



İNTRAABDOMİNAL ENFEKSİYONLARA BAKIŞ

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Sunum Planı

- Olgu Teorik/Pratik tanı ve tedavi yaklaşımı
- İntraabdominal infeksiyon riski
Lokalizasyon ? Reoperasyon ?
- Tedavi - Hangi antibiyotikler ?
Ne zaman ve nasıl bir cerrahi girişim
- Ülkemizde İntra-abdominal infeksiyonlar

Cerrah ve İnfeksiyoncu gözüyle...



Olgu

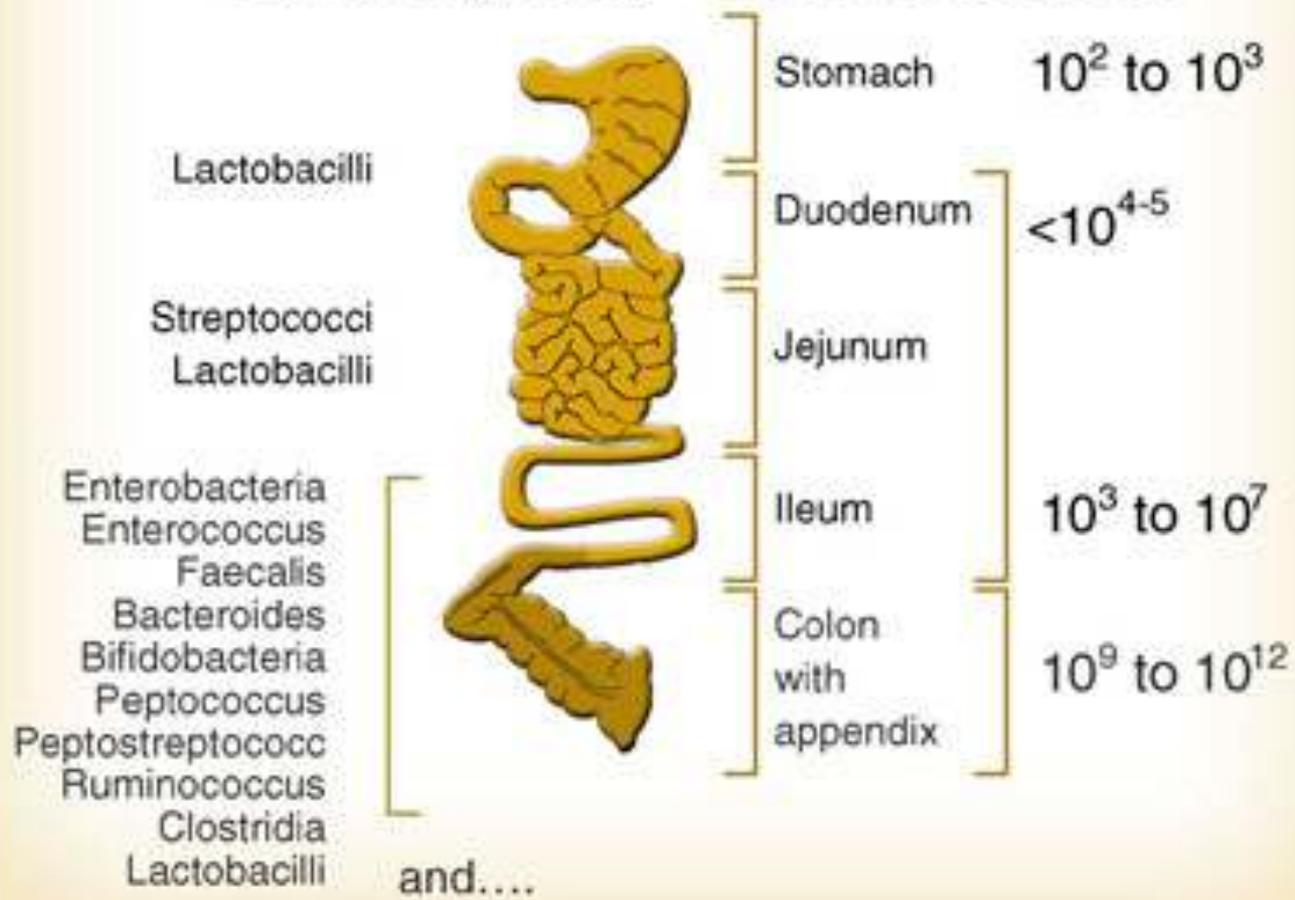
Teorik/Pratik tanı ve tedavi yaklaşımı



Intraabdominal infeksiyon riski
Lokalizasyon ?
Reoperasyon ?

