

Selected topics in Critical Care and Perioperative Medicine

Metabolismo

"Disfunzione cognitiva postoperatoria e delirium"



Marco Rossi

Istituto di Anestesia e Rianimazione

UOC Anestesia delle Chirurgie Specialistiche e Terapia del Dolore 1

Fondazione Policlinico Universitario IRCCS A. Gemelli

UCSC Roma



Neurocognitive disorders occur frequently in the community with 14 to 48% aged >70 yr suffering **mild cognitive impairment** and an additional 10% suffering **dementia**

> 30% of individuals >65yrs have surgery annually in western countries



Should general anaesthesia be avoided in the elderly?

C. Strøm,¹ L. S. Rasmussen² and F. E. Sieber³

Anaesthesia 2014, 69 (Suppl. 1), 35–44

Hospitalized older surgical patients are faced with many challenges during their **journey** through surgery and the recovery period



- fasting
- opioid analgesics
- anesthetic agents
- intraoperative blood loss
- postoperative pain
- nausea and vomiting
- unfamiliar hospital environment
- immobility during the perioperative period

The trajectories



The burden



1 week after surgery in adults >65yrs, with no difference in rates based on the type of surgery and/or anesthetic



What is postoperative delirium?

SUBTYPES OF DELIRIUM

Hyperactive delirium



Mixed form of delirium



More frequent Worst Prognostic bad

Hypoactive delirium



emergence delirium

mental disturbance during the recovery from general anesthesia into the recovery room/PACU primarily within the first 30 minutes of recovery time and typically resolves within an hour of onset

- Hallucinations
- Confusion
- restlessness
- moaning
- involuntary physical activity

- 5% to 10% of general surgery patients of all ages
- 41% of older adults

Pediatric Emergence Delirium





Predisposing and precipitating factors for postoperative delirium



How to diagnose delirium?

Confusion Assessment Method CAM

the most commonly used screening tool in research high sensitivity and specificity (between 90–100%)

long form with 10 sections

short form with 4 four domains commonly used in clinical practice

- acuity/fluctuation
- attention
- thinking
- level of consciousness



Clinical outcomes in older surgical patients with mild cognitive impairment



560 non-demented adults ≥70 years old

- **Delirium alone** associated with adverse postoperative outcomes and longer LOS
- MCI associated with increased risk of delirium after surgery and delirium severity
- **MCI without delirium** not associated with risk of poor postoperative outcomes
- **MCI with delirium** suggests synergistically higher risk of new cognitive impairments and more often discharged to a post-acute facility



healthcare costs

the 1 yr cost of care of patients **with delirium** estimated 2.5 times the cost of care for similar patients without delirium

Racine AM et al, Alzheimers Dement 2018; 14: 590–600

What is postoperative cognitive dysfunction?





significant reduction in **cognitive performance** from baseline following surgery and anesthesia, and diagnosed as subtle deficits in multiple core neurocognitive domains

- executive function
- attention
- verbal memory
- psychomotor speed
- visuospatial abstraction

in the **short** (7 days) and **medium** (1 to 3 months) term after anaesthesia and surgery, not fluctuating with lucid intervals

A problem of proper definition

methodological limitations of current literature



timing of administration of the tests

baseline status subjective complaints

lack of well-defined

rarely sought (ADLs)

significant heterogeneity in the type and number of tests administered

criteria or definition for change



Typical design of a study examining POCD with a profile of normal recovery and incomplete recovery across multiple postoperative time points

The diagnostic criteria for the perioperative cognitive changes should **not be differentiated** from neurocognitive disorders in the general population



The three DSM-5 pillars of diagnosis for neurocognitive disorders

- 1) Subjective complaint (participant, informant, clinician)
- Objective impairment/change (mild: 1 to 2 standard deviations below norms or controls; major: ≥2 standard deviations below norms or controls)
- Instrumental ADLs (for major NCD/dementia a decline in function is required)

Recommendations for the Nomenclature of Cognitive Change Associated with Anaesthesia and Surgery-2018

L. Evered, B. Silbert, D. S. Knopman, D. A. Scott, S. T. DeKosky, L. S. Rasmussen, E. S. Oh, G. Crosby, M. Berger, R. G. Eckenhoff, and The Nomenclature Consensus Working Group

