



Enterprise-Wide Medical Device Integration

First CIS Qatar International
Conference

*20 April, 2013
The Ritz-Carlton
Doha, Qatar*

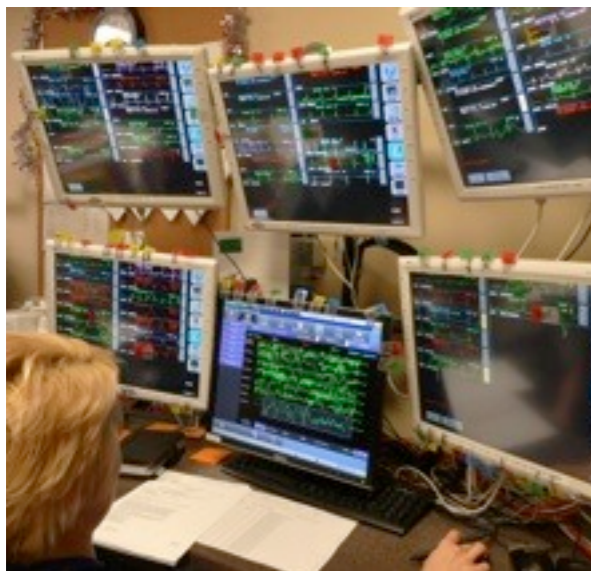


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The majority of clinical data originates from biomedical devices. For maximum efficiency, data quality and timeliness of data, the acquisition of biomedical device data must be automated.



Connectivity

Workflow automation through the integration of medical devices and information systems

Interoperability

The ability of two or more systems or components to exchange information *and to use the information that has been exchanged*

Manual Data Acquisition

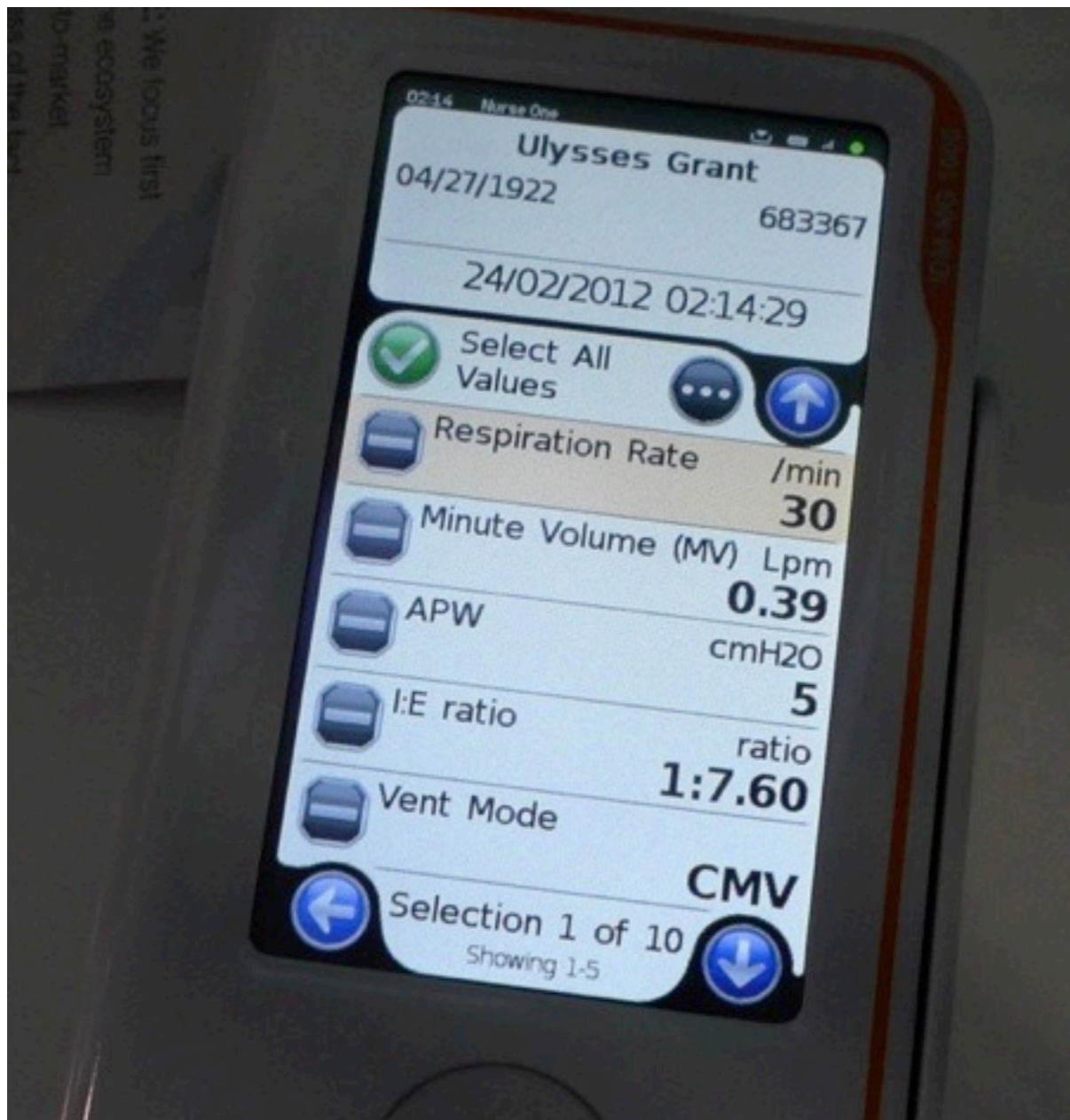
- ▶ Data collected or entered late or period missed: 15 min - 8 hours
- ▶ Typographical or transposition data entry errors
- ▶ Right data entered into wrong patient
- ▶ For nurse, entering data read off medical devices into EMRs is extra work for no value



