# Investigation of the Antibacterial Properties of Polychlorinated Cyclic Bisimides

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# Abstract

This work is devoted to the study of the influence of polychlorinated cyclic bisimides based on the Diels-Alder reaction for antibacterial properties and structure. As test cultura pseudomonas aeruginosa, intestinal bacillus, staphylococcus and Candida albicans have been used. The research of the antibacterial properties of polychlorinated cyclic bisimides has been studied in a series with dilution method. According to the antibacterial activities of the studied substances, N-[2-maleinimido] ethyl bisimide of endo. exo-1,2,3,4,11,11-[6.2.1.0<sup>5,10</sup>]undec-2-ene-7,8hexachlorotricyclo dicarboxylic acid and N-[4-maleinimidobenzyl] phenyl bisimide of 1,4,5,6,7,7*hexachlorobicyclo*[2.2.1]*hept-5-ene-2,3-dicarboxylic* acid have more active antimicrobial and fungicidal properties.

**Keywords** - antibacterial polymer additives, antibacterial composition materials, polychlorinated cyclic bisimides, antimicrobial and fungicidal properties

### I. INTRODUCTION

As it is known, during the exploitation, the polymer materials are subjected to biocorrosion under the influence of microbes and microfungi. In addition to their physical and mechanical properties, their quality also deteriorates. Typically, bactericidal and fungicidal additives are included in their composition to get rid of this. For this purpose, many antibacterial additives have been developed and widely used for polyolefins, polystyrene and styrene copolymers. For this purpose, many antibacterial additives for copolymers of polyolefins, polystyrene and styrene have been developed and widely used. Range of organic antibacterial supplements is quite broad. Currently, more than 3,000 antibacterial compounds have been tested in various polymeric materials [1]. All organic antibacterial additives contain halogen, thiazole, salicylates, acid amides and other biologically active groups.

It is widely disclosed in scientific papers published on the results of our previous studies on the acquisition of biomolecular active monomers of the salicylic group and their antibacterial polymers [2]. The purpose of this article is to study the antibacterial properties of polychlorinated cyclic bisimides, which are intended for use as additives when purchasing antibacterial polymeric materials that are new biologically active compounds on our part as a continuation of previous research.

#### **II. EXPERIMENTAL PART**

Cyclic bisimides obtained from the Diels-Alder reaction were synthesized by known methods [3-5] and have composition and structure as shown below:

I N, N '- (2-aminoethyl) bisimide of cis-cyclohex-4ene-1,2-dicarboxylic acid. Melting point 146-147 °C (benzene),  $R_f$  0.68. İG-spectrum sm<sup>-1</sup>: 1715, 1780 (C=O), 1608 (C=C), 680-750 (C-Cl).



**II** N, N '-(3,3-dimethoxy-4,4'-diphenylmethane) bisimide of 2,3,4,5-tetrachloro-1,3-cyclohexadiene-5,6-dicarboxylic acid. m.p. 205 °C (benzene),  $R_f$  0.61. İG-spectrum sm<sup>-1</sup>: 1720, 1780 (C=O), 1602 (C=C), 650-710 (C-Cl).



**III** N, N '- (4,4'-diphenylmethane) bisimide of 2,3,4,5,11,11-hexachlorotricyclo [6.2.1.0<sup>5,10</sup>] undec-2ene-7,8-dicarboxylic acid. m.p. 305 °C (hexane), R<sub>f</sub> 0.44. İG-spectrum sm<sup>-1</sup>: 1720, 1780 (C=O), 1603 (C=C), 660-750 (C-Cl).



**IV** N-[2-maleinimide] ethyl bisimide of endo, exo-1,2,3,4,11,11- hexachlorotricyclo [ $6.2.1.0^{5,10}$ ] undec-2-ene-7,8-dicarboxylic acid m.p. 400 °C (chloroform), R<sub>f</sub> 0.67. İG-spectrum sm<sup>-1</sup>: 1717, 1780 (C=O), 1600 (C=C), 650-770 (C-Cl).



**V** N-[4-maleinimidobenzyl] phenyl bisimide of 1,4,5,6,7,7-hexachlorobicyclo [2.2.1]hept-5-en-2,3-dicarboxylic acid m.p. 300 °C (benzene+hexane),  $R_f$  0.67. İG-spectrum sm<sup>-1</sup>: 1720, 1780 (C=O), 1600 (C=C), 660-740 (C-Cl).



The structure and degree of purity of monomers (I, II, III, IV, V) were proved by IG- and NMR spectrums [6]. In IG spectrums of monomers in 1720, 1780, 1602-1610 and 650-780 sm<sup>-1</sup> areas corresponding absorption zones belonging to C=O, C=C and C-Cl groups were recorded.

## **III. RESULTS AND DISCUSSION**

The research of antimiocrobe properties was studied in series by dilution method [7]. It was held in the following ratios (1:100, 1:200, 1:400,1:800). Test results were shown in Table I-II.

