

kinetik solutions



Lean London Forum



Lean Executives

executive search & selection for the lean enterprise

21 March 2012
Royal College of Surgeons

For more information, please email help@leanlondon.org.uk or telephone 0207 824 8448



We have some broad aims of the forum

- **Create the environment where Lean Solutions in the NHS are shared, discussed and acted upon by practitioners in the health service**
- **Engage in a debate about strengths and weakness of lean in the current NHS climate**
 - The QIPP agenda in reducing costs across the health system
 - Clinical Commissioning Groups that will redefine 'end to end' health systems processes
- **To network with colleagues and friends**



Agenda

- **1800 - 1810** **Welcome and introductions**
- **1810 - 1835** **The Path-ology to Lean Thinking - Dr Mathew Diggle & Suzanne Horobin**
- **1835 - 1900** **Pre-operative health evaluation - engagement with Primary Care - Dr Ahmed Chekairi, Whittington Hospital**
- **1900 - 1930** **Hot seat session**
- **1930 - 2000** **Networking and drinks**



Introductions



- **Your Name**
- **Your Role**
- **The one thing relating to Lean you are curious about?**



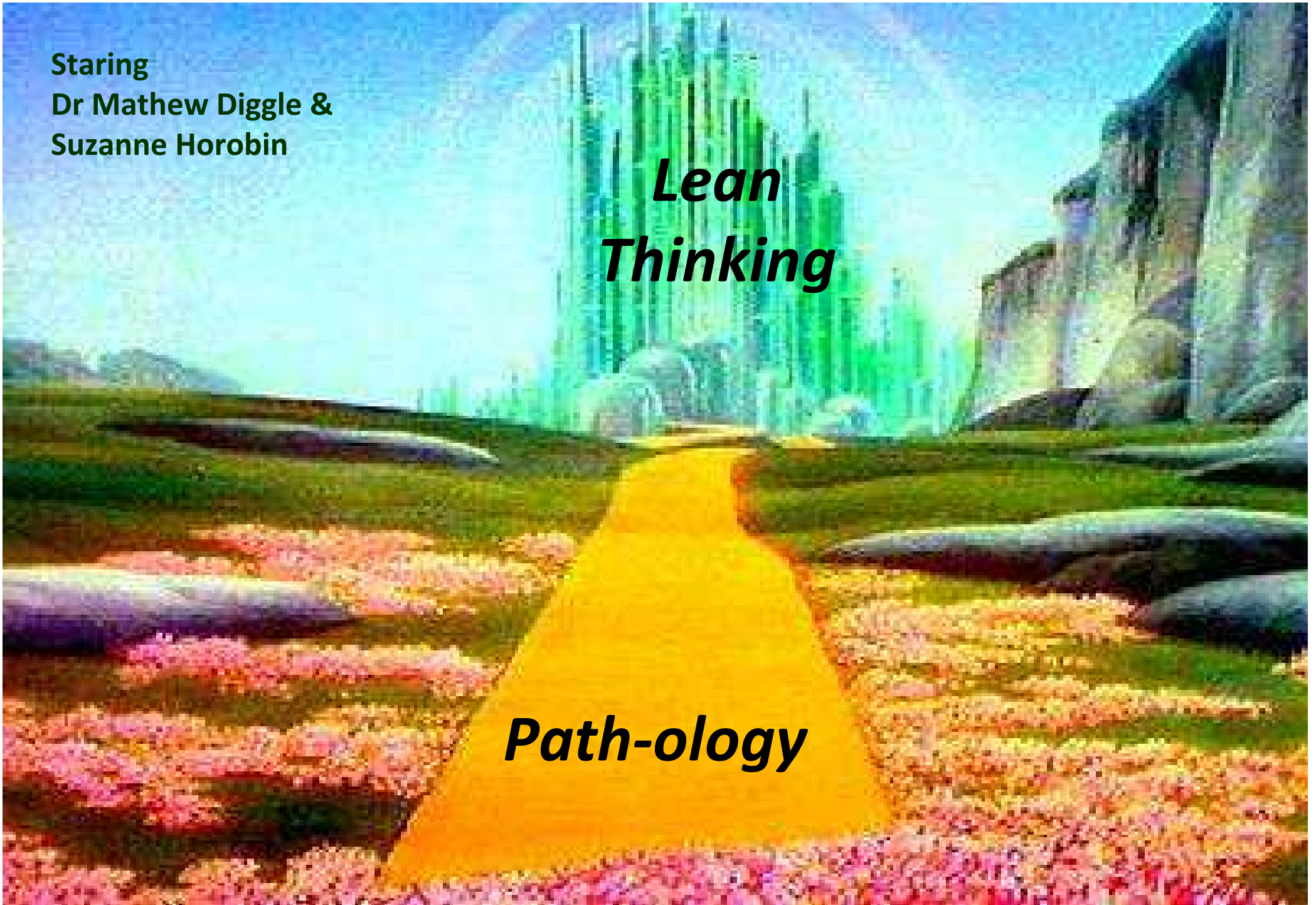
Recap – What is Lean?

- Focus on Value from a **Customer (Patient) point of view** on every step of process
- **Obsession on removing waste** within the ‘whole system’
- Bottom up approach in identifying value and waste – assumption that **much of waste and value is hidden**
- A true lean system would “flow” and need **little command and control**

Staring
Dr Mathew Diggle &
Suzanne Horobin

Lean Thinking

Path-ology



Background

- Diagnostic service: 24/7, 365 days per year
- Population served: >2.5 million
- Workload: 970,000 pa
- Isolation, identification and detection of medically important bacteria, viruses and parasites.



Background

- Screening and specialist service: expertise in biological agent detection
- Clinical advice -on the diagnosis, management & treatment of infections, with regular ward rounds on intensive care etc.
- Infection control- MRSA isolation and Clostridium difficile toxin screening



"All these years we battled the bugs, and all it took was a sign?"

