AVOIDING STUCK AND SLUGGISH FERMENTATIONS

PRIMARY CAUSES:

Nutrient deficiency (nitrogen, vitamins, minerals)

Competition from wild yeast, bacteria

Lack of survival factors (long-chain fatty acids, sterols)

Ethanol

Temperature

Osmotic shock

Toxins (medium-chain fatty acids, high volatile acidity, high SO₂)

Inadequate yeast population (low inoculation rate)

Highly clarified must

Pesticide residue

1. Nutrient deficiency

Nitrogen (ammonia and amino acid) deficiency.

Slows protein synthesis.

Diminishes Sugar transport activity.

Becomes limiting at <150 ppm initial level (generalization; depends on mix of amino acids and ammonia plus other growth factors).

N addition not related to kinetics at end of fermentation (may still stick w/late addition).

Above 10% alcohol, yeast will not take up N.

Ammonia N (DAP) vs. amino N (organic N).

Vitamin/mineral deficiency

Thiamin: increases viable yeast population, speeds fermentation; deficiency leads to acetic acid production.

Pantothenic acid: deficiency leads to acetic acid and H_2S production, even if N is adequate. 50 mg/hL limits acetic acid in stuck fermentations.

Biotin: aids viability at end of fermentation.

Inositol: aids membrane phospholipid synthesis.

Other growth factors: pyridoxine, nicotinamide, riboflavin (all found in dry yeast, inactivated yeast, etc.).

Minerals: Potassium, magnesium, phosphorus, zinc.

Oxygen

Yeast needs oxygen during growth.

Creates survival factors (sterols and fatty acids).

Increases effectiveness of N addition.

Action:

Monitor Yeast Available Nitrogen (YAN).

Use complex nutrients after lag phase and during growth phase (1/3 fermentation).

Augment with DAP if N is very low or sugar is high.

Aeration during growth phase.

2. Competition (wild yeast, fungi/mold and bacteria)

Depletes nutrients

Wild yeast can deplete thiamin within 3 hours.

Botrytis may decrease amino acid content 7-61%.

Action:

Avoid cold soak if microbial load is high.

If cold soak is used, add 50% of the yeast to the must at beginning of the cold soak.

Use sufficient SO₂ to knock down wild microbes.

In whites, if microbial load is high, clarify juice (boost turbidity afterwards with cellulose powder and/or inactivated yeast).

If competition is from lactic acid bacteria, lysozyme can be used to kill them.

If end is sluggish, low SO₂ addition will reduce competition without hurting yeast.

3. Sterols/fatty acids (survival factors) deficiency

Sterols and long-chain unsaturated FA's maintain membrane integrity.

Enables yeast cell to expel ethanol (EtOH).

EtOH can alter membrane, allowing entry of protons, which acidify cell, causing death.

Action:

Use active dry yeast (most are selected for high sterol and FA content).

Rehydrate yeast with rehydration nutrient high in survival factors.

Aeration during growth phase of yeast.

Supplement at 1/3 fermentation with complete nutrient containing survival factors.

4. Osmotic shock (high sugar)

High sugar must creates pressure across the yeast membrane from the cell toward the must. If pressure is high enough, yeast will die.

Action:

Use yeast selected to withstand osmotic stress.

Reduce sugar level by addition of water or low Brix must.

5. Ethanol

Alters cell membrane, allowing greater entry of protons, acidifying the cells; inhibits nutrient transport.

Toxic effects are exacerbated by high temperatures.

Action:

Pick grapes at a reasonable Brix level.

Use yeasts pre-conditioned for high alcohol

One can use weaker yeast at start of fermentation, adding the stronger yeast part way through the fermentation.

Build up sterols and FA's (above) to withstand the high alcohol.

Ferment at lower temperatures.

Increase yeast addition rate to accommodate potential alcohol.

6. Temperature

Above 90° F, puts high stress on yeast (stress increases at higher alcohol).

High temperatures (77-90° F, measured under cap) during exponential yeast growth favor acetic acid production.

Too low temperatures, yeast may be overly slow or stop fermenting and may form H₂S.

Action:

Control temperature (\sim 70-75° F during growth), especially in high Brix must.

The higher the potential alcohol, the lower the temperature should be.

For low temperature white fermentations, have sufficient turbidity or stirring/agitation to keep yeast in suspension.

7. Toxins (medium chain saturated FA's, volatile acidity, high SO₂)

Medium chain FA's created during fermentation may slow or stop fermentation.

