Request for Applications:
Advancing Behavioral Health Equity in Primary Care

OVERVIEW

In partnership with the California Health Care Foundation (CHCF), the Center for Care Innovations (CCI) is launching Advancing Behavioral Health Equity in Primary Care — a new learning collaborative to help California community health centers expand and improve behavioral health outcomes with a specific focus on advancing health equity and aligning behavioral health and social needs resources. The learning collaborative will use an evidence-based integration model (the Stepped Model), emphasize patient-centered access (virtual and in-person care), and test new approaches for reducing health inequities. By the end of the program, participants will be able to:

• Identify, manage, and treat mental health conditions and substance use disorders.
• Identify and address patients’ unmet social needs (e.g., food insecurity) through consistent screening, tracking, and robust referral processes.
• Stratify their data to identify and understand where inequities are greatest.
• Take effective action to reduce barriers to care — specifically, racism, discrimination, stigma, and trauma — by actively embracing health equity practices.
• Sustain and spread their successes.

WHAT WE’LL PROVIDE

Community health centers accepted into the program will receive grants of between $75,000 and $125,000 based on the number of sites enrolled, community health center size, and number of priority populations identified. Grant dollars can be used to offset staff time, travel costs, and other program-associated costs.

CCI will provide comprehensive technical assistance, including:

• A baseline and endline capability assessment tool to help participants set improvement goals and monitor their progress over time.
• Virtual and in-person learning sessions, including a mix of expert- and peer-led sharing sessions on topics participants identify as challenges.
• Monthly coaching sessions from professionals with expertise in quality improvement, behavioral health integration, health equity, and/or social needs.
• In-person site visits to exemplar organizations in California and across the country. (Site visits will be conducted virtually if there are COVID-19-related travel restrictions or concerns.)
• An online learning community with peer- and expert-generated tools and resources.
• Support in defining metrics to measure and evaluate participant progress and collecting and using data for improvement.

PROGRAM TIMELINE

The learning collaborative will run for 20 months, from September 27, 2021, to May 31, 2023.

Key Dates:
  • **Informational Webinar**: Wednesday, August 4, 2021, at 12:00 PM–1:00 PM (PT)
  • **Application Deadline**: Tuesday, August 17, 2021, at 5 PM (PT)
  • **Cohort Announced**: Monday, September 20, 2021
  • **Program Start**: Monday, September 27, 2021
  • **Baseline Capability Assessment**: Completed with team and reviewed with coach by October 15, 2021
  • **Virtual Program Kickoff Meeting**: Wednesday, November 3, 2021, at 12pm-1:30pm (PT)

ELIGIBILITY

Eligible organizations include California-based:
  • Federally Qualified Health Centers (FQHCs) and FQHC Look-Alikes.
  • Community clinics, rural health clinics, and free clinics.
  • Ambulatory care clinics owned and operated by county health systems or public hospitals.
  • Indian Health Services clinics.

Participants will be selected with the goal of achieving diversity in terms of geographic location, community health center size, and patient mix (specifically, Black, Latinx, Asian American / Native Hawaiian and Pacific Islander, American Indian and Alaskan Native, LGBTQ, those who are seriously and persistently mentally ill, and adolescents and transition-age youth).

APPLY ONLINE
**Click here to submit your application.**
LEARN MORE

Join us for an Informational Webinar on Wednesday, August 4 from 12:00 PM–1:00 PM (PT) to hear a detailed description of the program and ask questions. Register here.

For any other questions, please contact:
Juliane Tomlin, Senior Manager, Juliane@careinnovations.org
About the Program

BACKGROUND

People with behavioral health (BH) conditions often experience poor health overall, and those with a diagnosis of serious mental illness or substance use disorder (SUD) die on average 20 years earlier than others, often from preventable physical illnesses. Only one-third of people with any type of mental illness receive treatment, and just 10 percent of people with substance use disorders are treated. In California and nationally, there are dramatic racial and ethnic inequities in access to and utilization of care. Within Medi-Cal, for example, Black, Latinx, and Asian enrollees receive needed care at lower rates than their White counterparts. These unmet behavioral health needs are a health and health equity crisis in California.

While county behavioral health plans are primarily responsible for Medi-Cal services for serious mental illness and substance use disorder in California, people with all levels of behavioral health and social needs present in primary care settings. In response, community health centers have greatly expanded their behavioral health capacity, and today most community health centers routinely screen for distress using validated screening instruments, such as the PHQ-9, and refer patients for treatment. However, only a fraction of community health centers provide a full suite of behavioral health services and proactively manage patient outcomes on par with their management of physical illnesses, such as diabetes (e.g., by using a registry to track symptom reduction for patients with depression). Additionally, a lack of alignment between behavioral health services and social resources (e.g., housing assistance) is commonly cited as a pain point for patients, families, and providers alike.

