

How To Implement Credible QI Projects.....

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- Without large grants
- By leveraging interprofessional knowledge and skills
- In the conduct of routine work



My Personal Take on the “Science of Improvement”

- **Scientific** regardless of name:
 - Science of improvement
 - Health care delivery science
 - Implementation science
 - Systems strengthening
 - Systems engineering
- Scientific **methods** include
 - “Model for improvement” promulgated by IHI
 - Lean
 - Six Sigma
 - Lean Six Sigma



“Elevator Speech” on Key Attributes of Improvement Science (Model for Improvement Methodology)

- Clear, **measurable aim**
- A **measurement framework** in support of reaching the aim
- Clear description of the **ideas** (content) and how these ideas are expected to impact **results** (the **causal pathway** from changes to desired outcomes, and their attributable effect)
 - Conceptual or logic model, or “driver diagram”
- Clear description of the **implementation** strategy
 - What will be done to ensure reliable adoption of the content
- Dedication to rapid testing (PDSA) - **prediction and learning** from tests
- Understanding/describing/visualizing **systems** (process map, value stream)
- Learning from **variation** and **heterogeneity**
 - Use of time-ordered data to detect special cause and improvement
 - Understanding why results differ by ward, organization, region
- Application of **behavioral and social sciences**



Why Clinicians (Especially Physicians*) Are Skeptical About QI

- Many associate QI with old-style, punitive QA (profiling, pay-for-performance)
- QI gurus overemphasize the industrial quasi-”religious” origins of QI and use unfamiliar jargon
- QI experts tend to focus on non-clinical processes and outcomes rather than clinical outcomes of interest to clinicians
- Teams try to do QI “by the book” and get bogged down in tedious process and settle for small incremental improvements
- QI leaders are not up front about the fiscal agenda (“QI is free”)
- QI programs do not provide clinicians with the data they need to improve
- QI experts do not emphasize the academic potential of QI research

* I emphasize physicians because like it or not, improvement in clinical care requires that they be engaged



Why Physicians who Understand Laboratory Science Should be Comfortable with Rigorous QI

- My ten years working with a PhD scientist to develop a Staphylococcus vaccine...
- The “experimental method,” mice, theories/hypotheses, and PDSAs
 - While the “context” of lab work is far less complex than the “context” of the hospital or clinic, behavioral and contextual issues do come into play
- And why don't QI leaders keep a “lab book” to document exactly what they are doing and learning?



What to Do

- Leaders must not lose touch with work at the front line
 - Understand that clinicians have limited **capacity (time and energy)** to take on new QI projects and little **capability (skills)** in QI methodologies
 - A 'work smarter, not harder' approach to improving healthcare quality. Hayes CW, Batalden PB, Goldmann D;BMJ Qual Saf. 2015;24:100-2*
 - Clinician “burn out” is increasing, and generally additional time and payment are not provided for QI responsibilities
- Find and address the pain points – “what frustrates you the most?”
 - A personal story – Visits to radiology and the emergency department



What to Do

- Teamwork is essential
 - QI is inherently inter-professional, yet clinicians generally are not trained to work in teams, nurse-physician tensions remain, and work patterns and schedules are hard to synchronize
 - A gulf remains between clinicians at the bedside and critical support personnel, such as pharmacists, physical therapists, and social workers
- Imbed QI, including **data collection**, in real work – not separate and added on
 - Example: Frustration with physician “maintenance of certification” requirements related to QI in the US



