

Directorate of Legal Services

PRACTITIONERS IN LAW TO THE HEALTH & SOCIAL CARE SECTOR

2 Franklin Street, Belfast, BT2 8DQ DX 2842 NR Belfast 3

Your Ref: BMcL-0027-12 Our Ref: HYP/W50/2 Date:

26th February 2013

Mr B McLoughlin
Assistant Solicitor to the Inquiry
Inquiry into Hyponatraemia-related Deaths
Arthur House
41 Arthur Street
Belfast
BT1 4GB



Dear Sir

RE: INQUIRY INTO HYPONATRAEMIA RELATED DEATHS- RAYCHEL FERGUSON (PRELIMINARY)

I refer to the above matter and to your letter dated 13th November 2012, referenced above. I confirm having now received instructions from my client, as follows.

I am informed that, in its efforts to obtain the information requested in your above-mentioned letter, the Western Trust made various attempts to contact both of the authors of the presentation, Dr McMorrow and Dr Corrigan. Dr McMorrow was formerly employed by the legacy Trust in 2000 as a Junior Doctor. However, she is no longer employed by the Western Trust and the Trust is currently attempting to locate her whereabouts.

Dr Corrigan is a Consultant Paediatrician who is currently employed by the Trust at Altnagelvin Hospital. My client had passed on your request for information to Dr Corrigan, and he has only just been able to locate the relevant documentation which he has now forwarded to my client. Accordingly, I now enclose copies of the following documents, for your perusal:-

- Document entitled- 'Paediatric Hyponatraemia:Incidence Aetiology and Management'
- 2. 'Hyponatraemia in Paediatric Appendicitis'- Authors, McMorrow, Corrigan et alia.
- Presentation slides- 'Paediatric Hyponatraemia: Who Gets it and Why?'-Authors, McMorrow and Corrigan.

Providing Support to Health and Social Care







I also enclose a pro forma 'Checklist document' which has been provided to us by the Trust's Professional Audit Department, in connection with Dr Corrigan's Audit.

Yours faithfully

Angela Crawford Solicitor

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Paediatric Hyponatraemia: Incidence Aetiology and Management

Background and aims: Recent publicity has highlighted the potentially catastrophic results of paediatric hyponatraemia. Little data exists however on the overall incidence and course of uncomplicated hyponatraemia in general paediatric practice.

Methods: All children admitted to Altnagelvin Hospital over a 21-month period with hyponatraemia (<133) were identified using our laboratory database. Newborns and infants in NICU were excluded. Retrospective chart review enabled data collection regarding incidence, aetiology and management.

Results: 6,276 children were admitted to Altnagelvin Hospital over the study period. 153 patients (2.4%) had at least one documented episode of hyponatraemia (medical 68% vs surgical 32%). Of 105 medical children, the most common diagnoses were gastroenteritis (25%), viral infections (25%) and lower respiratory tract infection (21%). Of 21 general surgical children, acute appendicitis accounted for 50%. The incidence of hyponatraemia complicating acute appendicitis was 11%.

The primary pathophysiological trigger for the hyponatraemia was attributable to SIADH in 66% of children overall.

Hyponatraemia was a presenting feature in 88% of children. Intravenous fluid therapy was prescribed for 61% of children. 0.45% saline / dextrose (64%) and isotonic saline solutions (27%) comprised the majority of prescribed fluids with no children receiving solution 18. Fluid volumes were given as maintenance in 60%, restricted in 31% and potentially generous in 7%. No serious complications of hyponatraemia were identified.

Conclusion: Hyponatraemia is a common complication of acute illness in both medical and surgical children, with SIADH as the primary mechanism. Very hypotonic intravenous solutions are no longer being prescribed, in keeping with regional guidelines.

HYPONATRAEMIA IN PAEDIATRIC APPENDICITIS

AS McCoubrey, A McMorrow, N Corrigan, S Dace Altnagelvin Area Hospital, Londonderry, BT47 6SB

Introduction

Paediatric hyponatraemia is increasingly being recognised as a rare but significant cause of morbidity and mortality in ill children. An association with appendicitis is recognised however the incidence is unknown. This study aims to look at incidence and possible clinical markers predictive of hyponatraemia in children with appendicitis.

Methods

A retrospective chart review of all children less than 13 years admitted to a District General Hospital with appendicitis over a 24 month period (January 2004-December 2005) was undertaken. Data collection included serum sodium, length of presentation, examination findings, serum inflammatory markers, intraoperative pathology and fluid management. Significant hyponatraemia was defined as plasma sodium < 133mmol/l.

