

ORIGINAL RESEARCH

Study of BISAP Score in Evaluation of severity of Acute Pancreatitis in relation to MCTI score¹Dr. Shubham Agrawal, ²Dr. Jony Samria, ³Dr. S.S. Minhas¹Final year resident, ²Assistant Professor, ³Professor, Department of General Surgery, Maharishi Markandeshwar Medical College and Hospital, Kumarhatti, Solan, Himachal Pradesh India**Correspondence:**

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Abstract**Aim:** To assess severity of acute pancreatitis by means of BISAP score at the time of patient admission.**Material and Method:** This prospective study was conducted in the Department of General Surgery at Maharishi Markandeshwar Medical College and Hospital, Kumarhatti, Solan from May 2021 to April 2022 and included 37 patients of acute pancreatitis for period of 1 year. Following admission all patients underwent routine hematological and biochemical investigations and BISAP score was calculated. Modified CTSI score was calculated at 48 hours for assessing the severity of acute pancreatitis and this score was compared with BISAP score.**Results:** The mean age of the patients was 48.14±15.99 years with slight male preponderance. The main cause for acute pancreatitis was gall stones (70.3%) followed by alcohol (27%). Majority of patients had moderate disease (45.9%) while (32.4%) had mild disease and (21.6%) patients had severe disease. BISAP score of 3 and 4 was able to detect 100% patients with severe MCTSI and the association between MCTSI and BISAP was statistically significant ($p<0.001$).**Conclusions:** The BISAP score is a relatively good indicator of severity of acute pancreatitis and the cut-off value of ≥ 3 can be a good predictor according to present study for predicting severe acute pancreatitis.**Keywords:** Acute pancreatitis, BISAP score, Modified CTSI.**Introduction**

Acute pancreatitis is an acute inflammatory response resulting in local pancreatic damage caused by abnormal pancreatic enzyme activation. The process is initiated by co-localization of zymogen granules and lysosomes inside the acinar cells releasing cathepsin B in cytosol resulting in acinar cell death. This initial process is further aided by local inflammatory response which is initiated at the site of injury but can lead to systemic inflammation, if marked. In majority of the patients this inflammatory cascade is self-limited; however, in remaining patients it persists. The mortality in early phase of acute pancreatitis is resultant of this persistent immune response.

According to revised Atlanta classification, acute pancreatitis has been graded as mild, moderate and severe types¹. 80% of acute pancreatitis instances are mild, self-limiting, and have no long-term effects. But when necrosis occurs in portions of the pancreas and the surrounding tissues, serious illness often develops in 10-20% of patients. These individuals develop an initial

inflammatory reaction, which develops into a systemic inflammatory response syndrome, which results in multiorgan failure and death.^{2,3}

It might be challenging to predict how someone would react to pancreatic damage because it differs from person to person. Although the general death rate for acute pancreatitis is 2-5%, severe cases might have a mortality rate of up to 20-30%.^{4,5}

Because severe acute pancreatitis (SAP) has a high death rate, it is necessary to develop early and reliable methods for determining the disease's severity in order to start more active therapies that would lessen the disease's negative effects and high mortality. In order to forecast the SAP, meticulous continuous clinical examinations, a multi-factor grading system, and imaging investigations are needed.^{4,5}

A several multifactorial scoring systems comprising clinical and biochemical criteria for assessment of severity of acute pancreatitis have been in use for several years. The Glasgow score (8 criteria), MOSS score (12 criteria), BISAP score (5 criteria), the acute physiology and chronic health evaluation (APACHE II) score (14 criteria), and the 11 criteria outlined by Ranson et al. in the 1970s are among the. Depending on the cut-off value and the timing of scoring, the sensitivity and specificity of these scoring systems for determining severe acute pancreatitis ranges between 55% and 90%.⁶

The present study was conducted to assess the clinical outcomes of patients with acute pancreatitis and classify it according to BISAP score.

Material and methods

The Present single centric, prospective, observational and hospital-based study was conducted at Department of General surgery, Maharishi Markandeshwar Medical College and Hospital, Kumarhatti, Solan for period of 1 years. The sample size of 37 patients was undertaken for present study.

Selection of patients

Patients having clinical presentation of acute pancreatitis with ultrasound findings suggesting of acute pancreatitis with serum amylase or lipase level raised more than 3 times of normal cut-off value were admitted in surgery department.