Anesthesiology 2018; 129: 872-9





mild-NCD

- objective impairment of 1-2 SDs below controls/norms
- subjective complaint
- preserved ADL

Recommendations for POCD

modest (mild-NCD) or significant (major-NCD) cognitive decline from a previous level of performance in one or more cognitive domains up to **12 months**

- complex attention
- executive function
- · learning and memory
- language
- perceptual-motor
- social cognition





major-NCD

- objective impairment of ≥2 SDs below controls/norms
- subjective complaint
- impaired ADL



SHORT-TERM



LONG-TERM

Postoperative Delirium and Postoperative Cognitive Dysfunction

Retrospective cohort study of 560 older adults after noncardiac surgery in the Successful Aging after Elective Surgery study (SAGES)



- **Hypothesis:** Delirium is a significant risk factor for postoperative cognitive dysfunction
- Investigated the incidence of postoperative cognitive dysfunction within 6 months
- Evaluated the relationship between postoperative delirium and cognitive dysfunction



- During hospitalization, 134 of 560 participants (24%) developed delirium.
- Fewer than half (47%, 256 of 548) met the threshold for postoperative cognitive dysfunction at 1 month, but this proportion decreased at 2 months (23%, 123 of 536) and 6 months (16%, 85 of 528).

Agreement between delirium and postoperative cognitive dysfunction was poor (kappa < .08) and correlations were small (r < 0.16).

Postoperative cognitive dysfunction risk given history of postoperative delirium		
Post-operative interval	Relative risk; 95% CI	
1 month	1.34; 1.07-1.67	
2 months	1.08; 0.72-1.64	
6 months	1.21; 0.71-2.09	

Postoperative delirium and longer-term postoperative cognitive dysfunction may be different disorders.

Daiello LA, et al. ANESTHESIOLOGY. September 2019.

postoperative delirium is poorly predictive of **postoperative cognitive dysfunction** after 1 month of recovery and beyond, suggesting that POD and POCD are **separate clinical conditions**, rather than shared conditions reflecting two sides of the same coin



Trusted Evidence: Discovery to Practice

Potential theories on the development of delirium

most complementary rather than competing



synaptic impairment in susceptible patients, such as older adults or patients with previous head injury

brain region-specific vulnerability

enzymatic activation receptor activity

> Caspase GABAa Rs

neurotransmitter imbalance

altered proteins in the brain

alterations of synaptic

densities

- amyloid beta
- Tau protein
- Brain-derived neurotrophic factor

interruption of central cholinergic neurotransmission due to surgical stress and/or direct effect of anesthesia

Alzheimer's Disease Pathology as a risk factor for POD, POCD





Evered et al, Anesthesiology, 2016



110 surgical patients

Primary outcome- correlation between perioperative change in CSF AD biomarker (tau) and continuous cognitive change index (from preop to 6 weeks-1 year post-op)

Flow Cytometry Characterization of Cerebrospinal Fluid Monocytes in Patients With Postoperative Cognitive Dysfunction: A Pilot Study

Miles Berger, MD, PhD,*†‡ David M. Murdoch, MD,§ Janet S. Staats, BS, Cliburn Chan, MBBS, PhD, Jake P. Thomas,*¶ Grant E. Garrigues, MD,# Jeffrey N. Browndyke, PhD,‡** Mary Cooter, MSc,* Quintin J. Quinones, MD, PhD,* Joseph P. Mathew, MD, MHSc, MBA,* and Kent J. Weinhold, PhD ; for the MADCO-PC Study Team

Accepted for publication March 11, 2019

Flow cytometry panel to profile cerebrospinal fluid samples collected before and after major noncardiac surgery in 5 patients \geq 60 years of age who developed POCD and 5 matched controls who did not

Mono ratio = 0.41 Mono ratio = 0.70 Mono ratio = 0.40 Lymph Lymph Lymph 21.5 34.6 SSC FSC 0.0656 0.148 48.4 63.3 **CD16** 47 4 33.5 13.8 Mono:Lymph ratio CD192 (CCR2) ≥ 40000-Status 0.7 No POCD 0.6 POCD 1 30000 0.5 9 anla value 20000. 0.3 0.2 10000 0.1 Pre-op 24h 6wk Pre-op 24h 6wk postop postop postop postop

24 h postop

Preop

6 weeks postop

Patients who developed POCD

- increased CSF monocyte/lymphocyte ratio
- monocyte chemoattractant protein 1 receptor downregulation on CSF monocytes

24 hours after surgery

Where can we be active?



