HEDIS® Tip Sheet

Depression Remission or Response for Adolescents and Adults (DRR-E)

Measure Description

The percentage of members 12 years of age and older with a diagnosis of depression and an elevated PHQ-9 score, who had evidence of response or remission within 120-240 days (4-8 months) of the elevated score.

- Follow-up PHQ-9: The percentage of members who have a follow-up PHQ-9 score documented within 120-240 days (4-8 months) after the initial elevated PHQ-9 score.
 - o Elevated PHQ-9 scores are >9.
- Depression Remission: The percentage of members who achieved remission within 120-240 days (4-8 months) after the initial elevated PHQ-9 score.
 - Most recent PHQ-9 total score of <5 documented during the depression follow-up period.
- Depression Response: The percentage of members who showed response within 120-240 days (4-8 months) after the initial elevated PHQ-9 score.
 - Most recent PHQ-9 total score being at least 50% lower than the score associated with the initial elevated PHQ-9 total score >9 documented during the depression follow-up period.

Product Lines: Commercial, Medicaid, Medicare

Codes Included in the Current HEDIS® Measure

Codes to Identify Depression

Value Set Description	Code
Major Depression or Dysthymia	ICD-10: F32.0-F32.5, F32.9, F33.0-F33.3, F33.40-F33.42, F33.9, F34.1

Codes to Identify PHQ-9 Total Score

Direct Reference Code Display	Code
Patient Health Questionnaire 9 item (PHQ-9) total score [Reported]	LOINC: 44261-6
Patient Health Questionnaire-9: Modified for Teens total score [Reported PHQ Teen]	LOINC: 89204-2, 44261-6

Ways Providers can Improve HEDIS® Performance

- Establish and maintain follow-up with adult patients who have depression. Appropriate, reliable follow-up is highly correlated with improved response and remission scores.
- The PHQ-9 assessment does not need to occur during a face-to-face encounter; it may be completed over the telephone or through a web-based portal.
- Always offer general checkups and follow-ups, even if a behavioral health provider is following the patient.

Ways Health Plans can Improve HEDIS® Performance

- Educate providers to utilize PHQ-9 form in EMR to ensure included in electronic measure.
- Support providers in considering telemedicine visits when in-person visits are not available.
- Educate the member about the importance of timely follow-up and adherence to treatment recommendations.
- Assign care/case managers to members to ensure members keep follow-up appointments or reschedule missed appointments.
- Encourage providers to establish electronic file transfers to ensure documentation of evidence of response or remission within 120-240 days.

The Healthcare Effectiveness Data and Information Set ($HEDIS^{\circ}$) is a registered trademark of NCQA. The NCQA HEDIS measure specification has been adjusted pursuant to NCQA's *Rules for Allowable Adjustments of HEDIS*. The adjusted measure specification may be used only for internal quality improvement purposes. Updated 12/6/2024.

Exclusions

- Members who use hospice services or elect to use a hospice benefit any time during the measurement period.
- Members who die any time during the measurement period.
- Members with any of the following at any time during the Intake Period or during the measurement period:
 - o Bipolar disorder.
 - o Personality disorder.
 - o Psychotic disorder.
 - o Pervasive developmental disorder.



All summaries of the measures contained herein are reproduced with permission from HEDIS® Volume 2: Technical Specifications for Health Plans by the National Committee for Quality Assurance (NCQA).

The information presented herein is for informational and illustrative purposes only. It is not intended, nor is it to be used, to define a standard of care or otherwise substitute for informed medical evaluation, diagnosis and treatment which can be performed by a qualified medical professional. Molina Healthcare, Inc. does not warrant or represent that the information contained herein is accurate or free from defects.

COPYRIGHT NOTICE AND DISCLAIMER

HEDIS* is a registered trademark of the National Committee for Quality Assurance (NCQA). The HEDIS measures and specifications were developed by and are owned by NCQA. NCQA holds a copyright in these materials and may rescind or alter these materials at any time. Users of the HEDIS measures and specifications shall not have the right to alter, enhance or otherwise modify the HEDIS measures and specifications, and shall not disassemble, recompile or reverse engineer the HEDIS measures and specifications. Anyone desiring to use or reproduce the materials, subject to licensed user restrictions, without modification for an internal non-commercial purpose may do so without obtaining any approval from NCQA. Use of the Rules for Allowable Adjustments of HEDIS to make permitted adjustments of the materials does not constitute a modification. All other uses, including a commercial use (including but not limited to vendors using the measures and specifications with a product or service to calculate measure results), or any external reproduction, distribution and publication of the HEDIS measures or results ("rates") therefrom must be approved by NCQA and are subject to a license at the discretion of NCQA. Any use of the materials to identify records or calculate measure results, for example, requires a custom license and may necessitate certification pursuant to NCQA's Measure Certification Program.

