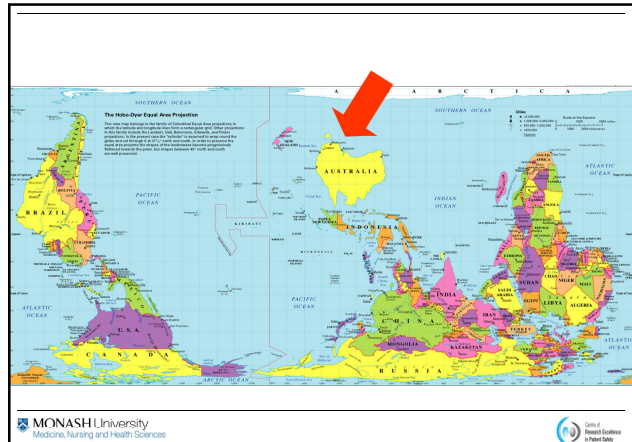


Application of Variable Life Adjusted Displays (VLAD) on Victorian Admitted Episodes Dataset (VAED)

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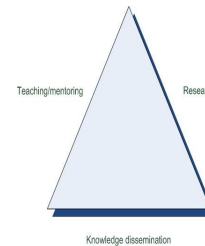
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Objectives of the CRE-PS

- To establish a national Centre of Research Excellence that will promote and develop resources to improve patient safety that will lead to:
 - Better identification of factors that affect patient safety
 - More effective use of data to monitor quality of care
 - Better understanding of human factors, organisational factors and system deficiencies leading to error
 - Use of clinical simulation to provide a safe environment for research, training and education
 - Development of mechanisms for ensuring uptake of research findings
 - Training of new patient safety researchers

- Research** to strengthen the evidence base for developing and implementing system improvements to reduce adverse events and improve safety and quality
- Teaching/ mentoring** to improve patient safety at clinical level and develop new researchers
- Knowledge dissemination** to enhance patient care



Introduction

- Control charts for quality control
- In-control vs out-of-control states for systems
- Many types e.g.
 - VLAD
 - Shewhart
 - Cumulative Sum (CUSUM)
 - Exponentially Weighted Moving Average (EWMA)
 - (Resetting) Sequential Probability Ratio Tests (SPRT and RSPRT)

Background

- Centre for Research Excellence in Patient Safety (CRE-PS) evaluated the VLAD methodology for Department of Human Services (DHS) Victoria
- In-hospital mortality for acute myocardial infarction (AMI), heart failure (HF) and stroke used for proof of concept
- Logistic regression risk model for each condition based on Queensland Hospital Admitted Patient Data Collection (QHAPDC) models

Background

Data management and initial analysis

↓
Logistic regression

↓
VLAD

↓
VLAD limits

↓
Implementation plan

VAED

- Victorian hospital admissions
- Over 3.9 million between 1st July 2004 – 30th June 2007
- De-identified hospital code
- Gender, age-group, admission date, separation date
- Primary admission diagnosis and up to 39 other co-morbidities captured as International Classification of Diseases (ICD 10) codes e.g.
 - AMI – I21 or I22, HF – I50, stroke I61-I64
- Separation mode e.g.
 - "D" = death, "T" = transfer

Separations - AMI

Criterion	N(%)
Total separations 2004-7	32550(100)
LOS <4 days and mortality = no	14814(45.5)
LOS >30 days	418(1.28)
Not admitted through hospital emergency department	8830(27.1)
Transfers out	10616(32.6)
Age <30 years	67(0.21)
Age ≥85 years	4661(14.3)
Eligible separations	8924(27.4)

Separations - HF

Criterion	N(%)
Total separations 2004-7	27389(100)
No overnight stay	3786(13.8)
LOS >30 days	649(2.4)
Not admitted through hospital emergency department	6293(23.0)
Transfers out	4510(16.5)
Age <30 years	141(0.5)
Age ≥85 years	7876(28.8)
Eligible separations	11124(40.6)

Separations - Stroke

Criterion	N(%)
Total separations 2004-7	22706(100)
LOS <4 days and mortality = no	6239(27.5)
LOS >30 days	1432(6.3)
Not admitted through hospital emergency department	4641(20.4)
Transfers out	8071(35.6)
Age <30 years	311(1.4)
Age ≥85 years	4768(18.4)
Carotid endarterectomy	102(0.5)
Eligible separations	5857(25.8)

