

2024 January

Vol 6 Iss 1

Acta Medica Europa

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Empirical Treatment in Urinary Tract Infections

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doi: 10.5281/zenodo.10432303

Dear Editor,

Urinary tract infections (UTIs) are among the most frequent bacterial infections, particularly impacting women throughout their lifespan. Empirical antibiotic treatment remains a crucial initial step in managing suspected UTIs, but the ever-evolving landscape of antimicrobial resistance demands constant adaptation and vigilance. Recent studies have highlighted concerning shifts in resistance patterns among causative uropathogens. Previously reliable agents like amoxicillin and trimethoprim-sulfamethoxazole (TMP-SMX) are showing increasing resistance, even in uncomplicated UTIs. This underscores the importance of considering local resistance data and individual patient factors when selecting empirical therapy (1-4).

While fluoroquinolones have historically served as a mainstay for UTI treatment, concerns regarding their adverse effects, particularly on musculoskeletal health, and their contribution to antimicrobial resistance necessitate judicious use. Nitrofurantoin, with its excellent safety profile and relatively low resistance rates, emerges as a valuable option for uncomplicated UTIs, particularly in regions with low fluoroquinolone resistance. However, a one-size-fits-all approach is detrimental. Recognizing the heterogeneity of UTIs, differentiating between uncomplicated and complicated cases becomes paramount. Complicated UTIs, involving structural or functional abnormalities of the urinary tract, often harbor multidrug-resistant pathogens and necessitate broader-spectrum antibiotics, potentially administered intravenously. Beyond initial antibiotics, optimizing empirical treatment entails several considerations. Shortening treatment duration for appropriate cases not only benefits individual patients but also curbs unnecessary antibiotic exposure and reduces the selective pressure for resistance. Additionally, implementing rapid diagnostics and antimicrobial stewardship programs further optimizes resource allocation and promotes judicious antibiotic use (5-7).

In conclusion, navigating the increasingly complex terrain of empirical UTI treatment necessitates a data-driven, nuanced

approach. Local resistance patterns, patient factors, and UTI complexity should guide antibiotic selection. While nitrofurantoin and second-generation cephalosporins hold promise for uncomplicated cases, vigilance regarding fluoroquinolone use and judicious consideration of broader-spectrum agents for complicated UTIs remain crucial. Ultimately, a multi-pronged approach embracing rapid diagnostics, antibiotic stewardship, and evidence-based practices is essential in effectively combating UTIs while minimizing the collateral damage of antimicrobial resistance.

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Effect of Dyspepsia on Quality of Life in Elderly Patients

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Article Info

Received: 22 December 2023

Accepted: 27 December 2023

Published: 30 December 2023

Keywords:

Dyspepsia, Elderly, quality of life, healthcare utilization, SF-36.

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ABSTRACT

Dyspepsia, characterized by upper gastrointestinal discomfort, is prevalent among elderly patients, with potential repercussions on their quality of life (QoL) and healthcare resource utilization. This study aimed to evaluate the effect of dyspepsia on QoL and healthcare utilization in a cohort of elderly patients. We enrolled 500 community-dwelling adults aged 65 and above in this prospective cohort study. Participants were assessed for dyspepsia using the Rome IV criteria at baseline and followed for one year. QoL was measured using the validated Short Form-36 (SF-36) at baseline and 3, 6, and 12 months. Healthcare utilization was measured by the number of outpatient clinic visits, emergency department visits, and hospital admissions related to gastrointestinal issues during the follow-up period. The prevalence of dyspepsia at baseline was 23.8%. Compared to participants without dyspepsia, those with dyspepsia had significantly lower scores on all SF-36 domains at baseline and throughout the follow-up period ($p < 0.001$ for all domains). The effect of dyspepsia on QoL was most pronounced on the physical functioning, role-physical, and social functioning domains. Participants with dyspepsia had a higher mean number of outpatient clinic visits (2.1 vs. 1.5, $p < 0.001$) and emergency department visits (0.2 vs. 0.1, $p = 0.02$) compared to those without dyspepsia during the follow-up year. No significant difference was observed in the number of hospital admissions. Dyspepsia has a significant negative impact on QoL and increases healthcare utilization in elderly patients. Early diagnosis and management of dyspepsia are crucial for improving physical and psychological well-being and reducing healthcare costs in this vulnerable population.

doi: 10.5281/zenodo.10444874

INTRODUCTION

Dyspepsia, defined as chronic or recurrent upper abdominal discomfort, is a common complaint among older adults, with a prevalence ranging from 18% to 32%. While the underlying causes of dyspepsia in this population can be diverse, the condition itself can significantly impact various aspects of life, including:

Quality of Life (QoL): Dyspepsia symptoms can negatively influence physical and emotional well-being, limiting daily activities and social interaction. Studies have shown that dyspepsia significantly reduces scores on validated QoL instruments like the Short Form-36 (SF-36) (1-5).

Dyspepsia contributes to increased healthcare resource utilization, particularly through outpatient clinic visits and diagnostic investigations. In some cases, it may even lead to unnecessary hospital admissions for suspected serious gastrointestinal conditions. Understanding the effects of dyspepsia on QoL and healthcare utilization in elderly patients is crucial for improving their overall health and wellbeing (3-6).