INTESTINAL MICROFLORA

10^{14} micro-organisms, >500 differentes species



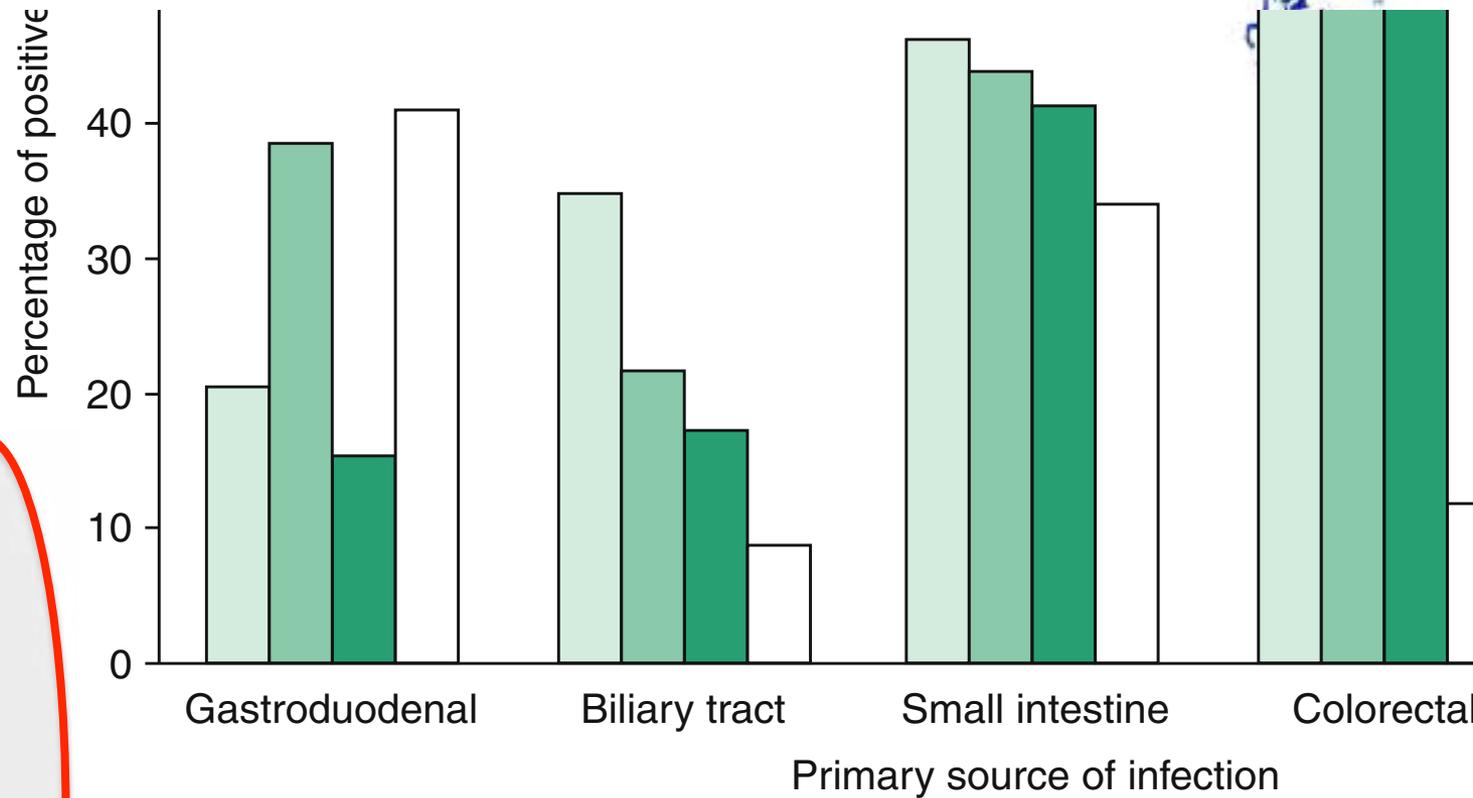


Fig. 1. Initial culture results in secondary and tertiary peritonitis: percentages of positive culture results for different primary sources of infection.^[19]

Konak özellikleri ? Hastalığın ciddiyeti



- >24 saat ilk müdahalenin gecikmesi
- APACHE II skoru ≥ 15 olması
- İleri yaş
- Ek hastalık varlığı ve organ yetmezliği
- Düşük albümin seviyesi
- Kötü beslenme
- Diffüz peritonit
- Kaynak kontrolünün başarısı ?
- Malignite varlığı

PUBMED ?



- Colorectal Cancer ± Reoperation ± infection
- Surgical site infection ± Reoperation/relaparotomy
- Wound infection ± Reoperation ± colorectal cancer
- Intraabdominal infection/sepsis ± reoperation
- Intraabdominal abscess ± cancer ± reoperation

Original research

Relaparotomy in colorectal cancer surgery – Do any factors influence the risk of mortality? A case controlled study[☆]

Michał Adamczyk, Radziszewski, Tarcinski

Department

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- Reope
- Anasto

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Particularly male gender, older age, poor general health were identified as factors of high mortality.

İlk operasyon sonrası 60 gün içinde

'On demand' 'Planned'

Kanama, Evisserasyon, İleus, anastomoz kaçağı, intraabdominal abse...

Abdominal reoperasyon % 1.1-4.4

Mortalite yüksek

Risk ?

Yaş (> 65)

Erkek cinsiyet

Genel durum bozukluğu

Anastomoz komplikasyonu

Some severe postoperative complications as consequent even elective colorectal cancer surgery may require reoperation. Relaparotomy is defined as an abdominal operation performed after an initial surgery within 60 days, and the decision is made on the basis of general reaction to surgical stress. The incidence of relaparotomy-requiring complications has been reported to be 1.1%–4.4% [1,2]. When the first (index) operation was performed

* The paper was accepted for poster presentation during: 8 International European Federation for Colorectal Cancer, Vienna, Austria, 4–6 April, 2013.

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1674 hasta elektif operasyon

121 (% 7.2) reoperasyon gereksinimi

Anastomoz yapılmayan grupta abse daha fazla

Mortalite - abdominal sepsis/septik şok (%68.7)

ter reoperations was so high. In a study of Sorensen et al. the incidence of relaparotomy in colorectal disease in medical history, malignancy and A

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Methods: Cohort analysis of 20,058 stage I to III colorectal cancer patients in the 1992 to 1996 SEER-Medicare database. *Independent variables:* sociodemographics, tumor characteristics, comorbidity, and acuity. *Primary outcome:* postoperative procedural intervention. *Analysis:* Logistic regression identified patient characteristics predicting postoperative procedures and the adjusted risk of 30-day mortality and prolonged hospitalization among patients with postoperative procedures.

Results: A total of 5.8% of patients required postoperative intervention. Patient characteristics had little impact on the frequency of postoperative procedures, except for acute medical conditions, including bowel perforation (relative risk [RR] = 3.0, 95% confidence interval [CI] = 2.5–3.6), obstruction (RR = 1.6; 95% CI = 1.4–1.8), and emergent admission (RR = 1.3; 95% CI = 1.1–1.4). After a postoperative procedure, patients were more likely to experience early mortality (RR = 2.4; 95% CI = 2.1–2.9) and prolonged hospitalization (RR = 2.2; 95% CI = 2.1–2.4). The most common interventions were performed for abdominal infection (31.7%; RR mortality = 2.9; 95%

Interest in measuring surgical quality is growing rapidly. Increasing recognition that surgical outcomes vary widely among patients are seeking more detailed information about provider performance prior to undergoing treatment. Providers are interested in assessing their own performance for quality improvement purposes. Payers are looking for better data by which to steer selected populations of surgical patients to high-quality providers. To meet these various interests, policy makers and health services researchers have redoubled their efforts to develop and implement quality indicators germane to surgery.

Unfortunately, however, current measures of surgical quality have major flaws which limit their usefulness. The most simple and direct, operative mortality is too uninformative to

sensitivity of claims was only 35%, although sensitivity of claims for reoperation was greater than 60%. We have minimized this limitation by using both ICD-9-CM and CPT codes to focus only on the subset of major surgical complications that require reoperation or other procedural intervention and are

that major complication rates after surgery were similar to the current population. Like our study, these investigators found that complications occur in a small subset of patients, usually as a result of surgically significant events that often

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Unfortunately, however, current measures of surgical quality have major flaws which limit their usefulness. Although simple and direct, operative mortality is too uncommon a measure for most procedures to allow consistent and reliable measurement of surgical quality and too blunt to direct quality improvement efforts. Volume of care has been correlated with

Mortalite riski 3 kat

Postop hastanede yatış süresi 2.7 kat



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We investigated complications of surgery in the context of colorectal cancer, a common and potentially fatal disease⁶ that is primarily treated with well-established surgical techniques.⁷ Our aim was to identify unplanned procedural interventions following colorectal cancer surgery that might be used as interme-



Tedavi -

Hangi antibiyotikler ?

Ne zaman ve nasıl bir Cerrahi girişim

Hangi antibiyotik ?

- Seftazidim / Sefepim
- Sefaperazon sulbaktam

• Seftriakson + Metronidazol

• Siprofloksasin/Levofloksasin + Metronidazol

• Moksifloksasin

• Piperasilin-tazobaktam

• Tigesiklin

• Karbapenemler



SIS-IDSA Rehberi-2010
WSES Rehberi-2012
SIS - 2017 revizyon

Konak özellikleri ?

