



# **PICANet Custom Audit Definitions**

# **Renal Dataset**

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Version 2.0 (March 2017)

# Renal Dataset Contents

<b>PICANet Custom Audit Definitions</b> .....	1
<b>Renal Dataset</b> .....	1
Version 2.0 (March 2017).....	1
<b>INTRODUCTION</b> .....	1
Background .....	1
Data Collection method .....	1
<b>RENAL DATASET</b> .....	2
<b>PATIENT DETAILS</b> .....	2
Family name or Surname .....	2
First name .....	2
Date of birth.....	3
NHS, CHI or H&C number.....	3
NHS, CHI or H&C number eligibility .....	3
Case note number.....	4
Admission number.....	4
<b>INITIATION DETAILS</b> .....	4
Date and Time of initiation of Continuous Renal Replacement Therapy (CRRT).....	4
Weight.....	4
Indication for Renal Replacement Therapy (RRT).....	5
Creatinine value (prior to commencement of continuous renal replacement therapy - CRRT): .....	6
Baseline creatinine value at PICU admission .....	6
Highest creatinine value during PICU admission prior to initiation of CRRT .....	6
Urine output.....	6
Ammonia value ( <i>if applicable</i> ) .....	7
Lactate.....	7
Net fluid balance.....	7
<b>PERITONEAL DIALYSIS</b> .....	8
Peritoneal dialysis .....	8
Total duration of peritoneal dialysis therapy.....	8
<b>EXTRA-CORPOREAL CRRT (<i>during this admission</i>)</b> .....	8
Continuous extracorporeal renal replacement therapy (CRRT) .....	8
CRRT machine used.....	9
Other CRRT machine specified.....	9
Site of Vein ( <i>on initiation</i> ) .....	9
Venous catheter ( <i>on initiation</i> ).....	10
Manufacturer name and type.....	10
Size of the venous catheter.....	10
Length of the venous catheter.....	10
Number of lumens of venous catheter .....	12
Blood transfusion.....	12

Continuous Venovenous Haemofiltration (CVVH) .....	13
Maximum blood flow .....	13
Maximum ultrafiltrate flow.....	13
Continuous Venovenous Haemodialysis (CVVHD) .....	14
Maximum blood flow .....	14
Maximum dialysate flow .....	14
Continuous Venovenous Haemodiafiltration (CVVHDF).....	14
Maximum blood flow .....	15
Maximum ultrafiltrate flow.....	15
Maximum dialysate flow .....	15
Type of Replacement fluid .....	15
Replacement fluid given ( <i>CVVH only</i> ) .....	16
Anticoagulant used .....	16
Type of anticoagulant ( <i>tick all that apply</i> ) .....	17
Target used .....	17
Total duration of Continuous Renal Replacement Therapies (CVVH/CVVHD/CVVHDF) .....	18
Total number of filters used.....	18
Number of elective filter changes.....	19
RRT STATUS AT DISCHARGE .....	19
Form completed by.....	20
Appendix A –Missing Values .....	21

# INTRODUCTION

## Background

In collaboration with the Continuous Renal Replacement Therapy (CRRT) Study Group of the Paediatric Intensive Care Society (PICS), UK and Dr Claire Westrope PICANet has developed the Renal Custom Audit of paediatric patients who receive renal CRRT in PICUs in the UK and Ireland.

This audit collects detailed data on each patient episode of CRRT in the paediatric intensive care unit; providing information about the use of CRRT in the critically ill child, an audit of current practice and an evidence base to potentially drive future research into best practice.

## Data Collection method

For units who agree to participate in this custom audit PICANet will enable access to the specific custom audit data collection tab on the data entry page.

1. A PICANet Renal custom audit form (Appendix B – page 21) is completed for all admissions who receive Continuous Renal Replacement Therapy (CRRT) during the PICU admission event.
2. When the PICU has entered or uploaded to PICANet Web, the admission event data for the named patient the Renal Dataset may be manually entered onto the **Renal** data collection tab.
3. To enter Renal data, open the admission event on PICANet Web for the named patient and select the **Renal** tab. Note that the **Renal** tab is visible for users in PICUs who have opted into this specific custom audit. It is visible for all admission events in participating units but completion is only required for patients who receive CRRT during the specified PICU admission event.

The screenshot shows a web-based data entry form for a Renal Audit. At the top, there are navigation buttons: 'Save', 'Cancel', 'Delete', '< Back', and 'Next >'. Below these are tabs for different data entry sections: 'Patient details', 'Admission details', 'Diagnoses and procedures', 'PIM', 'Daily interventions', 'Interventions', 'Trial + Growth', 'Discharge + Follow-up', 'Comments', 'Legacy data', and 'Renal Audit' (which is currently selected). The form is divided into several sections:

- Initiation Details:** Includes fields for 'Date and time of initiation of CRRT' (09/12/2014, 12:00), 'Weight' (80 kg), and a dropdown for 'Indicate if weight is' (Actual). It also has a list of checkboxes for 'Indication for RRT': Fluid overload (checked), Electrolyte imbalance (checked), High creatinine, Acidosis, Temperature control, Oliguria, and Toxin removal.
- Creatinine value prior to CRRT:** Includes 'Baseline at PICU admission' (248 µmol/l) and 'Highest during PICU admission' (312 µmol/l).
- Urine output:** Includes 'Total in 4 hrs prior to CRRT' (515 µmol/l).
- Ammonia value if applicable:** Includes 'Highest during PICU admission' (1012 µmol/l).
- Lactate:** Includes 'Highest during PICU admission' (15.8 µmol/l).
- Net fluid balance:** Includes 'PICU admission to initiation of CRRT' (15.8 µmol/l).
- Peritoneal dialysis during this admission:** Includes a checked checkbox for 'Peritoneal dialysis' and a field for 'Total duration' (34 hrs).
- CRRT During this admission:** Includes checkboxes for 'CRRT machines used': Aquarius, Prismaflex, and Other (specify) (checked).