Inclusion criteria

1. Patients with history of acute abdominal pain 24 hours prior to hospital admission.
2. Patients with increased serum amylase or lipase levels of more than 3 times.
3. Patients with ultrasonographic abdominal changes consistent with acute pancreatitis on hospitalization.

Exclusion criteria

1. Patients admitted to hospital with other diseases like coronary artery disease, diabetes mellitus, chronic kidney disease, chronic lung disease etc. along with acute pancreatitis at the time of hospital admission.
2. Patients with chronic pancreatitis and pancreatic malignancy.

History of the patient, general physical examination and clinical examination was done (Name, age, sex, duration of symptoms, history of epigastric pain, severity, nature, number of attacks of pain, whether radiated to back or not, does pain relieve on bending forward, history of alcohol intake, history of jaundice, history of gall stone, number of times admission required in the hospital in past, history of fever, dyspepsia, previous abdominal surgeries).

Routine hematological and biochemical investigations were done and BISAP score was calculated simultaneously. Modified CT score was calculated at 48 hours for assessing the

severity of Acute pancreatitis. It was considered to be gold standard and was compared with BISAP score.

Data analysis was done using SPSS ® Software version 26. The categorical variables were compared using Fisher's exact test and for group variables ANOVA test was used. p-value of <0.05 was considered to be significant and value of <0.001 considered to be highly significant.

Results

The mean age of the patients was 48.14 ± 15.99 years. There was slight male predominance with 51.4% of patients being male and 48.6% being females. The main cause for acute pancreatitis was gall stones (70.3%) followed by alcohol (27%) only 2.7% cases were due to idiopathic cause. (figure 1). Majority of patients according to MCTSI scoring had a moderate disease (45.9%) while 32.4% had mild disease (figure 2); 21.6% of patients had severe disease. 21 out of total 37 patients developed local complications. It was observed that acute necrotic collection was only present in 2 patients of moderate disease vs 4 patients of severe disease (11.8% vs 50%) while PP fluid collection was only present in 8 patients of moderate disease and 1 of severe (47.1% vs 12.5%). Pseudocyst was present in 3 patients of moderate disease (17.6%). While it was present in none of the patients of severe disease. WON was present in 3 patients of severe disease (37.5%) (table 1). 75% (6/8) patients of severe acute pancreatitis developed systemic complications while only 5.9% of moderate presented with systemic complications (ARDS; 1/17). Maximum patients in SAP had ARDS (37.5%) followed by multiorgan failure, paralytic ileus and renal failure (12.5% each). This association was statistically significant ($p < 0.001$) (table 2). In case of mild disease none of the patients needed ICU admission while in moderate disease 3(17.6%) patients needed ICU admission. In case of severe disease 6(75%) patients needed ICU admission. This association was statistically significant ($p < 0.05$) (table 3). Significant differences were observed in duration of hospital stay with hospital stay being highest in cases with severe pancreatitis followed by moderate and mild ($p < 0.001$) (table 4). 34 (91.9%) of the patients not needed referral while 3 (8.1%) patients needed referral (figure 3). Conservative approach was used to treat most of the patients. Only one patient with severe acute pancreatitis needed cysto-jejunostomy. This association was statistically not significant. The association between MCTSI and BISAP was statistically significant ($p < 0.001$). BISAP score of 3 and 4 was able to detect 100% patients with severe MCTSI thus the cut of ≥ 3 can be a good predictor according to present study for predicting SAP(table 4).

Figure 1: etiology of acute pancreatitis **Fig 2: classification of pancreatitis according to MCTSI**

