Inhibition of *Oenococcus oeni* during alcoholic fermentation by a selected Lactiplantibacillus plantarum strain

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Introduction

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The use of selected cultures of the species Lactiplantibacillus plantarum in Oenology has grown in prominence in recent years. While initial applications of this species centered around malolactic fermentation (MLF), it has been suggested that some strains can be harnessed for their bio-protective effects (Rubio-Bretón et al 2018). As little scientific literature exists on biological protection from this species in winemaking, the following work set out to investigate the effects of Lactiplantibacillus plantarum DSM27565 (trade name Viniflora[®] NoVA[™] Protect) on adventitious Oenococcus oeni, which can lead to unwanted MLF during alcoholic fermentation.

Results Methodology Samples of each vessel were taken and concentrations (copies/ml) of DSM27565 Crush grapes Add 25mg/L SO₂ (as PMS) Ensure YAN >200mg/L and *Oenococcus oeni* were measured by qPCR (Fig. 2). 1: + TA 3: - TA 4: - TA · DSM2756 - DSM27565 + DSM27565 + DSM27565 💙 Viniflora[®] NoVA[™] Protect concentration in wine Oenococcus oeni concentration in wine Add Tartaric acid 2-4g/L to achieve must pH of 3.50 No Acid Addition July Day 0 (PRE) Day 0 (POST) Day 2 EAF (Day 12) ■ Day 0 (PRE) ■ Day 0 (POST) ■ Day 2 ■ EAF (Day 12) Leave 2hrs to disperse then mix well 2,0 E+06 3.9 E+05 Inoculate Inoculate 3.6 E+05 1,8 E+06 DSM27565 DSM27565 No Inoculation No Inoculation mix to disperse mix to disperse S 3.3 E+05 1,6 E+06 \mathbf{V} 3.0 E+05 Maintain each vessel at ambient temperature for 24 hours 1,4 E+06 2.7 E+05 $\bigcirc \bigcirc$ 1 12 J.

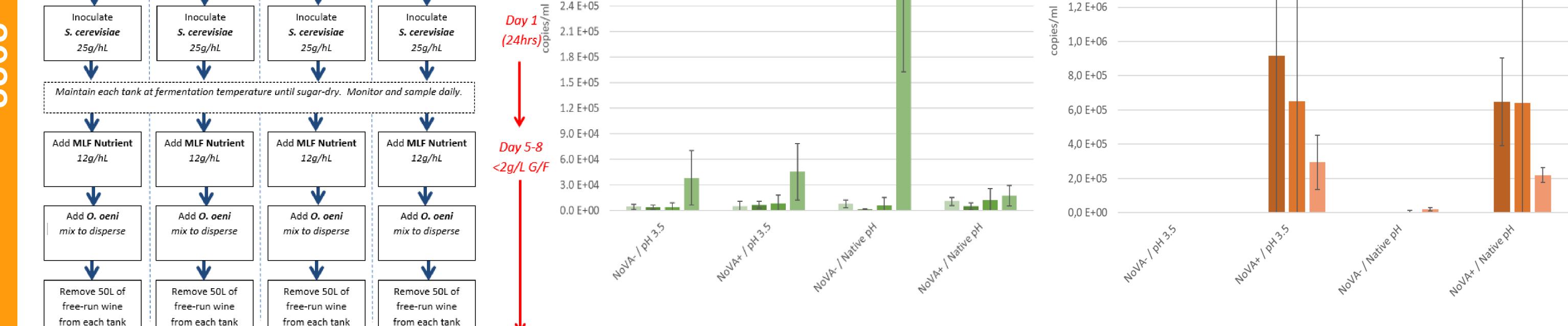


Figure 1. Winemaking Methodology.

1700kg of Cabernet Sauvignon (Grampians Region, Australia) was crushed and evenly distributed across eight fermentation vessels. Treatments were dosed with either 2g/L Tartaric Acid and/or DSM27565 at 2.0E+6 CFU/mL. The must had a native pH of 3.70 and 25mg/L of SO₂ was added at crushing.

Figure 2. Quantification of *O. oeni* and Viniflora[®] NoVA[™] Protect by qPCR. DNA was extracted from cells obtained by centrifugation from the grape must and used for quantification of the total O. oeni population and Viniflora[®] NoVA[™] Protect by qPCR. The qPCR results are shown as average of genome copies per ml of wine (copies/ml) calculated between the replicates of each treatment and the error bars represent the standard deviation between replicates (the DNA was extracted and quantified separately for each replicate). The different stages of alcoholic fermentation (AF) are indicated as: Day 0 (PRE) = beginning of AF before NoVA addition; Day 0 (POST) = beginning of AF after NoVA addition; Day 2 = two days after start of AF; EAF (Day 12) = End of AF (12 days after AF start).

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Conclusions

- Lactiplantibacillus plantarum DSM27565 implants well into red wine grape must, even with moderate levels of SO₂ and dominates red grape must at inoculation
- Excessive growth of adventitious *Oenococcus oeni* is suppressed in the presence of DSM27565
- Using Lactiplantibacillus plantarum DSM27565 shows an alternative to using tartaric acid as a biological control agent
- Further work is suggested to investigate the effects of DSM27565 on other spoilage organisms such as Acetobacter spp., and Hanseniaspora uvarum

References

Rubio-Bretón, P., Gonzalo-Diago, A., Iribarren, M., Garde-Cerdán, T., Pérez-Alvarez, E.P., 2018. Bioprotection as a tool to free additives winemaking: effect on sensorial, anthocyanic and aromatic profile of young red wines. LWT- Food Sci. Technol. 98,458–464.