Functionality

- ▶ Automatically acquires medical device data at specified periods
- ▶ Acquisition period definable by patient
- ▶ Provides clinician validation of data when required

Benefits



Acquired data reduces transcription and documentation errors - to zero if barcodes are used

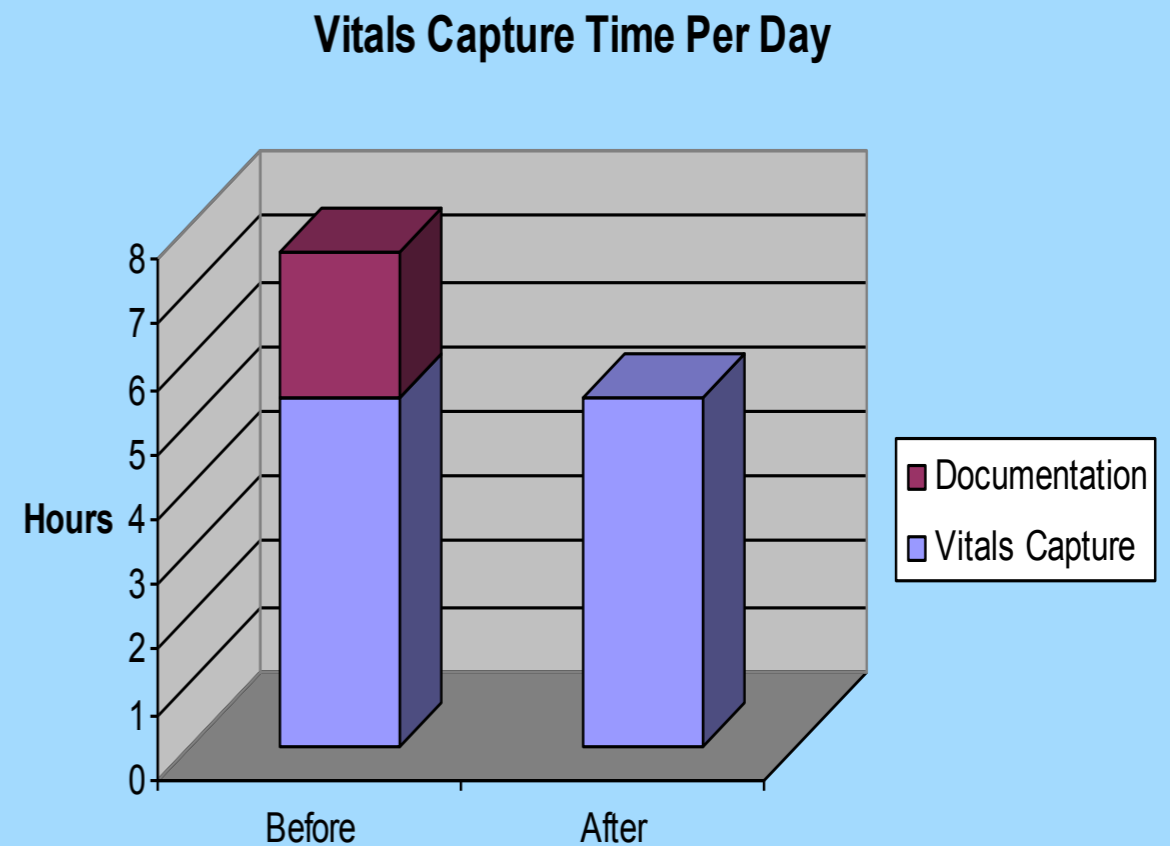
Data immediately available once acquired

Larger volume of consistent data

Increases adoption of EMRs and CPOE

Productivity Impact

- ▶ Average vitals capture time = 3.79 minutes
- ▶ Approx. 33% of taking vitals is admin
- ▶ Automating vitals saves 1.12 minute per vital event
- ▶ Time saved = 28 to 38 hours per day



Source: Automating the Vitals Capture Process, Michael Phelps, 2005, VA Medical Center, Minneapolis, MN