# RENAL DATASET

## PATIENT DETAILS

### Family name or Surname

<b>Definition</b>	The last or family name or surname given to the child as it would appear on the child's birth certificate or other appropriate document.
<b>Reason</b>	Family name provides an additional identifier that can aid patient tracking throughout the hospital.  Can help identify individuals who may have had multiple admissions to one or more PICUs.
<b>Format</b>	Free text (e.g. Brown).  If no family name available record as UNKNOWN and indicate why not available in the comments section.

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### First name

<b>Definition</b>	The first name given to the child as it would appear on the child's birth certificate or other appropriate document.
<b>Reason</b>	First name provides an additional identifier that can aid patient tracking throughout the hospital and PICANet Web.  Can help identify individuals who may have had multiple admissions to one or more PICUs.
<b>Format</b>	Free text (e.g. John).  If no first name available record as UNKNOWN and indicate why not available in the comments section.

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## Date of birth

<b>Definition</b>	The child's date of birth as recorded on the child's birth certificate or other appropriate document.
<b>Reason</b>	<p>Date of birth and date of admission are used to calculate age at admission to your unit.</p> <p>Date of birth provides an additional identifier that can aid patient tracking throughout the hospital and PICANet Web.</p> <p>Can help identify individuals who may have had multiple admissions to one or more PICUs.</p>
<b>Format</b>	<p>Date; dd/mm/yyyy.</p> <p>Date of birth should be between 01/01/1980 and Date of admission.</p> <p>If the child's date of birth is unobtainable, but the child is on your unit, use your judgement to estimate year of birth and record as 1 January of estimated year (e.g. 01/01/YYYY).</p> <p>If information is being extracted from notes and the child's date of birth is not recorded, or recorded as unavailable, leave the field blank and select DOB not known in the Indicate if date of birth is field at data entry.</p> <p>If it is necessary for Date of birth to be partly anonymised, enter the correct month and year and record 01 for the day (e.g. 01/MM/YYYY). Then select Anonymised in the Indicate if date of birth is field at data entry.</p>

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## NHS, CHI or H&C number

<b>Definition</b>	Unique identifying number enabling tracing of a patient through the NHS system in the United Kingdom. For English and Welsh patients the NHS number, for Scottish patients the CHI number and for Northern Ireland the H&C number is used as a unique numeric identifier.
<b>Reason</b>	<p>NHS, CHI or H&amp;C number gives a unique, identifiable variable that will allow other identifiable data items to be removed from the database.</p> <p>Can identify individuals who may have had multiple admissions to one or more PICUs.</p>
<b>Format</b>	Free text (e.g. 1463788990).

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## NHS, CHI or H&C number eligibility

<b>Definition</b>	The patient is not eligible for NHS, CHI or H&C number, he or she is an overseas national who is not ordinarily a resident in the UK and therefore does not have an allocated NHS, CHI or H&C number.
<b>Reason</b>	To enable effective audit of availability of NHS, CHI or H&C number and assessment of health services delivery.
<b>Format</b>	Tick box if patient is not eligible for an NHS, CHI or H&C number.

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## Case note number

<b>Definition</b>	Unique identifying number for an individual's hospital records at the treating unit.  Allocated on first admission to hospital.
<b>Reason</b>	Case note number provides a unique identifier that can aid patient tracking throughout the hospital.
<b>Format</b>	Free text (e.g. AB145C).

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## Admission number

<b>Definition</b>	Unique identifier assigned to each consecutive admission to your unit. As recorded in your unit admission book or clinical information system. Admission to your unit is defined as the physical admission and recording of that admission to a bed or cot in your unit.
<b>Reason</b>	Admission number provides a unique identifier for each admission to each unit participating in PICANet and thus allows identification of one set of admission data from another.
<b>Format</b>	Free text (e.g. 01/389).

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## INITIATION DETAILS

### Date and Time of initiation of Continuous Renal Replacement Therapy (CRRT)

<b>Definition</b>	The date and time that continuous renal replacement therapy was initiated during this admission to PICU.
<b>Reason</b>	Date and time of commencement of CRRT is required to calculate the length of stay in PIC prior to the commencement of this treatment.
<b>Format</b>	Date: (dd/mm/yyyy) Time: (24 hour clock); hh:mm.
<b>Validation rule</b>	Value expected Warning – date and time of commencement of RRT should not precede data and time of admission to PICU

<b>Variable name</b>	DateInitCRRT	<b>GroupID</b>	1	<b>AttributeID</b>	1
<b>Variable name</b>	TimeInitCRRT	<b>GroupID</b>	1	<b>AttributeID</b>	2

## Weight

<b>Definition</b>	Patient weight in kilograms rounded to nearest 1 decimal place as documented on admission to PICU.
<b>Reason</b>	To inform subsequent analyses
<b>Format</b>	Numerical value to 1 decimal place (e.g. 24.9) Units: kg; validation check if range outside 0.5 to 80.0
<b>Validation rule</b>	Value expected. Warning if range outside 0.5 to 80.0 Warning if missing