Brain protection

Preoperative recognition



Monitoring

Postoperative comprehensive rehabilitation Follow-up

Tailored anesthesia Environmental

JAMA Surgery | Review Enhanced Recovery After Surgery A Review

Olle Ljungqvist, MD, PhD; Michael Scott, MD; Kenneth C. Fearon, MD, PhD⁺



The journey

	Preadmission	Preoperative	Intraoperative	Postoperative
Surgery	Preadmission nutritional support Cessation of smoking Control alcohol intake	Selective bowel preparation	Minimal invasive surgery Minimize drains and tubes	Early removal of drains and tubes Stop intravenous fluids
Anesthesia	Medical optimization	Preoperative carbohydrates No NPO PONV prophylaxis	Regional analgesia Opioid-sparing anesthesia Balanced fluids Temperature control	Multimodal opioid-sparing Earl pain control Pain
Nursing	Preoperative information		Short acting	Early mobilization Early oral intake of fluids and solids Postdischarge follow-up

A combined strategy

Multimodal anesthesia



Multimodal analgesia





Active body warming



Depth of anesthesia monitoring



Recovery Rooom



Less tau proteins and caspase activation





attenuated excitotoxicity

- Inhibition of glutamate sensitive NMDA receptors
- Inhibition of postsynaptic glutamate receptor responses
- Activation of GABA A receptors



reducing sympathetic tone

Handbook of Clinical Neurology, Vol. 121 Neurologic Aspects of Systemic Disease Part III Jose Biller and Jose M. Ferro, Editors © 2014 Elsevier B.V. All rights reserved





MAC decline by $\approx 6\%$ per decade after 30 years of age



avoiding volatile anesthetic overdose by closely monitoring the **age-adjusted MAC fraction**



adjust end-tidal volatile anesthetic concentration during surgery

The shortcoming



major surgery and general anesthesia are unlikely to cause persistent POCD or incident dementia

Postoperative Cognitive Function Following General Versus Regional Anesthesia: A Systematic Review

Nicholas Davis, MD,* Melissa Lee, MD,* Albert Y. Lin, MD, MPH,* Lisa Lynch, MD,* Matthew Monteleone, MD,* Louise Falzon, PGDipInf,†[‡] Nighat Ispahany, MLS,[‡] and Susan Lei, MD*

References	Anesthetic Techniques	Outcomes at \geq POD 7
Asbjorn et al ¹¹	GA vs. EA	No difference in standard (<i>Z</i>) scores on psychological tests involving short-term, long-term, verbal, and visual memory, P > 0.05
Berggren et al ¹²	GA vs. EA	50% of EA group and 38% of GA group experienced mental status change, no significant difference $P > 0.05$
Bigler et al ¹³	GA vs. SA	No difference on abbreviated mental test between GA and SA groups, Student t test, $P > 0.05$
Campbell et al ¹⁴	GA vs. LA	No evidence of a change in the relationship between the LA and GA groups with time for all tests performed, analyzed using multivariate analysis of variance
Chung et al ¹⁵	GA vs. SA	No difference in mini-mental status exam score or geriatric mental exam score between or within groups, $P > 0.05$
Chung et al (1989) ¹⁶	GA vs. SA	No difference in mini-mental status exam score or geriatric mental exam score between or within groups, $P > 0.05$
Crul et al ¹⁷	GA vs. SA	<u>No</u> difference in 3 cognitive function tests using multivariate analysis of variance, P > 0.05
Ghoneim et al ¹⁸	GA vs. SA vs. EA	<u>No</u> difference in 17 cognitive function tests using multivariate analysis of variance, P > 0.05
Haan et al ¹⁹	GA vs. SA	No difference in 3 cognitive function tests using multivariate analysis of variance, P > 0.05
Hole et al ²⁰	GA vs. EA	0/29 patient in EA and 7/31 patients in GA with persistent mental changes, 2- sided Student <i>t</i> test or γ^2 test. $P < 0.01$
Jones et al ²¹	GA vs. SA	Of 7 cognitive function tests, significant difference in recognition and response times on choice reaction time test (improved in general anesthesia group, $P < 0.05$)
Karhunen and Jonn ²²	GA vs. LA	Two tests of memory were performed, patient in LA group scored significantly worse on Luria memory test, $P < 0.01$
Mandal et al ²³	GA vs. EA	Two cognitive function tests performed, gross difference at 7 d favoring regional anesthesia in the mini-mental status exam, $P < 0.01$
Nielson et al ²⁴	GA vs. SA	<u>No</u> difference in 8 cognitive function tests, P > 0.05
Riis et al ¹⁰	GA vs. EA vs. GEA	No difference in 13 cognitive function tests, data comparison via Z scores, P > 0.05
Williams- Russo et al ²⁵	GA vs. EA	<u>No</u> difference in 10 cognitive function tests, comparison via Student <i>t</i> test, P > 0.005



