

Pathogenic Activity of Methicillin-Resistant *Staphylococcus aureus* Strains Isolated from Colorectal Cancer Patients

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Abstract: There were studied pathogenic activity of methicillin-resistant *Staphylococcus aureus* (MRSA) strains isolated from colorectal cancer patients and were compared to non-MRSA strains isolated from patients with intestinal dysbiosis. The aim of study was to determine changes of biological properties of this important bacterium depend on the severity of disease. Study revealed that MRSA strains are widespread in patients with colorectal cancer. MRSA strains revealed higher pathogenic activity than non-MRSA strains. Some determinants of pathogenicity (urease, lecithinase production, hemolysis, proteolysis), also carbohydrate and mannitol fermentations in aerobic and anaerobic conditions are characterized with high activity in MRSA strains in comparison of non-MRSA strains. Coagulase- and catalase-positive were all strains from both groups.

Keywords: MRSA, colorectal cancer, *S. aureus*, pathogenic factors, enzymes, carbohydrate fermentation

1. Introduction

More than 100 trillion microorganisms live in the human intestine. They play an important role in health status and diseases, including cancer. Specific bacteria and gastrointestinal dysbiosis can potentiate the development and progression of gastrointestinal neoplasms by destroying DNA, activating oncogenic signaling pathways, producing metabolites that promote the development of tumors and suppressing antitumor immunity [8].

Changes in the interactions among the intestinal epithelial cells, intestinal microflora, and host immune system are associated with many diseases, including cancer [10].

Colorectal cancer represents an important disease as one of the major causes of death worldwide. Besides of many factors (diet, family history, age, ethnicity, others), metabolic products of intestinal microflora also influence and predispose to the development of colorectal cancer [9].

Staphylococcus aureus, although identified as a commensal, is also a common cause of human bacterial infections [4]. Colonization of the gastrointestinal system by *S. aureus* strains in hospitalized patients may lead to important clinical implications [4] and is able to acquire resistance to a variety of antibiotics [1,6].

In addition to its ability to outwit human immune system, its multi-drug resistance phenotype, particularly – in methicillin-resistant *S. aureus* (MRSA) strains makes it one of the most intractable pathogenic bacteria in the history of antibiotic chemotherapy [5,7].

Based on the above mentioned reasons we studied the activity of pathogenic factors of *S. aureus* isolated from colorectal cancer patients in comparison with the same

bacteria isolated from non-cancer patients to determine how change biological properties of this important bacterium can depend on the severity of disease.

2. Materials and Methods

There were studied 64 *S. aureus* strains. Sample materials – feces – were taken from patients with colorectal cancer (36 strains – I group) and from patients with intestinal dysbiosis (28 strains – control, II group). Samples were collected before surgical involvement or administration of therapeutic regimes. Cultivation of *S. aureus* were performed on 5% blood agar, egg yolk-salt agar, meat-peptone agar, meat-peptone broth, milk-salt agar of Eikman; Biochemical properties have been analyzed on results of carbohydrate fermentation in aerobic and anaerobic conditions; carotenoid pigment formation was tested on 37°C for 24 hours in milk-salt agar. Catalase activity was tested by using of hydrogen peroxide, hemolysis – on 5% blood agar, hemolytic zones around the colonies indicated positive results). Lecithinase activity was determined by egg-yolk saline medium. Turbidity around the colony with rainbow-like corona indicated positive result, coagulase (was demonstrated by mixing bacteria with blood plasma on a slide; in positive cases occurred clumping with fibrin formation), also were determined urease, proteolytic activities and novobiocin susceptibility tests [2].

Growth characteristics of cultivation were studied in meat-peptone broth. Equal turbidity in broth was one of the indicators of typical strains. Aerobic fermentation of carbohydrate determination was tested by using of 1% Peptone-Water (with glucose, mannitol, sucrose or lactose) in addition of 2ml Andrade reagent. After the incubation, the red color of medium indicated on carbohydrate fermentation. Bacterial culture was transferred in 2% peptone solution for

proteolytic activity. Additional test for *S. aureus* identification was novobiocin sensitivity (MIC- 2mg /ml).

3. Results

MRSA strains were discharged in both groups – 66.66±7.86% (I group) and 28.57±8.53% (II group). According of these data, non-MRSA strains were isolated in low frequency – I group - 33.33±7.85%, II group - 28.57±8.53% cases (Table N1).

Distribution of MRSA and non-MRSA strains in patients

Table N1

<i>S. aureus</i> strains	I group (n=36)		II group (n=28)	
	Abs.	%	Abs.	%
MRSA	24	66.66±7.86	8	28.57±8.53
Non-MRSA	12	33.33±7.85	20	71.42±8.53

Also were studied and compared pathogenic factors of MRSA and non-MRSA strains (table N2). As in table shown all strains were plasma coagulase- and catalase-positive in both groups. Ability of carbohydrate fermentation in anaerobic condition was higher in I group (83.33±7.6% vs. 75.00±12%).

Some pathogenicity factors of *S. aureus*

Table N2

№	Identity markers	MRSA strains n = 24		Non-MRSA strains n = 12	
		abs	%	abs	%
1	Plasma coagulase-positive	24	100	12	100
2	Catalase-positive	24	100	12	100
3	Urease -positive	21	87.50±6.75	6	50.00±14.43
4	Hemolytic activity (hemolysis)	24	100	9	75.00±12.50
5	Lecithinase activity	22	91.66±5.66	9	75.00±12.50
6	Proteolytic activity (production of H ₂ S)	22	91.66±5.66	8	66.66±13.60
7	Mannitol fermentation in anaerobic conditions	20	83.33±7.61	7	58.33±14.23
8	Carbohydrate (glucose, mannitol, lactose, saccharose) fermentation in aerobic conditions	23	83.33±7.61	9	75.00±12.50

Hemolytic activity revealed in all MRSA strains in comparison of non-MRSA strains (75.00±12.50%). Proteolytic (production of H₂S) and lecithinase activities were same in I group - 91.66±5.66%; these enzymes revealed more low activity in II group (75.00±12.50% and 66.66±13.60% - respectively). Mannitol fermentation in anaerobic condition was much higher in I group - 83.33±7.61% than in II group - 58.33±14.23%.

4. Conclusions

- 1) MRSA strains are widespread in patients with colorectal cancer.
- 2) Some determinants of pathogenicity (urease, lecithinase production, hemolysis, proteolysis), also carbohydrate and mannitol fermentations in aerobic and anaerobic conditions are characterized with high activity in

MRSA strains in comparison of non-MRSA strains. Coagulase- and catalase-positive were all strains from both groups.

- 3) MRSA strains revealed higher pathogenic activity than non-MRSA strains.

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