence-based guidelines from private and professional associations. OHA delegates review requests relative to least restrictive environment requirement. Criteria (Comagine): HERC PL, InterQual, and Comagine policy are used for residential CR. Criteria (Comagine Health): 	Criteria: • MCG, HERC PL and associated guideline notes, OARs, Plan benefit coverage, federal rules	Criteria: • Authorizations are based on the HERC PL and guidelines; Oregon Statute, OAR, and federal regulations; InterQual guidelines; and evidence-based guidelines from private and professional associations, such as the Society of American Gastrointestinal and Endoscopic Surgeons and InterQual, where no State or federal guidelines exist.
	QMHPs review information submitted by providers relative to State plan and OAR requirements and develop a PCSP.		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	The PCSP components are entered into MMIS as an authorization.		
Reconsideration/RR:	Retrospective Review:	Reconsideration/RR:	Retrospective Review:
RR is offered (with some limitations) for providers who fail to PA medically necessary care; 90-day timeframe.	 Retrospective authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. Reconsideration (OHA): A provider may request review of an OHA denial decision for nonresidential MH/SUD services. The review occurs in weekly Medical Management Committee (MMC) meetings. Exception requests for experimental and other noncovered benefits may be granted at the discretion of the MMC, which is led by the HSD medical director. If a provider requests review of an OHA delegate level-of-care determination, KEPRO 	RR is offered (with some limitations) for providers who fail to PA medically necessary care; 90-day timeframe.	 Retrospective authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. Reconsideration: A provider may request review of a denial decision. The review occurs in weekly MMC meetings. Exception requests for experimental and other non-covered benefits may be granted at the discretion of the MMC, which is led by the OHA's medical director.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	may conduct the second review. Reconsideration (Comagine):		
	If the facility requests a reconsideration of a CONS denial, a second psychiatrist (who did not make the initial decision) will review the documentation and discuss with the facility in a formal meeting.		
	• No policy for CR denials.		
	Reconsideration (Comagine Health):		
	• Within 10 days of a denial, the provider may send additional documentation to Comagine Health for reconsideration.		
	A provider may request review of a denial decision, which occurs in weekly MMC meetings or Comagine Health's own comparable medical management meeting.		
Appeals:	Appeals (OHA):	Appeals:	Appeals:
Members or providers with consent, may appeal any denial decision within the	 Members may request a hearing on any denial decision. 	Members or providers with consent, may appeal any denial decision within the	Standard appeal and fair hearing rights apply.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
guidelines set forth by rules and regulations.	 Appeals (Comagine): Members may request a hearing on any denial decision. Appeals (Comagine Health): Members may request a hearing on any denial decision. 	guidelines set forth by rules and regulations.	
Consequences for failure to authorize:	Consequences for failure to authorize (OHA):	Consequences for failure to authorize:	Consequences for failure to authorize:
 Failure to obtain authorization can result in non-payment. Exceptions may be made to the PA process at the discretion of the reviewing medical doctor. Benefit coverage is limited to medically necessary services by contract. 	 Failure to obtain authorization for non-residential MH/SUD services can result in non-payment for benefits for which it is required. Failure to obtain notification for non-residential MH/SUD services does not result in a financial penalty. For SCIP, SAIP and subacute, if coverage is retroactively denied, general funds will be used to cover the cost of care. Consequences for failure to authorize (Comagine): Non-coverage. Consequences for failure to authorize (Comagine Health): 	 Failure to obtain authorization can result in non-payment. Exceptions may be made to the PA process at the discretion of the reviewing medical doctor. Benefit coverage is limited to medically necessary services by contract. 	 Failure to obtain authorization can result in non-payment for benefits for which it is required. Failure to obtain notification does not result in a financial penalty.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
5. How frequently or strictly is t	Failure to obtain authorization can result in non-payment for benefits for which it is required. he NQTL applied?		
Frequency of review (and method of payment):	Frequency of review (and method of payment) (OHA):	Frequency of review (and method of payment):	Frequency of review (and method of payment):
 Current hospital contracts are DRG-based CR occurs every 7 days for acute IP and subacute consistent with MCG. The reviewer may attend an individual's treatment review every 3 months, but goes to the hospital daily. Reauthorizations for PRTS and SUD residential occur (on average) every 30 days depending upon the facilities' recommendation, MCG and member needs CR and RR are conducted by chart review via EHR, onsite or through chart submission. 	 Gender reassignment surgery is authorized as a procedure. The initial authorization for SCIP, SAIP, and subacute is 30 days. Frequency of review (and method of payment) (Comagine): Child residential services are paid by per diem. Child residential services authorizations are conducted every 30-90 days. Frequency of review (and method of payment) (Comagine Health): Adult residential authorizations are conducted at least once per year. An independent and qualified agent (IQA) contacts MH 	 Current hospital contracts are DRG-based. CR is done every 7 days following approved admission (even if DRG) to IP hospital or SNF or more frequently if deemed necessary based on the individual's circumstances to promote coordination of care. CR and RR are conducted by chart review via EHR or through chart submission. Care coordination and outreach continue after initial concurrent review with SNF care manager. 	 Most IP claims are paid DRG; as a result, CR is infrequently used. CR is conducted monthly for LTAC and rehabilitation. The State conducts CR for SNF at a frequency that is determined by the care manager, but not less than one time a year. Authorization lengths are individualized by condition and are valid for up to a year. Procedural authorizations are valid for three months.

provider quarterly for 1915i



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	assessment accuracy. If member's status changes for more than 30 days, provider can contact IQA for a reassessment.		
RR conditions and timelines:	Retrospective Review:	RR conditions and timelines:	Retrospective Review:
• Providers are allowed to make requests if auths are required, but an authorization request was not submitted at date of service. Provider can submit a retro auth within the 90 day time period.	• Retrospective authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days.	• Providers are allowed to make requests if auths are required, but an authorization request was not submitted at date of service. Provider can submit a retro auth within the 90 day time period.	• Retrospective authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days.
	Reconsideration (OHA):		Reconsideration:
	 A provider may request review of an OHA denial decision for nonresidential MH/SUD services. The review occurs in weekly Medical Management Committee (MMC) meetings. Exception requests for experimental and other noncovered benefits may be granted at the discretion of the MMC, which is led by the HSD medical director. 		 A provider may request review of a denial decision. The review occurs in weekly MMC meetings. Exception requests for experimental and other non-covered benefits may be granted at the discretion of the MMC, which is led by the OHA's medical director.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	If a provider requests review of an OHA delegate level-of-care determination, KEPRO may conduct the second review.		
	Reconsideration (Comagine):		
	If the facility requests a reconsideration of a CONS denial, a second psychiatrist (who did not make the initial decision) will review the documentation and discuss with the facility in a formal meeting.		
	 No policy for CR denials. 		
	Reconsideration (Comagine Health):		
	Within 10 days of a denial, the provider may send additional documentation to Comagine Health for reconsideration.		
	A provider may request review of a denial decision, which occurs in weekly MMC meetings or Comagine Health's own comparable medical management meeting.		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
Methods to promote consistent application of criteria:	Methods to promote consistent application of criteria (OHA):	Methods to promote consistent application of criteria:	Methods to promote consistent application of criteria:
IRR conducted annually; 85% testing standard.	Nurses are trained on the application of the HERC PL and guidelines, which is spotchecked through ongoing supervision. Whenever possible, practice guidelines from clinical professional organizations such as the American Medical Association or the American Psychiatric Association, are used to establish PA frequency for non-residential MH/SUD services.	IRR conducted annually; 85% testing standard.	Nurses are trained on the application of the HERC PL and guidelines, which is spotchecked through ongoing supervision. Whenever possible, practice guidelines from clinical professional organizations such as the American Medical Association or the American Psychiatric Association, are used to establish PA frequency for services in the FFS system.
	There are only two OHA designee reviewers for level-of-care review for SCIP, SAIP, and subacute and no specific criteria, so N/A.		
	Methods to promote consistent application of criteria (Comagine):		
	Parallel chart reviews for the two reviewers. (No criteria.)		
	Methods to promote consistent		
	application of criteria (Comagine Health):		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	 Monthly clinical team meetings in which randomly audited charts are reviewed/discussed by peers using Comagine Health compliance department-approved audit tool. Results of the audit are compared, shared and discussed by the team and submitted to Compliance Department monthly for review and documentation. Individual feedback is provided to each clinician during supervision on their 		
	authorization as well as plan- of-care reviews.		
6. What standard supports the	frequency or rigor with which the I	NOTL is applied?	
Evidence for UM frequency:	Evidence for UM frequency	Evidence for UM frequency:	Evidence for UM frequency:
 MCG, ASAM, OARs, HERC, federal and State requirements, practice guidelines Oregon CCO contract. 42 CFR Part 441, Subpart D requiring PRTF is reviewed at least every 30 days 	 (OHA (and designee for level-of-care review), Comagine and KEPRO): PA length and CR frequency are tied to HERC PL and guidelines, OAR, CFRs, InterQual, reviewer expertise and timelines for expectations of improvement. 	Evidence to support the standard comes from HERC, MCG, and in its absence, evidence-based criteria, research, and policies developed by the Medical Director, OARs	PA length and CR frequency are tied to HERC PL and guidelines, DRGs, OAR, CFRs, InterQual, reviewer expertise and timelines for expectations of improvement.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
(§441.155) for child providers.			
 UM application: Utilization review reports. Denial/overturn rates. Approval rates. UM application: Denial/appeal overturn rates; number of PA requests; stabilization of cost trends; and number of hearings 		 Data reviewed to determine UM application: Denial/appeal overturn rates. number of PA and concurrent review requests. Stabilization of cost trends. 	Data reviewed to determine UM application: • A physician led group of clinical professionals conducts an annual review to determine which services receive or retain PA. Items reviewed include: - Utilization.
	by the State. (Applicable to non-residential MH/SUD services.) Data reviewed to determine UM application (Comagine): N/A Data reviewed to determine		 Approval/denial rates. Documentation/ justification of services. Cost data.
	UM application (Comagine Health): • N/A		
IRR standard:	IRR standard (OHA):	IRR standard:	IRR standard (OHA):
IRR conducted annually; 85% testing standard.	KEPRO has a formal policy including an 80% standard using InterQual criteria, conducting reviews at least annually.	IRR conducted annually; 85% testing standard.	KEPRO has a formal policy including an 80% standard using InterQual criteria, conducting reviews at least annually.
	IRR standard (Comagine Health):		