no substantial differences in cognitive outcomes between regional and general anesthetic techniques

"Intravenous versus inhalational maintenance of anaesthesia for postoperative cognitive outcomes in elderly people undergoing non-cardiac surgery"

Miller D et al, Cochrane Database Syst Rev. 2018 Aug 21;8:CD012317





28 RCTs 4507 randomized participants different types of surgery

- Cardiovascular
- Laparoscopic
- Abdominal
- Orthopaedic
- Ophthalmic

Inhalational maintenance agents

- Sevoflurane 19 studies
- Isoflurane 8 studies
- Desflurane 3 studies

no evidence of a difference in incidences of postoperative delirium according to type of anaesthetic maintenance agents (OR 0.59)

uncertain whether maintenance with propofolbased TIVA or with inhalational agents affect incidences of postoperative delirium, mortality, or length of hospital stay because certainty of the evidence was very low Association of excessive depth of anesthesia with poor outcomes especially in high-risk patients







REPORTS OF ORIGINAL INVESTIGATIONS

Relation between bispectral index measurements of anesthetic depth and postoperative mortality: a meta-analysis of observational studies



Processed electroencephalogram and evoked potential techniques for amelioration of postoperative delirium and cognitive dysfunction following non-cardiac and nonneurosurgical procedures in adults (Review)

Punjasawadwong Y, Chau-in W, Laopaiboon M, Punjasawadwong S, Pin-on P



Cochrane Database of Systematic Reviews 2018, Issue 5. Art. No.: CD011283





results from three studies (2051 participants) indicate that this could also reduce the incidence of POCD at three months from 9.1% to 6.4%.



GUIDELINES

European Society of Anaesthesiology evidence-based and consensus-based guideline on postoperative delirium



Stress-diathesis model for postoperative delirium

away from the hypothesis that anesthetic toxicity is largely responsible for negative cognitive effects



a **very high risk** group for POD with readily identified diathesis markers, at risk irrespective of the anesthetic course, but potentially worsened by a **poorly conducted anesthetic**



a **low risk** group with no or few diathesis markers who do not manifest POD **irrespective of anesthetic exposure** or duration of procedure

an **intermediate risk** group with evidence of premorbid diatheses placed at risk by an **unstable anesthetic course**

depressionanxiety disorders

Proposed treatments for POCD

protecting neurons during and promoting neuronal health before surgery



preventing the oxidative component of inflammation

blocking inflammation by inhibiting inflammatory mediators

Evidence From Human Studies

Promising results in animal studies, more modest results when applied to human populations

Novel therapies for POCD

pharmacological and behavioural

Parecoxib/COX-II Inhibitors



Perioperative cognitive protection

Behavioural and Nonpharmacological Strategies for Prevention of Delirium



Re-habilitation

- 1. Sensory enhancement (ensuring glasses, hearing aids, or listening amplifiers)
- 2. Mobility enhancement (ambulating at least twice per day if possible)
- 3. Cognitive orientation and therapeutic activities (tailored to the individual)
- 4. Pain control with scheduled acetaminophen if appropriate
- 5. Cognitive stimulation (if possible, tailored to the individual's interests and mental status)
- 6. Simple communication standards and approaches to prevent the escalation of behaviours
- 7. Nutritional and fluid repletion enhancement