Given the growing mental health crisis in California, continuing to expand access to integrated care is a priority for California’s community health centers, policymakers, and other stakeholders. This learning collaborative was developed to help community health centers with existing commitment and capability for integrated care expand and strengthen their behavioral health program (e.g., offer new services or supports), align programs with social needs care/resources, and adopt health equity practices that can improve access, utilization, and outcomes for priority populations: Black, Latinx, Asian American / Native Hawaiian and Pacific Islander, American Indian and Alaskan Native, those with serious and persistent mental illness, transitional age youth, and LGBTQ populations.

Program Structure & Core Content

CURRICULUM TOPICS

The learning collaborative will address the following topics with an eye towards advancing health equity, patient and family engagement, and a culture of improvement.
• **Leadership & Organizational Commitment:** With dedicated leadership and a clear organizational strategy, participants will prioritize health equity in their efforts to integrate behavioral health, modeling this commitment by taking specific actions, such as:
  - Investing time in transparent hiring practices with a goal for leadership and staff to reflect the community they serve.
  - Integrating knowledge about trauma into policies, procedures, and practices.
  - Inviting patients and those with lived experience to guide strategy development.

• **Data Driven Systems and Decisions:** Participants will build and strengthen their abilities to deploy accurate and illustrative data in identifying and managing their population with behavioral health needs (e.g., registries), performance reporting, and decision making at the point of care. Participants will invest in collecting and stratifying data that can drive equitable outcomes for patients in need of behavioral health, substance use, and social needs services.

• **Access:** Participants will expand access to behavioral health by ensuring that all patients have access to timely, coordinated, culturally competent care.

• **Care Team/Care Delivery:** Participants will establish (or deepen) an integrated care team with clear roles. They will support care teams in providing evidence-based, trauma-informed, data-driven screening, assessment, and management. They will establish levels of care, clearly defining when and where patients will be referred to a higher level of care (e.g., specialty care).

• **Patient Activation/Self-Management:** Participants will create environments where patients and caregivers are engaged, informed, and empowered to participate in their care.

• **Community Partnerships:** Participants will identify and develop relationships with community-based organizations and specialty care providers; Participants will set up clear agreements regarding roles, services provided, workflows and referrals, shared data, and referral platforms.

**SUPPORTING ACTIVITIES**

• **Learning sessions** to provide timely content from experts and support peer sharing. We will offer a total of four learning sessions (two virtual and two in-person when it is safe to convene). In-person learning sessions will be one full day, while virtual sessions will be up to four hours. (Tentatively: Dec 2021, Q2-2022, Q4-2022, Q2-2023).
• **Site visits** to allow participants to visit an exemplar organization and make connections with peers. One to two one core team members will attend at least one site visit. Site visit locations will be offered in California and nationally. (Tentatively: Q2-2022, Q3-2022, Q4-2022). Site visits will be conducted virtually if there are COVID-19-related travel restrictions or concerns.

• **Quarterly webinars** to provide foundational content and spotlight best practices. Webinars will be between 60-90 minutes in length and will take place during months without other program offerings.

• **Monthly coaching sessions** to enable teams to apply program resources and learnings to their own projects. Coaches will help participants identify what they are trying to accomplish, what changes they will test/implement, and how they are using data to measure their success. Coaches will jointly problem solve challenges and connect teams to resources to advance their goals. (Coaching will start in October 2021 with the first session dedicated to reviewing and refining the baseline capability assessment.)

• **Online learning community:** The CCI Academy will be used to post program content, enable peer connection, and share peer- and expert-generated tools and resources between learning sessions and webinars.

• **Measures/data support:** CCI will help teams develop their own internal measures and data collection strategies to evaluate progress towards their self-identified goals, including a customized template and training on how to report data on internal measures.

• **Participant reporting of progress and challenges:** Participants will report their progress in quarterly progress reports, outlining promising practices, challenges, and where additional support from the CCI team would be helpful. Note this reporting requirement is in addition to the evaluation activities described below.

• **Final toolkit:** Participants will contribute relevant tools and resources to a final toolkit that will be published to help other community health centers advance behavioral health integration and health equity.

