URAC's IRO Accreditation: Demonstrating Your Excellence

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Before We Get Started







Message Lisa Silverman for any **tech issues** Use the chat box for questions and to introduce yourself

Explore resources we'll share in the chat box



Agenda for Today

Welcome and Introductions

About URAC

Updating Our Programs The Importance of Partnerships

Updated IRO Accreditation Standards

The IDR Designation

Questions and Answers



Opening Poll

What is your organization's relationship to URAC?



About URAC



Leader in Specialty Pharmacy, Telehealth and IRO/IME Accreditation

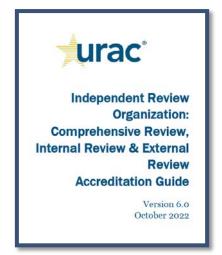
Our Clients Hospitals Health Plans Pharmacies Telehealth Providers Independent Review Population Health Utilization Management Credentials Verification Health Websites Case Management Disease Management





Client Relations Manager

People

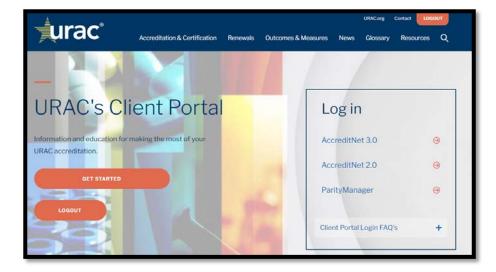


Program Guide



AccreditNet Guide





Client Information Hub





Accreditation Reviewer

ACCREDITATION PROCESS





Standards Revision

Process

current practice. Draft standards developed.

Two rounds of review with NAIRO to obtain input.

Worked closely with A&CS **IRO SMEs** to achieve a final standard set.

Presented to **Board of** Presented to **Directors** Health for **Standards** approval. **Committee** for

approval.

Program Release



urac

Research on industry

Identified

functions

of an IRO

and their

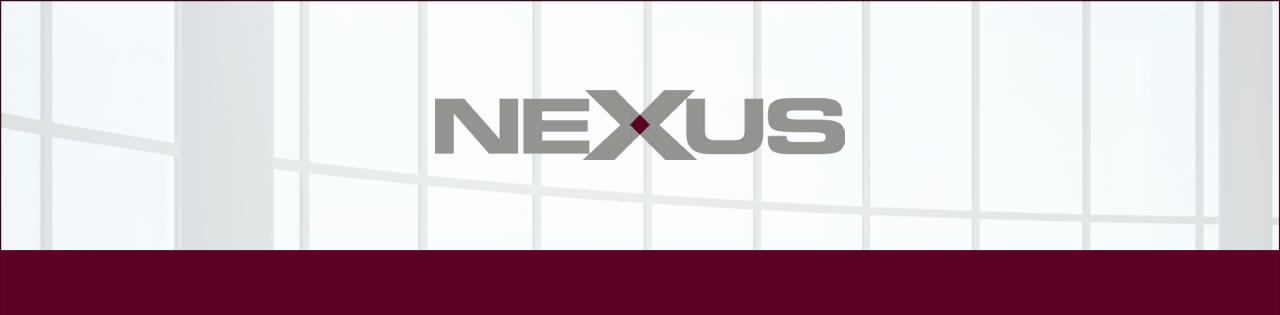
external

partners. Aligned

with **Focus** Areas.

main





URAC ACCREDITATION

THE PATHWAY TO SUCCESS



ACCREDITATION

Hello from Texas

Personal Experience to Professional Exposure

Framework for Excellence

The URAC/IRO Relationship – Synergistic not fearful or conflictual

NAIRO – Helping to protect the integrity of the clinical review business



ED BOLTON

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NEXUS

IRO Program 6.0 Accreditation Options

IRO Comprehensive (Internal & External Review)

IRO Internal

IRO External



Eligibility

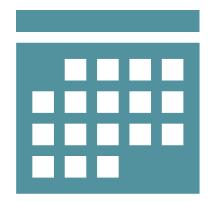
IRO-Comprehensive must provide both internal review, which include peer clinical review, internal appeals, and external review services.

IRO-Internal must provide internal review services, which include peer clinical review and internal appeals for health plans.

IRO-External must provide external review services, which are mandates by state and federal law.



Accreditation Timeline



Two Months

Desktop review

Four Months

Validation

review

Decision

Submit application



IRO v6.0 Revision Supports:

Moved to 2-4-8-G Scoring Methodology

Accreditation Guide includes information on scoring methodology

Standards in Six video series for understanding URAC's Foundational Focus Areas

Accreditation Timeline 6 months

Reduced # of required submissions by 25%

Single IRO:
Accreditation Guide
Standards Only
Standards-at- a- Glance

Rigor of Accreditation Process

Overall reduction in the number of Mandatory [M] standards:

Certain standards identified as "Not Met" require a follow-up validation review within six (6) months to verify full compliance.



Basic Revision Overview: IRO v6.0 Program Standards

Removed or Changed Concepts from v5.2:

Consistent Concepts:

New Concepts:

- Removed: Review Database (IR-TMR 1-1)
- Changed: Review of Additional Information (RP 2-1)
- While the requirements have been streamlined, the concepts remain highly similar due to the regulatory aspects of this program.
- During the program revision process, alterations in scoring were incorporated.
- Program Structure (IRO-PM 1-1)
- Summary Reports for Referring Entities (CMR-Int 1-1. v5.2 applied to External only)
- No delegation of the review rendered (COI 3-1 added to Reviewer Attestation)
- Inclusion on the List of Excluded Individuals/Entities (LEIE) maintained by the Office of Inspector General (OIG), if applicable. (CQP 2-1.)



Common Documents for Submission

Type of Evidence

What Your Organization Might Submit

Document(s)

Evidence

Sample

Job Description

Attestation

- Formal documentation (P/P, SOPs, Workflow Documents)
- What documents are you holding your employees accountable for following?
- Anything that demonstrates your organization is following the standard as specified in the language
- Meeting minutes are required when stated
- A sample document showing an example of how you do your work
- Only when specified
- We do not require job descriptions for all staff
- If the standard doesn't apply to your organization
- Risk Management



Common Language for Submission Documents

Outlining how

Typically, a mechanism, not a full process.

This is only one step of a normal process (who, what, when, where, why, how)

Outlining strategies

Exact methods for getting to a result

Describing a process

Full process (who, what, when, where, why, how)

Outlining the organization's requirement

Policy statement



URAC's New Independent Dispute Resolution Designation: Eligibility

- Must be located and operating in the U.S.
- Must have U.S. tax ID
- Must be U.S. corporation in good standing
- Must have license requirements in good standing

- Program specific eligibility:
- URAC IRO Accredited:
 - IRO Comprehensive or
 - IRO External

or

- Concurrently seeking URAC IRO Accreditation:
 - IRO Comprehensive or
 - IRO External



Independent Dispute Resolution Designation Focus Areas

Program Management Dispute Resolution Principles

Conflict of Interest

Professional Qualifications

Dispute Resolution Processes

Information Inclusions

Breach Notification

Administration



Where Can I Find Things?

Website

- Standards-at-a-Glance
- Standards for Purchase
- Are You Ready?
- Program Overviews
- Standards Interpretation Portal

Client Information Hub

- AccreditNet User Guide
- FAQs
- On-Demand Standards Education
- Link to Request Program Guide and Crosswalk

AccreditNet

- Program Guide
- Program Crosswalk
- Glossary
- AccreditNet User Guide



Questions?

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