Hastalığın ciddiyeti ?

Toplum / Hastane kökenli ?

Toplum / Hastane etken direnç özellikleri ?



Diagnosis and Management of Complicated Intra-abdominal Infection in Adults and Children: Guidelines by the Surgical Infection Society and the Infectious Diseases Society of America

Joseph S. Solomkin,¹ John E. Mazuski,² John S. Bradley,³ Keith A. Rodvold,^{7,8} Ellie J. C. Goldstein,⁵ Ellen J. Baron,⁶ Patrick J. O'Neill,⁹ Anthony W. Chow,¹⁶ E. Patchen Dellinger,¹⁰ Soumitra R. Eachempati,¹¹ Sherwood Gorbach,¹² Mary Hilfiker,⁴ Addison K. May,¹³ Avery B. Nathens,¹⁷ Robert G. Sawyer,¹⁴ and John G. Bartlett¹⁵

Regimen	Community-acquired infection in pediatric patients	Community-acquired infection in adults	
		Mild-to-moderate severity: perforated or abscessed appendicitis and other infections of mild-to-moderate severity	High risk or severity: severe physiologic disturbance, advanced age, or immunocompromised state
Single agent	Ertapenem, meropenem, imipenem-cilastatin, ticarcillin-clavulanate, and piperacillin-tazobactam	Cefoxitin, ertapenem, moxifloxacin, tigecycline, and ticarcillin-clavulanate	Imipenem-cilastatin, meropenem, doripenem, and piperacillin-tazobactam
Combination	Ceftriaxone, cefotaxime, cefepime, or ceftazidime, each in combination with metronidazole; gentamicin or tobramycin, each in combination with metronidazole or clindamycin, and with or without ampicillin	Cefazolin, cefuroxime, ceftriaxone, cefotaxime, ciprofloxacin, or levofloxacin, each in combination with metronidazole ^a	Cefepime, ceftazidime, ciprofloxacin, or levofloxacin, each in combination with metronidazole ^a



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Recommendations for Empiric Antimicrobial Therapy for Health Care–Associated Complicated Intra-abdominal Infection

Organisms seen in health care–associated infection at the local institution	Regimen				
	Carbapenem ^a	Piperacillin-tazobactam	Ceftazidime or cefepime, each with metronidazole	Aminoglycoside	Vancomycin
<20% Resistant <i>Pseudomonas aeruginosa</i> , ESBL-producing Enterobacteriaceae, <i>Acinetobacter</i> , or other MDR GNB	Recommended	Recommended	Recommended	Not recommended	Not recommended
ESBL-producing Enterobacteriaceae	Recommended	Recommended	Not recommended	Recommended	Not recommended
<i>P. aeruginosa</i> >20% resistant to ceftazidime	Recommended	Recommended	Not recommended	Recommended	Not recommended
MRSA	Not recommended	Not recommended	Not recommended	Not recommended	Recommended



Recommendations for intra-abdominal infections consensus report

Vildan Avkan-Oğuz¹, Nurcan Baykam², Selman Sökmen³, Rahmet Güner⁴, Fatih Ağalar⁵, Emine Alp⁶, Ahmet Doğrul⁷, Özge Turhan⁸, Canan Ağalar⁹, Behice Kurtaran¹⁰, İbrahim Ethem Geçim¹¹, Reşat Özaras¹², Gürdal Yılmaz¹³, Ayhan Akbulut¹⁴, İftihar Koksal^{13,15}

DERLEME / REVIEW

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Erişim: <http://dx.doi.org/10.4274/mjima.2016.7>



Intra-abdominal Enfeksiyonlar İçin Öneriler “Uzlaşma Raporu”

Recommendations for Intra-abdominal Infections “Consensus Report”

Vildan AVKAN OĞUZ¹, Nurcan BAYKAM², Selman SÖKMEN³, Rahmet GÜNER⁴, Fatih AĞALAR⁵, Emine ALP MEŞE⁶, Ahmet DOĞRUL⁷, Özge TURHAN⁸, Canan AĞALAR⁹, Behice KURTARAN¹⁰, İbrahim Ethem GEÇİM¹¹, Reşat ÖZARAS¹², Gürdal YILMAZ¹³, Ayhan AKBULUT¹⁴, İftihar KÖKSAL^{13,*}

İntra-abdominal enfeksiyon mu?

Ateş, karın ağrısı, rebound, defans, kusma, halsizlik vs. var mı?

Evet

Olası İAE

- Son 3 ay içinde 5 gün veya daha uzun süre hastane yatışı?
- Son bir ay içinde 2 gün veya daha uzun süre antibiyotik kullanımı?
- Son 3 ay içinde abdominal girişim var mı?

Du sorulardan herhangi birisi Evet ise;

Sağlık bakımı ilişkili İAE

!Bölgesel direnç verilerini gözden geçir!

Yanıt hepsi için Hayır ise;

Toplumdan edinilmiş İAE

1. **Yüksek riskli hasta mı? Konak özellikleri?**
İmmünyüpresyon, ciddi kardiyopulmoner hastalık, diyabetes mellitus vs.?
Yanıt EVET ise; yüksek riskli hasta

2. **Hastalığın ciddiyeti?**
APACHE II ≥ 15 ise ciddi İAE

3. **Tam: Olası enfeksiyon kaynağı;**
Laboratuvar (beyaz hücre, CRP, prokalsitonin)?
İntra-operatif örnek, kültür alımı
Görüntüleme?

4. **Tedavi;** sağlık bakımı ilişkili İAE -öneri 32-39
toplumdan edinilmiş İAE - Tablo 8 ve 9

Ampirik tedavinin 48-72. saatinde klinik ve laboratuvar değerlendirme yap:

- Kaynak kontrolü yapılmış ve ampirik tedaviye yanıt alınmışsa,
Tedavi 4-7 gün
- Kaynak kontrolü sağlanamadıysa ve tedavi yanıtı yoksa;
Tedavi süresi?

Radyolojik görüntüleme US, BT, MR, girişimsel yöntemlerle drenaj vs.
Farklı bir enfeksiyon odağı; pnömoni, üriner sistem enfeksiyonu vs.?
Enfeksiyon dışı benzer tablolar? (kompartment sendromu vs.)

