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Background

- Staphylococcus lugdunensis (S. lugdunensis) belongs to a group of coagulase negative Staphylococci (CoNS), which is present as a normal skin commensal in healthy individuals.
- Invasive infections caused by S. lugdunensis has been increasingly reported ¹⁻⁵.
- Whereas, there is no treatment guideline for *S. lugdunensis* bacteremia (SLB) due to a little clinical data.
- Owing to assess appropriate management for SLB, we conducted a retrospective case-control study to investigate characteristics, clinical courses and outcomes for SLB compared with those of Staphylococcus aureus (S. aureus) bacteremia and bacteremia due to Staphylococcus epidermidis (S. epidermidis), which is most common pathogen in CoNS-related bacteremia.

Materials and Methods

- **Setting, study design:** This retrospective case-control study was conducted at Kyoto University Hospital, a tertiary care 1141bed university hospital located in Japan, from January 1, 2005 to December 31, 2021.
- **Patients:** Patients who had at least one set of blood culture collection were included in this study. Of those, patients >=18 years of age with SLB with at least two sets of positive blood cultures, those who had S. aureus bacteremia (SAB) with at least one set of positive blood culture and those who had S. epidermidis bacteremia (SEB) with at least two sets of positive blood cultures were randomly selected in a 1:5:5 (SLB:SAB:SEB) ratio.
- Variables: Patients characteristics, illness severity, source of bacteremia, metastatic lesions, clinical managements (follow-up blood culture, echocardiography, early source control, days to appropriate treatment, early optimal therapy and duration of treatment) and outcomes were reviewed.
- **Statistical analysis:** When a p-value of <0.05 was revealed for comparisons among three groups, followed comparisons between SLB and SAB, and between SLB and SEB were performed.

References

- (1) Frank K.L., *et al.* Clin Microbiol Rev. 2008.
- (2) Aldman M.H., et al. Eur J Clin Microbiol Infect Dis. 2021.
- (3) Lourtet-Hascoet, J., et al. Int J Infect Dis. 2016. (4) Aegmi, X., et al. J Clin Microbiol. 2017. (5) Heilbronner, S., et al. Clin Microbiol Rev. 2021. (6) Aioda Y., et al. Jpn J Infect Dis. 2017. (7) Forsblom, E., et al PLOS ONE. 2021. (8) Lopez-Cortes, L. E., et al. Clin Infect Dis. 2013.

Poster Number: Characteristics and Outcomes in Patients with Staphylococcus lugdunensis bacteremia compared with Staphylococcus aureus and Staphylococcus epidermidis bacteremia

