

# HEDIS® Perinatal Depression Quality Measures: Field Test



NCQA is aiming to specify and test existing HEDIS®<sup>1</sup> depression care quality measures for prenatal/postpartum women. **We are seeking health plans and provider groups to participate in a field test.**

With funding from the **California Health Care Foundation** and the Colorado-based **Zoma Foundation**, NCQA will use its multi-stakeholder process to develop the measures. **Our overall goal** is to assess whether the existing depression measures can be stratified or adapted for perinatal women, or whether other options exist for capturing these quality issues.

## Measures

There are [three HEDIS measures](#) assessing depression screening and follow-up, monitoring and remission/response for adolescent and adult health plan members. These measures require reporting from electronic clinical data systems that include:

Claims data	<ul style="list-style-type: none"> <li>• Provide information on care received (e.g., psychotherapy visits, medication claims)</li> </ul>	<p>Plans report these measures using a structured technical specification that directs the collection of data from multiple sources in a consistent fashion.</p> <p>Measures using this reporting method have conformance rules requiring the sharing of this information with the providers at the point of care.</p>
Clinical databases (EHRs, registries)	<ul style="list-style-type: none"> <li>• Capture information on depression screening or assessments (e.g., whether a depression screen was positive and change in score over time)</li> </ul>	
Case management systems	<ul style="list-style-type: none"> <li>• Track members with depression over time to monitor symptoms, severity and access to appropriate follow-up services</li> </ul>	

## Project Phases and Anticipated Timeline

Testing Phase	Description	Anticipated Timeline
<b>Phase One: Patient-level data submission</b>	<ul style="list-style-type: none"> <li>• Test sites collect data on depression screening/care for prenatal/postpartum women and provide patient-level csv data file to NCQA.</li> <li>• Test sites participate in conference calls to discuss experiences with retrieving the requested data and specific challenges and lessons learned from the process.</li> </ul>	<p>Data collection: July-August 2018</p> <p>Feasibility interviews: September 2018</p>
<b>Phase Two: Certification of measure calculations</b>	<ul style="list-style-type: none"> <li>• Test sites program digital measure specifications</li> <li>• NCQA creates unique sets of sample data or "test decks" for each measure</li> <li>• Test sites process test decks through their downloaded measure code and return results for each patient in the test data</li> <li>• NCQA determines if measures compute correctly in the organization's system by comparing results for each patient to expected results.</li> </ul>	<p>Certification process: January-March 2019</p>

Plans and provider groups can participate in one or both testing phases and an honorarium will be provided.

For more information or to participate, contact Cindy Manaoat ([manaoat@ncqa.org](mailto:manaoat@ncqa.org)).

<sup>1</sup>HEDIS® is a registered trademark of the National Committee for Quality Assurance (NCQA)