Mortality - AMI

Year	N	%	Deaths	%	Mortality risk	Risk ratio
2004-5	2999	33.6	338	31.9	11.3%	1.00
2005-6	2977	33.4	371	35.0	12.5%	1.11
2006-7	2948	33.0	350	33.1	11.9%	1.05
2004-7	8924	100	1059	100	11.9%	-

Mortality - HF

Year	N	%	Deaths	%	Mortality risk	Risk ratio
2004-5	3598	32.3	213	34.6	5.9%	1.00
2005-6	3683	33.1	194	31.5	5.3%	0.89
2006-7	3843	34.6	209	33.9	5.4%	0.92
2004-7	11124	100	616	100	5.5%	-

Mortality - Stroke

Year	N	%	Deaths	%	Mortality risk	Risk ratio
2004-5	1922	32.8	540	30.9	28.1%	1.00
2005-6	1966	33.6	609	34.8	31.0%	1.10
2006-7	1969	33.6	599	34.3	30.4%	1.08
2004-7	5857	100	1748	100	29.8%	-

Comorbidities - AMI

Comorbidity	N	%	Deaths	%	Mortality risk	Risk ratio
Hypotension/shock	1393	15.6	437	41.3	31%	3.80
Dysrhythmias	2241	25.1	553	52.2	25%	3.26
CVD	179	2.0	64	6.0	36%	3.14
HF	1993	22.3	403	38.1	20%	2.14
Renal failure	1715	19.2	357	33.7	21%	2.14
Dementia	189	2.1	46	4.1	24%	2.10
Malignancy	196	2.2	43	4.1	22%	1.88
Diabetes	2876	32.2	323	30.5	11%	0.92
Hypertension	4693	53.6	449	42.4	10%	0.66
All separations	8924	100	1059	100	12%	-

Comorbidities - HF

Comorbidity	N	%	Deaths	%	Mortality risk	Risk ratio
Septicaemia	96	0.9	41	6.7	43%	8.19
Hypotension/shock	1031	9.3	195	31.7	19%	4.53
CVD	129	1.2	27	4.4	21%	3.91
Renal failure	3599	32.4	387	62.8	11%	3.53
Dementia	267	2.4	42	6.8	16%	2.98
Lower limb or decubitus ulcer	502	4.5	71	11.5	14%	2.76
Malignancy	307	2.8	43	7.0	14%	2.64
LRTI/ Influenza	1587	14.3	177	28.7	11%	2.42
IHD	2256	20.3	207	33.6	9%	1.99
Dysrhythmias	2720	24.5	223	36.2	8%	1.75
Hypertension	4605	41.4	218	35.4	5%	0.78
All separations	11124	100	616	100	6%	-

Comorbidities - Stroke

Comorbidity	N	%	Deaths	%	Mortality risk	Risk ratio
Septicaemia	72	1.2	49	2.8	68%	2.32
HF	229	3.9	145	8.3	63%	2.22
LRTI/ Influenza	250	4.3	130	7.4	52%	1.80
Malignancy	182	3.1	94	5.4	52%	1.77
Renal failure	543	9.3	216	9.3	40%	1.38
All separations	5857	100	1748	100	30%	-

Multiple logistic regression - AMI

2004-7 (N=8924)	OR	95% CI	P
Sex			
Male (reference)	1.00		
Female	1.21	1.04 – 1.40	0.013
Age - years			
30 – 54	0.55	0.37 – 0.81	0.002
55 – 59	0.54	0.34 – 0.85	0.008
60 – 64 (reference)	1.00	-	-
65 – 69	1.17	0.83 – 1.66	0.378
70 – 74	1.52	1.10 – 2.10	0.011
75 – 79	2.21	1.63 – 2.98	<0.0001
80 - 84	2.95	2.19 – 3.97	<0.0001

Multiple logistic regression – AMI (cont.)

