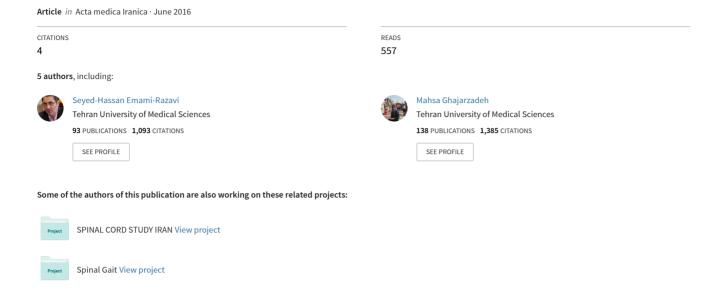
Incidence of Post-Operative Sepsis and Role of Charlson Co-Morbidity Score for Predicting Postoperative Sepsis



Incidence of Post-Operative Sepsis and Role of Charlson Co-Morbidity Score for Predicting Postoperative Sepsis

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Abstract- Sepsis and septic shock are among mortality causes following major surgeries. The Charlson comorbidity index consists of 19 weighted categories related to chronic health which measures the burden of co-morbidity. The goal of this study was to determine the incidence of postoperative sepsis in patients underwent gynecological and gastrointestinal cancer surgeries and predictive role of Charlson index for this situation. Two hundred and twenty-two patients who underwent gynecological and gastrointestinal cancer surgeries were evaluated. Sixty-four (28.6%) patients developed SIRS postoperatively. Forty-four (19.7%) patients developed sepsis postoperatively. Mean age, duration of hospitalization and surgery, the Charlson score were significantly higher in patients who developed sepsis than other cases. Blood transfusion and Charlson score were independent predictors of sepsis occurrence. Charlson co-morbidity index is a predictive factor for developing postoperative sepsis.

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Keywords: Sepsis; Post operative; Iran

Introduction

Sepsis and septic shock are among mortality causes following major surgeries (1).

Effects of anesthetic drugs, hormonal changes related to stress, rate of peri-surgical bleeding, transfusion, occurrence of ischemia—reperfusion, and duration of surgery are among post-surgical immune suppression factors (2).

Prophylactic therapies are widely used before procedures which have helped physicians to control post-operative sepsis and septic shock (3).

Factors like age, sex, underlying illness and comorbidities, emergent surgeries, the degree of bleeding during surgery, and transfusion of allogeneic blood play a critical role in developing post-operative shock (4-9). Systemic inflammatory response syndrome (SIRS), sepsis, severe sepsis, and septic shock are clinical features with increasing mortality from SIRS to septic shock (10). Early identification of patients at risk of developing post-operative sepsis could help applying strategies to reduce post-operative sepsis. The Charlson co-morbidity index consists of 19 weighted categories related to chronic health which measures the burden of

co-morbidity (11).

Higher levels of chronic co-morbidities are associated with higher risk of developing complications. So, application of this tool before major surgeries will help the physicians to identify high-risk cases and consider proper prophylactic therapies and strategies.

The goal of this study was to determine the incidence of postoperative sepsis in patients underwent gynecological and gastrointestinal cancer surgeries and predictive role of Charlson index for this situation.

Materials and Methods

In this study which conducted between September 2012 and September 2013 in Imam hospital (affiliated hospital of Tehran university of medical sciences), 250 patients who were a candidate for genitourinary or gastrointestinal cancer surgery were enrolled. Inclusion criteria were age more than 18 and candidate for surgery due to cancer of genitourinary or gastrointestinal tracts. Exclusion criteria were: age under 18, emergency surgery, evidence of sepsis prior to the surgery.

During the study period, a trained surgery resident recorded patient's data.

She calculated The Charlson co-morbidity score (CCS) for each patient preoperatively. Demographic characteristics (age,sex), duration of prophylactic antibiotic, everyday vital sign, laboratory findings, ICU (Intensive care unit) admission, units of blood transfusion and length of stay were recorded.

All cases followed up to 6 days after surgery. Temperature, respiratory rate, pulse rate and white blood cell counts recorded for all patients for 6 consecutive days by a trained nurse.

Systemic inflammatory response syndrome (SIRS) was sought from the observation charts and laboratory results daily. SIRS, sepsis, and septic shock were defined as per the American College of Chest Physicians/Society of Critical Care Medicine Consensus Conference criteria (12).

If the SIRS criteria were met, case notes, laboratory results, radiographs, computed tomography scan were re-evaluated to find the source.

