INTERNAL MEDICINE: Critical Care

ТЕХТВООК

Edited by O.Ya. BABAK, O.M. BILOVOL

RECOMMENDED

by the Academic Board of Kharkiv National Medical University as a textbook for students of higher education establishments — medical universities, institutes and academies

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The "Internal Medicine: Critical Care" provides the depth and breadth of coverage that reflects the complexity and expertise needed to practice emergency medicine successfully in today's fast—paced environments. It is an important contemporary clinical emergency care resource for students of higher education eshtablishments — medical universities, institutes and academies.

The textbook was published in the English language, illustrated with pictures and tables, which are easy to learn and to store in memory for a long time. This textbook also gives possibility to find answers quickly when you are faced with a difficult diagnosis or need the latest treatment recommendations, step-by-step guidelines and new pharmacologic considerations.

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Abbreviations

AAE	acquired angioedemoedema	CCS	Canadian Cardiovascular So-
ABG	artery bypass graft		ciety
ABG test	arterial blood gas test	CCU	critical care unit
ACC	American College of Cardiology	cGMP	cyclic guanosine monophos-
ACE	angiotensin-converting enzyme		phate
ACE-inhib	itor, ACEI angiotensin-conver-	CHF	chronic heart failure
	ting enzyme inhibitor	CHF	congestive heart failure
ACLS	advanced cardiac life support	CKD	chronic kidney disease
ACS	acute coronary syndrome	CMR	cardiac magnetic resonsnce
ACTH	adrenocorticotropic hormone	CNS	central nervous system
ADOI	acute dialysis quality initiative	CO,	carbon dioxide
AED	automated external defibrillator	COPD	chronic obstructive pulmonary
AF	atrial fibrillation		disease
AFL	atrial flutter	COX	cyclooxygenase
AHA	American Heart Association	СР	chest pain
AKI	acute kidney injury	CPAP	continuous positive airway pres-
ALT	alanine aminotransferase		sure
AMI	acute myocardial infarction	СРК	creatine phosphokinase
ANCA	antineutrophil cytoplasmic anti-	СРО	cardiogenic pulmonary oedema
	body	CPR	cardiopulmonary resuscitation
aPTT	activated partial thromboplastin	CRF	chronic renal failure
	time	CRT	continuous reaction time
ARB	angiotensin receptor blocker	CSF	cerebrospinal fluid
ARDS	adult respiratory distress syn-	СТ	computer tomography
	drome	CTI	cavotricuspid isthmus
AST	aspartate aminotransferase	cTn	cardiac-specific troponin
ATP	anti-tachycardia pacing	СТРА	computed tomographic pulmo-
ATP test	adenosine triphosphate test		nary angiography
ATV	automatic transport ventilator	CUS	compression ultrasonography
AVM	arteriovenous malformation	CVA	cerebrovascular accident
AV	atrioventricular	CVD	cardiovascular disease
AVNRT	atrioventricular node reentry	CVP	central venous pressure
	tachycardia	CVVHD	continuous venovenous haemo-
AVRT	atrioventricular reciprocating		diafiltration
	tachycardia	CXR	chest X-ray
BBB	bundle branch block	D5W	5 % dextrose in water
BiPAP	bi-level positive airway pressure	DBP	diastolic BP
BNP	B-type natriuretic peptide	DC cardioversion, DCC direct current car-	
BP	blood pressure		dioversion
BUN	blood urea nitrogen	DDAVP	desamino-D-arginine vasopres-
CAB	circulation, airway and brea-		sin
	thing	DIC	disseminated intravascular co-
CABG	coronary artery bypass grafting		agulation
C1-INH	C1-esterase inhibitor	DKA	diabetic ketoacidosis
CK-MB	creatine kinase MB	DM	diabetes mellitus
CAD	coronary artery disease	DSA	digital subtraction angiography
CBC	complete blood count	DVT	deep vein thrombosis
ССВ	calcium channel blocker	EAR	early allergic response