**EVALUATION ACTIVITIES**

A goal of the learning collaborative is to identify the practices, approaches, methods, and tools that facilitate progress, particularly with regards to health equity. CHCF has engaged the American Institutes for Research (AIR) to lead an independent evaluation of the learning collaborative’s impact and to identify lessons of use to the field. Participants will be required to submit data on a universal measure set at two points during the program (pre-implementation data may be requested, if available), complete surveys, participate in interviews and/or focus groups, and engage in other activities to understand and document change in the following domains: practice capability (e.g., delivery of trauma-informed care); patient impact (access, experience, and outcomes); equity; and staff experience. Refer to Appendix A for more information on the evaluation.
COMMUNICATIONS

Another goal is to tell the story of how the program is helping patients and transforming organizations. Participants will be expected to collaborate with CCI and CHCF on communications activities, which could include identifying patient stories and testimonials, helping to facilitate video shoots with patients and organization staff to capture both groups’ experiences in the program, and helping to facilitate photo shoots at community health centers to capture images that bring to life program operations and services.

Eligibility

WHO IS ELIGIBLE?

Organizations in California that provide comprehensive primary care services to historically underinvested populations are eligible to apply. Applicants must be nonprofit and tax-exempt under 501(c)(3) of the Internal Revenue Service Code or a governmental, tribal, or public entity.

Eligible organizations include:
- Federally Qualified Health Centers (FQHCs) and FQHC Look-Alikes.
- Community clinics, rural health clinics, and free clinics.
- Ambulatory care clinics owned and operated by county health systems or public hospitals.
- Indian Health Services Clinics.

Participants will be selected with the goal of achieving diversity in terms of geographic location, community health center size, and patient mix (Black, Latinx, Asian American / Native Hawaiian and Pacific Islander, American Indian and Alaskan Native, LGBTQ, those who are seriously and persistently mentally ill, and adolescents and transition-age youth).

MINIMUM EXPECTATIONS FOR BEHAVIORAL HEALTH INTEGRATION

CCI assumes participants enter the learning collaborative with core behavioral health integration capability already in place. This project is not suitable for health centers that do not currently have some degree of integrated care. To be eligible, applicants must be able to demonstrate:
- Co-located or fully-integrated care, at a minimum having behavioral health, primary care, and other health care providers work in the same facility or office space, where they communicate regularly about shared patients (by phone, email, or in-person); collaborate, driven by a need for each other’s services and more reliable referral, need for consultation and coordinated plans for patients, or as colleagues on the same care
team; meet to discuss cases (occasionally or regularly); and feel part of a larger team (formally or informally).

- Ability to identify populations and subpopulations via a registry, electronic health record, or population management software. Applicants must also have the capability to report data at the population level for common behavioral health measures such as number of patients screened, number of patients with behavioral health conditions, number of completed referrals (e.g., to social services, specialty care).

If you are unsure if your organization meets these requirements, please contact Juliane Tomlin (juliane@careinnovations.org) before proceeding with the application.

**What Makes a Strong Applicant?**

CCI is looking for applicants with the following characteristics:

- **Organizational commitment to advancing behavioral health integration as demonstrated by having:**
  - Influential behavioral health and primary care clinician champions who advocate for and provide clinical support to the participating team.
  - A commitment to protect the necessary time of core team members.
  - An administrative champion who supports your team and enables protected time to participate in the program and achieve project goals.

- **Commitment to expanding the range of behavioral health services or supports available to patients:** Care may address mental health and substance use conditions, health behaviors (including their contribution to chronic medical illnesses), life stressors and crises, stress-related physical symptoms, and ineffective patterns of health care utilization.

- **Commitment to health equity and racial justice such as:**
  - The organization invests in staff understanding of the impact of historical trauma and structural racism on patients and families.
  - The backgrounds of leadership and staff reflect the patient population.
  - Data are disaggregated and used to understand and act on inequities.
  - Patients and families who are representative of the patient population are engaged in efforts to understand and solve problems and improve care.

- **Demonstrated improvement capability:**
  - Successful applicants will show that they have achieved measurable improvements in care using human-centered design, performance improvement, or some other change method.
  - Applicants will show that they have the data collection and reporting capability to improve behavioral health care and health equity.
Participant Expectations

1. **Assure leadership buy-in:** Successful participants will have leadership that is committed to advancing equitable, integrated behavioral health care. We expect strong clinical and administrative leadership support from both medical and behavioral health disciplines (the chief clinical or medical officer/director, behavioral health officer or director, chief operating officer, or their equivalents), as demonstrated through a letter of leadership support committing to engage in the program in at least three ways:
   a. Join a coaching call in Year 1 focused on how to support the team’s action plan.
   b. Attend a virtual or in-person session by the end of Year 1 where the team shares work-to-date or a pitch for upcoming efforts (expected fall 2022, date TBD).
   c. Meet with the team in Year 2 to discuss spreading and sustaining the progress and accomplishments of program participants to date.