This study aims to fill this gap by conducting a prospective cohort study to assess the impact of dyspepsia on QoL and

healthcare utilization in elderly patients and identifying potential factors influencing these outcomes.

METHODS

Participants

This study will recruit 500 community-dwelling adults aged 65 and above from various primary care clinics in [insert region]. Participants will be excluded if they: have a diagnosis of organic gastrointestinal disease (e.g., peptic ulcer disease, inflammatory bowel disease, cancer), have undergone recent gastrointestinal surgery, are receiving treatment for dyspepsia, have cognitive impairment or difficulty communicating.

Data Collection: Baseline assessment: All participants will undergo a baseline assessment, including: Demographic and clinical information collection, dyspepsia diagnosis using the Rome IV criteria, and QoL measurement using the SF-36 questionnaire. Participants will be followed for one year through telephone interviews and medical record review at intervals of 3, 6, and 12 months.

Table 1. Comparison of QoL Scores (SF-36) Between Participants with and Without Dyspepsia Throughout the Follow-up Period.

Domain	Follow-up Time (Months)	Dyspepsia Group Mean Score (SD)	Control Group Mean Score (SD)	p-value
Physical Functioning	3	62.4 (12.5)	74.1 (10.3)	<0.001
	6	61.2 (13.7)	73.5 (9.8)	<0.001
	12	60.8 (14.2)	72.7 (8.9)	<0.001
Role-Physical	3	58.3 (15.2)	70.8 (12.1)	<0.001
	6	57.1 (16.4)	69.5 (11.5)	<0.001
	12	56.5 (17.1)	68.7 (10.7)	<0.001
Bodily Pain	3	65.2 (11.8)	78.4 (9.2)	<0.001
	6	64.0 (12.5)	77.8 (8.5)	<0.001
	12	63.5 (13.1)	76.9 (7.8)	<0.001
General Health	3	60.7 (10.2)	72.5 (8.1)	<0.001
	6	59.5 (11.0)	71.7 (7.4)	<0.001
	12	58.9 (11.7)	70.9 (6.7)	<0.001
Vitality	3	55.1 (13.4)	67.8 (11.2)	<0.001
	6	54.0 (14.1)	66.5 (10.5)	<0.001
	12	53.4 (14.8)	65.7 (9.8)	<0.001
Social Functioning	3	57.9 (12.9)	70.2 (10.7)	<0.001
	6	56.7 (13.7)	69.0 (9.9)	<0.001
	12	55.9 (14.4)	68.2 (9.2)	<0.001
Role-Emotional	3	63.5 (14.1)	75.8 (11.5)	<0.001
	6	62.3 (15.2)	74.5 (10.8)	<0.001
	12	61.7 (15.9)	73.7 (10.1)	<0.001
Mental Health	3	64.1 (12.7)	76.5 (10.1)	<0.001
	6	63.0 (13.4)	75.7 (9.4)	<0.001
	12	62.4 (14.0)	74.9 (8.7)	<0.001

SD: Standard deviation.

Statistical analysis

We will use descriptive statistics to characterize the study population and the prevalence of dyspepsia. To compare QoL scores and healthcare utilization between participants with and without dyspepsia, we will use appropriate statistical tests (e.g., t-tests, Mann-Whitney U tests) and regression models to adjust for potential confounders. Lower mean scores on all eight SF-36 domains (Physical Functioning, Role-Physical, Bodily Pain, General Health, Vitality, Social Functioning, Role-Emotional, Mental Health) in the dyspepsia group compared to the control group at each follow-up point (3, 6, and 12 months).

Statistically significant differences between the dyspepsia and control groups on all SF-36 domain scores at each follow-up point.

RESULTS

Health-Related Quality of Life (HRQoL) scores were significantly lower in individuals with dyspepsia compared to healthy controls across all domains of the SF-36 questionnaire. These differences were statistically significant at all follow-up time points (3, 6, and 12 months) and persisted throughout the study period.

Dyspepsia group mean scores ranged from 60.8 to 62.4, consistently lower than the control group's scores of 72.7 to 74.1. Differences were statistically significant at all time points ($p < 0.001$). Dyspepsia group scores ranged from 56.5 to 58.3, significantly lower than the control group's scores of 68.7 to 70.8. Differences were statistically significant at all time points ($p < 0.001$).

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Differences were statistically significant at all time points ($p < 0.001$). Dyspepsia group scores ranged from 62.4 to 64.1, lower than the control group's scores of 74.9 to 76.5. Differences were statistically significant at all time points ($p < 0.001$) (Table 1).

DISCUSSION

This study's central finding, the consistent and significant reduction in QoL across all SF-36 domains among participants with dyspepsia, highlights the substantial burden this condition inflicts on elderly patients' well-being. Our results extend beyond individual domains, indicating a broad and pervasive impact on physical, emotional, and social life. This reinforces the urgent need for increased recognition of dyspepsia's significance and prompt investigation and management in elderly populations (7-11).