MARCH 2013

THE VALUE OF MEDICAL DEVICE INTEROPERABILITY:

Improving patient care
with more than \$30 billion
in annual health care savings

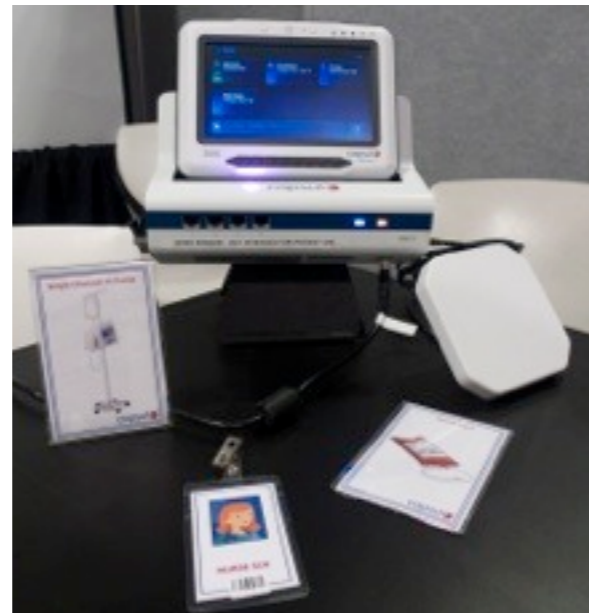


Impact on Patient Outcomes

Policy white paper
evaluating impact on U.S.
health care costs

Source: <http://www.westhealth.org/institute/interoperability>

Connectivity Impacts



Costs resulting from
redundant testing: \$3 Billion

Clinician time spent manually
entering data: \$12.4 Billion

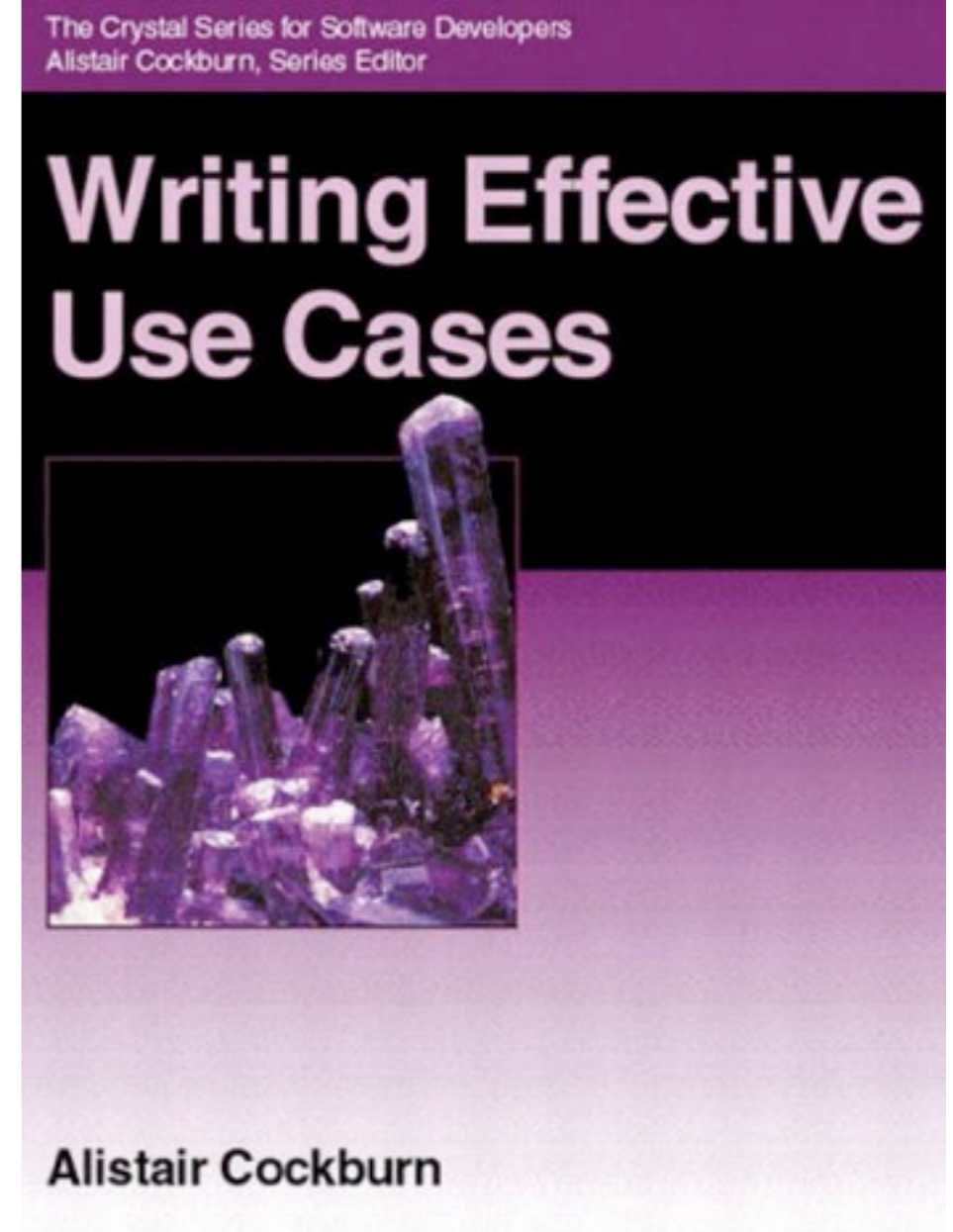
Increased length of stay:
\$17.8 Billion