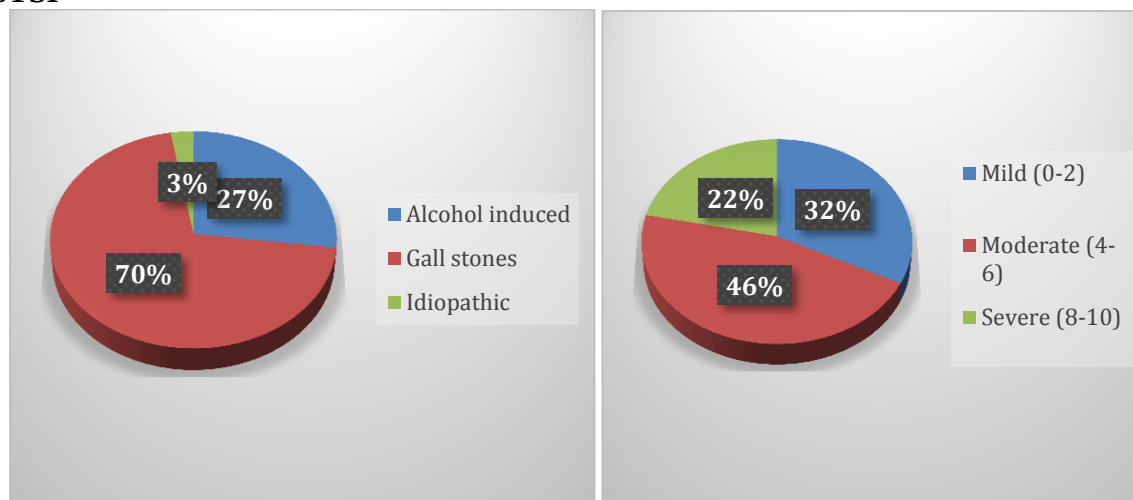


Table-1: Distribution of patients with local complications

Local complications	Moderate		Severe		P Value
	N	%	N	%	
Acute necrotic collection	2	11.8%	4	50.0%	0.0001***
PP fluid collection	8	47.1%	1	12.5%	
Pseudocyst	3	17.6%	0	0.0%	
WON	0	0.0%	3	37.5%	
None	4	23.5%	0	0.0%	
	17	100.0%	8	100.0%	

Table-2: Distribution of patients with systemic complications

Systemic complications	Moderate		Severe		P Value
	N	%	N	%	
ARDS	1	5.9%	3	37.5%	0.001**
MOF	0	0.0%	1	12.5%	
Paralytic ileus	0	0.0%	1	12.5%	
Renal Failure	0	0.0%	1	12.5%	
None	16	94.1%	2	25.0%	

Table-3: ICU stay

ICU Stay	Mild		Moderate		Severe		P Value
	N	%	N	%	N	%	
Yes	0	0	3	17.6	6	75	0.002**
No	12	100	14	82.4	2	25	

Table 4: Hospital stay

DOHS	N	Mean	Std. Deviation	p-value
Mild	12	4.42	1.621	<.001
Moderate	17	9.53	4.048	
Severe	8	17.13	5.167	
Total	37	9.51	5.900	

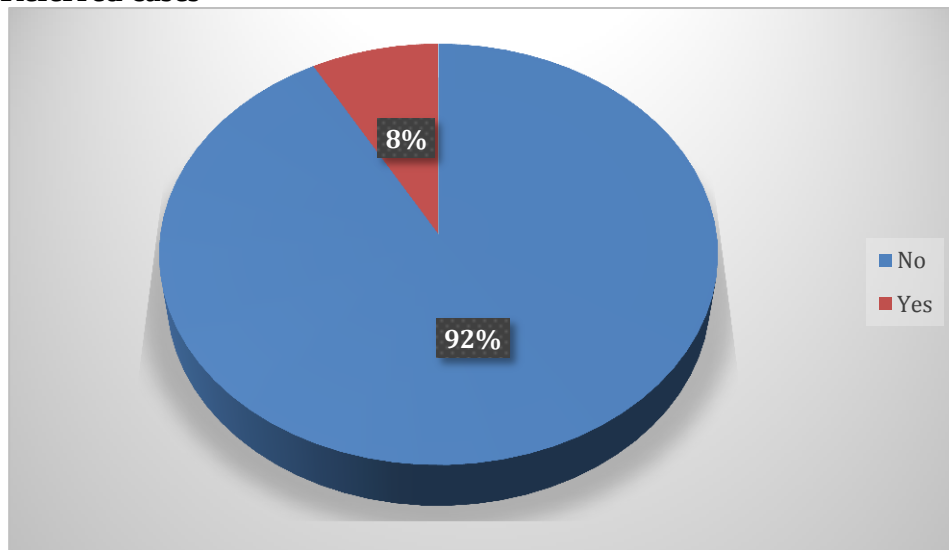
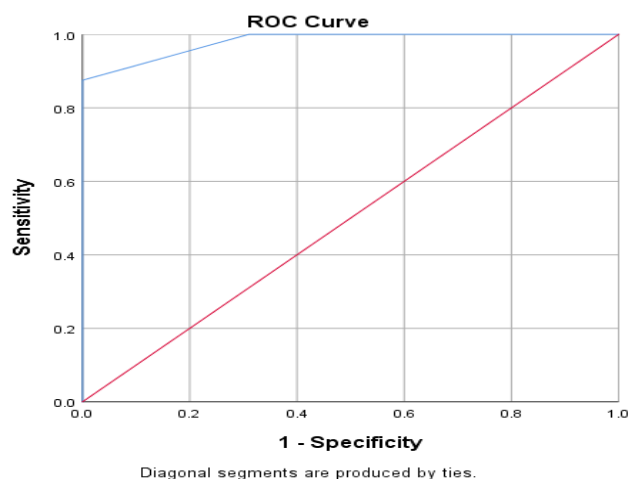
Figure 3: Referred cases

