REVIEW



Related factors of bloodstream infections associated with urinary tract infections and pathogenetic characteristics analysis



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Abstract

Objectives To explore correlations between biomarker indices and urosepsis severity, and investigate the prevalence of drug-resistant *Escherichia coli* in a patient population at the General Hospital of Ningxia Medical University in the Ningxia region of China.

Methods Patients with urinary tract infection-associated sepsis were categorized into three groups: a septic non-shock group (NSSPU), a septic shock group (USG), and a control group with non-sepsis cases of simple urinary tract infections (CG). The study analyzed various biomarkers, including the percentage of neutrophils (N%), neutrophil-to-lymphocyte ratio (NLR), and lactate (La), to assess their predictive value for urogenital sepsis severity.

Results The Kruskal–Wallis test showed significant differences in all measured biomarkers between the groups. ROC curve analysis identified N%, NLR, total protein (TP), albumin (ALB), and La as meaningful predictors of urosepsis severity. The combined detection indicators hold greater value in diagnosing uroseptic shock compared to individual test indicators. In addition, the study confirmed the prevalence of drug-resistant *E. coli* in cases of septic shock.

Conclusion The combined monitoring of N%, NLR, La, TP, and ALB proves beneficial in the clinical diagnosis of uroseptic shock. This study emphasizes the significance of monitoring *Escherichia coli* and its resistance patterns to decrease the occurrence of sepsis complications.

Keywords Urinary tract infection, Urosepsis, Biomarkers, Drug resistance, Escherichia coli

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Background

Urogenic sepsis is a severe complication of urinary tract infections that poses a significant threat to both individual and public health [1-3]. In high-risk groups, it can progress to septic shock, which has a very high mortality rate [4, 5]. Additionally, it places a significant burden on socioeconomic and family financial status [6]. As urologic diseases and surgeries become more common, the risk of urinary tract infections and resulting bloodstream infections also increases. The therapeutic challenge is exacerbated by the increasing resistance of causative organisms to conventional antibiotics. For instance, a high proportion of *Escherichia coli, Enterococcus, Pseudomonas aeruginosa*, and *Klebsiella* exhibit resistance to commonly used antibiotics [7, 8]. The European Guidelines



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for Urinary Tract Infections offer valuable guidance for diagnosing and treating various urinary tract infections. However, clinical management remains challenging due to regional differences in causative organisms and drug resistance. Therefore, investigating the influencing factors of UTI-associated bloodstream infections and local pathogen characteristics is crucial to enhance preventive and therapeutic strategies.

To address this issue, the aim of this study was to optimize the therapeutic strategy by analyzing the clinical data, pathogenetic composition of common strains and drug resistance in patients with urogenic sepsis admitted to the General Hospital of Ningxia Medical University.

Data and methodology Participants

This study included individuals diagnosed with UTIs complicated by sepsis at Ningxia Medical University General Hospital between January 1, 2020, and December 1, 2023 in the Ningxia region of China. Sepsis is diagnosed according to the criteria of the American Society of Critical Care Medicine and the European Society of Intensive Care Medicine's Campaign to Save Sepsis: International Guidelines for the Management of Sepsis and Septic Shock 2016, combined with urological practice and the diagnostic criteria for urinary septicaemia: (1) a clinically confirmed urinary tract infection (UTI); (2) a Sequential Organ Failure Assessment (SOFA) score of 2 or greater; (3) sustained hypotension post-adequate fluid resuscitation requiring vasopressors to maintain a mean arterial pressure (MAP) of \geq 65 mmHg and serum lactate levels>2 mmol/L are the criteria for sepsis and septic shock [9]. Exclusions included only initial diagnoses of sepsis or septic shock resulting in replicative hospital admissions, infections originating from alternate loci, individuals younger than 18, and incomplete clinical data sets.

Analytical approach

In this retrospective study, patients were organized into three cohorts for comparative analysis: a control group with non-sepsis cases of simple urinary tract infections (CG), a septic non-shock group (NSSPU), and a septic shock group (USG).

The study received ethical approval from the Institutional Review Board (IRB) of Ningxia Medical University General Hospital on January 17, 2024, under approval number KYLL-2024-066. Owing to the retrospective nature of the investigation, the necessity for informed consent was waived, with the study utilizing pre-existing clinical data that were anonymized in adherence to the ethical standards stipulated by the Declaration of Helsinki.