Typical Workflows

- ▶ Patient Management
- ▶ Patient Context
- ▶ Scheduling
- ▶ Order Workflow
- ▶ Data acquisition
- ▶ Data analysis
- ▶ Clinical documentation
- ▶ Surveillance
- ▶ Messaging
- ▶ Data management
- ▶ Report generation
- ▶ Device specific workflows

Use Cases

- ▶ A methodology used in systems analysis to identify, clarify workflow and organize system requirements
- ▶ Day-in-the-life story, formal very structured text, flow charts, sequence charts, etc.
- ▶ An agreement between users and product developers regarding how a system will operate

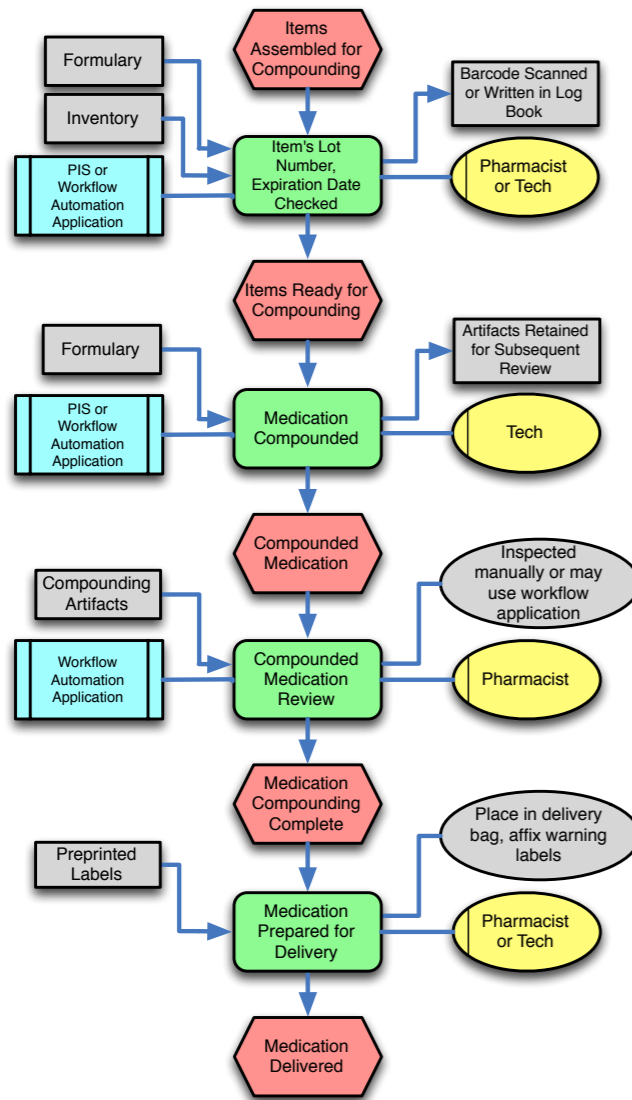


Use Cases Vary by Medical Device

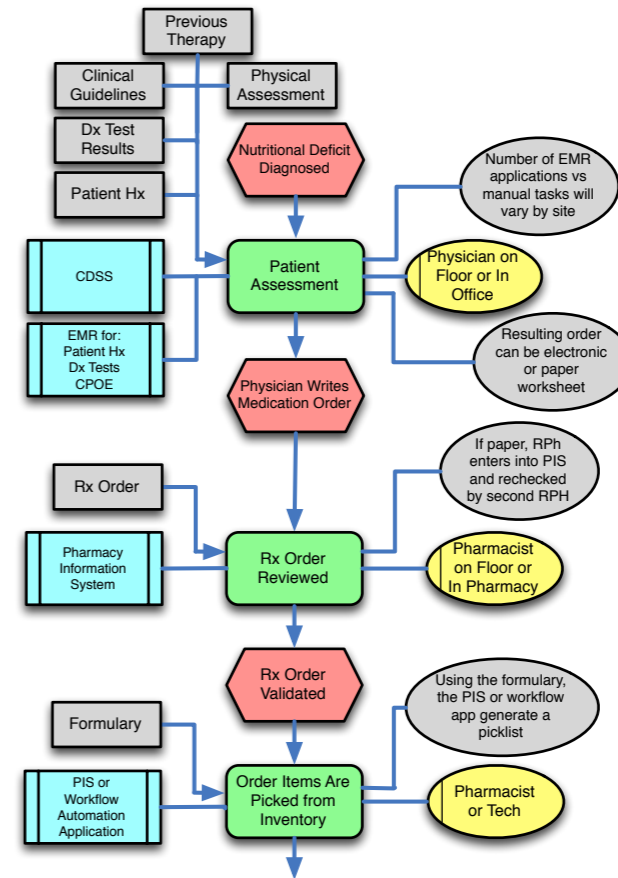
- ▶ Vital signs monitors - *spot check*
- ▶ Continuous patient monitors - *sampling of data stream*
- ▶ Infusion pumps - *limited EMR support*
- ▶ Ventilators - *limited EMR support*
- ▶ Workflows for the same device can vary across clinical areas - *e.g., ED, surgery, ICU or Med/ Surg units*



