New and Revised Requirements Related to Reducing Health Care Disparities

The Joint Commission has approved the following revisions for prepublication. While revised requirements are published in the semiannual updates to the print manuals (as well as in the online E-dition®), accredited organizations and paid subscribers can also view them in the monthly periodical The Joint Commission Perspectives®. To begin your subscription, call 800-746-6578 or visit http://www.jcrinc.com.

Please note: Where applicable, this report shows current standards and EPs first, with deleted language struck-through. Then, the revised requirement follows in bold text, with new language underlined.

APPLICABLE TO THE AMBULATORY HEALTH CARE ACCREDITATION PROGRAM

Effective January 1, 2023

Leadership (LD) Chapter

LD.04.03.08

Reducing health care disparities for the organization’s patients is a quality and safety priority.

Element(s) of Performance for LD.04.03.08

1. The organization designates an individual(s) to lead activities to reduce health care disparities for the organization’s patients.
   Note: Leading the organization’s activities to reduce health care disparities may be an individual’s primary role or part of a broader set of responsibilities.

2. The organization assesses the patient’s health-related social needs and provides information about community resources and support services.
   Note 1: Organizations determine which health-related social needs to include in the patient assessment. Examples of a patient’s health-related social needs may include the following:
   - Access to transportation
   - Difficulty paying for prescriptions or medical bills
   - Education and literacy
   - Food insecurity
   - Housing insecurity
   Note 2: Health-related social needs may be identified for a representative sample of the organization’s patients or for all the organization’s patients.

Key: □ indicates that documentation is required; ▪ indicates an identified risk area;
3. The organization identifies health care disparities in its patient population by stratifying quality and safety data using the sociodemographic characteristics of the organization’s patients. 
   Note 1: Organizations may focus on areas with known disparities identified in the scientific literature (for example, kidney disease, maternal care, diabetes management) or select measures that affect all patients (for example, experience of care and communication). 
   Note 2: Organizations determine which sociodemographic characteristics to use for stratification analyses. Examples of sociodemographic characteristics may include the following:
   - Age
   - Gender
   - Preferred language
   - Race and ethnicity

4. The organization develops a written action plan that describes how it will address at least one of the health care disparities identified in its patient population.

5. The organization acts when it does not achieve or sustain the goal(s) in its action plan to reduce health care disparities.

6. At least annually, the organization informs key stakeholders, including leaders, licensed practitioners, and staff, about its progress to reduce identified health care disparities.

Record of Care, Treatment, and Services (RC) Chapter

**RC.02.01.01**

The clinical record contains information that reflects the patient's care, treatment, or services.

**Element(s) of Performance for RC.02.01.01**

25. For organizations that elect The Joint Commission Primary Care Medical Home option: The clinical record contains the patient's:
   - Gender, race, and ethnicity
   - Family history
   - Work history (including any occupational risk factors or exposures)

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   - Gender
   - Family history
   - Work history (including any occupational risk factors or exposures)
31. The clinical record contains the patient’s race and ethnicity.

Rights and Responsibilities of the Individual (RI) Chapter

**RI.01.01.01**

The organization respects patient rights.

**Element(s) of Performance for RI.01.01.01**

29. The organization prohibits discrimination based on age, race, ethnicity, religion, culture, language, physical or mental disability, socioeconomic status, sex, sexual orientation, and gender identity or expression.