Campylobacter bacteraemia outcomes: a systematic review with meta-analysis

Verena Zerbato*, Stefano Di Bella, Stefano Guicciardi, Roberto Baldan, Chiara Fanelli, Lisa Fusaro, Alexandru Botan, Luigi Principe, Dan Alexandru Toc, Saveria Lory Crocè, Roberto Luzzati, Alberto Enrico Maraolo

Verena Zerbato: Infectious Diseases Unit, Trieste University Hospital (ASUGI), Trieste, Italy

Stefano Di Bella, Lisa Fusaro, Saveria Lory Crocè, Roberto Luzzati: Clinical Department of Medical, Surgical and Health Sciences, Trieste University, Trieste, Italy

Stefano Guicciardi: Health Directorate, Local Health Authority of Bologna, Bologna, Italy

Roberto Baldan: Department of Medicine, University of Verona, Verona, Italy

Chiara Fanelli: Department of Medicine, Surgery and Pharmacy, University of Sassari, Sassari, Italy.

Alexandru Botan, Dan Alexandru Toc: Department of Microbiology, Iuliu Hatieganu University of Medicine and Pharmacy, 400012 Cluj-Napoca, Romania Luigi Principe: Microbiology and Virology Unit, Great Metropolitan Hospital "Bianchi-Melacrino-Morelli", Reggio Calabria, Italy Alberto Enrico Maraolo: Department of Clinical Medicine and Surgery, Section of Infectious Diseases, University of Naples 'Federico II', Italy

*Corresponding author: Dr. Verena Zerbato, Infectious Diseases Unit, Trieste University Hospital (ASUGI), Piazza dell'Ospitale n°1, 34125 Trieste, Italy - verena.zerbato@gmail.com

Background

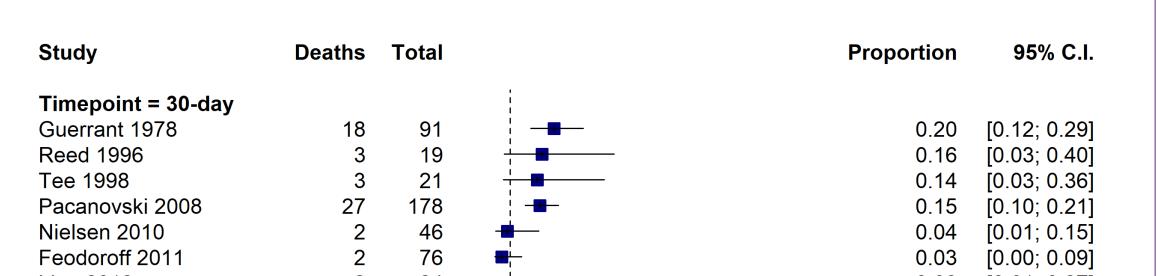
Campylobacter spp. is a common cause of acute enteric infections in humans. In immunocompromised or elderly patients, *Campylobacter* spp. can cause extraintestinal infections, including bacteremia. No specific international guidelines are present for campylobacteriosis management and treatment. The clinical significance of *Campylobacter* bacteremia is not yet fully understood [1].