Tablo 8. Ekstra-biliyer komplike intra-abdominal enfeksiyonun başlangıçtaki ampirik tedavisinde kullanılacak ajanlar ve rejimler^[5]

	Yetişkinlerde toplumdaki edinilmiş enfeksiyon	
Rejim	Hafif-orta enfeksiyon Perfore veya apseli apandisit ve diğer enfeksiyonlar	Ciddi enfeksiyon Yüksek risk veya ciddi fizyolojik bozukluk, ileri yaş veya bağışıklığın bozulması durumu
Tek ajan	Ertapenem Moksifloksasin Tigesiklin	Piperasilin-tazobaktam İmipenem-silastatin Meropenem
Kombinasyon	Sefazolin, Sefuroksim, Seftriakson, Sefotaksim, Siprofloksasin veya Levofloksasin ^a + Metronidazol	Sefepim, Seftazidim, Siprofloksasin veya Levofloksasin ^a + Metronidazol

Tablo 9. Yetişkinlerdeki biliyer enfeksiyonların başlangıçtaki ampirik tedavisinde kullanılacak ajanlar ve rejimler^[5]

Enfeksiyon	Rejim
Toplumdan edinilen hafif ile orta şiddette akut kolesistit	Sefazolin, Sefuroksim veya Seftriakson
Ciddi fizyolojik bozukluğa neden olan toplumdaki edinilmiş akut kolesistit, ileri yaş veya bağışıklığın bozulması durumu	İmipenem-silastatin, Meropenem, Piperasilin-tazobaktam, Siprofloksasin, Levofloksasin ^a veya Sefepim + Metronidazol

Kandida riski ???

Tablo 11. İntra-abdominal kandida enfeksiyonları için spesifik ve spesifik olmayan risk faktörleri sıralanmıştır^[10]

Spesifik risk faktörleri	Spesifik olmayan risk faktörleri
<ul style="list-style-type: none">- Laparoskopi dahil tekrarlayan cerrahi girişimler- 24 saat içinde üst gastrointestinal sistem, perforasyonları dahil tedavi edilemeyen ve/veya tekrarlayan perforasyonlar,- Özofagus dahil gastroduodenal cerrahi anastomoz kaçakları başta olmak üzere gastrointestinal anastomoz kaçakları	<ul style="list-style-type: none">- Akut renal yetmezlik,- Santral venöz kateter varlığı,- Parenteral nütrisyon ile beslenme,- Ciddi sepsis,- Diyabetes mellitus,- İmmünsüpresyon,- Uzun süre geniş spektrumlu antibiyotik kullanımının- Yoğun bakım yatışı

IAI - Antibiyotik tedavi süresi ???



Pharmacotherapy, 2018 Apr 20. doi: 10.1002/phar.2118. [Epub ahead of print]

Shortened courses of antibiotics for bacterial infections: A systematic review of randomized controlled trials.

Hanretty AM¹, Gallagher JC².

⊕ Author information

Abstract

BACKGROUND: Commonly prescribed durations of therapy for many, if not most, bacterial infections are not evidence-based. Misunderstandings by clinicians and patients alike influence perspectives on antibiotic use, including duration of therapy and its role in antibiotic resistance.

OBJECTIVE: To demonstrate that shorter durations of antibiotic therapy are as efficacious as longer durations for many infections.

DATA SOURCES: A systematic review of English language articles using PubMed were identified for inclusion. Additionally, infection-specific guidelines were identified for review of recommendations. Search terms included specific infection types, randomized controlled trial (RCT), duration of therapy, treatment duration, short course, and long course.

STUDY SELECTION: Only RCTs of single-agent antibiotic therapy for the treatment of bacterial infections in adults were included.

DATA EXTRACTION: Independent data extraction of articles was conducted by two authors using predefined guidance for article inclusion.

DATA SYNTHESIS: In total, 23 RCTs met our criteria for inclusion. All trials compared single-agent antibiotics for a short and long antibiotic course in six common infections: community-acquired pneumonia, ventilator-associated pneumonia, intra-abdominal infections, skin and soft tissue infections, uncomplicated cystitis, and complicated cystitis or pyelonephritis.

CONCLUSIONS: Clinicians can decrease net antibiotic use by recommending shorter courses where evidence supports them. Antimicrobial stewardship programs that systematically address treatment duration may significantly impact institutional antibiotic use without negatively affecting patient care. This article is protected by copyright. All rights reserved.



Tedavi -

Hangi antibiyotikler ?

Ne zaman ve nasıl bir Cerrahi girişim



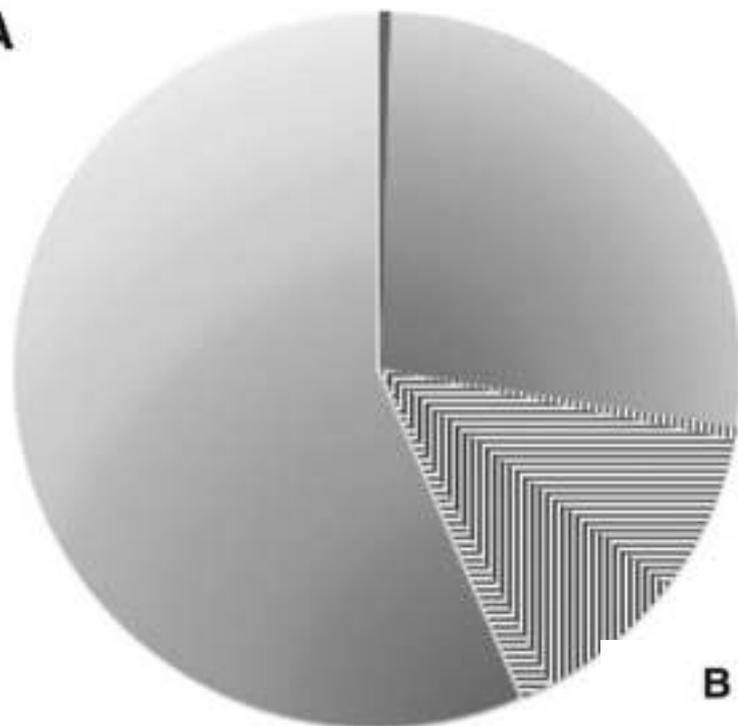
Abdominoperineal Resection, Pelvic Exenteration, and Additional Organ Resection Increase the Risk of Surgical Site Infection after Elective Colorectal Surgery: An American College of Surgeons National Surgical Quality Improvement Program Analysis