What is Lean?

LEAN

- Perfected by Toyota – from 1928
- ‘Lean’ coined by Jones & Womack in 1990s
- Lean is about ***improving flow and eliminating waste***
 - getting the right things to the right place,
 - at the right time,
 - in the right quantities,
- while minimising waste and being flexible and open to change.
- ***Customer*** at heart of the process
 - Driver for Quality and Safety

LEAN?



CHANGE



DANGER!



FEAR





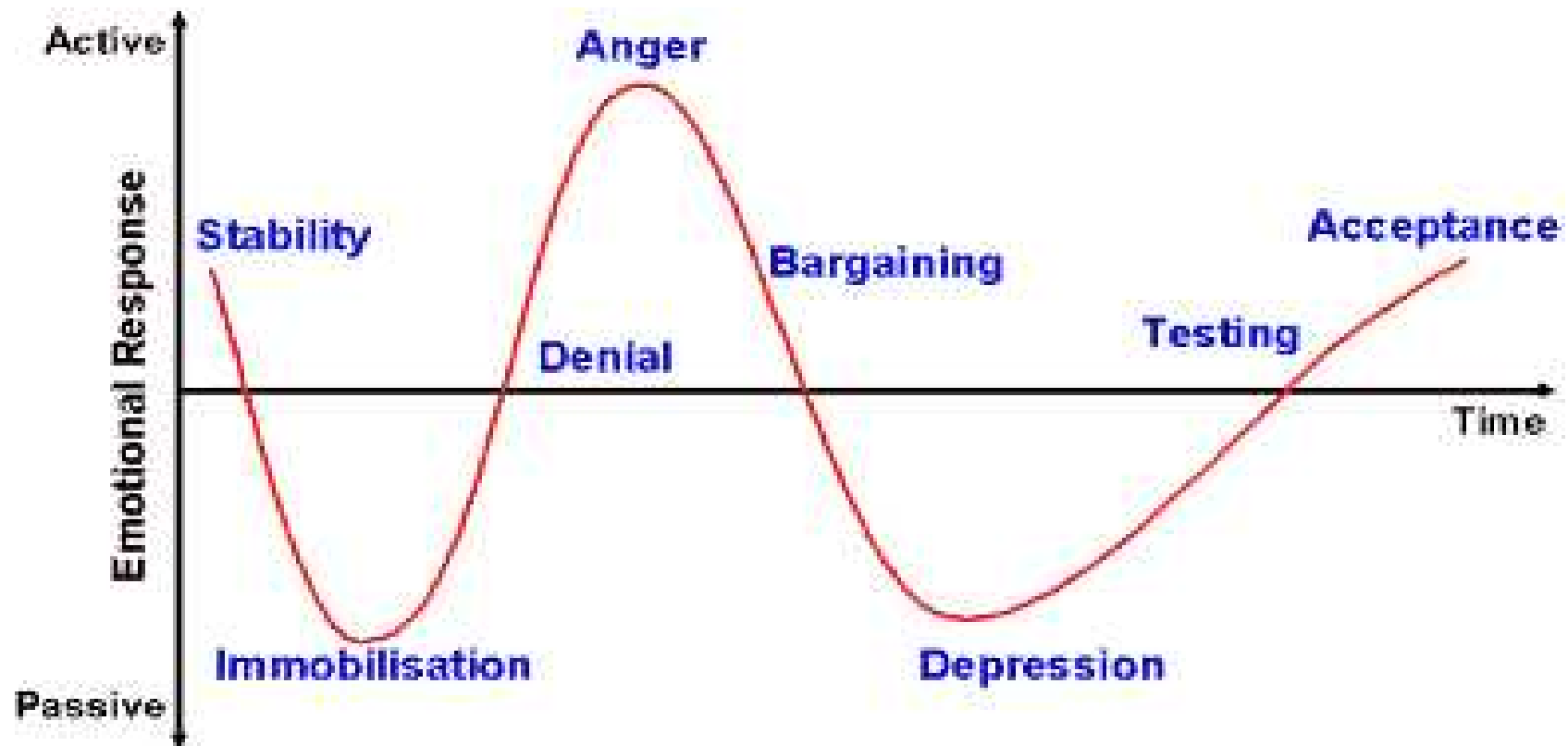
RESISTANCE

A large cargo ship is seen from a high angle, sailing on a dark, choppy sea. The ship's deck is visible, showing several large, reddish-brown rectangular sections. The sky is filled with heavy, dark clouds, with a bright, glowing area on the horizon where the sun is setting or rising, creating a dramatic silhouette effect. The word "LEAN" is overlaid in white, sans-serif capital letters on the left side of the image.

LEAN

Microbiology

The Path-ology



The Path-ology

- High Volume testing – Urines and MRSA screening
- Highly visible – Reception area
- Highly productive – What matters to me!
- Highly effective!

High Volume testing – Urines and MRSA screening

One piece flow

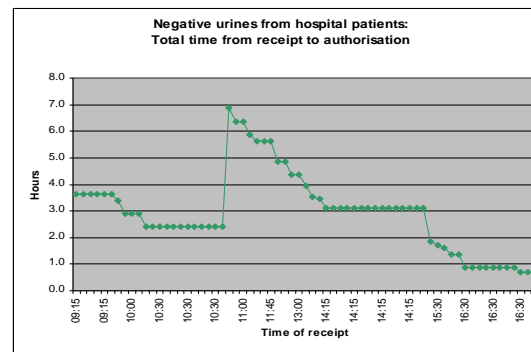
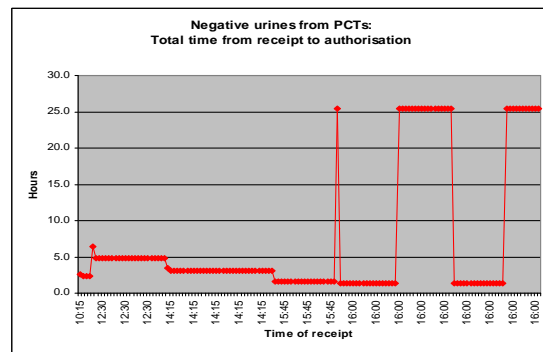


