A gap existed between physicians' perceptions and performance of pain, agitation-sedation and delirium assessments in Chinese intensive care units

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Additional file 1:

Case report form for cross-sectional investigation

Sedation practice in brain-injured patients in the intensive care units in China: a point prevalence study and a questionnaire survey

Case Report Form

Investigation site		
Investigator		
Screen No.		
Recruitment No.		
Medical record No.		
Abbreviation name		
Date	2010-00-00	

Illustrations

- This is a multicenter one-day point prevalence study combined with questionnaire survey.
- The protocol was approved by the Institutional Review Board of Beijing Tiantan Hospital, Capital Medical University. Written informed consent was obtained from each patient or their next of kin. During the study period, no attempt was made to alter or affect the routine clinical practice of each participating ICU.
- All recorded data must be timely, accurate and complete.
- Every page and every item of the CRF must be completed.
- Fill " $\sqrt{}$ " in the " \square " to indicate selection. "NA" to indicate "cannot provide" or "not applicable".
- Patient's abbreviated name should be recorded in the form of left aligned acronym of Chinese phonetic alphabet. For example: Zhang Wei should be recorded as Z W, and Zhang Xiao-Wei should be recorded as Z X W.
- Numeric data should be recorded with decimals provided as \Box . \Box in the form.
- If error(s) occurs in filling data, please use ONE strikethrough line and refill the correct data, and sign name of the corrector and date of correction.
 - Do not cover the original data.
 - Do not use the eraser or correction fluid.
 - Do not cross more than one strikethrough line.
- If investigators have any questions during the study, please contact Drs. Jian-Xin Zhou, Linlin Zhang:

• Dr. Jian-Xin Zhou: 010 59975098

Dr. Linlin Zhang: 010 59978451

Inclusion criterion: All adult patients admitted to the participating ICUs during the on-site screening,			
rega	ardless of the primary diagnosis.		
Excl	lusion criteria: please check the hospital records and	examine 1	the patient for exclusion:
•	Age < 18-years-old	☐ Yes	□No
•	Less than 24 hours of ICU stay before screening	☐ Yes	□No
•	Taking part in other studies	☐ Yes	□No
•	Refusing to participate	☐ Yes	□No

If there is no exclusion criterion, please go next page.

GCS at the ICU admission

APACHE II at the ICU admission

Case Report Form		
Investigator recruitment		
Data collection from hospital records		
Sex	□ male □ female	
Age	□□ years	
Height	□□□ cm	
Weight	$\square\square\square$ kg	
Date of admission to the hospital	201□/□□MM/□□DD	
Date of admission to the ICU	201□/□□MM/□□DD	
Primary diagnosis		
Brain injuries	☐ no ☐ yes, if yes, please tick one box bel	ow
Types of brain injuries	Traumatic brain injury	□yes □no
	Brain tumors	□yes □no
	Ischemic stroke	□yes □no
	Spontaneous intracerebral hemorrhage	□yes □no
	Subarachnoid hemorrhage	□yes □no
	Idiopathic epilepsy	□yes □no
	Intracranial infection	□yes □no
	Hypoxic-ischemic encephalopathy	□yes □no
	Others, please indicate	
Medical history	Hypertension	□yes □no
	Coronary artery disease	□yes □no
	Diabetes	□yes □no
	Chronic obstructive pulmonary diseases	□yes □no
	Chronic kidney diseases	□yes □no
	Ischemic stroke	□yes □no
	Alcohol abuse	□yes □no

History of smoke

 $\Box\Box$ points

Others, please indicate_

 $\Box\Box$ points: E (eyes) $\Box;$ V (verbal) $\Box;$ M (motor) \Box

□yes □no

Data collection from nursing records during the 24 hours prior to enrollment

Status

SOFA within 24 hours prior to enrollment	□□ points
Artificial airways	☐ oral ☐ nasal ☐ tracheostomy ☐others, please indicate:
Mechanical ventilation	□ no □ invasive □ non-invasive
	if yes, please tick one box below
	Date of start: $201\square/\square\squareMM/\square\squareDD$
	$\Box PCV \ \Box P-A/C \ \Box VCV \ \Box V-A/C \ \Box IPPV \ \Box PRVC \ \Box BIPAP$
	□APRV □P-SIMV □V-SIMV □PSV □CPAP □ others,
	please indicate:
	PEEP: □□cmH ₂ O; FiO ₂ : □□% VT: □□□ml
Arterial lines	□ yes □ no
Central venous catheters	□ yes □ no
ICP monitor	☐ no ☐ yes, please record: ☐☐mmHg
Any types of drainage tubes and colostomy	□ventricle □lumbar □epidural □subdural □thoracic
	□intraperitoneal □pelvic □mediastinal □gastrostomy
	□intestinal colostomy □bladder colostomy
	□others, please indicate:
Physical restraints	□ yes □ no
Body temperature control	☐ no ☐ physical cooling for hyperthermia
	☐ hypothermia therapy