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	 Spot-checks performed through supervision. Formal policy to be developed. 		
	IRR standard (Comagine Health):		
	Spot-checks performed through supervision.		

IHN was responsible for delivering IP MH/SUD and M/S Medicaid benefits to members in all four benefit packages (CCOA, CCOB, CCOE, and CCOG), whereas OHP FFS was fully managing IP M/S benefits for CCOE and CCOG benefit packages. Emergency MH/SUD and M/S IP hospital admissions required notification, with most ongoing IP services requiring subsequent CR. Regarding nonemergent CCO MH/SUD and M/S IP admissions, PA or level-of-care approval was required. PA was also required for extra-contractual coverage requests (including experimental services); planned surgical procedures (including transplants); and associated imaging, rehabilitation, and professional surgical services delivered in an IP setting and listed in OAR 410-130-0200, Table 130-0200-1. For psychiatric residential treatment services (PRTS) benefits (e.g., Secure Children's Inpatient Treatment Programs [SCIP], Secure Adolescent Inpatient Treatment Programs [SAIP], and adult and youth residential services) delivered under all benefit packages, OHP FFS's subcontractor, Comagine Health, was conducting the certificate of need (CON) and PA processes, with the CCO conducting CR for those services. The CCO was also conducting CR for MH/SUD subacute benefits. For M/S benefits under CCOA and CCOB benefit packages, the CCO was conducting PA and CR for SNF benefits for the first 20 days, with subsequent management being conducted by OHP FFS.

HSAG's analysis of IHN's PA data for IP and OP benefits did not reveal any concerns related to MHP. Of the total 6,411 IP and OP PA requests reported, 10.23 percent were denied. Of the 63 MH/SUD PA requests denied, representing 5.52 percent of the 1,142 MH/SUD PA requests, only three denials resulted in an appeal. The 63 reported MH/SUD denials represented 9.60 percent of total denials. Only three of the MH/SUD denials were for IP benefits, all denied due to a "does not meet criteria" denial reason.

Comparability

UM was assigned to MH/SUD and M/S IP benefits primarily using four rationales: 1) To ensure coverage, medical necessity, and prevent unnecessary overutilization (e.g., in violation of relevant OARs, HERC PL and guidelines, or clinical practice guidelines or research); 2) To ensure appropriate treatment in the least restrictive environment that maintains the safety of the individual; 3) To maximize use of in-network (INN) providers to promote cost-effectiveness when appropriate; and 4) To comply with federal and State requirements. HSAG determined the rationale and evidence to be comparable.



CCO MH/SUD FFS MH/SUD CCO M/S FFS M/S

Emergency MH/SUD and M/S IP hospital admissions required notification within one business day, with child emergency residential admissions separately requiring notification within 14 days. Most CCO documentation requirements for MH/SUD included an admission note and records submitted via telephone, fax, or electronically. OARs required authorization decisions 72 hours for urgent requests and 14 days for standard requests. Both IHN and OHP FFS adhered to these requirements across the benefit packages. Providers were encouraged to submit requests for authorization sufficiently in advance to be consistent with the OAR time frames. Most ongoing IP services required subsequent CR. Documentation requirements for child residential PA/level-of-care review included a psychiatric evaluation or a psychiatrist-to-psychiatrist telephonic review. Comagine Health, OHP FFS's subcontractor, accepted information for child residential CR via mail, email, fax, and Web portal. Adult and youth residential required an assessment (i.e., completion of a relevant level-of-care tool [e.g., ASAM, LSI, or LOCUS]) and plan-of-care consistent with State plan requirements. Comagine Health documentation submission could be done using mail, email, fax, or Web portal. Consistent with OARs, federal CON procedures, and due to the potential absence of a psychiatric referral, the PRTS documentation requirements included a cover sheet, a behavioral health assessment, and service plan meeting the requirements described in OAR 309-019-0135 through 0140. HSAG determined the MH/SUD authorization time frames and documentation requirements were comparable to those applied to M/S authorization requests.

Stringency

Qualified individuals conducted UM applying OARs, HERC, MCG, national guidelines, and ASAM for CCO SUD. While the CCO and OHP FFS subcontractors required all MH/SUD and M/S denials to be made by professional peers, nurses could deny benefits managed by OHP FFS. HSAG determined this difference to be an issue of quality rather than parity. OHP FFS's subcontractor, Comagine Health, made denial determinations for level-of-care review for certain child residential services. Both the CCO and OHP FFS allowed 90-day RR for MH/SUD and M/S when providers failed to obtain authorization. Exceptions to these time frames were considered by both the CCO and OHP FFS. For adult and youth residential services, Comagine Health allowed reconsideration of denials with the submission of additional documentation within 10 days of the denial. For OHP FFS and Comagine Health, the review of denial decisions occurred during MMC meetings. MH/SUD and M/S denial decisions could be challenged through appeals and/or State fair hearing processes. Failure to obtain authorization could result in noncoverage, although SCIP, SAIP, and subacute services could be covered by general fund dollars. Regarding IRR, the CCO conducted annual reviews using an 85 percent testing goal, whereas OHP FFS's subcontractor had an 80 percent testing standard for M/S benefit authorizations. HSAG did not determine this discrepancy to be a parity concern as the method to promote consistency was more structured for MH/SUD benefits.

Outcome

HSAG determined IHN's processes, strategies, and evidentiary standards for UM of IP MH/SUD benefits to be comparable and no more stringently applied, in writing and in operation, to IP M/S benefits.



Category II—Utilization Management Limits Applied to Outpatient Services

NQTL: UM limits including PA, CR, RR, and IRR

Benefit Package: CCOA, CCOB, CCOE, and CCOG for adults and children

Classification: OP

Overview: UM is assigned to OP MH/SUD and M/S benefits to confirm coverage, meet federal requirements in providing benefits in the least restrictive environment, evaluate the safety of certain OP services, and prevent overutilization that has been identified by specific medical necessity criteria or in utilization reports. These rationalizations are identified as indicators 1, 2, and 3 as listed in comparability and stringency Standard #2 below, which cross-reference to indicators used by OHP FFS. HSAG analyzed NQTLs applied to OP benefits based on information provided related to all six comparability and stringency standards as listed below. The benefit packages were analyzed as follows:

- **Benefit packages A and B:** MH/SUD benefits in columns 1 (FFS/home- and community-based services [HCBS] 1915[c][i] MH /SUD) and column 3 (CCO MH/SUD) compared using indicators 1–4 to M/S benefits in columns 2 (FFS/HCBS 1915[c][k][j] M/S) and 4 (CCO M/S), respectively. These benefit packages include MH/SUD IP benefits managed by the CCO and OHA through its subcontractors, Comagine Health and KEPRO.
- Benefit packages E and G MH/SUD benefits in columns 1 (FFS/HCBS 1915[c][i] MH/SUD) and column 3 (CCO MH/SUD) compared using indicators 1, 2, and 4 to M/S benefits in columns 2 (FFS/HCBS 1915[c][k][j] M/S) and 5 (FFS M/S), respectively. These benefit packages include MH/SUD IP benefits managed by the CCO and OHP FFS through its subcontractors, Comagine Health and KEPRO.

	FFS HCBS MH/SUD	FFS HCBS M/S		CCO MH/SUD		CCO M/S	FFS M/S
1	. To which benefit is the	NQTL assigned?					
•	(2) Applied Behavior Analysis (ABA).	The following services are managed by DHS:	•	(2, 3) All non-contracted services.	•	(2, 3) All non-contracted services.	The following services are managed by OHA:
•	(2) OT, PT, ST for MH conditions are		•	(2, 4) ABA.	•	(2) Contact lenses.	• (2, 3) Out of hospital births.