Results from Urines

% reported day of receipt			% reported within 24 hrs			% reported within 48 hrs		
Baseline Aug-11	Sept-11	Oct-11	Baseline Aug-11	Sept-11	Oct-11	Baseline Aug-11	Sept-11	Oct-11
40.9	39.8	39.2	58.7	66.1	77.4	85.9	86.9	95.7

Results from Reception

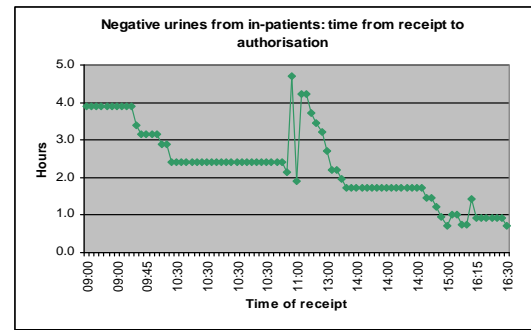
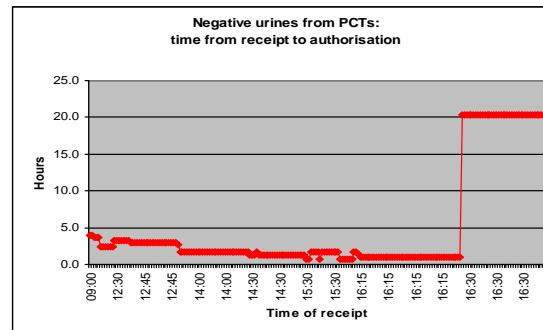
Then



Most GP specimens' turnaround time (TAT) < 5h (median=3.1h). Overnight storage results in TAT c. 25h

Most in-patient specimens' TAT <4h, but may be as high as 7h (median=3.1h).

Now



Most GP specimens' turnaround time (TAT) < 4h (median=1.7h). Overnight storage results in TAT c. 20h

i.e. **median TAT reduced by 45%**
(mean TAT reduced by 29%)

Majority of in-patient specimens' TAT <3h
(median=2.4h)

i.e. **median TAT reduced by 23%**
(mean TAT reduced by 23%)

Overall, **TAT for all negative samples has been reduced by 45%** (median 3.1h to 1.7h)

Number of samples processed via UF100 **increased by 11%** on dates sampled.

Highly visible – Reception area



Highly visible – Reception area



Improved process

- Post-Lean, there is **improved prioritisation of in-patient samples**, as the time from receipt to UF100 processing has been reduced by 50% (median 0.8h to 0.4h) and time from receipt to registration has been reduced by 25% (median 0.8h to 0.6h).
- **Removal of excessive checking** (demo-checking) has reduced the processing time for negative samples by 43% (from 2.3h to 1.3h).
- Positive feedback from staff as process runs more smoothly and calmly.

The Path - Key Challenges

- Maintaining momentum / energy / time
- Communication with all stakeholders
- Support laboratory staff and stakeholders



NHS Improvement
Diagnostics



Right First Time?



The cost of poor quality

Suzanne Horobin

National Improvement Lead -
Diagnostics



Who are NHS Improvement?

- DH partnership - aligned to support delivery of national strategies and priorities - support QIPP Pathology Programme
 - Key sponsors – Prof Sir Mike Richards, Prof Roger Boyle, Dr Ian Barnes, Dr Erika Denton, Prof Sue Hill
 - Over 10 years improvement experience, practical knowledge and “how to” approach to improve quality and productivity
 - Last 12 months – supported 240+ sites - improving Cancer, Diagnostics, Heart, Lung and Stroke services
- 
- 



NHS Improvement - Diagnostics

- Proven success in Pathology
 - 95% Histopathology results in 7 days, 50% in 3 days – Whipps Cross
 - 100% Cytology results in 14 days, >50% in 7 days – Phase 1/2 pilot sites
 - 50% improvement in Phlebotomy outpatient waiting times - Doncaster
 - 90% reduction in Microbiology TAT – Stoke on Trent
 - 43% reduction in Histopathology TAT – Calderdale & Huddersfield
 - 62% reduction in Haematology TAT – St James Hospital
 - 21% reduction of inappropriate demand urines – St Helens & Knowsley
 - Improvements in safety and quality
 - Increased capacity and space utilisation
 - Improved staff engagement
 - National Clinical Lead / Associate support
- 
- 

Right first time?



What's the problem?

- University Hospitals Coventry and Warwick Cytology – 47% forms missing sender code
 - Hull and East Yorkshire Hospitals NHS Trust – 154 cytology samples in 2 weeks received with errors
 - Taunton & Somerset NHS Foundation Trust – Musgrove Park Histopathology – 28% specimen receipt / data entry errors
 - Nottingham University Hospital NHS Trust – 8941 microbiology samples rejected due to defects (12.8%)
- 
- 

How do we know?