Results

Seventy-one appendicectomies were carried out over the study period in children aged 3-12 years. Seventeen patients were found to be significantly hyponatraemic giving an incidence of 23%. Predictive makers of hyponatraemia were a high CRP level (p=0.0025) and perforated appendix (p=0.003). Length of presenting symptoms, presence of peritonism and white cell count were not predictive of hyponatraemia.

Conclusion

Hyponatraemia is a common problem in children with appendicitis. Clinical markers associated with its presence or subsequent development include a markedly raised CRP level and a gangrenous or perforated appendix at time of operation. Electrolytes should be

carefully monitored throughout admission and acted on appropriately. $\ensuremath{\mathsf{P}}$

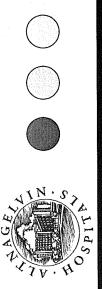
Clinical Audit Department Directorate of Nursing & Risk Management

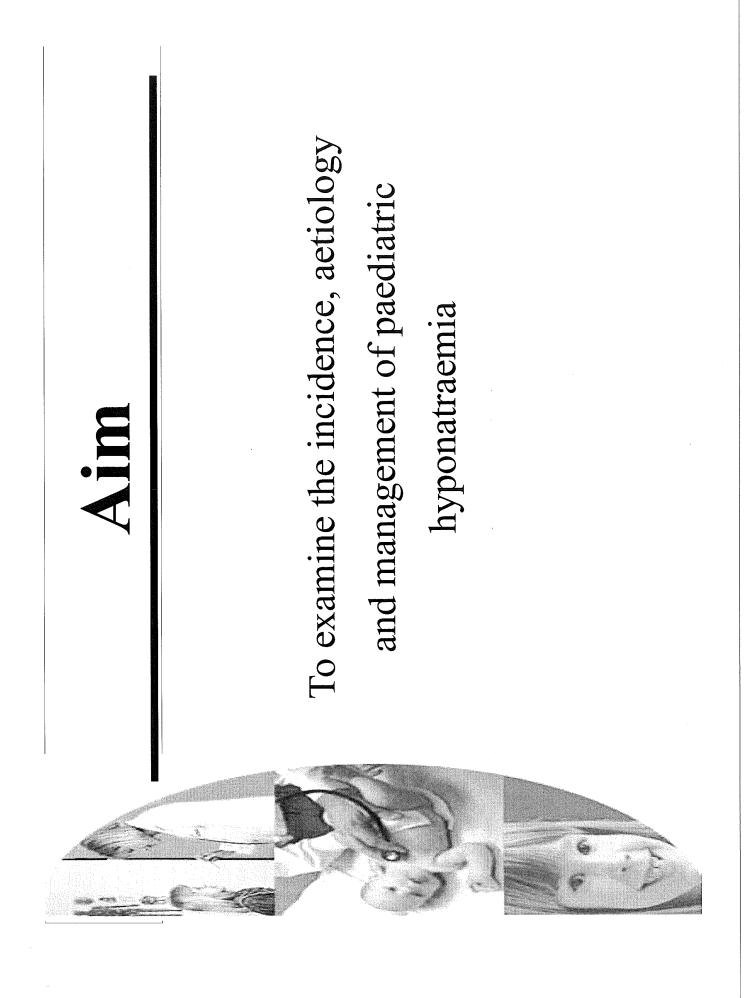
"Audits Undertaken Determines Improvement in Treatments"



Paediatric Hyponatraemia: Who Gets it and Why?







Methods

Retrospective chart review of all children admitted to Altnagelvin Hospital over a 21 month period with sodium <133