Abbreviations

ECG	electrocardiogram	
ECG	electrocardiography	
ECMO	extracorporeal membrane oxy-	
	genator	
ED	emergency department	
EDD	esophageal detector device	
EEG	electroencephalography	
EF	ejection fraction	
EGD	esophagogastroduodenoscopy	
EMS	emergency medical service	
EP	electrophysiological; electro-	
	physiology	
ERCP	endoscopic retrograde cholan-	
	giopancreatography	
ESC	European Society of Cardiology	
ESWL	extracorporeal shockwave litho-	
	tripsy	
FBAO	foreign-body airway obstruction	
FBC	full blood count	
FDA	Food and Drug Administration	
FEV1	forced expiratory volume in 1	•
	second	
FFP	fresh frozen plasma	
FHF	fulminant hepatic failure	
GABA	gamma-aminobutyric acid	
GCS	Glasgow coma scale	
GERD	gastroesophageal reflux	
GFR	glomerular filtration rate	
	gastrointestinal bleeding	
GpIIb/-III	a receptor glycoprotein IIb/-IIIa	1
	receptor	1
HAAF	hypoglycaemia-associated auto-	1
II. CEV	nomic failure	
HACEK	haemophilus actinobacillus car-	
	diobacterium eikenella and kin-	1
TIAE	gella	1
HAE	hereditary angioedemoedema	1
HDU	high dependency unit	
HE LI FADD	hepatic encephalopathy	
H-FABP	heart-type fatty acid-binding	
HFSA	protein Usert Feilure Seciety of Ameri	
пгза	Heart Failure Society of Ameri-	
HIT	ca	
1111	heparin-induced thrombocyto- paenia	1
HNC	hyperosmolar nonketotic coma	
HNS	hypothalamo-neurohypophysial	1
1113		1
носм	system hypertrophic obstructive cardio-	
1100.01	myopathy	1
HR	heart rate	
HTN	hypertension	
IABC	intra-aortic balloon catheter	
IABC	intra-aortic balloon pumping	
1/3/01	mara-aorae oanoon pumping	

ICD	implantable cardioverter-defi-
ICD	brillator
ICP	intracranial pressure
ICU	intensive care unit
IgE	immunoglobulin E
IGF	insulin growth factor
IHD	ischaemic heart disease
IHD	intermittent haemodialysis
n	iterleukin
IM	intramuscular
INR	international normalisation ratio
IPC	intermittent pneumatic com-
ISDN	pression isosorbide dinitrate
IU	international unit
IV	intravenous
IVC	inferior vena cava
IVP	intravenous push
IVP	intravenous pyelogram
IVU	intravenous urogram
JVD	jugulovenous distention
KUB	kidney-ureter-bladder
LAD	left axis deviation
LAR	late allergic response
LBBB	left bundle branch block
LGL-synd	rome Lown–Ganong–Levine
1 3 4 4	syndrome
LMA LMWII	laryngeal mask airway
LMWH LOLA	low-molecular-weight heparin L-ornithine L-aspartate
LULA	left ventricle
LVAD	left ventricular assist device
LVEDP	left ventricular end-diastolic
	pressure
LVEF	left ventricular ejection fraction
MAT	multifocal atrial tachycardia
MCS	mechanical circulatory support
МСТРА	multidetector CT pulmonary
	angiography
MDCT	multidetector computed tomog-
MDDD	raphy
MDRD	modification of diet in renal disease
MODS	multiple organ dysfunction syn-
MODS	drome
MR	mitral regurgitation
MRCP	magnetic resonance cholangio-
	pancreatography
MRI	magnetic resonance imaging
MRSA	methicillin-resistant staphylo-
	coccus aureus strains
MS	mental status
MTHFR	5,10-methylenetetrahydrofolate
	reductase
	F