Statistical analysis

Data analysis was facilitated using SPSS version 21.0. Normality assessments were conducted on the dataset. Median values were used to describe non-normally distributed enumerative data. The Kruskal-Wallis H test was used to discern disparities across groups for non-normally distributed metrics. Post hoc two-bytwo comparisons were made for factors with a *P*-value less than 0.05. Logistic regression analysis was utilized to investigate the correlation between the test indicators and sepsis, as well as to evaluate the impact of test indicators on the severity of the disease. Afterwards, a multi-factor analysis of statistically significant test indicators was performed using ROC curve analysis to assess the influence of relevant indicators on the severity of urogenital sepsis. This study evaluates the predictive value of relevant indicators for the severity of urosepsis by analyzing ROC curves and compares the diagnostic performance of various testing indicators. Categorical variables, such as gender, comorbidity profiles, and culture results, were presented as frequencies with percentages. Count data were analyzed using the χ^2 test with $R \times C$. Successive corrections were used when the theoretical frequency was less than 5. Fisher's exact probability method was used when the number of each cell less than 1. Statistically significant differences were considered at P < 0.05.

Results

Basic characteristics of urinary sepsis

This study analyzed clinical data from 111 patients diagnosed with sepsis, comprising 46 males and 65 females. Of these, 83 had positive urine cultures, while 28 had negative urine cultures. Moreover, 61 had positive blood cultures, and 50 had negative blood cultures. The age of the patients ranged from 18 to 97 years, with a median age of 67. The statistical analysis revealed that age was significantly associated with urosepsis (P < 0.05), individuals over the age of 65 are at a higher risk of developing urogenital sepsis (Table 1). Of the patients, 25 had a normal body temperature, while 86 had a fever. The median body temperature was 38.8 °C The median systolic blood pressure was 114 mmHg, and the median diastolic blood pressure was 70 mmHg. The median pulse rate was 90 beats per minute, and the median respiratory rate was 20 breaths per minute. Urine routine WBC $(+) \sim (++++)$. The study categorized patients with urinary sepsis into non-shock and shock groups. There were 59 cases in the non-shock group and 52 cases in the shock group. Additionally, 71 patients selected as a control group with non-sepsis cases of simple urinary tract infections (CG). Among these patients, death occurred in two cases,

Features	CG% (n=71)	NSSPU% (n = 59)	USG % (n=52)	P-value
Underlying disease				
Hypertension	1.41 (1)	20.60 (13)	37.50 (18)	< 0.001*
Diabetes mellitus	4.23 (3)	38.10 (24)	33.33 (16)	< 0.001*
Hypertension with diabetes mellitus	1.41 (1)	19.05 (12)	18.80 (9)	0.001*
Cardiovascular and cerebrovascular diseases	2.82 (2)	26.98 (17)	41.67 (20)	< 0.001*
Prostatic hypertrophy	5.63 (4)	6.35 (4)	10.42 (5)	0.763
Chronic kidney disease	7.04 (5)	41.27 (26)	62.50 (30)	< 0.001*
Urinary calculus	2.82 (2)	19.05 (12)	33.33 (16)	< 0.001*
Malignant tumor	0 (0)	14.29 (9)	14.58 (7)	0.002*
Post-urinary tract surgery	5.63 (4)	7.94 (5)	14.58 (7)	0.346
Hematologic disorders	0 (0)	19.05 (12)	33.33 (16)	< 0.001*
Malnutrition	0 (0)	20.63 (13)	39.58 (19)	< 0.001*
Fluid in the chest and abdomen	0 (0)	15.87 (10)	12.50 (6)	0.002*
Digestive tract disease	1.41 (1)	12.70 (8)	22.92 (11)	0.001*
Age distribution				
18–34	23.94 (17)	6.78 (4)	0 (0)	< 0.001*
35–49	5.63 (4)	5.09 (3)	7.69 (4)	
50-64	35.21 (25)	25.42 (15)	42.31 (22)	
>65	35.21 (25)	62.71 (37)	50.00 (26)	
Gender				
Male	22.54 (16)	40.68 (24)	42.31 (22)	0.031*
Female	77.46 (55)	59.32 (35)	59.69 (30)	
Blood culture				
Positive	4.23 (3)	50.85 (30)	59.62 (31)	< 0.001*
Negative	95.77 (68)	49.15 (29)	40.38 (21)	
Urine culture				
Positive	4.23 (3)	77.97 (46)	71.15 (37)	< 0.001*
Negative	95.77 (68)	22.03 (13)	28.85 (15)	

Table 1 Comparison of clinical and laboratory characteristics collected at hospital discharge in each study group

Values are represented as % (n) where n is the number of cases and (%) represents the percentage of the total within each column. If the superscript a or b is the same, there is no statistical difference between the two groups

*Indicates a statistically significant difference at P < 0.05

resulting in a mortality rate of 1.92% in the USG and 1.69% in the NSSPU.