2004-7 (N=8924)	OR	95% CI	P
Comorbidities			
Hypotension/shock	3.99	3.42 – 4.65	<0.001
Dysrhythmias	3.03	2.62 – 3.50	<0.001
Cerebrovascular disease	3.91	2.74 – 5.58	<0.001
HF	1.30	1.11 – 1.52	0.001
Renal Failure	1.62	1.37 – 1.91	<0.001
Dementia	1.52	1.04 – 2.23	0.031
Malignancy	2.16	1.48 – 3.16	<0.001
Diabetes	0.98	0.83 – 1.16	0.833
Hypertension	0.49	0.42 – 0.57	<0.001

Multiple logistic regression – AMI (cont.)

2004-7 (N=8924)	OR	95% CI	P
Year of separation			
2004-5 (reference)	1.00	-	-
2005-6	1.09	0.91 – 1.29	0.347
2006-7	1.09	0.92 – 1.30	0.311

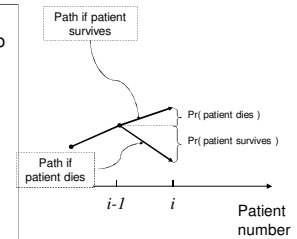
VLAD

- Plots cumulative expected (E) – observed (O) events
- E calculated from 2004-5 logistic models
- From 2005 onwards
- Calculated by de-identified hospital (N=11)

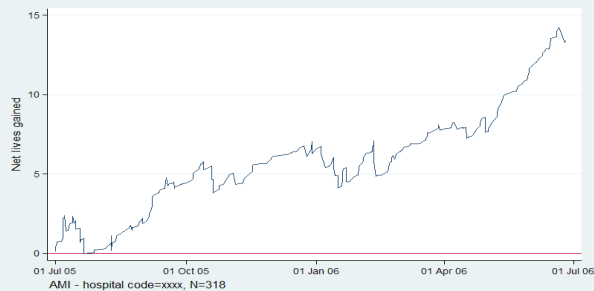
$$V_n = \sum_{i=1}^n E_i - \sum_{i=1}^n O_i$$

VLAD

- Type of quality control chart
- Counts excess of expected to observed events
- Expected events calculated using:
 - Case mix adjusted logistic regression model
 - Data from 2005-07 only
- 'Up' line – performance improving
- 'Down' line – performance declining
- 'Flags' identify marked changes in performance



VLAD



VLAD - Limits

- Upper and lower risk adjusted CUSUM limits
- ρ – odds ratio between risk of mortality under the alternative and null hypotheses
- h – control limit signifying when the CUSUM signals
- Resetting of limit with signal
- Methodology as per Sherlaw-Johnson¹

¹Sherlaw-Johnson C. (2005) A Method for Detecting Runs of Good and Bad Clinical Outcomes on Variable Life-Adjusted Display (VLAD) Charts. Health Care Management Science, 8: 61-65.

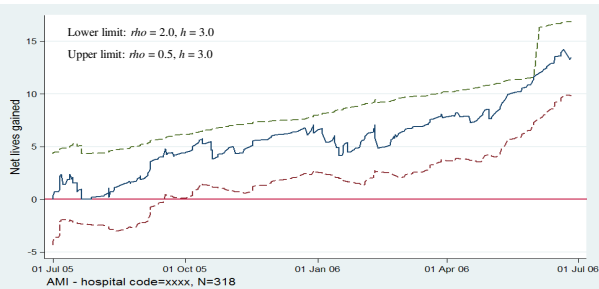
VLAD - Limits

- For the lower limit the CUSUM of the n th observation (C_n) with the corresponding weight is given by:
 - $C_0 = 0$ and $C_n = \max \{C_{n-1} + W_n, 0\}$
- For the upper limit the CUSUM of the n th observation (C_n) with the corresponding weight is given by:
 - $C_0 = 0$ and $C_n = \min \{C_{n-1} - W_n, 0\}$
- where
 - $W_n = O_n \log \rho - \log(1 + (\rho - 1)E_n)$

VLAD - Limits

- The lower VLAD limit (L_n) can then be calculated by the expression
 - $L_n = V_n + (C_n - h) / \log \rho$
- The upper VLAD limit (L_n) can then be calculated by the expression
 - $L_n = V_n - (C_n + h) / \log \rho$
- Should V_n intersect with L_n , the limit is reset to Z_n by
 - $Z_n = L_n + h / \log \rho$