Action:

Fining with yeast hulls or inactivated yeast to adsorb toxins (may need to reinoculate with yeast afterwards).

VA may be from bacteria (acetic and/or lactic acid bacteria) or wild yeasts, or by certain cultured yeasts under stress.

Conditions favoring VA production:

pH < 3.1

pH >4.0

Amino acid and vitamin deficiency.

High temperature during exponential growth of yeast.

Excessive must clarification.

Action:

Careful sorting in vineyard to reduce infected grapes.

Monitoring sanitary state of grapes in vineyard and winery.

Clarification of white juice to reduce microbial load (with some turbidity added back).

SO₂ addition as soon as possible.

Use lysozyme if problem is from lactic acid bacteria.

Small amounts of tainted wine can be incorporated into fresh must and the yeast will metabolize the VA.

Reduce other stresses on the yeast (nutrition, temperature, etc.)

High SO₂ may slow fermentation or kill yeasts.

In general, most cultivated yeasts can withstand moderately high SO₂.

Higher than 50 ppm SO₂ can deactivate thiamin.

Action:

Oxidation of must to reduce free SO₂ (not recommended).

Dilution with unsulfited juice (recommended).

8. Inadequate yeast population

Starting population should be $\geq 10^6$ cells per mL.

Recommendations of 20-30 g/hL (200-300 ppm) of active dry yeast are based on average European wines.

Action:

Increase inoculation rate to offset other stresses, particularly high potential alcohol..

9. Highly clarified must

Highly clarifying must can strip it of survival factors; may increase acetic acid.

Makes it harder to keep yeast in suspension.

Action:

Utilize rehydration nutrients and complex nutrients with survival factors.

Add cellulose powder, or cellulose and inactivated yeast.

Pre-fermentation, pre-settling skin contact can increase fermentability (with clean grapes)

10. Pesticide Residue

Certain chemicals, especially fungicides, applied too close to harvest, can interfere with yeast metabolism.

Action:

Apply pesticides to grapes only according to directions and extension recommendations. Some residues can be lowered by fining with inactivated yeast or yeast hulls.

GENERAL RECOMMENDATIONS:

Determine the source of the problem.

Test glucose/fructose, malic acid, volatile acidity, micro.

Stuck Fermentations (follow restart protocols: supplier websites, Scott Handbook, etc.):

Reds: Press must (helps lower bacteria population).

Light sulfiting to knock down competing microbes.

Fine with yeast hulls or inactivated yeast to remove toxins (20-30 g/hL).

Reinoculation with strong yeast at higher than normal levels per protocols.

Use rehydration nutrients and keep fermenting wine at 20-25° C.

Whites: Light sulfiting (20-30 ppm) when MLF is not desired.

Fine with yeast hulls or inactivated yeast.

Reinoculation with strong yeast at higher than normal levels per protocols.

Pro-Restart (alginate encapsulated yeast): pre-acclimated to alcoholic environment. Avoid serial inoculations:

After 5 generations, survival factors can be too depleted to complete fermentation.

ENOLOGICAL ENZYMES

Definition

Biological catalysts.

Naturally occurring vs. added.

Proper Use

Warmer temperature, higher activity. SO2 additions before or after, never with enzyme. Inactivation with bentonite.

White Wine Enzymes

Pectinases (Scottzyme Cinn-Free, Lallzyme C, Cuvee Blanc)
Break down pectin in flesh.
Increase juice extraction.
Release flavors and aromas.
Clarification and filterability.

Red Wine Enzymes

Pectinases with side activity (Scottzyme Color Pro, Lallzyme Ex, Color X)

Cellulase, hemicellulase, protease.

Break down structural elements in skins.

Side activity determines tannin extraction.

Fruit Wine

Concentrated pectinases, high levels of hemicellulase (Pec5L, Scottzyme HC, KS)

Pec5L + HC together break down most fruit.

High juice extraction from concentrated pectinases in Pec5L.

High hemicellulase activity for slimy grapes (Concord) and pomes (apple, pear) in HC.

KS—use only after solids are removed. Highest enzymatic activity.

Specialty Enzymes

Beta-Glycosidase (Scottzyme BG, Lallzyme Beta)
Cleaves glucoside bond to release terpenes.
Beta-Gluconase (Lallzyme MMX, VINStyle)
Breaks down glucans

Caution

Purified Enzymes—low levels of cinnamyl esterase.

Use benchtrials when using enzymes on finished wine, especially with B-glycosidase.