# Frailty in elderly patients with acute colonic diverticulitis is associated with worse in-hospital outcomes: A nationwide analysis

# Waqas Rasheed<sup>a</sup>, Anass Dweik<sup>b</sup>, Gnanashree Dharmarpandi<sup>c</sup>, Aamir Saeed<sup>d</sup>, Amir Humza Sohail<sup>e</sup>, Mohammad Baseem Shaikh<sup>f</sup>, Hassam Ali<sup>g</sup>, Sherif E. Elhanafi<sup>h</sup>

University of Kentucky, Lexington; University Health Sciences Center at Amarillo, Texas; Merit Health Wesley Hospital Hattiesburg, Mississippi; University of New Mexico Health Sciences, Albuquerque; East Carolina University/Brody School of Medicine, Greenville, North Carolina; Texas Tech University Health Sciences Center El Paso, USA

Abstract	<b>Background</b> Frailty has been identified as an independent predictor of mortality in the elderly. We investigated the effects of frailty status on in-hospital outcomes of acute colonic diverticulitis (ACD) in the elderly, using the Hospital Frailty Risk Score.
	<b>Methods</b> We used the National Inpatient Sample (NIS) databases from 2016-2020 to identify patients aged $\geq$ 75 years hospitalized with ACD. Using a 1:1 matching method, we created propensity-matched cohorts of frail (Hospital Frailty Risk Score $\geq$ 5) and non-frail (Hospital Frailty Risk Score $\leq$ 4) patients within the ACD population.
	<b>Results</b> We identified 53.3% ACD patients as frail. We matched 21,720 frail ACD patients to an equal number of non-frail ACD patients using propensity score matching. Frail patients exhibited significantly higher mortality rates, longer hospital stays, and greater median inpatient costs. Frail patients also experienced a greater number of complications, including abscess formation, intestinal perforation, gastrointestinal fistula formation, sepsis without shock, sepsis with shock, acute kidney injury, hypovolemic or hemorrhagic shock, need for blood transfusion, cardiac arrest, and need for intensive care (all P-values <0.001). Additionally, frail patients underwent open colectomy and colostomy procedures more frequently, while laparoscopic colectomies were performed less frequently (all P-values <0.001).
	<b>Conclusions</b> In this nationwide analysis, frailty in ACD is strongly associated with worse mortality, longer hospital stays and higher costs, as well as a greater incidence of local and systemic complications. Furthermore, frailty is linked to a greater need for open colectomy and colostomy procedures.
	Keywords Acute colonic diverticulitis, frailty, elderly, National Inpatient Sample
	Ann Gastroenterol 2024; 37 (XX): 1-7

Conflict of Interest: None

Correspondence to: Waqas Rasheed, MD, Department of Internal Medicine at University of Kentucky, 800 Rose St MN 150, Lexington, KY 40506, USA, e-mail: Wra232@uky.edu

Received 24 February 2024; accepted 15 May 2024; published online 12 July 2024

DOI: https://doi.org/10.20524/aog.2024.0904

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms

# Introduction

Acute colonic diverticulitis (ACD) refers to the inflammation of colonic diverticula, which can occur in approximately 10-25% of individuals with diverticulosis [1]. This condition can lead to various systemic and local complications, including sepsis, pericolonic abscesses formation, fistulas, bowel obstructions, bleeding, and perforations [2]. The risk of developing acute diverticulitis increases with age, ranging from 10% in individuals under 50 years old to 33% in those between 60 and 69 years old [3].

Like ACD, frailty is also believed to be highly prevalent in the elderly [4]. It is defined as a vulnerability to adverse health outcomes secondary to decreased resistance or reserve to stressors, resulting from a decline in the performance of multiple integrated physiological systems, and it is closely related to aging [5,6]. Various methods have been proposed to assess frailty based on objective performance (phenotypic frailty), such as the Fried Frailty Phenotype, or based on comorbidities, disabilities or social factors (deficit accumulation or index frailty) [5,7]. Various screening tools have been developed based on these methods; however, to date, there is no gold standard method of screening for frailty. This poses a major challenge to the development of successful interventions [8,9]. The presence of frailty can potentially affect inpatient outcomes related to conditions that preferentially affect the elderly, and the identification of frailty in these patients can provide an opportunity for early intervention in the population at risk.

Therefore, we investigated the effects of frailty on inhospital outcomes of ACD. For this purpose, we used the Hospital Frailty Risk Score, which is based on the International Classification of Diseases, Tenth Revision (ICD-10) codes and was developed using the Hospital Episode Statistics inpatient database [10]. Our study was performed using the National Inpatient Sample (NIS), the largest national inpatient database in the United States of America (USA).

