



Citation: Rezoagli E, Gatti S, Villa S, Villa G, Muttini S, Rossi F, et al. (2018) ABO blood types and major outcomes in patients with acute hypoxaemic respiratory failure: A multicenter retrospective cohort study. PLoS ONE 13(10): e0206403. https://doi.org/10.1371/journal.pone.0206403

Editor: Juan Carlos Lopez-Delgado, Hospital Universitari Bellvitge, SPAIN

Received: March 23, 2018

Accepted: October 14, 2018

Published: October 25, 2018

Copyright: © 2018 Rezoagli et al. This is an open access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Data Availability Statement: The owner of the data is the study sponsor, i.e. the hospital (San Gerardo Hospital, Azienda Socio Sanitaria Territoriale – Monza), data was collected, analyzed and submitted by the authors according to national and local rules. Data from this study will be available upon request to the Azienda Socio Sanitaria Territoriale – Monza, Ufficio Ricerca, Roberta Mazzoli (email: ufficioricerca@hsgerardo. org; r.mazzoli@hsgerardo.org), material owner of the data of the present study. This individual represents the hospital for reseach-related matter RESEARCH ARTICLE

ABO blood types and major outcomes in patients with acute hypoxaemic respiratory failure: A multicenter retrospective cohort study

Emanuele Rezoagli^{1,2,3}, Stefano Gatti¹, Silvia Villa¹, Giulia Villa¹, Stefano Muttini⁴, Fabio Rossi⁵, Loredana Faraldi⁶, Roberto Fumagalli^{1,6}, Giacomo Grasselli^{7,8}, Giuseppe Foti^{1,9}, Giacomo Bellani^{1,9,*}

 School of Medicine and Surgery, University of Milan-Bicocca, Monza, Italy, 2 Lung Biology Group, Regenerative Medicine Institute (REMEDI) at CÚRAM Centre for Research in Medical Devices, Biomedical Sciences Building, National University of Ireland Galway, Galway, Ireland, 3 Department of Anaesthesia and Intensive Care Medicine, Galway University Hospitals, SAOLTA University Health Group, Galway, Ireland, 4 Department of Emergency Medicine and Intensive Care, "Ospedale Civile" Vimercate, Vimercate, Monza Brianza, Italy, 5 Immunotransfusional Unit, San Gerardo Hospital, Monza, Italy, 6 Department of Anesthesia and Critical Care, ASST Grande Ospedale Metropolitano Niguarda, Milan, Italy, 7 Dipartimento di Fisiopatologia Medico-Chirurgica e dei Trapianti, Università degli Studi di Milano, Milan, Italy, 8 Dipartimento di Anestesia, Rianimazione ed Emergenza Urgenza, Fondazione IRCCS Ca' Granda—Ospedale Maggiore Policlinico, Milan, Italy, 9 Department of Emergency and Intensive Care, San Gerardo Hospital, Monza, Italy

* giacomo.bellani1@unimib.it

Abstract

Introduction

ABO blood type A was reported to correlate with an increased risk of acute respiratory distress syndrome (ARDS) in white patients with severe sepsis and major trauma compared with patients with other blood types. Information regarding ABO phenotypes and major outcomes in patients with ARDS is unavailable. The primary aim was to determine the relationship between ABO blood type A and intensive care unit (ICU) mortality in patients with acute hypoxemic respiratory failure (AHRF). The secondary aim was to describe the association between ABO blood type A and ICU length of stay (LOS) in this study population.

Methods

In a multicenter, retrospective cohort study, we collected the clinical records of patients admitted from January 2012 to December 2014 in five ICUs of Northern Italy. We included adult white patients admitted to the ICU who were diagnosed with AHRF requiring mechanical ventilation.

Results

The electronic records of 1732 patients with AHRF were reviewed. The proportion of patients with ABO blood type A versus other blood types was 39.9% versus 60.1%. ICU mortality (25%) and ICU LOS (median [interquartile range], 5 [2–12] days) were not different



and, along with other relevant bodies, provided the authors the authorization to extract the data from the records. The authors did not have any special access privileges that others would not have.

Funding: The authors received no specific funding for this work.