* CORE TEAM WALKED THE PATHWAY *
27 OCTOBER 2008

CURRENT STATE VALUE STREAM MAP

TACT: 72s
Time available: 410
350
Need to bring
Samples to team

350
Samples
per day

GP
SURGERY
CASH / MRP

HQMM
OFFICE

WYCSA

PATIENT

RESULT
RECEIVED



SAMPLE
TAKEN
01

349

RECEIPT
+
NUMBER
02

125

REGIST
CASE
07

1

PROCESSING
(LAB)
02

3

8

PRE
+
SCREENING
03

784

LAB
SIGN-OUT
01

334

RESULT
03
PRODUCTION
WYCSA

4

11

W
1.5 DAYS

W
5.2 DAYS

W
5 DAYS

96 min
C/S
< 1%
Looking for
Office

W
6 DAYS

5
11
<
to SW

W
1 DAY

1 1/2 min
16
< 1%
FURTHER
SCREENING

W
UP TO 5 DAYS

20 min
400-700
< 20 %
NEEDS LAB
WYCSA
TACT

W
2 DAYS

KEY

--- Delay

--- Pull

Δ - Waiting time

Δ - Inventory

⊖ - Pause at each
stop

☆

WYCSA

Looking for
Office

Looking for
Office

Looking for
Office

Looking for
Office

1 min

4 1/2 min

9 min

5-15 min

1 1/2 min

20 min

TOTAL TIME =
128-158 mins

9850
LEAD TIME

Insert the measurements - Gather **real data**
- this is your chance to **really understand**
what's happening.

START

Patient Referral

Sample Sorting

Cycle time =
Batch Size =
Defects =
Trigger=

Defects – What percentage is not right first time and requires further work?

- Manual data collection
- One week
- Tick every error that occurs
- Capturing lab number will enable root cause analysis later
- Errors as a % of total workload for the week

So what?



Result for patient?

- Repeat testing
- Delayed diagnosis
- Potential for incorrect diagnosis / treatment

Result for department?

Cost of poor quality - COPQ

Cost of poor quality - COPQ

Ideal – every case is booked in RIGHT FIRST TIME

- Dealing with a typical error requires a 5 minute search of the current laboratory database followed by a 5 minute telephone call to find the correct information = 10 mins per error
- 5% of cases have errors requiring this rework
- 5% of annual workload of 50,000 samples = 2500
- Handling time 2500 samples x 10 minutes = 25000 minutes
- Or 416 hours
- Or 55.5 WORKING DAYS
- At midpoint Band 3 = £5170.88

This is the cost of poor quality

What do we do about it?





“ALL USERS” email



DOESN'T WORK!!!

how many emails do you get every day?



GET MORE DATA



A weeks worth of data tells you
enough



GO SEE



Now we're getting there!



DiagnosticsImprovement

NHS Improvement **NHS**

Put label on each copy of form. Use a PAS label if available. Need NHS number if GP patient. Check hospital number is accurate. For accurate DOB check with patient. Check details match label on specimen pot.

Provide Clinical detail and any **antibiotic** treatment. Give detail of any **"Foreign Travel"** with date and countries visited.

Give GP details **NOT** initials and Drop number. If a copy required give full details. Requesting doctor **MUST** sign the form.

Give Full name and bleep number of requesting Dr this **MUST** be signed by requesting Dr. Always give full name, **NOT** initials, of requesting consultant under whose care patient is. Give ward in which patient is.

Indicate **Test** required and **Sample Type**. Specify site of the sample to look for appropriate micro-organisms. Do **NOT** mislead **CSU** (Catheter Specimen Urine with M&C (Microscopy and Culture).

Please specify: **DATE** and **TIME** of sample collection. It is recommended urines **SHOULD** be tested within 2 hours of its collection.

CLINICAL DETAILS
Drug and Antibiotic Therapy _____ essential

FOR GP USE ONLY - NAME & PRACTICE ADDRESS
GP _____ DROP NO. _____
SIGNATURE _____

HOSPITAL NUMBER _____
NHS NUMBER _____
SURNAME _____
FORENAME _____
DATE OF BIRTH _____ SEX _____
POST CODE _____
WARD/HOUSE No. _____
REQUESTING DR. _____
BLEEP NO. _____
DR. SIGNATURE _____
CONSULTANT _____
DATE OF REQUEST _____

Urgent Request send to station 999 ☐
Private ☐

CHEM PATH 5ml SST
☐ Renal
☐ Liver
☐ Bone
☐ Lipid Profile
☐ TSH
☐ PSA
☐ B12 Folate

HAEMATOLOGY 4ml
☐ FBC
☐ ESR
☐ GF Screen
☐ Sickie Screen
☐ HD Electrophoresis
☐ Antinuclear Ab 7.5ml
☐ RPR Citrate
☐ Coag Screen
☐ INR/PT and Urea/Vit D