Chest" was considered as the source of sepsis if there were one or more of purulent sputum, worsening hypoxemia, new infiltrates on chest radiograph or positive sputum culture.

"Abdomen" was considered as the source of sepsis if there was an imaging modality demonstrating an intraabdominal collection, or growth of pathogenic bacteria in any specimen of intra-abdominal drain fluid.

"Urinary tract" was considered as the site of infection if there was the positive growth of a pathogen

in a mid-stream specimen or catheter specimen of urine.

If blood culture was positive but microbiological culture from another site was negative or imaging results were negative, according to organism cultured from blood, type of surgery and clinical signs, the sepsis was attributed to the most likely source of primary infection.

Surgical site infection considered if surgical site culture was positive.

All data were analyzed using SPSS software version 18.0 (SPSS Inc., Chicago, IL, USA).

Student's t-test and Fisher's exact tests were used to compare continuous and categorical variables.

P-value less than 0.05 was considered as significant.

Results

Twenty-seven patients were excluded. Finally, 223 patients evaluated. The mean age of patients and mean duration of hospitalization were 49.8±9.9 years and 12.7±7.4 days. One hundred and twenty-two were male (54.7%), and 101 (45.3%) were female. Total colectomy, sub-total gastrectomy were the most common surgical procedures. The mean duration of surgery was 3 hours. One hundred and thirty-five (60.5%) received allogeneic blood intra-operatively, and median CHARLSON score was 3 (range 0-7).

The most common prophylaxis antibiotics were a combination of ciprofloxacin+ metronidazole (Table 1).

Table 1. Prophylactic antibiotics applied to

patients		
	Number (%)	
Cephazoline	14(6.2%)	
Cephazoline+Metronidazole	15(6.7%)	
Cephteriaxone+ Metronidazole	26(11.7%)	
Ciprofloxacin	2(0.9%)	
Ciprofloxacin+ Metronidazole	166(74.5%)	

Sixty-four (28.6%) patients developed SIRS postoperatively. Forty-four (19.7%) patients developed sepsis postoperatively.

The source of sepsis was urinary tract (6 patients), chest (2 cases), and abdomen (15 cases). In 15 patients, positive culture of surgical site recorded.

Thirty-five out of 44 patients who developed sepsis admitted to critical care unit (P<0.001), and 29 (65.9%) received allogeneic blood intraoperatively (*P*=0.005).

There was no significant difference between two sex groups in cases developed sepsis (23 male and 21 female, P=0.2). Nine patients (20%) who developed sepsis died.

Mean age, duration of hospitalization and surgery were significantly higher in patients who developed sepsis than other cases (Table 2). Blood transfusion and CHARLSON score were independent predictors of developing post operative sepsis (Table 3).

Table 2. Mean age, duration of hospitalization, surgery and vital sign in patients with and without sepsis