Abbreviations

MVP	mitral valve prolapse
MVT	monomorphic ventricular
	tachycardia
NCPO	non-cardiogenic pulmonary
	oedema
NIPPV	noninvasive positive pressure
	ventilation
NIV	noninvasive ventilation
NOAC	new oral anticoagulant
NSAIDs	non-steroidal anti-inflammatory
10011105	drugs
NST-ACS	non-ST-segment elevation acu-
	te coronary syndrome
NSTEMI	non-ST-segment elevation
	myocardial infarction
NTG	nitroglycerin
OD	overdose
PaCO,	partial pressure of carbon diox-
rac0 ₂	ide in arterial blood
PAD	
	peripheral artery disease
PaO ₂ or PC	D_2 partial pressure of oxygen
PCI	percutaneous coronary inter-
DOM	vention
PCNL	percutaneous nephrolithotomy
PCT	procalcitonin
PCWP	pulmonary capillary wedge
	pressure
PD	peritoneal dialysis
PE	peritoneal dialysys
PE	pulmonary embolism
PEEP	positive end-expiratory pressure
PEFR	peak expiratory flow rate
PEG	polyethylene glycol
PIOPED	prospective investigation of pul-
	monary embolism diagnosis
PMI	point of maximal impulse
po or p.o.	per os
PO	partial pressure of oxygen
PO	pulmonary oedema
PPI	proton pump inhibitor
prn	when necessary
PSE	portosystemic encephalopathy
PSS	partosystemic shunt
PSVT	paroxysmal supraventricular
	tachycardia
PTBR	peripheral-type benzodiazepine
	receptor
РТСА	percutaneous transluminal coro-
	nary angioplasty
PVT	polymorphic ventricular tachy-
	cardia
RAAS	renin-angiotensin-aldosterone
	system

RAD	right axis deviation
RAST	radioallergosorbent assay
RBBB	right bundle branch block
RMSF	Rocky mountain spotted fever
ROSC	return of spontaneous circula-
	tion
RRT	renal replacement therapy
RV	right ventricle
RVOT	right ventricular outflow tract
SA	sinoatrial
SBP	systolic blood pressure
SCD	sequential compression device
SCD SIRS	sudden cardiac death systemic inflammatory response
SIKS	
SLE	syndrome systemic lupus erythematosus
SOB	shortness of breath
SOB SRS-A	slow-release substance-anaphy-
510-11	laxis
STEMI	ST-segment elevation myocar-
012011	dial infarction
SVT	supraventricular tachycardia
ТВ	tuberculosis
TBI	traumatic brain injury
TIA	transient ischaemic attack
TIMI	thrombolysis in myocardial in-
	farction
TIPS	transjugular intrahepatic porto-
	systemic shunt
TNF	tumour necrosis factor
TnI	troponin I
TnT	troponin T
TOD	target organ damage
TOE	transoesophageal echo
TSH	thyroid-secreting hormone
TTP UA	thrombocytopenic purpura unstable angina
UA UA	urinalysis
UA	urine output
UFH	unfractionated heparin
V/Q scan	ventilation/perfusion scan
VC	vomiting centre
VF	ventricular fibrillation
VKA	vitamin K antagonist
VPB	ventricular premature beats
VSR	ventricular septal rupture
VT	ventricular tachycardia
VVS	vasovagal syncope
WBC	white blood cell count
WPW	Wolff-Parkinson-White syn-
	drome

Chapter 1

CARDIOPULMONARY RESUSCITATION

Definition

Cardiac arrest is a sudden cessation of effective cardiac pumping function as a result of either ventricular asystole (electrical or mechanical) or pulseless ventricular tachycardia or ventricular fibrillation. Cardiac arrest manifests clinically as sudden cardiac death. **Sudden cardiac death** is unexpected natural death from a cardiac cause within one hour of the onset of symptoms in a person without a previous condition that would appear fatal.

Epidemiology

Sudden cardiac death is a major clinical problem causing 300,000 to 400,000 deaths annually. The reported incidence of sudden cardiac death is 54 to 55 per 100,000 persons, accounting for 5.6 % of annual mortality. It was reported that 63 % of all cardiac deaths were caused by sudden cardiac death. Despite the overall decrease in cardiovascular mortality, the proportion of sudden cardiac death has remained constant.

Etiology and Pathogenesis

Cardiac arrest is strongly associated with coronary artery disease. This disease is present in 50 % to 80 % of patients older than 35 years of age with sudden cardiac death, either by history or autopsy. Weaver and colleagues demonstrated an 81 % prevalence of coronary artery disease on coronary angiography in CPR survivors. Other diseases associated with sudden cardiac death include aortic stenosis, congenital heart disease, Wolff–Parkinson–White syndrome and cardiomyopathies. The incidence of ventricular tachycardia or ventricular fibrillation presenting as cardiac arrest has been declining — only 21 % to 32 % of cardiac arrests present as ventricular tachycardia or ventricular fibrillation. Asystole and pulseless electrical activity are presenting more commonly as initial rhythms.