Competing interests: The authors have declared that no competing interests exist.

when stratified by ABO blood type (ICU mortality, overall p value = 0.905; ICU LOS, overall p value = 0.609). SAPSII was a positive predictor of ICU mortality (odds ration [OR], 32.80; 95% confidence interval [CI], 18.80–57.24; p < 0.001) and ICU LOS (β coefficient, 0.55; 95% CI, 0.35–0.75; p < 0.001) at multivariate analyses, whereas ABO blood type did not predict ICU outcome when forced into the model.

Conclusion

ABO blood type did not correlate with ICU mortality and ICU LOS in adult patients with AHRF who were mechanically ventilated.

Introduction

Acute hypoxemic respiratory failure (AHRF) is a frequent cause of intensive care unit (ICU) admission, requiring positive pressure ventilation and significantly contributing to morbidity and mortality. Its most severe form, acute respiratory distress syndrome (ARDS), is an under-recognized and severe cause of respiratory failure, with a high ICU mortality rate of up to 35% [1]. ARDS is characterized by a local inflammatory response with endothelial activation and microvascular injury contributing to the disruption of the alveolar–capillary barrier. The consequent increased permeability leads to a protein-rich edema fluid into the alveoli, causing hypoxemia [2].

The most common risk factors for ARDS are pneumonia, extrapulmonary sepsis, and aspiration [1]. However, "patient-related" risk factors, such as older age, non-Caucasian race, and specific genetic subsets, are also associated with an increased risk for developing ARDS [3].

Biomarkers of endothelial and alveolar epithelial injury, such as von Willebrand factor (vWF) and soluble intercellular adhesion molecule-1 (ICAM-1), have been associated with pathogenesis [4,5] and outcome [5–7] of patients with acute lung injury.

Clinical studies have suggested that the ABO blood type may influence the risk of both arterial [8–10] and venous vascular thrombosis [11,12], being correlated with different levels of vWF [13,14] and ICAMs [15,16], molecules highly associated with the clinical history of ARDS [4–7].

Reilly et al. recently observed that ABO blood type A is associated with an increased risk for ARDS in white patients with severe sepsis compared with those with other blood types. Similar findings were reported in critically ill white patients hospitalized after major trauma [17].

Based on the findings by Reilly et al., we proposed to understand if ABO blood type could be associated with ICU outcomes in a patient population fulfilling the criteria of acute noncardiogenic hypoxemic respiratory failure. We performed a retrospective study to investigate whether ABO blood type A was associated with ICU mortality and ICU length of stay (LOS) in a large cohort of critically ill patients with AHRF.

Materials and methods

This multicenter, observational, retrospective cohort study was conducted according to the Declaration of Helsinki and the Italian guidelines of good clinical practice and Data Protection Code.

Personal information of patients was recorded anonymously using an alpha-numeric code and was filed electronically.

We collected the clinical records of patients admitted from January 2012 to December 2014 in the general ICUs of five different Italian hospitals (San Gerardo Hospital, Monza; Alessandro Manzoni Hospital, Lecco; Vimercate Hospital, Vimercate; Niguarda Ca' Granda Hospital, Milano; and IRCCS Ospedale Maggiore Policlinico, Milano). The coordinating center was San Gerardo Hospital in Monza. The Ethical Committee of Monza-Brianza Province, ASST Monza-Ospedale San Gerardo-Anestesia e Rianimazione, Monza, Italy (Chairperson Prof. V. Locatelli), approved this study (Ethical Committee N° AB0-ARF (2319)). Informed consent was waived based on the retrospective observational nature of the investigation.

The inclusion criteria were adult white patients and presence of AHRF at admission, defined as a $PaO_2/FiO_2 < 300$ and need for invasive mechanical ventilation with a PEEP of ≥ 5 [1]. The exclusion criteria were patients admitted to ICU following elective surgery and cardiogenic shock and/or acute cardiogenic pulmonary edema as the primary cause of respiratory failure.

We collected the following clinical data from patient electronic medical records: sex, age, ABO blood type (A, B, AB, and O), Rh blood type (positive or negative), weight, height, simplified acute physiology score II (SAPSII) at admission [18], ICU LOS in patients who survived at discharge, and ICU mortality.

Statistical analysis

Normality of distribution was assessed using the Shapiro–Wilk test and by visual assessment of variable distribution using histogram. Continuous variables were described as mean \pm SD or median (interquartile range, IQR), as appropriate. Binary variables were described as count and proportion (%).