# **Materials and methods**

#### Data source and study population

The NIS database was searched for hospitalizations related to ACD between the years 2016 and 2020. NIS, recognized as the most extensive all-payer inpatient database in the USA, contains data from 48 states and the District of Columbia, covering over 98 percent of the entire USA population. NIS employs a stratified probability sampling technique, where the stratification is based on multiple factors, such as hospital bed capacity, teaching status, ownership, rural versus urban location, and geographical region. We used the ICD-10-CM classification, as well as procedural codes (ICD-10-PCS), to identify patients aged 75 years or older admitted to hospital with a primary diagnosis of ACD. Within the ACD patient group, further categorization was performed into a frail cohort, if the Hospital Frailty Risk Score was 5 or above, or a nonfrail cohort, if the score was less than 5. This scoring system, developed in 2018 using electronic hospital records, classifies

<sup>a</sup>Internal Medicine, University of Kentucky, Lexington, Kentucky (Waqas Rasheed); <sup>b</sup>Internal Medicine, University of Kentucky, Lexington, Kentucky (Anass Dweik); <sup>c</sup>Internal Medicine, Texas Tech University Health Sciences Center at Amarillo, Texas (Gnanashree Dharmarpandi); <sup>d</sup>Internal Medicine, Merit Health Wesley Hospital Hattiesburg, Mississippi (Aamir Saeed); <sup>c</sup>Department of Surgery, University of New Mexico health Sciences, Albuquerque, New Mexico (Amir Humza Sohail); <sup>f</sup>Internal Medicine, University of Kentucky, Lexington, Kentucky (Mohammad Baseem Shaikh); <sup>g</sup>Division of Gastroenterology, East Carolina University/Brody School of Medicine, Greenville, North Carolina (Hassam Ali); <sup>h</sup>Division of Gastroenterology, Texas Tech University Health Sciences Center El Paso, Texas (Sherif E. Elhanafi), USA patients as having low risk (<5), intermediate risk (5-15), or high risk (>15), using specific ICD-10 codes, with each code assigned a corresponding point value [10]. We used a cutoff point of 5 and above to identify frailty, in order to include both intermediate and high-risk patients in the frail cohort. Patients under 75 years of age were excluded from our analysis, as this scoring system was developed and validated on patients aged 75 years and older [10]. Additional information regarding the sampling methodologies employed by the NIS can be found on the official NIS website [11]. The ICD-10 codes used in our research are listed in Supplementary Table 1. Please refer to the Hospital Frailty Risk Score for a list of ICD-10 codes included in this scoring system [10].

# **Outcomes of interest**

Primary outcomes included a comparison of in-hospital mortality, median length of stay (LOS), and inflationadjusted median inpatient cost between frail and nonfrail patients hospitalized with ACD. Secondary outcomes included a comparison of biodemographic and hospital characteristics; local complications, such as abscess formation, intestinal perforation, intestinal obstruction, lower gastrointestinal bleeding, gastrointestinal fistula formation and paralytic ileus; as well as systemic complications, including sepsis without shock, sepsis with shock, acute kidney injury, hypovolemic/hemorrhagic shock, need for blood transfusion, acute respiratory distress syndrome, disseminated intravascular coagulation, cardiac arrest, and admission to a critical care unit. We also compared the procedures, including drainage procedures, colostomy and colectomy, between the 2 cohorts.

### **Statistical analysis**

The analyses were performed using STATA version 17.0 (StataCorp, College Station, Texas, USA). ACD patients were stratified into frail (Hospital Frailty Risk Score  $\geq$ 5) and nonfrail (Hospital Frailty Risk Score  $\leq 4$ ) cohorts. To mitigate the inherent selection bias in this retrospective study, we decided to perform a propensity-matched analysis. A propensity score was calculated for each hospitalization, based on biodemographic and hospital characteristics, as well as a list of comorbidities detailed in Tables 1 and 2. Frail patients were then matched to non-frail patients using a 1:1 matching method within 0.05 standard deviation of the calculated propensity score, and the covariate balance was analyzed using a covariance plot before and after matching. A receiver operating characteristic (ROC) plot was generated for evaluation of the performance of age combined with frailty, versus age only, in predicting in-hospital mortality (Fig. 1). Matched cohorts were analyzed for primary and secondary outcomes using two-tailed non-parametric tests, including Pearson's chi-squared test for categorical variables and the Wilcoxon rank-sum (Mann-Whitney) test for continuous variables. The categorical variables were reported as 

 Table 1 Biodemographic and hospital characteristics of acute colonic diverticulitis-related hospitalizations in the United States of America in 2016-2020 stratified by frailty status