VLAD - Limits



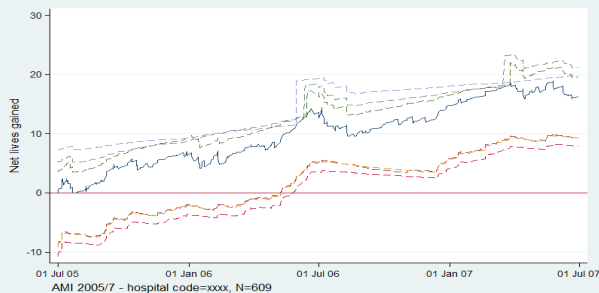
VLAD - Limits

- ARL = Average run length to false alarms
- Coory et. al.²

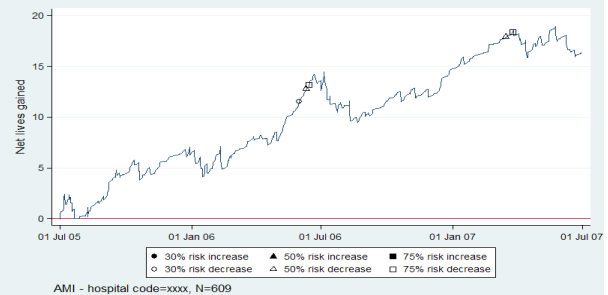
rho	improved performance			worse performance		
	0.70	0.50	0.25	1.30	1.50	1.75
h	2.6	3.6	4.9	2.8	3.7	5.0
ARL	229	682	2447	264	834	3118

²Coory M, Duckett S, Sketcher-Baker K. (2008) Using control charts to monitor quality of care with administrative data. International Journal for Quality in Health Care, 20: 31-39

VLAD - Multiple limits



VLAD - Multiple limits



VLAD - Signals

2005-7	Risk Decrease			Risk Increase		
	30%	50%	75%	30%	50%	75%
AMI						
Total number of signals	4	5	3	1	1	1
Hospitals signaling at least once	27%	27%	18%	9%	9%	9%
HF						
Total number of signals	3	5	5	1	1	1
Hospitals signaling at least once	18%	36%	36%	9%	9%	9%
Stroke						
Total number of signals	5	5	4	11	11	11
Hospitals signaling at least once	36%	36%	36%	45%	45%	45%

Implementation

- Upwards of 30 outcomes from VAED data
- Development of models by bootstrapping techniques
- Possibility of rolling logistic model
- Monthly availability of data with 3 month lag
- Fine tuning of ρ and h
- Validation of signals
- Response to signals and will this lead to change
- Initial trial with one or two well known outcomes in Victorian hospitals e.g. AMI

Summary

- VLAD may be a clinically appealing measure of quality control
- Monitoring will give an indication of potential signals to investigate
- False positives will occur
- Further work needed on methodology, implementation and validation for Victorian hospitals

Acknowledgements

- Department of Human Services, Victoria
- Associate Professor Michael Coory
- CRE-PS
- Department of Epidemiology and Preventive Medicine, Monash University

Multiple logistic regression - AMI

Year	N	Area under ROC curve	Hosmer-Lemeshow $\chi^2(8)$	P
2004-5	2999	0.8344	13.77	0.0880
2005-6	2977	0.8200	11.13	0.1943
2006-7	2948	0.8147	39.37	<0.0001
2004-7	8924	0.8207	43.94	<0.0001

Multiple logistic regression - HF

Year	N	Area under ROC curve	Hosmer-Lemeshow chi2(8)	P
2004-5	3598	0.8260	8.90	0.3506
2005-6	3683	0.8259	3.93	0.8631
2006-7	3843	0.7919	8.48	0.3881
2004-7	11124	0.8109	8.86	0.3540

Multiple logistic regression - Stroke

Year	N	Area under ROC curve	Hosmer-Lemeshow chi2(8)	P
2004-5	1922	0.6768	6.16	0.5208
2005-6	1966	0.6529	3.18	0.7862
2006-7	1969	0.6727	2.76	0.7370
2004-7	5857	0.6651	20.32	0.0092