
Evidence for sequential effects in reading from a million women's mammograms in the UK

Professor Sian Taylor-Phillips

Professor of Population Health

University of Warwick

David Jenkinson, Matthew Wallis, Aileen Clarke



Contents

1. Vigilance hypothesis
2. The CO-OPS trial
3. CO-OPS Results
4. Longer Batch results
5. Summary of findings and implications for future work



1. Vigilance Hypothesis



2. The CO-OPS Trial

Changing Case Order to Optimise Patterns of Performance in Screening

- Is there a vigilance decrement?
- Does reverse reading help?

Taylor-Phillips et al. JAMA. 2016;315(18):1956-1965.



The CO-OPS Trial

Pragmatic randomised controlled trial of a software intervention to change case order so that any vigilance decrement will occur for the first and second readers when examining different cases

Intervention ↓↑ or ↑↓

Control ↓↓ or ↑↑

SIRE Film Results for MU001 Wednesday 4 March 2009

Print Screen Film Result Client List What's This? Help

Reverse Order New

Time	SX	Name	Readers
15:03	KKE035002	BRADSHAW, CLAIRE	
14:49	KKE034999	HENSHAW, PAM	
14:42	KKE034998	LOINS, JONI	
14:35	KKE034996	SAVILLE, MARGE	
14:28	KKE034995	FULLER, BRENDA MAY	
14:14	KKE030140	GREY, B	
14:07	KKF030138	SPFARS, DIANE	



	<u>Control</u> <u>arm</u>	<u>Intervention</u> <u>arm</u>
Reader 1: Forwards		
Reader 2: Forwards		
Both readers together		
	Number of disagreements = Low Recall Rate = Low Cancer Detection Rate = Low	Number of disagreements = High Recall Rate = High Cancer Detection Rate = High



The CO-OPS Trial

Participation:

- 46 English Breast Screening Centres completed the trial
- One year of data collection



The CO-OPS Trial

Primary Analysis

- Multi-level model of the predictors of cancer detection rate

Secondary Analyses

- Multi-level model of the predictors of recall rate and rate of disagreements
- Patterns of performance with time since a break



Randomized
(n=1,207,633)
in 37,724 batches

Intervention Arm

(n=603,528) in 18,797 batches

Received allocated intervention
(n=524,971, 87%)

Lost to follow-up 0.07%

Analysed
(n=596,642, 98.9%)

Control Arm

(n=604,105) in 18,927 batches

Received allocated intervention
(n=560,760, 93%)

Lost to follow-up 0.06%

Analysed
(n=597,505, 98.9%)