Characteristics	Acute colonic diverticulitis					
	Unmatched cohorts			Propensity matched cohorts		
Patient and hospital characteristics	Non-frail patients	Frail patients	P-value	Non-frail patients	Frail patients	P-value
Number of hospitalizations (n)	241095 (46.7%)	275630 (53.3%)		21720	21720	
Median age, years (IQR)	81 (78-86)	83 (79-88)	< 0.001	82 (78-87)	82 (78-87)	0.039
Sex Male Female	90580 (37.6%) 150470 (62.4%)	94155 (34.2%) 181445 (65.8%)	<0.001	7910 (36.4%) 13810 (63.6%)	7965 (36.7%) 13755 (63.3%)	0.580
Race White Black Hispanic Asian or Pacific Islander Others	182475 (77.5%) 25835 (11.0%) 17955 (7.6%) 3880 (1.6%) 5420 (2.3%)	208465 (77.4%) 33295 (12.4%) 17645 (6.6%) 4415 (1.6%) 5450 (2.0%)	<0.001	16991 (78.2%) 2495 (11.5%) 1359 (6.3%) 400 (1.8%) 475 (2.2%)	16930 (77.9%) 2570 (11.8%) 1350 (6.2%) 385 (1.8%) 485 (2.2%)	0.800
Median household income, national quartile for patient ZIP Code \$1-\$43,999 \$44,000-\$55,999 \$56,000-\$73,999 \$74,000 or more	62685 (26.3%) 61085 (25.7%) 59750 (25.1%) 54440 (22.9%)	72665 (26.7%) 71195 (26.1%) 67370 (24.7%) 61290 (22.5%)	<0.001	5605 (25.8%) 5676 (26.1%) 5380 (24.8%) 5059 (23.3%)	5535 (25.5%) 5930 (27.3%) 5200 (23.9%) 5055 (23.3%)	0.028
Insurance type Medicare Medicaid Private including HMO Others	224875 (94.1%) 1675 (0.7%) 11645 (4.9%) 835 (0.3%)	258340 (94.6%) 1635 (0.6%) 12355 (4.5%) 835 (0.3%)	<0.001	20410 (94.0%) 135 (0.6%) 1100 (5.1%) 75 (0.3%)	20350 (93.7%) 140 (0.6%) 1140 (5.2%) 90 (0.4%)	0.520
Median Elixhauser Comorbidity Index (IQR)	3 (2-4)	4 (3-6)	< 0.001	3 (2-5)	3 (2-5)	0.884
Median hospital frailty risk score (IQR)	1.8 (0.7-3)	8 (5.6-12.3)	< 0.001	2.3 (1.5-3.2)	7.2 (5.2-11.2)	< 0.001
Hospital size Small Medium Large	58435 (24.2%) 75880 (31.5%) 106780 (44.3%)	64405 (23.4%) 87515 (31.8%) 123710 (44.9%)	<0.001	5495 (25.3%) 6775 (31.2%) 9450 (43.5%)	5485 (25.3%) 6825 (31.4%) 9410 (43.3%)	0.870
Hospital teaching status Non-teaching hospital Teaching hospital	88915 (36.9%) 152180 (63.1%)	96360 (35.0%) 179270 (65.0%)	<0.001	7020 (32.3%) 14700 (67.7%)	6840 (31.5%) 14880 (68.5%)	0.064
Region of hospital Northeast Midwest South West	52660 (21.8%) 51645 (21.4%) 97695 (40.5%) 39095 (16.2%)	53915 (19.6%) 64325 (23.3%) 110700 (40.2%) 46690 (16.9%)	<0.001	4524 (20.8%) 4545 (20.9%) 9031 (41.6%) 3620 (16.7%)	4575 (21.1%) 4635 (21.3%) 8855 (40.8%) 3655 (16.8%)	0.380

SE, standard error; IQR, interquartile range

frequency (N) and percentage (%), while continuous variables were reported as median and interquartile range (IQR). The inpatient cost was adjusted for inflation up to January 2023 using the consumer price index, in order to calculate the inflation-adjusted median inpatient cost [12]. A P-value of 0.05 or less was set as the threshold for statistical significance, and all P-values were 2-sided. The study was exempt from institutional review board approval or patient consent, as the NIS databases contain de-identified patient information and are available publicly. The study findings are reported in accordance with the Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) guidelines [13].

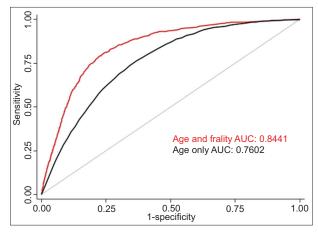
# Results

A total of 516,725 patients were identified and included in our study after application of the inclusion and exclusion criteria; of these, 241,095 (46.7%) were categorized as non-frail 

 Table 2 Elixhauser comorbidities among acute colonic diverticulitis-related hospitalizations in the United States in 2016-2020 stratified by frailty status

