CENTRAL VENOUS CATHETER ASSOCIATED BLOOD STREAM INFECTION IN NEONATAL INTENSIVE CARE UNIT - PREDICTION AND PREVENTION

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Introduction

Central venous catheters (CVCs)

 Common form of vascular access in the NICU - Parenteral nutrition, complex care

Major source of late onset sepsis and the most common cause of hospital acquired infection

Introduction

Implications of neonatal CVC-BSI

- Mortality (4% to 20%) *
- Morbidity (increased ventilation and respiratory supports)
- Prolonged hospital stay (average 7 days)
- Poor growth and adverse neurodevelopmental outcome at 2 years of age

Importance in identifying modifiable risk factors to reduce infection rate

Saint S, Veenstra DL, Lipsky BA. The clinical and economic consequences of nosocomial central venous catheter- related infection: are antimicrobial catheters useful? Infect Control Hosp Epidemiol. 2000; 21(6):375-380

AIM

Determine the incidence of CVC-BSI in our NICU.
 Identify potential factors contributing to CVC-BSI.

Review the effectiveness of implementing a CVC-BSI prevention bundle.

Study methods

Definitions

CVC - Catheters inserted into a central vein - terminated at or close to the heart or in one of the great vessels (inferior vena cava, superior vena cava, brachiocephalic, internal jugular vein or subclavian vein)

- Laboratory confirmed CVC- associated blood stream infection (LC CVC-BSI) -Positive blood cultures while the CVC was insitu and the cultured organism was not related to infection from an alternative source.
- Clinical CVC- BSI development of temperature instability, tachypnoea, apnoea, lethargy, abdominal distension, an increase in C reactive protein (CRP) level, or subtle nonspecific signs in keeping with probable sepsis, but in the absence of a positive blood culture and clinical signs attributable to any particular organ system.

Design

- Retrospective analysis of prospectively collected data over 9 years (2006–2014)
- Pooled data Analysed for LC CVC BSI and clinical CVC BSI pooled together

Inclusion criteria:

- All CVCs inserted
- First hospitalisation to NICU, Royal Hobart Hospital

Exclusion

Umbilical line

CVC-BSI rate per 1000 catheter days - number of CVC-BSI divided by the number of catheter days and multiplying the result by 1000.

Variables studied

- Birth weight
- Intrauterine growth restriction
- Gestational age

- Type of catheter (PICC, 3F to 5F)
- Number of lumens
- Site of insertion
- Operator seniority
- Line adjustment after insertion to locate the tip in appropriate position
- Breaches in catheter to deliver medications
- Nature of Infusate
- Presence of previous catheters
- Presence of multiple CVC's concurrently
- Underlying disease

Methodology

- Baseline data extracted from NICU data base
- Study Parameters from DMR and charts review
- Microsoft excel and NCSS software for analysis
- Continuous data Median, interquartile ranges Mann Whitney test
- Categorical data- Chi square test
- Associations Logistic regression

Univariate analysis done- Association of parameters with catheter infection

Multivariate analysis model- for factors found predictive on univariate analysis

P value < 0.05 taken significant

CVC BSI rates

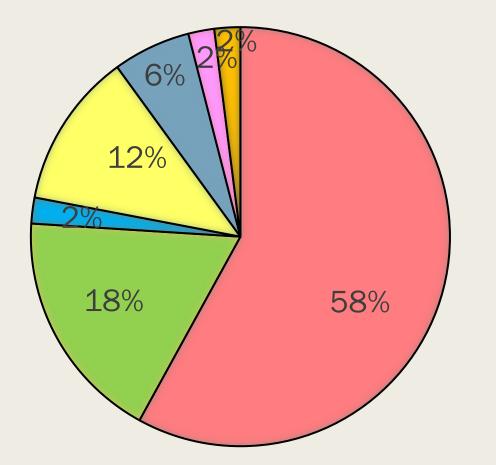
- Pooled 19.2/1000 catheter days
- LC CVC-BSI 8.2/ 1000 catheter days

Table 1: Patient demographics

	CVC-BSI	No CVC-BSI	P value
Number of patients (n)	119	534	
Birth weight (g)	930* (860-1085)	1200* (901-1880)	<0.01
Gestation (weeks)	27* (26-28)	29* (27-34)	<0.01
Males (n)	68 (57%)	295 (55%)	0.7
Females(n)	51 (42%)	239 (45%)	
Male: Female	1.3:1	1.2:1	

Table 1: Demographics of neonates admitted to the NICU who had CVC inserted. Data presented as median (IQR) or n (%). CVC-BSI: Central venous catheter associated blood stream infection. P value considered statistically significant if P < 0.01.