Clinical practice use cases



Product Use Cases

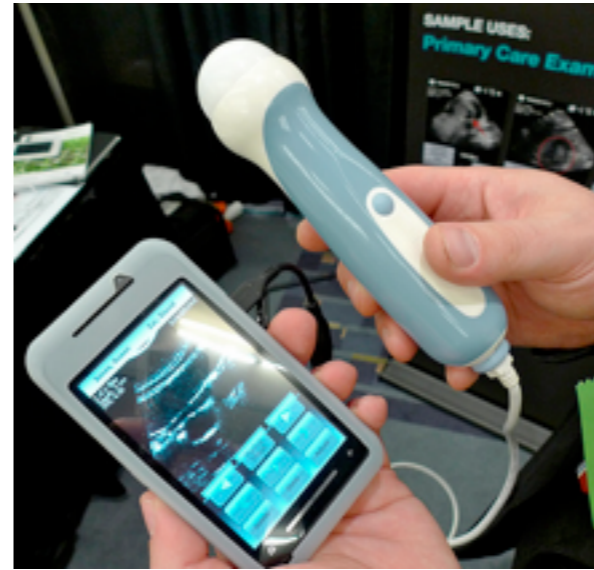


Workflow matches
 Workflow gaps
 Flexibility - configurability
 Evaluate trade-offs

Recommendations

- ▶ Document your workflow with use cases
- ▶ Require vendors to disclose their uses cases
- ▶ Include use case requirements in purchase agreements to ensure vendor compliance





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Patient Management

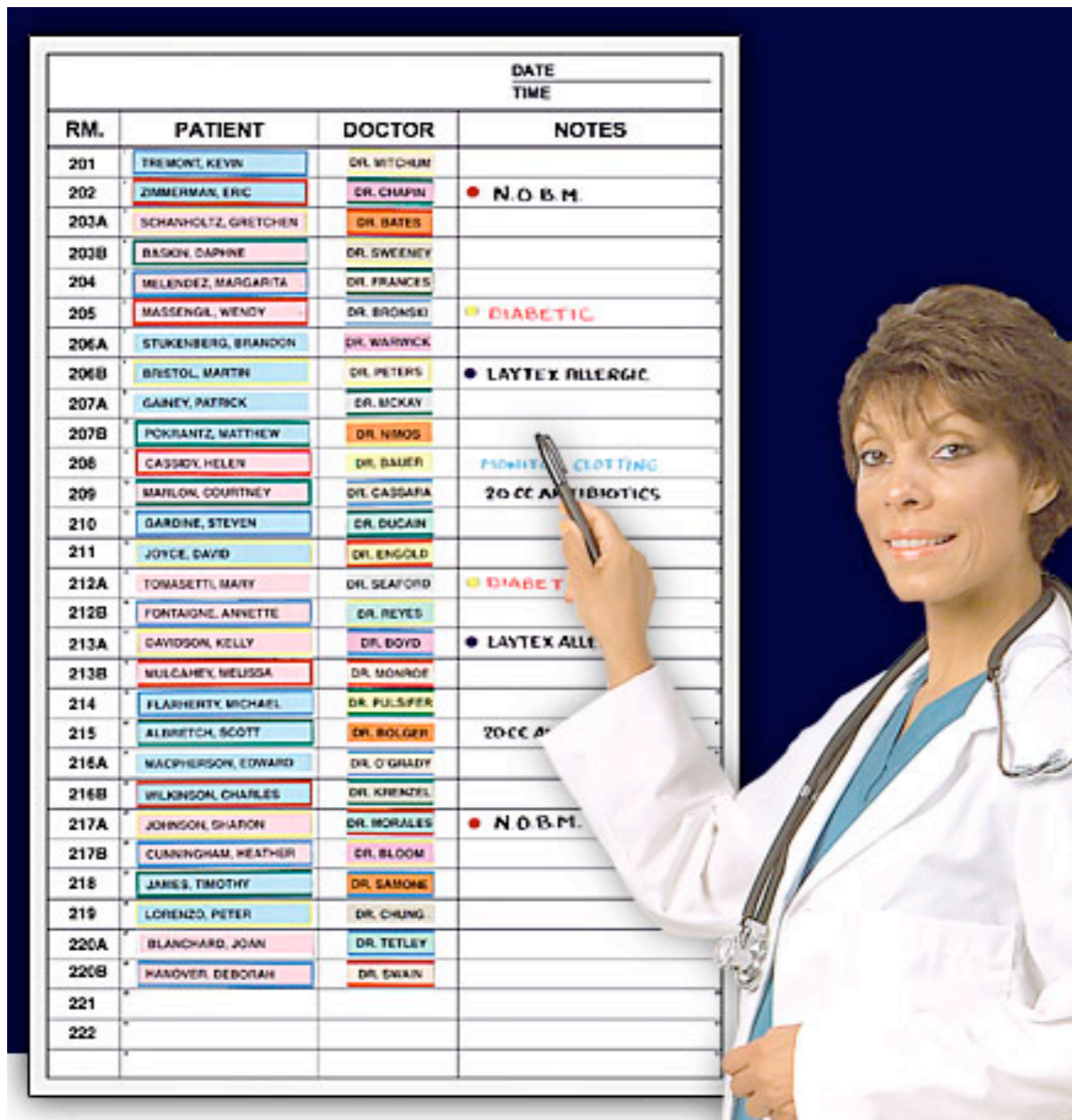
Database of relevant patient demographics