Elixhauser comorbidities	Acute colonic diverticulitis						
	Unmatched cohorts			Propensity matched cohorts			
	Non-frail patients	Frail patients	P-value	Non-frail patients	Frail patients	P-value	
Number of hospitalizations (n)	241095	275630		21720	21720		
Congestive heart failure	7385 (18.4%)	14870 (27.8%)	< 0.001	5010 (23.1%)	5055 (23.3%)	0.610	
Cardiac arrhythmias	13875 (34.6%)	22520 (42.2%)	< 0.001	8396 (38.7%)	8395 (38.7%)	0.990	
Valvular disease	4265 (10.6%)	6525 (12.2%)	< 0.001	2625 (12.1%)	2605 (12.0%)	0.770	
Pulmonary circulation disorders	1440 (3.6%)	3600 (6.7%)	< 0.001	1070 (4.9%)	1055 (4.9%)	0.740	
Peripheral vascular disorders	4235 (10.6%)	7260 (13.6%)	< 0.001	2720 (12.5%)	2680 (12.3%)	0.560	
Hypertension, uncomplicated	22200 (55.3%)	21665 (40.6%)	< 0.001	10749 (49.5%)	10585 (48.7%)	0.120	
Hypertension, complicated	55 (0.1%)	360 (0.7%)	< 0.001	50 (0.2%)	55 (0.3%)	0.630	
Paralysis	965 (2.4%)	6050 (11.3%)	< 0.001	895 (4.1%)	925 (4.3%)	0.470	
Other neurological disorders	7935 (19.8%)	13635 (25.5%)	< 0.001	4866 (22.4%)	4920 (22.7%)	0.540	
Chronic pulmonary disease	5920 (14.8%)	5610 (10.5%)	< 0.001	2865 (13.2%)	2850 (13.1%)	0.830	
Diabetes, uncomplicated	4795 (12.0%)	11650 (21.8%)	< 0.001	3520 (16.2%)	3510 (16.2%)	0.900	
Diabetes, complicated	8485 (21.1%)	13215 (24.7%)	< 0.001	4961 (22.8%)	5070 (23.3%)	0.210	
Hypothyroidism	6480 (16.2%)	20305 (38.0%)	< 0.001	5576 (25.7%)	5645 (26.0%)	0.450	
Renal failure	1335 (3.3%)	2385 (4.5%)	< 0.001	840 (3.9%)	810 (3.7%)	0.450	
Liver disease	510 (1.3%)	1010 (1.9%)	< 0.001	385 (1.8%)	355 (1.6%)	0.270	
Peptic ulcer disease excluding bleeding	10 (<1%)	5 (<1%)	0.063	0 (0%)	0 (0%)	-	
AIDS/HIV	380 (0.9%)	765 (1.4%)	< 0.001	225 (1.0%)	240 (1.1%)	0.480	
Lymphoma	675 (1.7%)	1110 (2.1%)	< 0.001	380 (1.7%)	400 (1.8%)	0.470	
Metastatic cancer	1730 (4.3%)	2935 (5.5%)	< 0.001	1045 (4.8%)	1055 (4.9%)	0.820	
Solid tumor without metastasis	1685 (4.2%)	2560 (4.8%)	< 0.001	935 (4.3%)	940 (4.3%)	0.910	
Rheumatoid arthritis/collagen vascular diseases	2780 (6.9%)	5585 (10.5%)	< 0.001	1920 (8.8%)	1865 (8.6%)	0.350	
Coagulopathy	4625 (11.5%)	6665 (12.5%)	< 0.001	2540 (11.7%)	2570 (11.8%)	0.660	
Obesity	1735 (4.3%)	5580 (10.4%)	< 0.001	1350 (6.2%)	1270 (5.8%)	0.110	
Weight loss	4670 (11.6%)	29135 (54.5%)	< 0.001	4414 (20.3%)	4485 (20.6%)	0.400	
Fluid and electrolyte disorders	1260 (3.1%)	1820 (3.4%)	0.024	755 (3.5%)	745 (3.4%)	0.790	
Blood loss anemia	2080 (5.2%)	4035 (7.6%)	< 0.001	1425 (6.6%)	1390 (6.4%)	0.500	
Deficiency anemia	370 (0.9%)	780 (1.5%)	< 0.001	250 (1.2%)	250 (1.2%)	1.000	
Alcohol abuse	175 (0.4%)	305 (0.6%)	0.004	70 (0.3%)	90 (0.4%)	0.110	
Drug abuse	70 (0.2%)	210 (0.4%)	< 0.001	55 (0.3%)	25 (0.1%)	< 0.001	
Psychoses	3340 (8.3%)	7850 (14.7%)	< 0.001	2450 (11.3%)	2380 (11.0%)	0.290	
Depression	10355 (25.8%)	24845 (46.5%)	< 0.001	7716 (35.5%)	7770 (35.8%)	0.590	

and 275,630 (53.3%) as frail patients, as detailed above. Frail patients were older, and presented a higher burden of comorbidities as indicated by their higher Elixhauser Comorbidity Index. The majority of patients in both cohorts were white females, hospitalized in large teaching hospitals in the southern USA (Table 1). Our matching process resulted in no significant differences in biodemographics, hospital characteristics or major comorbidities among the matched cohorts, as indicated by a P-value >0.05 (Table 1 and Table 2). The difference in median hospital frailty risk score between the 2 cohorts remained statistically significant after matching (7.2 vs. 2.3, P<0.001), as the matching process was



**Figure 1** ROC plots for age and frailty combined (red) vs. age only (black) as primary predictors of in-hospital mortality ROC, receiver operating characteristic; AUC, area under the curve

performed after the initial patient stratification. The ROC plot showed a better prediction of in-hospital mortality using age and frailty combined compared to age alone (Fig. 1). Matched frail patients showed significantly greater in-hospital mortality (1.1% vs. 0.2%, P<0.001), median LOS (4 days vs. 3 days, P<0.001) and inflation-adjusted median inpatient cost (10442 vs. 9028 USD, P<0.001) compared to their non-frail counterparts. Frailty in ACD was also significantly associated with higher rates of complications, including abscess formation, intestinal perforation, gastrointestinal fistula formation, sepsis without shock, sepsis with shock, acute kidney injury, hypovolemic or hemorrhagic shock, need for blood transfusion, cardiac arrest, and need for intensive care (all P-values <0.001). However, frail patients experienced less frequent gastrointestinal bleeding compared to non-frail patients.

