# **PEAL**

Prevalence of **P**ulmonary complications after **E**mergency **A**bdominal **L**aparo-tomy/scopy. An international prospective multicentre observational cohort study.

Identifier			
HOSPITAL			
PATIENT IDENTIFICATION			
RESEARCHER 1			
RESEARCHER 2			

CASE REPORT FORM (CRF) Version 01.0 05-12-2022

CONFIDENCIAL

HOSPITAL		SUBJECT	
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## **PREOPERATIVE DATA**

DEMOGRAPHIC DATA					
Age (years):	Male Female	Heigth(cm):			
weight (kg):	IMC (kg/m²):	Ideal body weight (kg/m²):			
Admission date (dd/mm/yyyy):	Surgery date (dd/mm/yyyy):	Date of hospital discharge (dd/mm/yyyy):			

Inclusion criteria	YES	NO
Age equal to or older than 18 years		
Emergency laparo-tomy/scopy		
Informed consent		

Exclusion criteria	None

	Informed Consent			
Yes	Indicate date/time of getting the informed o	onsent.		
	/(dd/mm/yyyy) hour:			
No S	No Specify the reason:			
Waive	Waived by local ethics committee			
Reject	ted by patient or relatives Rejected by the	physician Absence of investigat	or	

CO-MORBIDITIES	YES	NO	CO-MORBIDITIES	YES	SI
Arterial hypertension			Dyslipemia		
Ischemic cardiopathy		OSA			
Diabetes mellitus II		COPD			
smoker	Chronic renal failure				
Ex smoker (> 3 months)	Chronic liver failure				
Alcohol consumption (more than two drinks per day)		Oncological			
Neuromuscular disease		Inmunosuppresion			

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Surgery	
Surgery time starts:h:min (24h) ends: _	h: min(24h)
Type of surgery	
Laparotomy	Laparoscopy
General Surgery	
Apendicectomy	Colecistectomy
Hemoperitoneum (gastrointestinal, gynecological, urological, vascular)	Adhesiolisis
Colorectal resection	Small bowel resection
Gastrectomy	Grastrointestinal perforation
Abdominal abscess	Hysterectomy
Gastrointenstinal obstruction	Gastrointenstinal ischaemia
Gastrointestinal dehiscence	Incarcerated hernia
Other Gastrointestinal:	Other Urological:
Other Vascular:	Other Gynecological:
Other:	9.50
3	

PREOPERATIVE DATA					
Primary diagnosis:	Primary diagnosis:				
ASA I II III IV					
SpO <sub>2</sub> (FiO <sub>2</sub> 0.21)	%	Preoperative Hb (g/dl)			
Lung infection during the last month yes No					
Clinical Frailty Scale (from 1 to 9): Charlson:					

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### **INTRAOPERATIVE DATA**

INTRAOPERATIVE DATA					
VARIABLE	то	T1	T2		
	(10 min after intubation)	(60 min after intubation)	(pre-extubation)		
PEEP (cmH <sub>2</sub> O)					
FR					
VT (ml)					
FiO <sub>2</sub> (%)					
*PaO₂ (mmHg)					
*PaO <sub>2</sub> /FiO <sub>2</sub> (mmHg)					
*PaCO₂ (mmHg)					
*рН					
Plateau pressure (cmH <sub>2</sub> O)					
Cdyn (ml/cmH₂O)					
PAM (mmHg)					
IC (ml/min/m²)					
Air-Test (0.21 FiO <sub>2</sub> during 5 min or SpO <sub>2</sub> 97%)					
SpO <sub>2</sub> (%) a FiO <sub>2</sub> 21%					

<sup>\*</sup>Only if arterial catheterization

Fluids (ml)					
		T			
Crystalloids			Red blood cells		
Colloids			Others outputs		
Estimated blood loss			Urinary output		
Other inputs					
Additional information		l		,	
Duration of surgery (min	)		Duration of MV (min)		
Surgical position.	Supine	Trend	Reverse trend		
Surgical Incision.	Periferic	Superior Abdominal	Intra-thoracic		
Use of vasoactive drugs  Yes No Drug/dose: (separated with commas):					
Dose of vasoactive drugs: (in the same order as previously listed, separated with commas):					
Pneumoperitoneum pressure (mmHg):					

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Calculate ARISCAT (Automatic calculated field in	online C	CRF):			
Anesthetic management			,		
Hypnotic manteinance Intravenous	Hal	ogenated	Antibiotic prophylaxis.	Yes	No
Neuromuscular blockade	Yes	No	Epidural Analgesia	Yes	No
Quantitative Neuromuscular Monitorization	Yes	No	Temperature monitoring	Yes	No
NMB reversion	Yes	No	Depth of anesthesia monitoring	Yes	No
Recruitment maneuvers	Yes	No	Individualized PEPP	Yes	No
TOFr >0,9 before extubation	Yes	No			