Add, edit and delete patients and associated data

Supports flow of patients' contact with medical device system

Data captured prior to, or at the time medical device is applied to patient

Analogs: department marker boards, dashboards



Patient Context



Associates medical device data with the patient attached to the device

Creates and breaks down association with application and removal of device

Maintains reliable association during use - wireless makes this challenging

Essential step requires backup methods

Best managed at point of care

Scheduling

Patient and resource scheduling

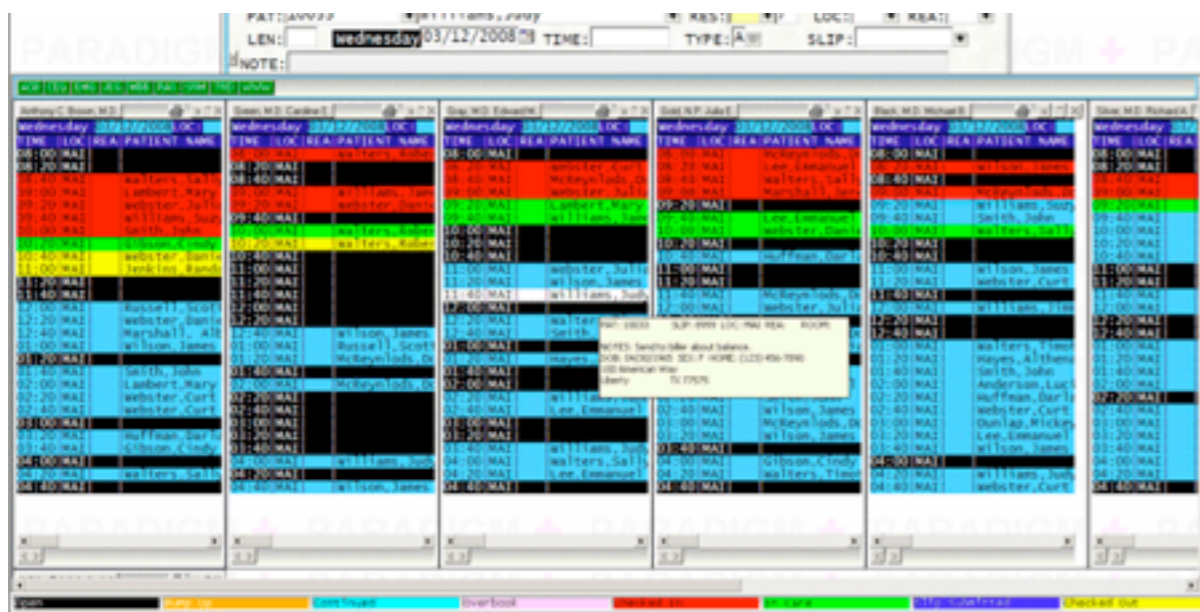
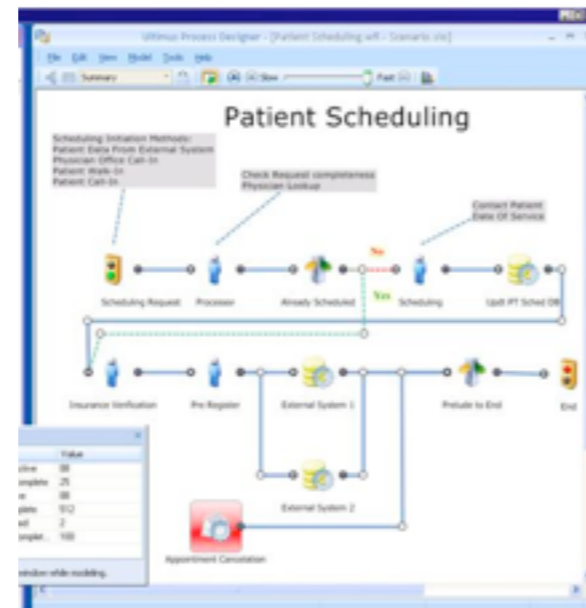
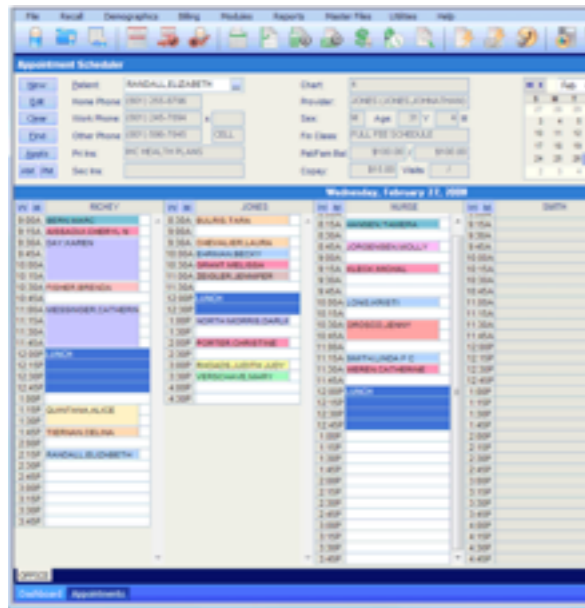
Can be time-specific or a work list