There were also noteworthy differences between the two cohorts in the need for colectomy and colostomy. The frail ACD cohort exhibited a higher prevalence of open colectomy and colostomy (both P-values <0.001), whereas the rate of laparoscopic colectomy was lower in frail patients compared to their matched counterparts (P<0.001) (Table 3).

# Discussion

Diverticular disorder is a common condition, and its prevalence increases with advancing age. The impact of age is particularly evident, with rates ranging from 10% among individuals under 40 years of age to a staggering 66% among those aged 80 and above [14]. Approximately 10-25% of individuals with diverticular disease will eventually experience ACD [1]. Given the growth of the aging population, it is anticipated that the incidence of ACD will continue to rise [15]. Numerous published studies have investigated the relationship between age and outcomes related to ACD [15-18]. In our research, we investigated the impact of frailty on elderly patients hospitalized with ACD; to the best of our knowledge, no previous studies have investigated this.

Frailty is closely linked to the aging process and is characterized by an increased vulnerability to stress. This vulnerability, in turn, elevates the risks of adverse outcomes such as sickness, falls, hospitalization, delirium and disability, as well as mortality. Therefore, it is a significant concern in terms of susceptibility to poor outcomes [5,6,19]. Frailty does not have a single etiology; rather, it is a product of an imbalance in the intricate interactions between the body's different physiological systems, contributing to compromised homeostasis [5]. The observation that a subset of frail patients lacks significant comorbidities suggests that these imbalances can result either from age-related decline, such as age-related anorexia or loss of muscle mass, or as a consequence of comorbidities [5]. In our effort to contribute to the ongoing discussion, we present a comprehensive 5-year analysis based on the NIS database, shedding light on clinical outcomes among both frail and non-frail patients with ACD. We used the Hospital Frailty Risk Score for our analysis, as, like the NIS database, it uses ICD-10 codes and aligns closely with our methodology.

Based on the Hospital Frailty Risk Score, the prevalence of frailty in our study population was found to be over 50%. The prevalence of frailty varies depending on the assessment method used. While previous studies that included patients older than 65-70 years reported a prevalence between 4% and 16.3%, our patient population exhibited a significantly higher prevalence [20-22]. The patients in our study were older than the patients included in the aforementioned studies, as we specified a minimum age of 75 years as an inclusion criterion. However, the high frailty prevalence of over 50% in our study patients might still suggest that elderly patients with ACD are at particularly higher risk, and should receive more tailored care to prevent adverse inhospital outcomes.

The other notable findings of our study are a higher inhospital mortality, a longer LOS, and a higher inpatient cost associated with frailty status. Additionally, frail patients experienced a higher risk of local and systemic complications of ACD requiring admission to critical care units, and a greater need for inpatient laparoscopic and open colectomy or colostomy procedures. This indicates a greater severity of ACD in frail patients compared to their propensity scorematched non-frail counterparts. When we compared age combined with frailty to age alone, age combined with frailty was better at predicting in-hospital mortality, as indicated by the area under the ROC curve (Fig. 1). Numerous published articles report age as an independent factor for elevated mortality and complications in ACD patients [15-18]. As far as we are aware, no published article currently available investigates frailty as an independent factor within this context.

This study, based on the NIS database, possesses both strengths and limitations. An important strength lies in its large sample size, which adds to the statistical power and generalizability of the study findings. Furthermore, the NIS database provides a comprehensive dataset, including patient demographics, diagnoses and procedures, allowing analysis of various variables in depth. A limitation is that the retrospective

Table 3 Primary and secondary	y outcomes of acute colonic diverticulitis-related hos	spitalizations in the United States in 2016-2020 stratified	by frailty status
-------------------------------	--	---	-------------------