Analysis of underlying diseases in the NSSPU and USG

Patients with urinary sepsis often have underlying diseases. In the NSSPU group, the most common underlying diseases were chronic kidney disease (41.27%), diabetes mellitus (38.1%), cardiovascular and cerebrovascular diseases (26.98%), malnutrition (20.63%), hypertension (20.6%), hypertension with diabetes mellitus (19.05%), and hematologic disorders (19.05%). In the USG, common underlying diseases included chronic kidney disease (62.5%), cardiovascular and cerebrovascular diseases (41.67%), malnutrition (39.58%), hypertension (37.5%), diabetes mellitus (33.33%), hematologic disorders (33.33%), hypertension with diabetes mellitus (18.8%), and post urinary tract surgery (14.58%). The statistical analysis revealed that the incidence of urinary sepsis was significantly associated with certain underlying diseases, such as hypertension and diabetes mellitus, but not with prostatic hypertrophy or post-urologic surgery (Table 1).

Normality test

To test the normality of the three groups of test indexes, skewness and kurtosis were used. It was found that leukocyte count (WBC), neutrophils (N%), neutrophil-to-lymphocyte ratio (NLR), platelet (PLT), D-dimer, total protein (TP), albumin (ALB), urea (URE), serum creatinine (SCR), and lactate (La) did not follow a normal distribution simultaneously.

Univariate analysis

After WBC, N%, NLR, PLT, D-Dimer, TP, ALB, URE, SCR, La, and other factors were subjected to the

Kruskal–Wallis test. The test results indicated statistically significant differences between the control group, the sepsis non-shock group, and shock group. For qualitative data, such as blood culture, urine culture, and gender distribution, significant differences between groups

 Table 2
 Comparison of test indicators across study groups

Test indicators	CG (<i>M</i>)	NSSPU (<i>M</i>)	USG (<i>M</i>)	Statistics (H)	P-value
WBC	8.35 ^a	13.02 ^b	42.61 ^b	15.90	< 0.001*
N%	71.90 ^a	87.40 ^b	92.80 ^c	90.38	< 0.001*
NLR	4.25 ^a	13.34 ^b	23.78 ^c	85.14	< 0.001*
La	1.10 ^a	2.01 ^b	2.65 ^c	89.28	< 0.001*
PLT	244.00 ^a	166.00 ^b	126.00 ^b	34.28	< 0.001*
D-dimer	0.82 ^a	2.01 ^b	5.10 ^b	83.38	< 0.001*
URE	5.09 ^a	8.73 ^b	9.96 ^b	44.52	< 0.001*
SCR	58.50 ^a	115.40 ^b	123.55 ^b	44.19	< 0.001*
TP	71.40 ^a	63.60 ^b	56.35 ^c	47.04	< 0.001*
ALB	41.30 ^a	34.10 ^b	29.35 ^c	59.71	< 0.001*

Data are presented as median (M)

*P-value < 0.05 is considered statistically significant. If the superscript a, b or c is the same, there is no statistical difference between the two groups

Table 3 Multiple logistic regression analysis of factors related to sepsis

were found using the Chi-square test (Table 1). Therefore, it can be inferred that patients who test positive for blood and urine cultures and are female are more likely to develop urogenic septic shock.

A post hoc two-by-two comparison was conducted to analyze the test indicators between different groups

The comparison revealed significant differences in N%, NLR, TP, ALB, and La between the CG and NSSPU, CG and USG, and NSSPU and USG. The test indices' median results in different groups indicate that TP and ALB values decrease with increasing sepsis severity, while N%, NLR, and La values increase (Table 2).

Analysis of the indicators for the progression of urinary tract infection to septic shock using multiple logistic regression

Compared to the non-sepsis cases of simple urinary tract infections, NLR and La have significant effects on the septic non-shock and septic shock. An increase in NLR and La levels increases the probability of septic nonshock to 1.3 times and 3.95 times, respectively, compared to when these levels are not increased. The probability of

Progression	Variable	В	S.E	Wald χ^2	Р	OR	95% CI
NSSPU	WBC	0.010	0.075	0.018	0.893	1.010	0.872-1.169
	Ν	0.010	0.044	0.049	0.825	1.010	0.926-1.102
Progression NSSPU	NLR	0.229	0.087	6.919	0.009*	1.257	1.060-1.492
	PLT	-0.010	0.004	7.322	0.007*	0.990	0.983-0.997
	La	1.373	0.433	10.069	0.002*	3.946	1.690-9.212
	TP	0.002	0.064	0.001	0.977	1.002	0.884-1.135
	ALB	-0.082	0.087	0.898	0.343	0.921	0.777-1.092
	SCR	0.011	0.006	3.034	0.082	1.011	0.999-1.023
	D-dimer	0.073	0.093	0.623	0.430	1.076	0.897-1.290
	URE	-0.137	0.090	2.349	0.125	0.872	0.731-1.039
USG	WBC	-0.020	0.083	0.060	0.807	0.980	0.833-1.153
	Ν	0.012	0.070	0.031	0.860	1.012	0.883-1.161
	NLR	0.318	0.092	12.054	0.001*	1.375	1.149–1.645
	PLT	-0.006	0.004	2.162	0.141	0.994	0.986-1.002
	La	2.106	0.473	19.794	< 0.001*	8.211	3.248-20.761
	TP	-0.071	0.083	0.722	0.395	0.932	0.792-1.097
	ALB	-0.071 0.117		0.368	0.544	0.932	0.741-1.171
	SCR	0.011	0.007	2.864	0.091	1.011	0.998-1.024
	D-dimer	0.094	0.096	0.962	0.327	1.099	0.910-1.327
	URE	-0.112	0.100	1.265	0.261	0.894	0.735-1.087