Pre-habilitation

- 8. Sleep enhancement (daytime sleep hygiene, relaxation, nonpharmacological sleep protocol, and nighttime routine)
- 9. Medication review and appropriate medication management
- 10. Daily rounding by an interdisciplinary team to reinforce the interventions

American Geriatrics Society Expert Panel on Postoperative Delirium in Older Adults. Postoperative delirium in older adults: best practice statement from the American Geriatric Society. J Am Coll Surg. 2015;220:136-148

avoidance or reduction of benzodiazepines and anticholinergics

Best Practices for Postoperative Brain Health: Recommendations From the Fifth International Perioperative Neurotoxicity Working Group

Miles Berger, MD, PhD,* Katie J. Schenning, MD, MPH,† Charles H. Brown IV, MD, MHS,‡ Stacie G. Deiner, MD,§ Robert A. Whittington, MD,|| and Roderic G. Eckenhoff, MD,¶ for the Perioperative Neurotoxicity Working Group

Anesth Analg 2018;127:1406-13

Consensus Statement

"More studies are needed to evaluate the efficacy, feasibility, and cost-effectiveness of various strategies to assess short and long-term cognitive outcomes after hospital discharge, to optimally manage these disorders, and to clarify who should follow patients after surgery for these disorders and what patients should be told about the current understanding regarding recovery from these disorders"



Brief Co	ognitive Screening Tool	Is			
Tool/Test	Advantage	Disadvantage	Sensitivity ^a	Specificity ^a	Time to Administer
Minicog ¹⁴⁻¹⁸	Brief, minimal language, education, race bias	Use of different word lists may affect scoring	76–100 (54–100)	54-85.2 (43-88.4)	2-4 min
MoCA ¹⁹⁻²²	Can identify mild cognitive impairment, available in multiple languages	Education bias, limited published data	n/a	n/a	10–15 min
MMSE ^{18,23,24}	Widely used and studied	Subject to age and cultural bias, ceiling effects	88.3 (81.3–92.9)	86.2 (81.8–89.7)	7–10 min
Clock-drawing test ^{18,25}	Very brief	No standards for administration and scoring	67–97.9 (39–100)	69–94.2(54–97.1)	<2 min
Verbal Fluency Test18,26	Brief	Cut point not obvious	37-89.5 (19-100)	62-97 (48-99)	2–4 min
CODEX ^{27–29}	Brief	Less well studied	81%-93%	81%-85%	≤3 min



Valutazione del rischio di insorgenza del *delirium post*-operatorio (POD) nei pazienti sottoposti a chirurgia per *hip fracture*



Una complicanza maggiore dell'*hip fracture* nell'anziano è il *delirium* post operatorio (POD) con un'incidenza variabile, a seconda delle stime, tra il 13 e il 70%.

Endpoints

- L'endpoint primario dello studio è la comparsa di POD entro i primi 7gg.
- Gli endpoints secondari sono lo sviluppo di POCD, demenza di qualunque tipo di nuova insorgenza a 12 mesi, mortalità a 30gg, tempo di degenza postoperatoria (comprendente la riabilitazione eseguita all'interno del Policlinico).



Test neuropsicologi e scale usate nello studio

• **RASS: La Richmond Agitation-Sedation Scale** misura il livello di agitazione o sedazione di un paziente; ottenere un RASS score risulta il primo punto per somministrare il Confusion Assessment Method (CAM). Lo score va da +4 (combattivo o violento), passando per 0 (calmo e vigile, presta attenzione al caregiver) a -5 (non risvegliabile alla voce né alla stimolazione fisica)

• **CAM-ICU: il** *Confusion Assessment Method* – *intensive care unit* è uno strumento neuropsicologico che valuta la presenza di Delirium nei pazienti critici

• **MoCA: il** *Montreal Cognitive Assessment* è un test rapido di screening del deterioramento cognitivo lieve. Permette di verificare diverse aree funzionali: attenzione, concentrazione, funzioni esecutive, memoria, linguaggio, abilità visuo-costruttive, astrazione, calcolo, orientamento.