Differences in continuous baseline characteristics between blood type A and other blood types were tested with unpaired Student *t*-test, Mann–Whitney U test, or Chi-squared test.

ICU mortality and ICU LOS were compared between different ABO blood types with the Pearson Chi-squared test and ordinary one-way analysis of variance, respectively. Multiple comparisons *post-hoc* analyses were performed using pairwise Chi-squared tests (ICU mortality) and Bonferroni correction (ICU LOS).

Univariate logistic regression was used to examine the association between survival and blood type A versus other blood types in all patients (primary outcome).

Univariate linear regression was used to examine the association between ICU LOS and blood type A versus other blood types in patients who survived at discharge (secondary outcome).

Variables that were not normally distributed were inserted into the models after log transformation (log).

The association between clinical data, participating centers, and patient outcomes was further tested at the univariate analysis.

Each single variable that correlated with survival or with hospital LOS with p < 0.10 was eligible for inclusion in a multivariate logistic or linear regression, respectively. Backward elimination was performed to finalize the independent predictors of the multivariate models.

The variable defining the presence of blood type A versus other blood types was forced into the multivariate model to explore the study hypothesis. Multicollinearity among predictors of outcomes in the multivariate models was tested using the study of variance inflation factors.

Statistical significance was achieved when p was <0.05 (two-tailed). Statistical analyses were performed using STATA-14/MP (StataCorp LP, College Station, TX, USA) and Microsoft Excel for Mac 2017, Version 15.32.

The study was conducted and reported based on the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guidelines for observational studies [19].

Results

Baseline characteristics

A cohort of 1732 patients was included in the study. The ABO blood type distribution was blood type A: 39.9% and other blood types: 60.1%. Baseline patient characteristics are summarized in Table 1 and S1 Table.

Study outcomes: ICU mortality and ICU LOS

More than a quarter (25.5%; n = 441) of patients died in the hospital. No difference in mortality was found among patients with different ABO blood types (ABO blood type, deceased/all patients; frequency: 1. type A, 181/691, 26.2%; 2. type B, 53/223, 23.8%; 3. type AB, 20/77, 26.0%; 4. type 0, 187/741, 25.2%; overall p value = 0.905; Fig 1A).

The median ICU LOS of patients who survived at discharge was 6 days (IQR, 2–12 days) and was not different among patients with different ABO blood types (type A, 6 days [2–12 days]; type B, 6 days [2–13 days]; type AB, 5 days [2–15 days]; type 0, 6 days [2–12]; overall p value = 0.609; Fig 1B).

Major outcomes stratified by the institution of patient admission are summarized in <u>S2</u> <u>Table</u> and <u>S3 Table</u>.

Predictors of ICU mortality

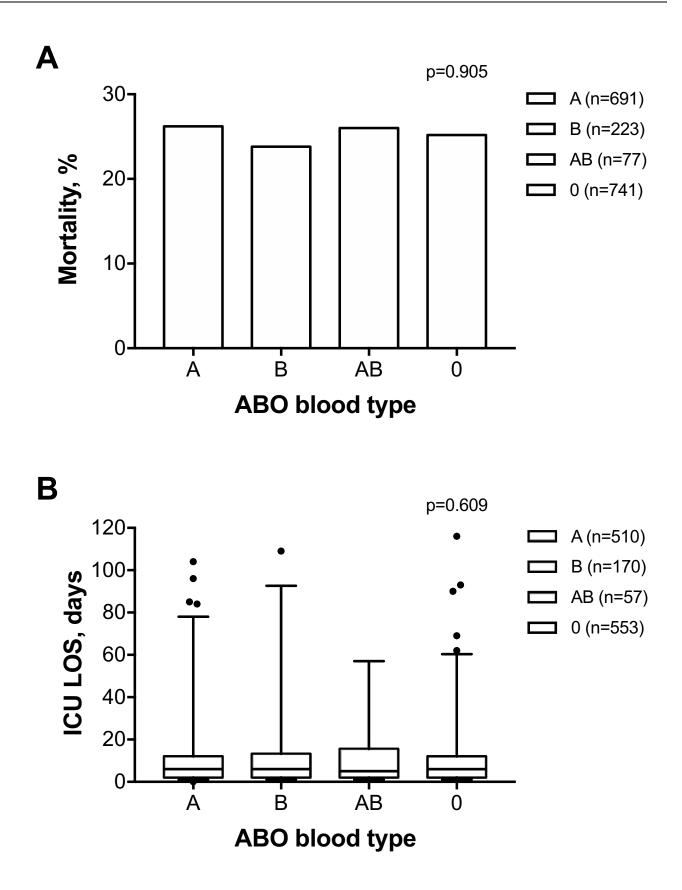
In the univariate analysis, two clinical variables at admission were associated with a higher ICU mortality: age (OR, 3.02; 95% CI, 1.94–4.70; p < 0.001) and SAPS II (OR, 32.55; 95% CI, 18.68–56.72; p < 0.001). Higher Glasgow coma score (GCS) score (OR, 0.27; 95% CI, 0.20–0.37; p < 0.001), longer ICU LOS (OR, 0.84; 95% CI, 0.76–0.92; p < 0.001), and higher weight