Multiple logistic regression analysis was performed to identify factors associated with different stages of sepsis severity. B represents the regression coefficient for each independent variable

S.E standard error

Wald χ^2 is the Chi-squared statistic from the Wald test. *P*-values show the level of statistical significance (**P*<0.05). The OR (odds ratio) value represents the degree of influence one variable has on another in the relationship between two categorical variables. Confidence intervals (CI) are provided at the 95% level

septic shock increases to 1.4 times and 8.2 times, respectively, as shown in Table 3.

Predictive value of N%, NLR, TP, ALB and La for urosepsis

We evaluated the diagnostic value of N%, NLR, TP, ALB and La levels in diagnosing sepsis by comparing them between the non-sepsis control group (CG) and sepsis groups (NSSPU and USG). Receiver operating characteristics (ROC) were also plotted (Fig. 1). At the maximum Youden index, the corresponding critical value was determined. The results showed that an increase of N% had diagnostic value for the severity of urogenital sepsis (AUC=0.913, 95% CI=0.871-0.956, P=0.000). The optimal threshold was 80.8%, at which the Youden index was 0.718, with a sensitivity of 90.1% and specificity of 81.7%. An increase in NLR% is a diagnostic indicator of the severity of urogenital sepsis (AUC=0.929, 95% CI=0.893-0.966, P=0.000), with an optimal cut-off value of 9.23, and a Jordon's index of 0.71. The sensitivity and specificity of this indicator are 86.5% and 84.5%, respectively. Similarly, elevated levels of La (AUC = 0.891, 95% CI=0.831-0.933, P=0.000) was found to be correlated with the severity of urogenic sepsis and showed some diagnostic value. However, severity of urinary sepsis correlated with changes in TP and ALB with some but not much specificity. Comparing the levels of N%, NLR and La between the septic non-shock group and the septic shock group, the cut-off values for diagnosing septic shock were identified as N% at 90.15%, NLR at 18.44, and

Compare the ROC curves of N%, NLR, TP, ALB, La, and the combined indicator to assess the differences in diagnostic performance for predicting uroseptic shock

80.80%, and 83.10%, respectively (Table 4).

Comparison of the predictive value differences for N%, NLR, TP, ALB, La, and the combined indicator in predicting uroseptic shock showed that there were significant differences in predictive value among N–TP, N–ALB, N–PRE, NLR–TP, NLR–ALB, NLR–PRE, La– TP, La–ALB, La–PRE, TP–PRE, and ALB–PRE. It is evident that the diagnostic performance of the combined indicator is significantly superior to that of the individual tests. Furthermore, the diagnostic performance of N, NLR, and La individually is higher than that of TP and ALB individually (Table 5).

Distribution characteristics of pathogenic bacteria

In the analysis of blood and urine samples, a total of 195 strains were identified, of which 155 were Gramnegative bacteria, accounting for 79.5% of the total number of isolates. Additionally, 29 Gram-positive bacteria (14.87%) and 11 fungi (5.60%) were also identified. Among patients with urinary sepsis, the most



Fig. 1 ROC curve analysis of N%, NLR, La, TP and ALB. **A** A larger test result indicates a more positive test; **B** smaller test results indicate a more positive test. **A** Orange solid line—La; purple solid line—TP; black solid line—ALB; steel blue solid line—N; green solid line—NLR; red solid line—baseline. **B** Steel blue solid line—TP; green solid line—ALB; orange solid line—baseline. *X*-axis: specificity; *Y*-axis: sensitivity; scale: the range for both the *x*-axis (specificity) and the *y*-axis (sensitivity) extends from 0.0 to 1.0, representing 0% to 100%. *AUC* area under the receiver operating characteristic curve

Test index		AUC	P value	95% CI	Threshold	Sensitivity (%)	Specificity (%)
Urosepsis	N%	0.913	< 0.001*	0.871-0.956	80.80	90.10	81.70
	NLR	0.929	< 0.001*	0.893-0.966	9.23	86.50	84.50
	La	0.891	< 0.001*	0.831-0.933	1.71	79.30	93.00
	TP	0.767	< 0.001*	0.699–0.836	39.40	100.00	0
	ALB	0.821	< 0.001*	0.758-0.885	18.65	100.00	0
USG	N%	0.773	< 0.001*	0.684-0.863	90.15	71.20	72.90
	NLR	0.775	< 0.001*	0.686-0.863	18.44	73.10	76.30
	La	0.772	< 0.001*	0.687-0.858	2.00	100.00	49.20
	TP	0.696	< 0.001*	0.598-0.795	39.40	100.00	0
	ALB	0.661	0.002*	0.558-0.763	18.20	100.00	0
	PRE	0.863	< 0.001*	0.795-0.931	42.10	80.80	83.10