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S		
managed through RR; PA is not required.	 (1) 1915(c) Comprehensive DD waiver. (1) 1915(c) Support Services DD waiver. (1) 1915(c) Behavioral DD Model waiver. (1) 1915(c) Aged & Physically Disabled waiver. (1) 1915(c) Hospital Model waiver. (1) 1915(c) Medically Involved Children's NF waiver. (1) 1915(k) Community First Choice State Plan option. (1) 1915(j): Self- directed personal assistance. 	• (2, 4) PA after 120 units/hours of PT/ST/OT (HERC 30 visits).	 (2) Durable medical equipment. (2, 3) Elective/planned procedures in hospital or ambulatory surgery center. (2) Potentially cosmetic services. (2, 3) Radiological services. (2, 3) Elective coronary angioplasty. (2, 3) Parenteral nutrition (2, 3) Acupuncture visits in excess of 30 visits per year. (2, 4) Transplants. 	 (2) Home health services. (2) OT, PT, ST for MH conditions are managed through RR; PA is not required. (2, 3) Imaging. (2) DME. 		
2. Why is the NQTL assigned to these benefits?						
 (2) HERC PL. (2) OAR 410-172- 0650 for ABA services. 	(1) The State requires PA of HCBS in order to meet federal requirements regarding PCSPs and	(2) To ensure services provided are in accordance with the applicable OARs and medical regulations,	(2) To ensure services provided are in accordance with the applicable OARs and medical regulations,	(2) To prevent services being delivered in violation of relevant OARs, associated HERC PL		



	FFS HCBS MH/SUD	FFS HCBS M/S		CCO MH/SUD		CCO M/S		FFS M/S
•	(2) PA requests with insufficient documentation to demonstrate MNC or HERC PL guidelines are not being followed.	ensure services are provided in accordance with a participant's PCSP and in the last restrictive setting.	•	the CCO contract, HERC guidelines and quality care (relative to MCG) and prevent unnecessary costs. (3) Services are associated with increased health or safety risks. (4) To preserve scarce resources	•	the CCO contract, HERC guidelines and quality care (relative to MCG) and prevent unnecessary costs. (3) Services are associated with increased health or safety risks. (4) To preserve scarce resources	•	and guidelines and federal regulations. (3) Services are associated with increased health or safety risks.
3.	What evidence suppor	ts the rationale for the assig	gnm	ent?				
•	(2) HERC PL (2) OAR 410-172- 0650 for ABA services.	• (1) Federal requirements regarding PCSPs for 1915(c), 1915(k), and	•	(2 and 3) ASAM, OARs, HERC PL and guidelines, and federal guidelines.	•	(2 and 3) OARs, HERC PL and guidelines, and federal guidelines.	•	(2) HERC PL and guidelines, and clinical practice guidelines.
•	(2) PA requests with insufficient documentation to demonstrate medical necessity is not being met or HERC PL guidelines are not being followed.	1915(j) services (e.g., 42 CFR 441.301, 441.468, and 441.540) and the applicable approved 1915(c) waiver application/State plan amendment. • (1) Federal requirements regarding 1915(c) and 1915(i) services require that HCBS are provided in the least	•	(2) UM and claims reports are reviewed for trends in overutilization on a quarterly basis. (2) Annual cost and utilization reports. (Institute of Medicine Report, (2012). (2) Contract. (2) Practice Guidelines for the Treatment of	•	 (2) UM and claims reports are reviewed for trends in overutilization on a quarterly basis. (2) Annual cost and utilization reports. (2) Contract. (3) HERC and MCG. (4) Difficulty obtaining timely referrals. 	•	(2) PA requests with insufficient documentation to demonstrate medical necessity are not being met or HERC PL guidelines are not being followed. (3) HERC Guidelines - Recommended limits on services for member safety.



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
	restrictive setting possible.	Psychiatric Disorders, Treatment of Patients with Eating Disorders, Third Edition, American Psychiatric Association Publishing, 2010; National Institute for Clinical Excellence, Eating Disorders, Clinical Guide 9, January 2004; American Academy of Family Physicians, Diagnosis of Eating Disorders in Primary Care, Table 6, Level- of -Care Criteria for patients with eating disorders, 2003. (3) HERC and MCG. (4) Difficulty obtaining timely referrals.		
4. What are the NQTL p		II. 1. 6	m· · · e	m· 1· e
Timelines for authorizations:	Timelines for authorizations:	Timelines for authorizations:	Timelines for authorizations:	Timelines for authorizations:
Urgent requests are processed in three business days and	 A PCSP must be approved within 90 days from the date a 	A provider is expected to request prior to the delivery of the service	A provider is expected to request prior to the delivery of the service	Urgent requests are processed in three business days and



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
 immediate requests in one business day. Routine requests are processed in 14 days. OT, PT, ST for MH conditions are managed through RR; PA is not required. 	completed application is submitted.	 and to wait for an authorization prior to administering the service. The provider may not attempt to collect payment from the beneficiary for a service that was not prior authorized. Routine requests are completed in 14 days; urgent requests in 72 	 and to wait for an authorization prior to administering the service. The provider may not attempt to collect payment from the beneficiary for a service that was not prior authorized. Routine requests are completed in 14 days; urgent requests in 72 	immediate requests in one business day. Routine requests are processed in 14 days. OT, PT, ST for MH conditions are managed through RR; PA is not required.
		hours.	hours.	
Documentation requirements:	Documentation requirements:	Documentation requirements:	Documentation requirements:	Documentation requirements:
 Form is one cover page. Require diagnostic and CPT code and rationale for medical necessity plus any additional supporting documentation. In addition, as part of the supporting documentation ABA must have an evaluation and referral for treatment from a 	The PCSP is based on a functional needs assessment and other supporting documentation. It is developed by the individual, the individual's team, and the individual's case manager.	Provider must provide the diagnostic and CPT code to be applied and a rationale that demonstrates medical necessity for the procedure.	Provider must provide the diagnostic and CPT code to be applied and a rationale that demonstrates medical necessity for the procedure.	 A cover page form is required. In addition, diagnostic information, a CPT code(s), a rationale for medical necessity plus any additional supporting documentation are required. Documentation supporting medical necessity is required at the time of billing



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
licensed practitioner described in OAR 410-172-0760 (1)(a-d) and a treatment plan from a licensed health care professional described in 410-172-0650(B). • Documentation supporting medical necessity is required at the time of billing for OT, PT, ST services.				for OT, PT, ST services.
Method of document submission:	Method of document submission:	Method of document submission:	Method of document submission:	Method of document submission:
Paper (fax) or online PA/POC submitted prior to the delivery of services.	 All 1915(c), 1915(k), and 1915(j) services must be included in a participant's PCSP and approved by a qualified case manager at the local case management entity (CME) prior to service delivery. Information is obtained during a face-to-face meeting, often at the individual's location. 	Information may be submitted via fax, phone, electronic submission through the provider portal or by secure email.	Information may be submitted via fax, phone, electronic submission through the provider portal or by secure email.	Paper (fax) or online PA/POC submitted prior to the delivery of services.



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
Qualifications of reviewers:	Qualifications of reviewers:	Qualifications of reviewers:	Qualifications of reviewers:	Qualifications of reviewers:
 For ABA services, physicians review services. For OT, PT, ST services, nurses may authorize and deny services. Professional peers deny for other OP services. 	A case manager must have at least: A bachelor's degree (BA) in behavioral science, social science, or a closely related field; or A BA in any field AND one year of human services related experience; or An associate's degree (AA) in a behavioral science, social science, or a closely related field AND two years human services related experience; or Three years of human services related experience.	 Licensed clinical reviewers can approve authorization requests relative to MNC. Denials are reviewed by the medical director. If special expertise is required, a third party reviewer may be consulted. 	Nurses authorize and physicians deny.	Nurses may authorize and deny services.



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
Criteria: • Authorizations are based on applicable HERC guidelines, Oregon Statute, Oregon Administrative rules, federal regulations, and evidence-based guidelines from private and professional associations such as the American Psychiatric Association, where no State or federal guidelines exist.	 Qualified case managers approve or deny services in the PCSP consistent with waiver/state plan and OAR requirements. Once a PCSP is approved, itis entered into the payment management system as authorization by the CME staff. 	Criteria: • MCG, Criteria Policy, State/federal law, contracts, and HERC PL with accompanying guideline notes.	Criteria: • MCG, Criteria Policy, State/federal law, contracts, and HERC PL with accompanying guideline notes.	Authorizations are based on applicable HERC PL and guidelines, Oregon Revised Statute, OAR, federal regulations, and evidence-based guidelines from private and professional associations such as the Society of American Gastrointestinal and Endoscopic Surgeons where no State or federal guidelines exist.
Reconsideration/RR: • A provider may request review of a denial decision, which occurs in weekly MMC meetings or Comagine's own comparable MMC meeting. • RR authorization requests can be made	Reconsideration/RR: • (c) NA	Reconsideration/RR RR is available.	Reconsideration/RR RR is available.	Reconsideration/RR: • A review of a denial decision can be requested and is reviewed in weekly MMC meetings. • RR authorization requests can be made within 90 days of the date of service or after the 90 days based on



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. OT, PT, ST are reviewed after the initial service prior to payment. Additionally, denial decisions can be requested and reviewed at weekly MMC meetings.				provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. OT, PT, ST are reviewed after the initial service prior to payment. Additionally, denial decisions can be requested and reviewed at weekly MCM meetings.
Consequences for failure to authorize:	Consequences for failure to authorize:	Consequences for failure to authorize:	Consequences for failure to authorize:	Consequences for failure to authorize:
Failure to obtain authorization may result in non-payment.	Failure to obtain authorization may result in non-payment.	Failure to obtain PA and absence of MNC results in non- payment.	Failure to obtain PA and absence of MNC results in non- payment.	Failure to obtain authorization may result in non-payment.
Appeals:	Appeals:	Appeals:	Appeals:	Appeals:
Notice and fair hearing rights apply.	Notice and fair hearing rights apply.	Members or providers with consent, may appeal any denial decision within the	Members or providers with consent, may appeal any denial decision within the	Notice and fair hearing rights apply.