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### **POSTOPERATIVE DATA**

NOTE: Before the Air-Test a VAS < 4 must be guaranteed

Air Test after 15-30 min at PACU	SpO2:	%

	POSTOPERATIVE DATA								
Acute	postope	rative respiratory failur	e at PACU						
Yes	No	If yes, indicate	treatment: Inc	crease in FIO <sub>2</sub>	HFNT	СРАР	NIMV	IMV	
Was t	he patier	nt extubated in the OR*							
Yes	No	If not, indicate:	Respiratory	Hemodyn	amic	Neurolog	ical	Others:	
iCU د	due to M\	/ requirement?	Yes No	If yes, in	dicate: Tin	ne until extı	ubation (m	nin):	

Day 1					
Does the patient have any pulmonary complication	on until the first day	after surgery?	Yes No		
Mild acute respiratory failure	Severe acute re	espiratory failure	Weaning failure		
ARDS mild. ARDS moderate. ARDS severe	Respiratory inf	ection	Pleural effusion		
Atelectasis	Pneumothorax		Bronchoespasm		
Aspiration pneumonitis	Pulmonary eden	na	Pulmonary embolism		
Imaging technique:					
Chest X-ray	LUS		СТ		
Does the patient have any systemic complication	?		Yes No		
Yes No					
Surgical site infection		Urinary infection			
Septic shock. Sepsis		AKII AKII	I AKI III		
Cardiac failure		Myocardial ischemia			
De novo Arrythmia		Delirium			
Multiorgan failure		Paralytic ileus			
Postoperative hemorrhage		Anastomotic leakag	е		

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ICU admission?	Yes No
Yes No	
Per protocol	Respiratory
Septic shock. Sepsis	Multiorgan failure
renal failure	Hemodynamic failure
Others:	ICU length of stay (hours):
Re-intervention	Yes No
□ Bleeding	☐ Anastomotic leakage
□ Infection	□ Others:

Day 3					
Does the patient have any pulmonary complication	on until the first day	after surgery?	Yes No		
Mild acute respiratory failure	Severe acute re	espiratory failure	Weaning failure		
ARDS mild. ARDS moderate. ARDS severe	Respiratory infe	ection	Pleural effusion		
Atelectasis	Pneumothorax		Bronchoespasm		
Aspiration pneumonitis	Pulmonary eden	na	Pulmonary embolism		
Imaging technique:					
Chest X-ray	LUS		СТ		
Does the patient have any systemic complication	?	Yes No			
Surgical site infection		Urinary infection			
Septic shock. Sepsis		AKI I AKI II AKI III			
Cardiac failure		Myocardial ischemia			
De novo Arrythmia		Delirium			
Multiorgan failure		Paralytic ileus			
Postoperative hemorrhage		Anastomotic leakage			
ICU admission?			Yes No		
Per protocol		Respiratory			

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Septic shock. Sepsis	Multiorgan failure
Fallo renal	Hemodynamic failure
Others:	ICU length of stay (hours):
Re-intervention	Yes No
☐ Bleeding	☐ Anastomotic leakage
☐ Infection	☐ Others:

	Da	v 5	
Does the patient have any pulmonary complication			Yes No
Mild acute respiratory failure	Severe acute respiratory failure		Weaning failure
ARDS mild. ARDS moderate. ARDS severe	Respiratory inf	ection	Pleural effusion
Atelectasis	Pneumothorax		Bronchoespasm
Aspiration pneumonitis	Pulmonary edema		Pulmonary embolism
Imaging technique:			
Chest X-ray	LUS		СТ
Does the patient have any systemic complication	?		Yes No
Surgical site infection		Urinary infection	
Septic shock. Sepsis		AKI I AKI II	AKI III
Cardiac failure		Myocardial ischemia	
De novo Arrythmia		Delirium	
Multiorgan failure		Paralytic ileus	
Postoperative hemorrhage		Anastomotic leakage	
ICU admission?			Yes No
Per protocol		Respiratory	
Septic shock. Sepsis		Multiorgan failure	
Fallo renal		Hemodynamic failure	
Others:		ICU length of stay (hour	s):

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Re-intervention	Yes No
□ Bleeding	☐ Anastomotic leakage
□ Infection	☐ Others:

Day 7			
Does the patient have any pulmonary complication until the first day after surgery? Yes No			
Mild acute respiratory failure	Severe acute re	espiratory failure	Weaning failure
ARDS mild. ARDS moderate. ARDS severe	Respiratory info	ection	Pleural effusion
Atelectasis	Pneumothorax		Bronchoespasm
Aspiration pneumonitis	Pulmonary eden	na	Pulmonary embolism
Imaging technique:			
Chest X-ray	LUS		СТ
Does the patient have any systemic complication	?		Yes No
Surgical site infection		Urinary infection	
Septic shock. Sepsis		AKI I AKI II	AKI III
Cardiac failure		Myocardial ischemia	
De novo Arrythmia		Delirium	
Multiorgan failure		Paralytic ileus	
Postoperative hemorrhage		Anastomotic leakage	
ICU admission?			Yes No
Per protocol		Respiratory	
Septic shock. Sepsis		Multiorgan failure	
Fallo renal		Hemodynamic failure	
Others:		ICU length of stay (hours	s):
Re-intervention		I	Yes No
☐ Bleeding		☐ Anastomotic leakage	
□ Infection		☐ Others:	

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Signed (Local investigator):		
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Name and family name:	Data:	

### NOTE:

At the end of the study, a copy of the CRF will be collected on paper completed and signed by the Investigator.