Table 4 ROC curve analysis of	test Indicators in the	urosepsis
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The PRE variable can be used to assess the likelihood of an individual having a certain disease

PRE predicted probability, AUC area under the receiver operating characteristic curve

P-values show the level of statistical significance (*P<0.05). Confidence intervals (CI) are provided at the 95% level

Table 5 Comparative analysis of ROC curves for different diagnostic in	indicators in predicting uroseptic shock
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Test index	AUC difference	S.E	Statistics (Z)	Р	95% CI	
N–NLR	-0.001	0.297	-0.043	0.966	-0.119-0.058	
N-LA	0.001	0.299	0.016	0.987	-0.240-0.121	
N-TP	0.470	0.312	6.057	< 0.001*	0.318-0.622	
N-ALB	0.434	0.315	5.648	< 0.001*	0.283-0.585	
N-PRE	- 0.090	0.280	-2.416	0.016*	-0.145-(-0.017)	
NLR-LA	0.002	0.299	0.036	0.971	-0.249-0.127	
NLR-TP	0.471 0.312		6.138	< 0.001*	0.321-0.622	
NLR-ALB	0.435 0.315		5.639	< 0.001*	0.284-0.587	
NLR-PRE	-0.088	0.279	-2.673	0.008*	-0.129-(-0.024)	
LA-TP	0.469	0.307	6.842	< 0.001*	0.335-0.603	
LA-ALB	0.433	0.310	6.271	< 0.001*	0.298-0.568	
LA-PRE	-0.091	0.277	-2.270	0.023*	-0.157-(-0.012)	
TP-ALB	-0.036	0.314	-1.218	0.223	-0.116-0.022	
TP-PRE	-0.559	0.294	-7.828	< 0.001*	-0.281-(-0.419)	
ALB-PRE	-0.524	0.297	- 7.265	< 0.001*	-0.283-(-0.382)	

The PRE variable can be used to assess the likelihood of an individual having a certain disease

PRE predicted probability, AUC area under the receiver operating characteristic curve, S.E standard error

P-values show the level of statistical significance (*P<0.05). Confidence intervals (CI) are provided at the 95% level

commonly isolated Gram-negative bacteria were *Escherichia coli* (51.42%), followed by *Klebsiella pneu-moniae* (7.14%), and *Acinetobacter baumannii* (2.86%). Among Gram-positive bacteria, *Enterococcus faeca-lis* (14.29%) and *fungi* (2.86%) were the most common causative agents. In cases of urinary tract infections related to urosepsis, *Escherichia coli* and *Klebsiella pneumoniae* were the most commonly isolated pathogens, with detection rates of 65.75% and 9.59% (Table 6). Among patients with urogenic septic shock,

Escherichia coli showed significant antibiotic resistance, with rates exceeding 66.67% for antibiotics such as penicillin, cefazolin, ceftriaxone, and levofloxacin. The bacterial strain exhibited resistance rates of 61.9% to Aztreonam and 38.1% to cefepime. Among the cases of urinary tract infection, *Escherichia coli* showed the highest resistance rate to ampicillin at 88.57%, followed by ciprofloxacin and piperacillin at 68.57% and 62.86%, respectively. The resistance rate to levofloxacin was 61.90% (Table 7).

Features	Total (n = ⁻	182)	CG (n = 71)	NSSPU (n	= 59)	USG (n=52)		
	Strain (<i>n</i>)	Proportion (%)	Strain (<i>n</i>)	Proportion (%)	Strain (<i>n</i>)	Proportion (%)	Strain (<i>n</i>)	Proportion (%)	
Gram-negative bacilli	155	85.17	63	88.74	56	94.91	37	71.16	
E. coli	105	57.69	48	67.61	36	61.02	21	40.39	
Klebsiella pneumoniae	15	8.24	7	9.86	5	8.47	3	5.77	
Proteus mirabilis 7		3.85	1	1.41	2	3.39	4	7.69	
Acinetobacter baumannii	4	2.20	0	0	2	3.39	2	3.85	
Other	24	13.19	7	9.86	11	18.64	7	13.46	
Gram-positive bacilli	29	15.94	8	11.28	12	20.34	9	17.3	
Enterococcus faecium	21	11.54	3	4.23	10	16.95	8	15.38	
Enterococcus faecalis	5	2.75	3	4.23	2	3.39	0	0	
Other	3	1.65	2	2.82	0	0	1	1.92	
Fungus	11	6.04	2	2.82	2	3.39	7	13.46	