Results

6,276 admissions

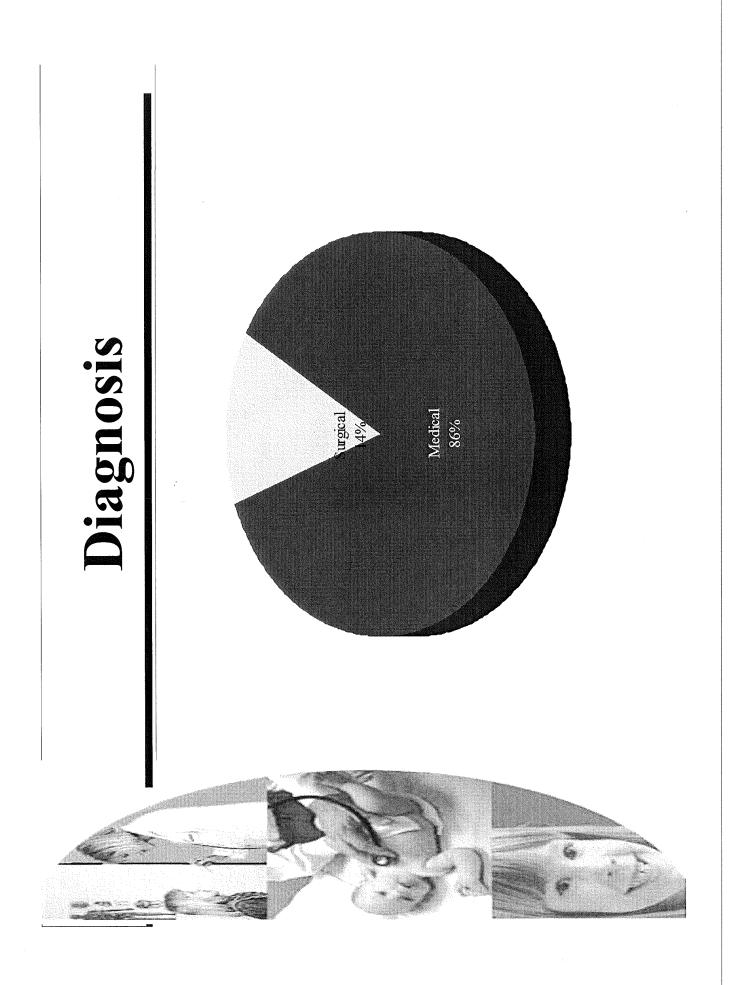
147 children (2.3%) identified

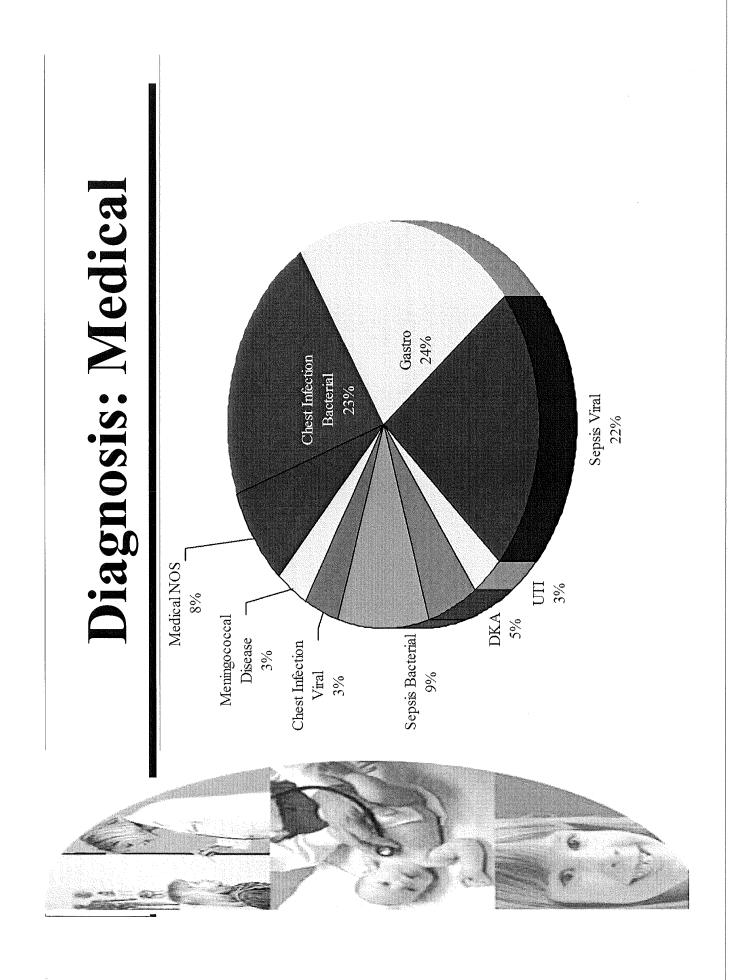
136 charts reviewed

Newborns and infants in NICU

excluded