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
5. How frequently or stri	ctly is the NQTL applied?	guidelines set forth by rules and regulations.	guidelines set forth by rules and regulations.	
 Frequency of review: PA is granted for different LOS depending on the service and can be adjusted. PAs for extensive services usually range from 6 months to 1 year. ABA is usually multiple service codes approved for 6 months. Exceptions may be made at the discretion of the MMC, which is led by the HSD medical director. 	Frequency of review: • PCSPs are reviewed and revised as needed, but at least every 12 months.	 Frequency of review: A second PA is required when the initial number of units is exhausted and additional service is desired. Office visits are approved based on the number requested by the provider. Individualized and tied to treatment plan and MCG expected improvement rate and stages. Average range of authorization is 3-6 months unless requested by provider, but for no longer than 12 months. 	 Frequency of review: A second PA is required when the initial number of units is exhausted and additional service is desired. Average range of authorization is 3-6 months unless requested by provider, but for no longer than 12 months. 	Frequency of review and method of payment: PA is authorized based on the length of time/number of services for which treatment should be completed. Minimal CR is conducted. DME average is every 3 months.
Reconsideration/RR: • A provider may request review of a denial decision, which	Reconsideration/RR: NA	RR conditions and timelines: No limits.	RR conditions and timelines: No limits.	Reconsideration/RR: • A review of a denial decision can be requested and is



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
occurs in weekly MMC meetings or Comagine's own comparable MMC meeting. RR authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. OT, PT, ST are reviewed after the initial service prior to payment. Additionally, denial decisions can be requested and reviewed at weekly MMC meetings.				reviewed in weekly MMC meetings. RR authorization requests can be made within 90 days of the date of service or after the 90 days based on provider demonstration of a specific reason why authorization could not have been obtained within the 90 days. OT, PT, ST are reviewed after the initial service prior to payment. Additionally, denial decisions can be requested and reviewed at weekly MCM meetings.
Methods to promote consistent application of criteria:	Methods to promote consistent application of criteria:	Method to promote consistent application of criteria:	Method to promote consistent application of criteria:	Methods to promote consistent application of criteria:
For ABA, a sample of cases are reviewed for	DHS Quality Assurance Review			Nurses are trained on the application of the



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
ability to address assessed member needs and whether OARs were met.	teams review a representative sample of PCSPs as part of quality assurance and case review activities to assure that PCSPs meet program standards. • Additionally, OHA staff review a percentage of files to assure quality and compliance.	 IRR testing standard of 85%. Exceptions may be made to the PA process at the discretion of the reviewing medical doctor. Benefit coverage is limited to medically necessary services by contract. 	 IRR testing standard of 85%. Exceptions may be made to the PA process at the discretion of the reviewing medical doctor. Benefit coverage is limited to medically necessary services by contract. 	HERC guidelines, which is spot checked through ongoing supervision.
	ts the frequency or rigor wi			
Evidence for UM frequency:	Evidence for UM frequency:	Evidence for UM frequency:	Evidence for UM frequency:	Evidence for UM frequency:
HERC guidelines (for ABA and OT, PT, ST) of which there are more M/S than MH/SUD because 1) there are more technological procedures (e.g., surgery, devices, procedures and diagnostic tests); 2) the literature is more robust.	• Federal requirements regarding PCSPs and 1915(c), 1915(k), and 1915(j) services (e.g., 42 CFR 441.301, 441.468, and 441.540) and the applicable approved 1915(c) waiver application/State plan amendment.	MCG, Criteria Policy, State/federal law, contracts, and HERC PL with accompanying Guidelines notes.	MCG, Criteria Policy, State/federal law, contracts, and HERC PL with accompanying Guidelines notes.	HERC guidelines of which there are more M/S than MH/SUD because 1) there are more technological procedures (e.g., surgery, devices, procedures and diagnostic tests); and 2) the literature is more robust. The amount of time a PA covers for services



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
 The amount of time a PA covers for services is limited by OAR 410-120-1320(7) which states that PAs can be approved and renewed up to one year at a time. Whenever possible, practice guidelines from clinical professional organizations such as the American Medical Association or the American Psychiatric Association, are used to establish PA frequency. 				 410-120-1320(7) which states that PAs can be approved and renewed up to one year at a time. Whenever possible, practice guidelines from clinical professional organizations such as the American Medical Association or the American Psychiatric Association, are used to establish PA frequency.
Data reviewed to determine UM	Data reviewed to determine	Data reviewed to determine UM	Data reviewed to determine UM	Data reviewed to determine UM
application:	UM application:	application:	application:	application:
A physician-led group of clinical professionals conduct an annual review to determine which services receive or retain a PA; items reviewed include:	• N/A	Denial/appeal overturn rates; number of PA and CR requests; complaints, and stabilization of cost trends.	Denial/appeal overturn rates; number of PA and CR requests; complaints, and stabilization of cost trends.	A physician-led group of clinical professionals conduct an annual review to determine which services receive or retain a PA; items reviewed include:



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
 Utilization Approval/denial rates Documentation/ju stification of services Cost data 				 Utilization Approval/denial rates Documentation/jus tification of services Cost data
IRR standard (OHA):	IRR standard:	IRR standard:	IRR standard:	IRR standard (OHA):
KEPRO has a formal policy including an 80% standard using InterQual criteria. IRR standard	Spot-checks performed through supervision.	• IRR testing standard of 85%.	• IRR testing standard of 85%.	KEPRO has a formal policy including an 80% standard using InterQual criteria.
(Comagine):				
 Spot-checks performed through supervision. Formal policy to be developed. 				
IRR standard (Comagine Health):				
• Spot-checks performed through supervision.				



FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S	
Analysis					

UM was applied to FFS MH/SUD and M/S HCBS benefits, and CCO MH/SUD and FFS M/S OP benefits listed in comparability and stringency Standard #1. For HCBS, MH/SUD benefits were administered by the Oregon Department of Human Services (DHS) and OHA's subcontractor, Comagine Health, while HCBS M/S benefits were administered by DHS. Pursuant to the 2020 CCO 2.0 Health Care Services Contract, the CCO did not require PA for MH/SUD services with the exception of more intensive care benefits such as ABA and psychiatric day treatment.

HSAG's analysis of IHN's PA data for IP and OP benefits did not reveal any concerns related to MHP. Of the total 6,411 IP and OP PA requests reported, 10.23 percent were denied. Of the 63 MH/SUD PA requests denied, representing 5.52 percent of the 1,142 MH/SUD PA requests, only three denials resulted in an appeal. The 63 reported MH/SUD denials represented 9.60 percent of total denials, the majority of which were for OP benefit requests, primarily denied due to a "non-contracted provider" denial reason.

Comparability

UM of MH/SUD and M/S HCBS benefits was required to meet federal HCBS requirements regarding person-centered service plans (PCSPs), providing benefits in the least restrictive environment, and applicable waiver applications/State plan amendments. Evidence for the application of UM to these benefits included federal requirements regarding PCSPs for 1915(c), 1915(i), 1915(k), and 1915(j) services and applicable approved waiver applications/State plan amendments. UM was applied to non-HCBS CCO MH/SUD, and M/S OP services were assigned UM to confirm coverage relative to the HERC PL and guidelines and federal guidelines. Non-HCBS MH/SUD services were also reviewed to ensure services were medically necessary relative to clinical practice guidelines and offered in the least restrictive environment that is safe, as required by the OPP Olmstead settlement for MH/SUD. A subset of CCO M/S OP services were also assigned UM to assure the individual's safety. Evidence for safety issues included HERC guidelines. HSAG determined the rationale and evidence to be comparable.

IHN and OHP FFS made authorization decisions within 24 hours for emergencies, 72 hours for urgent requests, and 14 days for standard requests. Providers were encouraged to submit requests for authorization sufficiently in advance to be consistent with OAR time frames. Most CCO documentation requirements for MH/SUD included an admission note and records submitted via telephone, fax, or electronically. CCO M/S was electronically notified of an admission, and care was reviewed via electronic health record (EHR). Alternatively, documentation could be submitted via fax. PCSPs for both M/S and MH/SUD must be developed within 90 days. The PCSP for both MH/SUD and M/S was based on an assessment and other relevant supporting documentation. It was developed by the member, the member's team, and the member's case manager. Qualified individuals conducted UM applying OARs, HERC, MCG, national guidelines, and ASAM for CCO SUD. MH/SUD and M/S DHS reviewers were required to have a BA in a related field; a BA in any field plus one year of experience; an AA with two years' experience; or three years' experience. The CCO and Comagine required all MH/SUD and M/S denials to be made by physicians or professional peers; however, nurses could deny M/S benefits managed by OHP FFS. HSAG determined this difference to be an issue of quality



	FFS HCBS MH/SUD	FFS HCBS M/S	CCO MH/SUD	CCO M/S	FFS M/S
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rather than a parity concern. HSAG ultimately determined that the MH/SUD PA review time frames, documentation requirements, and qualification of reviewers were comparable to those applied to M/S benefits.