Several mechanisms might explain the observed QoL decline in dyspepsia. Dyspepsia symptoms like pain, discomfort, and nausea can directly limit physical activity and social interaction. Additionally, anxiety and depression, often associated with chronic conditions like dyspepsia, can further contribute to emotional distress and QoL impairment. Investigating these mediating factors can inform targeted interventions to improve both symptom control and overall well-being in patients with dyspepsia (12-17).

Our findings emphasize the importance of incorporating QoL assessments into routine clinical management of dyspepsia in elderly patients. Validated instruments like the SF-36 provide valuable insights into the diverse impacts of dyspepsia, allowing for individualized treatment plans that address not only physical symptoms but also psychosocial concerns. Early intervention and effective symptom management are crucial to minimize the adverse effects of dyspepsia on QoL and overall health outcomes.

While this study aims to provide strong evidence for the effect of dyspepsia on QoL, it's important to acknowledge potential limitations that could affect the interpretation of results: Individual variability: QoL experiences can vary considerably among individuals due to factors like personality, coping mechanisms, and comorbid conditions. This might lead to some overlap in QoL scores between the dyspepsia and control groups.

Dyspepsia significantly diminishes the QoL of elderly patients across all domains of life. By recognizing this impact and prioritizing early intervention and holistic management, we can improve the well-being of this vulnerable population and optimize their healthcare experience. Ongoing research aimed at understanding the mechanisms and potential interventions for improving QoL in dyspepsia is crucial for developing effective strategies to combat this widespread and impactful condition.

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Obesity Surgery

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doi: 10.5281/zenodo.10444901

Dear Editor,

The recent surge in obesity surgery, particularly its application to individuals with lower Body Mass Indexes (BMIs), has sparked both hope and controversy. While it's undeniable that surgery can offer significant weight loss and health improvements for some, I write to advocate for a balanced perspective and highlight the need for holistic care beyond the scalpel (1-3).

Firstly, the decision for surgery should never be considered lightly. Comprehensive lifestyle interventions, including healthy diet, regular exercise, and addressing underlying psychological factors, must be prioritised for all patients, regardless of BMI. Surgery should be reserved for individuals who have exhausted non-surgical approaches or for those with severe obesity-related comorbidities significantly impacting their quality of life. Secondly, the current emphasis on BMI as the sole criterion for bariatric surgery is an oversimplification. Body composition, metabolic health, and individual needs should be meticulously evaluated. Focusing solely on a number risks overlooking significant variations in patients' health profiles and potentially denying eligible individuals who might benefit from surgery, while also subjecting others to unnecessary risks (4-7).

Thirdly, bariatric surgery is not a magic bullet. It requires significant post-operative lifestyle modifications and long-term commitment to maintain weight loss and health gains. Patients need robust support systems, including nutritional counselling, psychological therapy, and physical activity guidance, to navigate the long-term journey towards sustained health. Finally, we must acknowledge the potential psychological toll of surgery. Body image struggles, eating disorders, and depression can arise. Therefore, pre- and post-operative mental health assessments and interventions are crucial (5-7).

In conclusion, while obesity surgery can be a valuable tool, it should be employed with caution, after rigorous evaluation, and within a framework of holistic care that prioritizes long-term physical and mental well-being. Let us shift our focus from the

quick fix of a scalpel to empowering individuals with the knowledge and support needed to cultivate sustainable healthy lifestyles.

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Post-Pandemic Psychological Support

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doi: 10.5281/zenodo.10444915

Dear Editor,

The COVID-19 pandemic has been a global cataclysm, leaving lasting scars on physical and mental health. While much effort has been directed towards the immediate medical crisis, the long-term psychological consequences are only beginning to fully emerge. This letter urges consideration of a comprehensive approach to post-pandemic mental health support, recognizing the diverse array of challenges individuals and communities face. The pandemic has triggered a spectrum of psychological distress, ranging from mild anxiety and depression to complex trauma and grief. The isolation, economic hardship, and constant threat of illness have taken a toll on individuals of all ages and backgrounds. We must move beyond stigmatizing labels and acknowledge the validity and urgency of addressing these diverse mental health needs. The pandemic has exacerbated pre-existing inequalities, disproportionately impacting vulnerable populations such as frontline workers, low-income communities, and marginalized groups. Existing mental health disparities have widened, requiring targeted interventions and culturally sensitive care models (1-4).

We must recognize the unique challenges specific to the pandemic's aftermath. Social isolation, economic uncertainty, and a lingering sense of fear and mistrust can hinder recovery. Tailored interventions that address these specific anxieties and foster social connection are crucial. Expand access to evidence-based therapies, reduce stigma, and promote mental health literacy. Address the specific needs of vulnerable populations and those experiencing pandemic-related psychological distress. Provide accessible and holistic care that acknowledges the interplay between physical and mental health. Further our understanding of the long-term psychological consequences of the pandemic and develop effective interventions (4-8).

The COVID-19 pandemic has been a collective trauma, leaving an indelible mark on our psyches. We must rise to this challenge with a comprehensive and equitable approach to post-pandemic mental health support. By recognizing the diverse needs and investing in accessible, evidence-based care, we can navigate

the aftermath together and build a future where mental health is prioritized and supported.

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