May 8, 2015

NQRN[®] National Quality Registry Network

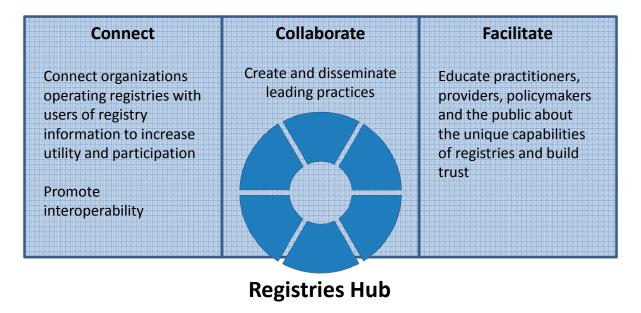
CMSS ITG: NQRN Update and Discussion

Seth Blumenthal, MBA | NQRN Program Manager American Medical Association

This presentation is a general overview of NQRN[®]. For more information, please visit **nqrn.org**.

NQRN: A Learning Hub for Registry Stewards

The National Quality Registry Network (NQRN[®]) is a voluntary network of organizations operating registries and others interested in increasing the usefulness of clinical registries to measure and improve patient health outcomes.



NQRN Steering Committee

Co-chairs: David Shahian, MD, Lewis Sandy, MD

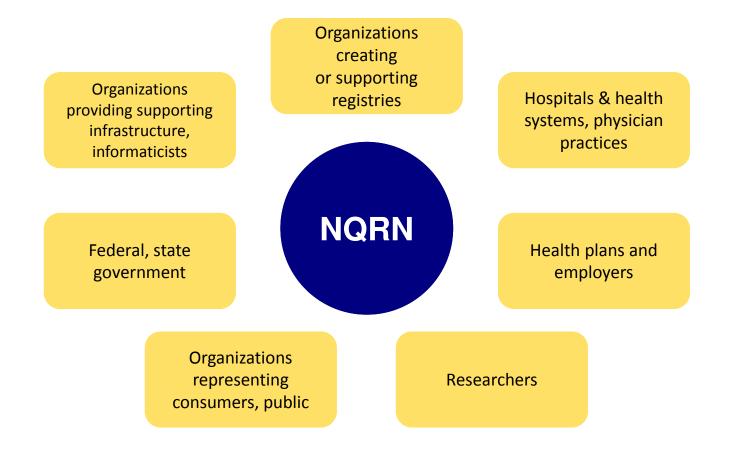
Representative	Organization
Kevin Bozic, MD, MBA	California Joint Replacement Registry
Richard Dutton, MD, MBA	American Society of Anesthesiologists
Kathleen Hewitt, RN, MSN	American College of Cardiology
Tim Jackson	Centers for Medicare & Medicaid Services
Mike Kappell, MBA	National Coalition for Cancer Survivorship
Art Levin, MPH	Center for Medical Consumers
Frank Opelka, MD	Physician Consortium for Performance Improvement
Dana Gelb Safran, ScD	Blue Cross Blue Shield of Massachusetts
Lewis Sandy, MD	United Health Group
John Santa, MD, MPH	Consumer Reports
David Shahian, MD	Society of Thoracic Surgeons
Modena Wilson, MD	American Medical Association

AMA professional staff: Kathleen Blake, MD, MPH, Dana Richardson, RN, MHA, Seth Blumenthal, MBA

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NQRN Council

More than 80 organizations participate in the NQRN Council





NQRN Learning Activities

Leading Practices Webinars

2015 to date:



- The Swedish registries
- Registry of Patient Registries (RoPR)
- Watch for announcements on the next webinars in the series!

2014:

- Vendor selection
- Data security
- Participant recruitment
- Device surveillance

2013:

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- Quality Oncology Practice Initiative (QOPI)
- National Cardiology Data Registry (NCDR)

National Quality Registry Network

Conferences

2015 annual meeting:

- Uses of registry information including QI collaboratives and value-based payment
- Measure development
- Public reporting

2014 annual meeting

- HIPAA and the Common Rule grey area for registries
- Electronic quality measures (eCQMs) accelerating the ability of registries to improve outcomes
- Early learnings from CMS Qualified Clinical
 Data Registries

2013 meeting: Maturational Framework

• Advance a clinical registry maturational framework to evaluate registry capabilities

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Knowle	edge Center	NQRN overview 2 NQRN coordinating task force recommendation 2011 2 National Quality Registry Network					rk				
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Conne	ct with the PCPI	The NQRN Steering Committee consists of 12 members, representing medical (NQRN®) provides the									
Nation Netwo	al Quality Registry rk	professionals, registry stewards, experts in registry technology, payers, and consumer/patient users. The Steering Committee provides strategic direction NQRN by placing a clickable NQR						the			
Wet	pinars	and oversight for NQRN activities. — If you wish to participat					participate in t	his			
Mee	tings	The NQRN Council is composed of members, plus Federal Government program, please						se <u>contact us</u> .			
Join	the NQRN	liaisons, who represent registry stewards and participants, non delivery system registry users, and supporting technologies. <u>NORN C</u>					NQRN Collabo	ration Portal			
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Learning Resources:

- NQRN meeting presentations
- Webinar recordings
- Registry inventory
- QCDR guide
- "What is a registry?"
- Maturational framework public version
- Registry FAQ
- Registry vendor assessment
- Additional information about NQRN

Collaboration Portal:

Ask questions and get answers on registry related topics. Collaborate with your colleagues on new NQRN resources!

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2015 – 2017 Strategic Goals

Connect, Collaborate, Facilitate

Goal	Objectives
Goal 1: Become a hub for creating and disseminating leading practices and guidance to the registry enterprise	 Objective A: Resources to assist registry organizations Objective B: Measure how well NQRN meets participants' needs Objective C: Help participants network, and share leading practices
Goal 2: Encourage active participation in registries by physicians, other health care professionals, hospitals, health systems and patients	 Objective A: Promote registries to external audiences Objective B: Support the QCDR program
Goal 3: Support the increased use of registries by health plans, employers, patients, consumers, federal and state government by increasing their value for these stakeholders	 Objective A: Support public reporting using Objective B: Assist organizations operating or planning registries in evaluating registry maturity
Goal 4: Advocate for a supportive regulatory environment and interoperability	 Objective A: Assist registries in supporting multiple uses while protecting privacy through compliance Objective B: Promote interoperability
Goal 5: Educate policymakers and the public about the unique capabilities of clinical registries	 Objective A: Federal government representatives participate in NQRN committees and meetings Objective B: Create or disseminate information about registries reaching a broad audience
Goal 6: Secure external funding to achieve the goals of the NQRN and ensure long term sustainability	 Objective A: Create a sustainable business plan Objective B: Seek grant and sponsorship opportunities

Join the NQRN!

Contact:

Seth Blumenthal American Medical Association 330 N. Wabash Ave., Suite 39300 Chicago, IL 60611

(312) 464-4049 seth.blumenthal@ama-assn.org

Visit: <u>nqrn.org</u>

