

Allegheny Health Network

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BACKGROUND

Endoscopic ultrasound (EUS) transmural par pseudocyst drainage is increasingly being perfe patients with clinical symptoms. Limited data is about readmissions rates of such patients when compared to other procedural techniques. We assess outcomes and unplanned readmission pancreatic pseudocyst drainage based on type procedural intervention.

METHODS

The NRD database was used to identify all patients in 2016 with pancreatic pseudocyst who underwent endoscopic, percutaneous (IR), or surgical drainage. Patient selection was based on ICD-10 CM coding. Inpatient outcomes were calculated for all patients based on procedural type. Bivariate and multivariate logistic regression analysis was performed to identify independent predictors multi-day readmission rates. P value of ≤ 0.05 denotes statistical significance.

RESULTS

Of the 32139 discharges for pancreatic pseudocyst, 2220 patients underwent pseudocyst drainage - 36.2% were endoscopic, 51.6% were percutaneous, and 12.2% were surgically drained. Of these cases 4.23% required unplanned readmission within ≤30 days - 29.8% were endoscopic, 61.7% were percutaneous, and 8.5% were surgical drainage, p = 0.126.

30-day readmission rate for EUS-guided drainage was 3.5% and 5.1% for IR-guided drainage, p=0.096. Regression analysis showed index intervention by EUSguided drainage had decreased risk for unplanned readmission at \leq 60 days (OR 0.639, p= 0.034) and \leq 90 days (OR 0.626, p=0.02) when compared to the IRguided group. Adjusted multivariable regression analysis showed patients with endoscopic pseudocyst drainage (aOR 0.591, p=0.031) had an independent decreased risk of unplanned 30-day readmission, Table 2. Regression analysis showed no statistical significance of inpatient mortality when comparing EUS-guided and IRguided pseudocyst drainage (p= 0.108).

Readmission Rates Following Endoscopic, Percutaneous, and Surgical Pseudocyst Drainage

RESULTS

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			Percutaneous Imaging-guided (IR)	Surgical	Dyalua
		EUS-guided (n=803)	(n=1146)	(n=271)	Pvalue
Age, median (IQR), years		55	57	52	0.001
		(43 – 66)	(45 – 68)	(43 – 64)	
Gender, %	Male	59.9	57.4	57.9	0.542
	Female	40.1	42.6	42.1	
Hospital size, %	Small	4.2	7.7	8.2	<0.001
	Medium	15.5	22.4	25.3	_
	Large	80.3	69.9	66.4	
Weekend admission, %		20.2	20.5	12.0	0.003
Teaching hospital, %		86.2	79.2	79.3	<0.001
Length of stay, median (IQR),		7	9	11	<0.001
days		(4–15)	(5–19)	(6–19)	
Total cost, median (IQR), \$		73 332	77 340	94 281	<0.001
		(40843 – 139944)	(40096 – 162009)	(48098 – 228544)	
In-patient death, %		1.1	2.9	4.1	0.007
Subsequent admission	Endoscopic, %	56.9	29.7	14.3	0.001
procedure	IR, %	28.8	58.4	57.1	_
	Surgical, %	11.5	11.9	28.6	
Readmissions	30-day, n	28	58	8	0.126
	60-day, n	34	74	9	0.030
	90-day, n	38	84	11	0.021

Table 2: Readmission	rates and regression analysis comparing	g endoscopic	versus IR pseudocyst drainage				
Endoscopic-guided Pseudocyst Drainage			IR-guided Pseudocyst Drainage	Р			
30-day readmission, %	3.5		5.1	0.096			
60-day readmission, %	4.2		6.5	0.033			
90-day readmission, %	4.7		7.3	0.017			
	Endoscopic vs IR drainage Bivariable Regression OR (95%CI)	Ρ	Endoscopic vs IR Drainage Multivariable Regression* OR (95%CI)	Р			
30-day readmission	0.675 (0.427 – 1.069)	0.094	0.591 (0.367 – 0.952)	0.031			
60-day readmission	0.639 (0.421 – 0.968)	0.034	0.590 (0.384 – 0.907)	0.016			
90-day readmission	0.626 (0.423 – 0.928)	0.020	0.570 (0.380 – 0.857)	0.007			
In-patient mortality	0.380 (0.181 – 0.799)	0.011	0.533 (0.871 – 4.041)	0.108			
*Adjusted for age, gender, income, comorbidities, hospital size, teaching hospital, primary expected payer							

CONCLUSIONS

Thirty-day readmissions after index hospitalization in patients undergoing EUS-guided pseudocyst drainage were at lower risk when compared to those receiving IR-guided pseudocyst drainage. Endoscopic intervention had decreased risk of unplanned readmission at day 60 and 90 after initial discharge.

EUS therapy was shown to have an associated shorter hospital stay and decreased healthcare cost. Further multicenter RCT will be needed to further examine and validate these findings.