New Disease Reports

First report of strobilurin resistance in *Cercospora* beticola in sugar beet (*Beta vulgaris*) in Michigan and Nebraska, USA

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Cercospora leaf spot (CLS) caused by Cercospora beticola Sacc. is the most important foliar disease of sugar beet (Beta vulgaris) worldwide (Jacobsen & Franc, 2009). CLS is controlled mainly with fungicides, including strobilurins (FRAC group 11). Resistance to strobilurins in C. beticola has not been reported in the field (Secor et al., 2010) but insensitive mutations have been artificially developed (Malandrakis et al., 2011). In 2011, fields from several areas in Michigan, USA treated with strobilurins had severe CLS and diminished control was also noted in small plot trials (Fig. 1). Individual leaf spot lesions were sampled from leaves and grown on sugar beet leaf extract agar (SBLEA). A conidium germination bioassay was done on SBLEA covered with water agar amended with pyraclostrobin, azoxystrobin or trifloxystrobin at 0, 0.001, 0.01, 0.1, 1, 10, or 100 $\mu\text{g/ml},$ supplemented with salicylhydroxamic acid (SHAM) to block the alternate oxidation pathway (Olaya et al., 1998). After 24 h incubation at 22°C, under ambient light, percentage germinated conidia (n = 50) was calculated from three replicates per treatment. Germination was recorded as positive when the germ tube was at least half the length of the conidium. A representative wild type isolate was unable to germinate over the 0.01 μ g/ml concentration. EC50 values for each isolate were calculated by regression analysis of percentage growth inhibition vs. the log fungicide concentration using Sigmaplot Version 9.01 (Systat Software, Chicago). The EC50 for the sensitive isolate was <0.01 µg/ml. Isolates from several counties in Michigan had uninhibited germination and EC50 values exceeded the highest concentration tested. Isolates also grew on spiral gradient dilution plates (Förster et al., 2004) amended with the three strobilurins (Fig. 2, for illustration of resistance response only). Two isolates were obtained from Nebraska and each showed similar response to strobilurin fungicides in amended plate assays.

Pure cultures of four resistant isolates were grown in potato dextrose broth at 125 rpm, and DNA extracted. A fragment of the cytochrome b (*CYTB*) gene was amplified by PCR using the *C. beticola* primers of Malandrakis *et al.* (2011) to amplify the region of the *CYTB* gene likely to contain resistance mutations (Malandrakis *et al.*, 2011). This fragment was sequenced at the Genomics Technology Support Facility (MSU, East Lansing, MI) and showed 99% identity with both the *C. beticola* cytochrome b mRNA, partial sequence (GenBank Accession No. EF176921.1) and the *C. kikuchii* mitochondrial gene for cytochrome b partial sequence (AB231863.1). Sequence results revealed that each resistant isolate contained a change in codon 143 that predicts to a



substitution of G143A, which was demonstrated to confer QoI resistance in several other fungi (Ma & Michailides, 2005). All four isolates with the G143A mutation germinated at 100 μ g/ml pyraclostrobin (50% of conidia), while sensitive isolates that lacked the mutation failed to grow. Isolates that contained the G143A mutation included representatives from Michigan and Nebraska, USA. These findings reveal that reduced Cercospora leaf spot control in some commercial sugar beet fields may be due to the development of resistance to strobilurins.

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References

Förster H, Kanetis L, Adaskaveg JE, 2004. Spiral gradient dilution, a rapid method for determining growth responses and 50% effective concentration values in fungus-fungicide interactions. *Phytopathology* **94**,163-170. [http://dx.doi.org/10.1094/PHYTO.2004.94.2.163]

Jacobsen BJ, Franc GD, 2009. Cercospora leaf spot. In: Harveson RM, Hanson LE, Hein GL, eds. *Compendium of Beet Diseases and Pests*, 2nd edn. St. Paul, MN, USA: American Phytopathological Society, 7-10.

Ma Z, Michailides TJ, 2005. Advances in understanding molecular mechanisms of fungicide resistance and molecular detection of resistant genotypes in phytopathogenic fungi. *Crop Protection* **24**, 853-863. [http://dx.doi.org/10.1016/j.cropro.2005.01.011]

Malandrakis AA, Markoglou AN, Nikou DC, Vontas JG, Ziogas BN, 2011. Molecular diagnostic for detecting the cytochrome b G143S - QoI resistance mutation in *Cercospora beticola*. *Pesticide Biochemistry and Physiology*, 87-92. [http://dx.doi.org/10.1016/j.pestbp.2011.02011]

Olaya G, Zheng D, Köller W, 1998. Differential responses of germinating *Venturia inaequalis* conidia to kresoxim-methyl. *Pesticide Science* **54**, 230-236.

[http://dx.doi.org/10.1002/(SICI)1096-9063(1998110)54:33.0.CO;2-O]

Secor GA, Rivera VV, Khan MFR, Gudmestad NC, 2010. Monitoring fungicide sensitivity of *Cercospora beticola* of sugar beet for disease management decisions. *Plant Disease* **94**, 1272-1282. [http://dx.doi.org/10.1094/PDIS-07-09-0471]



Figure 1

Figure 2

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