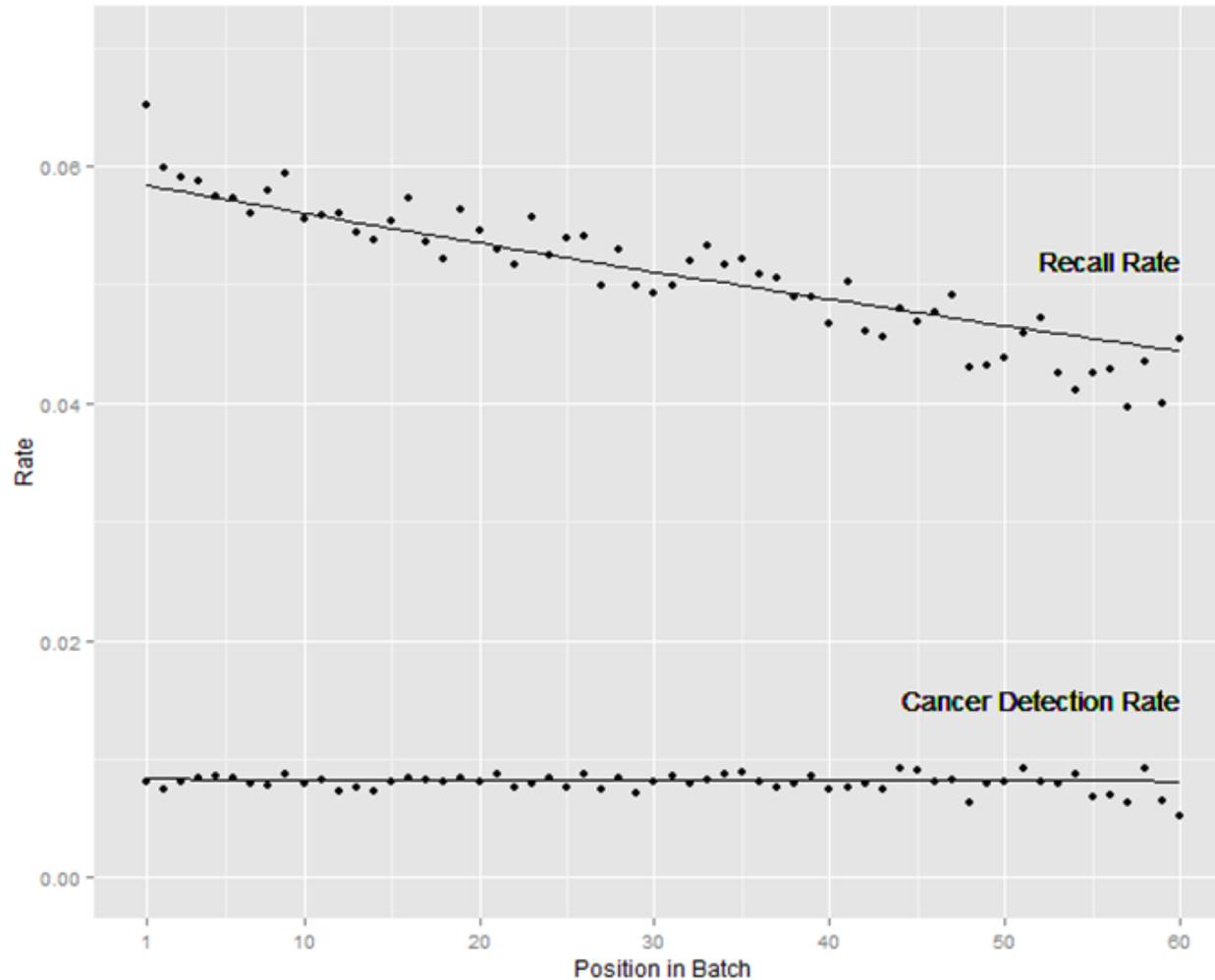


3. CO-OPS trial results

	Intervention	Control
Cancer Detection Rate (95%CI)	0.89% (0.86% to 0.91%)	0.88% (0.85% to 0.90%)
Recall Rate (95%CI)	4.14% (4.09% to 4.19%)	4.17% (4.12% to 4.22%)
Rate of disagreements (95%CI)	3.43% (3.39% to 3.48%)	3.48% (3.43% to 3.53%)