Data collection from nursing records during the 24 hours prior to on-site investigation

Assessments

Pain assessment	☐ no ☐ yes, please indicate		
	□ СРОТ	□□ points	□□/d
	□VAS	□□ points	□□/d
	□ FPS	□□ points	□□/d
	□NRS	□□ points	□□/d
	☐ others, please indicate:	□□ points	□□/d
Agitation and sedation	☐ no ☐ yes, please indicate		
assessments	□ RASS	□□ points	□□/d
		□□ points	□□/d
	□ Ramsay	□□ points	□□/d
	□ others, please indicate:	□□ points	□□/d
Delirium assessment	☐ no ☐ yes, please indicate		
	□ CAM-ICU	□ yes □ no	□□/d
		□ yes □ no	□□/d
	□ others, please indicate:	□ yes □ no	□□/d

Data collection from nursing records during the 24 hours prior to on-site investigation

Medications

Analgesics	☐ no ☐ yes, please indicate			
	☐ Fentanyl	□ IV □ IM □ PO	□□□□µg/d	
	☐ Sufentanil	□ IV □ IM □ PO	□□□□µg/d	
	☐ Remifentanil	□ IV □ IM □ PO	□□□□μg/d	
		□ IV □ IM □ PO	□□mg/d	
	☐ Pethidine	□ IV □ IM □ PO	□□mg/d	
	☐ Tramadol	□ IV □ IM □ PO	□□mg/d	
	☐ Dezocine	□ IV □ IM □ PO	□□mg/d	
	☐ Butorphanol	□ IV □ IM □ PO	□□mg/d	
	☐ Flupiprofen ester	□ IV □ IM □ PO	□□mg/d	
	☐ Paracetamol oxycodone	□ IV □ IM □ PO	□□mg/d	
	☐ Acetaminophen	□ IV □ IM □ PO	□□mg/d	
	☐ others, please indicate:	□ IV □ IM □ PO	□□mg/d	
	☐ others, please indicate:	□ IV □ IM □ PO	□□mg/d	
Sedatives	□ no □ yes, please indicate			
	☐ Midazolam	□ IV □ IM □ PO	□□mg/d	
	☐ Propofol	□ IV □ IM □ PO	□□mg/d	
	☐ Dexmedetomidine	□ IV □ IM □ PO	□□µg/d	
	☐ Diazepam	□ IV □ IM □ PO	□□mg/d	
	□ Lorazepam	□ IV □ IM □ PO	□□mg/d	
	☐ Estazolam	□ IV □ IM □ PO	□□mg/d	
	☐ others, please indicate:	□ IV □ IM □ PO	□□mg/d	
	☐ others, please indicate:	□ IV □ IM □ PO	□□mg/d	
Muscle relaxants	☐ no ☐ yes, please indicate	•	•	
	☐ Rocuronium	IV	□□mg/d	
	☐ Vecuronium	IV	□□mg/d	
	☐ Pancuronium	IV	□□mg/d	
	☐ Cis-atracurium	IV	□□mg/d	
	☐ others, please indicate:	IV	□□mg/d	
Anti-delirium	☐ no ☐ yes, please indicate	•	•	
	☐ Haloperidol	□ IV □ IM □ PO	□□mg/d	
	☐ Chlorpromazine	□ IV □ IM □ PO	□□mg/d	
	☐ Promethazine	□ IV □ IM □ PO	□□mg/d	
	☐ Olanzapine	□ IV □ IM □ PO	□□mg/d	
	□ Risperidone	□ IV □ IM □ PO	□□mg/d	
	☐ others, please indicate:	□ IV □ IM □ PO	□□mg/d	

Outcome measures	
Type of follow-up (please fill " $$ " one)	
☐ Discharge from the hospital	
☐ Death in the ICU	
☐ Death in the hospital	
☐ At 60 days after enrollment	

Please review the ICU and hospital records

During ICU stay	Accidental removal of the catheter	□no □yes, please indicate: □□times	
after enrollment Duration of mechanical ventilation		□□□days	
	Hospital acquired infections	□Pneumonia □Urinary tract infection	
		□Intracranial infection □Surgical site infection	
		□Bacteremia □CRBSI	
		□others, please indicate:	
	Sepsis	□no □Sepsis □Septic shock	
	Survival when discharge from ICU	□yes □no □have not discharge yet	
	Date of discharge from ICU	If yes, please indicate :201□/□□MM/□□DD	
	ICU length of stay (LOS)	□□□□days	
During hospital	Survival when discharge from hospital	□yes □no □have not discharge yet	
stay after	Date of discharge from hospital	If yes, please indicate: 201□/□□MM/□□DD	
enrollment	Hospital LOS	□□□□days	
	Hospital costs	□□□□□□□.□□RMB	
	GOS	If brain-injured, please indicate: □points	