

---

# ***Marrying Technology to the Chronic Care Model***

**Neil A. Solomon, MD**

President, NAS Consulting Services

Faculty Director, Breakthroughs in Chronic Care Program

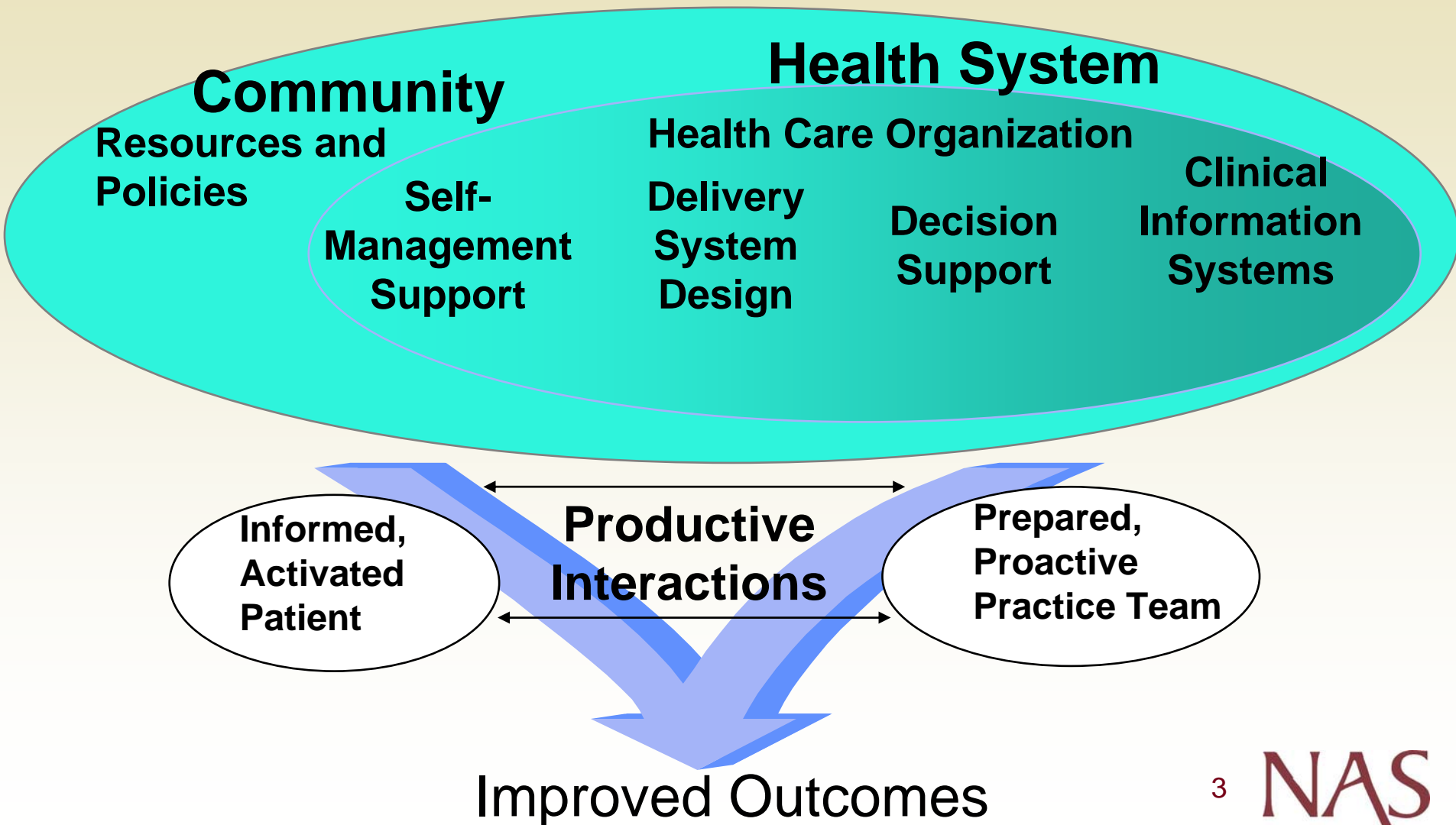
415-836-6777

August 22, 2006

# The Quality Chasm



# What Do We Do With the CCM?

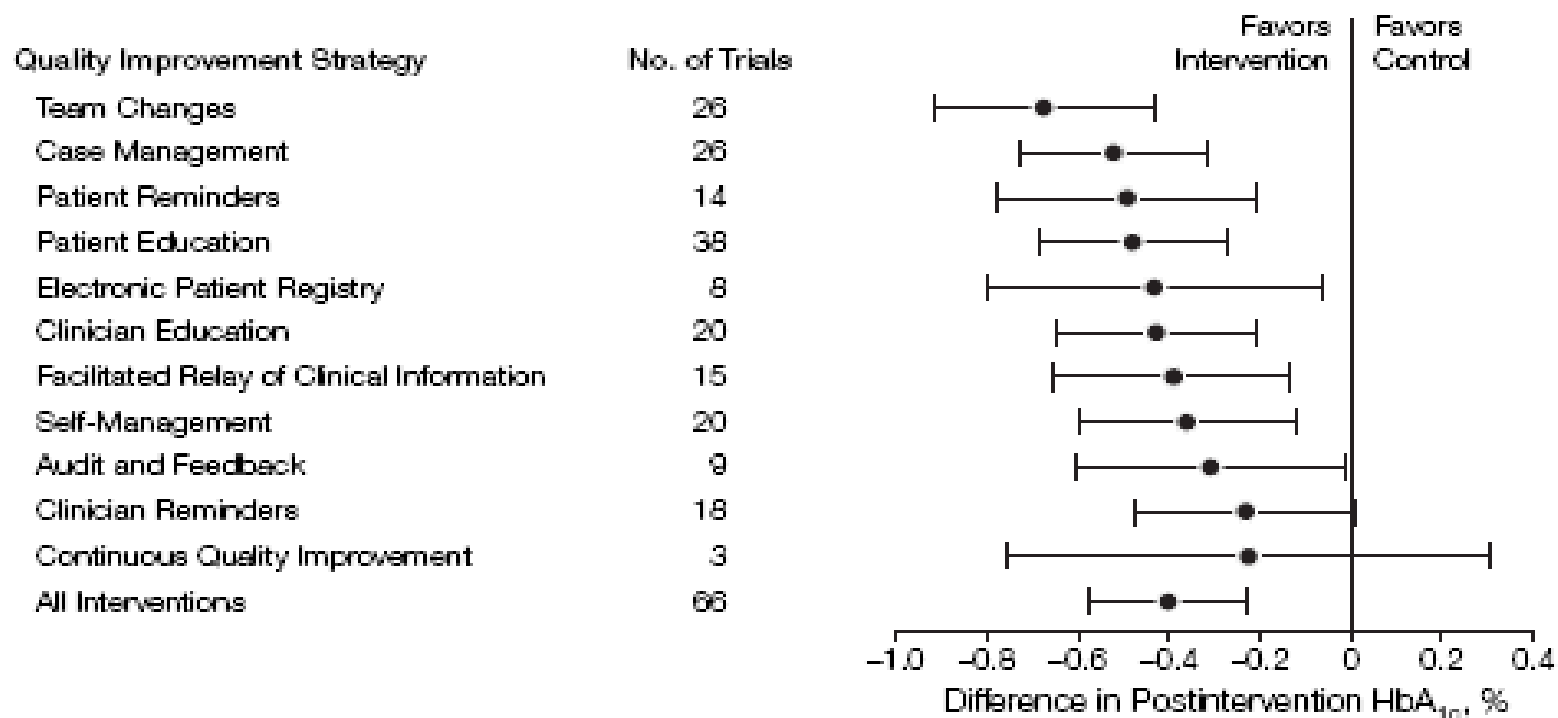


# Linking QI Process to Outcomes

---

- Shojania, et.al., JAMA July 26, 2006
- Meta-regression analysis for HBA1c control
- 66 eligible controlled trials, mostly RCTs
- Considered 11 QI categories: case mgmt, team care, registry, CME, clinician reminders, facilitated relay of clinical info, patient ed, self-mgmt support, patient reminders, CQI

**Figure 2.** Postintervention Differences in Serum HbA<sub>1c</sub> Values After Adjustment for Study Bias and Baseline HbA<sub>1c</sub> Values



Negative estimates favor intervention groups over control groups for the indicated quality improvement (QI) strategy. These estimates were derived from a meta-regression model with adjustment for the effects of study size (at least as many patients as the median among all included studies vs not) and a dichotomous variable that equaled 1 if the mean baseline glycosylated hemoglobin (HbA<sub>1c</sub>) value was  $\geq 8.0\%$ . Thus, the point estimate of  $-0.67\%$  for team changes indicates that, given a large study (ie, having at least as many participants as the median of 180 patients) in which the mean baseline HbA<sub>1c</sub> values were  $8.0\%$  or higher, the intervention group would have a follow-up HbA<sub>1c</sub> value  $0.67\%$  lower than the follow-up value in the control group.