	ABO blood type A (n = 691)	ABO blood type non-A (n = 1041)	P-value
Sex, M/F (%)	242/368 (35/65)	449/673 (43/57)	0.888
Age, years, median (IQR)	68 (56–77)	68 (56–77)	0.643
SAPSII, median (IQR)	42 (31–56)	43 (32–59)	0.294
Weight, kg, median (IQR)	75 (65–85)	73 (65–85)	0.595
Height, cm, median (IQR)	170 (163–175)	170 (165–175)	0.623
Rh, n (%)			0.437
• Positive	600 (86.8)	917 (88.1)	
• Negative	91 (13.2)	124 (11.9)	
Admission GCS, median (IQR)	15 (11-15)	15 (10–15)	0.722
Admission Hospital, n (%)			0.832 (overall comparison)
• San Gerardo, Monza	114 (16.5)	152 (14.6)	
• A. Manzoni, Lecco	251 6.3)	390 (37.5)	
Vimercate Hospital	55 (8.0)	82 (7.9)	
• Niguarda Ca' Granda, Milano	89 (12.9)	145 (13.9)	
• IRCCS Ospedale Maggiore Policlinico, Milano	182 (26.3)	272 (26.1)	

Table 1. Patient baseline characteristics.

Abbreviations: n = number; M = male; F = female; IQR = interquartile range; SAPSII = simplified acute physiologic score II; GCS = Glasgow Coma Scale.

https://doi.org/10.1371/journal.pone.0206403.t001



PLOS ONE

Fig 1. Frequency of ICU mortality (A) and ICU length of stay in patients who survived at discharge (B) stratified by ABO blood type. Data in panel A are expressed as frequency (%) and in panel B as box and whiskers plots (1–99 percentile) with outliers (dots).

https://doi.org/10.1371/journal.pone.0206403.g001

(OR, 0.54; 95% CI, 0.31–0.96; p = 0.035) were associated with lower ICU mortality (Table 2). In the multivariate analysis, only SAPS II retained a significant association with mortality (OR, 32.80; 95% CI, 18.80–57.24; p < 0.001) (Table 3).

Predictors of ICU LOS

In univariate analysis, SAPS II was positively associated with ICU LOS (β coefficient, 0.56; 95% CI, 0.36–0.75; p < 0.001). In contrast, age (β coefficient, -0.67; 95% CI, -0.88 to -0.47; p < 0.001), GCS (β coefficient, -0.40; 95% CI, -0.65 to -0.16; p = 0.001), and male sex (β coefficient, -0.17; 95% CI, -0.31 to -0.04; p = 0.010) were negatively associated with ICU LOS. These results were confirmed by the multivariate analysis for SAPS II (β coefficient, 0.02; 95% CI, 0.01–0.03; p < 0.001) (Table 4 and Table 5). Age and GCS are components of the SAPS II score, and moderate collinearity could not be excluded and were not added to the model.

Discussion

In this retrospective observational study of a large cohort of patients with AHRF undergoing mechanical ventilation, we did not observe any significant correlation between ABO blood type A with ICU mortality compared with other ABO blood types. Furthermore, ABO blood type A did not predict ICU LOS in our statistical model.