HEDIS measures and specifications are not clinical guidelines, do not establish a standard of medical care and have not been tested for all potential applications. The measures and specifications are provided "as is" without warranty of any kind. NCQA makes no representations, warranties or endorsements about the quality of any product, test or protocol identified as numerator compliant or otherwise identified as meeting the requirements of a HEDIS measure or specification. NCQA also makes no representations, warranties or endorsements about the quality of any organization or clinician who uses or reports performance measures. NCQA has no liability to anyone who relies on HEDIS measures and specifications or data reflective of performance under such measures and specifications.

Limited proprietary coding is contained in the measure specifications for convenience. Users of the proprietary code sets should obtain all necessary licenses from the owners of these code sets. NCQA disclaims all liability for use or accuracy of any coding contained in the specifications.

CPT* codes, descriptions and other data are copyright 2024 American Medical Association (AMA). All rights reserved. CPT is a trademark of the AMA. Fee schedules, relative value units, conversion factors and/or related components are not assigned by the AMA, are not part of CPT, and the AMA is not recommending their use. The AMA does not directly or indirectly practice medicine or dispense medical services. The AMA assumes no liability for the data contained herein. Applicable FARS/DFARS restrictions apply to government use.

Health Care Provider Taxonomy Code Set codes copyright 2024 AMA. The codes are published in cooperation with the National Uniform Claim Committee (NUCC) by the AMA. Applicable FARS/DFARS restrictions apply.

The American Hospital Association holds a copyright to the Uniform Billing Codes ("UB") contained in the measure specifications. The UB Codes in the HEDIS specifications are included with the permission of the AHA. All uses of the UB Codes may require a license from the AHA. Specifically, anyone desiring to use the UB Codes in a commercial product to generate HEDIS results, or for any other commercial use, must obtain a commercial use license directly from the AHA. To inquire about licensing, contact ub04@aha.org.

The American Dental Association (ADA) holds a copyright to the Current Dental Terminology (CDT) codes contained in certain measure specifications. The CDT codes in the HEDIS specifications are included with the permission of the ADA. All uses of the CDT codes require a license from the ADA. No alteration, amendments, or modifications of the CDT or any portion thereof is allowed. Resale, transmission, or distribution of copies of the CDT or other portions of the CDT is also not allowed. To inquire about licensing, contact CDT-SNODENT@ada.org.

Some measure specifications contain coding from LOINC® (https://loinc.org/). The LOINC table, LOINC codes, LOINC panels and form file, LOINC linguistic variants file, LOINC/RSNA Radiology Playbook, and LOINC/IEEE Medical Device Code Mapping Table are copyright © 1995–2024 Regenstrief Institute, Inc. and the Logical Observation Identifiers Names and Codes (LOINC) Committee and are available at no cost under the license at https://loinc.org/kb/license/.

"SNOMED" and "SNOMED CT" are registered trademarks of the International Health Terminology Standards Development Organisation (IHTSDO).

The CDC Race and Ethnicity code system was developed by the U.S. Centers for Disease Control and Prevention (CDC). NCQA's use of the code system does not imply endorsement by the CDC of NCQA, or its products or services. The code system is otherwise available on the CDC website at no charge.

Certain NullFlavor codes are owned and copyrighted by Health Level Seven International (HL7°); 2024. "HL7" is a registered trademark of Health Level Seven International.

RadLex copyright 2014, The Radiological Society of North America (RSNA), all rights reserved. Licensed under RadLex License Version 2.0. You may obtain a copy of the license at: http://www.rsna.org/radlexdownloads/ This work is distributed under the above noted license on an "AS IS" basis, WITHOUT WARRANTIES OF ANY KIND, either express or implied. Please see the license for complete terms and conditions.

No part of this publication may be reproduced or transmitted in any form or by any means, electronic or mechanical, including photocopy, recording or any information storage and retrieval system, without the written permission of NCQA.

© 2024 by the National Committee for Quality Assurance 1100 13th Street NW, Third Floor, Washington, DC 20005