Valutazione dello stato di coscienza e monitoraggio del delirium

Fase 1 – Livello di Coscienza: RASS

Punteggio	Definizione	Descrizione		
+4	COMBATTIVO	Chiaramente combattivo, violento, imminente pericolo per sé o per lo staff	•	
+3	MOLTO AGITATO	Aggressivo, rischio evidente di rimozione invasività	SSE	
+2	AGITATO	Frequenti movimenti afinalistici, disadattamento alla ventilazione meccanica	RVAZ	
+1	IRREQUIETO	Ansioso ma senza movimenti aggressivi o vigorosi	ION	
0	SVEGLIO E TRANQUILLO	Comprende i periodi di sonno fisiologico) "	
-1	SOPOROSO	Non completamente sveglio, apre gli occhi allo stimolo verbale, mantiene il contatto visivo > 10 secondi) < v	
-2	LIEVEMENTE SEDATO	Brevi risvegli allo stimolo verbale, contatto visivo < 10 secondi	ERBAL	
-3	MODERATAMENTE SEDATO	Movimenti o apertura degli occhi allo stimolo verbale (ma senza contatto visivo)) ^{m o}	
Se F	RASS ≥ - 3 🗪 somminis	tra CAM-ICU (il paziente ha delirium oppure no?)		
-4	SEDAZIONE PROFONDA	Nessuna risposta allo stimolo verbale, movimenti o apertura occhi alla stimolazione fisica		
-5	NON RISVEGLIABILE	Nessuna risposta alla stimolazione tattile o dolorosa		
Se RASS ≤ -4 → RIVALUTA più tardi (paziente attualmente incosciente)				

Sessler, et al. AJRCCM 2002; 166:1338-1344.² Ely, et al. JAMA 2003; 289:2983-2991.³

CAM-ICU: Scheda di lavoro

Punto 1: Alterazione Acuta o Fluttuazione dello Stato Mentale	Punteggio	Segna se presente
Il paziente si presenta in modo diverso dal suo stato mentale di base? <u>OPPURE</u> Il paziente ha presentato fluttuazioni dello stato mentale nelle ultime 24 ore come evidenziato da una variazione in una scala di sedazione (i.e., RASS), di stato di coscienza (GCS), o in un precedente assessment sul delirium?	Se almeno una risposta è SI →	
Punto 2: Disattenzione		
Test 'Lettere' (in alternativa consulta il manuale per il test 'Immagini') Indicazioni. Dire al paziente: "Sto per leggerle una serie di 10 lettere. Mi stringa la mano quando dico la lettera A". Leggere le lettere dalla seguente lista con un tono di voce normale e costante ad intervalli di 3 secondi. S A V E A H A A R T Viene contato un errore quando il paziente non stringe la mano sulla lettera "A", o quando la stringe in risposta alle altre lettere	Numero di errori > 2 →	
Punto 3: Alterato Livello di Coscienza		
Il paziente è agitato, sedato o incosciente?	RASS \neq 0 \rightarrow	
Punto 4: Pensiero Disorganizzato		
Domande a cui si può rispondere solo Si/No, come ad esempio: 1. Un sasso galleggia nell'acqua? 2. Ci sono pesci nel mare? 3. Un chilo pesa più di due chili? 4. Si può usare il martello per piantare un chiodo? Errore: quando il paziente risponde in maniera scorretta alla domanda. Ordine semplice 5. Dire al paziente: "Mi mostri queste dita" (mostrare 2 dita); "Ora faccia lo stesso con l'altra mano" (senza mostrarle) se il paziente non riesce a muovere entrambe le braccia dire: "Aggiunga un altro dito" Errore: quando il paziente non è in grado di completare l'intero esercizio.	Numero totale di errori > 1 →	

Punto 1 +	Almeno uno (+)	Soddisfazione dei criteri \rightarrow	CAM-ICU Positivo (presenza di Delirium)
Punto 2 (+)	fra punto 3 e 4	Criteri non soddisfatti \rightarrow	CAM-ICU Negativo (assenza di Delirium)



