Patient study number	Patient Initials		



# APACHE II DATA FOR NON ICNARC (CMP) SITES

If your ICU participates in the ICNARC Case Mix Programme the patient's CMP number should be entered on page 7 in CRF 1. This booklet does not need to be completed.

Patient study number	Patient Initials
APACHE II DATA (NOTES)	

These data relate to the patient's first 24 hours in the Intensive Care Unit (ICU)

# **Past Medical History**

N.B There must be documented evidence that the condition existed or that the patient received therapy for the condition in the six months prior to admission to ICU.

Portal hypertension - Presence of oesophageal or gastric varices demonstrated by surgery, imaging or endoscopy or the demonstration of retrograde splenic venous flow by ultrasound. Do <u>not</u> include gastrointestinal bleeding without evidence of portal hypertension.

Hepatic encephalopathy - Episodes of hepatic encephalopathy, Grade 1 or greater

Grade 0 = No abnormality detected, Grade 1 = Slowness in cerebration, intermittent mild confusion and euphoria, Grade 2 = Confused most of the time, increasing drowsiness, Grade 3 = Severe confusion, rousable, responds to simple commands, Grade 4 = Unconscious, responds to painful stimuli.

Very severe cardiovascular disease - Fatigue, claudication, dyspnoea or angina at rest, where any activity increases symptoms. Symptoms must be due to myocardial or peripheral vascular disease. Functionally, this patient cannot stand alone, walk slowly or dress without symptoms.

Severe respiratory disease- Permanent shortness of breath with <u>light activity</u> due to pulmonary disease. Functionally this patient is unable to work and has SOB performing most normal activites of daily living e.g. walking 20m on level ground, walking slowly in the house, climbing one flight of stairs, dressing or standing.

Home ventilation - Has used or uses home ventilation within 6 months of admission.

Ventilation defined as where all, some, or a portion of the breaths (pressure support) are delivered by a mechanical device. Ventilation can be simply defined as a treatment where some or all of the energy required to increase lung volume during inspiration is supplied by a mechanical device. <a href="CPAP">CPAP is excluded</a>.

Chronic renal replacement - Currently requires chronic renal replacement therapy (either chronic haemodialysis, chronic haemofiltration or chronic peritoneal dialysis) for irreversible renal disease.

HIV - definite diagnosis of HIV infection - positive HIV test confirmed by an accredited microbiology laboratory

AIDS -HIV positive and has had an AIDS-defining illness (e.g. Pneumocystis carinii (P.jiroveci) pneumonia, Karposi's sarcoma, lymphoma, tuberculosis, toxoplasma infection).

Steroid treatment - Has received ≥ 0.3 mg kg<sup>-1</sup> prednisolone or an equivalent dosage of another corticosteroid, daily for the 6 months prior to admission to your unit.

Radiotherapy Has received externally administered radiotherapy, <u>excluding</u>: radiotherapy for non-invasive skin tumours; enteral or parenteral radioisotope therapy; radioactive implants; radiotherapy for prevention of heterotopic bone formation.

Chemotherapy - Has received drug treatment resulting in a lower resistance to infection (e.g drug treatment for malignancy, vasculitides, rheumatoid arthritis, inflammatory bowel disease). Excludes corticosteriods alone.

Metastatic disease - Distant (not regional lymph node) metastates, documented by surgery, imaging or biopsy.

Acute myelogenous/lymphocytic leukaemia or multiple myeloma - evident in the 6 months prior to admission to your ICU.

Chronic myelogenous/lymphocytic leukaemia - evident in the six months prior to admission to your ICU.

Lymphoma - Has active lymphoma, documented by surgery, imaging or biopsy.

Congential immunohumoral or cellular immune deficiency – documented state. Examples include Common Variable Immunodeficiency (CVID), agammaglobulinaemia including X linked (XLA), severe combined immunodeficiency (SCID), Chronic Granulomatous Disease, IgA deficiency, IgG deficiency, functional antibody deficiency, hyper IgE syndrome, Wiskott Aldrich syndrome, Chronic Mucocutaneous Candidiasis (CMCC), DiGeorge syndrome, Ataxia Telangiectasia, Leucocyte Adhesion Defect, Complement deficencies, C1 Esterase inhibitor deficiency, Kostmann's syndrome.