Several studies have shown that ABO blood type is a relevant genetic factor playing a key role in regulating hemostasis [12,13,20,21]. Furthermore, two meta-analyses of the literature reported that the non-O blood group was a potential genetic risk factor for venous thrombosis [13], whereas patients with blood type O group had a higher risk of bleeding compared with those with non-O blood type [21]. Non-O blood group individuals have higher levels of vWF [22] and higher vWF adhesive activity [23]. vWF may be a useful biomarker in identifying patients with ARDS at higher risk for mortality [24–26] and prolonged ventilation [26,27]. Despite the reported association between ABO blood type A and the risk for developing ARDS, we did not observe any correlation between ABO blood group and relevant clinical outcomes in our patient population. Our results were obtained from a heterogeneous population of patients with AHRF from different causes, and we cannot exclude an association

	OR	p	95% CI
Male	1.07	0.539	0.86-1.35
Age	3.02	<0.001	1.94-4.70
SAPSII	32.55	<0.001	18.68-56.72
Weight	0.54	0.035	0.31-0.96
Height	0.99	0.990	0.96-1.04
ABO A-blood type	1.07	0.569	0.86-1.33
Rh positive	0.97	0.833	0.70-1.34
GCS	0.27	<0.001	0.20-0.37
ICU LOS	0.84	<0.001	0.76-0.92

Table 2. Univariate logistic regression of patient admission variables associated with AHRF ICU mortality.

Abbreviations: SAPSII = simplified acute physiologic score II; GCS = Glasgow Coma Scale; ICU LOS = length of stay in Intensive Care Unit.

https://doi.org/10.1371/journal.pone.0206403.t002

	OR	Р	95% CI
SAPSII	32.80	<0.001	18.80-57.24
ABO A-blood type	1.14	0.454	0.80-1.63

Table 3. Multivariate logistic regression of patient admission variables associated with AHRF ICU mortality.

Abbreviations: SAPSII = simplified acute physiologic score II.

https://doi.org/10.1371/journal.pone.0206403.t003

between ABO blood type and clinical outcomes in specific subgroups of patients with ARDS, as reported by Reilly et al. for patients with severe sepsis and major trauma.

We observed a significant correlation among SAPS II, age, and GCS with major outcomes in our population, at univariate analysis. These correlations were confirmed for SAPS II in the multivariate models. SAPS II is a robust and validated score, which provides an estimate of the risk of death in a generalized population of medical and/or surgical ICU patients [18]. Of note, the mortality in our population (25.5%) matched the predicted mortality estimated by Le Gall et al., considering a median SAPS II score of 43 in our patients. Older age is accounted among the non-modifiable factors associated with hospital mortality from ARDS, as reported by Laffey et al., in a secondary analysis of the LUNG SAFE study [28]. Furthermore, GCS score is a potentially useful predictor of mortality in general intensive care patients [29].

Interestingly, at the univariate analysis, weight tended to be negatively correlated with mortality, but this finding was not confirmed at the multivariate analysis. Despite the lack of a clear pathophysiologic mechanism, obesity has been associated with lower mortality in patients with ARDS [30], the so-called "obesity paradox" [31]. However, obese patients are prone to develop collapse of the lung-dependent zones by the mechanical weight of the abdominal content, which displaces the diaphragm toward the lung bases [32,33]. Although this mechanism leads to atelectasis-correlated hypoxemia [32] because of a right shift of the pressure--volume relationship of the respiratory system [33], it can be easily reversed by a recruitment maneuver followed by PEEP [32].

This study has several limitations. First, although we analyzed a large cohort of patients with AHRF, we acknowledge the disadvantages of the retrospective nature of the study. Second, we do not have information on the actual presence of ARDS in these patients and of its severity [1], etiology [34], and phenotype [35]. Hence, we cannot exclude an association between ABO blood type and patient outcome in different subcategories of ARDS. Third, we included only patients with AHRF requiring mechanical ventilation in the study; thus, we cannot disclose a possible role of ABO blood type in patients with mild hypoxemia managed only

	β	р	95% CI
Age	-0.67	<0.001	-0.880.47
Male	-0.17	0.010	-0.310.04
ABO A-blood type	-0.02	0.784	-0.15-0.11
Rh positive	-0.03	0.775	-0.22-0.17
SAPSII	0.56	<0.001	0.36-0.75
Weight	0.23	0.201	-0.12-0.59
Height	-0.01	0.377	-0.03-0.01
GCS	-0.40	0.001	-0.650.16

Table 4. Univariate linear regression of admission variables of AHRF patients associated with ICU length of stay in patients who survived at discharge (n = 1291).