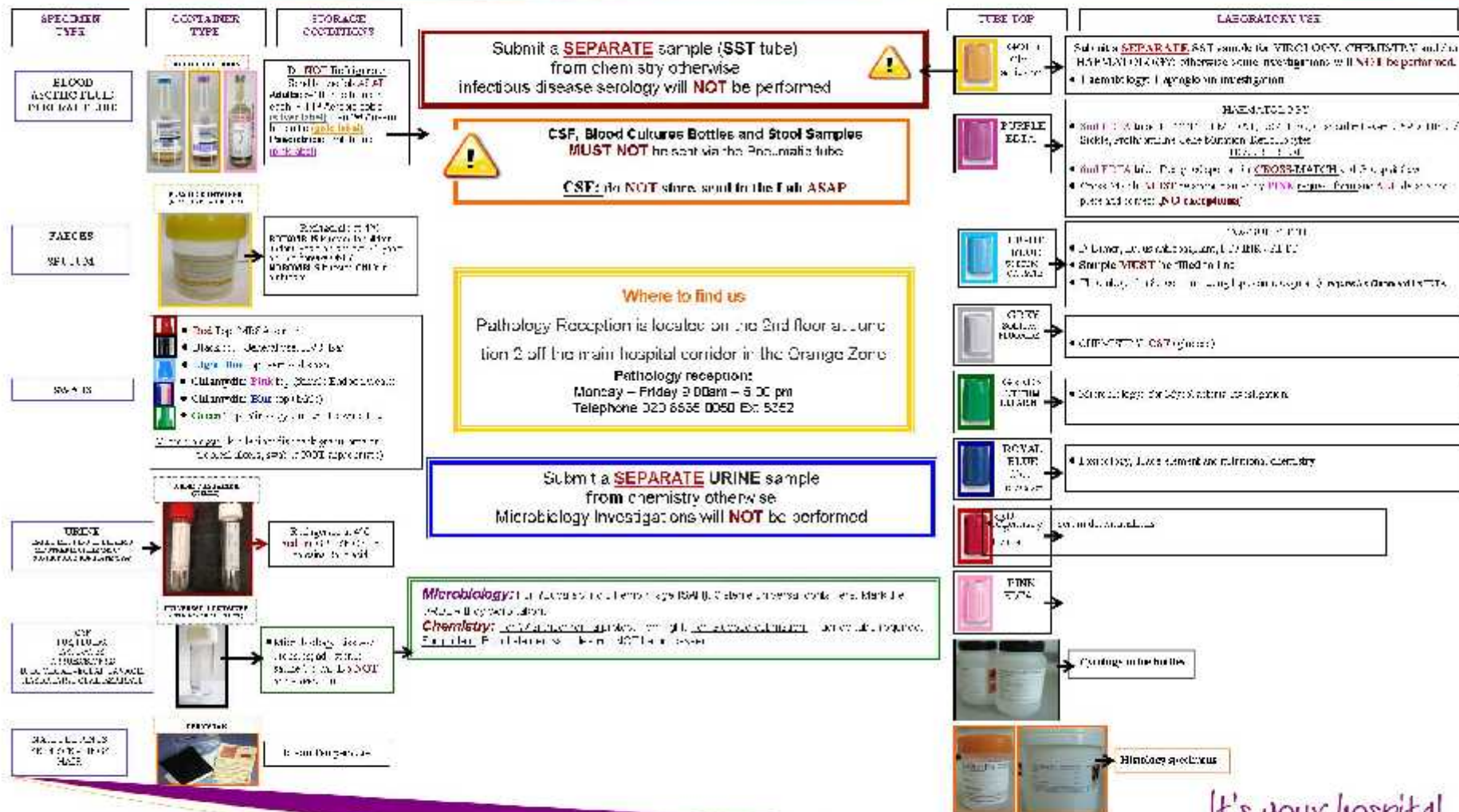
ROUTINE MICROBIOLOGY
Urine: MS/CSU Micro & Cult
Genital Swab Culture
Site _____ Chlamydia
Swab Culture
Site _____ Culture
Pus/Spec - other Culture
Fluid Culture
Respiratory specimen Routine
Tissue AFB

SEROLOGY (5ml SST)
☐ Antenatal Screen
☐ Sexual Health Screen
☐ Hepatitis A - Acute
☐ Hepatitis B - Immunity
☐ HIV
☐ VZV - Immunity
☐ VZV - Contact
☐ Other

SAMPLE DETAILS
Date _____ Time _____
Fasting ☐ Taken by _____
SST _____ EDTA _____ Citrate _____
Heparin _____
Contact Date _____
Indicate if Pregnant ☐ No. of _____

