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review results of evaluation of compliance with individual accreditation standards, noted deficiencies, corrective action plans, and summaries of unmet accreditation requirements.

§ 438.364 External quality review results.

- (a) Information that must be produced. The State must ensure that the EQR produces at least the following information:
- (1) A detailed technical report that describes the manner in which the data from all activities conducted in accordance with §438.358 were aggregated and analyzed, and conclusions were drawn as to the quality, timeliness, and access to the care furnished by the MCO or PIHP. The report must also include the following for each activity conducted in accordance with §438.358:
 - (i) Objectives.
- (ii) Technical methods of data collection and analysis.
 - (iii) Description of data obtained.
 - (iv) Conclusions drawn from the data.
- (2) An assessment of each MCO's or PIHP's strengths and weaknesses with respect to the quality, timeliness, and access to health care services furnished to Medicaid recipients.
- (3) Recommendations for improving the quality of health care services furnished by each MCO or PIHP.
- (4) As the State determines, methodologically appropriate, comparative information about all MCOs and PIHPs.
- (5) An assessment of the degree to which each MCO or PIHP has addressed effectively the recommendations for quality improvement made by the EQRO during the previous year's EQR.
- (b) Availability of information. The State must provide copies of the information specified in paragraph (a) of this section, upon request, through print or electronic media, to interested parties such as participating health care providers, enrollees and potential enrollees of the MCO or PIHP, recipient advocacy groups, and members of the general public. The State must make this information available in alternative formats for persons with sensory impairments, when requested.
- (c) Safeguarding patient identity. The information released under paragraph

(b) of this section may not disclose the identity of any patient.

§ 438.370 Federal financial participation.

- (a) FFP at the 75 percent rate is available in expenditures for EQR (including the production of EQR results) and EQR-related activities set forth in \$438.358 conducted by EQROs and their subcontractors.
- (b) FFP at the 50 percent rate is available in expenditures for EQR-related activities conducted by any entity that does not qualify as an EQRO.

Subpart F—Grievance System

§ 438.400 Statutory basis and definitions.

- (a) Statutory basis. This subpart is based on sections 1902(a)(3), 1902(a)(4), and 1932(b)(4) of the Act.
- (1) Section 1902(a)(3) requires that a State plan provide an opportunity for a fair hearing to any person whose claim for assistance is denied or not acted upon promptly.
- (2) Section 1902(a)(4) requires that the State plan provide for methods of administration that the Secretary finds necessary for the proper and efficient operation of the plan.
- (3) Section 1932(b)(4) requires Medicaid managed care organizations to establish internal grievance procedures under which Medicaid enrollees, or providers acting on their behalf, may challenge the denial of coverage of, or payment for, medical assistance.
- (b) *Definitions*. As used in this subpart, the following terms have the indicated meanings:

Action means—

In the case of an MCO or PIHP—

- (1) The denial or limited authorization of a requested service, including the type or level of service;
- (2) The reduction, suspension, or termination of a previously authorized service;
- (3) The denial, in whole or in part, of payment for a service;
- (4) The failure to provide services in a timely manner, as defined by the State:
- (5) The failure of an MCO or PIHP to act within the timeframes provided in \$438.408(b); or