Abbreviations: SAPSII = simplified acute physiologic score II; GCS = Glasgow Coma Scale.

https://doi.org/10.1371/journal.pone.0206403.t004

Table 5. Multivariate linear regressi		les of AHRF patients assoc	clated with ICU length of
stay in patients who survived at disc	harge (n = 1291).		

β	P	95% CI
0.55	<0.001	0.35-0.75
-0.03	0.677	-0.20-0.13

Abbreviations: SAPSII = simplified acute physiologic score II.

https://doi.org/10.1371/journal.pone.0206403.t005

with non-invasive ventilation. In contrast, the strength of this study is that we collected complete data on survival and ABO blood group in 1732 patients with AHRF, with 40% ABO blood type A. Considering a level of significance (α) of 0.05 (two-tailed), we had a power (1- β) of 0.80 to estimate a difference in intra-hospital mortality as low as 7% between patients with blood type A and those with other blood types.

Conclusions

11 - 14 14

In a large multicenter, retrospective cohort study of adult patients on mechanical ventilation for AHRF, ABO blood type did not correlate with ICU mortality and ICU LOS. Future large prospective studies are required to confirm our findings, also considering specific subcategories of patients with ARDS.

Supporting information

S1 Table. Patient baseline characteristics. (DOC)

S2 Table. Frequency of mortality in patients who survived at discharge stratified by hospital admission. Pearson's chi-squared was used to test the overall difference in mortality among different institutions. (DOC)

S3 Table. Frequency ICU length of stay in patients who survived at discharge stratified by hospital admission. Ordinary one-way ANOVA was used to test the overall difference in ICU length of stay among different institutions. Post-hoc comparisons were performed using Bonferroni's correction. *p<0.05 versus Monza. Abbreviations: IQR = interquartiles. (DOC)

Author Contributions

Conceptualization: Giacomo Bellani.

Data curation: Emanuele Rezoagli, Silvia Villa, Giacomo Bellani.

Formal analysis: Emanuele Rezoagli.

Investigation: Emanuele Rezoagli, Stefano Gatti, Giulia Villa, Stefano Muttini, Loredana Faraldi, Giacomo Bellani.

Methodology: Emanuele Rezoagli, Giacomo Bellani.

Project administration: Giacomo Bellani.

Supervision: Roberto Fumagalli, Giacomo Grasselli, Giuseppe Foti, Giacomo Bellani.

Writing – original draft: Emanuele Rezoagli.

Writing – review & editing: Stefano Muttini, Fabio Rossi, Giacomo Grasselli, Giacomo Bellani.

