[Grant-in-Aid for Scientific Research(S)]

Biological Sciences (Agricultural sciences)



Title of Project: Mechanism of disease caused by deficiency of degradation of tRNA intron in mitochondria

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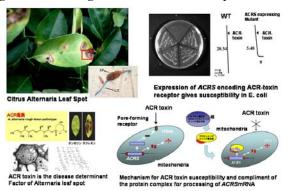
Research Area: Plant Pathology

Keyword: Gene, Plant, Mitochondria, Host-selective toxin

[Purpose and Background of the Research]

The host specificity of the rough lemon and tangerine pathotypes of *Alternaria alternata* depends upon the production of host-selective toxins (HSTs). The toxin from the rough lemon pathotype was named ACR-toxin I (MW 496).

ACR-toxin causes uncoupling similar to classic protonophores with a loss of membrane potential causing a leakage of co-factor, NAD+ from the TCA cycle. We identified the gene (ACRS) confers ACR-toxin sensitivity, and also identified that the mechanism of specificity in plants is alternative transcript processing of the gene conferring ACR-toxin sensitivity



In this study, promoter analysis of gene encoding *ACRS*mRNA binding 30kD protein, identification of components of *ACRS*mRNA processing protein complex, and genomics of the gene cluster of ACR- and ACT-toxin biosynthesis will be performed for elucidation of specificity in plant and microbe interactions.

[Research Methods]

Transcript of gene encoding *ACRS*mRNA binding 30kD protein can be detected in mRNA from ACR-toxin-insensitive cultivars but not from the toxin sensitive cultivar, and thus promoter analysis of this gene in different cultivars will be performed.

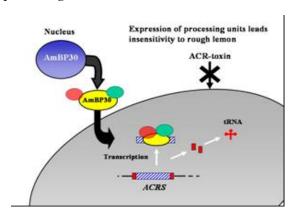
The 30kD protein is considered to be a recognition component of the processing protein complex, and thus identification of the entire complex will be performed.

Conditionally dispensable chromosome

carrying biosynthesis gene clusters for ACR- and ACT-toxin will be also sequenced.

[Expected Research Achievements and Scientific Significance]

Research purpose of this study is that elucidation of specificity by identification of *ACRS*mRNA processing protein complex as well as gene clusters responsible for the biosynthesis of ACR- and ACT-toxins. Addition of *ACRS*mRNA processing protein complex in the toxin sensitive cultivar might lead resistance against both the toxin and the pathogen producing the toxin in future.



[Publications Relevant to the Project]

- · Ohtani K, Yamamoto H, Akimitsu K. (2002) Sensitivity to Alternaria alternata toxin in citrus because of altered mitochondrial RNA processing. Proc Natl Acad Sci USA. 99:2439-2444.
- Miyamoto, Y., et al. (2008) Functional Analysis of a Multicopy Host-Selective ACT-Toxin Biosynthesis Gene in the Tangerine Pathotype of *Alternaria alternata*Using RNA Silencing. Mol Plant Microbe Interac 21:1591-1599.

[Term of Project] FY2009 - 2013

[Budget Allocation] 82,200 Thousand Yen

[Homepage Address and Other Contact Information]

http://www.ag.kagawa-u.ac.jp/plantpathology/index.html