Outcomes	Acute colonic diverticulitis					
	Unmatched cohorts			Propensity matched cohorts		
	Non-frail patients	Frail patients	P-value	Non-frail patients	Frail patients	P-value
Number of hospitalizations	241095 (46.7%)	275630 (53.3%)		21,720	21,720	
In-hospital mortality	765 (0.3%)	5360 (1.9%)	< 0.001	45 (0.2%)	245 (1.1%)	< 0.001
Median LOS, days (IQR)	3 (2-5)	4 (3-7)	< 0.001	3 (2-5)	4 (2-6)	< 0.001
Median inpatient cost, USD (IQR)	8119 (5430-13394)	10508 (6770-18282)	< 0.001	9028 (5991-14868)	10442 (6847-17774)	< 0.001
Local complications Abscess Perforation Bleeding Obstruction Fistula Paralytic ileus	32950 (13.7%) 32845 (13.6%) 121155 (50.3%) 3405 (1.4%) 6480 (2.7%) 280 (0.1%)	41895 (15.2%) 41680 (15.1%) 135325 (49.1%) 4135 (1.5%) 8455 (3.1%) 625 (0.2%)	<0.001 <0.001 <0.001 0.009 <0.001 <0.001	2991 (13.8%) 2971 (13.7%) 12348 (56.9%) 490 (2.3%) 628 (2.9%) 25 (0.1%)	3361 (15.5%) 3336 (15.4%) 11559 (53.2%) 490 (2.3%) 835 (3.8%) 30 (0.1%)	<0.001 <0.001 <0.001 1.000 <0.001 0.500
Systemic complications Sepsis without shock Sepsis with shock Acute kidney injury Hypovolemic/hemorrhagic shock Blood transfusion ARDS DIC Cardiac arrest Admission to critical care unit	$\begin{array}{c} 210 \ (0.1\%) \\ 65 \ (<1\%) \\ 11960 \ (5.0\%) \\ 1375 \ (0.6\%) \\ 33480 \ (13.9\%) \\ 10 \ (<1\%) \\ 35 \ (<1\%) \\ 745 \ (0.3\%) \\ 385 \ (0.2\%) \end{array}$	$\begin{array}{c} 2425\ (0.9\%)\\ 3050\ (1.1\%)\\ 80140\ (29.1\%)\\ 3715\ (1.3\%)\\ 50935\ (18.5\%)\\ 165\ (0.1\%)\\ 245\ (0.1\%)\\ 1790\ (0.6\%)\\ 3565\ (1.3\%)\end{array}$	<0.001 <0.001 <0.001 <0.001 <0.001 <0.001 <0.001 <0.001	$\begin{array}{c} 20 \; (0.1\%) \\ 10 \; (0.05\%) \\ 1340 \; (6.2\%) \\ 135 \; (0.6\%) \\ 3511 \; (16.2\%) \\ 0 \; (0.0\%) \\ 0 \; (0.0\%) \\ 70 \; (0.3\%) \\ 15 \; (0.1\%) \end{array}$	$150 (0.7\%) \\ 165 (0.8\%) \\ 5580 (25.7\%) \\ 210 (1.0\%) \\ 3826 (17.6\%) \\ 10 (<1\%) \\ 5 (<1\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 120 (0.6\%) \\ 100 $	<0.001 <0.001 <0.001 <0.001 0.002 0.025 <0.001 <0.001
Colectomy Laparoscopic Robotic Open Colostomy	33190 (13.8%) 585 (0.2%) 13980 (5.8%) 110 (<1%)	28655 (10.4%) 280 (0.1%) 19415 (7.0%) 265 (0.1%)	<0.001 <0.001 <0.001 <0.001	3432 (15.8%) 45 (0.2%) 1162 (5.3%) 60 (0.2%)	2714 (12.5%) 30 (0.1%) 1263 (5.8%) 85 (0.4%)	<0.001 0.083 0.035 0.038

LOS, length of stay; IQR, interquartile range; ARDS, acute respiratory distress syndrome; DIC, disseminated intravascular coagulation

nature of the study may have introduced selection bias and limited the ability to establish causal relationships. However, the propensity-matched technique was used to eliminate these biases and unmeasured confounders. Additionally, the reliance on administrative codes for diagnoses and procedures within the NIS database could potentially lead to inaccuracies and misclassifications [23]. NIS also lacks detailed clinical information regarding vital signs, laboratory results and longitudinal follow up [24]. Despite these limitations, the study offers valuable insights into the subject matter, which can be further explored and substantiated through additional research.

In summary, this study found that frailty contributes independently to adverse inpatient outcomes related to ACD in the elderly, including higher mortality rates, prolonged hospital stays, greater healthcare costs, and higher complication rates. While age remains an important factor, frailty emerges as a distinct and substantial determinant of patient outcomes. These findings underscore the importance of factoring in frailty when clinically managing elderly ACD patients, and offer valuable insights for future research and interventions aimed at improving the care and outcomes of this vulnerable population.

#### **Summary Box**

# What is already known:

- Acute colonic diverticulitis is a common condition, with significant healthcare resource utilization
- Acute colonic diverticulitis preferentially affects
   older patients
- Frailty is known to independently predict mortality in the elderly

# What the new findings are:

- Frailty is highly prevalent in elderly patients with acute colonic diverticulitis
- Frail patients with acute colonic diverticulitis had higher in-hospital mortality and healthcare resource utilization, and greater local and systemic complications
- In acute colonic diverticulitis, frailty combined with age was found to be a better predictor of mortality compared to age alone