LAB

PATHOLOGY SAMPLES

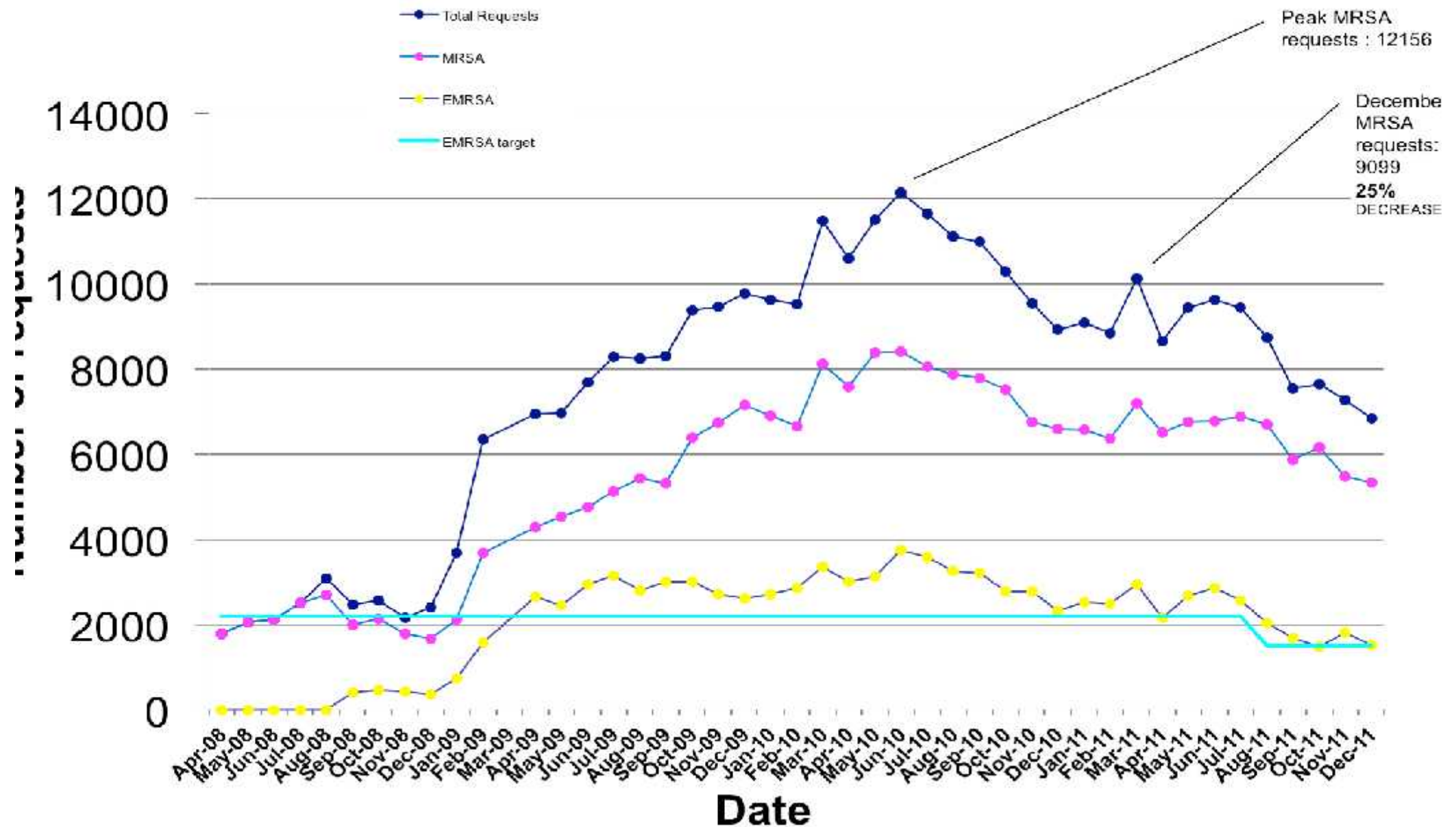


It's your hospital

Does it really work?

- University Hospitals Coventry and Warwick Cytology – reduction in errors reduced booking in time saving **1789 hours of office time per annum (240 working days)**
- Hull and East Yorkshire Hospitals NHS Trust – reduction in returned samples freed up **10 hours per month**
- Taunton & Somerset NHS Foundation Trust – Musgrove Park Histopathology – errors down from 29% to 8% so far
- Nottingham University Hospital NHS Trust – 8941 microbiology samples rejected due to defects (12.8%) – work in progress!

Total MRSA requests 2008-2011



Pre-Operative Care Assessment at Primary Care

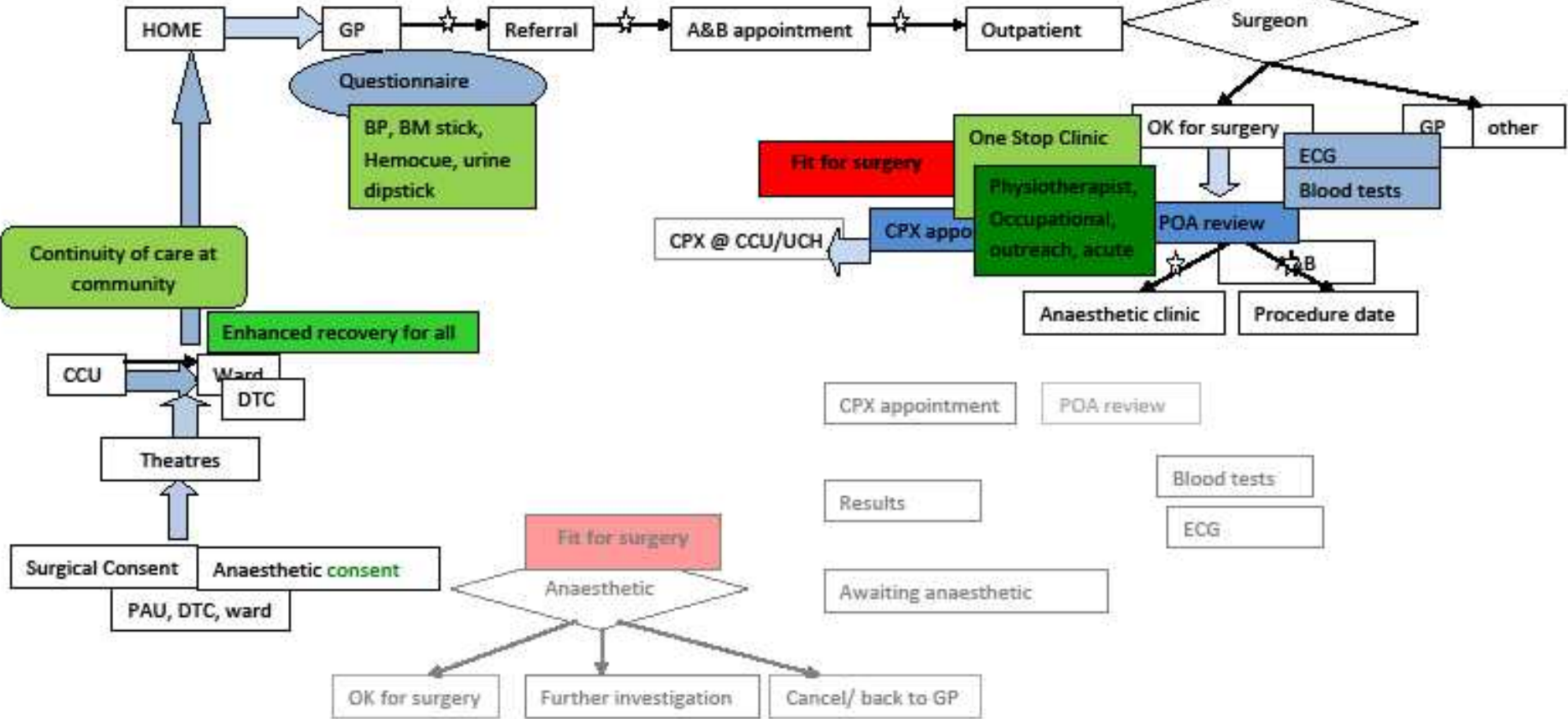
Dr Ahmed CHEKAIRI
MD, FRCA, PGDip
Consultant Anaesthetist
Whittington Health, London

Lean London
The Royal College of Surgeons
21 March 2012




Old Paradigm

- Pre-operative assessment is the process by which patients who are booked in for an elective surgical procedure undergo an anaesthetic assessment to evaluate their fitness for surgery

VSM: Proposed PERIOPERATIVE CARE PATHWAY for majority of patients



Information movement (letters, phones, emails, notes, test results)