Table-4: Association of BISAP score to MCTSI

BISAP score	Modified CTSI Score						Total		p-value
	Mild		Moderate		Severe				
	n	%	n	%	N	%	N	%	
One	7	58.30%	0	0.00%	0	0.00%	7	18.90%	<0.001
Two	5	41.70%	8	47.10%	0	0.00%	13	35.10%	
Three	0	0.00%	9	52.90%	1	12.50%	10	27.00%	
Four	0	0.00%	0	0.00%	7	87.50%	7	18.90%	
Total	12	100.00%	17	100.00%	8	100.00%	37	100.00%	

ROC curve

Area Under the ROC Curve					
Test Result Variable(s)	Area	Std. Error ^a	Asymptotic Sig. ^b	Asymptotic 95% Confidence Interval	
				Lower Bound	Upper Bound
BISAP score	.978	.026	.000	.927	1.030

b. Null hypothesis: true area = 0.5

The AUC for BISAP score was .978 which was almost similar to MCTSI with statistical significant difference. The results of cut-off value of ≥ 3 are given below:

Results

Statistic	Value
Sensitivity	100.00%
Specificity	68.97%
Positive Predictive Value (*)	47.06%
Negative Predictive Value (*)	100.00%

Discussion

The present single centric, prospective, observational and hospital-based study was conducted at Maharishi Markandeshwar Medical College and hospital, Kumarhatti, Solan, H.P. It included 37 patients with acute pancreatitis and the severity of acute pancreatitis was determined by using BISAP, Modified CTSI score. The mean age of the patients was 48.14 ± 15.99 years. Maximum patients were present in age group 56-70 years (35.1%) followed by 26-40 and 41-55 years with 27% in each group; least patients (8.1%) were ≤ 25 years or > 70 years. The results of the present study were similar to Manjunath B. D. et al⁷ who reported the mean age to be 54.5 years. In a study by Kaushik MR et al⁸ (2017) the mean ($\pm$ SD) age was 43.74 ± 16.85 years.

There was slight male predominance with 51.4% of patients being male and 48.6% being females. In a study by Vaidya et al⁹ 89% of patients were male. Study by Veena P et al¹⁰ in 2020, 90% were males and 10% were females. Males to females ratio was 9:1 in a study by Kaushik et al⁸ in which 82% cases were male and 18% were female. 43% of the patients were males in a study by Katta et al¹⁶. In a study by Arif et al¹¹ 39.3% patients were male while 60.7% patients were female.

The main cause for acute pancreatitis was gall stones (70.3%) followed by alcohol (27%) only 2.7% cases were due to idiopathic cause. In a study by Januja et al¹² most common cause of AP was gallstones in 73 (68.9%), idiopathic in 20 (18.9%) patients and hypertriglyceridemia in 6 (5.7%) patients. According to the study by Chandrashekhar et al¹³ the most common etiology in this study was revealed to be consumption of alcohol which was found in 33(66%) patients, followed by Gall stone disease, which was attributed in 10(20%) patients.

75% (6/8) patients of severe acute pancreatitis developed systemic complications while only 5.9% of moderate presented with systemic complications (ARDS; 1/17). Maximum patients in SAP had ARDS(37.5%) followed by multiorgan failure, paralytic ileus and renal failure (12.5% each). This association was statistically significant ($p < 0.001$). It shows that the rate of cardiac, MODS and ARDS complication was comparable with the studies by chittipotula et al¹⁴, Venkatapuram MR et al¹⁵ and Veena P et al¹⁰. However a study by Katta et al¹⁶ the organ failure was much more than present study. In a study by Manjunath PD On CECT, 14(28%) patients were found to have pancreatic necrosis, 19(38%) were found to have peri pancreatic fluid collection, 26(52%) had ascites and 30(60%) had pleural effusion. In present study 16.2% patients had acute necrotic collection, 24.5% have PP fluid collection, 8.1% each have pseudocyst and WON.