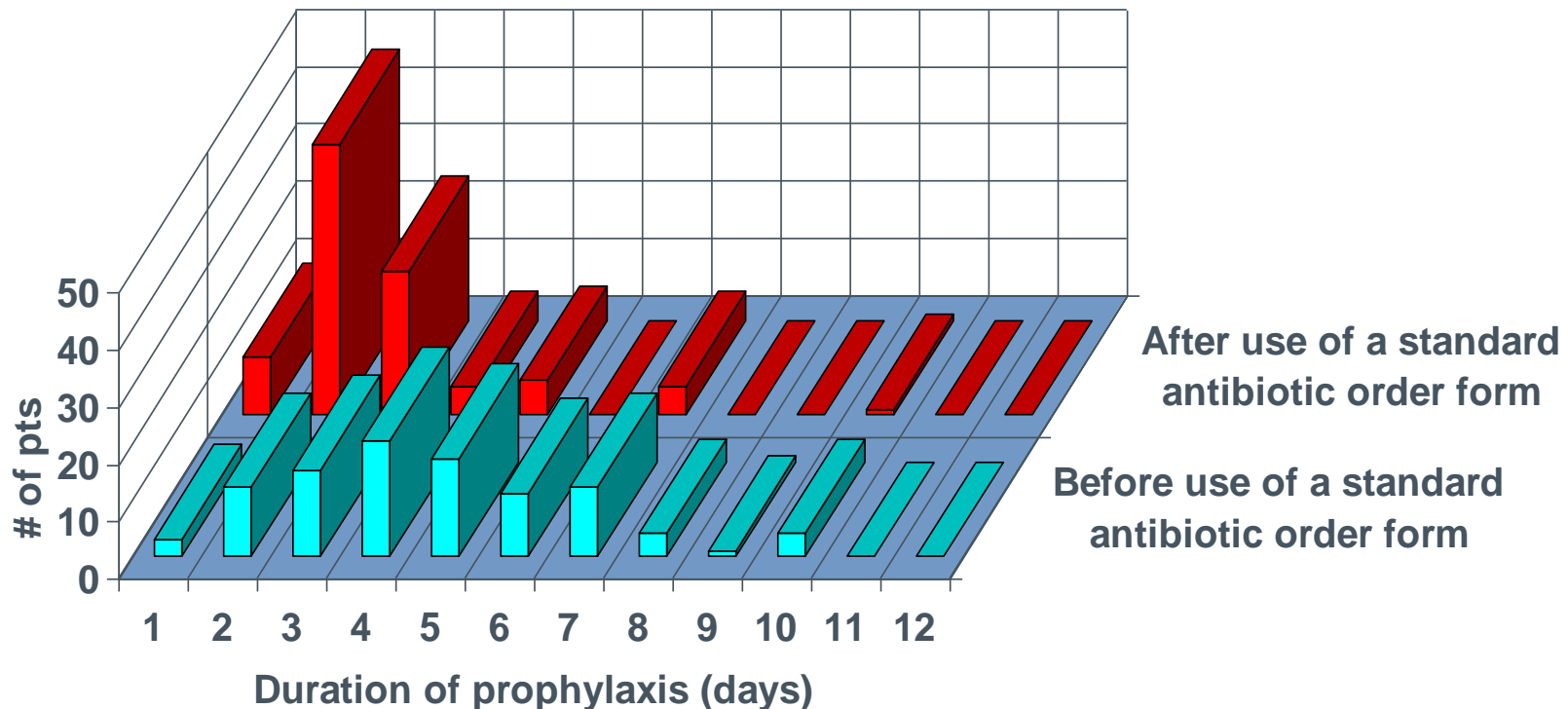
Rigorous, Even Publishable QI Is
Possible Almost Anywhere –
Without Grants (!)