References

- Bellani G, Laffey JG, Pham T, Fan E, Brochard L, Esteban A, et al.; LUNG SAFE Investigators; ESICM Trials Group. Epidemiology, Patterns of Care, and Mortality for Patients With Acute Respiratory Distress Syndrome in Intensive Care Units in 50 Countries. *JAMA* 2016; 315:788–800.
- Thompson BT, Chambers RC, Liu KD. Acute Respiratory Distress Syndrome. N Engl J Med 2017; 377:562–572. https://doi.org/10.1056/NEJMra1608077 PMID: 28792873
- 3. Rezoagli E, Fumagalli R, Bellani G. Definition and epidemiology of acute respiratory distress syndrome. *Ann Transl Med* 2017; 5:282. https://doi.org/10.21037/atm.2017.06.62 PMID: 28828357
- 4. Fremont RD, Koyama T, Calfee CS, et al. Acute lung injury in patients with traumatic injuries: utility of a panel of biomarkers for diagnosis and pathogenesis. *J Trauma* 2010; 68:1121–7. <u>https://doi.org/10.1097/TA.0b013e3181c40728</u> PMID: 20038857
- Calfee CS, Eisner MD, Parsons PE, Thompson BT, Conner ER Jr, Matthay MA, et al.; NHLBI Acute Respiratory Distress Syndrome Clinical Trials Network. Soluble intercellular adhesion molecule-1 and clinical outcomes in patients with acute lung injury. *Intensive Care Med* 2009; 35:248–57. https://doi. org/10.1007/s00134-008-1235-0 PMID: 18670758
- Ware LB, Conner ER, Matthay MA. von Willebrand factor antigen is an independent marker of poor outcome in patients with early acute lung injury. Crit Care Med 2001; 29:2325–31. PMID: 11801836
- Ware LB, Eisner MD, Thompson BT, Parsons PE, Matthay MA. Significance of von Willebrand factor in septic and non septic patients with acute lung injury. *Am J Respir Crit Care Med* 2004; 170:766–72. https://doi.org/10.1164/rccm.200310-1434OC PMID: 15201135
- Meade TW, Cooper JA, Stirling Y, Howarth DJ, Ruddock V, Miller GJ. Factor VIII, ABO blood group and the incidence of ischaemic heart disease. Br J Haematol 1994; 88:601–607. PMID: 7819072
- Whincup PH, Cook DG, Phillips AN, Shaper AG. ABO blood group and ischaemic heart disease in British men. BMJ 1990; 300:1679–1682. PMID: 2390546
- He M, Wolpin B, Rexrode K, Manson JE, Rimm E, Hu FB, et al. ABO blood group and risk of coronary heart disease in two prospective cohort studies. *Arterioscler Thromb Vasc Biol* 2012; 32:2314–20. https://doi.org/10.1161/ATVBAHA.112.248757 PMID: 22895671
- Schleef M, Strobel E, Dick A, Frank J, Schramm W, Spannagl M. Relationship between ABO and Secretor genotype with plasma levels of factor VIII and von Willebrand factor in thrombosis patients and control individuals. *Br J Haematol* 2005; 128:100–7. <u>https://doi.org/10.1111/j.1365-2141.2004.05249.x</u> PMID: 15606555
- Dentali F, Sironi AP, Ageno W, Turato S, Bonfanti C, Frattini F, et al. Non-O blood type is the commonest genetic risk factor for VTE: results from a meta-analysis of the literature. *Semin Thromb Hemost* 2012; 38:535–48. https://doi.org/10.1055/s-0032-1315758 PMID: 22740183
- Franchini M, Capra F, Targher G, Montagnana M, Lippi G. Relationship between ABO blood group and von Willebrand factor levels: from biology to clinical implications. *Thromb J* 2007; 5:14. <u>https://doi.org/</u> 10.1186/1477-9560-5-14 PMID: 17894864
- Bowen DJ. An influence of ABO blood group on the rate of proteolysis of von Willebrand factor by ADAMTS13. J Thromb Haemost 2003; 1:33–40. PMID: <u>12871537</u>
- 15. Paré G, Chasman DI, Kellogg M, Zee RY, Rifai N, Badola S, et al. Novel association of ABO histo-blood group antigen with soluble ICAM-1: results of a genome-wide association study of 6,578 women. *PLoS Genet* 2008; 4:e1000118. https://doi.org/10.1371/journal.pgen.1000118 PMID: 18604267
- Barbalic M, Dupuis J, Dehghan A, Bis JC, Hoogeveen RC, Schnabel RB, et al. Large-scale genomic studies reveal central role of ABO in sP-selectin and sICAM-1 levels. *Hum Mol Genet* 2010; 19:1863– 72. https://doi.org/10.1093/hmg/ddq061 PMID: 20167578
- Reilly JP, Meyer NJ, Shashaty MGS, Feng R, Lanken PN, Gallop R, et al. AB0 blood type A is associated with increased risk of ARDS in whites following both major trauma and severe sepsis. *Chest* 2014; 145:753–761. https://doi.org/10.1378/chest.13-1962 PMID: 24385226
- Le Gall JR, Lemeshow S, Saulnier F. A new Simplified Acute Physiology Score (SAPS II) based on a European/North American multicenter study. JAMA 1993; 270:2957–63. PMID: 8254858
- von Elm E, Altman DG, Egger M, Pocock SJ, Gøtzsche PC, Vandenbroucke JP; STROBE Initiative. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. *Ann Intern Med* 2007; 147:573–7. PMID: 17938396