<b>Variable name</b>	RenalWeight	<b>GroupID</b>	1	<b>AttributeID</b>	3
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## Indicate if weight is

<b>Definition</b>	Specifies whether the weight recorded is actual or estimated
<b>Reason</b>	Estimated weights may reduce the power of subsequent analyses
<b>Format</b>	Choose from one of the following: Actual Estimated

**Validation rule** Expected value if weight is recorded

<b>Variable name</b>	WeightActEst	<b>GroupID</b>	1	<b>AttributeID</b>	5
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## Indication for Renal Replacement Therapy (RRT)

<b>Definition</b>	The clinical indication(s) for the decision to commence renal replacement therapy
<b>Reason</b>	To inform the clinical indications for the decision to commence CRRT
<b>Format</b>	Tick all that apply:

**Fluid Overload** - where the indication for RRT is the removal of excess fluid from the patient or to facilitate increased fluid intake

**Electrolyte imbalance** - where the indication for RRT is to correct deranged electrolytes including sodium, potassium, urea, creatinine, phosphate, calcium and urate

**Acidosis** - where the indication for CRRT is metabolic acidosis including lactic acidosis

**Oliguria** - where the indication for CRRT is reduced urine output < 0.5ml/hr for previous four hours

### High creatinine

Where the indication for CRRT is a creatinine value that is above the normal range for age, or the creatinine value has risen from baseline at admission to PICU

### Temperature Control

Where the initiation of CRRT is to facilitate control of hypothermia or hyperthermia

### Toxin removal

Where initiation of CRRT is to facilitate the removal of external toxins such as poisons, sepsis or the removal of toxic metabolites in inborn errors of metabolism e.g. ammonia (tick box if toxin is ammonia)  
Specify the type of toxin.

**Validation rule** At least one option is expected

<b>Variable name</b>	IndicRRTFluid	<b>GroupID</b>	1	<b>AttributeID</b>	8
<b>Variable name</b>	IndicRRTElectrolyteImb	<b>GroupID</b>	1	<b>AttributeID</b>	10
<b>Variable name</b>	IndicRRTHiCreatine	<b>GroupID</b>	1	<b>AttributeID</b>	11
<b>Variable name</b>	IndicRRTAcidosis	<b>GroupID</b>	1	<b>AttributeID</b>	12
<b>Variable name</b>	IndicRRTTempCont	<b>GroupID</b>	1	<b>AttributeID</b>	13
<b>Variable name</b>	IndicRRTOliguria	<b>GroupID</b>	1	<b>AttributeID</b>	14
<b>Variable name</b>	IndicRRTToxRem	<b>GroupID</b>	1	<b>AttributeID</b>	15

## Toxin removal specify

<b>Definition</b>	The type of toxin if CRRT is commenced for specified toxin removal such as poisons, sepsis or the removal of toxic metabolites in inborn errors of metabolism e.g. ammonia
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<b>Reason</b>	To inform the clinical indications for the decision to commence CRRT
<b>Format</b>	Free text
<b>Validation rule</b>	Expected value if toxin removal is selected in Indication for RRT

<b>Variable name</b>	IndicRRTToxRemReas	<b>GroupID</b>	1	<b>AttributeID</b>	16
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## Creatinine value (prior to commencement of continuous renal replacement therapy - CRRT):

### Baseline creatinine value at PICU admission

<b>Definition</b>	The first recorded creatinine value following admission to PIC.
<b>Reason</b>	Rise in creatinine from baseline required to calculate Acute Kidney Injury Score
<b>Format</b>	Creatinine value in micromol/litre [ $\mu\text{mol/L}$ ]. Expected range 50-115
<b>Validation rule</b>	Value expected Validation check if zero or >500 Warning if missing

<b>Variable name</b>	BaseCreatAtAdm	<b>GroupID</b>	1	<b>AttributeID</b>	18
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### Highest creatinine value during PICU admission prior to initiation of CRRT

<b>Definition</b>	The highest recorded creatinine value during this PIC admission and prior to the commencement of continuous renal replacement therapy. This value may be the same as the baseline result.
<b>Reason</b>	Rise in creatinine from baseline required to calculate Acute Kidney Injury Score
<b>Format</b>	Creatinine value in micro mol/litre [ $\mu\text{mol/L}$ ]. Expected range 50-115
<b>Validation rule</b>	Value expected Validation check if zero or >500 Warning if missing

<b>Variable name</b>	BaseCreatAtAdm	<b>GroupID</b>	1	<b>AttributeID</b>	19
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### Urine output

<b>Definition</b>	Total urine output in the 4 hours immediately prior to the initiation of continuous renal replacement therapy (CRRT)
<b>Reason</b>	Urine output in ml/kg/hr is required to calculate Acute Kidney Injury Score
<b>Format</b>	Record as millilitres in whole numbers (e.g. 55).
<b>Validation rule</b>	Value expected Can be zero Validation check if > 1000 Warning if missing

<b>Variable name</b>	UrineOutput	<b>GroupID</b>	1	<b>AttributeID</b>	22
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### Ammonia value (if applicable)

<b>Definition</b>	The highest recorded ammonia value measured prior to the initiation of continuous renal replacement therapy (this may be a measurement from the previous day) <b>Record only if toxin removal due to raised ammonia</b> is recorded in indications for RRT
<b>Reason</b>	To assess/establish threshold value for initiation of RRT for hyperammonia
<b>Format</b>	Micromols /litre [ $\mu\text{mol/L}$ ]. Whole number (e.g. 545)
<b>Validation rule</b>	Value expected only if toxin removal is ticked in indications for RRT and ammonia is specified as type of toxin Validation check if outside range 100-2000