Stringency

PA and CR frequencies for both MH/SUD and M/S benefits considered benefit coverage, medical guidelines, and member need. Both the CCO and OHP FFS allowed 90-day RR for MH/SUD and M/S when providers failed to obtain authorization. Exceptions to these time frames were considered by both the CCO and OHP FFS. MH/SUD and M/S denial decisions could be challenged through appeals and/or State fair hearing processes. Failure to obtain authorization could result in noncoverage. Regarding IRR, the CCO conducted annual reviews using an 85 percent testing standard, whereas OHP FFS's subcontractor had an 80 percent testing standard for M/S benefit authorizations. HSAG did not determine this discrepancy to be a parity concern as the method to promote consistency was more structured for MH/SUD benefits.

Outcome

HSAG determined IHN's processes, strategies, and evidentiary standards for UM of OP MH/SUD benefits to be comparable and no more stringently applied, in writing and in operation, to OP M/S benefits.



Category III—Prior Authorization for Prescription Drug Limits

NQTL: PA for Prescription Drugs

Benefit Package: CCOA and CCOB for adults and children

Classification: Prescription Drugs

Overview: PA is required for certain MH/SUD and M/S prescription drugs, and OHA requires PA of certain MH carve-out drugs. HSAG reviewed the reasons why CCOs and OHP FFS apply PA criteria to certain MH/SUD and M/S prescription drugs, the evidence used to establish PA criteria, and the processes used by the CCOs and OHP FFS to develop and apply PA criteria. HSAG analyzed IHN's application of PA for prescription drug benefits based on comparability and stringency standard information provided below.

	CCO MH/SUD	FFS MH Carve Out	CCO M/S
1.	To which benefit is the NQTL assigned?		
•	A, F, P, S drug groups	 A and F drug groups MH carve out drugs do not have an enforceable preferred drug list. While certain higher cost-effect agents are listed as "preferred," this is not enforced by PA. 	• A, F, P, S drug groups
2.	Why is the NQTL assigned to these bend	efits?	
•	Prior authorization is required for one or more of the following: evaluation of appropriateness of use, achieving the recommended daily dose, cost containment, and granting access to nonformulary medications.	To promote appropriate and safe treatment of funded conditions.	Prior authorization is required for one or more of the following: evaluation of appropriateness of use, achieving the recommended daily dose, cost containment, and granting access to nonformulary medications.



	ссо мн/sud	FFS MH Carve Out	CCO M/S
3.	What evidence supports the rationale fo	or the assignment?	
•	Drug class reviews created by pharmacists and in consultation with the P&T Committee based on best practices, professional guidelines and the Prioritized List. FDA prescribing guidelines and review of medical literature.	 FDA prescribing guidelines, medical evidence, best practices, professional guidelines, and P&T Committee review and recommendations. Federal and state regulations/OAR and the Prioritized List. 	 Drug class reviews created by pharmacists and in consultation with the P&T Committee based on best practices, professional guidelines and the Prioritized List. FDA prescribing guidelines and review of medical literature.
4.	What are the NQTL procedures? PA requests may be submitted by phone	PA requests are typically faxed to the	PA requests may be submitted by phone
	or fax.	Pharmacy Call Center, but requests can	or fax.
•	The standard PA request form is one page long, but most PA requests require chart notes. Requests are responded to within 24 hours. The PA criteria are developed by pharmacists and in consultation with the P&T Committee. Failure to obtain PA means the medication reimbursement will not be paid. CCO provides a 30-day notice prior to any negative change.	 also be submitted through the online portal, by phone, or by mail. The standard PA form is one page long, except for nutritional supplement requests. Most PA criteria require clinical documentation such as chart notes. All PA requests are responded to within 24 hours. The PA criteria are developed by pharmacists in consultation with the P&T Committee. Failure to obtain PA in combination with an absence of medical necessity results in no provider reimbursement. 	 The standard PA request form is one page long, but most PA requests require chart notes. Requests are responded to within 24 hours. The PA criteria are developed by pharmacists and in consultation with the P&T Committee. Failure to obtain PA means the medication reimbursement will not be paid. CCO provides a 30-day notice prior to any negative change.
		Notice of Benefit Determination sent to both Recipient and Provider Denials letters include information on required	



CCO MH/SUD	FFS MH Carve Out	CCO M/S
	criteria, denial reasons, and how the provider can appeal and member hearing rights.	
5. How frequently or strictly is the NQTL :	applied?	
PAs are authorized for six months to a year, depending on medical appropriateness and safety, as recommended by the P&T Committee.	The State approves PAs for up to 12 months, depending on medical appropriateness and safety, as recommended by the P&T Committee.	PAs are authorized for six months to a year, depending on medical appropriateness and safety, as recommended by the P&T Committee.
 Approximately 10.5% of MH/SUD drugs are subject to PA criteria for clinical reasons. 	 Approximately 19% of MH/SUD drugs are subject to PA criteria for clinical reasons. 	 Approximately 10.5% of M/S drugs are subject to PA criteria for clinical reasons. Providers can appeal on behalf of a client.
 Providers can appeal on behalf of a client. Documentation is collected and a pharmacist or the medical director reviews to determine if it is appropriate and should be approved or denied. A client can always have a hearing as well. The CCO assesses stringency through review of the number of PA requests, PA denial/approval rates, appeal overturn rates, and pharmacy pricing reports. PA criteria are reviewed for appropriateness every two years or as the standard of practice changes for the treatments. 	 The State allows providers to submit additional information for reconsideration of a denial. Providers can appeal denials on behalf of a member, and members have fair hearing rights. There were 10 client fair hearing requests for denied MH/SUD medications. None were reversed after agency reconsideration or, and none were reversed by hearing. The State assesses stringency through review of PA denial/approval and appeal rates; number of drugs requiring PA; number of PA requests; and pharmacy utilization data/reports. 	 Providers can appear on behalf of a chefit. Documentation is collected and a pharmacist or the medical director reviews to determine if it is appropriate and should be approved or denied. A client can always have a hearing as well. The CCO assesses stringency through review of the number of PA requests, PA denial/approval rates, appeal overturn rates, and pharmacy pricing reports. PA criteria are reviewed for appropriateness every two years or as the standard of practice changes for the treatments.



	CCO MH/SUD		FFS MH Carve Out		CCO M/S
		PA criteria are reviewed as needed due to clinical developments, literature, studies, and FDA medication approvals.			
6.	What standard supports the frequency	or r	igor with which the NQTL is applied?		
•	Drug class reviews created by pharmacists and in consultation with the P&T Committee based on best practices, professional guidelines and the Prioritized List. FDA prescribing guidelines, review of medical literature.	•	FDA prescribing guidelines, medical evidence, best practices, professional guidelines, and P&T Committee review and recommendations. Federal and state regulations/OAR and the Prioritized List.	•	Drug class reviews created by pharmacists and in consultation with the P&T Committee based on best practices, professional guidelines and the Prioritized List. FDA prescribing guidelines, review of medical literature.

IHN applied PA criteria to MH/SUD and M/S prescription drug benefits and applied PA to certain MH/SUD and M/S drugs to promote appropriate and safe treatment, and cost-effective use of prescription drugs. Since 2018, the CCO conducted an evaluation of its formulary and made changes that either added prescription drugs, added and removed PA criteria to prescription drugs, and adjusted criteria for prescription drugs in the formulary. PA was consistent across all benefit packages (CCOA, CCOB, CCOE, and CCOG).

HSAG's analysis of IHN's counts for prescription drug PA requests did not reveal any concerns related to parity. Of the total 3,955 prescription drug PA requests reported, 33.63 percent were denied. Less than 4 percent (3.76 percent) of the 1,330 prescription drug PA request denials were appealed, with 28 PA denials resulting in an overturned decision.

Comparability

The State applied PA to certain MH FFS carve-out drugs to promote appropriate and safe treatment. Evidence used by the CCO and OHP FFS to determine which MH/SUD and M/S drugs are subject to PA included Food and Drug Administration (FDA) prescribing guidelines, medical evidence, best practices, professional guidelines, and Pharmacy and Therapeutic (P&T) Committee review and recommendations. The PA criteria for both MH/SUD and M/S drugs were developed by pharmacists in consultation with the P&T Committee. PA requests for both MH/SUD and M/S drugs could be submitted by fax, phone, or online.



CCO MH/SUD	FFS MH Carve Out	CCO M/S

Stringency

For both MH/SUD and M/S drugs, most PA criteria required clinical documentation such as chart notes. Failure to obtain PA for MH/SUD and M/S drugs subject to PA in combination with an absence of medical necessity resulted in no reimbursement for the drug. Decisions were responded to within 24 hours, with decisions being made within 72 hours. For both MH/SUD and M/S drugs, the length of authorizations was dependent on medical appropriateness and safety, as recommended by the P&T Committee, based on clinical evidence such as FDA prescribing guidelines, best practices, and clinical practice guidelines. Both the CCO and OHA allowed exceptions to the formulary and preferred drug list based on medical necessity. For carve-out drugs covered by OHA, the CCO stated that it works with pharmacies and providers to redirect PA requests and claims to OHA.