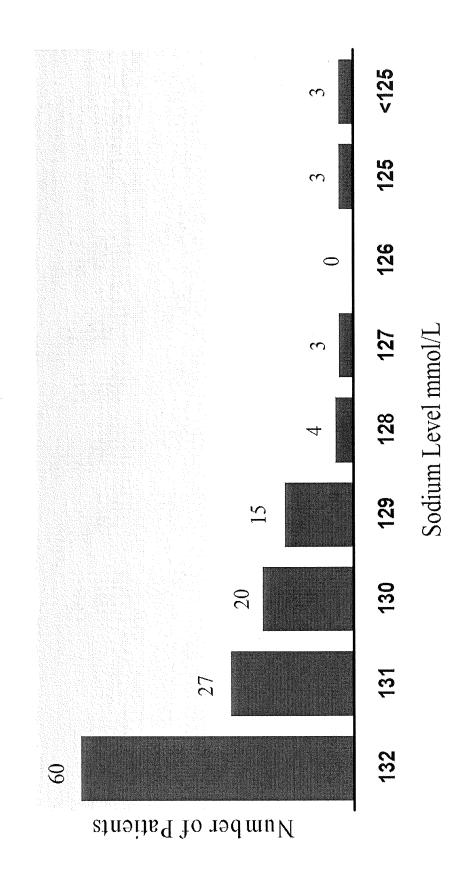




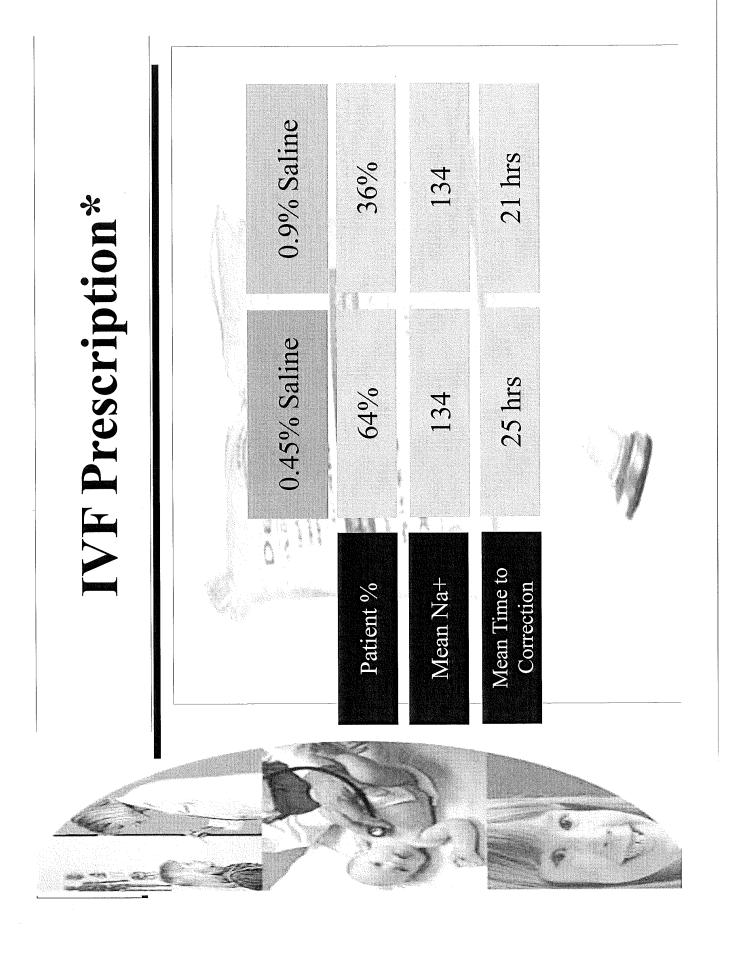


Diagnosis: Surgical Appendicitis 47% Other Surgical NOS 53%

Distribution of Lowest Na+

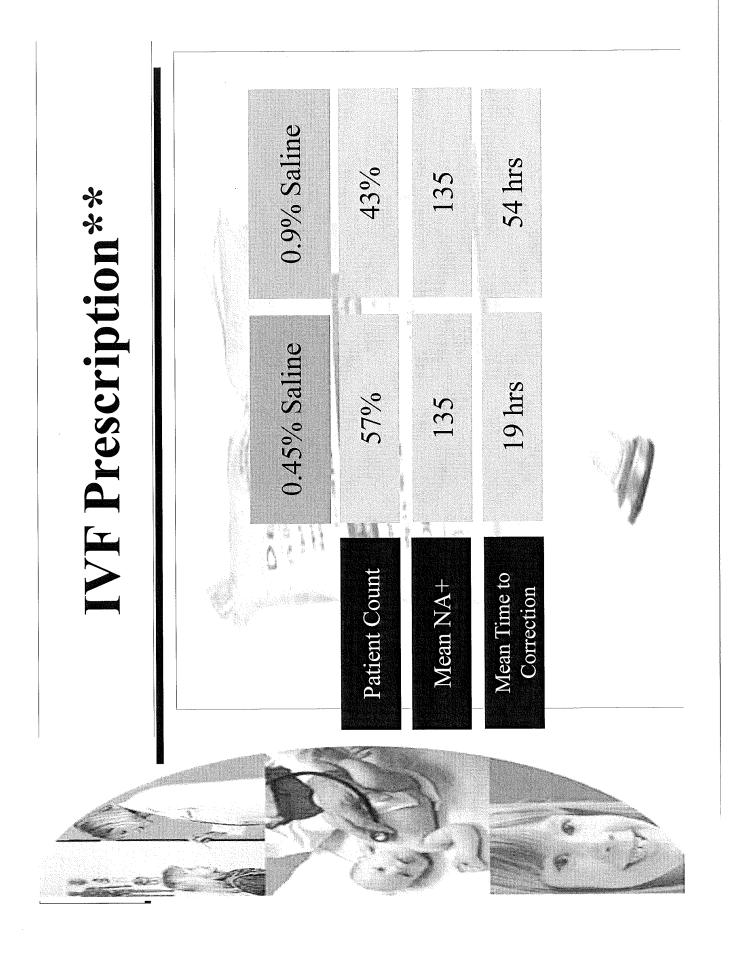