<b>Variable Name</b>	AmoniaMax	<b>GroupID</b>	1	<b>AttributeID</b>	24
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### Lactate

<b>Definition</b>	The most recent lactate value measured during current PICU admission prior to initiation of continuous renal replacement therapy (this may be a measurement from the previous day)
<b>Reason</b>	To assess/establish threshold value for initiation of RRT for hyperlactaemia
<b>Format</b>	Value rounded up to nearest 1 decimal place in millimol/l (e.g. 13.5) Range 0.1 to 20.0
<b>Validation rule</b>	Value expected Validation check if zero or >30.0 Enter 88.8 if out of range reading

<b>Variable name</b>	LactateMax	<b>GroupID</b>	1	<b>AttributeID</b>	26
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### Net fluid balance

<b>Definition</b>	The total net fluid balance from the time of admission to PICU to initiation of CRRT
<b>Reason</b>	To calculate percentage fluid overload at initiation of CRRT using net fluid balance divided by weight on admission
<b>Format</b>	+/- millilitres e.g. - 1300 or + 400
<b>Validation rule</b>	Value expected Validation check if outside range -10000 to +10000

<b>Variable name</b>	NetFluidBallance	<b>GroupID</b>	1	<b>AttributeID</b>	21
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## PERITONEAL DIALYSIS

This section is applicable to patients who have received peritoneal dialysis therapy during their PICU admission event

### Peritoneal dialysis

<b>Definition</b>	Peritoneal dialysis used during this PICU admission Includes patients who have a peritoneal catheter inserted and used for the purpose of peritoneal dialysis. Do not include patients who have a peritoneal catheter inserted for drainage of ascites A patient may receive both peritoneal dialysis and CRRT during the same PICU admission
<b>Reason</b>	To record the use of peritoneal dialysis therapy
<b>Format</b>	Tick box if peritoneal dialysis used at any time during this PICU admission

**Validation rule** None

<b>Variable name</b>	PeritonealDialysis	<b>GroupID</b>	1	<b>AttributeID</b>	28
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### Total duration of peritoneal dialysis therapy

<b>Definition</b>	The total number of hours spent on peritoneal dialysis therapy. Applies to the number of hours spent on actual peritoneal dialysis therapy. Do not include any hours where the patient is not actually receiving peritoneal dialysis e.g. where there is complication stopping peritoneal dialysis or treatment is stopped to assess renal recovery and then restarted
<b>Reason</b>	To assess the number of hours of peritoneal dialysis received
<b>Format</b>	Hours rounded to nearest hour, e.g. 15. Enter 9999 if not known

**Validation rule** Value expected value only if Peritonea Dialysis is selected

<b>Variable name</b>	PDTtotalDuration	<b>GroupID</b>	1	<b>AttributeID</b>	29
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## EXTRA-CORPOREAL CRRT (*during this admission*)

This section is applicable to patients who have had continuous extracorporeal renal replacement therapy during their PICU admission event

### Continuous extracorporeal renal replacement therapy (CRRT)

<b>Definition</b>	Continuous extracorporeal renal replacement therapy used during this PICU admission.
<b>Reason</b>	To record the use of extracorporeal renal replacement therapy
<b>Format</b>	Tick box if extracorporeal renal replacement therapy used at any time during this PICU admission
<b>Validation rule</b>	None

<b>Variable name</b>	ExtraCorporealCRRT	<b>GroupID</b>	1	<b>AttributeID</b>	78
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## CRRT machine used

<b>Definition</b>	The name of the machine used to deliver CVVH, CVVHD or CVVHDF
<b>Reason</b>	To assess if machine used to deliver CVVH/CVVHD or CVVHDF influences performance, complication rates or patient outcomes
<b>Format</b>	Tick all that apply Aquarius (Edwards Life Sciences) Prismaflex (Gambro) Other ( <i>specify</i> ) If Other, record name

**Validation rule** Expect one box to be ticked if any of CVVH, CVVHD or CVVHDF are true

<b>Variable name</b>	CRRTAquarius	<b>GroupID</b>	1	<b>AttributeID</b>	31
<b>Variable name</b>	CRRTPrismaflex	<b>GroupID</b>	1	<b>AttributeID</b>	32
<b>Variable name</b>	CRRTOther	<b>GroupID</b>	1	<b>AttributeID</b>	33

## Other CRRT machine specified

<b>Definition</b>	The name of the other CRRT machine used to deliver CVVH, CVVHD or CVVHDF Record if other CRRT machine used is selected
<b>Reason</b>	To assess if machine used to deliver CVVH/CVVHD or CVVHDF influences performance, complication rates or patient outcomes
<b>Format</b>	Free text
<b>Validation rule</b>	If other CRRT machine used is selected expect the name of the machine to be completed

<b>Variable name</b>	CRRTOtherDesc	<b>GroupID</b>	1	<b>AttributeID</b>	34
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## Site of Vein (*on initiation*)

<b>Definition</b>	The site of the vein into which vascular access is inserted, on initiation, for the purpose of facilitating continuous renal replacement therapy  Choose from Internal jugular –Left or Right Subclavian - Left or Right Femoral – Left or Right Umbilical ECMO Other -specify
<b>Reason</b>	To assess the influence of insertion site of vascular access on the application of renal replacement therapy
<b>Format</b>	Tick one box only Internal jugular vein – Left = 1 Internal jugular vein – Right = 2 Subclavian vein – Left = 3 Subclavian vein – Right = 4 Femoral vein – Left = 5 Femoral vein – Right = 6 Umbilical vein = 7 ECMO - tick if RRT circuit connected via ECMO circuit = 8 Other- specify = 9