### References

- Lanas A, Abad-Baroja D, Lanas-Gimeno A. Progress and challenges in the management of diverticular disease: which treatment? *Therap Adv Gastroenterol* 2018;11:1756284818789055.
- Fugazzola P, Ceresoli M, Coccolini F, et al. The WSES/SICG/ACOI/ SICUT/ACEMC/SIFIPAC guidelines for diagnosis and treatment of acute left colonic diverticulitis in the elderly. *World J Emerg Surg* 2022;17:5.
- 3. Ubaldi E, Grattagliano I, Lapi F, Pecchioli S, Cricelli C. Overview on the management of diverticular disease by Italian General Practitioners. *Dig Liver Dis* 2019;**51**:63-67.
- Collard RM, Boter H, Schoevers RA, Oude Voshaar RC. Prevalence of frailty in community-dwelling older persons: a systematic review. J Am Geriatr Soc 2012;60:1487-1492.
- Fried LP, Tangen CM, Walston J, et al; Cardiovascular Health Study Collaborative Research Group. Frailty in older adults: evidence for a phenotype. J Gerontol A Biol Sci Med Sci 2001;56:M146-M156.
- 6. Clegg A, Young J, Iliffe S, Rikkert MO, Rockwood K. Frailty in elderly people. *Lancet* 2013;**381**:752-762.
- Rockwood K, Mitnitski A. Frailty in relation to the accumulation of deficits. J Gerontol A Biol Sci Med Sci 2007;62:722-727.
- Rockwood K, Stadnyk K, MacKnight C, McDowell I, Hébert R, Hogan DB. A brief clinical instrument to classify frailty in elderly people. *Lancet* 1999;353:205-206.
- Winograd CH. Targeting strategies: an overview of criteria and outcomes. J Am Geriatr Soc 1991;39:25S-35S.
- Gilbert T, Neuburger J, Kraindler J, et al. Development and validation of a Hospital Frailty Risk Score focusing on older people in acute care settings using electronic hospital records: an observational study. *Lancet* 2018;**391**:1775-1782.
- 11. Agency for Healthcare Research and Quality. H-CUP User Support. Available from: https://hcup-us.ahrq.gov/db/nation/nis/ nisdbdocumentation.jsp [Accessed 1 July 2024].
- U.S. Bureau of Labor Statistics. CPI Home. Available from: https:// www.bls.gov/cpi/[Accessed 1 July 2024].
- 13. STROBE Checklists. Available from: https://www.strobe-

statement.org/checklists/[Accessed 1 July 2024].

- Peery AF, Keku TO, Martin CF, et al. Distribution and characteristics of colonic diverticula in a United States screening population. *Clin Gastroenterol Hepatol* 2016;14:980-985.
- 15. Covino M, Rosa F, Ojetti V, et al. Acute diverticulitis in elderly patients: does age really matter? *Dig Dis* 2021;**39**:33-41.
- Lidsky ME, Thacker JK, Lagoo-Deenadayalan SA, Scarborough JE. Advanced age is an independent predictor for increased morbidity and mortality after emergent surgery for diverticulitis. *Surgery* 2012;**152**:465-472.
- Ünlü, van de Wall BJ, Gerhards MF, et al. Influence of age on clinical outcome of acute diverticulitis. J Gastrointest Surg 2013;17:1651-1656.
- Horesh N, Shwaartz C, Amiel I, et al. Diverticulitis: does age matter? J Dig Dis 2016;17:313-318.
- 19. Walston J, Hadley EC, Ferrucci L, et al. Research agenda for frailty in older adults: toward a better understanding of physiology and etiology: summary from the American Geriatrics Society/National Institute on Aging Research Conference on Frailty in Older Adults. *J Am Geriatr Soc* 2006;54:991-1001.
- 20. Cawthon PM, Marshall LM, Michael Y, et al; Osteoporotic Fractures in Men Research Group. Frailty in older men: prevalence, progression, and relationship with mortality. *J Am Geriatr Soc* 2007;**55**:1216-1223.
- Woods NF, LaCroix AZ, Gray SL, et al; Women's Health Initiative. Frailty: emergence and consequences in women aged 65 and older in the Women's Health Initiative Observational Study. *J Am Geriatr* Soc 2005;53:1321-1330.
- 22. Kiely DK, Cupples LA, Lipsitz LA. Validation and comparison of two frailty indexes: the MOBILIZE Boston Study. *J Am Geriatr Soc* 2009;**57**:1532-1539.
- O'Malley KJ, Cook KF, Price MD, Wildes KR, Hurdle JF, Ashton CM. Measuring diagnoses: ICD code accuracy. *Health Serv Res* 2005;40:1620-1639.
- 24. Khera R, Krumholz HM. With great power comes great responsibility: "Big Data" research from the National Inpatient Sample. *Circ Cardiovasc Qual Outcomes* 2017;**10**:e003846.