Personal Experience



Baby Steps: Effect of Standard Antibiotic Order Form on Duration of Prophylaxis



Durbin et al. JAMA 1981;246:1796

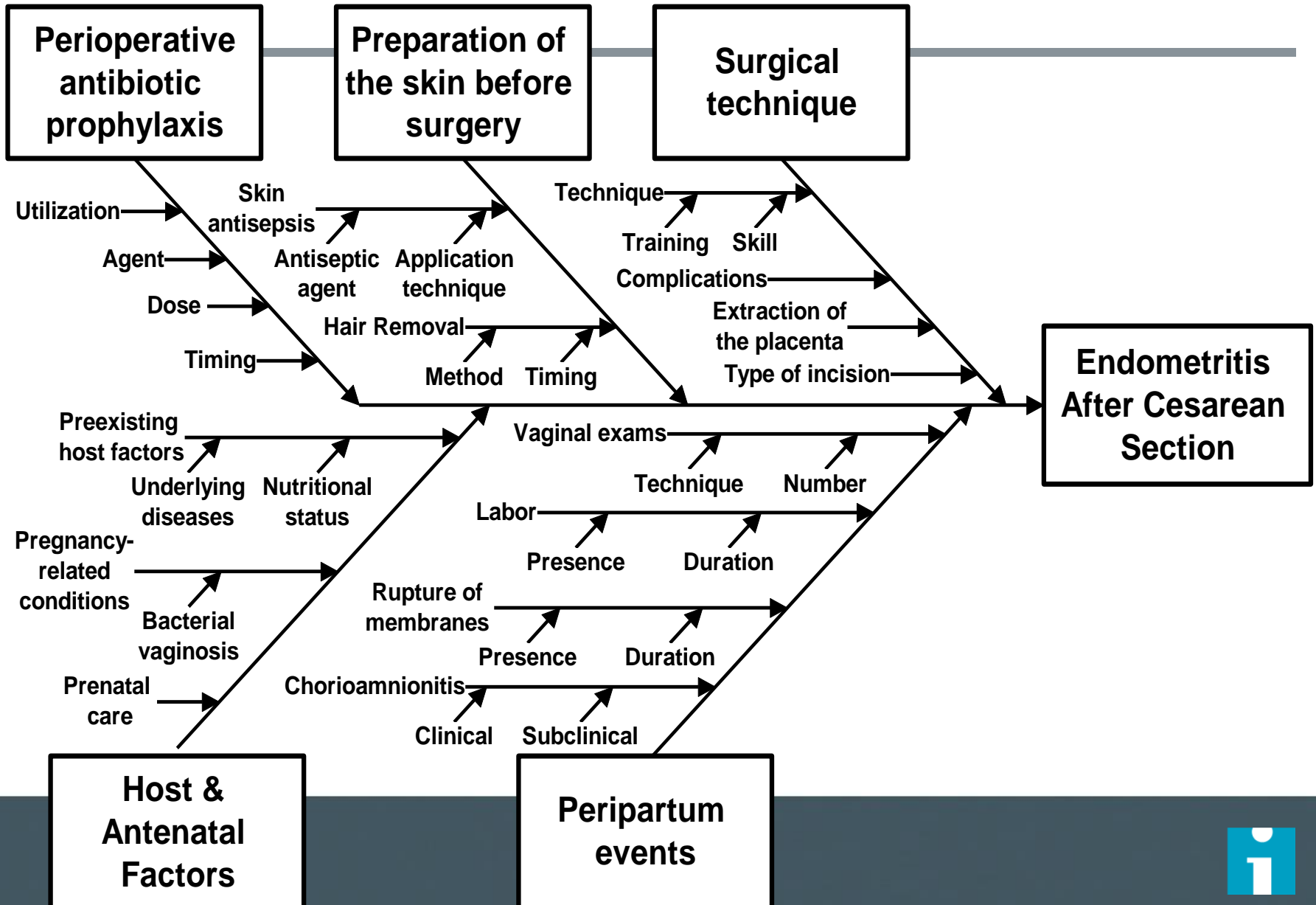


If They Can Do It in Bogotá during
Civil Conflict with Constrained
Resources...