Coronary artery disease and cardiomyopathies account for 90 % to 95 % of cases of sudden cardiac death in the United States. Both diseases

Chapter 1

act as structural substrates underlying the functional abnormality of arrhythmias. Therefore, underlying risk factors for coronary artery disease act as risk factors for sudden cardiac death over time. Transient risk factors, such as myocardial ischaemia, hypoxaemia, acidosis, electrolyte imbalances and toxic effects of certain drugs act on the underlying structural abnormalities to produce ventricular tachycardia or ventricular fibrillation.

As time passes, ventricular tachycardia or ventricular fibrillation deteriorates to asystole, which has a dismal prognosis. Patients with ventricular tachycardia or ventricular fibrillation are more responsive to resuscitative efforts than are those with asystole or pulseless electrical activity. Weisfeldt and Becker proposed a 3-phase time-sensitive model of cardiac arrest. These phases are the electrical, circulatory and metabolic phases, lasting 0 to 4 minutes, 4 to 10 minutes and more than 10 minutes, respectively, from the time of cardiac arrest. Each requires specific treatments. During the electrical phase of cardiac arrest, defibrillation is the most effective treatment, whereas in the circulatory phase, good-quality CPR gains increasing importance, along with defibrillation. In the third and finalmetabolic phase, global ischaemic injury occurs and therapeutic strategies that focus on metabolic derangements are critical.

Survival rates among patients in out-of-hospital cardiac arrest vary from 5 % to 18 %. Survival also depends on the presenting rhythm, with a low 0.9 % survival rate for pulseless electrical activity and asystole and a 9.5 % to 41 % reported survival rate for ventricular tachycardia or ventricular fibrillation.

Clinical Presentation

Early recognition is a key step in the early treatment of cardiac arrest. It is important to determine the most accurate method of diagnosing cardiac arrest.

Rescuers should start CPR if the victim is unconscious (unresponsive), not moving and not breathing. Even if the victim takes occasional gasps, rescuers should suspect that cardiac arrest has occurred and should start CPR. The rescuer should not spend more than 10 seconds checking for a pulse, and if a pulse is not definitely felt within 10 seconds, should begin CPR and use AEDs when available.

Early CPR and Defibrillation

Cardiopulmonary resuscitation provides artificial circulation and ventilation until advanced life support can be provided and spontaneous circulation and ventilation can be restored. The 2010 AHA Guidelines for CPR and ECC recommend a change in the basic life support a sequence of steps from A-B-C (Airway, Breathing, Chest compressions) to C-A-B (Chest compressions, Airway, Breathing) for adults, children and infants (excluding the newly born).

CPR is divided into three support stages:

Basic life support:

- C Circulation support.
- A Establishment of an Airway.
- **B B**reathing support.

Advanced life support:

- **D D**iagnosis and **D**rugs.
- **E E**lectrocardiography.
- **F F**ibrillation control

Prolonged life support:

- **G G**auging the patient's response.
- H Hopeful measures for the brain.
- I Intensive care.

Basic Life Support

Initial steps of resuscitation may include:

- 1. The lone rescuer should begin CPR with 30 compressions.
- 2. Opening the airway.
- 3. Giving 2 breaths.

Chest Compressions. Several components of chest compressions can alter effectiveness: hand position, position of the rescuer, position of the victim, depth and rate of compression, decompression and duty cycle. Evidence for these techniques was reviewed in an attempt to define the optimal method.

Chest Compression Technique. It is reasonable for laypeople and healthcare professionals to be taught to position the heel of their dominant hand in the centre of the chest of an adult victim, with the nondominant hand on top.

It is reasonable for lay rescuers and healthcare providers to perform chest compressions for adults at a rate of at least 100 compressions per min and to compress the sternum by at least 5 cm. Rescuers should allow complete recoil of the chest after each compression. When feasible, rescuers should frequently alternate the "compressor" duties, regardless of whether they feel fatigued, to ensure that fatigue does not interfere with the delivery of adequate chest compressions. It is reasonable to use a duty cycle (i. e. the ratio between compression and release) of 50 %.