2. **Develop a team:**
   a. Core Team: Four to six individuals are required to be committed members of a core program team. The core team should include the following roles and include both primary care and behavioral health representatives:
      i. A project lead who is responsible for day-to-day activities and can coordinate the work of the team.
      ii. A behavioral health care provider and leader to champion changes in their clinic and influence change across the organization.
      iii. A physical health care provider and leader to champion changes in their clinic and influence change across the organization.
      iv. At least one frontline staff member (e.g., medical assistant, care manager, care coordinator, call center representative) who will co-develop, test, and implement changes.
      v. Other team members could include those with expertise in data, systems management/electronic health record, etc.

   b. Extended Team: One to two individuals are required to be committed members of the extended program team:
      i. A senior administrative leader sponsor (e.g., chief executive officer, executive director, chief operating officer) who can ensure protected time for team members to participate in program activities and lead change within their organization.
      ii. A clinical leader, such as a medical director or chief medical officer, with decision-making authority over behavioral health care efforts in the organization.
      iii. Clinical and front-line champions and staff from participating clinics should actively participate in relevant coaching calls and learning sessions.
c. Recommended, if available, but not required:
   i. Family specialist, navigator, peer support specialist, community health
      worker, patient, or family advisor(s).

3. **Involve patients and community representatives:**
   a. Successful participants will involve patients and community members’ perspectives
      and experiences in multiple ways, for example, in goal setting, development of
      workplans, and/or review of data.
   b. Participants will participate in communications activities to capture their patient
      experience in the learning collaborative (e.g., patient testimonials or videos).

4. **Actively participate in program events and activities** (see above for details) to learn,
   teach, and collaborate transparently with other program participants (participation as
   outlined below will be tied to grant funding):
   a. Core Team must commit to the following:
      i. Full and active participation at collaborative learning sessions, webinars, and
         monthly coaching calls. At least one Core Team member will attend a site
         visit.
      ii. Commit to action: Take what is learned from the learning collaborative,
          share, and implement in your organization, and report back on those efforts.
      iii. Share approaches, lessons learned, and failures in improvement efforts,
          including documenting tools, materials, and solutions to share with a public-
          facing audience, with attribution (e.g., through final toolkit).
      iv. Prioritize connection: Build connections during learning sessions and continue
          collaboration between learning sessions (e.g., contribute on the virtual
          learning platform).
      v. Complete defined pre-work assignments for virtual and in-person sessions
         and share lessons learned by presenting examples of project successes and
         challenges.
      vi. Complete the capability assessment with your team at the baseline, midpoint,
          and endpoint of the program, and leverage for goal setting and tracking.
      vii. Participate in the data collection and evaluation activities described in
           Appendix A.
   b. The Extended Team must participate in one learning session a year and at least
      three coaching sessions a year.

**How to Apply**

**STEP 1: ATTEND THE INFORMATIONAL WEBINAR.** Applicants are encouraged to
participate in an informational webinar on August 4, 2021, at 12:00 PM–1:00 PM (PT) to hear a
detailed description of the program and ask questions. Register here.
**STEP 2: APPLY ONLINE.** Your proposal and budget must be submitted online [HERE](#) by Tuesday, August 17, 2021, at 5 PM (PT).

Applications should include the following:
1. Application Narrative (see below)
2. Budget Template: [Download template](#)
3. Tax Status Documentation
4. Letter of Leadership Support from the chief medical officer, chief operating officer, or chief executive officer: see “Participant Expectations” above for requirements.

**APPLICATION NARRATIVE QUESTIONS**

Please limit responses to no more than 7 pages (shorter preferred), using at least 11-point font and 1-inch margins. See suggested length for each item.