Patterns of recall: Single reader

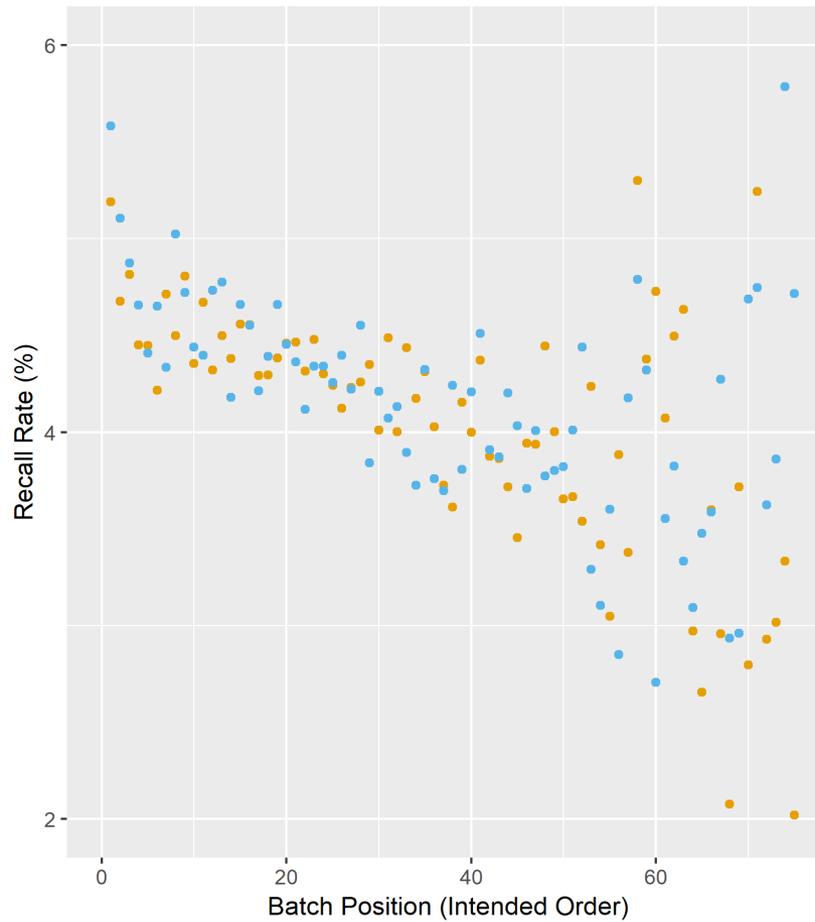


Is this pattern real?

- Reading order isn't random
- One batch = one day on one machine (30-60)
- Order is alphabetised by GP practice
- Women can rearrange appointment time but rare



Patterns of recall: Single reader



Direction
● Forward
● Reverse

Direction ● Forward ● Reverse



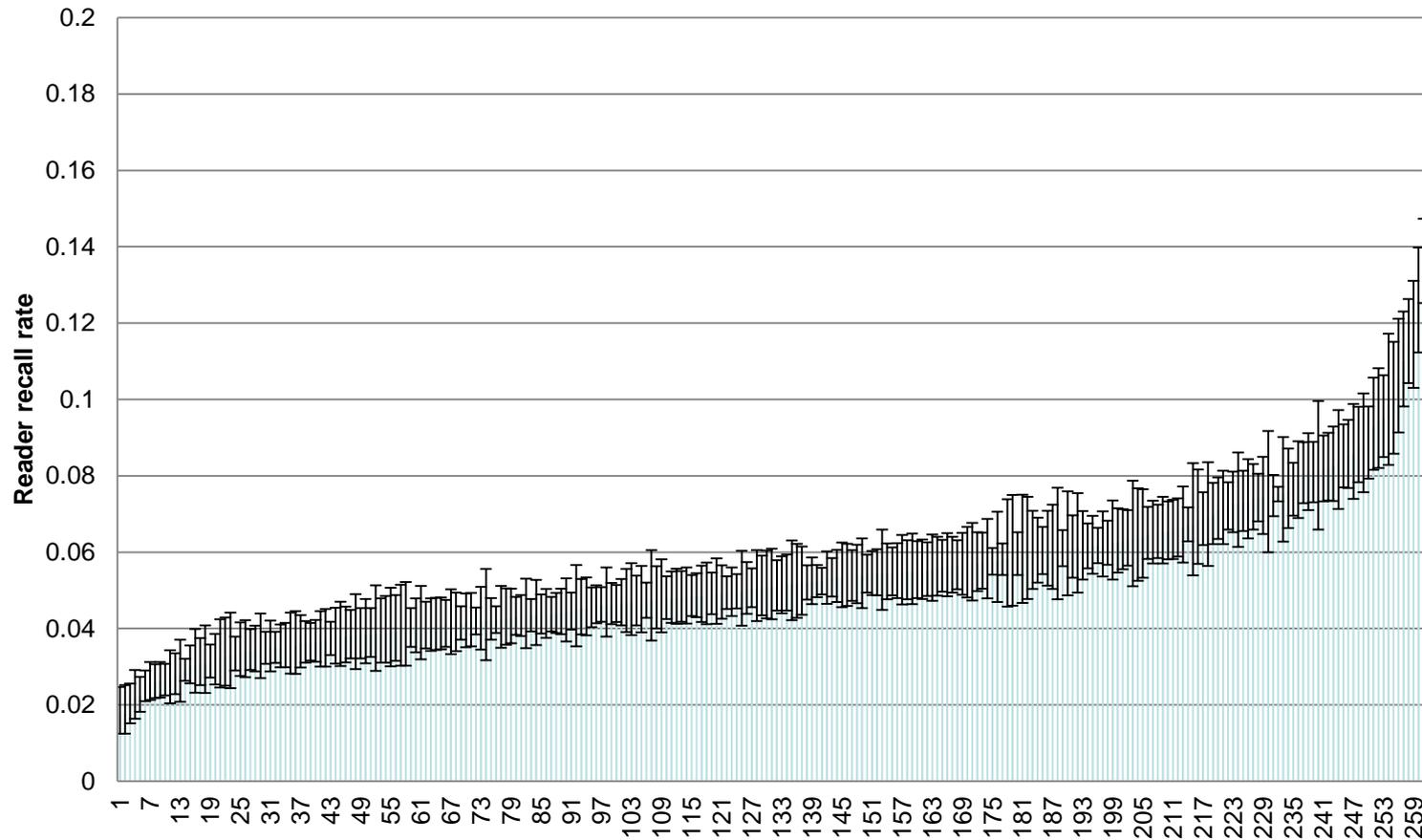
Why doesn't it change overall recall rate?