# What Roles For Technology?

---

- Facilitate care teams
- Case management support
- Home-based care assistance
- Links between patient and clinicians
- Physician reporting and feedback
- Clinician prompts and reminders

# What Do the Tools Look Like?

---

- Registries
- Electronic Health Records
- Personal Health Records
  - ◆ Putting the control into the hands of the patients
- Web-based communication tools



# What is a Registry?

---

- An electronic tool that captures and tracks data for a patient population with a particular disease or health state.
- Enables population-based care for entire group of patients, not just those that elect to come into the office.

# Chronic Disease Registries

---

- Identify, stratify and track populations
- Interconnect members of the care team
- Find patients “falling through the cracks”
- Provide point-of-care support
- Help the organization prioritize

# Starting a Registry

---

## Key considerations:

1. Product selection: Buy vs. build
2. Data issues
3. Provider involvement

# Registry Options

---

- Home grown—built off claims, labs, rx
- Vendor product
  - ◆ Public domain software (Access database)
  - ◆ Commercial software on client hardware
  - ◆ Data supplier delivers regular reports
- ASP Model
  - ◆ Runs remotely; MDs and managers access over secure internet connection

# Buy vs. Build

- Decision depends on organizational resources, characteristics
- Buy:
  - + Speed to implementation
  - + Less internal staff needed to maintain
  - + Some products can track multiple conditions
  - ◆ Reliance on external vendor for customization
    - New features
    - Integration with EHR or other local needs
- Build:
  - + Retain control over functionality and data
  - + Can customize to meet needs
  - ◆ Requires knowledge in database management, IT understanding of chronic care management

# Data Quality

---

- Data accuracy important to success
  - ◆ Relevant recommendations, comparisons
  - ◆ Credibility with physicians, patients
- Consider implications of errors
  - ◆ Cohort: false positives, false negatives
  - ◆ Interventions/measures
- Timeliness of data refresh
- On-going quality control, maintenance

# Registry in Action

---

- NP reviews report of all patients overdue for a test or out of control
  - ◆ Calls highest risk patients to check in and schedule visit or tests
  - ◆ Generates letters to patients mildly out of adherence
- MA prints snapshot of patient's status and clips to chart for MD seeing patient
  - ◆ May show most recently lab values, when meds last filled, any recent hospital admissions, etc.
  - ◆ May provide prompts if patient overdue for evals



# Electronic Health Records

---

- Acquire and assemble data – lab, radiology, etc.
- Connect to colleagues – team care, consultants
- Provide clinician reminders/decision support
  - ◆ Identify patients overdue for routine care – prompts
  - ◆ Drug-drug; drug-dx; drug-lab; dose checking
  - ◆ Tagged literature to support decision-making
- Support clearer and fuller documentation
- All patients, all parts of their care

# EHR in Action

---

- Delivery system EHR allows case manager to review MD notes and manage patient with primary physician
- Chronic disease templates allow rapid documentation of key findings
- Embedded decision support prompts and reminds clinician during encounter

# EHR vs. Registry for Chronic Disease

---

## ➤ EHR positives

- ◆ Useful for all patients, all data
- ◆ Complete visit documentation
- ◆ Strong R&D and support (major vendors)

## ➤ Registry positives

- ◆ Population based
- ◆ Easier to stratify, target, track patients
- ◆ Less expensive, easier to implement

# EHR vs. Registry for Chronic Disease

---

## ➤ EHR negatives

- ◆ Handle one patient, one problem at a time
- ◆ Weak population management functions, little or no care/case management
- ◆ Expensive, hard to implement and maintain, usually takes a long time to get to chronic dz fxns

## ➤ Registry negatives

- ◆ Limited data—patients, and clinical info for them
- ◆ Can't document for entire clinical note
- ◆ Limited R&D to expand capabilities
- ◆ Weak implementation support