# Supplementary material

Acute colonic diverticulitis	K5700 K5701 K5710 K5711 K5712 K5713 K5720 K5721 K5730 K5731 K5732 K5733 K5740 K5741 K5750 K5751 K5752 K5753 K5780 K5781 K5790 K5791 K5792 K5793
Abscess	K651 K6811 K6812 K6819 K51514 K630 K6811 K6819 K5720 K5721 K5740 K5741 K5780 K5781
Intestinal perforation	K631 P780 K5720 K5721 K5740 K5741 K5780 K5781
Diverticulitis with bleeding	K5721 K5731 K5733 K5741 K5751 K5753 K5781 K5791 K5793
Intestinal obstruction	K50012 K50112 K50812 K50912 K51012 K51212 K51312 K51412 K51512 K51812 K51912 K56600 K56601 K56609 K56690 K56691 K56699 K9130 K9131 K9132
Gastrointestinal fistula	K51913 K603 K604 K605 K632 N321 N822 N823 N824
Paralytic ileus	K560
Sepsis without shock	R6520
Sepsis with shock	R6521
Acute kidney injury	N170 N171 N172 N178 N179
Hypovolemic/hemorrhagic shock	R571
Blood transfusion	30230H1 30230N1 30233N1 30240H1 30240N1 30243H1 30243N1
ARDS	J80
DIC	D65
Cardiac arrest	I462 I468 I469 I97120 I97121 Z8674
Percutaneous drainage	0D9E30Z 0D9E3ZX 0D9E3ZZ 0D9F30Z 0D9F3ZX 0D9F3ZZ 0D9G30Z 0D9G3ZX 0D9G3ZZ 0D9H30Z 0D9H3ZX 0D9H3ZZ 0D9K30Z 0D9K3ZX 0D9K3ZZ 0D9L30Z 0D9L3ZX 0D9L3ZZ 0D9M30Z 0D9M3ZX 0D9M3ZZ 0D9N30Z 0D9N3ZX 0D9N3ZZ 8E0WXBG 8E0WXBH
Laparoscopic drainage	0D9E40Z 0D9E4ZX 0D9E4ZZ 0D9E80Z 0D9E8ZX 0D9E8ZZ 0D9F40Z 0D9F4ZX 0D9F4ZZ 0D9F80Z 0D9F8ZX 0D9F8ZZ 0D9G40Z 0D9G4ZX 0D9G4ZZ 0D9G80Z 0D9G8ZX 0D9G8ZZ 0D9H40Z 0D9H4ZX 0D9H4ZZ 0D9H80Z 0D9H8ZX 0D9H8ZZ 0D9K40Z 0D9K4ZX 0D9K4ZZ 0D9K80Z 0D9K8ZX 0D9K8ZZ 0D9L40Z 0D9L4ZX 0D9L4ZZ 0D9L80Z 0D9L8ZX 0D9L8ZZ 0D9M40Z 0D9M4ZX 0D9M4ZZ 0D9M80Z 0D9M8ZX 0D9M8ZZ 0D9N40Z 0D9N4ZX 0D9N4ZZ 0D9N80Z 0D9N8ZZ
Open drainage	0D9E0ZX 0D9E0ZZ 0D9F0ZX 0D9F0ZZ 0D9G0ZX 0D9G0ZZ 0D9H0ZX 0D9H0ZZ 0D9K0ZX 0D9K0ZZ 0D9L0ZX 0D9L0ZZ 0D9M0ZX 0D9M0ZZ 0D9N0ZX 0D9N0ZZ
Colostomy	0D1E074 0D1E0J4 0D1E0Z4 0D1E474 0D1E4J4 0D1E4K4 0D1E4Z4 0D1E874 0D1E8J4 0D1E8K4 0D1E8Z4
Laparoscopic colectomy	0DBE4ZX 0DBE4ZZ 0DBE8ZX 0DBE8ZZ 0DBF4ZX 0DBF4ZZ 0DBF8ZX 0DBF8ZZ 0DBG4ZX 0DBG4ZZ 0DBG8ZX 0DBG8ZZ 0DBGFZZ 0DBH4ZX 0DBH4ZZ 0DBH8ZX 0DBH8ZZ 0DBK4ZX 0DBK4ZZ 0DBK8ZZ 0DBH4ZZ 0DBL4ZX 0DBL4ZZ 0DBL8ZX 0DBL8ZZ 0DBLFZZ 0DBM4ZX 0DBM4ZZ 0DBM8ZZ 0DBM8ZZ 0DBMFZZ 0DBN4ZX 0DBN4ZZ 0DBN8ZX 0DBN8ZZ 0DBNFZZ 0DTE8ZZ 0DTE4ZZ 0DTF4ZZ 0DTF8ZZ 0DTG4ZZ 0DTG8ZZ 0DTGFZZ 0DTH4ZZ 0DTH8ZZ 0DTK4ZZ 0DTK8ZZ 0DTL4ZZ 0DTL8ZZ 0DTLFZZ 0DTM4ZZ 0DTM8ZZ 0DTMFZZ 0DTN4ZZ 0DTN8ZZ 0DTNFZZ
Robotic colectomy	8E0W0CZ 8E0W3CZ 8E0W7CZ 8E0WXBZ 8E0WXCZ
Open colectomy	0DBE0ZX 0DBE0ZZ 0DBF0ZX 0DBF0ZZ 0DBG0ZX 0DBG0ZZ 0DBH0ZX 0DBH0ZZ 0DBK0ZX 0DBK0ZZ 0DBL0ZX 0DBL0ZZ 0DBM0ZX 0DBM0ZZ 0DBN0ZX 0DBN0ZZ 0DTE0ZZ 0DTF0ZZ 0DTG0ZZ 0DTH0ZZ 0DTK0ZZ 0DTL0ZZ 0DTM0ZZ 0DTN0ZZ