Molecular profiling of the sugar beet pathogen Fusarium secorum

Subidhya Shrestha ^{a,b}, Roshan Sharma Poudel ^a, Ronnie de Jonge ^c, Kimberly Webb ^d, Gary Secor ^a, Melvin D. Bolton^{a,b,†}

^a Department of Plant Pathology, North Dakota State University, Fargo, ND 58102

^bU.S. Dept. Agriculture – Agricultural Research Service, Northern Crop Science Laboratory,

Fargo, ND 58102

^c Plant-Microbe Interactions, Department of Biology, Science4Life, Utrecht University, Utrecht,

NL

^d U. S. Dept. of Agriculture – Agricultural Research Service, Soil Management and Sugar Beet

Research Unit, Fort Collins, CO 80523

[†] Corresponding author: M. D. Bolton; <u>melvin.bolton@usda.gov</u>

Fusarium yellowing decline caused by the fungal pathogen *Fusarium secorum* has become an emerging problem in the sugar beet industry. Since *F. secorum* is not closely related to other *Fusarium* pathogens of sugar beet, little is known of the virulence mechanisms of this pathogen. To that end, we utilized whole-genome sequencing of the pathogen, xylem sap mass spectrometry, and transcriptome analysis of *F. secorum* infected-sugar beet plants to understand the molecular basis of virulence of this pathogen. *Fusarium secorum* showed an increased genome size due to an increased number of introns and repetitive elements. Additionally, we successfully developed a CRISPR-Cas9 ribonucleoprotein mediated gene-editing technique to disrupt target genes encoding candidate effector proteins in the pathogen. There were five gene targets, and among them, one gene target (*Fsec2*) was identified as a virulence factor of *F. secorum*. This study provides valuable genomic resources and a better understanding of the virulence strategies of an important pathogen of sugar beet.