<b>Validation rule</b>	Expected value. Only one site can be selected				
<b>Variable name</b>	SiteOfVein	<b>GroupID</b>	1	<b>AttributeID</b>	35

### Other site (*specify*)

**Definition** The other site into which vascular access is inserted, on initiation, for the purpose of facilitating continuous renal replacement therapy  
Record only if other site is selected as site of vein on initiation

**Reason** To assess influence of vascular access on application of renal replacement therapy

**Format** Free text

**Validation rule** Expected value if other site of vein is selected

<b>Variable name</b>	OtherSiteOfVein	<b>GroupID</b>	1	<b>AttributeID</b>	36
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### Venous catheter (*on initiation*)

#### Manufacturer name and type

**Definition** The manufacturer name and type of the specialised central venous catheter inserted into patient for the purpose of continuous renal replacement therapy on initiation  
For example Vygon Double Lumen Vascath or Cook Triple Lumen Central Venous Line  
Not required if site of vein is umbilical or ECMO

**Reason** To assess influence of vascular access on application of renal replacement therapy

**Format** Free text

**Validation rule** Value expected if site of vein is internal jugular left or right, subclavian left or right or femoral left or right or other

<b>Variable name</b>	VCManufacturer	<b>GroupID</b>	1	<b>AttributeID</b>	38
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#### Size of the venous catheter

**Definition** The size, recorded as the external diameter of specialised central venous catheter inserted into the patient for initiation of continuous renal replacement therapy  
Not required if site of vein is umbilical or ECMO

**Reason** To assess influence of vascular access size on application of renal replacement therapy, particularly blood flow rates and filter life

**Format** Record in French Fr (to one decimal place), e.g. 12.4

**Validation rule** Value expected if site of vein is internal jugular, subclavian or femoral and manufacturer name and type is completed  
Validation check if outside range 4.0 to 20

<b>Variable name</b>	VCDiameter	<b>GroupID</b>	1	<b>AttributeID</b>	39
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#### Length of the venous catheter

**Definition** Total length of the specialised central venous catheter (vas cath) used for initiation of continuous renal replacement therapy

Do **NOT** record the length inserted into the patient

**Reason** To assess influence of vascular access on application of renal replacement therapy

**Format** Record in centimetres (whole number), e.g. 25cm

**Validation rule** Value expected if site of vein is internal jugular, subclavian or femoral and manufacturer name type is completed  
Validation check if outside range 5 to 30  
Warning if value missing

<b>Variable name</b>	VCLength	<b>GroupID</b>	1	<b>AttributeID</b>	40
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## Number of lumens of venous catheter

<b>Definition</b>	The total number of lumens of the specialised central venous inserted into patient for initiation of continuous renal replacement therapy  Not the number of lumens used for CRRT
<b>Reason</b>	To assess influence of vascular access on application of renal replacement therapy
<b>Format</b>	Record whole number, e.g. 4
<b>Validation rule</b>	Expected value if site of vein is internal jugular, subclavian or femoral and manufacturer type is completed Warning if number is outside normal range 1-4 Warning if missing value

<b>Variable name</b>	VCNoOfLumens	<b>GroupID</b>	1	<b>AttributeID</b>	41
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## Blood transfusion

<b>Definition</b>	Whether the patient received a blood transfusion at initiation of continuous renal replacement therapy. Either via blood priming of the circuit and /or by administration of a blood transfusion to the patient during the first hour of continuous renal replacement therapy Choose from <b>To circuit:</b> if CRRT circuit was primed with blood prior to connection to the patient or <b>To patient:</b> if patient received a blood transfusion during first hour of continuous renal replacement therapy <b>Both:</b> the CRRT circuit was primed with blood prior to connection to the patient and the patient received a blood transfusion during the first hour of continuous renal replacement therapy <b>None:</b> if the CRRT circuit was clear primed and patient was not exposed to a blood transfusion during the first hour of continuous renal replacement therapy
<b>Reason</b>	To assess incidence of blood exposure to patients receiving continuous renal replacement therapy
<b>Format</b>	Tick one <b>To circuit</b> <b>To patient</b> <b>Both</b> <b>None</b>
<b>Validation rule</b>	Warning if None is selected and the patient received CVVH, CVVHD or CVVHDF

<b>Variable name</b>	BloodTransfusion	<b>GroupID</b>	1	<b>AttributeID</b>	42
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## Continuous Venovenous Haemofiltration (CVVH)

<b>Definition</b>	Continuous venovenous haemofiltration used during this PICU admission.
<b>Reason</b>	To review current practice, inter-unit variability and outcomes by modality
<b>Format</b>	Tick box if CVVH mode used during this PICU admission
<b>Validation rule</b>	If type of CRRT machine is specified one or more modes of CRRT (CVVH, CVVHD, CVVHDF) are expected

<b>Variable name</b>	CVVH	<b>GroupID</b>	1	<b>AttributeID</b>	43
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## Maximum blood flow

<b>Definition</b>	If CVVH given state the maximum blood flow recorded during all periods of CVVH, during this PICU episode
<b>Reason</b>	To establish usual blood flow rates used/achieved during CVVH
<b>Format</b>	ml/min rounded up to nearest whole value, e.g. 350.
<b>Validation rule</b>	Value expected value if CVVH selected. Validation check if outside range 20 to 430 Warning if missing value.