Mary R. Kwaan,¹ Genevieve B. Melton,¹ Robert D. Madoff,¹ and Jeffrey G. Chipman²

TABLE 1. ADDITIONAL ORGAN RESECTIONS (N = 17,839) AND ASSOCIATED UNADJUSTED WOUND INFECTION RATES

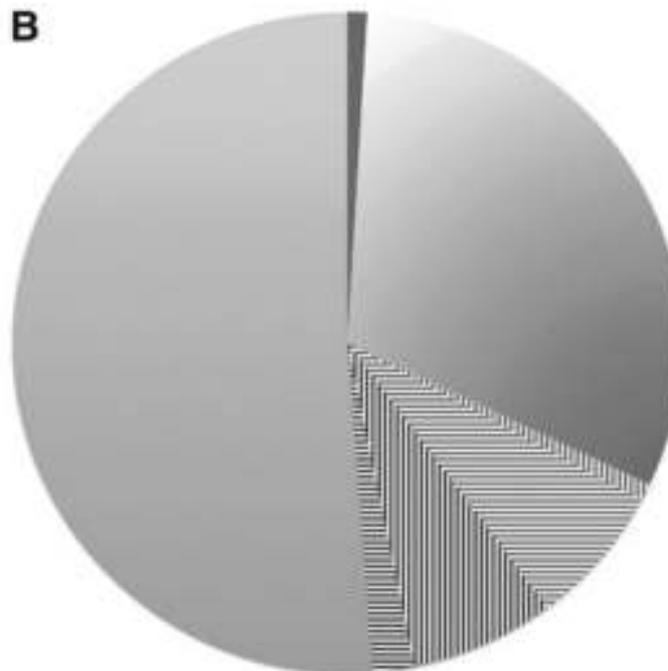
	n	Proportion of cohort	Surgical site infection %
Additional segmental colectomy	14,046	12.5 %	11.6%
Small bowel resection	2,133	2.7 %	13.7%
Hysterectomy	1,120	1 %	12.4%
Bladder resection	667	0.6 %	16.5%
Partial vaginectomy	334	0.3 %	17.1%
Gastric resection	153	0.14%	7.2%
Splenectomy	150	0.13%	18 %
Major vascular repair	86	0.08%	18.6%
Laparoscopic hysterectomy	44	0.04%	6.8%
Diaphragm resection	35	0.03%	17.1%
Pelvic exenteration	510	0.45%	19.4%

A



- Pelvic exenteration
- Proctectomy
- ||| Proctectomy with small bowel resection
- ≡ Proctectomy and additional organ resection
- ||| Colectomy and additional organ resection

B



- Pelvic exenteration
- Proctectomy
- ||| Proctectomy with small bowel resection
- ≡ Proctectomy and additional organ resection
- ||| Colectomy and additional organ resection
- ≡ Colectomy with small bowel resection
- Colectomy

35–39.9	2.24	[2.08 –2.42]	2.11	[1.96 –2.28]
>40	2.72	[2.49 –2.96]	2.51	[2.30 –2.74]
Pre-operative inpatient hospitalization	1.11	[1.03 –1.18]	1.14	[1.07 –1.22]
Smoker	1.26	[1.19 –1.32]	1.24	[1.18 –1.31]
Diabetes mellitus	1.13	[1.07 –1.20]	1.13	[1.06 –1.19]
Pre-operative weight loss		NS		NS
Pulmonary comorbidities	1.21	[1.14 –1.28]	1.22	[1.15 –1.3]
Cardiac comorbidity	1.09	[1.01 –1.18]	1.11	[1.03 –1.2]
Peripheral vascular disease		NS		NS
Disseminated cancer	1.19	[1.08 –1.31]	1.14	[1.04 –1.26]
Immunosuppressive medications or steroid use		NS		NS
Radiation therapy within 90 d of surgery	1.13	[1.01 –1.26]	1.12	[1.0 –1.25]
Chemotherapy within 30 d of surgery		NS		NS
Serum albumin <3.0 mg/dL		NS		NS
Hematocrit >37%		NS		NS
Hematocrit 29%–37%		NS		NS
Hematocrit 26%–28%		NS		NS
Hematocrit <26%		NS		NS
Benign neoplasm		Reference group		
Colon cancer	0.99	[0.92 –1.07]	0.97	[0.90 –1.04]
Rectal cancer	1.05	[0.95 –1.15]	0.94	[0.85 –1.03]
Diverticular disease	1.11	[1.03 –1.21]	1.10	[1.01 –1.19]
Inflammatory bowel disease	1.24	[1.1 –1.4]	1.12	[0.99 –1.26]
Bowel obstruction	1.47	[1.30 –1.67]	1.40	[1.24 –1.59]
Colonic volvulus	0.99	[0.8 –1.23]	1.04	[0.83 –1.29]
Surgical site infection classification 3 or 4	1.21	[1.14 –1.28]	1.18	[1.11 –1.25]
Stoma procedure included		NS		NS
Urostomy		NS		NS
Total proctocolectomy		NS		NS
Abdominoperineal resection	1.62	[1.46 –1.79]	1.58	[1.42 –1.75]
Pelvic exenteration	1.38	[1.08 –1.78]	1.07	[0.84 –1.38]
Rectal resection		NS		NS
Laparoscopic approach	0.56	[0.53 –0.59]	0.55	[0.52 –0.57]
Additional organ resection	1.19	[1.12 –1.26]	1.08	[1.02 –1.15]
Soft tissue flap procedure	1.26	[1.00 –1.59]	0.88	[0.70 –1.11]
Operative duration (minutes)		Not included	1.002	[1.002–1.002]

COMPLEX COLORECTAL SURGERY AND SSI

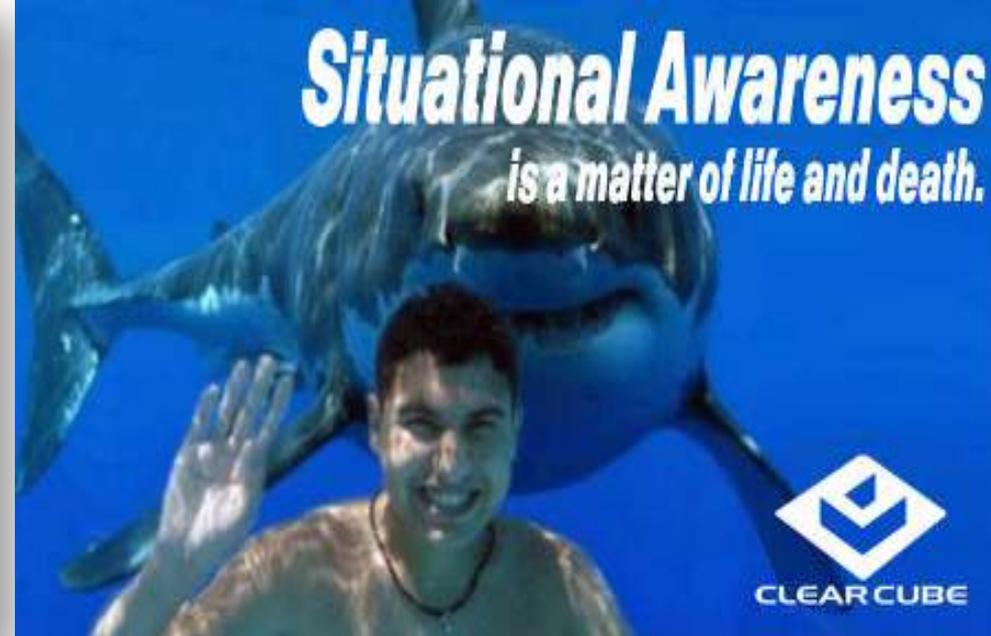
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TABLE 4. RATE OF WOUND INFECTION BY OPERATIVE DURATION

	Surgical site infection %	Proportion of the entire cohort n (%)	Procedure subsets stratified by operative time n (%)		
			Abdominoperineal resection	Exenteration	Colorectal resection with additional organ resection
<1 h	5.6%	3,573 (3.2%)	15 (0.3%)	3 (0.6%)	155 (.9%)
1–2 h	6.7%	29,082 (26.4%)	191 (3.9%)	1 (0.2%)	2,287 (13.1%)
2–3 h	8.8%	34,198 (31.0%)	895 (18.4%)	19 (3.8%)	4,986 (28.5%)
3–4 h	9.9%	21,674 (19.7%)	1,245 (25.5%)	44 (8.7%)	4,485 (25.6%)
4–5 h	12.2%	10,936 (9.9%)	975 (20 %)	70 (13.9%)	2,659 (15.2%)
5–6 h	13.6%	5,287 (4.8%)	640 (13.1%)	71 (14.1%)	1,373 (7.8%)
>6 h	15.1%	5,373 (4.9%)	911 (18.7%)	296 (58.7%)	1,537 (8.8%)

- Familiarity of the situation
- Focusing of attention
- Information quantity
- Information quality
- Instability of the situation
- Concentration of attention
- Complexity of the situation
- Variability of the situation
- Arousal
- Spare mental capacity

The Situation Awareness Rating Technique uses 10 different dimensions.