Figure 1: Organisms cultured on LC CVC-BSI



CONS
Staph Aureus
Enterococcus
Enterobacter Species
Candida
Pseudomonas
Acinetobacter

Table 2: Predictive factors for CVC-BSI

	OR (95% CI)	P value
Gestation < 28 weeks	2.9 (1.8-4.7)	<0.01
Saphenous vein insertion	1.99 (1.3-3.09)	<0.01
Parenteral nutrition	2.3 (1.5-3.8)	<0.01
Multi-lumen CVCs	5.6 (2.8-11.4)	<0.01
Underlying GIT disease	2.1 (1.3-3.5)	0.05

Table 2: Factors examined and found predictive of neonatal CVC-BSI. Results presented as odds ration (95% confidence interval). GIT: Gastrointestinal tract. P value considered statistically significant if P < 0.01

Table 3: Factors not predictive of CVC-BSI

			P value
Duration of CVC in	≤10 days	84/442 (19%)	0.4
situ	>10 days	32/179 (17.9%)	
CVC adjustment		26/141 (18.4%)	0.6
CVC breaches		20/117 (17%)	0.7
	24 G	8/46 (17.4%)	0.7
CVC size	28 G	104/486 (21.4%)	0.3
	Others	7/52 (13.5%)	
Multiple CVCs		11/60 (18.3%)	0.9
Personnel inserting	SMO	64/397 (16.1%)	
CVC	JMO	55/256 (21.5%)	0.08

Table 3: Factors examined and demonstrated to be not predictive of neonatal CVC-BSI. G: gauge. Multiple CVCs reflects the presence of multiple CVC inserted in a neonate at any given time. SMO: Senior Medical Officer. JMO: Junior Medical Officer. P values are considered statistically significant if P < 0.01. **Results in context**

Operator related factors

Veins used

- Saphenous vein
- Below diaphragm veins (femoral and saphenous) more likely to be infected compared to above diaphragm venous lines

Operator seniority

- No difference in infection risk
- Senior registrar, consultants and registrars

Results in context

Catheter related

- Multilumen CVC's
- Infusate Parenteral nutrition (?Lipids in the TPN predisposes to infection)

Results in context

Patient related factors- Underlying GIT

GIT disease predisposed to infection

- Definite NEC, Gastroschisis, Omphalocele, intestinal obstruction

CVC prevention bundle

Study methods

CVC prevention bundle

Data for the subsequent 12-months was prospectively collected following the introduction of a CVC-BSI prevention bundle and the infection rates were compared over the two epochs.

CVC insertion by designated team of trained physicians who have successfully completed

- CVC training package
- Aseptic non-touch technique (ANTT)
- Hand hygiene modules, and
- Accreditation by a neonatal consultant after supervision of two satisfactory CVC insertions

Study methods

CVC prevention bundle

- CVC insertion checklist
- A dedicated nurse observing CVC insertion to ensure asepsis
- Appropriate structural barriers around the patient (e.g. screens)
- Dedicated CVC insertion trolley and
- Standardized dressing, redressing and CVC access procedures

CVC BSI rates

- Pooled 19.2/1000 catheter days
- LC CLABSI 8.2/ 1000 catheter days

After CVC bundle – 1.2 per 1000 catheter days

CVC BSI rates across countries

- CVC BSI rate 1.7 to 2.3 cases / 1000 catheter days*
- Spain 6.7/1000
- Brazilian and Latin American hospitals (1.6 to 44.6/1000 catheter days)
- Brazilian study (15/1000 in 2010 to 13/1000 in 2011)**

* Safdar N, maki DG.Risk of catheter-related blood stream infection with peripherally inserted central venous catheters used in hospitalized patients. Chest.2005;128(2):489-495

**Rosado V, Camargos P, Clemente W, Romanelli R. Incidence of infectious complications associated with central venous catheters in pediatric population. Am J Infect Control 2013;41:e81-e84