<b>Variable name</b>	CVVHMaxBloodFlow	<b>GroupID</b>	1	<b>AttributeID</b>	44
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## Maximum ultrafiltrate flow

<b>Definition</b>	If CVVH given state the maximum ultrafiltrate flow recorded during all periods of CVVH, during this PICU episode
<b>Reason</b>	To establish usual ultrafiltration rates achieved during CVVH and to assess dose effects on outcome
<b>Format</b>	ml/hour rounded up to the nearest whole value; e.g. 450
<b>Validation rule</b>	Expected value if CVVH selected Validation check if outside range 10-2000ml/hr Warning if missing value

<b>Variable name</b>	CVVHMaxUltrafiltrateFlow	<b>GroupID</b>	1	<b>AttributeID</b>	45
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## Continuous Venovenous Haemodialysis (CVVHD)

<b>Definition</b>	Continuous venovenous haemodialysis used during this PICU admission.
<b>Reason</b>	To review current practice, inter-unit variability and outcomes by modality
<b>Format</b>	Tick box If CVVHD mode used during this PICU admission
<b>Validation rule</b>	If type of CRRT machine is specified one or more modes of CRRT (CVVH, CVVHD, CVVHDF) are expected

<b>Variable name</b>	CVVHD	<b>GroupID</b>	1	<b>AttributeID</b>	46
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## Maximum blood flow

<b>Definition</b>	If CVVHD given state the maximum blood flow recorded during all periods of CVVHD, during this PICU admission.
<b>Reason</b>	To establish usual blood flow rates used/achieved during CVVHD
<b>Format</b>	ml/min rounded up to nearest whole value, e.g. 350
<b>Validation rule</b>	Expected value if CVVHD selected Validation check if outside range 20-450 Warning if missing value

<b>Variable name</b>	CVVHDMaxBloodFlow	<b>GroupID</b>	1	<b>AttributeID</b>	47
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## Maximum dialysate flow

<b>Definition</b>	If CVVHD given state the maximum dialysate flow recorded during CVVHD, during this PICU admission.
<b>Reason</b>	To establish usual dialysis rates achieved during CVVHD and to assess dose effects on outcome
<b>Format</b>	ml/hr rounded up to the nearest whole value, e.g. 450.
<b>Validation rule</b>	Expected value if CVVHD selected Validation check if outside range 10 to 8000 Warning if missing value

<b>Variable name</b>	CVVHDMaxUltrafiltrateFlow	<b>GroupID</b>	1	<b>AttributeID</b>	48
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## Continuous Venovenous Haemodiafiltration (CVVHDF)

<b>Definition</b>	Continuous venovenous haemodiafiltration used during this PICU admission.
<b>Reason</b>	To review current practice, inter-unit variability and outcomes by modality
<b>Format</b>	If CVVHDF mode used during this PICU admission tick box
<b>Validation rule</b>	If type of CRRT machine is specified one or more modes of CRRT (CVVH, CVVHD, CVVHDF) are expected

<b>Variable name</b>	CVVHDF	<b>GroupID</b>	1	<b>AttributeID</b>	49
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### Maximum blood flow

<b>Definition</b>	If CVVHDF given state the maximum blood flow recorded during all periods of CVVHDF, during this PICU admission.
<b>Reason</b>	To establish usual blood flow rates used/achieved during CVVHDF
<b>Format</b>	ml/min rounded up to nearest whole value, e.g. 350
<b>Validation rule</b>	Expected value if CVVHDF selected Validation check if outside range 20-450 Warning if missing value

<b>Variable name</b>	CVVHDFMaxBloodFlow	<b>GroupID</b>	1	<b>AttributeID</b>	50
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### Maximum ultrafiltrate flow

<b>Definition</b>	If CVVHDF given state the maximum ultrafiltrate flow recorded during all periods of CVVHDF, during this PICU admission.
<b>Reason</b>	To establish usual ultrafiltration rates achieved during CVVHDF and to assess dose effects on outcome
<b>Format</b>	ml/hr rounded up to the nearest whole value, e.g. 450
<b>Validation rule</b>	Expected value if CVVHDF selected Validation check if outside range 10-2000 Warning if missing value

<b>Variable name</b>	CVVHDFMaxUltrafiltrateFlow	<b>GroupID</b>	1	<b>AttributeID</b>	51
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### Maximum dialysate flow

<b>Definition</b>	If CVVHDF given state the maximum dialysate flow recorded during all periods of CVVHDF, during this PICU admission
<b>Reason</b>	To establish usual ultrafiltration rates achieved during CVVHDF and to assess dose effects on outcome
<b>Format</b>	ml/hr rounded up to the nearest whole value, e.g. 450
<b>Validation rule</b>	Expected value if CVVHDF selected Validation check if outside range 10-8000 Warning if missing value

<b>Variable name</b>	CVVHDFMaxDialysateFlow	<b>GroupID</b>	1	<b>AttributeID</b>	52
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### Type of Replacement fluid

<b>Definition</b>	The type of replacement fluid used during CVVH, CVVHHD or CVVHDF
<b>Reason</b>	To establish usual practice and inter-PICU variability
<b>Format</b>	Tick all that apply Haemosol BO Primasol 4 Accusol Other -specify
<b>Validation rule</b>	At least one value expected if CRRT (CVVH, CVVHD, CVVHDF) given Warning if missing value