In present study we observed that in mild disease none of the patient needed ICU admission while in moderate disease 3(17.6%) patients needed ICU admission. In case of severe disease 6(75%) patients needed ICU admission. This association was statistically significant ($p < 0.05$)

The results were similar to the study by Januja et al¹² where patients having modified CTSI score of 8 to 10 (severe acute pancreatitis) 4(100%) patients required ICU admission, 4(100%) patients developed complications and mortality in 4(100%) patients occurred. The results were in discordance with a study by Sahu et al¹⁷ who showed similar results, except that they did not find significant association of these scores with the length of ICU stay. This may be partly due to the relatively small number of our patients (19/60) who had a stay in the ICU (the rest of the patients were managed in the ward).

In present study for mild disease hospital stay was 4.42 day for moderate it was 9.53 day while for severe disease the DOHS was 17.13 days. Significant differences were observed among them ($p < 0.001$). The results of the study were in accordance to Sahu et al¹⁷ who showed positive correlation between hospital stay and severity of patients according to MCTSI. Padu et al¹⁸ in their study observed that patients with severe AP had longer duration of hospital stay (mean 20 days) when compared with mild AP (mean stay of 6.56 days). Similar result was found in the study conducted by Gürleyik *et al.*¹⁹ who noted a mean hospital stay of 10.3 days in mild cases and a mean hospital stay of 21.4 days in severe cases, respectively.

The association between MCTSI and BISAP was statistically significant ($p < 0.001$). BISAP score of 3 and 4 was able to detect 100% patients with severe acute pancreatitis thus the cut-off of ≥ 3 can be a good predictor according to present study for predicting SAP.

Chen et al²⁰ compared BISAP with other scoring systems in predicting pancreatic necrosis, organ failure, and mortality in patients with SAP. BISAP was comparable to other scoring systems in predicting the prognoses of SAP. They demonstrated that the best cutoff value for BISAP was 2 for predicting pancreatic necrosis and organ failure, and 3 for predicting mortality. A study by

Qiu, et al.²² indicated that the AUC (0.604), sensitivity (0.905), and specificity (1.000) of BISAP were higher in the prediction of SAP compared to MCTSI.

The advantage of BISAP score is its relative ease with acquiring data and can be calculated within 24 hours of presentation. Patients with a BISAP score of equal to or greater than 4 have high mortality.¹²

Previous studies have well emphasized that the BISAP score is an accurate means for risk-stratification in patients with AP. Its components are clinically relevant and easy to obtain. The prognostic accuracy of BISAP is similar to that of the other scoring systems, and BISAP score would continue to be in clinical practice until any further breakthrough in the field of acute pancreatitis.^{23,24}

Conclusion

BISAP score is effective in predicting severity in patients with acute pancreatitis. It is an accurate tool to classify patients into mild and severe disease; it is easy to perform and can be done on the bedside of patients with acute pancreatitis in every setup.