Page 3 ADAPT-Sepsis APACHE II Version1.0 19-Dec-2017

	number  ATA: FIRST 24 HOURS IN ICU	Patien	t Initials
	e this section if your ICU <u>does not</u> contribute to the ICNARC case mix p lete for first 24 hours in your ICU ( <u>not</u> the first 24 hours in the trial)	rogramn	ne.
12.1 Date and	time of admission to Critical Care Unit		
D D -	M M M - 2 0 Y Y H H : M M		
12.2 Was the	patient admitted to ICU directly from the operating theatre/recovery area in	n your ho	spital?
	Yes No		
If Yes cross or	ne box only:		
	Emergency (resuscitation is simultaneous with surgical treatment)		
	Urgent (surgery as soon as possible after resuscitation)		
	Scheduled (early surgery but not immediately life saving)		
	Elective (surgery at a time to suit both patient and surgeon)		
12.3 Does the	patient have a past medical history of one or more of the conditions listed be	low?	
condition in th	e aix mantha naigh ta admission to your ICLL This documentary evidence r		
recorded in the	e six months prior to admission to your ICU. This documentary evidence renotes either prior to admission or at admission to your ICU.  Disse listed below, cross here:		
recorded in the	e notes either prior to admission or at admission to your ICU.	Cross one	for each
recorded in the	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?		
recorded in the  If 'No' to all the  Past medical  Biopsy prove  Portal hypert	e notes either prior to admission or at admission to your ICU.  be listed below, cross here:  history present?  n cirrhosis ension	Cross one	for each
recorded in the  If 'No' to all the  Past medical  Biopsy prove  Portal hypert  Hepatic ence	e notes either prior to admission or at admission to your ICU.  Discontinuous listed below, cross here:  history present?  n cirrhosis ension phalopathy	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence	e notes either prior to admission or at admission to your ICU.  Disse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi	e notes either prior to admission or at admission to your ICU.  be listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila	e notes either prior to admission or at admission to your ICU.  be listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila	e notes either prior to admission or at admission to your ICU.  be listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS	e notes either prior to admission or at admission to your ICU.  Disse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr	e notes either prior to admission or at admission to your ICU.  be listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap Metastatic dis	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)  // // // // // // // // // // // // /	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap Metastatic dia Acute myelog	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)  y sease genous / lymphocytic leukaemia or multiple myeloma	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap Metastatic dia Acute myelog	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)  // // // // // // // // // // // // /	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap Metastatic di Acute myelog Chronic mye Lymphoma	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)  y sease genous / lymphocytic leukaemia or multiple myeloma	Cross one	for each
Past medical Biopsy prove Portal hypert Hepatic ence Very severe Severe respi Home ventila Chronic rena HIV AIDS Steroid treatr Radiotherapy Chemotherap Metastatic dia Acute myelog Chronic mye Lymphoma Congenital in	e notes either prior to admission or at admission to your ICU.  pse listed below, cross here:  history present?  n cirrhosis ension phalopathy cardiovascular disease ratory disease tion I replacement  ment (daily for six months)  // // // // // // // // // // // // /	Cross one	for each

## **NOTES (CONT.)**

## This is data relating to the patients first 24 hours in the Intensive Care Unit (ICU).

#### Arterial Blood Gas (ABG) with lowest PaO<sub>2</sub>

Lowest PaO<sub>2</sub> and associated values from the same ABG measured and recorded in the first 24 hours in your ICU.

#### Arterial Blood Gas (ABG) with lowest pH (or highest H+)

Lowest pH (or highest H+) values with their associated PaCO<sub>2</sub> value from the same ABG measured and recorded in the first 24 hours in your ICU.

Intubated is defined as a laryngeal mask, an endotracheal, endobronchial or tracheostomy tube in place.

## **Temperature**

Central Sites: tympanic membrane, nasopharyngeal, oesophageal, rectal, pulmonary artery, bladder. All other sites are regarded as non-central.

#### **Blood pressure (BP)**

Enter paired systolic and diastolic readings at the most extremes. (If there is a decision to be made between 2 readings take the one that gives the most extreme MAP). If BP is unmeasurable / undetectable enter zero.

#### Heart Rate (HR)

For patients who are paced, record the actual measured ventricular rate. HR should not be recorded during periods of iatrogenic disturbance eg. physiotherapy, turning, periods of crying etc. If undetectable enter zero.

# Non-ventilated/Ventilated Respiratory Rate

Ventilated: when all, some, or a portion of the breaths (pressure support) are delivered by a mechanical device.

CPAP and ECMO are considered <u>non – ventilated</u>. Ventilated respiratory rate should be the sum of both ventilated and spontaneous breaths in a minute.

#### Serum sodium / potassium / creatinine / bicarbonate / haemoglobin / white blood cell count

Must be <u>measured values</u> from <u>laboratory test results only</u>, *not* estimated from blood gas analyser. If no laboratory value is available from first 24 hours of admission, a pre-admission value measured within the 4 hours prior to admission may be used. For white blood cell count, the effects of steroids, inotropes and splenectomy are ignored.

#### **Urine output**

No account is taken of the effect of diuretics.