1. **Commitment to Equity (<1.5 pages):** CCI strives to ensure that “everyone has fair, just, and inclusive opportunities to be healthy.” Similarly, “CHCF is working with a wide range of partners to break down structural barriers to care and create a health care system that is designed to redress, not perpetuate, the inequities brought about by barriers to care such as race or ethnicity, immigration status, income, or ZIP code.” Throughout this program, participants will rigorously examine and address barriers within their system, considering anti-racism and trauma-informed practices. Please address the following:
   a. Describe steps your organization has taken to mitigate the impact of racism, stigma, and discrimination on the patients and community you serve and reduce barriers to access, engagement, and outcomes. These might include changes to clinical care, workflows, patient outreach, staffing models, or services provided by your organization or through community partners.
   b. How do you identify patients at greatest risk for care gaps or poor health outcomes, and how do you address their needs and preferences?
   c. How are you addressing equity within your organization?
   d. What priority subpopulations do you plan to focus on in the learning collaborative? Participants should identify one to four subpopulations such as: Black, Latinx, Asian American / Native Hawaiian and Pacific Islander, American Indian and Alaskan Native, LGBTQ, those who are seriously and persistently mentally ill, and adolescents and transition-age youth). (Note: This program is not focusing on children as a subpopulation.)

2. **Behavioral Health Integration in Primary Care (<1.5 pages):** Please refer to [this graphic](#) to answer the following questions.
   a. Describe your current level of behavioral health integration. Include the titles and roles of those on your behavioral health integration team and how you
communicate, collaborate, and co-manage your patients. Describe how integration has changed in response to the COVID-19 pandemic.

b. Which behavioral health conditions do you currently treat (e.g., depression, anxiety, PTSD, SUD, trauma, domestic violence)?

c. Which social needs (e.g., food insecurity, housing instability) do you routinely address?

d. What level of integration do you hope to reach by the end of this 20-month program?

e. What services, supports, approaches, or treatments do you hope to add to your current program?

3. **Strategy Alignment (<0.5 pages):**
   a. How do the goals of this program align with your organization’s current goals on behavioral health integration, social needs, and health equity? What specific aspects of the learning collaborative do you think will help you be successful?
   b. What other current initiatives and grants do you hope to align or integrate with this learning collaborative so you can concentrate your efforts?

4. **Spread and Scale (<0.5 pages)**
   a. How many sites (physical locations) does your health center operate?
   b. Which sites will participate in the learning collaborative, either initially or as spread locations?
   c. How many patients are currently served at each of these sites?

5. **Quality Improvement Experience (<0.75 pages):**
   a. Please describe a clinical or operational improvement project you implemented in your organization over the past one to two years. What was the aim statement? What measurable level of improvement did you achieve?
   b. What data did you collect to assess progress and how often did you collect it?
   c. Describe one small test or prototype you did and what you learned.
   d. If you use a consistent performance improvement, human centered design, or change management method, what is it?

6. **Data Collection and Measure Reporting (<1 page):**
   a. Please describe the data you currently collect to identify and manage your patients with behavioral health needs. Are these data standardized by any operational process? If so, please list the names of the systems you use.
   b. What behavioral health measures are you regularly reporting to funders or state or federal agencies?
   c. Please list the names of any data systems you use to manage your behavioral health and/or social need data and referrals (e.g., registry names, electronic health record vendor names, population management solution systems).
d. What information are you collecting on social needs screening and referrals? Please list any specific measures or tools that you use?

e. What measures, if any, are you currently using to assess equity in care delivery?

f. What types of patient characteristics do you use to assess equity? Where are characteristics data stored? Briefly describe how these data are used in identifying and managing patients, and if relevant, in your internal or external reporting activities.

**Next Steps**

CCI and our program partners will review applications. We may contact you if we have questions about your application. The cohort will be announced via email by Monday, September 20, 2021.
Appendix A: Overview of Program Evaluation

AIR’s evaluation will assess the impact of the learning collaborative on participating grantees and their providers, staff, and patients. The evaluation will address a series of research questions, such as:

- What improvements in organizational structure and practices, staff and patient experiences, and clinical outcomes are we observing among participating organizations and what is driving these improvements?
- To what extent, and how, are participants able to identify inequities in practice (e.g., prescription practices, screening, diagnosis, referral, and treatment) and address these inequities?

To answer the research questions, AIR will collect data from grantees at two points in the learning collaborative (pre-implementation data may be requested, if available). AIR anticipates collecting approximately 10–15 standard clinical quality measures and 10–15 evaluation team developed measures. These measures will be collected at the patient population level as well as stratified by race, ethnicity, language, and sexual orientation/gender identification, as available. Measures and indicators under consideration for the evaluation are shown in Table A-1. Note that these indicators and measures are being considered for inclusion in the evaluation and are not yet final. The final list of measures and indicators will be determined based on the cohort of participants, their selected priority populations, and the information shared in the responses to this request for applications.

