

Exploring genome diversity of 6 strains of the model arbuscular mycorrhizal symbiont Rhizophagus irregularis

Emmanuelle Morin, Eric C H Chen, Alan Kuo, Denis Beaudet, Igor V Grigoriev, Bernard Henrissat, Christophe Roux, Nicolas Corradi, Francis M Martin

▶ To cite this version:

Emmanuelle Morin, Eric C H Chen, Alan Kuo, Denis Beaudet, Igor V Grigoriev, et al.. Exploring genome diversity of 6 strains of the model arbuscular mycorrhizal symbiont Rhizophagus irregularis. JOBIM, Jul 2018, Marseille, France. hal-03324326

HAL Id: hal-03324326 https://hal.science/hal-03324326v1

Submitted on 23 Aug 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Aix*Marseille

Exploring genome diversity of 6 strains of the model arbuscular mycorrhizal symbiont Rhizophagus irregularis

Emmanuelle Morin¹, Eric C.H. Chen², Alan Kuo³, Denis Beaudet², Igor V. Grigoriev³, Bernard Henrissat⁴, Christophe Roux⁵, Nicolas Corradi² and Francis M. Martin¹



1 : Institut National de la Recherche Agronomique (INRA), Unité Mixte de Recherche 1136 Interactions Arbres/ Microorganismes, Laboratoire D'excellence ARBRE, Centre INRA-Grand Est-Nancy, Champenoux 54280, France

2 : Department of Biology, University of Ottawa, Ottawa, ON K1N9A7, Canada

3 : US Department of Energy Joint Genome Institute, Walnut Creek, CA 94598, USA

4 : Architecture et Fonction des Macromolecules Biologiques, CNRS, Aix-Marseille Université, Marseille 13288, France 5 : Laboratoire de Recherche en Sciences Végétales, UPS, CNRS 24 Chemin de Borde Rouge-Auzeville, Université de Toulouse Castanet-Tolosan 31326, France





- fungal transporters in the extr mycelium (red or green symbol - transport through the hyphae

Background and data

Arbuscular mycorrhizal fungi (AMF) are members of the subphylum Glomeromycotina and are involved in a widespread symbiosis with the roots of most land plants, AMF are known to improve plant fitness through the establishment of mycorrhizal symbioses. Genetic and phenotypic variations among closely related AMF isolates can significantly affect plant growth, but are unclear.

To elucidate the genomic changes underlying this variability, we improved the genome assembly and gene annotation of the model strain Rhizophagus irregularis DAOM197198, and compared its gene content with five isolates of R. irregularis sampled in the same field.

Genome annotation and portals

JGI Annotation Pipeline:

- integrates an array of tools for gene prediction : protein-coding gene models are predicted using a combination of gene predictors of three general kinds:
 - ab initio : Fgenesh and GeneMark
 - homology-based : Fgenesh+ and Genewise
 - transcriptome-based : models are predicted from assembled RNAsequencing data derived from the cultured fungus
- for each genomic locus, the best representative gene model is selected based on a combination of protein homology and transcriptome support
- all predicted proteins are then functionally annotated :
 - protein alignments to the NCBI's NR, Swiss-Prot, KEGG and KOG DBs detection of sequence motifs responsible for protein localization : signal P and TMHMM
 - prediction of functional domains from Pfam and other databases : InterProScan
 - expert annotation of different categories of proteins : CAZymes, proteases (MEROPS), secondary metabolism proteins JGI 🕅



For this study, we generated a portal for each strain, which provides different tools to browse, annotate and compare

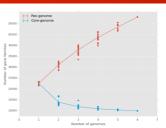


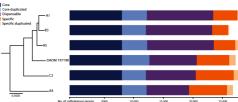


Pan and core genome curves

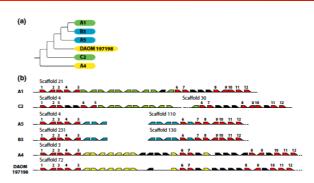
Prediction of secreted proteins was performed using combined characteristics: protein structure : presence of a signal peptide detected with SignalP and no

- transmembrane helix or one overlapping the signal peptide was found by TMHMM
- protein subcellular localization : subcellular localization assigned as





Intraspecific genome diversity



A divergent region between the Rhizophagus irregularis isolates. (a) Phylogenetic relationships of the isolates based on rRNA sequences, mitochondrial genes, and most orthologous nuclear genes.

(b) Gene order and sequence comparison along a 67 kb syntenic region present in all isolates investigated. The red color represents flanking syntenic genes that extend beyond what is presented in the figure, with the number indicating which genes are orthologs to each other; black represents genes showing no evidence of synteny; other colors represent the sequence conservation and synteny unique to pairs of isolates.

Conclusion

All isolates harbor striking genome variations, with large numbers of isolate-specific genes, gene family expansions, and evidence of interisolate genetic exchange Overall, these findings indicate a substantial divergence in the functioning capacity of isolates harvested from the same field, and thus their genetic potential for adaptation to biotic and abiotic changes. Our data also provide a first glimpse into the genome diversity that resides within natural populations of these symbionts.

Chen E.C.H, Morin E. et al. High intraspecific genome diversity in the model arbuscular mycorrhizal symbiont Rhizophagus irregularis (2018) New Phytologist, doi: 10.1111/nph.14989



This work was also supported by the Laboratory of Excellence ARBRE (ANR-11-LABX-0002-01),

secretory pathway using TargetP and as