

DENTAL NETWORK VERSION 7.2

ORGANIZATIONAL STRUCTURE

CORE 1: Organizational Structure

CORE 2: Organization Documents

POLICIES AND PROCEDURES

CORE 3: Policy and Procedure Maintenance, Review and Approval

REGULATORY COMPLIANCE

CORE 4: Regulatory Compliance

INTER-DEPARTMENTAL COORDINATION

CORE 5: Inter-Departmental Coordination

OVERSIGHT OF DELEGATED FUNCTIONS

CORE 6-8: N/A

CORE 9: Delegation Management

MARKETING AND SALES COMMUNICATIONS

CORE 10: Review of Marketing and Sales Materials

BUSINESS RELATIONSHIPS

CORE 11: Written Business Agreements

CORE 12: Client Satisfaction

INFORMATION MANAGEMENT

CORE 13: Information Management

CORE 14: Business Continuity

CORE 15: Information Confidentiality and Security

CORE 16: Confidentiality of Individually-Identifiable Health Information

QUALITY MANAGEMENT

CORE 17: Quality Management Program

CORE 18: Quality Management Program Resources

CORE 19: Quality Management Program Requirements

CORE 20: Quality Management Committee

CORE 21: Quality Management Documentation

CORE 22: Quality Improvement Projects

CORE 23: Quality Improvement Project Requirements

CORE 24: Quality Improvement Projects: Consumer Organizations

STAFF QUALIFICATIONS

CORE 25: Job Descriptions

CORE 26: Staff Qualifications

STAFF MANAGEMENT

CORE 27: Staff Training Program

CORE 28: Staff Operational Tools and Support

CORE 29: Staff Assessment Program

CLINICAL STAFF CREDENTIALING & OVERSIGHT ROLE

CORE 30: Clinical Staff Credentialing

CORE 31: Senior Clinical Staff Requirements

CORE 32: Senior Clinical Staff Responsibilities

CORE 33: Financial Incentive Policy

CORE 34: Access to Services

CORE 35: Consumer Complaint Process

HEALTH CARE SYSTEM COORDINATION

CORE 36: Coordination with External Entities

CONSUMER PROTECTION AND EMPOWERMENT

CORE 37: Consumer Rights and Responsibilities
CORE 38: Consumer Safety Mechanism
CORE 39: Consumer Satisfaction
CORE 40: Health Literacy

NETWORK MANAGEMENT

DN-NM 1: Scope of Services
DN-NM 2: Provider Network Access and Availability
DN-NM 3: Provider Selection Criteria
DN-NM 4: Out-of-Network and Emergency Services
DN-NM 5: Participating Provider Representation
DN-NM 6: Participating Provider Relations Program
DN-NM 7: Participating Provider Written Agreements
DN-NM 8: Participating Provider Written Agreement Exclusions
DN-NM 9: Written Agreement Inclusions
DN-NM 10: Written Agreement Subcontracting
DN-NM 11: Other Participating Provider Agreement Documentation
DN-NM 12: Provider Network Disclosures
DN-NM 13: Participating Provider Violation Mechanism
DN-NM 14: General Requirements for Provider Dispute Resolution Mechanisms
DN-NM 15: Disputes Concerning Professional Competence or Conduct
DN-NM 16: Disputes Involving Administrative Matters
DN-NM 17: Participating Provider Suspension Mechanism for Consumer Safety

CREDENTIALING

DN-CR 1: Practitioner and Facility Credentialing
DN-CR 2: Credentialing Program Oversight
DN-CR 3: Credentialing Committee
DN-CR 4: Credentialing Program Plan
DN-CR 5: Credentialing Application
DN-CR 6: Credentialing Confidentiality
DN-CR 7: Review of Credentialing Information
DN-CR 8: Credentialing Communication Mechanisms
DN-CR 9: Primary Source Verification
DN-CR 10: Consumer Safety Credentialing Investigation
DN-CR 11: Credentialing Application Review
DN-CR 12: Credentialing Time Frame
DN-CR 13: Credentialing Determination Notification
DN-CR 14: Participating Provider Credentials Monitoring
DN-CR 15: Recredentialing
DN-CR 16: Recredentialing and Participating Provider Quality Monitoring
DN-CR 17: Credentialing Delegation



Re: New Regulatory Compliance Standard

Dear URAC Prospects and Clients,

As the nation's leading health care accreditor, URAC values patient care and safety first. When we discover an improvement opportunity that could enhance patient safety within our standards, we work quickly to incorporate the additional knowledge into our programs.

Therefore, URAC has introducing a new regulatory compliance Standard. Effective immediately, this new Standard will apply to all current, non-deemed applications, accreditations and certifications. Although we expect immediate compliance with this Standard, we will not be asking for any additional documentation at this time. As new versions of programs are released, this Standard will be built into the program requirements and documentation will be submitted on Desktop.

Attached you will find the Standard language (see Attachment A). As a requirement of this Standard, organizations must remain in good standing with any issuing body/agency for all permits, licenses, registrations and/or charters held by the organization. If at any time during the accreditation or certification cycle this Standard is identified as "Not Met," the finding(s) will be presented to the Accreditation Committee for review and final determination of status.

If you have any further questions, please reach out to Product Development Department at productdevelopment@urac.org.

Sincerely,

Jenn Richards, PharmD, JD, CSP

Product Development Principal

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Attachment A: URAC's New Regulatory Compliance Standard

Standard: Regulatory Compliance

The organization maintains compliance with applicable jurisdictional laws and regulations.

Regulatory Compliance

The organization:

- a. Maintains compliance with applicable laws, regulations and requirements from any relevant jurisdictions