## What, Why \& When

A bench trial is a small-scale trial meant to simulate the addition of an additive or fining agent to a larger volume of wine.

The idea is that by trying an addition or fining out on a small scale, you can try a range of dosages, or even different products, without having to treat all of your wine. This allows you to accurately determine the exact process and dosage that will have the optimal impact on your wine allowing you to move forward and perform this process on your whole batch.
Ideally, we'd be doing a bench trial ahead of the addition of any product that has a dosage range rather than a fixed dosage that is appropriate. Indeed, this is most additions in winemaking. For example, most of us have seen how fining agents like bentonite come with a range of potential dosages on the package rather than a predetermined "correct" dosage level. You may have also read about how there is a danger of stripping away much of a wine's character when performing a fining, including bentonite. Wouldn't it be nice to know exactly how much bentonite will take care of your hazy wine, while having the least amount of impact on the flavor, color and aroma of your wine? Enter the bench trial.
Bench trials should be performed immediately ahead of a fining or addition procedure. Your wine is always changing. If you wait too long after a trial to make the actual addition or fining, the effect may be different from what you experienced in the trial itself. This means that you might schedule a bench trial for egg white fining a week ahead of the actual treatment, but plan on doing a TA addition trial just 2-3 days ahead of when you plan on making the addition to the whole batch of wine.

## How To

The bench trial process can be broken down into 6 basic steps: 1) Determining your sample volume, 2) Determining your range of dosages for the trial, 3) Scaling the dosages down to your sample size, 4) Creating a model solution, 5) Dosing the samples and 6) Evaluating the results. Let's go through these step by step.

1. Determine your sample volume: There is no perfectly correct sample volume for a trial. The correct volume depends on how much wine you have to work with, how small the dosages of the product you're trying are and how exact you can be with a pipet and scale. So, select a larger sample size if you have plenty of wine to
work with, if the additive or fining agent needs a very small dosage or if you feel that you may not be too accurate with measuring out the test dosages. Remember that the smaller the sample and/or dosage are, the more significant any small error on your part becomes. If you are trying to measure out 10 mL of liquid and are off by 1 mL , that is a $10 \%$ error - pretty significant. However, if you are measuring out 100 mL of liquid and are off by 1 mL , a $1 \%$ error is not such a concern. We recommend sample sizes anywhere from 50 mL up through 500 mL .
2. Determine your range of dosages: Most additives and fining agents have a recommended range of dosages for treating wines. For instance, MoreWine!'s unique, pre-soaked bentonite product Albumex can be added in a range between 1 and $3 \mathrm{~g} / \mathrm{L}$ of wine. Your first step is deciding how many samples you are going to run. We typically run 4 or 5 samples. Remember to always keep one untreated sample off to the side as a control. Also, it is a good idea to try to keep the step between the dosages uniform. If we were running a trial with Albumex, 5 samples and a control would be a good idea. The dosages for the samples would be $1 \mathrm{~g} / \mathrm{L}, 1.5 \mathrm{~g} / \mathrm{L}, 2 \mathrm{~g} / \mathrm{L}, 2.5$ $\mathrm{g} / \mathrm{L}$ and $3 \mathrm{~g} / \mathrm{L}$. You may have noticed that we are working with the metric system here. While it may take some getting used to, it's the best and easiest way to do this. If you're having trouble wrapping your head around it, we suggest that you print out our handy conversions sheet from the MoreManuals section of our website to use while you're working out your dosages.
3. Scale down the dosages to your sample size: As we mentioned in the previous step, the dosages that you choose will likely be in terms of grams/liter, or perhaps in oz / gallon, though metric units are recommended. No matter what units you use, it is unlikely that you'll be running the trial on samples as big as 1 liter (or 1 gallon). You'll need to do a little math to scale down the dosage to match your sample size. The basic idea here is that you ask yourself the following question: "If I want to achieve a dosage of $1 \mathrm{~g} / \mathrm{L}$, how much product do I weigh out for my 50 mL sample?" Since 50 mL is $5 \%$ of 1 L , you also need $5 \%$ of 1 g , which is 0.05 g . The easiest way to set up the math is as follows: (Dosage $) \times($ Conversion Factor $) \times($ Sample Size $)=($ Amt of Product for Sample). For example; (1g/L)x(1L/1000mL) $x(50 \mathrm{~mL})=0.05 \mathrm{~g}$. Notice how both the terms for $L$ and mL cancel out leaving only $g$ in the end. If you ever wind
up with a unit related to volume at the end of the equation then you know that you've made a mistake somewhere and need to go back to the start.
4. Create a model solution: Most of us don't have a scale that can weigh down to 0.05 g accurately. Even a scale that claims to have a resolution of 0.1 g will not weigh out accurately until you have at least 0.5 g on the scale, unless you are using a very advanced laboratory scale that costs thousands of dollars. How do we get the small amount of product that we need for the trial? The answer is to create a solution of the product you are trying and add a measured amount of it to each sample. This is also very straightforward. The first step is to take a look at the range of dosages: is the increment between each dose more or less than $50 \%$ of the first dosage? In our Albumex example it is exactly $50 \%$ : the first dose is $1 \mathrm{~g} / \mathrm{L}$ and the dose rises by 0.5 g at each step. When the increment between doses is $50 \%$ or more of the original dose, you want to set up the solution so that 1 mL of the solution is equivalent to the smallest dosage itself. For Albumex this means that you'd create a solution where 1 mL of the solution would add 0.05 g of Albumex to your 50 mL sample (equivalent to $1 \mathrm{~g} / \mathrm{L}$ addition). To make the second sample, where you need 0.075 g of Albumex ( $50 \%$ increase over the first dose, $1.5 \mathrm{~g} / \mathrm{L}$ ), you then just add 1.5 mL of your model solution. In order to create the model solution you must first determine how much of the product you are testing you wish to have contained in 1 mL of the model solution. In our case the answer is 0.05 g . Weigh out 1 g of your product and then add enough water to make up a final volume of 10 mL . Note that a graduated cylinder is the best tool for this. Now you have a model solution where each mL contains 0.1 g of Albumex: $(1 \mathrm{~g} / 10 \mathrm{~mL}) x(1 \mathrm{~mL})=0.1 \mathrm{~g}$. To reach your desired $0.05 \mathrm{~g} / \mathrm{mL}$ then simply cut this solution with another 10 mL of water, cutting the amount of Ambumex in each mL in half to 0.05 g . Now, If your interval between doses is less than $50 \%$ of the initial dose(dosages of $0.5 \mathrm{~g} / \mathrm{L}, 0.6 \mathrm{~g} / \mathrm{L}, 0.7 \mathrm{~g} / \mathrm{L}$, etc, for example) then it is best to make a solution where each mL will contain enough to make up the interval rather than enough to make the initial dosage.
5. Dose your samples: Now that you have made up your model solution it is time to add the doses to the samples. This is probably the easiest part of doing the bench trial. Simply add enough mLs of the model solution to each sample in order to achieve the dosage rate that you are looking for. In our example with Albumex, this would mean adding 1 mL to the first sample, 1.5 mL to the second, then $2 \mathrm{~mL}, 2.5$ and finally 3 mL to the last sample. The easiest way to do this is with a pipette. However, you must take care not to add too much model solution or you will have to discard the sample and start over. A good technique for this is to fill a pipette to a given level, then seal the end with your finger tip. Since you
have to push pretty hard to get a good seal, it is possible to allow liquid out of the pipette by simply reducing the pressure you're using to keep it sealed. You should not have to actually take your finger off the pipette in order to allow liquid to flow. Try practicing this with water a bit and you'll get the hang of it pretty quickly.