Conclusions

- 1. Below diaphragm central line insertion preferably avoided
- 2. Judicious use of multilumen catheters
- 3. CVC prevention bundles
- 4. CVC BSI Surveillance

Acknowledgements

- Prof Peter Dargaville (supervisor)
- Dr. Kathleen Lim
- Dr. Sanoj Ali
- Dr.Hamish Jackson
- Dr. Tony DePaoli
- Dr. Sheena Kaul
- Pacific Venant

- Gemma Plottier
- Dr. Tim Jones
- Karen Butterly
- Kate Herbert
- Dr. Reeshma Pattan
- Michael Bourne



CLABSI rates (Per 1000 PICC line days)				
CIABSI rate for LC confirmed	8.2			
infection				
CLABSI rate for LC and clinical				
infection				
CLABSI rate for SGA babies (11.3			
Combined)				
CLABSI rate in SGA babies LC	5.3			
confirmed				
SGA babies in less than 28 weeks (8			
Combined)				
SGA babies in less than 28 weeks(2			
LC confirmed)				

Central venous catheter analysis results

Veins used

Saphenous	54	170	224	24.1	0.005
Antecubital fossa	33	207	240	13.8	0.02
Femoral	12	47	59	20.3	
Jugular	1	9	10	10	
Forearm	14	49	63	22	
Scalp	5	48	53	9.4	
Others(Scalp,subclav ian,wrist)	0	4	4	0	

CVC Manipulation

Line note manipulated	93(78 %)	458(78 %)	551	16.9	0.6
Line manipulated	26(22 %)	115(22 %)	141	18.4	

SGA					
SGA	17	129	146	11.6	0.02
AGA	102	405	507	20.1	

SGA in less than 28 weeks					
SGA less than 28	12	41	53	22.6	0.002
weeks SGA more than	5	88	93	5.4	
28 weeks					

Infusate						
TPN/IL	78	274	352	22.2	0.007	
Multiple infusions	41	260	301	13.6		

CVC breaks						
No Line breaks	99	437(82 %)	536	18.5	0.7	
Line breaks	20	97(18%)	117	17		
Total	119	534	653			
Total line breaks - 17.9						

	Duration of line							
Median		7(6-9)	8(5-12)			0.4		
	Less than 5	39	146	185	21			
	6 to 10	45	212	257	17.5	(for lines		
	11 to 15	17	98	115	14.8	less than 10		
	16 to 20	15	49	64	23.4	versus more		
	More than 20	3	29	32	9.4	than 10 and also when analyzed less than 5 versus more than 5)		

Number of lumen for CVC						
1	99	504(94%)	603	16.4	0.00	
More than 1	20	30(6%)	50	40		
Total	119	534	653			

Previous PICCS								
Previous PICC	54	59	113	47.8	0.00			
None 65 312 377 17.2								

Personnel inserting the PICC						
Senior Registrar	21	147	168	12.5	0.04	
Consultant	43	186	229	18.8	0.6	
Others	47	183	230	20.4		
Total	111	516	627			
Not recorded	8	18	26			

Type of catheter					
24G	8	41	46	17.4	0.7
+28G	104	444	486	21.4	0.3
Others	7	49	52	13.5	
Total	119	534	653		

Underlying severe disease						
GIT	45	106	151	29.8	0.00	
Non GIT	74	428	502	14.7		

Multiple CVC's							
No	108	485 (91%)	593	18.2	0.9		
Yes	11	49(9%)	60	18.3			
Total	119	534	653				

Outcome						
Normal	60(50.4 %)	391(73%)	451	13.3	0.00	
Abnormal (Dev delay, Stoma)	59(49.6 %)	143(27%)	202	29.2		
Total	119	534	653			
Death	14	24	38	36.8	0.00	

Total Saphenous vein %	34.3%						
Total Antecubital fossa %	36.7%						
Total line manipulated – 21.6%							
Management basis							
Clinical	35(70%)						
Clinical and septic blood	11(22%)						
Septic blood	4(8%)						
Consequence of infection							
No increase in supports	24(48%)						
Intervention	26(52%)						
Underlying medical problem							
Total % with GIT problems	23						
Total % without GIT	77						
problems	0.0						
Total % of Multiple PICCS	9.2						