

Book of Abstracts

Oral & Poster Abstracts







ACKNOWLEDGEMENT

WITH THANKS

























CONTENTS

- 1. Welcome
- 2. Scientific Committee
- 3. <u>Local Organising Committee</u>
- 4. <u>Scientific Programme</u>
- 5. Oral Abstracts
 - a. Opening Event L1 L2
 - b. <u>Topic A: Environmental interactions L3 L12</u>
 - c. Topic B: Grapevine Development L 13 L22
 - d. Topic C: Fruit Composition L23 L33
 - e. Topic D: Computational Resources and Data Integration L34 L38
 - f. Topic E: Technological Advances L39 L46
- 6. Poster Abstracts

Poster Index

- a. Topic A: Environmental interactions P1 P33
- b. Topic B: Grapevine Development P34 P59a
- c. Topic C: Fruit Composition P60 P70
- d. Topic D: Computational Resources and Data Integration P71 P76
- e. Topic E: Technological Advances P77 P99
- 7. <u>Integrape Workshop (COST Action CA17111)</u>
 - a. Data Integration, The Power of Omics & Grape Improvement
- 8. <u>List of Participants</u>





NEXT





L41

Whole genome resequencing and custom genotyping unveil clonal lineages in `Malbec´ grapevines (Vitis vinifera L.)

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Grapevine cultivars are clonally propagated to preserve their varietal attributes. However, genetic variations accumulate due to the occurrence of somatic mutations. This process is anthropically influenced through plant transportation, clonal propagation and selection. Malbec is a cultivar that is well-appreciated for the elaboration of red wine. It originated in Southwestern France and was introduced in Argentina during the 1850s. In order to study the clonal genetic diversity of Malbec grapevines, we generated whole-genome resequencing data for four accessions with different clonal propagation records. A stringent variant calling procedure was established to identify reliable polymorphisms among the analyzed accessions. The latter procedure retrieved 941 single nucleotide variants (SNVs). A reduced set of the detected SNVs was corroborated through Sanger sequencing and employed to custom-design a genotyping experiment. We successfully genotyped 214 Malbec accessions using 41 SNVs and identified 14 genotypes that clustered in two genetically divergent clonal lineages. These lineages were associated with the time span of clonal propagation of the analyzed accessions in Argentina and Europe. Our results show the usefulness of this approach for the study of the scarce intra-cultivar genetic diversity in grapevines. We also provide evidence on how human actions might have driven the accumulation of different somatic mutations, ultimately shaping the Malbec genetic diversity pattern.

Keywords: Malbec, genetic diversity, clonal lineages, propagation history, whole genome resequencing







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