# Personal Health Records

---

- Same data as EHR, different presentation format to non-MD
- Some PHRs allow for patient data entry (e.g. home BP readings)
- Good tools pre-populate all key fields
- Potential to motivate and provide self-management support for patients
- Can dramatically improve office and organizational efficiency
- Still in infancy

# Patient Communication Tools

---

- Web-based secure e-mail or similar
- Focused reminders to patients (outbound)
- Succinct updates and questions/ concerns from patients (inbound)
- Referrals to sources of “information therapy”

# Types of Secure Communication

---

- Stand-alone programs—e.g. Relay Health, Medem
- Provided by delivery system
- Integrated into EHR

# Common Concerns

---

- HIPAA, privacy, confidentiality
- Lack of reimbursement
- Malpractice fears
- Time sink

# Recap

---

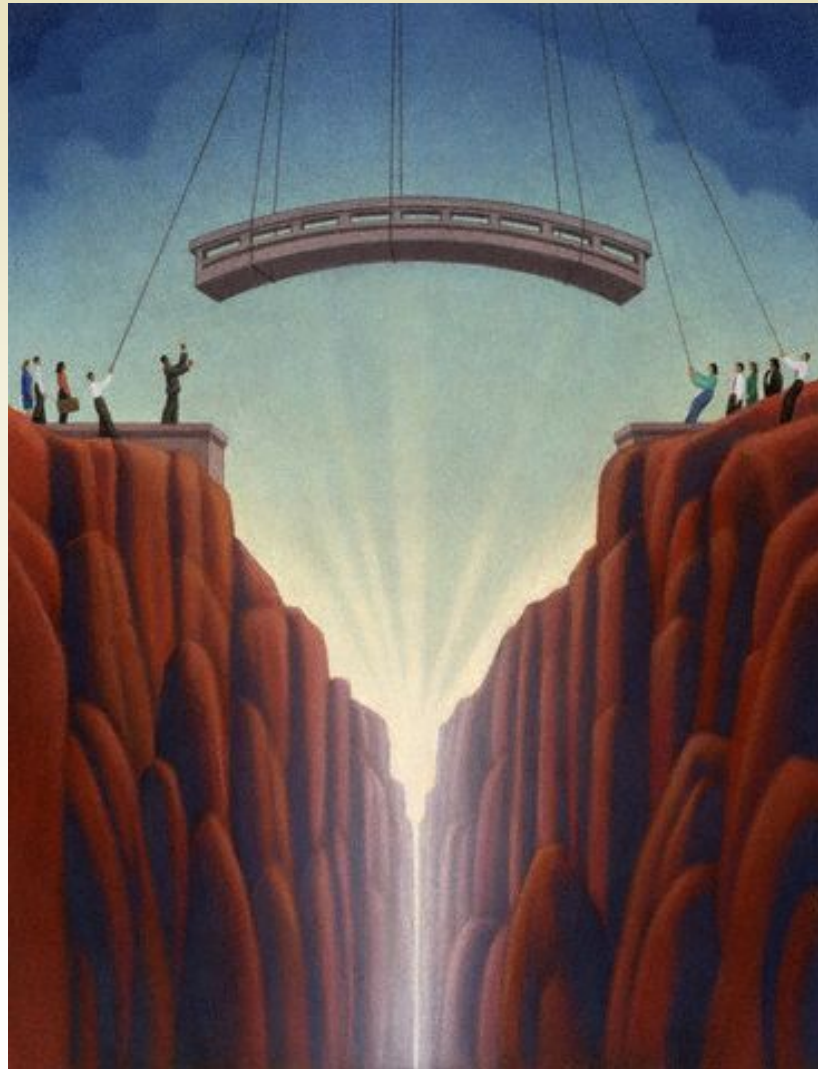
## ➤ Top QI changes

- ◆ Team care, case management, patient reminders and patient education

## ➤ Key tools

- ◆ Registry, EHR, PHR, secure communications
- ◆ All are evolving, some interrelate
- ◆ Implementation more important than what you choose to use

# Its All About the Team



# References

---

- IT Tools for Chronic Disease Management: How do they Measure Up? Jantos and Holmes, CHCF, 2006
- Effects of Quality Improvement Strategies for Type 2 Diabetes on Glycemic Control. Shojania, et.al., JAMA July 26, 2006
- Improving Chronic Illness Care web site: [improvingchroniccare.org](http://improvingchroniccare.org)