Anesthesiology, 2013 Mar;118(3):729-42. doi: 10.1097/ALN.0b013e318280a40f.

Situation awareness in anesthesia: concept and research.

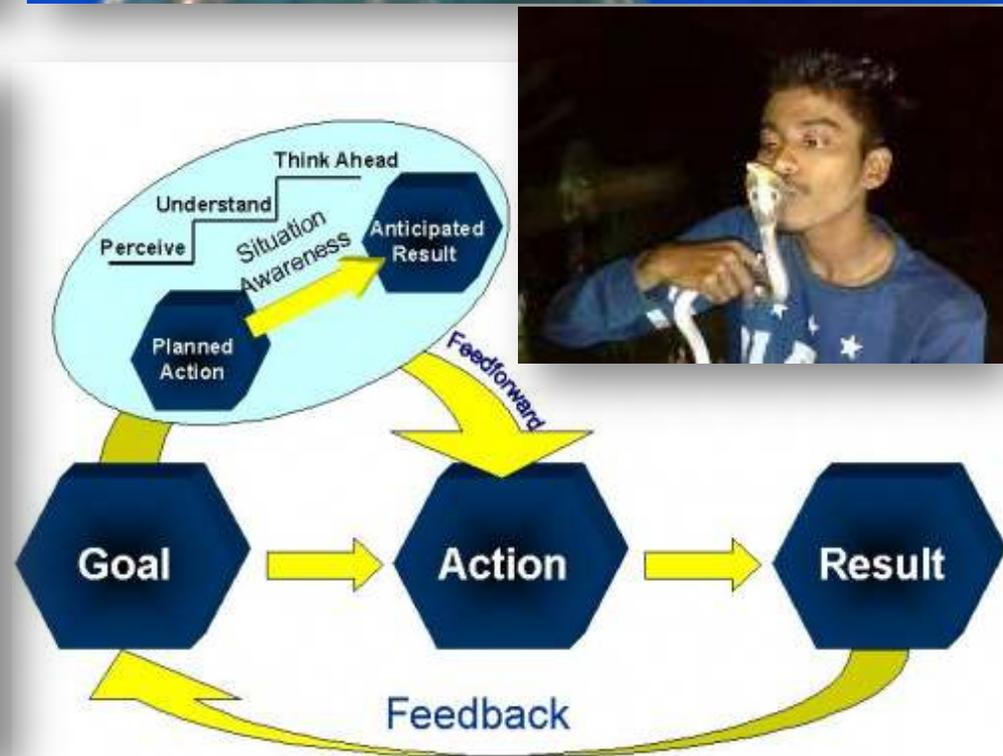
Schulz CM¹, Endsley MR, Kochs EF, Gelb AW, Wagner KJ.

Author information

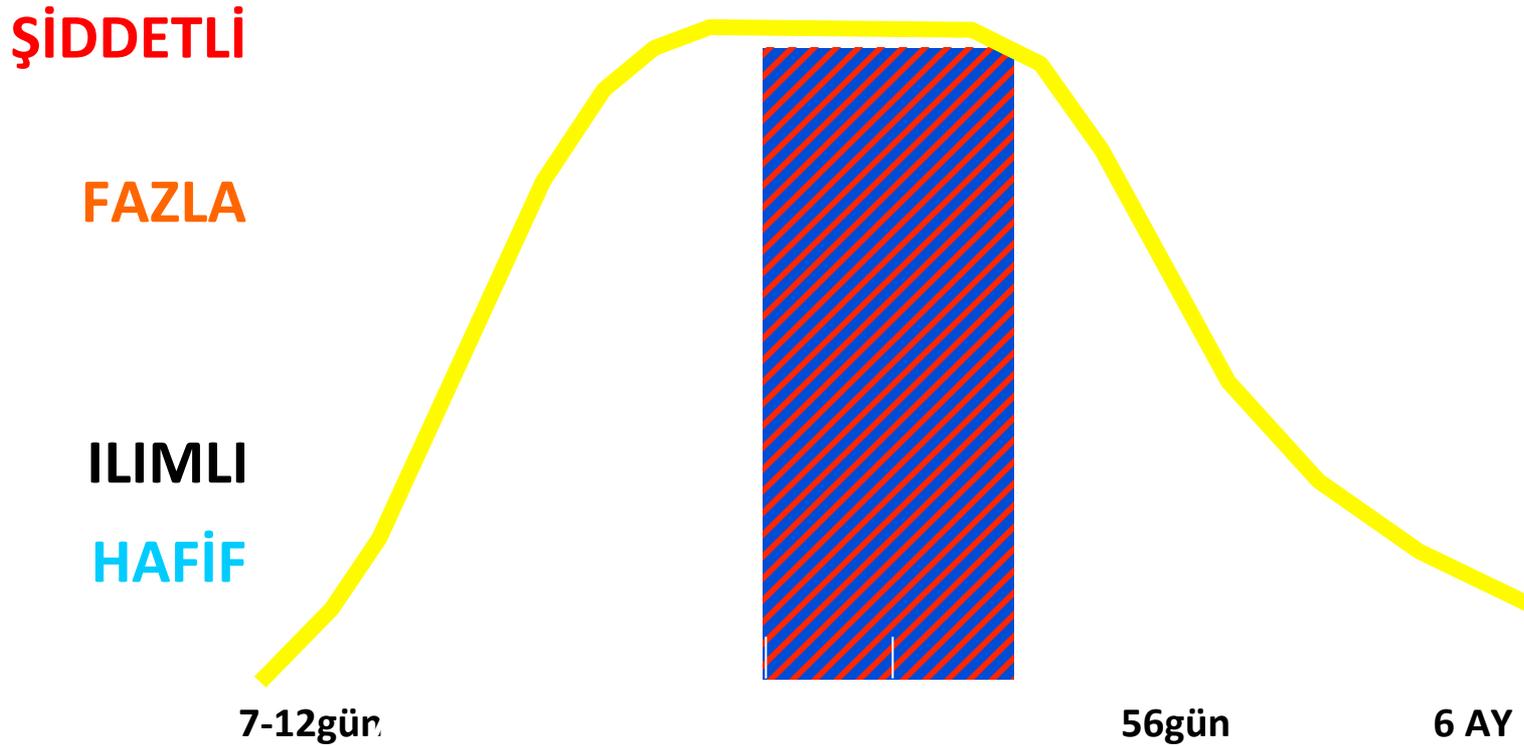
Abstract

Accurate situation awareness (SA) of medical staff is integral for providing optimal performance during the treatment of patients. An understanding of SA and how it affects treatment of patients is therefore crucial for patient safety and an essential element for research on human factors in anesthesia. This review describes the concept of SA in the anesthesia environment, including the interaction with associated medical teams. Different approaches for its assessment in the work environment of anesthesia are provided. Factors contributing to expertise in SA are described and approaches for the training of SA in anesthesia are discussed, as are types of errors that occur during the development of SA. Finally, the authors briefly present strategies to improve SA during daily anesthesia practice through altered designs of monitor displays.