A word about sample bottles and their handling: All samples should be handled so as to minimize their contact with oxygen. Ideally, this would mean flushing out your sample bottles with $\mathrm{CO}_{2}$ or Argon gas before you fill them with wine. If you don't have access to inert gas for this purpose it is best to fill your sample container completely to the rim. Be aware than many flasks, bottles, etc are not designed to be filled 100\% completely. A 50 mL Erlenmeyer flask is a good example of this - with 50 mL in the flask the liquid does not come all the way up to the top, or even all the way to the bottom of a stopper. To account for this you can either A) determine what the "full" volume of your container is and adjust the rest of your math to this new sample volume, or B) prepare your samples in an even larger container and then fill the smaller container as much as you can. With method A you will get accurate results but the math may become complicated with lots of decimal points, etc. Method B should be pretty straightforward though: Following the example of our 50 mL flask, which holds some small amount more than 50 mL , simply adjust your sample size and model solution so as to make 100 mL samples instead - our $40 z(120 \mathrm{~mL})$ bottle works great for this. Make up a 100 mL sample then fill the 50 mL flask as much as you can and discard the rest of what you made. It represents such a small amount of wine and product cost that there should be no issues with this method.
6. Taste your samples: Now for the fun bit. After allowing enough time for the product you're testing to work, you want to come back and taste the samples to see which dosage (if any of them) you liked the best. It is a good idea to do this tasting with one or two other people there as well - many palates are better than just one.

Here's an important note: ideally you would be able to leave your samples for as long as it normally takes the product to work, but in many cases this is not possible - for additives that take several months to fully integrate into the wine try to wait at least 2 weeks before tasting your samples. It's also important to recognize that while a bench trial will give you a really good idea as to what is going to happen with your wine when you make the addition, real life works in such a way that the model is never exactly parallel to that which it is modeling. What I mean is, the results when you add the product to your whole volume of wine will almost always be somewhat different than what you experienced in the trial. We prefer in all applications a metered approach. Once you've decided on a dosage rate from the trial, add only $1 / 2$ to $2 / 3$ of the total addition to your wine. Wait the full period of time it takes for that product to be effective, then evaluate and see if you feel like you need the rest. We can assure you now that in some cases you will not.

## Tips and Tricks

Finally, here are a few general tips and tricks to keep in mind:

1. We'll keep saying it: get comfortable with the metric system. It makes all of the math for scaling up and down much easier, as everything works on the same base 10 system.
2. Invest in a decent scale that measures in grams and has a resolution down to 0.1 g . Our MT351A is a perfect choice.
3. Erlenmeyer flasks are not marked exactly and should not be trusted for exact measurements of volume. They typically have a $+5 \%$ error. Measuring your liquid with a pipette is always best. We have pipettes that measure up to 50 mL with very high accuracy.
4. A 50 mL or 100 mL graduated cylinder is pretty much required if you want to do this correctly. You may come up with another way to measure volume with a high degree of accuracy, but graduated cylinders are cheap and highly accurate.
5. Remember that accuracy is key here. Being off by 0.1 g in 100 mL is the same as being off by 10 g in 100 L . That can represent a pretty significant degree of error considering the dosages that we work with most of the time.
