

## ANTIFUNGAL DEPSIPEPTIDE COMPOUNDS FROM *Paenibacillus polymyxa* HY96-2

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*Paenibacillus polymyxa* can produce many peptide antibiotics, including polymyxins [1], gatavalin [2], jolipeptin [3], tridecaptins [4], the peptide complex named LI-F [5, 6], gavaserin [7], saltavalin [7], fusaricidins A–D [8–10], and so on. Many of them have been described to exhibit antimicrobial activities against certain human pathogens but only a minor number of plant pathogens.

In the course of our search for agricultural antibiotics, we have isolated three antifungal depsipeptides from the culture broth of *P. polymyxa* HY96-2 by bioassay-guided isolation techniques.

The culture broth (30 L) was acidified at pH 3.0 and heated at 80°C for 15 min. The acidified broth was centrifuged, and antifungal compounds were extracted from the supernatant with 3 volumes of *n*-BuOH. Then the *n*-BuOH layer was concentrated, and triple volumes of EtOAc were added to precipitate the crude antifungal extract. The crude extract was applied to a silica gel column eluted with CHCl<sub>3</sub>–MeOH and a Sephadex LH-20 column eluted with MeOH, then purified by preparative RP-HPLC with 32% CH<sub>3</sub>CN in 0.1% TFA. Finally, we obtain compound **1** (20 mg) and a mixture of compounds **2** and **3** (10 mg).

On the basis of spectrum analysis, compounds **1**, **2**, and **3** were identified as fusaricidin A, C, and D, respectively. Fusaricidin A showed strong and broad-spectrum antimicrobial activity against 16 tested plant pathogenic strains. The minimal inhibitory concentrations (MICs) of fusaricidin A against *Fusarium oxysporum*, *Thanatephorus cucumeris* (Frank) Donk., *Rhizoctonia solani*, *Erwinia carotovora* var. *carotovora*, and *Bacillus subtilis* are 12.5 µg/mL, 25 µg/mL, 25 µg/mL, 6.25 µg/mL, and 6.25 µg/mL (Table 1), respectively. The results widened the antifungal spectrum and showed good prospects for the application of fusaricidin A in agriculture.

TABLE 1. The Antimicrobial Activity of Fusaricidin A

Indicator strain	MIC, µg/mL	Indicator strain	MIC, µg/mL
<i>Fusarium semitectum</i>	50	<i>Penicillium thomii</i>	50
<i>Verticillium albo-atrum</i>	25	<i>Curvularia lunata</i>	25
<i>T. cucumeris</i> (Frank) Donk.	25	<i>Alternaria alternata</i> (Fries) Keissler	25
<i>Alternaria longipes</i>	25	<i>Colletrichum lagenarium</i>	25
<i>Fusarium graminearum</i> Schw.	50	<i>Sclerotium</i> sp.	25
<i>Alternaria</i> sp.	25	<i>B. subtilis</i> *	6.25
<i>R. solani</i>	25	<i>E. carotovora</i> var. <i>carotovora</i> *	6.25
<i>F. oxysporum</i> f. sp. <i>cucumerinum</i>	12.5	<i>Escherichia coli</i> *	> 200
<i>F. oxysporum</i> sp. <i>cubense</i> (E. F. Simth) Snyder ethansen	12.5	<i>Xanthomona oryzae</i> *	> 200
<i>Alternaria Solani</i>	25		

Medium – PDA, \*Medium – LB.

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## REFERENCES

1. Zhiping Zhang, *Antibiotics and Bioactive Substance from Microorganism* [in Chinese], Chemical Industry Press, Beijing, 2005, p. 274.
2. N. Nakajima, S. Chihara, and Y. Koyama, *J. Antibiot.*, **25**, 243 (1972).
3. M. Ito and Y. Koyama, *J. Antibiot.*, **25**, 305 (1972).
4. J. Shoji, H. Hino, R. Sakazaki, Y. Wakisaka, M. Mayama, S. Matsuura, and H. Miwa, *J. Antibiot.*, **7**, 646 (1978).
5. K. Kurusu and K. Ohba, *J. Antibiot.*, **40**, 1506 (1987).
6. J. Kuroda, T. Fukai, M. Konishi, J. Uno, K. Kurusu, and T. Nomura, *Heterocycles*, **53**, 1533 (2000).
7. B. Pichard, J. P. Larue, and D. Thouvenot, *FEMS Microbiol. Lett.*, **133**, 215 (1995).
8. Y. Kajimura and M. Kaneda, *J. Antibiot.*, **49**, 129 (1996).
9. Y. Kajimura and M. Kaneda, *J. Antibiot.*, **50** (3), 220 (1997).
10. P. H. Beatty and S. E. Jensen, *Can. J. Microbiol.*, **48**, 1533 (2002).