PMID: 23291626 DOI: 10.1097/ALN.0b013e318280a40f



ADEZYONLARIN ŞİDDETI



Ne zaman ameliyat edilmeli?

- 4-6 ay bekle
- KARIN DUVARI mobilitesi (fazio manevrası)
 - Her iki el hastanın karnının her iki yanına konur
 - Güçlü bir şekilde ileri-geri hareket ettirilir
 - Rijid('düşman karın') veya HAREKETLİ!
 - Hareketli ise laparotomi daha kolay olacak demektir(adezyonlar yumuşamıştır)

"BEKLENMEYENİN" önlenmesi

- Düzenini kur
- Laboratuvar çalışmaları
- Düzeltilebilir eksiklikleri düzelt
- Doku planlarını ve anatomiye haritalayabileceğin radyolojik tetkikler
- O gün için başka program yapma
- Üreterik stentler, es/tdp temini, aydınlatma ve thompson retraktör sistemi

Adezyolizis ve relaparotomi tekniği

- Abdomene en kolay yerinden gir
- Ekspozur yeterli olmalı
- Bağırsağı dekomprese et; özellikle interlup
- Birbirine yapışmış lupları karın dışına al
- Üreteri pelvis içinde arama
- Arkanda dolaşımı bozuk yağ dokusu bırakma



Adezyolizis ve relaparotomi tekniđi «durumsal farkındalık»

- Keskin diseksiyon
- Sf enjeksiyonu (hidrodiseksiyon)
- Ekstrafasiyal diseksiyon
- Serozal yırtıkları onar
- Emilebilen dikişler kullan
- En zor kısmı en sona bırak
- Pudra ve benzeri kontaminasyon olmamalı

Reoperatif cerrahide risk faktörleri

- Büyük tümörler
- Işınlanmış bağırsak
- Anatomik varyantlar
- Kısa rektal güdük
- Kronik abdominopelvik sepsis
- Yüksek riskli hasta
- Tecrübesiz cerrah



Ülkemizde İntra-abdominal enfeksiyonlar

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REVIEW

Open Access



Antimicrobials: a global alliance for optimizing their rational use in intra-abdominal infections (AGORA)

Massimo Sartelli^{1*}, Dieter G. Weber², Etienne Ruppé³, Matteo Bassetti⁴, Brian J. Wright⁵, Luca Ansaloni⁶, Fausto Catena⁷, Federico Coccolini⁸, Fikri M. Abu-Zidan⁹, Raul Coimbra¹⁰, Ernest E. Moore¹¹, Frederick A. Moore¹², Ronald V. Maier¹³, Jan J. De Waele¹⁴, Andrew W. Kirkpatrick¹⁵, Ewen A. Griffiths¹⁶, Christian Eckmann¹⁷, Adrian J. Brink¹⁸, John E. Mazuski¹⁹, Addison K. May²⁰, Rob G. Sawyer²¹, Dominik Mertz²², Philippe Montravers²³, Anand Kumar²⁴, Jason A. Roberts²⁵, Jean-Louis Vincent²⁶, Richard R. Watkins²⁷, Warren Lowman²⁸, Brad Spellberg²⁹, Iain J. Abbott³⁰, Abdulrashid Kayode Adesunkanmi³¹, Sara Al-Dahir³², Majdi N. Al-Hasan³³, Ferdinando Agresta³⁴, Asma A. Althani³⁵, Shamshul Ansari³⁶, Rashid Ansumana³⁷, Goran Augustin³⁸, Miklosh Bala³⁹, Zsolt J. Balogh⁴⁰, Oussama Baraket⁴¹, Aneel Bhangu⁴², Marcelo A. Beltrán⁴³, Michael Bernhard⁴⁴, Walter L. Biffi⁴⁵, Marja A. Boermeester⁴⁶, Stephen M. Brecher⁴⁷, Jill R. Cherry-Bukowiec⁴⁸, Otmar R. Buyne⁴⁹, Miguel A. Cainzos⁵⁰, Kelly A. Cairns⁵¹, Adrian Camacho-Ortiz⁵², Sujith J. Chandy⁵³, Asri Che Jusoh⁵⁴, Alain Chichom-Mefire⁵⁵, Caroline Colijn⁵⁶, Francesco Corcione⁵⁷, Yunfeng Cui⁵⁸, Daniel Curcio⁵⁹, Samir Delibegovic⁶⁰, Zaza Demetrashvili⁶¹, Belinda De Simone⁶², Sameer Dhingra⁶³, José J. Diaz⁶⁴, Isidoro Di Carlo⁶⁵, Angel Dillip⁶⁶, Salomone Di Saverio⁶⁷, Michael P. Doyle⁶⁸, Gereltuya Dorj⁶⁹, Agron Dogjani⁷⁰, Hervé Dupont⁷¹, Soumitra R. Eachempati⁷², Mushira Abdulaziz Enani⁷³, Valery N. Egiev⁷⁴, Mutasim M. Elmangory⁷⁵, Paula Ferrada⁷⁶, Joseph R. Fitchett⁷⁷, Gustavo P. Fraga⁷⁸, Nathalie Guessennd⁷⁹, Helen Giamarellou⁸⁰, Wagih Ghnnam⁸¹, George Gkiokas⁸², Staphanie R. Goldberg⁷⁶, Carlos Augusto Gomes⁸³, Harumi Gomi⁸⁴, Manuel Guzmán-Blanco⁸⁵, Mainul Haque⁸⁶, Sonja Hansen⁸⁷, Andreas Hecker⁸⁸, Wolfgang R. Heizmann⁸⁹, Torsten Herzog⁹⁰, Adrien Montcho Hodonou⁹¹, Suk-Kyung Hong⁹², Reinhold Kafka-Ritsch⁹³, Lewis J. Kaplan⁹⁴, Garima Kapoor⁹⁵, Aleksandar Karamarkovic⁹⁶, Martin G. Kees⁹⁷, Jakub Kenig⁹⁸, Ronald Kiguba⁹⁹, Peter K. Kim¹⁰⁰, Yoram Kluger¹⁰¹, Vladimir Khokha¹⁰², Kaoru Koike¹⁰³, Kenneth Y. Y. Kok¹⁰⁴, Victory Kong¹⁰⁵, Matthew C. Knox¹⁰⁶, Kenji Inaba¹⁰⁷, Arda Isik¹⁰⁸