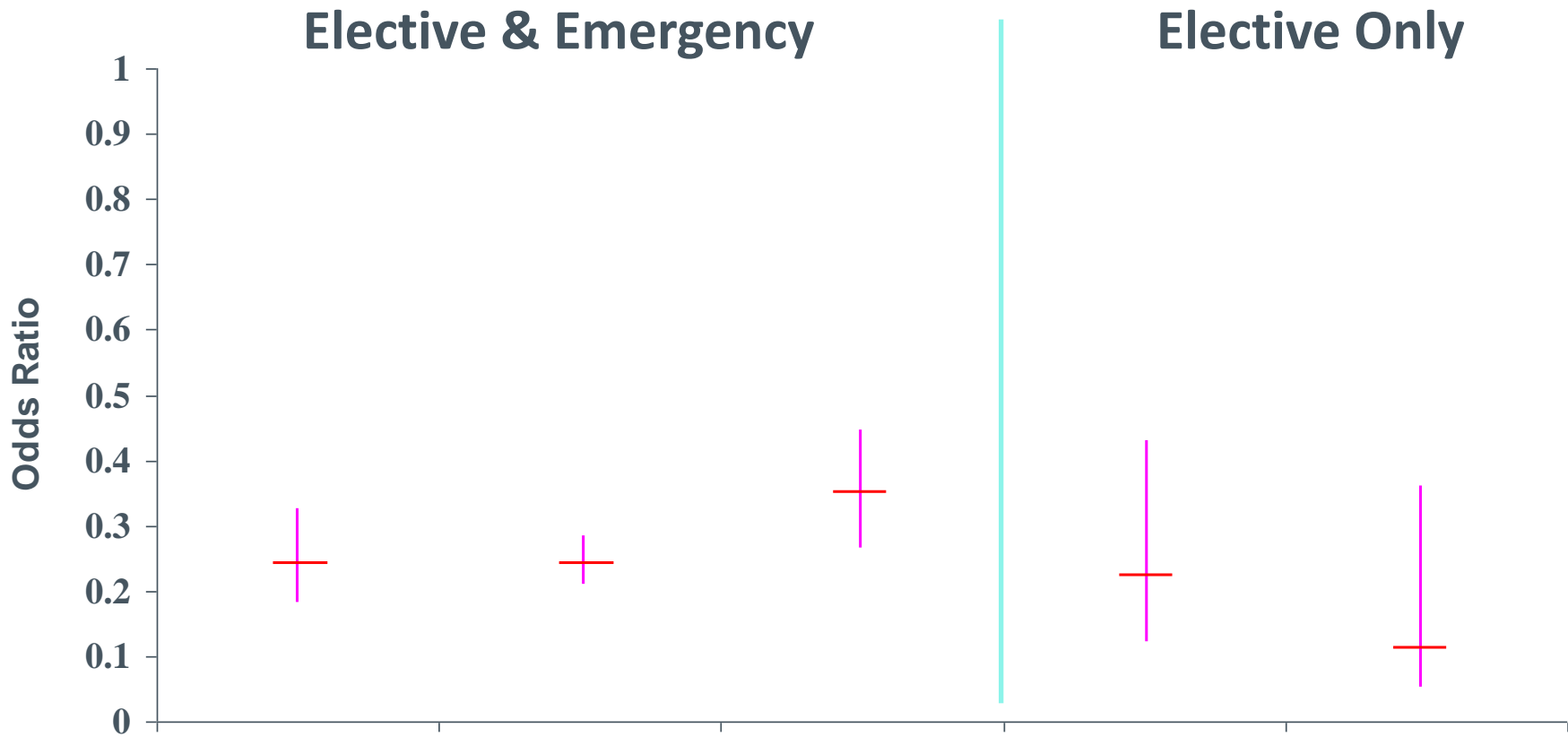
Reducing Post-Caesarian Infections



Cause and Effect Diagram



Meta-Analysis the Effect of Antibiotic Prophylaxis on Infection Rates after Cesarean Section



Priority Matrix

| Factor | Importance | Within the capacity of hospital personnel to improve | Timeframe for improvement |
|-------------------------------|-------------------|---|----------------------------------|
| Antibiotic prophylaxis | 4 | 4 | short |
| Skin preparation | 3 | 4 | short |
| Surgical technique | 4 | 4 | medium |
| Antenatal factors | 3 | 1 | long |
| Peripartum events | 4 | 2 | medium |

Utilization and Timing of Antibiotic Prophylaxis for Cesarean Section

| | % receiving prophylaxis | % receiving prophylaxis ≤ 1 hour after delivery |
|--|--------------------------------|--|
|--|--------------------------------|--|