Table 6 Distribution of strains and their proportions across different groups

The total counts for each bacterial and fungal strain were calculated from the data collected across all patient groups within the study

Discussion

Urogenic sepsis is a type of sepsis caused by a bacterial infection in the urinary tract that spreads to other organs or tissues through the bloodstream, resulting in new lesions. According to Nagao et al., urogenic sepsis is more common in male patients [10, 11], particularly in the elderly population, and is typically caused by Gramnegative bacteria. The study indicates that patients aged over 65 are more prone to developing sepsis following a urinary tract infection. Bou-Anton et al. studied the prevalence and risk factors of Escherichia coli bacteremia in the United Kingdom over a 2-year period. The study found that both age and female gender were associated with an increased likelihood of *E. coli* bacteremia [12, 13]. Furthermore, we found that the prevalence of sepsis was lower in male patients (25.27%) than in female patients (35.71%). This gender difference is likely to be related to region-specific health conditions and management [14]. When analyzing the underlying diseases of sepsis, it was found that patients in the non-shock group had mainly chronic kidney disease (41.27%), diabetes mellitus (38.1%), and cardiovascular disease (26.98%). In contrast, patients in the shock group had chronic kidney disease (62.5%), cardiovascular disease (41.67%), and malnutrition (39.58%) more commonly. In addition to prostatic hypertrophy and urological surgery, other underlying diseases were found to be statistically significant in the development of urogenic sepsis and may increase the risk of sepsis [15, 16]. Positive urine and blood cultures were significantly associated with the development of urogenital sepsis. It can be inferred that positive urine and blood culture results may increase the risk of sepsis.

The field of critical care medicine has made considerable advancements in the diagnosis and management of septic shock. The "Campaign to Save Sepsis: International Guidelines for the Management of Sepsis and Septic Shock 2016", endorsed by the American College of Critical Care Medicine and the European Society of Intensive Care Medicine, identifies a serum lactate level above 2 mmol/L as a critical marker for septic shock diagnosis. Studies demonstrate that this threshold boasts a sensitivity of 82.5% and a specificity of 22.4%, with a 95% confidence interval (CI) ranging from 0.816 to 0.834 [17]. Additionally, the NLR has emerged as an important biomarker in detecting systemic inflammation [18–20]. In research conducted by Wu et al., it was found that an NLR value of 11.57 could serve as a diagnostic threshold for septic shock, exhibiting a sensitivity and specificity of 100%. This study explored the differences in N% across control groups, the septic shock group and non-shock group, finding that N% is statistically significant with an area under the curve (AUC) of 0.885. A cut-off value of 77% for N% in urinary sepsis diagnosis was established, yielding a sensitivity of 93.8% and specificity of 69.6% [21]. While in the present study, the receiver operating characteristic (ROC) curve analysis of patients diagnosed with septic shock showed that the AUC for La, NLR and N% were an impressive 0.772, 0.775 and 0.773, with corresponding sensitivities of 100%, 73.10% and 71.20%, and specificities of 49.20%, 76.30% and 72.90%. The 95% CIs, when setting diagnostic thresholds at 2.00 mmol/L for La, 18.44 for NLR and 90.15% for N%, ranged between 0.687-0.858, 0.686-0.863 and 0.684-0.863, respectively. The diagnostic threshold for La reported in this study aligns with prior research. Nonetheless, the broad confidence interval, possibly due to a small sample size for septic shock incidents, limits the precision, highlighting the need for larger sample sizes to enhance the accuracy.