74		Synthesized new compounds																			
ures	Exposure time (minute	Ι				II			III			IV			V						
Test cult		1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4	1	2	3	4
S	10	-	-	+	+	-	-	+	+	-	+	+	+	-	-	+	+	-	-	+	+
reu	20	-	-	+	+	-	-	+	+	-	+	+	+	-	-	+	+	-	-	+	+
.au	40	-	-	+	+	-	-	+	+	-	+	+	+	-	- + - + - + - + - +	+	-	-	+	+	
St.	60	-	-	+	+	-	-	+	+	-	+	+	+	-	-	+	+	-	-	+	+
Ps.aerugi noza	10	-	-	+	+	-	-	-	+	-	•	+	+	-	-	+	+	-	-	-	+
	20	-	-	+	+	-	-	-	+	-	•	+	+	-	-	-	+	-	-	•	+
	40	-	-	-	+	-	-	-	+	-	-	+	+	-	-	-	+	-	-	-	+
	60	-	-	-	+	-	-	-	+	-	-	+	+	-	-	-	+	-	-	-	-
E.coli	10	-	-	+	+	-	-	+	+	-	-	+	+	-	-	-	+	-	-	+	+
	20	-	-	-	+	-	-	+	+	-	-	+	+	-	-	-	-	-	-	+	+
	40	-	-	-	+	-	-	+	+	-	-	+	+	-	-	-	-	-	-	+	+
	60	-	-	-	+	-	-	+	+	-	-	+	+	-	-	-	-	-	-	+	+
	10	-	-	+	+	-	-	+	+	-	-	+	+	-	-	-	+	-	-	-	+
ida ans	20	-	-	+	+	-	-	+	+	-	-	+	+	-	-	-	+	-	-	-	-
<b>Candi</b> albica	40	-	-	-	+	-	-	+	+	-	-	+	+	-	-	-	+	-	-	-	-
	60	-	-	-	+	-	-	-	+	-	-	+	+	-	-	-	+	-	-	-	-

 TABLE I

 Antimicrobial effects of new synthesized compounds and controls

**Symbols:** Diluted in proportion 1(1:100), 2 (1:200), 3 (1:400), 4 (1:800). "+"shows full end, "-" does not end

Test cultures	Exposure time (minute)	Control Ingredients												
			Riv	anol			Eth	anol		Nitrofungin				
		1	2	3	4	1	2	3	4	1	2	3	4	
St.aureus	10	+	+	+	+	-	+	+	+					
	20	+	+	+	+	-	+	+	+					
	40	+	+	+	+	-	+	+	+					
	60	-	-	+	+	-	+	+	+					
Ps.aerugi noza	10	+	+	+	+	+	+	+	+					
	20	+	+	+	+	-	+	+	+					
	40	+	+	+	+	-	+	+	+					
	60	-	-	-	+	-	+	+	+					
E.coli	10	-	-	-	+	+	+	+	+					
	20	-	-	-	+	-	+	+	+					
	40	-	-	-	+	-	+	+	+					
	60	-	-	-	+	-	+	+	+					
Candida albicans	10	+	+	+	+	+	+	+	+	+	+	+	+	
	20	+	+	+	+	+	+	+	+	+	+	+	+	
	40	+	+	+	+	-	+	+	+	-	+	+	+	
	60	+	+	+	+	-	+	+	+	-	+	+	+	

Table II Antimicrobial effects of Control Ingredients

As it is seen from Table I newly synthesized substances influence to various microorganisms differently and when diluted in 1:100 ratio these substances show strong antimicrobe activity against all tested test culturas. Antimicrobe impact of these substances was studied comparatively with alcohol, rivanol and nitrofurgine that are widely used in medicine.

## **IV. CONCLUSIONS**

Tested substances N-[2-maleinimido] ethyl bisimide endo, exo-1,2,3,4,11,11of [6.2.1.0<sup>5,10</sup>]undec-2-ene-7,8hexachlorotricyclo dicarboxylic acid (IV) and N-[4-maleinimidobenzyl] phenyl bisimide of 1,4,5,6,7,7-hexachlorobicyclo [2.2.1]hept-5-ene-2,3-dicarboxylic acid (V) are more active. So, substance (IV) has killed intestinal bacillus in 1:800 ratio dilution during 20 minute, intestinal bacillus and Candida in 1:400 ratio during 10 minute, pseudomonas aeruginosa during 20 minute. Substance (V) keep from developing pseudomonas aeruginosa in 1:400 ratio during 10 minute, Candida in 1:800 ratio for 10 minute. Substances (I, II, IV, V) keep from developing staphylacoccus in 1:200 ratio for 10 minute, but substance (III) could not influence to staphylacoccus.

Examined compounds can be suggested as antimicrobial substances.

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