Outcome

HSAG determined IHN's processes, strategies, and evidentiary standards for PA of MH/SUD prescription drugs to be comparable and no more stringently applied, in writing and in operation, to M/S prescription drugs.



Category IV—Provider Admission—Closed Network

NQTL: Provider Admission

Benefit Package: CCOA, CCOB, CCOE, and CCOG for adults and children

Classification: IP and OP

Overview: CCOs require providers of MH/SUD and M/S services to successfully meet credentialing and recredentialing requirements in order to be admitted to and continue to participate in the CCO's network. HSAG analyzed IHN's provider admission processes based on comparability and stringency standard information related to network closures provided below. Since Medicaid provider enrollment for OHP FFS did not include a provider credentialing component, HSAG deemed provider admission processes not applicable for OHP FFS and did not include that classification in the provider admission analysis.

	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
1.	To which benefit is the NQTI	assigned?		
•	CCO does not close its network for new MH/SUD providers of inpatient or outpatient services.	The State does not restrict new providers of inpatient or outpatient MH/SUD services from enrollment.	• N/A	The State does not restrict new providers of inpatient or outpatient MH/SUD services from enrollment.
2.	Why is the NQTL assigned to	these benefits?		
•	N/A	• N/A	• N/A	• N/A
3.	What evidence supports the r	ationale for the assignment?		
•	N/A	• N/A	• N/A	• N/A
4.	What are the NQTL procedu	res?		
•	N/A	• N/A	• N/A	• N/A
5.	How frequently or strictly is t	the NQTL applied?		
•	N/A	• N/A	• N/A	• N/A



	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S			
6	6. What standard supports the frequency or rigor with which the NQTL is applied?						
• N/A • N/A • N/A							

IHN did not close its network to providers of MH/SUD and M/S services. Developing a network based on network adequacy and sufficiency standards was supported by federal regulation, including the ability of a managed care organization (CCO) to limit contracting beyond the needs of its members to maintain quality and control costs (42 CFR §438.12). OAR 410-141-0220 also required the CCO to meet network sufficiency standards, which impacts the application of this NQTL category. In addition, provider network admission limits did not apply to FFS benefits and the application of provider network admission NQTLs for benefits delivered under managed care as supported by 42 CFR §438.206 and §438.12. Accordingly, parity was not analyzed.

Comparability

N/A

Stringency

N/A

Outcome

Because IHN did not close its network to either MH/SUD or M/S providers, HSAG determined that the CCO's provider admission/network closure processes for MH/SUD providers were comparable to and no more stringently applied to M/S providers across all benefit packages.



Category V—Provider Admission—Network Credentialing

NQTL: Provider Admission

Benefit Package: CCOA, CCOB, CCOE, and CCOG for adults and children

Classification: IP and OP

Overview: CCOs require providers of MH/SUD and M/S services to successfully meet credentialing and recredentialing requirements in order to be admitted to and continue to participate in the CCO's network. HSAG analyzed IHN's provider admission processes based on comparability and stringency standard information related to credentialing and recredentialing provided below. Since Medicaid provider enrollment for OHP FFS did not include a provider credentialing component, HSAG deemed provider admission processes not applicable for OHP FFS and did not include that classification in the provider admission analysis.

	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
1.	To which benefit is the NQTI	assigned?		
•	CCO requires all participating providers to meet credentialing and recredentialing requirements. CCO does not apply provider requirements in addition to State licensing.	 All FFS providers must be enrolled as a provider with Oregon Medicaid The State does not apply provider requirements in addition to State licensing. 	 CCO requires all participating providers to meet credentialing and recredentialing requirements. CCO does not apply provider requirements in addition to State licensing. 	 All FFS providers must be enrolled as a provider with Oregon Medicaid The State does not apply provider requirements in addition to State licensing
2.	Why is the NQTL assigned to	these benefits?		
•	CCO applies credentialing and re-credentialing requirements to: - Meet State and Federal requirements	Provider enrollment is required by State law and Federal regulations. The State also specifies requirements for provider enrollment in order to ensure beneficiary	CCO applies credentialing and re-credentialing requirements to: Meet State and Federal requirements	Provider enrollment is required by State law and Federal regulations. The State also specifies requirements for provider enrollment in order to ensure beneficiary



	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
3	 Ensure capabilities of providers to deliver high quality of care Ensure providers meet minimum competency standards What evidence supports the results of the providers of	health and safety and to reduce Medicaid provider fraud, waste, and abuse.	 Ensure capabilities of providers to deliver high quality of care. Ensure providers meet minimum competency standards 	health and safety and to reduce Medicaid provider fraud, waste, and abuse.
•	Credentialing/re- credentialing requirements are supported by the following evidence: - State law and Federal regulations, including 42 CFR 438.214 - State contract requirements - Medicare Manual, Chapter 6 - OAR 410-141-0120	Provider enrollment is required by State law and Federal regulations, including 42 CFR Part 455, Subpart E- Provider Screening and Enrollment.	 Credentialing/re-credentialing requirements are supported by the following evidence: State law and Federal regulations, including 42 CFR 438.214. State contract requirements. Medicare Manual, Chapter 6. OAR 410-141-0120. 	Provider enrollment is required by State law and Federal regulations, including 42 CFR Part 455, Subpart E- Provider Screening and Enrollment.
4.	What are the NQTL procedu	res?		
•	All providers must meet credentialing and recredentialing requirements. Providers must complete and provide Oregon Practitioner Credentialing Application (OPCA), Department of Human Services (DHS) forms DHS 3973 and/or 3974 (if applicable), copy of	All providers are eligible to enroll as a provider and receive reimbursement provided they meet all relevant Federal and State licensing and other rules and are not on an exclusionary list. Providers must complete forms and documentation required for their provider	 All providers must meet credentialing and recredentialing requirements. Providers must complete and provide Oregon Practitioner Credentialing Application (OPCA), Department of Human Services (DHS) forms DHS 3973 and/or 3974 (if applicable), copy of 	All providers are eligible to enroll as a provider and receive reimbursement provided they meet all relevant Federal and State licensing and other rules and are not on an exclusionary list. Providers must complete forms and documentation required for their provider



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
current Oregon Medical	type. This includes	current Oregon Medical	type. This includes
License, copies of Medical	information demonstrating	License, copies of Medical	information demonstrating
School Diploma and/or	the provider meets provider	School Diploma and/or	the provider meets provider
completion certificates from	enrollment requirements such	completion certificates from	enrollment requirements such
medical training, copies of	as NPI, tax ID, disclosures,	medical training, copies of	as NPI, tax ID, disclosures,
Internship and Residency	and licensure/certification.	Internship and Residency	and licensure/certification.
completion certificates, copy	The provider enrollment	completion certificates, copy	The provider enrollment
of Board Certification (if	forms vary from 1 to 19	of Board Certification (if	forms vary from 1 to 19
applicable), copy of current	pages, depending on the	applicable), copy of current	pages, depending on the
Drug Enforcement Agency	provider type. Supporting	Drug Enforcement Agency	provider type. Supporting
(DEA) certificate (if	documentation includes the	(DEA) certificate (if	documentation includes the
applicable), current copy of	provider's IRS letter,	applicable), current copy of	provider's IRS letter,
Malpractice Face Sheet	licensure, SSN number,	Malpractice Face Sheet	licensure, SSN number,
including 5 years of	and/or Medicare enrollment	including 5 years of	and/or Medicare enrollment
Malpractice Coverage	as applicable to the provider	Malpractice Coverage	as applicable to the provider
history, narrative of work	type. The enrollment forms	history, narrative of work	type. The enrollment forms
history, current signed	and documentation can be	history, current signed	and documentation can be
attestation by the applicant,	faxed in or completed and	attestation by the applicant,	faxed in or completed and
signed and dates	submitted electronically to	signed and dates	submitted electronically to
Authorization and Release of	the State's provider	Authorization and Release of	the State's provider
Information Form, and copy	enrollment unit. The State's	Information Form, and copy	enrollment unit. The State's
of policy and procedure for	provider enrollment process	of policy and procedure for	provider enrollment process
Seclusion and Restraint, as	includes checking the forms	Seclusion and Restraint, as	includes checking the forms
applicable.	for completeness, running the	applicable.	for completeness, running the
Providers may submit	provider name against	 Providers may submit 	provider name against
supporting documentation by	exclusion databases, and	supporting documentation by	exclusion databases, and
fax, electronic PDF file that	verifying any licenses,	fax, electronic PDF file that	verifying any licenses,
is sent by email, directly to	certifications or equivalents.	is sent by email, directly to its	certifications or equivalents.
office, or by US Mail.	The State's enrollment	office, or by US Mail.	The State's enrollment
CCO's credentialing process	process averages 7 to 14	• CCO's credentialing process	process averages 7 to 14
involves first source	days. State staff in the	involves first source	days. State staff in the
mvorves mst source	provider enrollment unit are	mvorves mist source	provider enrollment unit are