Hospital A

70%

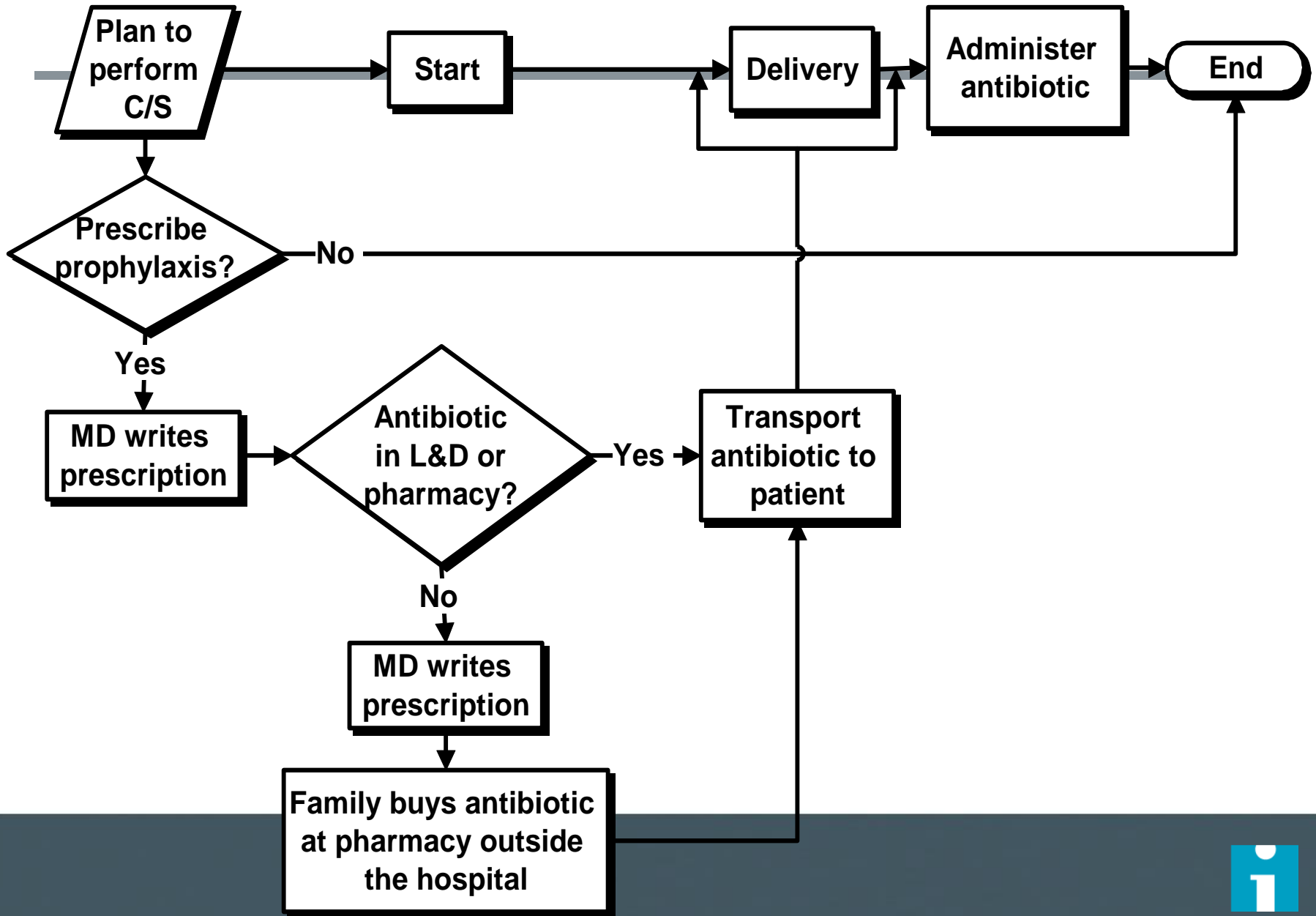
31%

Hospital B

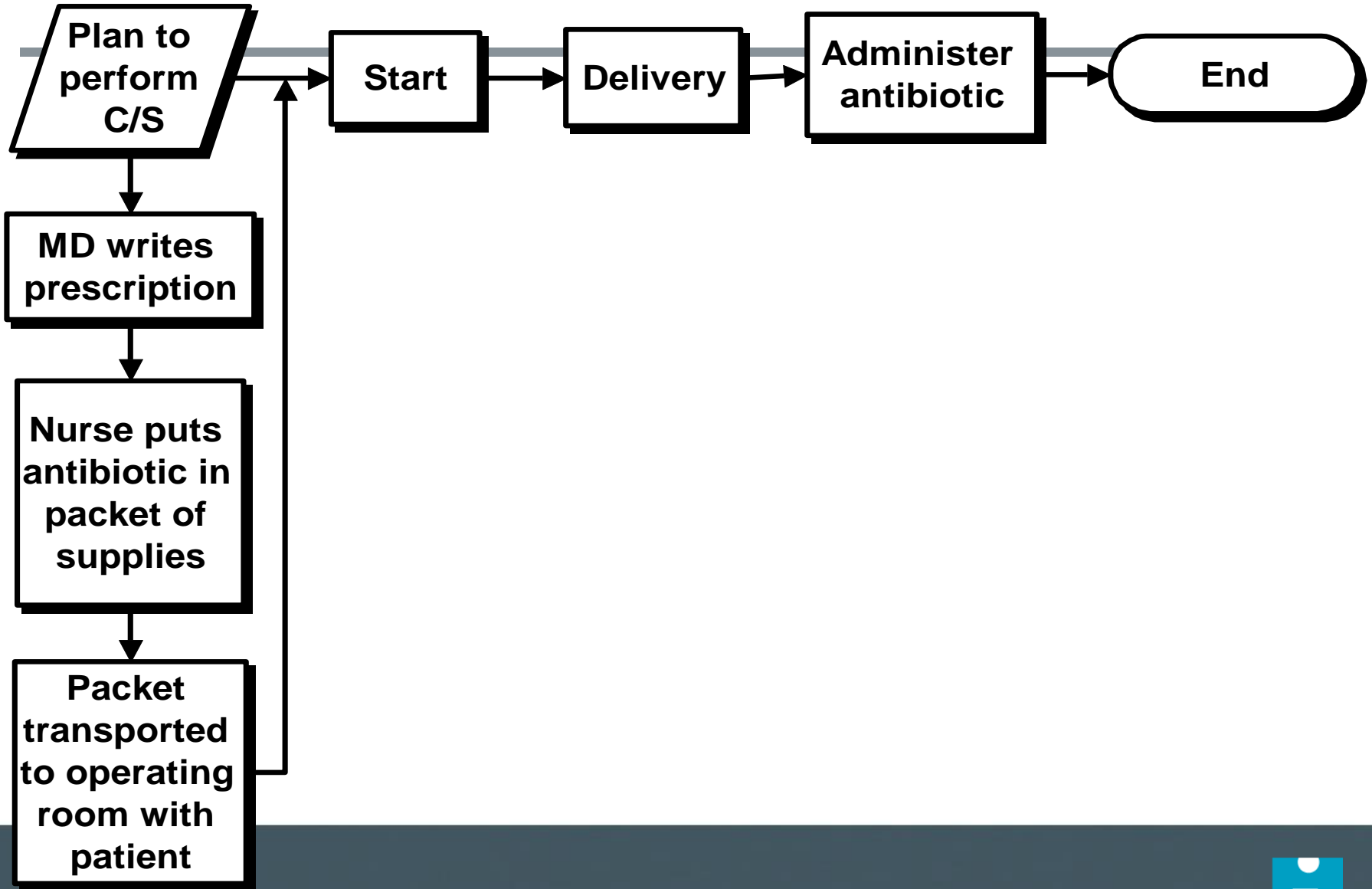
32%

70%

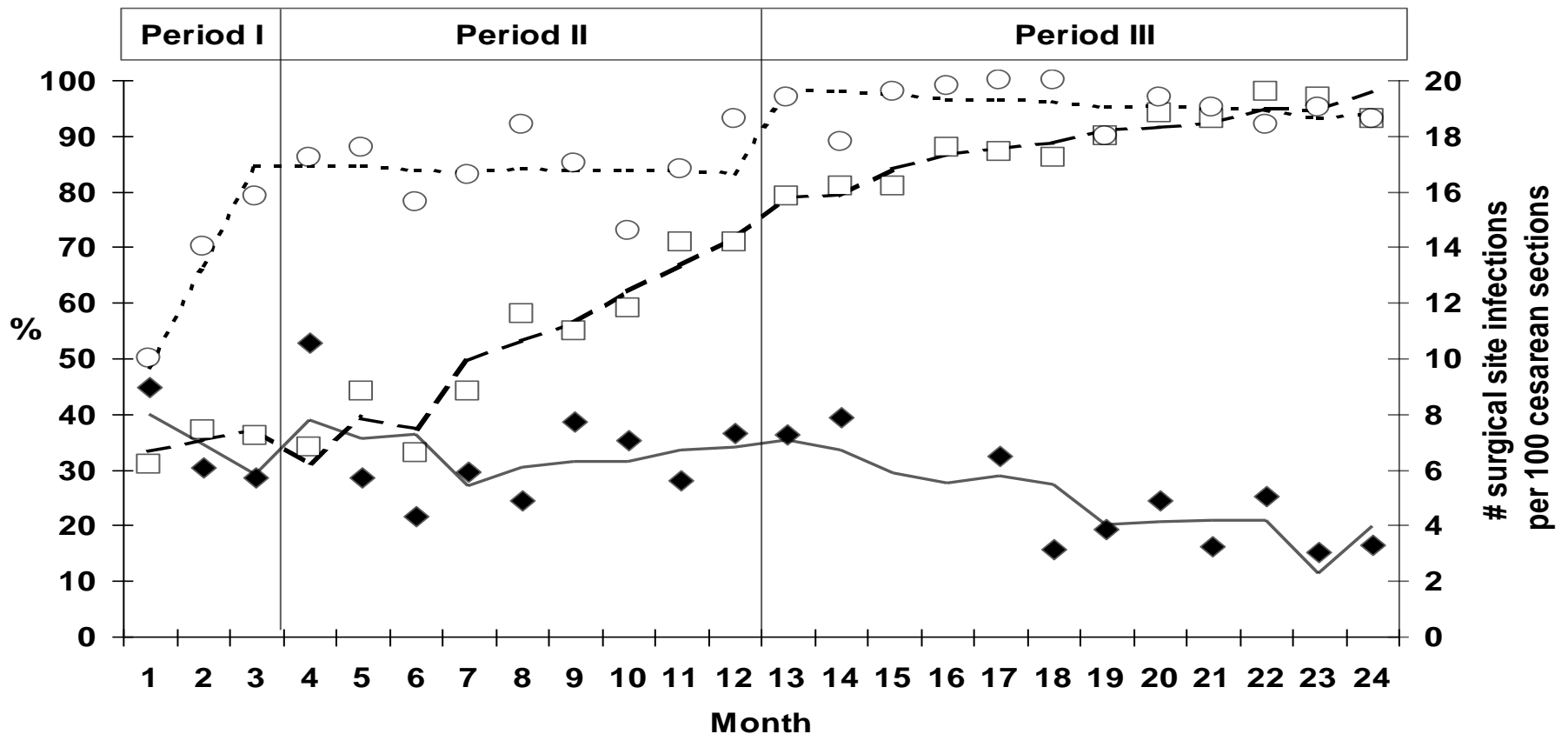
Hospital A: Existing System



Hospital A: Revised System



Utilization and Timing of Perioperative Antibiotic Prophylaxis & Surgical Site Infections After Cesarean Section



Experiential Learning – Making Rigorous QI Part of Routine Work at the Point of Care



Monitoring Patient Safety

- Voluntary event reporting
- Morbidity and mortality conferences/reports
- Chart auditing
 - IHI Global Trigger Tool
- Automated data mining
 - Patient Safety Indicators (AHRQ PSIs)
 - Automated trigger tools
- **Random Safety Audit**