<b>Variable name</b>	ReplacementFluidType	<b>GroupID</b>	1	<b>AttributeID</b>	53
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### Other Type of Replacement fluid (specify)

<b>Definition</b>	If other is specified the name of the other type of replacement fluid used during CVVH, CVVHHD or CVVHDF
<b>Reason</b>	Record the brand name and type of fluid given. If custom made state CUSTOM To establish usual practice and inter-PICU variability
<b>Format</b>	Free text
<b>Validation rule</b>	Expected if type of replacement fluid is other Warning if missing value

<b>Variable name</b>	OtherReplacementFluid	<b>GroupID</b>	1	<b>AttributeID</b>	54
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### Replacement fluid given (CVVH only)

<b>Definition</b>	During CVVH only: was replacement fluid given pre filter (predilution), post filter (post dilution) or a combination of both
<b>Reason</b>	To establish usual practice and inter-PICU variability and to assess effect on circuit survival
<b>Format</b>	Tick one box Pre dilution Post dilution Both
<b>Validation rule</b>	Expected value if CVVH given Warning if missing value

<b>Variable name</b>	ReplacementFluidGiven	<b>GroupID</b>	1	<b>AttributeID</b>	55
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### Anticoagulant used

<b>Definition</b>	Any medication given during CRRT episode in order to anticoagulate the CRRT circuit
<b>Reason</b>	To establish usual practice and inter-PICU variability and to assess the effect of anticoagulation practice on circuit survival.
<b>Format</b>	Tick yes or no
<b>Validation rule</b>	Expected value if CVVH, CVVHD, CVVHDF given Warning if missing and CVVH, CVVHD, CVVHDF given

<b>Variable name</b>	AnticoagulantUsed	<b>GroupID</b>	1	<b>AttributeID</b>	56
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### Type of anticoagulant *(tick all that apply)*

<b>Definition</b>	If anticoagulation used record the name of the anticoagulant(s) given Choose from Heparin Protacyclin Citrate Other - specify
<b>Reason</b>	To establish the efficacy and effects of different anticoagulation methods on CRRT circuit survival
<b>Format</b>	Tick all that apply: Heparin Protacyclin Citrate Other - specify
<b>Validation rule</b>	Expected value if Anticoagulant used is yes Warning if missing and Anticoagulant used is yes

<b>Variable name</b>	AnticoagulantHeparin	<b>GroupID</b>	1	<b>AttributeID</b>	58
<b>Variable name</b>	AnticoagulantProtacyclin	<b>GroupID</b>	1	<b>AttributeID</b>	59
<b>Variable name</b>	AnticoagulantCitrate	<b>GroupID</b>	1	<b>AttributeID</b>	60
<b>Variable name</b>	AnticoagulantOther	<b>GroupID</b>	1	<b>AttributeID</b>	61

### Other type of anticoagulant used

<b>Definition</b>	If other type of anticoagulant is specified record the type of anticoagulant used.
<b>Reason</b>	To establish usual practice and inter-PICU variability and to assess the effect of anticoagulation practice on circuit survival.
<b>Format</b>	Free text
<b>Validation rule</b>	Expected value if type of anticoagulant used is other Warning if value is missing and type of anticoagulant used is other

<b>Variable name</b>	OtherAnticoagulantName	<b>GroupID</b>	1	<b>AttributeID</b>	62
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### Target used

<b>Definition</b>	Record the anticoagulation target used, the value used to titrate, adjust and maintain anticoagulation treatment Choose from APTT (activated prothrombin time) ACT (activated clotting time) Ionized Calcium Other – specify
<b>Reason</b>	To establish the efficacy and effects of different anticoagulation methods on CRRT circuit survival
<b>Format</b>	Tick all that apply APTT (activated prothrombin time) ACT (activated clotting time) Ionized Calcium Other – specify
<b>Validation rule</b>	Expected value if Anticoagulant used is yes Warning if value is missing and Anticoagulant used is yes

<b>Variable name</b>	TargetAPTT	<b>GroupID</b>	1	<b>AttributeID</b>	64
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<b>Variable name</b>	TargetACT	<b>GroupID</b>	1	<b>AttributeID</b>	65
<b>Variable name</b>	TargetIonizedCalcium	<b>GroupID</b>	1	<b>AttributeID</b>	66
<b>Variable name</b>	TargetOther	<b>GroupID</b>	1	<b>AttributeID</b>	67

### Other target used

<b>Definition</b>	If other target is selected record the type of target used.
<b>Reason</b>	To establish the efficacy and effects of different anticoagulation methods on CRRT circuit survival
<b>Format</b>	Free text
<b>Validation rule</b>	Expected value if target used is selected as other Warning if value is missing and if target used is selected as other

<b>Variable name</b>	OtherTargetName	<b>GroupID</b>	1	<b>AttributeID</b>	68
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### Total duration of Continuous Renal Replacement Therapies (CVVH/CVVHD/CVVHDF)

<b>Definition</b>	The total duration of extracorporeal CRRT therapies (CVVH/CVVHD/CVVHDF) administered during single PICU admission, recorded in hours. Applies to the total number of hours spent on extra-corporeal CRRT therapies CVVH/CVVHD/CVVHDF only. Do not include hours where the patient is disconnected from the machine or there is machine malfunction while the patient is connected. Do not include hours spent on peritoneal dialysis
<b>Reason</b>	To establish relationship between duration of CRRT and patient outcome, and to evaluate cost implications of renal support in PICU
<b>Format</b>	Hours rounded to nearest hour, e.g. 15
<b>Validation rule</b>	Expected value if CVVH/CVVHD/CVVHDF selected Warning if value is missing and CVVH/CVVHD/CVVHDF selected