In addition to clinical quality measures, AIR will collect data on the advancement of equity and use it to calculate indicators of progress toward the learning collaborative’s goals. These data may be collected via document reviews (e.g., of protocols/standard operation procedures/job aides, screening tools), organizational surveys, a standardized capability assessment (e.g., to gain an understanding of payer mix; behavioral health treatment services provided; number, credentials, demographics of providers and staff; and patient demographics collected through self-attestation), and site visits with participating grantees.

Last, dependent upon feasibility and availability of pre-existing data, the evaluation team may also conduct surveys, interviews, or focus groups with providers, staff, and patients to assess their experiences with practice changes and services.

All data collection will occur twice over the implementation period to allow for pre-post comparisons; pre-implementation data may be requested, if available. The evaluation team will analyze these data to assess performance relative to benchmarks as well as changes over time. The following table outlines the types of measures under consideration and the anticipated data source or reporting method for the measure. The AIR team will provide technical support and, if needed, templates for collecting these indicators at the population level, and stratifying by race, ethnicity, language, and sexual orientation/gender identification.
<table>
<thead>
<tr>
<th>Measure or Indicator Name*</th>
<th>Reporting Method</th>
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<tbody>
<tr>
<td>Health Care and Outcomes</td>
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<tr>
<td>Depression Utilization of the PHQ-9 Tool (NQF 0712e)</td>
<td>EHR or paper medical records</td>
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<tr>
<td>Depression Response at Six Months- Progress Towards Remission (NQF 1884)</td>
<td>EHR or paper medical records</td>
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<tr>
<td>Initiation and Engagement of Alcohol and Other Drug Dependence Treatment (NQF 0004)</td>
<td>Claims</td>
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<tr>
<td>Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention (NQF 0028e)</td>
<td>EHR</td>
</tr>
<tr>
<td>Substance Use Screening and Intervention Composite (NQF 2597)</td>
<td>EHR</td>
</tr>
<tr>
<td>SBIRT Unhealthy Alcohol Use Screening and Brief Counseling (NQF 2152)</td>
<td>EHR or Registry</td>
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<tr>
<td>Closing the Referral Loop: Receipt of Specialist Report</td>
<td>EHR</td>
</tr>
<tr>
<td>Number/ percentage of patients that are referred to treatment by health need/diagnosis of mental health or substance use disorder</td>
<td>EHR</td>
</tr>
<tr>
<td>AAFP Social Needs Screening Tool Utilization and Need Prevalence</td>
<td>EHR</td>
</tr>
<tr>
<td>Mental Health and Substance Use Disorder Diagnoses;</td>
<td>EHR</td>
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<tr>
<td>Behavioral Health Care Utilization (defined based on applicant focus areas)</td>
<td>Claims data</td>
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<tr>
<td>Patient Experience</td>
<td></td>
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<tr>
<td>Gains in Patient Activation (PAM) Scores at 12 Months (NQF 2483)</td>
<td>Instrument-Based Data</td>
</tr>
<tr>
<td>CAHPS Experience of Care and Health Outcomes Measures:</td>
<td>Patient survey</td>
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<tr>
<td>• Q7 Get appointment as soon as wanted</td>
<td></td>
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<tr>
<td>• Q12 Clinicians explain things</td>
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<td>• Q17 Told about side effects of medication</td>
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<td>• Q18 Involved as much as you wanted in treatment</td>
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<td>• Q19 Talk about including family and friends in treatment</td>
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<td>• Q21 Told about different treatments that are available for condition</td>
<td></td>
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<tr>
<td>• Q27 Care responsive to cultural needs</td>
<td></td>
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<tr>
<td>Care Coordination Quality Measure for Primary Care (CCQM-PC)</td>
<td>Patient survey</td>
</tr>
<tr>
<td>Care Delivery, Cultural Competency, and Organizational Commitment</td>
<td></td>
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<tr>
<td>Number and percentage of clinic staff with workforce training competencies completed</td>
<td>Clinic Data</td>
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<tr>
<td>Competency trainings include CLAS Standards (cultural humility, recognition and addressing of implicit bias), SUD stigma education, and others</td>
<td>Clinic Data</td>
</tr>
<tr>
<td>Care team’s perception of trainings as culturally competent/informative for care delivery.</td>
<td>Key Informant Interviews or Staff Survey</td>
</tr>
</tbody>
</table>

*Outcome data on key metrics can be disaggregated by race/ethnicity, sexual orientation and gender identity, languages spoken, ZIP code.