Patient reminders

Driven by order set

Order-specific prerequisites

Complex rules-driven scheduling algorithms



Order Workflow



Order initiates workflow; can initiate patient management
Specifies how medical device is to be used

Monitoring or therapy orders define device configuration

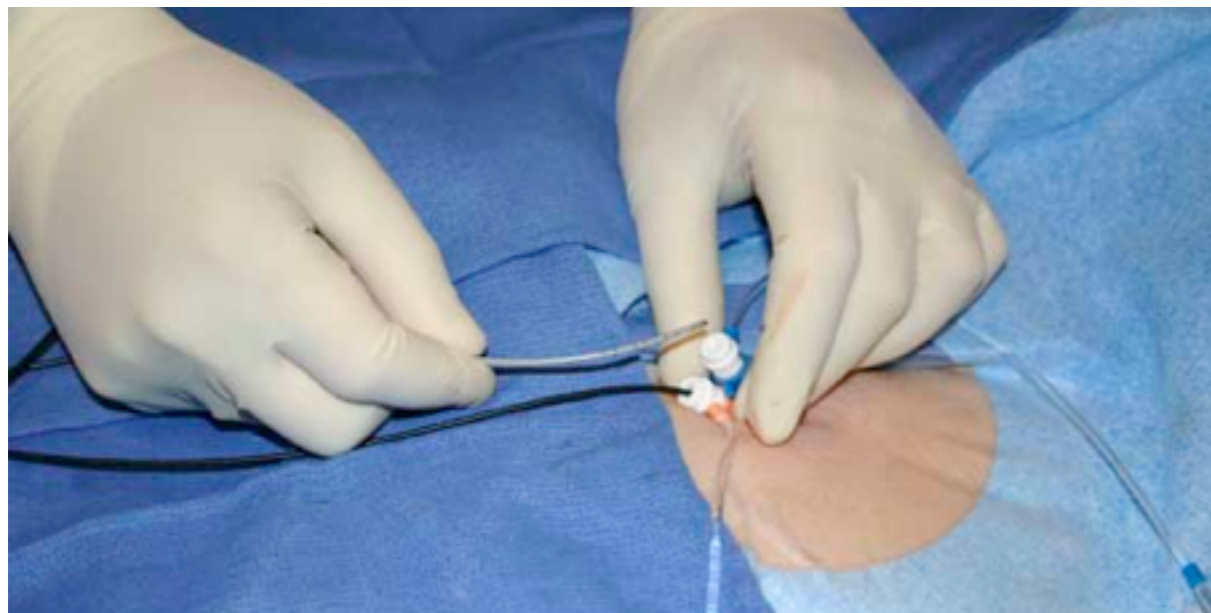
Diagnostic orders define device configuration and procedure