Random Safety Audit

- Translated from industry (banking and random process audits *via* Paul Plesk)
- Real time by the front line
- Data and feedback virtually immediate
 - Reliability of key safety processes evident immediately
 - Motivating, enabling, reinforcing; builds self-efficacy and social norms (key elements of behavioral change theory)
- Combines audit and feedback with iterative PDSAs
 - Even better than “what can I try by next Tuesday”



Random Safety Audit

- Systematically monitors a subset of error-prone points in the system that have the potential to harm patients
- Items selected randomly to be addressed either on
 - On multi-disciplinary rounds (provider input required)
 - Any time during day (provider input not needed)
- Deck can be “packed”
- 20 items developed by expert consensus for testing in NICU (21st item added later)
- 4X6 “cards” include yes/no data form; trivia question on back



Staff Perceptions of the Random Safety Audit

- 84% of staff participated in rounds on which audit performed
- 100% agreed or strongly agreed that this improved quality and safety
- 95% agreed/strongly agreed that it increased knowledge of clinical guidelines and safety goals
- Only 9% agree with statement “asking a safety question of rounds took up too much time”



Three Simple Examples of Interprofessional QI Involving Junior Doctors

- Do you know who your doctor is?
- Understanding drug usage and reducing unnecessary prescriptions
- Learning how to look for medical errors as part of routine work



Try Something Yourself

- At work – “You are always late for rounds!”
- At home – avoiding the weed whacker
(<http://www.ihl.org/education/IHIOpenSchool/resources/Pages/Activities/PDSACyclesFromCLABSIsToCucumbers.aspx>)



Ten Tips for Incorporating Rigorous Quality Improvement into Everyday Work

BMJ Qual Saf. 2011 Apr;20 Suppl 1:i69-72



Tip 1

- Select projects that really will make a difference to providers and patients
 - Focus on clinically relevant projects that substantially improve those processes of care that are tightly linked to the outcomes of interest to providers and patients
 - Think of a headline the CEO or CMO would want to feature on the organization's website



Tip 2

- Set bold, clear, measureable aims and a specific timeline for achieving them
 - Think of fundamental advances that will measurably impact care and outcomes and engage clinical staff



Tip 3

- Assemble a multi-disciplinary team including providers, stakeholders, and methodologists, tailored to the specific aim of the project
 - Be agnostic with respect to disciplines and titles when assigning roles and rewards
 - If publication is anticipated, define roles and authorships very early on
 - Giving appropriate first authorships to non-MDs does not jeopardize publication in leading journals



Tip 4

- Be creative in recruiting experts
 - Behavioral scientists, sociologists, economists, epidemiologists, statisticians, qualitative researchers, and other experts often are looking for opportunities to partner with clinical researchers, especially if there is a prospect of co-authorship



Tip 5

- Adopt the most rigorous study design possible without disrupting routine work unduly
 - Incorporate data collection into usual activities of professional staff (eg: infection control, clinical pharmacists)



Tip 6

- Do everything possible not to sacrifice data quality and completeness
 - Develop simple data collection tools that also simplify and increase reliability of daily work
 - Checklists and standardized order sets are especially useful



Tip 7

- Take advantage of emerging certification requirements for clinical staff and make improvement academically viable
 - MOC requirements can be satisfied by improvement activities (eg: Vermont Oxford's NICQ collaborative)
 - Morph “good citizen” work, such as CPG development and evaluation, into publications and other CV-worthy work products



Tip 8

- Do not assume that substantial external grant funding is required to perform credible quality improvement work
 - Leverage institutional resources
 - Encourage development of institutional small grant awards for quality improvement
 - Consider support from payers, industry, and professional societies
 - Look for “free” hands, such as graduate students



Tip 9

- Pay careful attention to the ethics of quality improvement work, but try to craft projects that are unlikely to require formal IRB approval
 - Remember
 - Poorly designed projects are unlikely to yield useful knowledge and arguably are not ethical
 - Patients have a right to expect that unexpected consequences will be considered and monitored



Tip 10

- Anticipate publication
 - Apply the SQUIRE guidelines
 - Write a “dummy” abstract and construct “dummy” tables and figures
 - Be clear about authorships
 - Make the most of “negative” studies