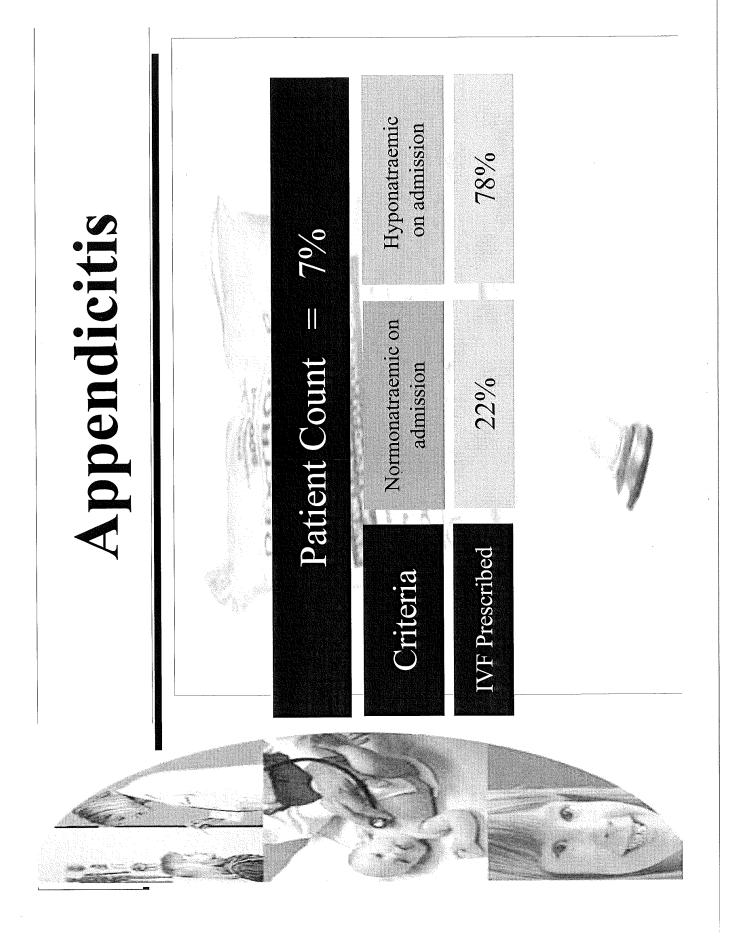
Hyponatraemia On Admission No IVF 22 hrs 32% 92% %8 130 IVF Prescribed 23 hrs %89 91% 134 %6 Mean Time to Correction Mean Na+ Patient % Surgical Medical