	with a	nd without seps	515	
	SIRS	Mean	Std. deviation	P.value
ACE	YES	53.7895	8.55873	0.007
AGE	NO	49.0117	10.05631	0.007
Duration of	YES	22.8000	12.23530	< 0.001
hospitalization(day)	NO	10.4463	2.47458	<0.001
D ()	YES	195.4167	50.64407	<0.001
Duration of surgery (min)	NO	151.5515	42.21524	< 0.001
CHARLCON	YES	4.1026	1.09532	<0.001
CHARLSON	NO	2.7045	1.32802	< 0.001
PR1	YES	100.7750	2.75948	< 0.001
IKI	NO	95.6978	5.91766	٥.001
RR2	YES	19.1000	2.71558	< 0.001
KKZ	NO	16.7747	2.15397	-0.001
RR3	YES	18.6000	2.75309	< 0.001
KKS	NO	17.0604	12.37453	<0.001
DD4	YES	18.7000	3.16390	< 0.001
RR4	NO	15.7747	1.96624	\0.001
DD5	YES	19.2500	3.46965	<0.001
RR5	NO	15.6429	1.81733	< 0.001
DD4	YES	100.1351	5.13789	0.002
PR6	NO	90.0000	4.00000	0.002
RR1	YES	19.4750	2.68889	< 0.001
Turi	NO	17.3901	2.61033	0.001
RR2	YES	19.1000	2.71558	< 0.001
	NO	16.7747	2.15397	
RR3	YES	18.6000	2.75309	0.4
	NO	17.0604	12.37453	
RR4	YES	18.7000	3.16390	< 0.001
	NO	15.7747	1.96624	*****
RR5	YES	19.2500	3.46965	< 0.001
	NO	15.6429	1.81733	
RR6	YES	19.3784	3.49087	0.1
KKU	NO	16.0000	1.00000	0.1
m4	YES	38.0500	.15852	0.001
T1	NO	37.9258	.23160	0.001
TPA	YES	37.9875	.12848	<0.001
T2	NO	37.6011	.31395	< 0.001
Т3	YES	37.9025	.25165	< 0.001
13	NO	37.2687	.29350	\0.001
T4	YES	45.4000	47.46841	0.01
	NO	37.1049	.27336	0.01
T5	YES	38.0750	.29936	< 0.001
	NO	37.0324	.24696	0.001
T6	YES	38.0811	.20526	< 0.001
10	NO	37.2250	.38622	\0.001
****	YES	12830.0000	2237.92486	0.001
WBC1	NO	10367.1429	2258.93427	< 0.001
WBC2	YES	12440.0000	2342.12264	<0.001
WBC2	NO	9393.9560	1490.12097	< 0.001
WBC3	YES	11685.0000	2256.79032	< 0.001
	NO	8778.5714	1100.74417	-0.001
WBC4	YES	11667.5000	2392.65838	< 0.001
HUCT	NO	8529.6703	1010.22338	~0.001
WBC5	YES	12005.0000	2685.66470	< 0.001
H DC3	NO	8344.5055	914.67322	\0.001
WBC6	YES	12275.6757	2699.11564	< 0.001
= •••	NO	6451.5000	4353.26035	001

PR: pulse rate RR: Respiratory rate T: temperature WBC: White blood cells

Table 3. Logistic regression analysis considering sepsis as dependent and other variables as independent factors

	OR(95% CI)	P.VALUE	
Sex (male/female)	0.8(0.3-1.8)	0.6	
Blood transfusion	3.4(1.2-9.7)	0.02	
CHARLSON score	0.3(0.2-0.5)	< 0.001	
Age	0.9 (0.9-1)	0.1	

Discussion

The result of the current study showed that postoperative sepsis occurred in 19% of cases who underwent GI cancer surgery while 28% developed SIRS postoperatively. This rate was lower than the rate reported by Hampshire *et al.*, .

They evaluated 101 patients who underwent elective major surgery and reported postoperative sepsis in 27% and SIRS in 58% (1). In another study, Mokart *et al.*, evaluated 93 cases who underwent major oncological surgery and investigated that 19 patients (20.4%) developed a severe sepsis after surgery (13). seven (36%) of the septic patients died. Nine patients (20%) who developed sepsis died in our study. In current study sex was not an independent predictor of postoperative sepsis while in Mokart (13) male gender was an independent predictor (OR 4.7, 95% CI between 1.5 and 15.5, *P*<0.01). Offiner *et al.*, reported postoperative sepsis rate as 40% (14). These different rates could be due different inclusion and exclusion criteria and different settings in different studies.

We found that Charlson score was significantly different in cases with and without SIRS development. This finding is compatible with Mokart *et al.*, findings and against Hampshire *et al.*, findings (1,13). It has been reported that Charlson score equal or more than 6 is predictive of developing postoperative severe sepsis. As the results show, the Charlson score was significantly different between SIRS-positive and negative cases. In this score, for instance, metastatic cancer is heavily weighted which could strongly contribute to development of postoperative sepsis.

Charlson score is a simple, objective scoring system which consists of 19 weighted categories related to chronic health that measures the burden of co-morbidity (11).

It could be used easily and completely in a short time, but it has some disadvantages.

First, it does not include all co-morbidities. Second, some co-morbidities included in the score are not directly related to developing SIRS (15).

In the current study, mean age of patients who developed SIRS was significantly higher than the mean age of cases who did not develop SIRS. In Mokart *et al.*, and Hampshire *et al.*, studies, age was not significantly different between septic and not septic groups (1,13).

People older than 65 years old are more likely to have chronic conditions which could contribute to developing SIRS and sepsis. We also investigated that mean duration of hospitalization and meant duration of surgery were significantly higher in SIRS group which is compatible with Hampshire *et al.*, findings (1).

This study had some limitations. First, it conducted in a referral hospital. Second, all cases who underwent all types of surgery were not included.

Multicenter studies with larger sample sizes are recommended.

Charlson co-morbidity index is a predictive factor for developing postoperative sepsis.

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