References

1. Krishna D, Sajjan SS, Sethi BK. Comparative Study Between BISAP Scoring System and C-Reactive Protein Analysis in Predicting Severity of Acute Pancreatitis. *New Indian Journal of Surgery*. 2020;11(1):23-34.
2. Bhatia M, Wong FL, Cao Y, Lau HY, Huang J, Puneet P, et al. Pathophysiology of acute pancreatitis. *Pancreatology*. 2005;5(2-3):132-44.
3. Cho Y-S, Kim H-K, Jang E-C, Yeom J-O, Kim S-Y, Yu J-Y, et al. Usefulness of the Bedside Index for severity in acute pancreatitis in the early prediction of severity and mortality in acute pancreatitis. *Pancreas*. 2013;42(3):483-7.
4. Singh VK, Wu BU, Bollen TL, Repas K, Maurer R, Johannes RS, et al. A prospective evaluation of the bedside index for severity in acute pancreatitis score in assessing mortality and intermediate markers of severity in acute pancreatitis. *Official journal of the American College of Gastroenterology| ACG*. 2009;104(4):966-71.
5. Brown A, Orav J, Banks PA. Hemoconcentration is an early marker for organ failure and necrotizing pancreatitis. *Pancreas*. 2000;20(4):367-72.
6. Khanna AK, Meher S, Prakash S, Tiwary SK, Singh U, Srivastava A, et al. Comparison of Ranson, Glasgow, MOSS, SIRS, BISAP, APACHE-II, CTSI Scores, IL-6, CRP, and procalcitonin in predicting severity, organ failure, pancreatic necrosis, and mortality in acute pancreatitis. *Hpb Surgery*. 2013;2013.
7. Manjunath BD, Ali MA, Razack A, Harindranath HR, Avinash K, Kavya T, et al. Comparison between Ransons score and Modified CTSI in predicting the severity of acute pancreatitis based on modified atlanta classification 2012. *Int Surg J* 2019;6(5):1596-600.
8. Kaushik M, Dubey A, Jain R, Rathore A, Pathak A. Prospective evaluation of the BISAP score and its correlation with Marshall score in predicting severity of organ failure in acute pancreatitis. *Int J Adv Med*. 2017;4(2):534-9.
9. Vaidya Y, Vaithianathan R, Manickam R. Comparative Evaluation of the BISAP Score with CT Severity Index in Predicting the Severity of Acute Pancreatitis. *Ind J Surg*. 2018 Aug;80(4):353-8.
10. Veena LP, Kumar KP. A Prospective study of BISAP score in Acute Pancreatitis. *Narayana Medical Journal*. 2020 Dec; 9(2):9-15.
11. Arif A, Jaleel F, Rashid K. Accuracy of BISAP score in prediction of severe acute pancreatitis. *Pak J Med Sci*. 2019;35(4):1008–1012.
12. Janjua SS, Zaman F, Qamar T. Comparison of Ranson's Score, BISAP, and CTSI in Predicting the Severity of Acute Pancreatitis. *JIMDC*. 2018;7(4):255- 9.

13. Chandrashekar S, Rajput SK. Evaluation of Bisap Scoring System in Predicting Severity and Prognosis of Acute Pancreatitis Patients. IOSR-JDMS 2020 Jan19;1(2): 47-52.
14. Chittipotula B, Ch MB, Rao SS, Teja PA. Study of BISAP Score in Evaluation of Acute Pancreatitis and Its Severity. JMSCR. 2020; 8(1): 1042-1048.
15. Venkatapuram MR, Sateesh S, Batchu D. A prospective study of BISAP score in assessing severity of acute pancreatitis. Int Surg J. 2018;5:1785-91.
16. Katta VR, Kongara R, Prasad A, Kumar SK, Kumar A. Evaluation of BISAP score in predicting severity of acute pancreatitis. Int J of Cur Res.2020;8(3): 28624-28627.
17. Sahu B, Abbey P, Anand R, Kumar A, Tomer S, Malik E. Severity assessment of acute pancreatitis using CT severity index and modified CT severity index: Correlation with clinical outcomes and severity grading as per the Revised Atlanta Classification. Indian J Radiol Imaging. 2017 Apr-Jun;27(2):152-160.
18. Padu G, Lal P, Vindal A. Comparison of Modified Atlanta Classification With Modified CT Severity Index in Acute Gallstone Pancreatitis. MAMC J Med Sci 2019;5(2):63-8
19. Gurleyik G, Emir S, Kiliçoglu G, Arman A, Saglam A. Computed tomography severity index, APACHE II score, and serum CRP concentration for predicting the severity of acute pancreatitis. Jop. 2005 Nov 10;6(6):562-7.
20. Chen L, Lu G, Zhou Q, Zhan Q. Evaluation of the BISAP score in predicting severity and prognosis of acute pancreatitis in Chinese patients. Int Surg. 2013;98:6–12.
21. Aggarwal A, Mathur AV, Verma RK, Gupta M, Raj D. Comparison of BISAP and Ranson's score for predicting severe acute pancreatitis and establish the validity of BISAP score. Int Surg J 2020;7(5): 1-8.
22. Qiu L, Sun RQ, Jia RR, Ma XY, Cheng L, Tang MC, et al. Comparison of existing clinical scoring systems in predicting severity and prognoses of hyperlipidemic acute pancreatitis in Chinese patients: a retrospective study. Medicine. 2015;94(23).
23. Papachristou GI, Muddana V, Yadav D, O'connell M, Sanders MK, Slivka A, et al. Comparison of BISAP, Ranson's, APACHE-II, and CTSI scores in predicting organ failure, complications, and mortality in acute pancreatitis. Official journal of the American College of Gastroenterology| ACG. 2010;105(2):435-41.
24. Mounzer R, Langmead CJ, Wu BU, Evans AC, Bishehsari F, Muddana V et al. Comparison of existing clinical scoring systems to predict persistent organ failure in