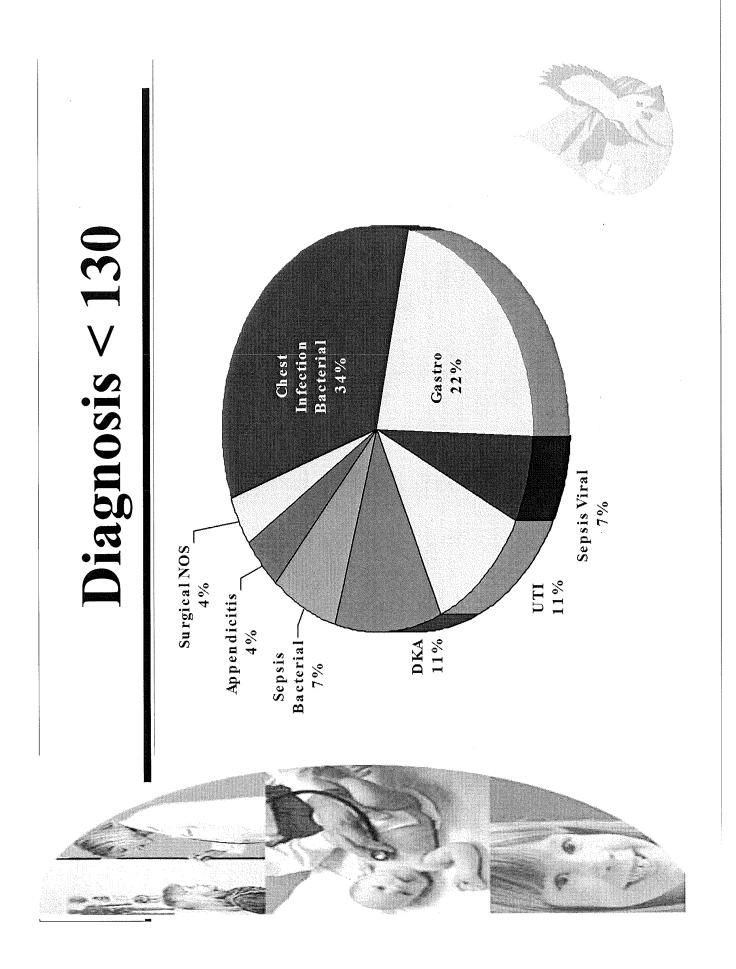
Normonatraemic On Admission No IVF 25% 75% 25% IVF Prescribed 92% 50% 75% 20% %8 Prescribed 0.45% Saline Prescribed 0.9% Saline Patient Count Surgical Medical

Became Hyponatraemic No IVF 25% %19 5 hrs 33% 136 During Admission IVF Prescribed 49 hrs 50% 75% 50% 135 Mean Time to Mean Na+ Correction Patient % Surgical Medical





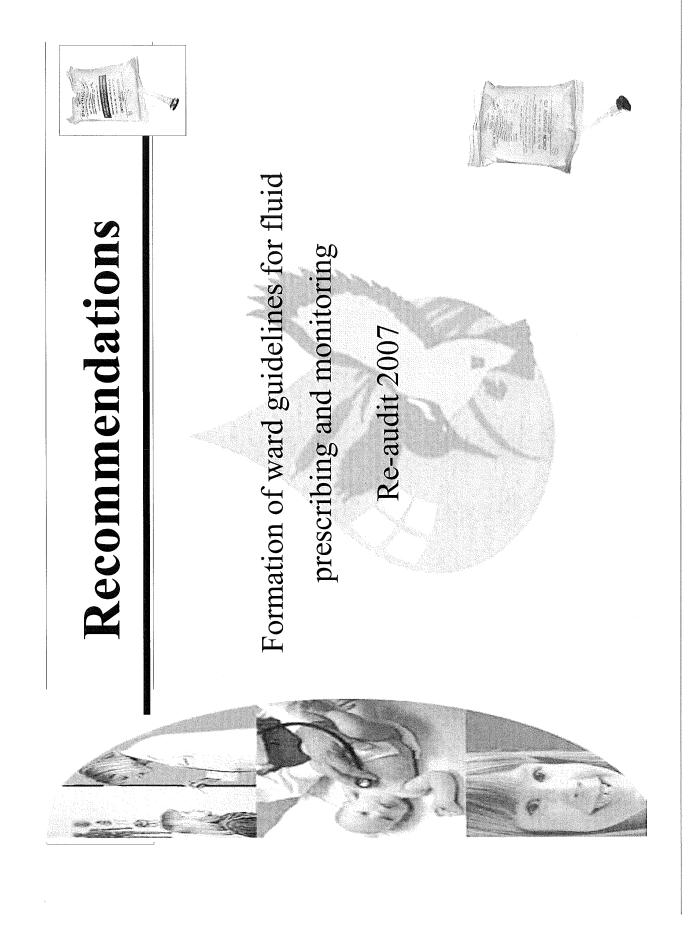
Sodium Level < 130 44% 85% 52% 4% Patient Count = 20% Received IVF 0.90% Saline 0.45% Saline Hartman's