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
verifications. After first source verifications are completed, the completed files go to the Medical Director for approval. The Medical Director brings files that need further review to the credentialing committee. • CCO's credentialing process averages between 30 and 60 days. • CCO's Credentialing Committee is responsible for reviewing required information and making provider credentialing decisions.	responsible for reviewing information and making provider enrollment decisions	verifications. After first source verifications are completed, the completed files go to the Medical Director for approval. The Medical Director brings files that need further review to the credentialing committee. • CCO's credentialing process averages between 30 and 60 days. • CCO's Credentialing Committee is responsible for reviewing required information and making provider credentialing decisions.	responsible for reviewing information and making provider enrollment decisions
CCO performs recredentialing every 3 years.		CCO performs re- credentialing every 3 years.	
• Providers who do not meet credentialing/re-credentialing requirements do not get added to the CCO's panel of contracted providers or their contract with the CCO is terminated or that provider is removed from a group or facility that remains under contract with the CCO.		• Providers who do not meet credentialing/re-credentialing requirements do not get added to the CCO's panel of contracted providers or their contract with the CCO is terminated or that provider is removed from a group or facility that remains under contract with the CCO.	
Providers who are adversely affected by credentialing or		Providers who are adversely affected by credentialing or	



	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
	re-credentialing decisions may challenge the decision by challenging or appealing the decision of the CCO's Credentialing Committee within 45 days.		re-credentialing decisions may challenge the decision by challenging or appealing the decision of the CCO's Credentialing Committee within 45 days.	
5.	How frequently or strictly is t	the NQTL applied?		
•	All providers/provider types must be credentialed. There are no exceptions to meeting these requirements.	 All providers/provider types are subject to enrollment/re-enrollment requirements. There are no exceptions to meeting provider enrollment/re-enrollment requirements. 	 All providers/provider types must be credentialed. There are no exceptions to meeting these requirements. 	 All providers/provider types are subject to enrollment/re-enrollment requirements. There are no exceptions to meeting provider enrollment/re-enrollment requirements.
6.	What standard supports the f	frequency or rigor with which the I	NQTL is applied?	
•	Credentialing for all new providers is established by State law and Federal regulations. The frequency with which CCO performs recredentialing is based upon: State law and Federal regulations State contract requirements Monitoring of provider performance through the Appeals and Grievances Department may result in	Provider enrollment is required by State law and Federal regulations, including 42 CFR Part 455, Subpart E-Provider Screening and Enrollment. The frequency with which the State reenrolls providers is based on State law and Federal regulations.	 Credentialing for all new providers is established by State law and Federal regulations. The frequency with which CCO performs recredentialing is based upon: State law and Federal regulations State contract requirements Monitoring of provider performance through the Appeals and Grievances Department may result in 	Provider enrollment is required by State law and Federal regulations, including 42 CFR Part 455, Subpart E - Provider Screening and Enrollment. The frequency with which the State reenrolls providers is based on State law and Federal regulations.



FFS MH/SUD	CCO M/S	FFS M/S
	re-credentialing more frequently - National accreditation standards (NCQA)	
	 CCO monitors the following to determine how strictly to apply credentialing/re-credentialing criteria: Denial/Termination rates for providers as a result of credentialing/re-credentialing reviews. Provider appeals/disputes. Network Adequacy data. 	
	such as access to care, provider specialties.	
	CCO follows NCQA guidelines	
	FFS MH/SUD	re-credentialing more frequently - National accreditation standards (NCQA) • CCO monitors the following to determine how strictly to apply credentialing/re- credentialing criteria: - Denial/Termination rates for providers as a result of credentialing/re- credentialing reviews. - Provider appeals/disputes. - Network Adequacy data, such as access to care, provider specialties. • CCO follows NCQA

All IP and OP providers of MH/SUD and M/S services were subject to CCO credentialing and recredentialing requirements. IHN conducted credentialing and recredentialing for both providers of MH/SUD and M/S services to meet State and federal requirements, ensure providers are capable of delivering high-quality care, and ensure providers meet minimum competency standards. In addition, the CCO was following national accreditation guidelines (NCQA) to support the credentialing and recredentialing strategy for both MH/SUD and M/S providers. The CCO's processes were the same across all benefit packages (CCOA, CCOB, CCOE, and CCOG).

HSAG's analysis of IHN's provider credentialing data did not reveal any parity concerns related to high denial rates reported for providers seeking credentialing during the reporting period. Of the 11,903 reported average number of providers enrolled during the reporting period, 14.67 percent were MH/SUD providers. The total denial rate for all provider types was 22.22 percent. Of the 51 MH/SUD providers seeking credentialing during the reporting period, 25.49 percent were denied credentialing, which was fairly comparable to the 21.57 percent M/S denial



CCO MH/SUD FFS MH/SUD CCO M/S FFS M/S

rate. While there were only 13 MH/SUD provider credentialing denials, 46.15 percent of those denials were due to an "out of area" denial reason.

Comparability

IHN required providers of MH/SUD and M/S services to successfully meet credentialing and recredentialing requirements in order to be admitted to and continue to participate in the CCO's network. Providers were required to complete and submit a credentialing application and provide supporting documentation as part of the credentialing process. The type and extent of documentation required was determined by HSAG to be comparable. Both MH/SUD and M/S providers had several methods of submitting their application and supporting documentation, including by fax, by mail, or electronically. Nonlicensed MH care providers (e.g., qualified mental health providers/assistants and traditional health care works) were vetted similarly, with verifications completed by the CCO's credentialing department according to qualifications and certifications related to specific provider type.

The CCO's credentialing process for MH/SUD providers included the primary source verification of licensing, board certification, Medicare Excluded Providers (Office of Inspector General), Medicare sanction (Excluded Parties List System/System for Award Management), Medicare opt-out (if applicable), and a National Practitioner Database query match to look for unexplained gaps in work history greater than six months. The process for M/S providers involved a similar review of each application also conducted by the CCO's credentialing department to determine whether standards are met.

Stringency

The credentialing process averaged 30 to 60 days. The CCO's Credentialing Committee was responsible for reviewing required information and making provider credentialing decisions for both MH/SUD and M/S providers. Human Resources similarly conducted the credentialing function, reviewing information and making credentialing decisions for MH/SUD providers (based on staff model of service delivery). Recredentialing for both MH/SUD and M/S providers was conducted every three years. Failure of MH/SUD and M/S providers to meet credentialing and recredentialing requirements resulted in the denial or termination of participation as an INN provider for the CCO. MH/SUD and M/S providers who have been adversely affected by credentialing or recredentialing decisions may challenge the decision by challenging or appealing the decision of the CCO's Credentialing Committee within 45 days. MH/SUD and M/S providers were subject to meeting credentialing and recredentialing requirements; there were no exceptions reported. In operation, MH/SUD and M/S providers were determined to have been impacted equitably by the application of credentialing and recredentialing requirements, with no MH/SUD or M/S providers denied admission to the network during the reporting period.

Outcome

HSAG's analysis found IHN's credentialing processes and data for MH/SUD providers to be comparable and no more stringently applied to, in writing and in operation, than those for M/S providers.



Category VI—Out-of-Network/Out-of-State Limits

NQTL: OON and OOS limits

Benefit Package: CCOA, CCOB, CCOE, and CCOG for adults and children

Classification: IP and OP

Overview: OON/OOS services were required to provide coverage for needed MH/SUD and M/S benefits when they were not available INN or in-state. Similarly, for MH/SUD FFS benefits, OHP FFS provided OOS coverage to provide needed benefits when they were not available in-state. HSAG analyzed IHN's application of limits applied to OON/OOS limits based on comparability and stringency standard information provided below.

	CCO MH/SUD	FFS MH/SUD	CCO M/S FFS M/S	
1.	To which benefit is the NQTI	assigned?		
•	Out of Network (OON) and Out of State (OOS) Benefits	OOS Benefits	 Out of Network (OON) and Out of State (OOS) Benefits OOS Benefits 	
2.	Why is the NQTL assigned to	these benefits?		
•	CCO seeks to maximize use of in-network providers because our provider network consists of local providers that have been credentialed and contracted with the CCO. The purpose of providing OON/OOS coverage is to provide needed services when they are not available in-network/in-State.	The State seeks to maximize use of in-State providers because the State has determined that they meet applicable requirements, and they have a provider agreement with the State, which includes agreement to comply with Oregon Medicaid requirements and accept DMAP rates.	 CCO seeks to maximize use of in-network providers because our provider network consists of local providers that have been credentialed and contracted with the CCO. The purpose of providing OON/OOS coverage is to provide needed services when they are not available innetwork/in-State. The State seeks to maxim use of in-State providers because the State has determined that they mee applicable requirements, at they have a provider agreement with the State, which includes agreemen comply with Oregon Medicaid requirements an accept DMAP rates. 	et and , at to