- Film reader variability?
- Arbitration of discordant cases removing excess recalls at beginning?
- Changes to who is recalled but not number recalled?



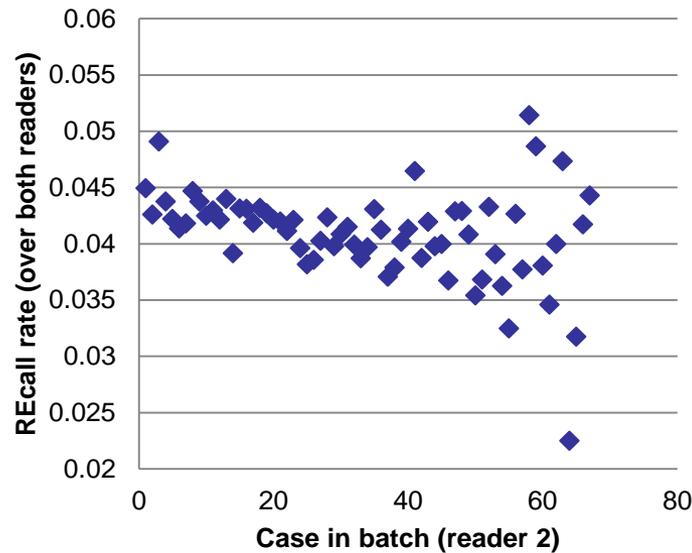
Film Reader Variability

Recall rate by reader (only including those who read >1000)

