

LALVIN[®] ICV OKAY[®]

ORIGIN AND APPLICATION

For young fresh and aromatic rosé, white and red wines. Lalvin ICV OKAY® offers fermentation security whilst producing very low levels of acetaldehyde, H₂S and SO₂.

The development of **Lalvin ICV OKAY**[®] was associated with a PhD related to the identification of a new mechanism responsible for the control of SO₂ and H₂S production by wine yeast . A genetic study identified the molecular basis of these properties. Using marker-assisted selection techniques, Lallemand and ICV have selected, with the collaboration of INRAE and Sup' Agro Montpellier (France), **Lalvin ICV OKAY**[®].

Lalvin ICV OKAY® exhibits a special ability to produce very low levels of H₂S and SO₂. Moreover, the low acetaldehyde production of **Lalvin ICV OKAY®** will be a good asset to stabilize most wines with moderate SO₂ levels. This yeast also offers fermentation security, completing fermentation in a large range of fermentation conditions. Tends to contribute good fruit intensity.

MICROBIAL AND OENOLOGICAL PROPERTIES

- Recommended for white, rosé and red wine production.
- Saccharomyces cerevisiae
- Alcohol tolerance to 16% v/v *subject to fermentation conditions.
- Temperature tolerance 12°C 30°C
- Low relative nitrogen demand.
- Short lag phase with steady fermentation kinetics.
- Very low potential for SO₂ production.
- Very low acetaldehyde production.
- Very low H₂S production.
- Killer factor active.
- Compatible with malolactic fermentation.
- Low foam producer.























FURTHER READING (Please request this booklet from your Lallemand representative).

Lallemand Winemaking Update – Number 1 2008: 'The YSEO® Process'

Evaluation of the YSEO® Process to prepare dried winemaking yeast – Summary of a study done by Washington State University and Lallemand.

Lallemand FOCUS paper : Yeast options for fruit wine and cider making.

INSTRUCTION FOR USE

Dosage Rate:

- 25g/hL of Active Dried Yeast (this will provide an initial cell population of approximately 5 x10⁶ viable cells/mL)
- 30g/hL of Go-Ferm Protect® / Go-Ferm Protect Evolution™
- Nitrogen source from the Fermaid range

Procedure for 1000L ferment.

- 1) Add 300g of Go-Ferm Protect[®] / Go-Ferm Protect Evolution[™] to 5L of 40-43°C clean, chlorine free water. Stir until an homogenous suspension free of lumps is achieved.
- 2) When the temperature of this suspension is between 35-40°C, sprinkle 250g of yeast slowly and evenly onto the surface of the water, whilst gently stirring. Ensure any clumps are dispersed.
- 3) Allow to stand for 20 minutes before further gently mixing.
- A) Mix the rehydrated yeast with a little juice, gradually adjusting the yeast suspension temperature to within 5-10°C of the juice/must temperature.
- 5) Inoculate into the must.

Further Notes

- Steps 1-5 should be completed within 30 minutes.
- It is best to limit first juice/must volume addition to one tenth the yeast suspension volume and wait 10 minutes before the addition to juice.
- To minimize cold shock, ensure temperature changes are less than 10°C.
- It is recommended that juice / must be inoculated no lower than 18°C.
- It is recommended to use complex nutrition nitrogen source, such as either **Fermaid AT™** or **Fermaid O™**.

PACKAGING AND STORAGE

• All Active Dried Yeast should be stored dry, best pratice between 4-12°C and the vacuum packaging should remain intact.

of technological properties of wine yeast (Jessica Noble, Advisor: Bruno Blondin, 2011). This work resulted in a patent application filed by INRA and Montpellier SupAgro: «Method of control on the production of sulfites, hydrogen sulfur and acetaldehyde by yeast (Variants MET₂ / SKP₂) «This approach has enabled the development of an innovative selection technique for yeast which produces low levels of S0₂, H₂S and acetaldehyde. «

The selection of these yeasts was largely made possible through a collaborative study between the ICV Group, Lallemand Oenology, SupAgro and INRA Montpellier. This study, using the QTL technique (Quantitative Trait Locus), was used during the thesis: Identification of the molecular basis

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