 Patient movement
  time in days
  time in weeks
 Green: new introduction
 Blue: movement upward the stream

Reversible Causes

- Blood pressure
- Diabetes
- Infections
- Weight??
- ANAEMIA
- Smoking

New Pre-assessment Service

New Paradigm: Pre-Operative Health Evaluation and Optimisation

New set up

- One stop clinic: Health evaluation on the day decision to operate is made
- 8000 travels/year less, More than 4000 patients would go straight to surgery
- Standardised documentation with traffic light system, triggers for right investigations and right referrals
- Improving patients' safety by including checklists alerts
- Criteria-based Decision making algorithms based on best evidence and in response to National and international Governance bodies
- Systematic risk stratification= better consent=better planning=appropriate use of Intensive care



Promoting good health beyond surgery

- Pre assessment is Health MOT opportunity for surgical patients
- Improving nutritional state pre-operatively and at long-term
- Ensuring good diabetes and blood pressure control & promoting smoking cessation and sensible alcohol intake

Better patient's experience, Less travels to the Hospital, Better quality of care, Less waste

Benefits

- Active treatment of anaemia pre-operatively using intra venous iron
 - Right investigations (more than 20% unnecessary investigations last year)
 - Better diagnosis and more inclusive screening (Sleep apnea and mental state screenings)
 - Promoting day surgery and Enhanced Recovery Care for all patients
 - Systematic assessment of mobility = better post operative planning
 - Minimising cancellation on day
- Systematic optimization of reversible conditions such as diabetes ,HBP, infection & others



Staff

- CPD programme for nurses
- Training programme for trainees
- Consultant anaesthetists run clinics
- Continuous measurement

A patient story

G.H, a self-employed, attended the Pre assessment service on 01/03/12. This was the day of the new pre assessment service trial. He was seen **soon** after surgeon told him about the need for surgery.

Health evaluation revealed one medical concern that required blood tests and an ECG. These were done on the spot in POA clinic.

He was also seen by a Consultant anaesthetist at same visit.

Patient was advised of the risks of his condition and the need for treatment at long term; he was also advised to contact his GP. Then was given a **suitable** date for his surgery

Similar patients in the past would have been added to the waiting surgical list and given 2 dates. One for surgery and another one for pre-assessment. Time interval between the 2 dates is variable. They may also have a 3rd date to see an anaesthetist consultant.

G.H would have lost 3 working days as unpaid leave even before getting his operation.

Future (near future)

- Connect and cooperate with GPs
- Align peri-operative goals between Hospital and Primary care: 'optimise as you refer'

Ahmed.chekairi@nhs.net for more information

Anaesthesia

Journal of the Association of Anaesthetists of
Great Britain and Ireland

Editorials

- 103 The role of steroids in treating septic shock *V. Grover and J. M. Handy*
106 Surgery and cardiovascular outcomes: an untapped public health benefit that potentially saves lives *B. M. Biccard*