#### Assessment of Glasgow Coma Score (GCS)

Only GCS assessed when the patient is free from the effects of sedative and/or paralysing or neuromuscular blocking agents are valid. For patients sedated or paralysed for part of the first 24 hrs, give the lowest GCS prior to sedation or during the periods they were free of drug effects. For patients sedated for whole 24 hr ICU period, give lowest in-hospital GCS in the 24hr period prior to sedation.

The best motor response		The best verbal response		
Obeys verbal command	6	Orientated and converses	5	
Localises pain	5	Disorientated and converses	4	
Flexion withdrawal	4	Inappropriate words	3	
Flexion-abnormal/decorticate rigidity	3	Incomprehensible sounds	2	
Extension/decerebrate rigidity	2	No response	1	
No response	1	If a patient is <b>intubated</b> , use clinical judgement to score verbal response as follows:		
The best eye opening response				
Spontaneous	4	Appears orientated	5	
To verbal command	3	Responsive but ability to converse questionable	3	
To pain	2	Generally unresponsive	1	
No response	1		l	

Page 5 ADAPT-Sepsis APACHE II Version1.0 19-Dec-2017

Patient study n	umber			Patient Initi
Please comple	te for the first 24 hou	rs in	ICU (regardless of date of ra	ndomisation)
15.4 Arterial	blood gas with lowest Pa	<b>aO</b> <sub>2</sub> (i	f more than one, take sample with	highest FiO <sub>2</sub> )
	Lowest PaO <sub>2</sub>		kPa	
	FiO <sub>2</sub>			
	PaCO <sub>2</sub>		kPa OR	mmHg
	pH or [H+]	pH OR [H+] nmol I-1		
	Patient intubated? (✓)	Yes	: No:	
15.5 Arterial	blood gas with lowest pl	H or h	nighest [H+] (if more than one, take	sample with lowest PaCO <sub>2</sub> )
	Lowest pH (or highest [H+])		pH OR	[H+] nmol l-1
	PaCO <sub>2</sub>		kPa OR	mmHg
[if only one value of	available, record in lowest b	ox]	Lowest	Highest
15.7 Non-centra	-		° c	°c
	lood Pressure / paired		°c	
15.9 Heart Rate	ı		(beats min <sup>-1</sup> )	(beats min <sup>-1</sup> )
15.10 <b>Non-vent</b>	ilated Respiratory Rate		(breaths min <sup>-1</sup> )	(breaths min <sup>-1</sup> )
15.11 Ventilated	d Respiratory Rate (tota	al)	(breaths min <sup>-1</sup> )	(breaths min <sup>-1</sup> )
15.12 <b>Serum Sc</b>	odium		(mmol I <sup>-1</sup> )	(mmol I <sup>-1</sup> )
	ntassium		. (mmol I <sup>-1</sup> )	(mmol I <sup>-1</sup> )
15.13 <b>Serum Po</b>	nassium			
15.13 <b>Serum Po</b>			(μmol I <sup>-1</sup> )	(µmol l-1)
	reatinine		(μmol I <sup>-1</sup> )	
15.14 <b>Serum Cr</b>	reatinine carbonate			(μmol l <sup>-1</sup> )

Page 6

Patient study number		Patient Initials
15.18 <b>Total Urine Output</b> (ml):	Total for first 24 hrs	(ml)
	OR Total if stay < 24hrs	(ml)
15.19 Lowest pre-sedation Glasgow Coma Score (GCS)	Lowest GCS	
(If patient sedated for whole 24 hr ICU period, give lowest in-hospital GCS in the 24hr period pre-sedation)	Insufficient data (✓)  (GCS will be scored as 15)	
15.20 Please describe the primary real	ason for admission to the	ICU
As assessed during first 24hrs (not necessarily detailed description and/or identify:	diagnosis). What prompted their a	dmission to ICU? Please provide a
<ul> <li>Body system: eg. Respiratory, cardiova</li> <li>Anatomical site: eg. Lungs, upper airw</li> <li>Physiological/pathological process: epoisoning, self-intoxication or poisoning</li> <li>Condition: eg. Lung collapse or atelecte myxoma, trauma to aortic valve etc.</li> </ul>	ay and trachea, coronary arteries, eg. Haemorrhage, infection, traum, , inflammation, obstruction etc.	a, accidental intoxication or
Body system:		
Anatomical site:		
Physiological/pathological process:		
Condition:		
Office use only: ICNARC coding method:		
Name of Investigator completing this form		
Signature PRINT NAME	Date form compl	
Page 7 ADA	APT-Sepsis APACHE II	Version1.0 19-Dec-2017



Page 8 ADAPT-Sepsis APACHE II Version1.0 19-Dec-2017