Antibiotic	CG (n=48)		NSSPU (n=3	36)	USG (n=21)		Total (n = 105)		
	Drug resistance rate% (<i>n</i>)	Sensitivity % (n)	Drug resistance rate% (n)	Sensitivity % (n)	Drug resistance rate% (n)	Sensitivity % (n)	Drug resistance rate% (n)	Sensitivity % (n)	
Ampicillin	87.50 (42)	10.42 (5)	81.08 (30)	16.67 (6)	100.00 (21)	0 (0)	88.57 (93)	10.48 (11)	
Piperacillin	54.17 (26)	12.50 (6)	69.44 (25) 25.00 (9)		71.43 (15)	14.29 (3)	62.86 (66)	17.14 (18)	
Ampicillin/sul- bactam	43.75 (21)	29.17 (14)	41.67 (15)	25.00 (9)	38.10 (8)	19.05 (4)	41.90 (44)	25.71 (27)	
Cefazolin	45.83 (22)	54.17 (26)	55.56 (20)	44.44 (16)	71.43 (15)	23.81 (5)	54.29 (57)	44.76 (47)	
Cefuroxime	43.75 (21)	43.75 (21)	55.56 (20)	38.89 (14)	66.67 (14)	28.57 (6)	52.38 (55)	39.05 (41)	
Cefoperazone/ sulbactam	6.25 (3)	89.58 (43)	0 (0)	100.00 (36)	9.52 (2)	80.95 (17)	4.76 (5)	91.43 (96)	
Ceftazidime	31.25 (15)	56.25 (27)	33.33 (12)	58.33 (21)	42.86 (9)	38.10 (8)	34.29 (36)	53.33 (56)	
Cefepime	27.08 (13)	68.75 (33)	27.78 (10)	52.78 (19)	38.10 (8)	42.86 (9)	29.52 (31)	58.10 (61)	
Ceftriaxone	47.92 (23)	52.08 (25)	55.56 (20)	55.56 (20) 44.44 (16)		80.95 (7)	54.29 (57)	45.71 (48)	
Ciprofloxacin	66.67 (32)	22.92 (11)	63.89 (23)	25.00 (9)	80.95 (17) 9.52 (2)		68.57 (72)	20.95 (22)	
Levofloxacin	58.33 (28)	8.33 (4)	58.33 (21)	30.56 (11)	76.19 (16)	76.19 (16) 19.05 (4)		18.10 (19)	
Amikacin	4.17 (2)	95.83 (46)	2.78 (1)	97.22 (35)	4.76 (1)	95.24 (20)	3.81 (4)	96.19 (101)	
Gentamicin	43.75 (21)	56.25 (27)	30.56 (11)	69.44 (25)	57.14 (12)	42.86 (9)	41.90 (44)	58.10 (61)	
Imipenem	0 (0)	100 (48)	0 (0)	100 (36)	0 (0)	100.00 (21)	0 (0)	100.00 (105)	
Meropenem	0 (0)	100 (48)	0 (0)	100 (36)	0 (0)	100.00 (21)	0 (0)	100.00 (105)	
Cotrimoxazole	47.92 (23)	52.08 (25)	58.33 (21)	41.67 (15)	76.19 (16)	23.81 (5)	57.14 (60)	42.86 (45)	
Piperacillin– tazobactam	4.17 (2)	89.58 (43)	2.78 (1)	97.22 (35)	14.29 (3)	80.95 (17)	5.71 (6)	90.48 (95)	
Aztreonam	33.33 (16)	66.67 (32)	44.44 (16)	55.56 (20)	61.90 (13)	38.10 (8)	42.86 (45)	57.14 (60)	
Tobramycin	12.50 (6)	58.33 (28)	8.33 (3)	69.44 (25)	23.81 (5)	80.95 (7)	13.33 (14)	57.14 (60)	
Furantoin	4.17 (2)	87.50 (42)	2.78 (1)	86.11 (31)	19.05 (4)	76.19 (16)	6.67 (7)	84.76 (89)	
Ticarcillin/cla- vulanic acid	8.33 (4)	81.25 (39)	5.56 (2)	66.67 (24)	9.52 (2)	85.71 (18)	7.62 (8)	77.14 (81)	
Tigecycline	0 (0)	100.00 (48)	0 (0)	100.00 (36)	0 (0)	100.00 (21)	0 (0)	100.00 (105)	
Minocycline	10.42 (5)	75.00 (36)	8.33 (3)	80.56 (29)	28.57 (6)	57.14 (12)	13.33 (14)	73.33 (77)	
Doxycycline	20.83 (10)	50.00 (24)	16.67 (6)	66.67 (24)	38.10 (8)	38.10 (8)	22.86 (24)	53.33 (56)	
Colistin	4.17 (2)	93.75 (45)	0 (0)	83.33 (30)	0 (0)	90.48 (19)	1.90 (2)	89.52 (94)	

Tal	ble 7	Ana	vsis o	f druc	a resistanc	e in	Esci	herich	<i>піа со</i> і	i iso	lated	from	different	aroup	s [strains	(%))]
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Both the drug resistance rate and sensitivity rate are expressed as percentages, with the number in parentheses indicating the corresponding number of bacterial strains. All data have been analyzed according to standardized statistical procedures

n number of bacterial strains analyzed

The slightly elevated diagnostic threshold for the NLR may be attributed to a swift rise in neutrophil counts, a response often observed during episodes of acute stress, pain, or trauma [22]. Furthermore, the surge in endogenous cortisol and catecholamine levels can exacerbate this effect, leading to increased neutrophil and decreased lymphocyte counts. The influence of other hormones and cytokines on these parameters cannot be discounted. Utilizing logistic regression analysis, a significant positive correlation was established between the levels of blood La, and NLR in the context of urogenic sepsis, with respective regression coefficients of 1.373 and 0.229, all bearing statistical significance with *P*-values less than 0.05. These findings underscore the value of lactate and

NLR not simply as diagnostic tools for septic shock but also as indicators to gauge the condition's severity and monitor its trajectory [23].