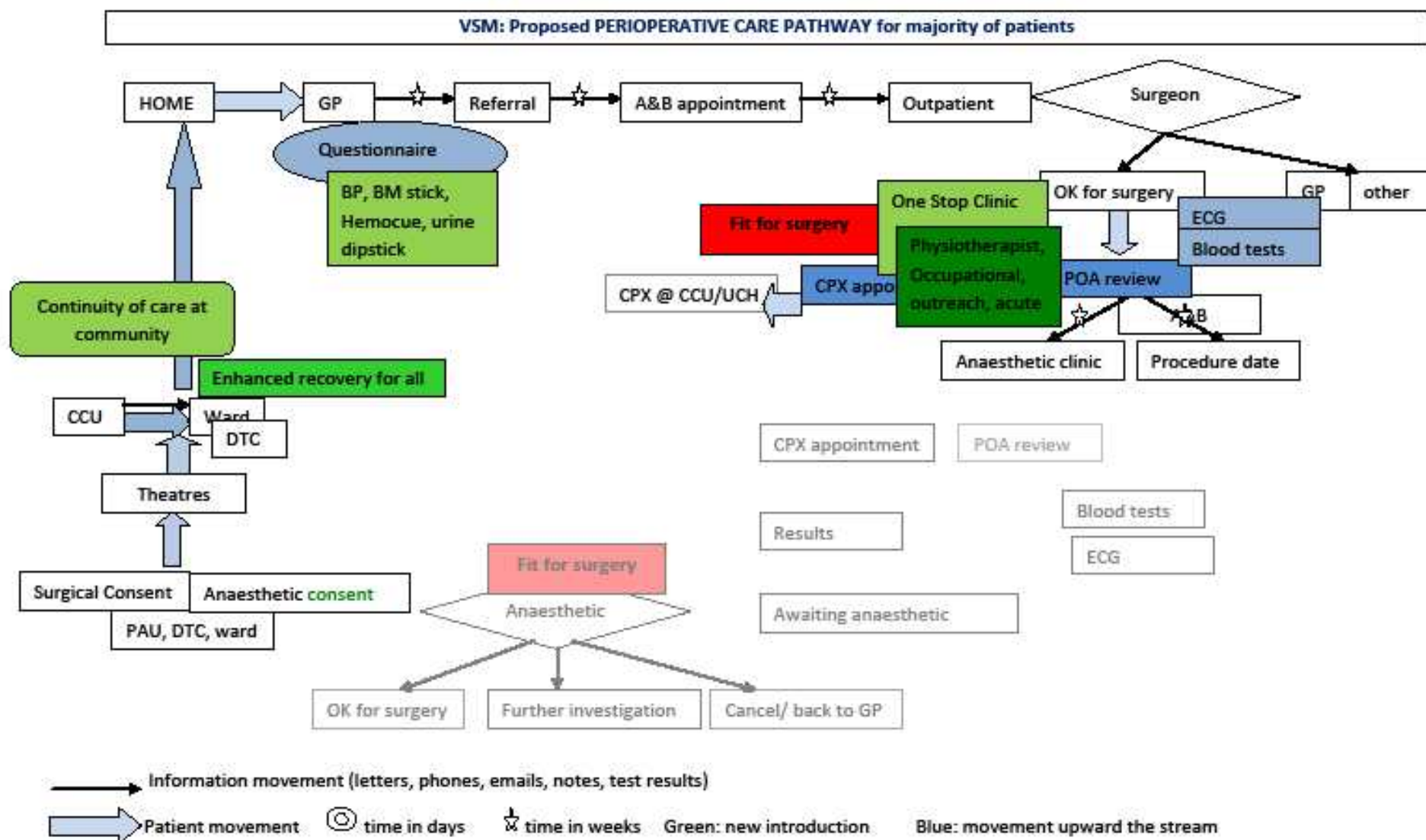
Original articles

- 110 The influence of surgery on the onset of symptomatic coronary artery disease
H. J. McFarlane, L. Girdwood, A. Bhaskar, D. Clark and N. R. Webster
115 Adjacent central venous catheters can result in immediate aspiration of infused
drugs during renal replacement therapy *K. Y. R. Kam, J. M. Mori and T. J. Wigmore*
122 The accuracy of transoesophageal echocardiography in estimating pulmonary
capillary wedge pressure in anaesthetised patients *M. M. Ali, A. G. Royse,
K. Connelly and C. F. Royse*
132 Easy and difficult nasal intubation – a randomised comparison of Macintosh vs
Airtraq® laryngoscopes *G. St. Mont, I. Biesler, R. Pförtner, C. Mohr and H. Groeben*
139 A randomised trial comparing the laryngeal mask airway Supreme™ with the
laryngeal mask airway Unique™ in children *N. Jagannathan, L. E. Sohn,
A. Sawardekar, E. Chang, K. E. Langen and K. Anderson*
145 Limited maximal flow rate of target-controlled remifentanyl infusion and induced
cough *S. K. Min, D. H. Kim, H. B. Cho, B. K. Moon and J. Y. Kim*
149 The influence of pre-admission hypoglycaemic therapy on cardiac morbidity and
mortality in type 2 diabetic patients undergoing major non-cardiac surgery:
a prospective observational study *D. Bolliger, M. D. Seeberger, G. Lurati Buse,
P. Christen, E. Seeberger, W. Ruppen and M. Filipovic*
158 Development of workplace-based assessments of non-technical skills in anaesthesia
G. V. Crossingham, P. J. A. Sice, M. J. Roberts, W. H. Lam and T. E. F. Gale



Table 2. Individual index operation types vs matched controls and incidences of myocardial infarction (MI) and acute coronary syndrome (ACS) in index surgery patients and controls studied for a minimum of 3 years after surgery. Values are number (proportion).

	Cases	Control
Joint surgery		
n	43 183	43 183
MI/ACS within 3 years	781 (1.8%)	830 (1.9%)
Subsequent MI/ACS	1900 (4.4%)	2057 (4.8%)
Deaths (all cause)	6034 (14.0%)	10 949 (25.4%)
Disc surgery		
n	1676	1676
MI/ACS within 3 years	34 (2.0%)	31 (1.8%)
Subsequent MI/ACS	83 (5.0%)	85 (5.1%)
Deaths (all cause)	205 (12.2%)	324 (19.3%)
Laparoscopic Cholecystectomy		
n	8485	8485
MI/ACS within 3 years	107 (1.3%)	141 (1.7%)
Subsequent MI/ACS	286 (3.4%)	339 (4.0%)
Deaths (all cause)	682 (8.0%)	1759 (20.7%)
Herniorraphy		
n	20 544	20 544
MI/ACS within 3 years	364 (1.8%)	519 (2.5%)
Subsequent MI/ACS	979 (4.8%)	1243 (6.1%)
Deaths (all cause)	2802 (13.6%)	5525 (26.9%)
Upper GI endoscopy		
n	109 350	109 350
MI/ACS within 3 years	2153 (2.0%)	2268 (2.1%)
Subsequent MI/ACS	5146 (4.7%)	5621 (5.1%)
Deaths (all cause)	27 203 (24.9%)	28 618 (26.2%)



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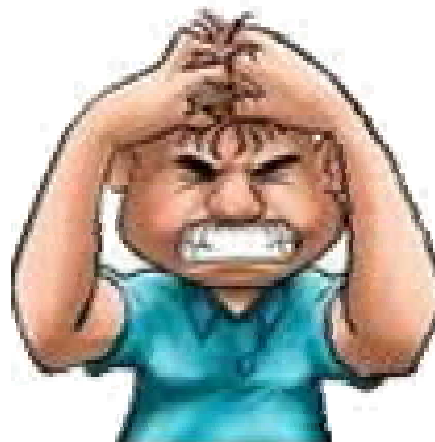
Your Questions on Lean

We have low morale and no one is willing to change, what can I do?

Can Lean fail?, and for what reason?

What the one thing that makes the biggest difference?

What is the impact on CCG on end to end transformation?



We have no budget to start Lean – what can we do?



What's Next?

- **Today's presentation and feedback survey sent out by email within 48 hours**
- **The Next Lean London Forum will be held in September 2012.**
 - Register at www.leanlondon.org.uk
 - We will send out reminders to all participants from today
 - We have a Lean Midland Forum on 12 June 2012 taking place in Birmingham. Register at www.leanmidland.org.uk
 - If you'd like to take up one of our presentation slots, please do let us know. We are keen to hear from Community Trust and GP Groups
- **Find us on Linked In and Twitter - LeanNHS**





Big Thanks To Our Presenters

Suzanne Horobin

Mathew Diggle

Ahmed Chekairi

..and to you all for attending





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