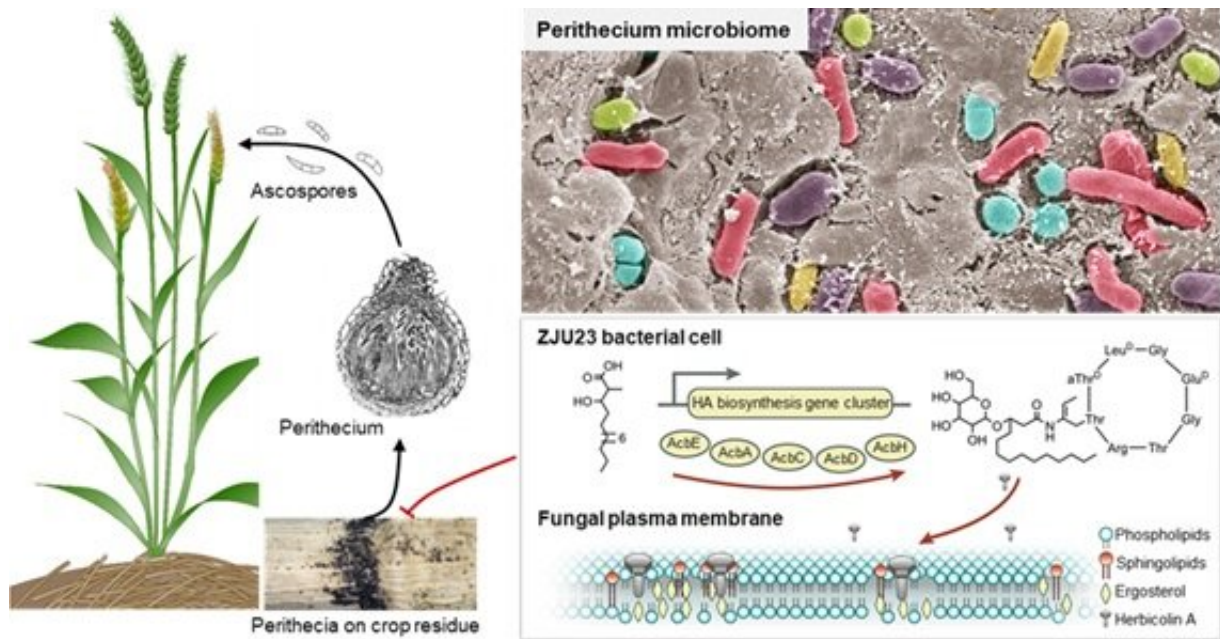


# Harnessing functions of microbiota to combat fungal pathogens

May 31 2022, by Zhang Nannan



Proposed model for the mode of action of herbicolin A secreted by ZJU23 in the fungal perithecium microbiome. Credit: IGDB

*Fusarium graminearum* is a widespread pathogenic fungus that causes Fusarium head blight (FHB) in cereal crops worldwide, especially in wheat. Between 2000 and 2018, more than 4.5 million hectares were annually affected by FHB in China, or around 20% of the total planted area of wheat. This has resulted in annual production losses of more than 3.41 million tons.

Researchers led by Prof. Bai Yang from the Institute of Genetics and Developmental Biology (IGDB) of the Chinese Academy of Sciences, Prof. Chen Yun and Prof. Yu Yunlong at Zhejiang University, have recently found that *F. graminearum* perithecia provided a specific ecological niche for bacteria that could play an important role in disease establishment.

Over 2,000 [bacterial strains](#) were isolated from the native microbiota of *F. graminearum*, and subsequently identified and screened for antagonistic activity. The researchers successfully recovered 113 isolates that showed antagonistic activity against the pathogen. One specific bacterial isolate, termed as ZJU23, was identified as *Pantoea agglomerans*, and had the strongest inhibitory ability against *F. graminearum*. Subsequently, they found that herbicolin A secreted by ZJU23 was responsible for the observed suppression of *F. graminearum*.

Although herbicolin A was identified about four decades ago, its biosynthetic gene cluster and mode of action against [fungi](#) were not known. In this study, the researchers set out to investigate the biosynthetic gene cluster and mode of action against various fungi by combining various approaches, including transposon mutagenesis, liquid chromatography-mass spectrometry, atomic force microscopy and confocal microscopy.

By comparing the metabolic profiles of ZJU23 and four deletion mutants of a potential biosynthesis gene cluster, they found that herbicolin A was synthesized by the *AcbA-AcbJ* cluster. They then proceeded to uncover the mode of action of herbicolin A against various fungi.

It is important to note that the modes of action of cyclic lipopeptides against fungi are mostly unknown. It's surprise that herbicolin A was shown to disrupt [lipid rafts](#) by interacting with ergosterol, which resulted

in the formation of abnormal cell membranes, and, ultimately, caused cell death.

In addition, herbicolin A was found to inhibit the growth of *Candida albicans* and *Aspergillus fumigatus*, and was more effective than the clinical fungicides Amphotericin B and fluconazole. This provided important evidence for potential applications of herbicolin A beyond agriculture, for example in medicine.

Overall, the researchers have deciphered the mechanism of the inhibitory effect of herbicolin A on fungi and identified its biosynthetic gene [cluster](#). This could support future developments for sustainable management of Fusarium head blight worldwide.

This work, titled "Fusarium fruiting body microbiome member *Pantoea agglomerans* inhibits fungal pathogenesis by targeting lipid rafts," was published in *Nature Microbiology*.

**More information:** Sunde Xu et al, Fusarium fruiting body microbiome member *Pantoea agglomerans* inhibits fungal pathogenesis by targeting lipid rafts, *Nature Microbiology* (2022). [DOI: 10.1038/s41564-022-01131-x](#)

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