<b>Variable name</b>	TotalDurationOfCRRT	<b>GroupID</b>	1	<b>AttributeID</b>	69
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### Total number of filters used

<b>Definition</b>	The total number of CRRT filters used during the delivery of CRRT therapies CVVH/CVVHD/CVVHDF during the PICU admission (include elective and non-elective filter changes)
<b>Reason</b>	Number of filter changes is a surrogate for CRRT circuit survival and allows us to assess impact of both patient and circuit factors on CRRT circuit survival
<b>Format</b>	Whole number, e.g. 23
<b>Validation rule</b>	Expected value if CVVH/CVVHD/CVVHDF selected. Warning if outside the range 1-20 or value is missing and CVVH/CVVHD/CVVHDF selected

<b>Variable name</b>	TotalNumberOfFiltersUsed	<b>GroupID</b>	1	<b>AttributeID</b>	70
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## Number of elective filter changes

<b>Definition</b>	The total number of CRRT filters changed electively The number of filters that were changed for elective reasons during CVVH/CVVHD/CVVHDF during this PICU admission e.g. patient moved to or from theatre, CRRT paused for Therapeutic Plasma exchange, CRRT paused for elective procedure
<b>Reason</b>	Number of filter changes is a surrogate for CRRT circuit survival.  The number of filters /circuits which failed may be calculated from the number of filters changed for elective reasons and the total number of filters used
<b>Format</b>	Whole number, e.g. 18
<b>Validation rule</b>	Expected value if CVVH/CVVHD/CVVHDF selected Warning if outside the range 1-20 or value is missing and CVVH/CVVHD/CVVHDF is selected

<b>Variable name</b>	NumElecFilterChanges	<b>GroupID</b>	1	<b>AttributeID</b>	71
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## RRT STATUS AT DISCHARGE

<b>Definition</b>	The status of the patient at time of discharge from paediatric intensive care Complete for all patients who received renal replacement therapy Choose from <b>No renal replacement therapy - patient discharged alive.</b> The patient is not receiving ongoing RRT and is discharged from PICU alive <b>No renal replacement therapy - patient died.</b> The patient died during this PICU admission <b>Ongoing renal replacement therapy</b> – select the type of ongoing RRT, choose from <b>Peritoneal dialysis</b> - record if ongoing RRT is peritoneal dialysis <b>Intermittent haemodialysis</b> – record if ongoing RRT is peritoneal dialysis <b>Other ongoing RRT</b> –specify type
<b>Reason</b>	To establish the relationship between need for renal replacement therapy and patient outcome; both in terms of survival and need for on-going renal support
<b>Format</b>	Tick appropriate box(s) <b>No RRT-patient discharged alive</b> <b>No RRT-patient died: patient died</b> <b>Ongoing RRT</b> – select and tick box for type of RRT <b>Peritoneal dialysis</b> <b>Intermittent haemodialysis</b> <b>Other</b> –specify
<b>Validation rule</b>	Required to select from No RRT or Ongoing RRT If Ongoing RRT required to select type of RRT Error if missing value

<b>Variable name</b>	RRTStatusAtDischarge	<b>GroupID</b>	1	<b>AttributeID</b>	72
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## Other Ongoing RRT

<b>Definition</b>	If other ongoing RRT is selected in RRT status at discharge specify the type of ongoing RRT
<b>Reason</b>	To establish the relationship between need for renal replacement therapy and patient outcome; both in terms of survival and need for on-going renal support
<b>Format</b>	Free text
<b>Validation rule</b>	Expected value if other Ongoing RRT-Other is selected Error if missing value

<b>Variable name</b>	OtherRRTAtDischarge	<b>GroupID</b>	1	<b>AttributeID</b>	76
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## Form completed by

<b>Definition</b>	Insert the name of the person completing this form
<b>Reason</b>	For local use to assist with the follow-up of data queries
<b>Format</b>	Free text

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## Appendix A –Missing Values

### Missing PICANet Renal Dataset Values

For patients receiving continuous renal replacement therapy data for all fields included in the renal dataset should be available. Missing refers to data which has never been recorded and cannot be calculated from other available data; therefore it will never be available for this patient admission event.

When the following exception values are recorded on PICANet Web a validation query will be produced asking you to check the notes and charts to identify the missing information or provide confirmation that the data will never be available.

The information in the table below refers only to the summary renal dataset:

Initiation details	Field Description:	If missing record:
	Weight	999.9
	Baseline creatinine value	9999
	Highest creatinine value	9999
	Urine output	9999
	Ammonia value	9999
	Lactate	99.9
	Net fluid balance	99999
<b>Peritoneal Dialysis</b>	Total duration	9999
<b>Extra-corporeal CRRT</b>	Size of venous catheter	99.9
	Length of the specialised CVC	99
	Total no. of lumens	9
CVVH/CVVHD/CVVHDF	Maximum blood flow	999
	Maximum ultrafiltrate flow	999
CVVHDF only	Maximum dialysate flow	9999
	Total duration of CRRT therapies	9999
	Total no. of CRRT filters used	99
	No of elective filter changes	99

All other missing data values will show as missing data. Please complete as soon as possible.



## Appendix B –Data Collection Form

Placeholder for Data Collection Form