- Preston AE, Barr A. The plasma concentration of factor VIII in the normal population. Br J Haematol 1964; 10:238–245. PMID: 14141622
- Dentali F, Sironi AP, Ageno W, Bonfanti C, Crestani S, Frattini F, et al. Relationship between ABO blood group and hemorrhage: a systematic literature review and meta-analysis. *Semin Thromb Hemost* 2013; 39:72–82. https://doi.org/10.1055/s-0032-1329550 PMID: 23299820
- ill JC, Endres-Brooks J, Bauer PJ, Marks WJ Jr, Montgomery RR. The effect of ABO blood group on the diagnosis of von Willebrand disease. Blood 1987; 69:1691–1695. PMID: 3495304
- **23.** Sarode R, Goldstein J, Sussman II, Nagel RL, Tsai HM. Role of A and B blood group antigens in the expression of adhesive activity of von Willebrand factor. *Br J Haematol* 2000; 109:857–864. PMID: 10929042
- Ware LB, Conner ER, Matthay MA. von Willebrand factor antigen is an independent marker of poor outcome in patients with early acute lung injury. *Crit Care Med* 2001; 29:2325–2331. PMID: <u>11801836</u>
- **25.** Ware LB, Eisner MD, Thompson BT, Parsons PE, Matthay MA. Significance of von Willebrand factor in septic and nonseptic patients with acute lung injury. *Am J Respir Crit Care Med* 2004; 170:766–72 https://doi.org/10.1164/rccm.200310-1434OC PMID: 15201135
- Calfee CS, Gallagher D, Abbott J, Thompson BT, Matthay MA; NHLBI ARDS Network. Plasma angiopoietin-2 in clinical acute lung injury: Prognostic and pathogenetic significance. *Crit Care Med* 2012; 40:1731–1737. https://doi.org/10.1097/CCM.0b013e3182451c87 PMID: 22610178
- Ware LB, Koyama T, Billheimer DD, Wu W, Bernard GR, Thompson BT, et al.; NHLBI ARDS Clinical Trials Network. Prognostic and pathogenetic value of combining clinical and biochemical indices in patients with acute lung injury. *Chest* 2010; 137:288–96. <u>https://doi.org/10.1378/chest.09-1484</u> PMID: 19858233
- Laffey JG, Bellani G, Pham T, Fan E, Madotto F, Bajwa EK, et al.; LUNG SAFE Investigators and the ESICM Trials Group. Potentially modifiable factors contributing to outcome from acute respiratory distress syndrome: the LUNG SAFE study. *Intensive Care Med* 2016; 42:1865–1876. https://doi.org/10. 1007/s00134-016-4571-5 PMID: 27757516
- Bastos PG, Sun X, Wagner DP, Wu AW, Knaus WA. Glasgow coma scale score in the evaluation of outcome in the intensive care unit: findings from the acute physiology and chronic health evaluation III study. *Crit Care Med* 1993; 21:1459–65. PMID: 8403953
- Ni YN, Luo J, Yu H, Wang YW, Hu YH, Liu D, et al. Can body mass index predict clinical outcomes for patients with acute lung injury/acute respiratory distress syndrome? A meta-analysis. *Crit Care* 2017; 21:36. https://doi.org/10.1186/s13054-017-1615-3 PMID: 28222804
- Zhi G, Xin W, Ying W, Guohong X, Shuying L. "Obesity Paradox" in Acute Respiratory Distress Syndrome: Asystematic Review and Meta-Analysis. *PLoS One* 2016; 11:e0163677. https://doi.org/10.1371/journal.pone.0163677 PMID: 27684705
- Reinius H, Jonsson L, Gustafsson S, Sundbom M, Duvernoy O, Pelosi P, et al. Prevention of atelectasis in morbidly obese patients during general anesthesia and paralysis: a computerized tomography study. *Anesthesiology* 2009; 111:979–87. https://doi.org/10.1097/ALN.0b013e3181b87edb PMID: 19809292
- 33. Behazin N, Jones SB, Cohen RI, Loring SH. Respiratory restriction and elevated pleural and esophageal pressures in morbid obesity. *J Appl Physiol (1985)* 2010; 108:212–8.
- Gattinoni L, Pelosi P, Suter PM, Pedoto A, Vercesi P, Lissoni A. Acute respiratory distress syndrome caused by pulmonary and extrapulmonary disease. Different syndromes? *Am J Respir Crit Care Med* 1998; 158:3–11. https://doi.org/10.1164/ajrccm.158.1.9708031 PMID: 9655699
- Rezoagli E, Magliocca A, Scalia Catenacci S, Bellani G, Fumagalli R. Identification of biological phenotypes in acute respiratory distress syndrome: from biomarkers to clinical outcome. *Am J Resp Crit Care Med* 2018; 1; 197(9):1209–1211. https://doi.org/10.1164/rccm.201708-1713RR PMID: 29406790