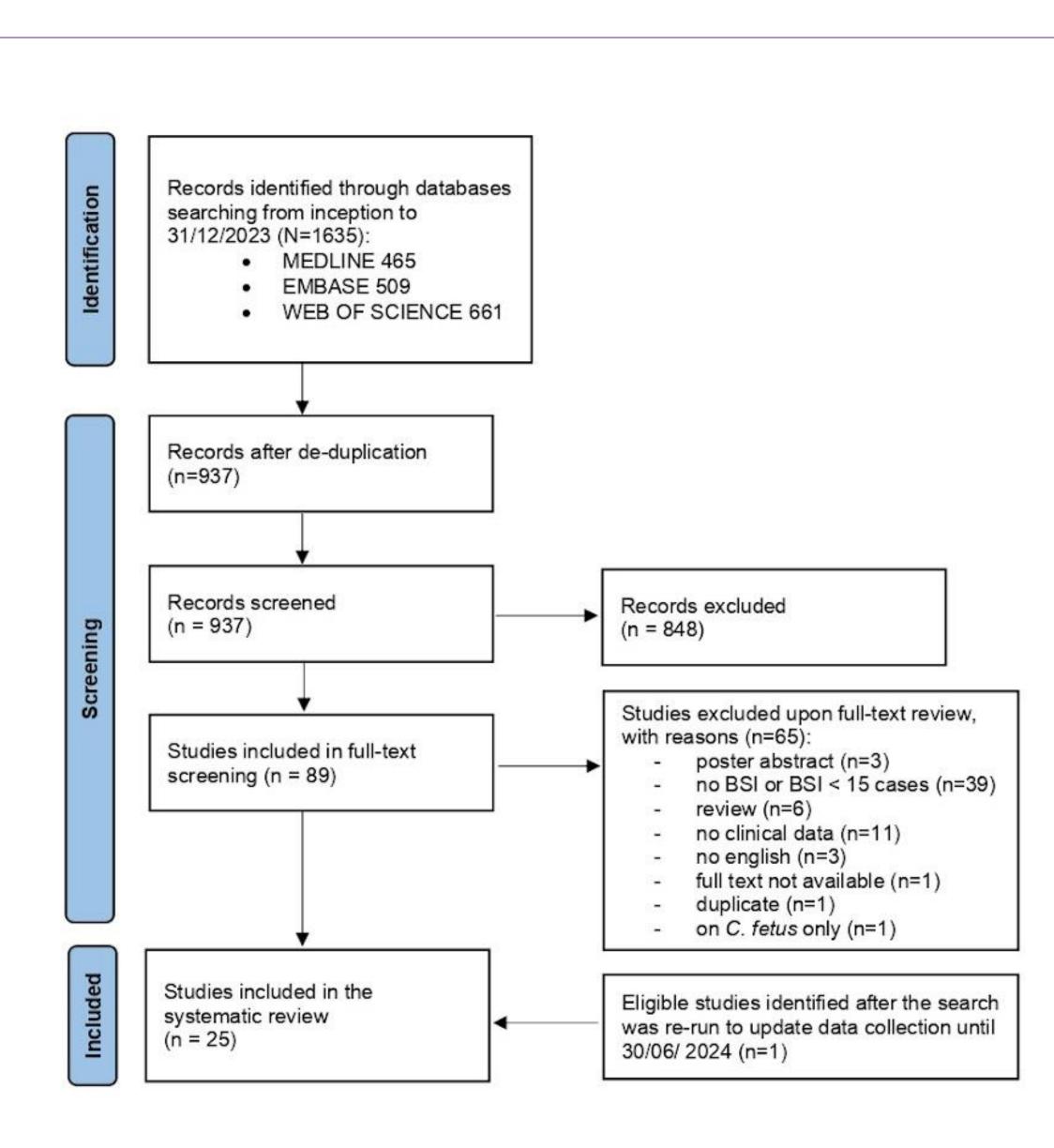
Methods

We conducted a systematic review on bloodstream infections (BSI) caused by *Campylobacter* spp. The searches covered studies in humans published from inception until December 31, 2023. The search was re-run to update the data collection until June 2024.

Results

The included studies reported a total of 2,480 patients, with a mean age ranging from 1 to 70 years. Male patients accounted for 62.45% of the cases. The pooled prevalence of *Campylobacter* species was as follows: *C. jejuni* 60% [95% CI 0.45-0.73], *C. coli* 8% [95% CI 0.04-0.13], *C. fetus* 7% [95% CI 0.03-0.15], and other species 9% [95% CI 0.04-0.16]. Mortality was reported as the primary outcome in 22 studies. The overall pooled case-fatality risk associated with *Campylobacter* spp. bacteremia was 5% [95% CI 0.03-0.15] (Figure 2).





Liao 2012	2	24		0.08	[0.01; 0.27]
Hussein 2016	3	65	- -	0.05	[0.01; 0.13]
Liu 2019	3	56	- +	0.05	[0.01; 0.15]
Tinevez 2021	69	592	—	0.12	[0.09; 0.15]
Tau 2022	9	76		0.12	[0.06; 0.21]
Graham 2024	1	34	a	0.03	[0.00; 0.15]
Sunnerhagen 2024	0	29		0.00	[0.00; 0.12]
Random effects model	142	1307		0.08	[0.05; 0.13]
Heterogeneity: $I^2 = 46\%$, $p = 0$	0.03				
Timepoint = in-hospital					
Baek 2023	14	108		0.13	[0.07; 0.21]
Otsuka 2023	0	39		0.00	[0.00; 0.09]
Random effects model	14	147		0.03	[0.00; 1.00]
Heterogeneity: $I^2 = 0\%$, $p = 1$.	00				
Timepoint = not specified					
Skirrow 1993	1	394		0.00	[0.00; 0.01]
Schonheyder 1995	1	15	- 	0.07	[0.00; 0.32]
Pigrau 1997	6	58	÷ =	0.10	[0.04; 0.21]
Fernandez-Cruz 2010	11	68	— —	0.16	[0.08; 0.27]
Ben-Shimol 2013	0	76	■ -¦	0.00	[0.00; 0.05]
O'Hara 2017	2	41	-	0.05	[0.01; 0.17]
Moffatt 2021	0	25		0.00	[0.00; 0.14]
Random effects model	21	677		0.02	[0.00; 0.13]
Heterogeneity: $I^2 = 67\%$, $p < 0$	0.01				
Random effects model	177	2131		0.05	[0.03; 0.09]
Prediction interval					[0.01; 0.36]
Heterogeneity: $I^2 = 50\%$, $p < 0$					_
Test for subgroup differences:	$\chi^2_2 = 3.2$	0, df = 2 (<i>p</i> = 00.20)0.2 0.4 0.6 0.8 1		
	_		Proportion of deaths		

Figure 2. Forest plot illustrating the pooled analysis of mortality outcome

Meta-regression analysis on mortality revealed that *C. fetus* species (b=3.22, [95% CI 0.63-5.80], p=0.018) and immunocompromised status (b=3.22, [95% CI 0.63-5.80], p=0.018) were associated with an increased risk of death in

Fig 1: Literature selection procedure

Results

A total of 25 retrospective observational studies (with at least 15 cases), published between 1978 and 2024, were included (Figure 1). The geographic distribution of the studies was as follows: 12 from Europe, 3 from Israel, 4 from Asia, 3 from Australia, 1 from the United States, and 2 from Africa.

patients with *Campylobacter* bacteremia. Regarding complications associated with *Campylobacter* spp. bacteremia, the pooled prevalence of secondary localizations was 9% [95% CI 0.04-0.18], while the pooled prevalence of endocarditis was 5% [95% CI 0.01-0.03].

Conclusions

In conclusion, the results confirm the clinical relevance of *Campylobacter* spp. bacteremia, highlighting a significant risk of mortality and complications.

References

1. Fitzgerald C. Campylobacter. Clin Lab Med. 2015 Jun;35(2):289-98. doi: 10.1016/j.cll.2015.03.001. PMID: 26004643.

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