Quantification of the *Phytophthora infestans* population densities in upland soils in Japan using real-time PCR

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To estimate the *Phytophthora infestans* population density in soil using real-time PCR, we used a modified CTAB-beadbeating method to extract DNA from upland soils in Japan. The quantity data obtained using real-time PCR were compatible with the symptom development in a non-control potato field. Furthermore, there was a correlation between the quantity of *P. infestans* DNA measured by real-time PCR and the inoculum potential in soil. Therefore, the quantification of the population density of *P. infestans* using real-time PCR may be a guide to preventing potato storage rot.

Objectives

Result 1



Tuber rot can be minimized by decreasing harvesting injuries and/or the population density of *P. infestans* in soil [1]. However, it is impossible to harvest tubers without injury. Thus, we developed the real-time PCR assay to estimate *P. infestans* densities in soils and investigated the relationship between DNA quantity and inoculum potential.

Materials & Methods

Sample soils



The quantity data were compatible with the symptom development.





Extraction & Real-time PCR



Templates were analyzed using real-time PCR system

System: StepOnePlus (Applied biosystems) Primers, Probe: PinfTQF/PinfTQR, PinfTQPR [2]

Inoculum potential



Method: Sato's test [3] Inoculum potential (%) $= \frac{\text{Rotten tuber pieces}}{\text{Total tuber pieces}} \times 100$

Result 2



between the quantity of *P. infestans* DNA and the inoculum potential.

Conclusion & Future work

Quantification of the DNA in soil may be a guide to preventing storage rot. Another factor also should be investigated how much it affects tuber rot.

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