Hyponatraemic on admission Aetiology: SIADH 41% 62% Patient Count = 66%Normonatraemic on admission 83% %0 Received IVF IVF Volume Restricted

Hyponatraemia on IVF Volume: Generous Admission 83% 17% 5% Normal Serum Na⁺on Admission 100% %0 %9 Patient Count Surgical Medical

Very hypotonic solutions no longer being Common complication in acute medical and surgical conditions Conclusion prescribed





Acknowledgement

H&SST through the Clinical Audit The Author / Authors of this audit resources provided by Altnagelvin project wish to acknowledge the Department in support of this piece of work. · compon

Hosp No: AH442134	Patient 14 0
Hyponatraemia on admission: If NO Go to Section A:	Table 1
Section A: General Management:	Dalua Dalua
Was Fluid Yes No Bolus Volume Yes No Appropriate:	Bolus Yes No Solution Appropriate:
Were IV Fluids Yes No No Input Yes administered: No No Recorded:	No Output Yes No Recorded:
How was it recorded: Standard Urine Napples Weighed	Catheterised Not recorded
Initial 0.5% N saline N saline (N) Hartmans (H Prescription:) Sol 18 (SO) Other (O)
Initial Restricted Appropriately SIADH General	erous Too little Other
appropriate clinically: reassessment hypo	The state of the s
(hours) havin	ng had normal al sodium level?
Was Fluid Yes No Bolus Volume Yes No Bolus required: Appropriate:	Bolus Yes No solution appropriate:
Were IV fluids Yes No No Input Yes administered: of recorded:	No Output Yes No recorded:
How was it recorded: Standard Urine Napples weighed	Catheterised Not recorded
Initial 0.5% Fluid prescription N saline N saline (N) Hartman for Hyponatraemia: + dextrose (SD)	ns (H) Sol 18 (SO) Other (O)
Initial Restricted appropriately SIADH General Prescribed:	nerous
Repeat 0.5% Fluid prescription N saline N saline (N) Hartmar + dextrose (SD)	ns (H) Sol 18 (SO) Other (O)
Fluid Volume appropriately SIADH Prescribed:	nerous Too Little Other
Were fluids Yes No Time to prescribed If patie reassessment relevant to surgic of fluids probable were ti (Hours) ongoing actiology referre of hyponatraemia:	al
Is there record Yes No Is there record in chart of discussion in chart of discussion	
4 sugas gatio hall gibli gibli 1601	





Clinical Audit Department Directorate of Nursing & Risk Management

"Audits Undertaken Depresines Improvement in Treatments" Patient Hyponatraemia Study Data Sheet: ID: December 2005 Age: Gender: Male Name: length of Adm Time: stay (days) Date: U&E Time (hours) to Time to 1st post treatment: documented Tot number of U&E's checked: <6h \ <12h normal sodium: Daily [> 24 h Lowest Lowest serum osmolality: (= 2Na + glucose + urea) Highest Highest urine osmolality: CRP: WČC: Aetiology if surgical: Pre-op Post-op N/A Directorate: Medical Surgical Diagnosis/code: Surgical: Infective: Medical Other: Appendicitis (A) Chest infection - bacterial (CB) DKA (D) Other Surgical NOS (SU Other medical NOS (MN) Chest infection - viral (CV) Orthopaedic (OR) ENT infective (EI) UTI (U) ENT other (EO) Septicaemia/meningococcal disease/meningitis (M) Sepsis - bacterial NOS (SB) Sepsis - Viral NOS (SV) Gastroenteritis (G) Aetiology <u>Inappropriate</u> <u>Other</u> Inappropriate IV Inappropriate IV <u>Drugs</u> SIADH Excessive Losses Replacement Volume 1