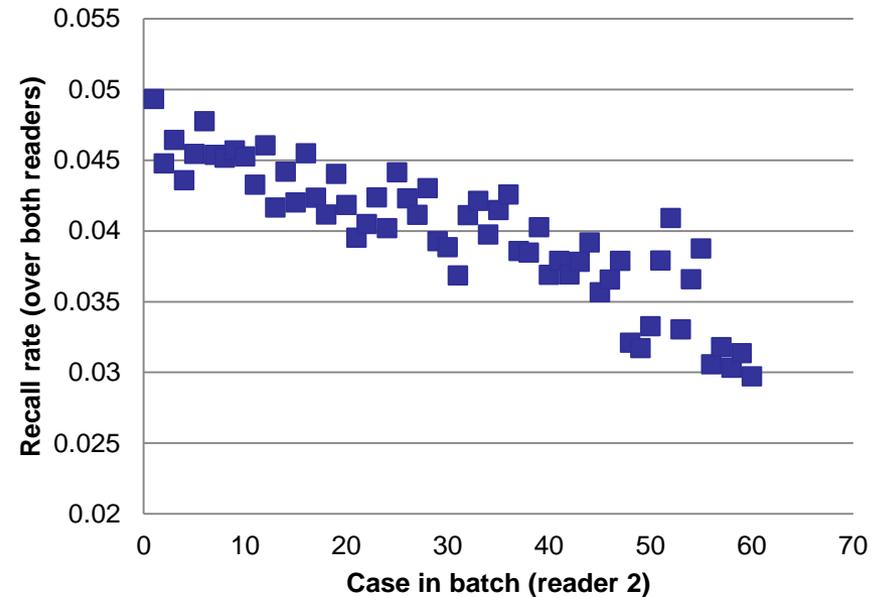


Recall rate combining both readers

recall rate intervention



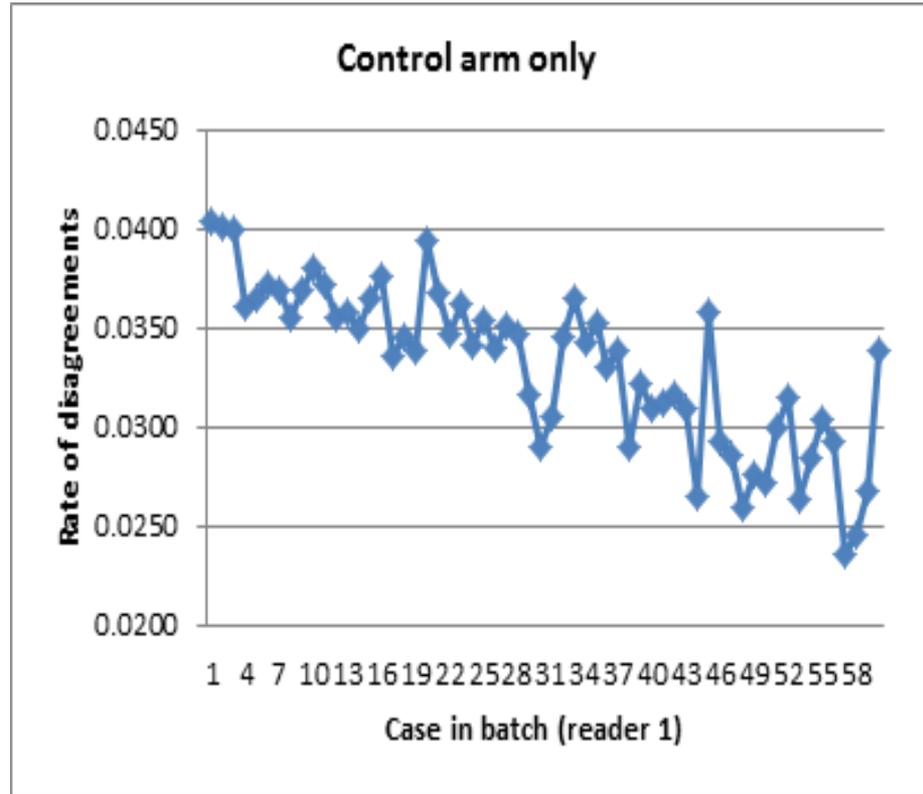
recall rate control



Arbitration of discordant cases removing excess recalls at beginning? No
Changes to who is recalled but not number recalled? Maybe



Disagreements



Excess of recall at beginning of the batch, in the intervention arm this is spread over different cases



4. Analysis of longer time on task

- What about sessions longer than 60 cases? UK film readers often examine several batches back to back
- Analysed patterns of performance over time for each reader
 - Time stamps for decision on each case
 - Break defined as 10, 20, 60, 180, or 480 minutes without a decision
 - Excluded cases moved from intended order
 - Excluded first case after break



Follow-on analysis

- Reader 1 analysed, because reader 2 not blinded
- Multi-level models, levels: woman, reader1, centre
- Adjusted for woman's age and whether she has previously attended screening
- Linear models with knots at 20 and 40 cases
- Distribution: logistic for recall and cancer detection, gamma for time taken



Follow-on analysis

- Outcomes
 - Reader 1 recall yes/no
 - Reader 1 cancer detected yes/no
 - (Reader recalls and cancer detected in follow up biopsy)
 - Reader 1 time taken to examine the case
 - (Time stamp at end of case minus time stamp at end of previous case)
 - Sensitivity and specificity of reader 1
 - (reference standard is cancer detected at screening or symptomatically in 3 years following screening)



Summary of findings

- There appear to be an excess of recalls at the beginning of a reading session
- The reverse reading intervention doesn't affect recall rate overall, but does appear to affect who is recalled.
- Reading speed appears to increase and recall rate appears to decrease with time on task
 - Good fit with evidence that batch reading improves specificity.
- Cautious interpretation of post-hoc analyses of large observational dataset



Next Steps in Research

- What do we want to detect in breast cancer screening?
- Characteristics of cancers/pre-cancers detected.
- Maximise morbidity and mortality benefits, and minimise overdiagnosis.
- What are the long term outcomes after previous changes to breast screening?
 - Two readers
 - Reader test threshold (recall rate)
 - Age of eligibility



Further Information

Email: s.taylor-phillips@warwick.ac.uk

Website: <https://warwick.ac.uk/fac/sci/med/research/hscience/pet/screening/breast/>

Twitter: @siantphillips2



This work was funded by an NIHR Career Development Fellowship for Professor Taylor-Phillips. The views expressed are those of the author(s) and not necessarily those of the NHS, NIHR, or Department of Health.