The total protein in the serum is made up of two main categories: ALB and globulin. These components are important for assessing nutritional status and diagnosing various diseases. ALB maintains osmotic pressure balance in plasma, which is essential for normal bodily functions. Hypoproteinemia is frequently observed in patients with sepsis at the onset of the condition. Several studies have confirmed a strong correlation between hypoproteinemia and poor prognosis in critically ill patients, indicating its significance as a prognostic indicator [16, 24]. In cases of sepsis where the initial concentration of ALB in the plasma drops below 20 g/L, failure to replenish ALB immediately can result in life-threatening complications. The study analyzed predictive curves to determine the area under the curve (AUC) for TP in relation to the severity of sepsis. The results showed that a TP threshold of 39.4 g/L had a sensitivity of 100% for diagnosing sepsis, but a specificity of 0%. The AUC for ALB was also analyzed and found to be 0.179. The diagnostic cut-off for ALB was set at 18.65 g/L, achieving a sensitivity of 100%. However, its specificity was as low as 1.4%. Therefore, while TP and ALB levels can predict sepsis to some extent, their effectiveness as prognostic tools is notably limited. This study only included two fatal cases, thus could not provide a more comprehensive prognosis of patients. Multiple studies have shown that nutritional assessment can predict mortality rates [25].

Kiiru et al. conducted research that identified Escherichia coli as the predominant pathogen in UTIs [26]. Literature reviews indicate that the primary pathogens causing UTIs are Escherichia coli, Klebsiella pneumoniae, Pseudomonas aeruginosa, and Acinetobacter baumannii [27]. The hospital's surveillance data show that Escherichia coli has the highest detection rate in UTIs, standing at 53.85%. This is consistent with the range of 42% to 69.3% reported by Nicolle et al. [28]. The most prevalent Gram-negative bacteria isolated from patients with urosepsis in our hospital are Escherichia coli (51.42%), Klebsiella pneumoniae (7.14%), and Acinetobacter baumannii (2.86%). The Gram-positive bacteria most commonly isolated from these patients are Enterococcus faecalis (14.29%) and Enterococcus faecium (2.86%), along with fungi (2.86%). In septic shock patients of urologic origin, Escherichia coli shows high resistance rates to a majority of antibiotics, including penicillin, cefazolin, ceftriaxone, and levofloxacin, all exceeding a resistance rate of 66.67% [2]. Additionally, this bacterial strain exhibits a 61.9% resistance rate to aztreonam and a 38.1% resistance rate to cefepime. In the case of urinary tract infections, *Escherichia coli* displays the highest resistance rate to ampicillin, at 88.57%, which is slightly higher than the 80% rate found among E. coli isolates studied by Lau, Peng, and others [29, 30].

Conclusion

Blood markers are essential for assessing the severity of urogenital sepsis; elevated levels of WBC, N%, NLR, D-dimer, URE, SCR, and La suggest worsening conditions, whereas decreased PLT, TP, and ALB levels indicate disease progression. Especially in diagnosing uroseptic shock, combined testing of multiple indicators has greater diagnostic value compared to single indicators. *E. coli* plays a critical pathogenic role in sepsis, particularly its drug-resistant strains. Considering risk factors such as age, chronic diseases, and low socioeconomic status, the necessity of resistance monitoring is highlighted to guide therapy and mitigate antibiotic resistance issues.

Strengths and limitations

This study reports the diagnostic data, pathogenetic distribution, and drug resistance of urogenic sepsis in the General Hospital of Ningxia Medical University in the Ningxia region of China. However, there are some limitations to this study, including a low number of deaths (only two) and a lack of prognostic causes for analysis.

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Author contributions

Jia Wei—participated in the revision, responsible for the overall study planning, design and implementation, approved the final release version Gang Li—verify the feasibility of collecting cases, participate in discussions and revisions, provide experimental equipment, technical and financial support Yanxia Shao—writing thesis manuscript, case collection, processing data and analyzing data, iconography, reviewing literature.

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Data availability

No datasets were generated or analysed during the current study.

Declarations

Ethics approval and consent to participate

This survey was conducted in accordance with the Declaration of Helsinki and the ethical standards of the national and institutional guidelines. The ethical approval for the study protocol was granted by the Medical Research Ethics Committee of the General Hospital of Ningxia Medical University [Approval No. KYLL-2024-066].

Consent to publication

The author confirms and consents that all materials included in the submitted manuscript, such as photographs and tables, have received the proper ethical clearance and informed consent from the relevant participants. The author has presented the content of the manuscript to the participants and has obtained their agreement. The author is committed to cooperating and will promptly provide all necessary materials to ensure the seamless publication of the manuscript.

Competing interests

The authors declare no competing interests.

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