DRAFT

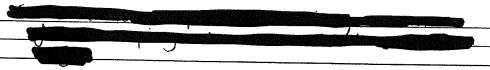
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NAME OF PARTICIPANTS:D	r Neil Corrigan,	Dr Aoife N	IcMorrow	- Approved autside
PRIMARY CONTACT: N Con	rigan	CO	NTACT NUMI	2/12/05 · BER: Bleep 8279
AUDIT TITLE The mana	agement of hypon	The second second second second	TO A STREET PROPERTY OF THE	to AAH over a 21 month period
METHOD: PROSPECTIVE	·	IF QUESTI	ONNAIRE PL	EASE INDICATE:
RETROSPECTIV	$E\begin{bmatrix}x\end{bmatrix}^{\frac{1}{2},\frac{1}{2}} x \xrightarrow{x} \overset{x}{\xrightarrow{x}} \overset{x}{\xrightarrow{x}}$	PATIENT [STATT
STANDARDS / GUIDELINES	* (0) 4	l College l Hospital		CREST Professional College
Please attach guideline/standar	d Other	please specify	DHSS	
OURCE OF AUDIT: ALT	NAGELVIN [REGIO	NAL [NATIONAL [
Line Manager Signature:	Dr Murray Quinn			
(To state that he/she has appro- N.B. THIS FORM WILL NOT	ved this audit wi BE PROCESSE	u take place) D UNLESS S	IGNED BY LI	NEMANACER
WHERE DO YOU INTEND TO		25.27.2000.275.2647.13509.2	- 大学では、「製力庫」でも、だり	iety / local audit
f you intend to present your audit outsic			sion from Line Me	nety / local audit
CLINICAL AUDIT DEPART! The Clinical Audit Department is	MENT ASSISTA happy to assist w	NCE ath one or mo	re of these field	
QUESTIONNAIRE/ PROFORMA DESIGN		k required as LYSIS	sistance	
PRESENTATION		NOTES/X-R		♥ MANY? 153
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OMMENCEMENT DATE: ASAP		√Vai	TION DATE:	Ceb 2006

If you have not requested assistance with your questionnaire, please enclose a copy of audit questionnaire you intend to use. Failure to do so may delay the start of your audit

The Clinical Audit Committee will discuss all requests for audit assistance, the schedule of meetings are detailed below. You will receive a letter within 2-3 days after the meeting informing you of assistance that will be available to you.

In EXCEPTIONAL circumstances, the Clinical Audit Committee can give approval for a study to commence and audit assistance to begin.

If you consider you need an urgent audit commenced before the next meeting, please indicate why?



Please return this form to Clinical Audit Manager, Clinical Audit Office, Altnagelvin Hospital.

Meeting Date		Closing Date for Entry				
	4 TH JANUARY 2005	Closing date 22 nd December 2004				
	1 ST MARCH 2005	Closing date 22 nd February 2005				
**************************************	3 RD MAY 2005	Closing date 26 th April 2005				
	5 TH JULY 2005	Closing date 28th June 2005				
	6 TH SEPTEMBER 2005	Closing date 30 th August 2005				
	1 ST NOVEMBER 2005	Closing date 25th October 2005				
	3 RD JANUARY 2006	Closing date 22 nd December 2005				

DATE RECEIVED:

AUDIT COMMITTEE APPROVAL: YES! No.

Hyponatraemia study data sheet November 2005

Further da	Further data to be collected:								
Identifier:	Identifier:								
Hyponatraemia									
Serial resu	Serial results								
day					T	T.			
sodium									
			***************************************		<u> </u>		L		
Other electrolyte abnormalities y/n Other electrolyte abnormalities – define									
WCC		CRP	Mary polyton nymotronia y s		, , , , , , , , , , , , , , , , , , ,			Approximate and any any and any any and any	
Aetiology:									
	Surgical versus medical S/M Hyponatraemic at admission Y/N Actiology								
Aetiology	ISADH	Excessive Losses		ropriate ement	Drugs		napprop Oth luids		
·				LIPINA II.					
Diagnosis: Diagnosis classification: infective resp/git/neuro/other Surgical appx/other git/other									
Manageme	Management:								
Input/output recorded Y/N How recorded Standard/urine measured/nappies weighed/catheterised/other									
IV fluids administered Y/N									
Type of fluid									
Type of	0.5%N	N salin	ne	Hartma	ns	Sol 18	O	lher	
fluid	saline +		·						
Type of flui	d appropriat	te clinically	Y/N	,					

Fluid prescription:

	rigid brescrib	tion.				
	Maintenance	Restricted appropriately ISADH	Restricted with concomitant orals	Generous	Too little	
-				,		

If generous how much extra given %

Became hyponatraemic on IV fluids:

Y/N