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Vilnius University BLOODSTREAM INFECTIONS IN COVID-19 POSITIVE PATIENTS HOSPITALIZED IN THE UNIVERSITY HOSPITAL, LITHUANIA

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Background Bacterial, viral and fungal co-infections in COVID-19 patients are associat therapy. This study aimed to describe bloodstream infections (BSIs) an positive adult patients.	ated with increased morbidity and mortality, requiring timely diagnosis and d to evaluate risk factors for developing BSIs in hospitalized COVID-19
Methods COVID-19 positive adults hospitalized in Vilnius University Hospital San between March 2020 and May 2021. Depersonalized data were retrieved BSI was defined as the growth of non-skin flora commensal on one or m (coagulase-negative staphylococci, <i>Micrococcus</i> spp., <i>Propionibacteriu</i> contaminants. Only one BSI episode was counted when several BC sets To explore the risk factors associated with BSIs, multivariable logistic regu	taros Klinikos, Lithuania, were included in this retrospective cohort study from electronic medical records. ore blood cultures (BCs). Bacteria belonging to the commensal skin flora <i>um</i> spp., <i>Corynebacterium</i> spp.) growing in BC sets were defined as were positive with the same microorganism for a patient. ression models were created. <i>p</i> -value < 0.05 was considered significant.
 Results A total of 2844 patients were included in this study. Baseline characteristics of hospitalized COVID-19 patients and patients according to BSI group are provided in Table 1. Total 3031 BCs were obtained from 1905 patients. Overall, 1611 patients had BC collected at less than 48 hours from admission to hospital and 768 patients – ≥ 48 hours from admission. A total of 56 community-acquired BSIs were documented in 51 (1.8%) patient, 142 hospital-acquired infections were documented in 102 (3.6%) patients. Seven patients that were admitted with community-acquired infection got BSI caused by another pathogen during hospitalization. The most frequent pathogens of community-acquired BSIs were <i>Esherichia coli</i> (26.8%), <i>Staphylococcus spp.</i> (23.2%) and <i>Klebsiella pneumoniae</i> (14.3%) (Figure 1). The most frequent microorganisms causing hospital-acquired BSIs were <i>Acinetobacter baumanii</i> (23.2%), <i>Staphylococcus spp.</i> (14.8%), <i>Klebsiella pneumoniae</i> (13.4%) and <i>Enterococcus faecium</i> (10.6%) (Figure 1). The distribution of pathogens in BCs of hospitalized COVID-19 patients are shown in Figure 1. Age (OR 1.03; 95%CI 1.01–1.06, p=0.005) and having atrial fibrillation (OR 3.08; 95%CI 1.57–6.01, p=0.001) increased odds of having community-acquired BSI (Figure 2). Risk factors for hospital-acquired BSI were identified to be obesity (OR 2.43; 95%CI 1.15–5.11, p=0.020), previous stroke (OR 10.83; 95%CI 1.02–3.7.56, p=0.001), invasive ventilation/logh-flow oxygen therapy (OR 1.99; 95%CI 1.06–3.76, p=0.033) (Figure 2). 	Acinetobacter baumanii Klebsiella pneumoniae Staphylococcus spp. Enterococcus factoria coli Proteus spp. Canietobacter spp. Enterococcus spp. Klebsiella spp. Citrobacter spp. Streptococcus spp. Fungi Citrobacter spp. Streptococcus spp. Fungi Citrobacter freundii Pseudomonas spp. Fungi Citrobacter spp. Klebsiella spp. Fungi Citrobacter freundii Pseudomonas spp. Risk factors associated with hospital-acquired bloodstream infections Figure 1. Distribution of pathogens in blood cultures of hospitalized COVID-19 patients. Risk factors associated with community-acquired BSI (100 - 376), p=0.001 (100 - 376), p=0.
Conclusions	- The analysis of the second s

Bloodstream infections' rate in hospitalized COVID-19 patients was quite low accounting 1.8% of community-acquired infections and 3.6% of hospital-acquired infections. Age and atrial fibrillation were identified as risk factors for community-acquired bloodstream infections, while obesity, previous stroke, the need of invasive and non-invasive ventilation/high flow oxygen therapy were associated with increased risk of hospital-acquired bloodstream infections in COVID-19 patients.

Characteristic	Total patients, n (%)	Patients with community–acquired bloodstream infection, n (%)	Patients with hospital-acquired bloodstream infection, n (%)
	N=2844	N=51	N=102
ge in years, median (IQR)	59 (48 – 70)	70 (58 – 78)	62 (55.75 – 70.25)
male	1301 (45.7)	25 (49.0)	39 (38.2)
underlying condition	1335 (46.9)	40 (78.4)	74 (72.5)
erial hypertension	1037 (36.5)	30 (58.8)	54 (52.9)
onary artery disease	105 (3.7)	1 (2.0)	3 (2.9)
ngestive heart failure	221 (7.8)	6 (11.8)	18 (17.6)
al fibrillation	286 (10.1)	18 (35.3)	24 (23.5)
petes	385 (13.5)	11 (21.6)	25 (24.5)
etes with no complications	111 (3.9)	1 (2.0)	2 (2.0)
etes with complications	275 (9.7)	10 (19.6)	23 (22.5)
ity	129 (4.5)	6 (11.8)	21 (20.6)
nic obstructive pulmonary disease	45 (1.6)	1 (2.0)	3 (2.9)
nic kidney disease	216 (7.6)	8 (15.7)	18 (17.6)
ous stroke	39 (1.4)	2 (3.9)	6 (5.9)
sive ventilation	227 (8.0)	10 (19.6)	61 (59.8)
-invasive ventilation / high flow oxygen therapy	273 (9.6)	9 (17.6)	42 (41.2)
ospital mortality	359 (12.62)	21 (41.2)	56 (54.9)
with of been italiantion (in days)	11 (7 – 16)	14 (10 – 27)	32 (15.75 - 53)