VALUTAZIONE DEL RISCHIO DI INSORGENZA DEL DELIRIUM NEI PAZIENTI CON FRATTURA DI FEMORE PROSSIMALE



Fondazione Policiinico Universitario A. Gemelli Università Cattolica del Sacro Cuore

SCHEDA .	DI ARRUOL	AMENTO
----------	-----------	--------

	TER (DI)	101 050500101115	DOCT IS DO INVITO DOCCO.
SPAZIO ETICHETTA	(FARMACO e POSOLOGIA)	ADL PERFORMANCE	POST-IC, PZ INVIATO PRESSO:
		(0-6)	TIPO RR
	·		
(criterio <u>esclusione</u> all'arruolamento:	•		Valore Hb, SpO2 e NRS post- I.C.
eta < 65 anni)		(0-8)	g/dL %
	·		
		Tempo dall'arrivo in PS al I MOCA (ORE)	Complicanze post-operatorie (se SI, specificare quali):
		8-20 20-8	specificare quanj.
	• •	Tempo dall'arrivo in PS all'intervento	•
		chirurgico (ORE)	•
Indicare livello scolarità:	·		
		Valore Hb e SpO2 pre-operatorio	23. Valore Hb (mg/dL):
TRAUMA CRANICO		g/dL %	PARELE. PROMA GEORGIAN PARELE. TERES GEORGIAN Parel-LE.
SI NO			
SCOMPENSO	PROVENIENZA	TRASFUSIONE PRE-I.C.?	24. SpO2 all'uscita RR/TIPO. Necessita di ossigenoterania all'arrivo in reparto?
CUORE/POLMONE	CASA ALTRO RESIDENZA PROPRIA OSPEDALE PROTETTA	SI NO SESI, SPECIFICARE	% SI/NO
ACO IO (specificare)	CT (TO DI C (TTTT		
	GENERALE	ARRIVO IN SO	INDTROPI/VASOCOSTRITTORTIN INFUSIONE INTRA E POST-OPERATORI
		(0-10)	(se SI, specificare farmaco e posologia)
DEMENZA	• IA SI NO		SI NO
MODERATA-SEVERA		ANTIDOLORIFICI PRE-OPERATORI	
SI NO	• DM s NO	(se SI, specificare prescrizione e	
ASSENZA CAREGIVER	 Insulino-dipendente 	posologia)	•
SI NO	SI NO	•	
TABAGISMO e/o	CREATININA	•	Riabilitazione motoria post-I.C.
ADDICTION SI NO	>1.5 mg/dL	•	(specificare dove svolta e tipologia)
	SI NO		
HCV S NO			
PREGRESSO	• OSAS	Intervento svolto in sala urgenze?	Farmaci psicotropi peri-I.C.
TIA/STROKE		SI NO	:
(evenio/anno//eliquali)	 FA CRONICA 	<u> </u>	•
	SI NO		•
	 PORTATORE 	Anestesia eseguita (descrizione)	
	PM/ICD (se si,	Blocco fascia?	TEMPO DI DEGENZA post-I.C. E
	spiegare tipologia)		Numero gioni
	SI NO	SI NO	
DD D C D D C C C	-		
PREGRESSA	Ricovero pre-operatorio in	Necessità di trasfusione durante o nel	MORTALITA' BOST LC
CARDIOVASCOLARE O	(medico/chirurgico)	post-IC	MORIALITA POSI-I.C.
NECROCHIRCRGICA (tipe/anne)		SI NO	dopo
			giorni anno primo
		Se si, specificare:	anno anno
I	l	1	



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The tailoring

rigorous and routine preoperative cognitive screening

identifying preexisting cognitive impairment determining if the patient has a history of previous postoperative confusion or delirium documenting other episodes of delirium or confusion

assessing for low education

use of specific anesthetic regimens

- general versus regional anesthesia
- specific anesthetic agents
- blood transfusions
- systemic arterial pressure monitoring
- · depth of anesthesia
- normothermia



ART

ERAS postoperative protocols

