Comparative genomics of *Spongospora subterranea* f. sp. *subterranea* isolates for effector mining

<u>Maria de la O Leyva-Pérez¹</u>, Jack H.Vossen², Dan Milbourne¹, Denis Griffin¹, Marga van Gent-Pelzer³, Ueli Merz⁴, Leah Tsror⁵, Jacquie van der Waals⁶, Rudolph Strydom⁶, Shay Phelan¹, Colm McDonnell⁷, Kiwamu Tanaka⁸, Natalia Moroz⁸, Yuan Zeng⁹, Ana Cristina Fulladolsa¹⁰, Brian Alexander Charlton¹¹, Nicole Rabbiosi¹¹, Ronald Hutten², Calum Wilson¹², Ivette Acuña¹³, Camila Sandoval¹³, and Theo van der Lee³

¹ Teagasc, Ireland

² Plant Breeding, Wageningen University and Research, The Netherlands

- ³ Biointeractions and Plant Health, Wageningen University and Research, The Netherlands
- ⁴ ETH Zurich, Switzerland
- ⁵ ARO, Volcani Center, Israel
- ⁶ University of Pretoria, South Africa
- ⁷ IPM Potato Group Limited, Ireland
- ⁸ Washington State University, United States of America
- ⁹ Virginia Tech, United States of America
- ¹⁰ Colorado State University, United States of America
- ¹¹ Oregon State University, United States of America
- ¹² Tasmanian Institute of Agriculture (TIA). University of Tasmania, Australia
- ¹³ Institute of Agricultural Research (INIA), Chile

Breeding potato varieties resistant to Spongospora subterranea f. sp. subterranea (Sss) is impaired by phenotyping methods, which can give variable results due to environmental conditions, inoculum instability, and isolate-specific tolerance/resistance in host lines. Screening based on effector response ("effectoromics") could replace difficult phenotyping assays, and has been used to identify R genes in potato that confer resistance to Phytophthora infestans and Synchytrium endobioticum. Effectoromics relies on potential effectors identified in predicted secretomes. For Sss. however, no specific effector signatures are known. This study aims to identify new effectors and corresponding signatures, and select secretome genes which could have resulted from previous Solanum/Sss interactions and are under diversified selection pressure. To capture the required genomic diversity for selection, Sss genomic DNA was prepared from sporosori from more than 20 different isolates originating from 15 countries across five continents. These samples were assessed for Sss DNA proportions using Tagman-qPCR, and were subjected to Illumina paired-end sequencing. Initial comparison of the isolates showed at least two groups, one being very different from the current reference genome (SSUBK13), as indicated by regions that were either highly polymorphic or absent in this group when mapped to the reference. To help assemble polymorphic regions through *de novo* assemblies, two isolates representing the two groups were selected for long-read Nanopore sequencing. Characteristics of their assemblies will be described, including insights into genome-wide diversity. There is potential to develop an Sss pangenome knowledge base as a resource for plant pathology and potato breeding research.