Introduction

The Enterobacteriaceae, including *Escherichia coli*, *Klebsiella pneumoniae*, *Klebsiella oxytoca*, *Proteus mirabilis*, *Citrobacter* spp. and *Enterobacter* spp., is the family most commonly implicated in the aetiology of both urinary tract infections (UTIs) and intra-abdominal infections (IAIs).¹⁻³ Over the last decade, a dramatic increase has occurred in the worldwide prevalence of ESBL-positive Enterobacteriaceae, which frequently show resistance to several antibiotic classes, including fluoroquinolones and aminoglycosides,⁴⁻⁶ and thus represent a challenge for practitioners, further

limiting antibiotic therapy choices and adversely impacting patient outcomes.⁶⁻⁸

The choice of empirical antimicrobial therapy is guided by knowledge of the bacterial spectrum and the extent of antimicrobial resistance.^{9,10} Given the global increase in antimicrobial resistance, particularly for Gram-negative bacteria, the variability of antimicrobial resistance in different geographical regions and over time, and the paucity of novel antibiotics in development, continuous surveillance of the pathogen prevalence including the presence of ESBL producers, as well as of the emergence and trend

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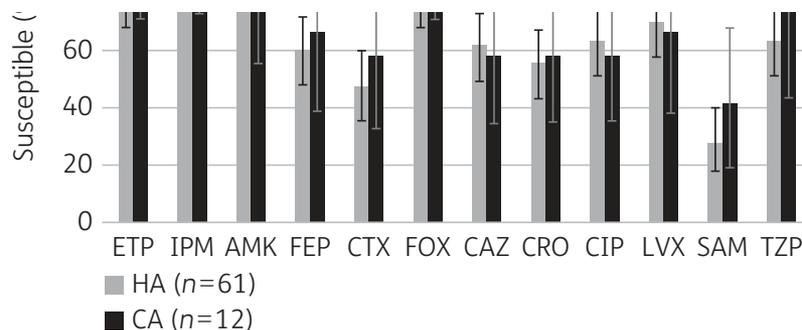


Figure 1. Susceptibility (with 95% CI) for (a) overall Gram-negative isolates and for (b) *E. coli* and (c) *K. pneumoniae* in HA versus CA UTIs in 2011–12. ETP, ertapenem; IPM, imipenem; AMK, amikacin; FEP, cefepime; CTX, cefotaxime; FOX, ceftazidime; CAZ, ceftazidime; CRO, ceftriaxone; CIP, ciprofloxacin; LVX, levofloxacin; SAM, ampicillin/sulbactam; TZP, piperacillin/tazobactam.

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Infectious Complications after Cytoreductive Surgery and Hyperthermic Intra-Peritoneal Chemotherapy

Naciye Cigdem Arslan,¹ Selman Sokmen,² Vildan Avkan-Oguz,³ Funda Obuz,⁴
Aras Emre Canda,² Cem Terzi,² and Mehmet Fuzun²

Abstract

Background: The aim of this study was to review the post-operative and infectious complications and determine the risk factors associated with infections in cytoreductive surgery (CRS) and hyperthermic intra-peritoneal chemotherapy (HIPEC).

Patients and Methods: Between October 2007 and December 2013, patients who underwent CRS and HIPEC with a

MICRO-ORGANISMS ISOLATED FROM CULTURES

Microorganisms	Infections (n, %)			
	Surgical site	Blood stream	Pulmonary	Urinary tract
<i>Escherichia coli</i> /ESBL+	19 (53)/9	4 (44)/4	-	10 (71)/3
<i>Klebsiella pneumoniae</i> /ESBL+	6 (17)	4 (44)/1	-	2 (14)
<i>Enterococcus spp</i>	10 (28)	1 (11)	-	-
<i>Staphylococcus spp</i>	-	1 (11)	2 (23)	-
<i>Acinetobacter spp</i>	-	2 (23)	2 (23)	-
<i>Pseudomonas aeruginosa</i>	2 (6)	1 (11)	-	2 (14)
<i>Candida albicans</i>	6 (17)	6 (67)	3 (27)	6 (43)
Others	2 (6)	1 (11)	-	-



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INFECTIONS IN PERITONEAL CARCINOMATOSIS TREATMENT

TABLE 5. DETAILS OF THE PATIENTS WITH PERI-OPERATIVE MORTALITY

Patient	Complications	Culture
#1	Pneumonia, nephrotoxicity	<i>Pseudomonas aeruginosa</i>
#2	Anastomotic leak, urinary leak, hematotoxicity (febrile neutropenia), SSI, BSI	<u><i>C. albicans</i></u> , <i>K. pneumoniae</i> (ESBL+)
#3	SSI, BSI, nephrotoxicity, hematotoxicity (febrile neutropenia)	<u><i>C. albicans</i></u> , <i>E. coli</i> (ESBL+), MR-CoNS, <i>K. pneumoniae</i>
#4	Anastomotic leak, SSI, BSI, UTI	<u><i>C. albicans</i></u> , <i>E. coli</i> (ESBL+), <i>Klebsiella</i> , <i>S. aureus</i>
#5	Pneumonia, UTI	<u><i>C. albicans</i></u> , <i>Acinetobacter sp.</i>
#6	Pneumonia, BSI, SSI, nephrotoxicity, hematotoxicity (febrile neutropenia)	<u><i>Acinetobacter sp.</i></u> , <i>E. coli</i>
#7	Pleural effusion, SSI, BSI	<u><i>C. albicans</i></u> , <i>E. coli</i>
#8	Pneumonia, iatrogenic small bowel injury, SSI, BSI, hematotoxicity (febrile neutropenia)	<u><i>C. albicans</i></u> , <i>E. coli</i>
#9	Pneumonia, nephrotoxicity, hematotoxicity	NA
#10	Pneumonia, urinary leak	NA
#11	Stroke	-



SONUÇ - I

- Multidisipliner ekip birlikte öğrenir.
- Toplum kaynaklı - Sağlık bakımı ilişkili ?
- 72. saatte antimikrobiyal tedavi yanıtı değerlendirilmeli
- Ağır yaygın kompleks kanser olgularında komplikasyon (İnfeksiyöz + Cerrahi) BEKLENİR

SONUÇ

- Bir merkez/ekibin onkolojik başarı ölçütü komplikasyonları tedavi edebilmesi ile değerlendirilir.
' Failure to rescue ' kavramı
- Reoperasyon, tekrarlayan girişimler ;
Mortalite ve hastanede yatış süresini uzatır
İnfeksiyon riski ? Maliyet ?
- İnfeksiyon - Önce olay yeri incelenmeli
(Abdominal pnömoni aranmalı !!!)