Can include establishing patient context

Dependent on patient management

Includes a formal end step

Data Acquisition



Medical device data is acquired

Can be continuous, periodic or both

Acquired data can include multiple data sets

Can include data annotation - both automatic and manual

Acquired data is often edited for reporting or clinical documentation

Data Analysis



Proprietary algorithms extract physiological data and improve data quality

Data analysis to render proposed diagnosis

Can be done in embedded device or via network

Can be automated or user directed

Clinical Documentation



EMR or procedure documentation

Subset of medical device data is often used

Combination of automated or user-generated data

May require data validation step

Surveillance



Display of device data in locations beyond the medical device

Continuous stream of data

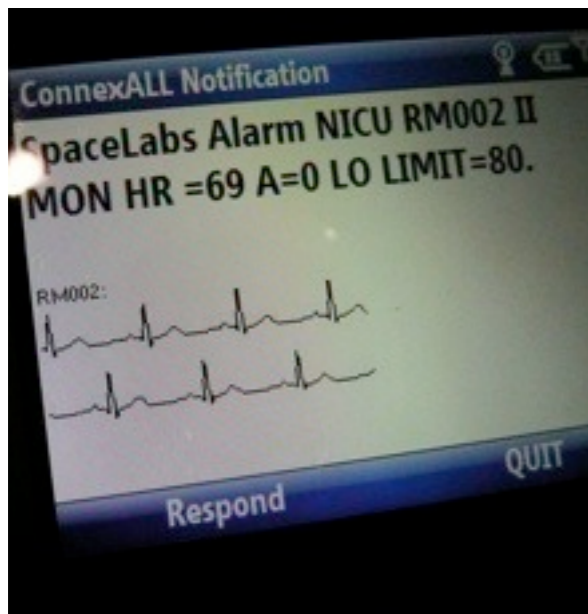
Near real time requirement

Highly variable transmission distances

Wide variety of client devices

Implemented as “slave” monitor or part of broader workflow

Messaging



Can include audio, text data and waveforms

Broadcast or narrow cast

Contextual clinical data often required

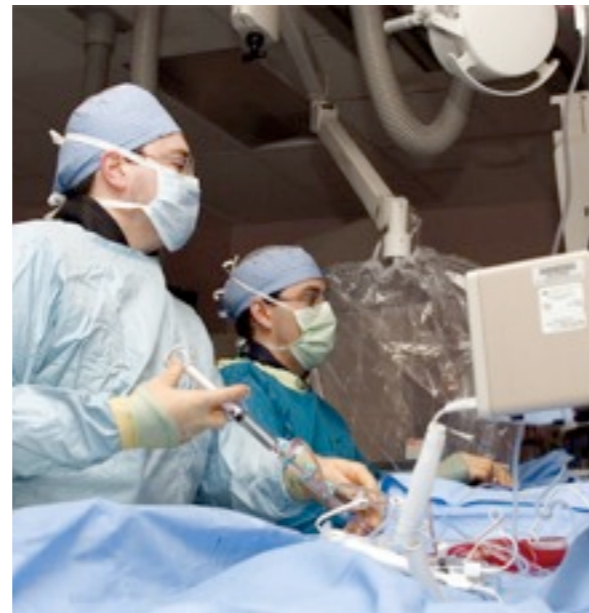
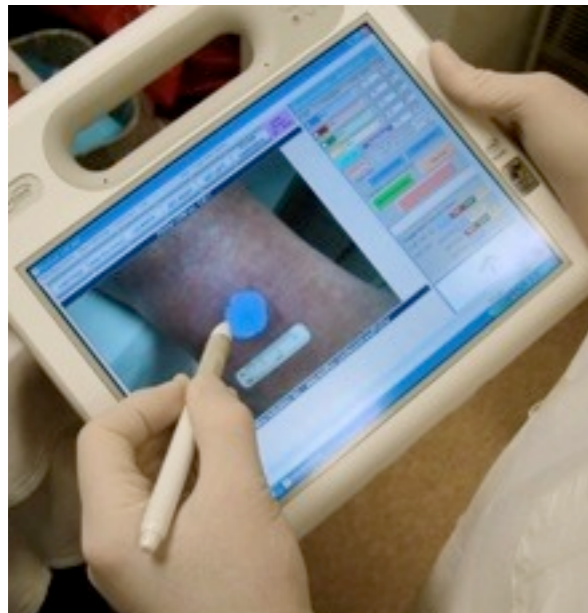
May require escalation

May track event driven messages to conclusion

May include complex rules-driven

Wide variety of client devices

Report Generation



- Fulfillment of order workflow
- Diagnostic report generation
- Procedure documentation
- Report generation, review and approval workflow
- Structured or unstructured data
- Management reporting

Data Management



Supports patient management and order workflow

Event review

May include annotation or editing of data

Data queueing for review or analysis

Storage and retrieval

Data archiving with appropriate retrieval

Teaching file generation

Clinical trial and research support

Device Specific Workflows



Basic templates: diagnostic, monitoring, and therapeutic

Device specific variations and combinations of preceding workflows

IV pump Drug Error Reduction Systems (DERS)

QA requirements