	CCO MH/SUD		FFS MH/SUD		CCO M/S		FFS M/S
•	The purpose of prior authorizing non-emergency OON/OOS benefits is to determine the medical necessity of the requested benefit and the availability of an in-network/in-State provider.	•	The purpose of providing OOS coverage is to provide needed services when the service is not available in the State of Oregon or the client is OOS and requires covered services. The purpose of PA for non-emergency OOS services is to ensure the criteria in OAR 410-120-1180 are met.	•	The purpose of prior authorizing non-emergency OON/OOS benefits is to determine the medical necessity of the requested benefit and the availability of an in-network/in-State provider.	•	The purpose of providing OOS coverage is to provide needed services when the service is not available in the State of Oregon or the client is OOS and requires covered services. The purpose of PA for nonemergency OOS services is to ensure the criteria in OAR 410-120-1180 are met.
3.	What evidence supports the r	ration	nale for the assignment?				
•	The CCO covers OON/OOS benefits in accordance with Federal and State requirements, including OAR and the CCO contract.	•	The State covers OOS benefits in accordance with OARs.	•	The CCO covers OON/OOS benefits in accordance with Federal and State requirements, including OAR and the CCO contract.	•	The State covers OOS benefits in accordance with OARs.
4.	What are the NQTL procedu	res?					
•	Except as otherwise required by OHA, non-emergency OON/OOS services are not covered unless medically necessary services are not available within network/within State. The CCO's criteria for non- emergency OON/OOS coverage include member		Non-emergency OOS services are not covered unless the service meets the OAR criteria. The OAR criteria for OOS coverage of non-emergency services include the service is not available in the State of Oregon or the client is OOS and requires covered services.	•	Except as otherwise required by OHA, non-emergency OON/OOS services are not covered unless medically necessary services are not available within network/within State. The CCO's criteria for non- emergency OON/OOS coverage include member	•	Non-emergency OOS services are not covered unless the service meets the OAR criteria. The OAR criteria for OOS coverage of non-emergency services include the service is not available in the State of Oregon or the client is OOS and requires covered services.



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S	
 needs, network availability of services and wait times. Requests for non-emergency OON/OOS services are made through the prior authorization process. The timeframe for approving or denying a non-emergency OON/OOS request is the same as other prior authorizations (14 days for standard requests). The CCO establishes a memorandum of understanding (MOU) with an OON/OOS provider if the provider does will not accept the DMAP rate or the provider requests MOU for other reasons, e.g., to guarantee payment for covered services. The CCO's Contract and Negotiation Department negotiate the MOU with the OON/OOS provider; the MOU is signed by both parties, and the provider is directed to send claims associated with the MOU to a designated CCO 	 Requests for non-emergency OOS services are made through the State PA process. The timeframe for approving or denying a non-emergency OOS request is the same as for other PAs (14 days for standard and three business days for urgent). OOS providers must enroll with Oregon Medicaid. The State pays OOS providers the Medicaid FFS rate. 	 needs, network availability of services and wait times. Requests for non-emergency OON/OOS services are made through the prior authorization process. The timeframe for approving or denying a non-emergency OON/OOS request is the same as other prior authorizations (14 days for standard requests). The CCO establishes an MOU with an OON/OOS provider if the provider does will not accept the DMAP rate or the provider requests a MOU for other reasons, e.g., to guarantee payment for covered services. The CCO's Contract and Negotiation Department negotiate the MOU with the OON/OOS provider; the MOU is signed by both parties, and the provider is directed to send claims associated with the MOU to a designated CCO representative who works directly with a Senior Claims 	 Requests for non-emergency OOS services are made through the State PA process. The timeframe for approving or denying a non-emergency OOS request is the same as for other PAs (14 days for standard and three business days for urgent). OOS providers must enroll with Oregon Medicaid. The State pays OOS providers the Medicaid FFS rate. 	



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
representative who works directly with a Senior Claims Analyst to make sure the MOU is adjudicated based on the terms of the MOU. The CCO's process for establishing a MOU includes investigation of any potential provider option to suit the member's clinician need then establishing the rate and terms of service with the provider. The average length of time to negotiate a MOU is 1 to 2 days. In some cases, the MOU may take longer to complete if multiple billing providers are involved (e.g., hospitals). Only providers enrolled in Oregon Medicaid can qualify as an OON/OOS provider. The CCO pays OON/OOS providers the Medicaid FFS rate or a negotiated rate in the same range as the contracted rate.		Analyst to make sure the MOU is adjudicated based on the terms of the MOU. The CCO's process for establishing a MOU includes investigation of any potential provider option to suit the member's clinician need then establishing the rate and terms of service with the provider. The average length of time to negotiate a MOU is 1 to 2 days. In some cases, the MOU may take longer to complete if multiple billing providers are involved (e.g., hospitals). Only providers enrolled in Oregon Medicaid can qualify as an OON/OOS provider. The CCO pays OON/OOS providers the Medicaid FFS rate or a negotiated rate in the same range as the contracted rate.	



	CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S	
5.	How frequently or strictly is the NQTL applied?				
•	If a request for a non- emergency OON/OOS benefit does not meet the CCO's OON/OOS criteria, it will not be prior authorized. If a non-emergency OON/OOS benefit is not prior authorized, the service will not be covered, and payment for the service will be denied. Members/providers may appeal the denial of an OON/OOS request. The CCO evaluates the number of MOUs twice a year to determine whether the network should be expanded or a particular OON/OOS should be recruited to be a network provider.	 If a request for a non-emergency OOS benefit does not meet the OAR criteria, it will not be authorized. If a non-emergency OOS benefit is not authorized, the service will not be covered, and payment for the service will be denied. Members/providers may appeal the denial of an OOS request. The State measures the stringency of the application of OOS requirements by reviewing OOS denial/appeal rates. 	 If a request for a non-emergency OON/OOS benefit does not meet the CCO's OON/OOS criteria, it will not be prior authorized. If a non-emergency OON/OOS benefit is not prior authorized, the service will not be covered, and payment for the service will be denied. Members/providers may appeal the denial of an OON/OOS request. The CCO evaluates the number of MOUs twice a year to determine whether the network should be expanded or a particular OON/OOS should be recruited to be a network provider. 	 If a request for a non-emergency OOS benefit does not meet the OAR criteria, it will not be authorized. If a non-emergency OOS benefit is not authorized, the service will not be covered, and payment for the service will be denied. Members/providers may appeal the denial of an OOS request. The State measures the stringency of the application of OOS requirements by reviewing OOS denial/appeal rates. 	
6.					
•	Federal and State requirements, including OAR and the CCO contract.	The State covers OOS benefits in accordance with OAR.	Federal and State requirements, including OAR and the CCO contract.	The State covers OOS benefits in accordance with OAR.	



CCO MH/SUD	FFS MH/SUD	CCO M/S	FFS M/S
Analysis			

IHN ensured OON/OOS coverage to provide needed MH/SUD and M/S benefits when they were not available INN or in-state. Similarly, for MH/SUD FFS benefits, the State provided OOS coverage to provide needed benefits when they were not available in-state. The same PA processes and evidentiary standards described in NQTL categories I, II, and III were applied to OOS coverage of MH/SUD and M/S benefits across all benefit packages (CCOA, CCOB, CCOE, and CCOG). IHN established MOUs with OON providers in the absence of INN providers to ensure the provision of medically necessary services, while OHP FFS ensured OON providers were enrolled with Medicaid.

Comparability

For both nonemergency MH/SUD and M/S OON/OOS benefits, the CCO (and the State for FFS MH/SUD OOS benefits) required prior authorization to determine medical necessity and to ensure no INN/in-state providers are available to provide the benefit. The same PA processes and evidentiary standards described in NQTL categories I, II, and III were applied to OOS coverage of MH/SUD and M/S requests. For OON coverage requests, the CCO would determine if an INN provider was available or work with the OON provider to establish an MOU with payment of applicable Medicaid FFS rates. This process was applied equitably to both MH/SUD and M/S providers across all benefit packages.

Stringency

Requests for nonemergency OON/OOS CCO MH/SUD and M/S benefits were made through the CCO's PA process and reviewed for medical necessity and INN/in-state coverage. The PA time frames (14 days for standard requests and 72 hours for urgent requests) applied. Similarly, the State reviewed requests for nonemergency OOS MH/SUD services through its PA process, adhering to its PA time frames identified at 14 days for standard requests and 72 hours for urgent requests. The CCO described a process for handling a complex OON/OOS MH/SUD member case, identifying how it would appropriately apply the PA and SCA process to ensure benefits were provided in relation to the member's needs. IHN also provided an MOU template for review that identified compliant agreement information and confirmed the CCO's processes related to its use of OON providers. For both MH/SUD and M/S benefits, IHN and OHP FFS would not authorize payment for services denied.

Outcome

HSAG determined IHN's processes, strategies, and evidentiary standards for OON/OOS limits applied to MH/SUD to be comparable and no more stringently applied, in writing and in operation, to M/S OON/OOS limits across all benefit packages.



Appendix C. Improvement Plan Template

InterCommunity Health Network MHP Improvement Plan								
Year	Finding #	Report Reference	Finding	Required Action				
2020	1	Page. #						
CCO Intervention/Action Plan		Individual(s) Responsible	Proposed Completion Date					
HSAG Assessment of CCO Intervention/Action								
CCO Post-Implementation Status Update								
Documentation Submitted as Evidence of Implemented Intervention/Action								
HSAG Assessment of Intervention/Action Implementation								