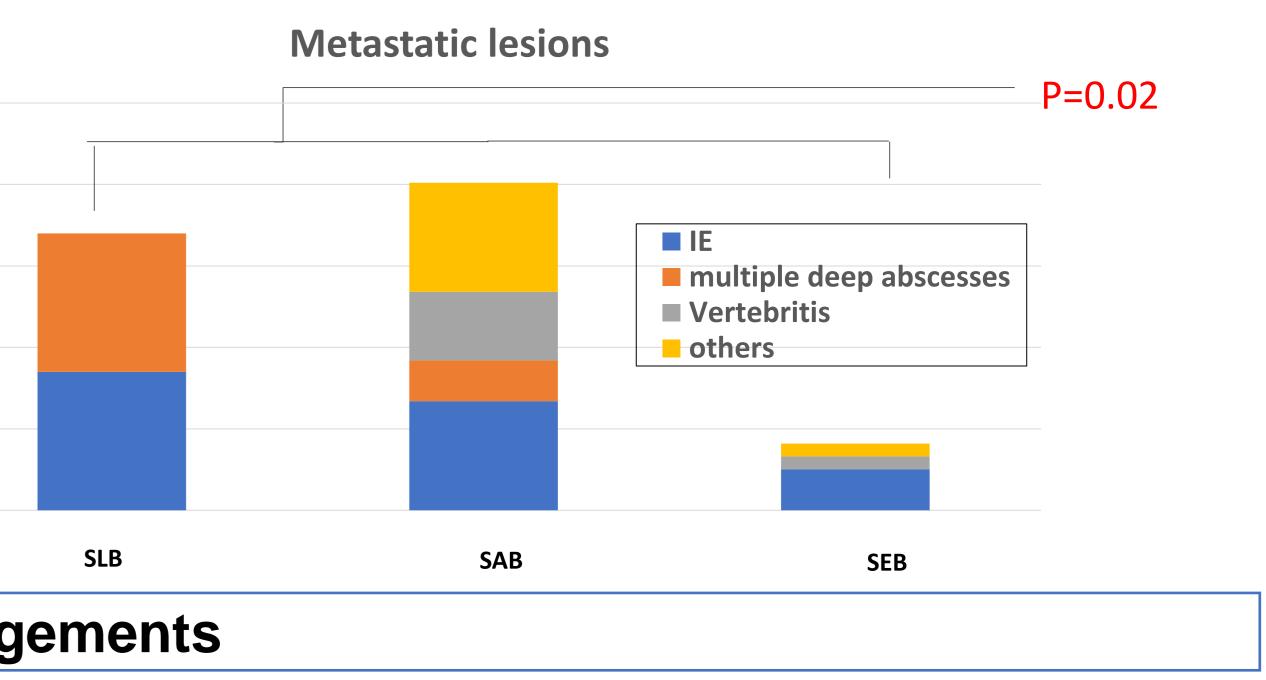
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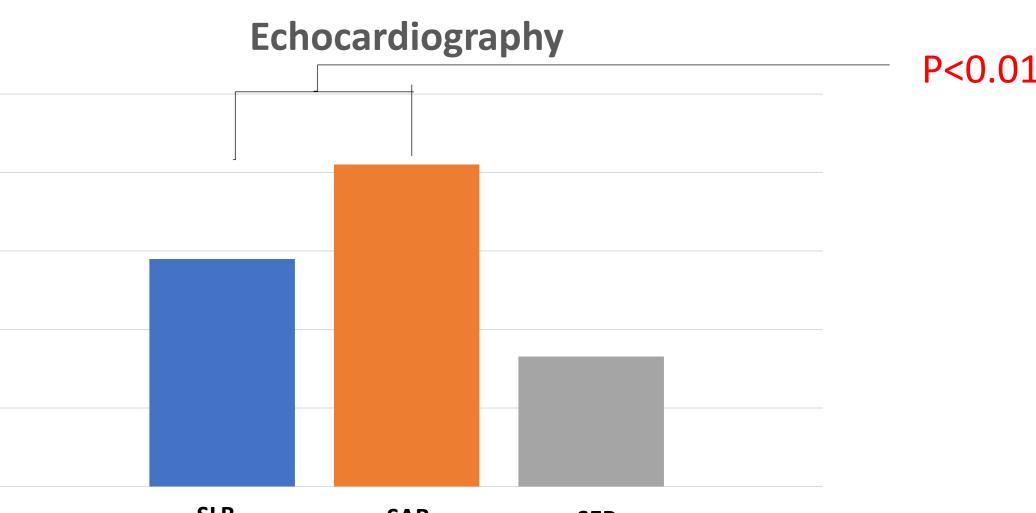
						Res	SU	lts						
Figure; Flow diagram of patients in this study							2	. Source of ba	acteren	nia, metas	static les	ions (Co	ontinue	
								(%) 25		Meta	static lesions			– P=0.02
Patients with Stap	Stapl aureu	Patients with <i>Staphylococcus</i> <i>aureus</i> positive bl culture: n = 966		Patients withStaphylococcusloodepidermidis positblood culture:n = 1090		ive	20				 IE multiple of Vertebriti others 	deep abscesses		
Exclue • Age		y 1 set		nly ed from			5 0	SLB		SAB		SEB		
• pat	only 1 set			s who =18 years		3.	Clinical man		nts	575		JED		
	blood culture positive n = 45			old				(%) 100			diography		P<0.0)1
A total of 24 patients SEB were included.		, 120 patie	ents with SA	⁺ ∖B, and	120 patie	nts with		80 60						
. Patients Characteristics								40 20						
	SLB (n=24)	SAB (n=120)	SEB (n=120)	P value among three groups	P value SLB– SAB	P value SLB– SEB	•	0 Echocardiograp Days to start th	e approp	performed lo priate therap	y from blo	ntly in SLE		
	N (%)	N (%)	N (%)					than in SEB [0				rtian of fol	llow up bl	
je (years), median R)	69 (60, 75)	68 (49, 77)	65 (51, 73)	0.28			•	There were no early source co	U		• •		•	
ale ealthcare setting	13 (54)	67 (56)	75 (63)	0.33 <0.01	0.86	<0.01	<u> </u>	The duration of		• •		Ŭ	Ŭ	jups.
	5 (21)	18 (15)	2(1.5)				1	4. Mortality						
ealthcare associated ospital acquired	6 (25) 12 (54)	25 (21)	9 (7.5)									P value	P value	P value
thicillin resistant	13 (54) 6 (25)	77 (64) 41 (34)	109 (91) 99 (83)	<0.01	0.38	<0.01			SLB	SAB	SEB	among	SLB-	SLB-
ymicrobial	6 (25) 5 (21)	8(7)	25 (21)	< 0.01	0.03	1.0			(n=24)	(n=120)	(n=120)	three	SAB	SEB
morbidities	5 (21)	0(7)	20 (21)	\U.U1	0.00	1.0						groups		
abetes	12 (50)	22 (18)	23 (19)	0.23					N (%)	N (%)	N (%)			
emodialysis	6 (25)	13 (11)	8 (7)	0.02	0.06	<0.01		7-day mortality	2 (8)	8 (7)	1 (0.5)	0.04	0.77	0.019
alignancy	13 (54)	54 (45)	· · · ·	0.02	0.41	0.44		30-day mortality	4 (17)	21 (18)	12 (10)	0.23		
ver cirrhosis	1 (4)	12 (10)	7 (6)	0.38				Hospital mortality	6 (25)	32 (27)	23 (19)	0.38		
nmunosuppressants	8 (33)	33 (28)		<0.01	0.56	0.2	,	 Seven-day model 	ortalitv wa	as similar h	etween SI	B and SA	B. hiaher	in SLB
hemotherapy	6 (25)	23 (19)	47 (39)	<0.01	0.51	0.19		than in SEB.					_,	
harlson index, nedian (IQR)	5 (3, 6)	3 (2, 6)	3 (2, 4)	0.52			(Thirty-day and SAB. 	d hospita	I mortalities	were com	parable be	etween S	LB and
. Source of bact	teremia,	metasta	tic lesior	IS					Summ	nary an	d Con	clusio	ns	

- Intravascular catheter related bloodstream infection was the most common in each group as source of bacteremia except for unknown focus.
- The proportion of metastatic lesions was comparable between SLB and SAB, higher in SLB than in SEB.

- patients with SLB.

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SLB is associated with complicated infections and high mortality. Echocardiography was performed less frequently in SLB than in SAB. Clinical outcome for SLB was worse, as same as that of SAB. Appropriate evaluation and treatment that are recommended for SAB maybe warrant for