

Customer Prepared Library Submission Guidelines

DNA Quantity

Libraries must be submitted at a minimum concentration of 5 ng/ μ L as measured by Qubit or 15 ng/ μ L as measured by Nanodrop, with a 260/280 ratio >1.7. Minimum library volume needed for QC and sequencing is 10 ng, though more is preferred.

Upstream Library Prep considerations

- 1. Please provide all index **SEQUENCES** (the same index number from different library preparation kits do not always refer to the same sequence).
- 2. The A_{260}/A_{280} should be 1.7-1.9. Ratios that deviate from this significantly suggest that DNA quantification is not accurate and your library may not have been successful.
- 3. If possible, please provide an expected size for your library—this is especially important for amplicon libraries that do not always size correctly on the Agilent Bioanalyzer. A gel image is not necessary, but can be helpful.
- 4. Samples should be in 10 mM Tris-HCl or nuclease-free water. Please note the type of buffer on the submission form. EDTA should be avoided as it retards downstream enzymatic reactions.
- 5. Upon receipt, all libraries will be QC'd by Agilent Bioanalyzer and qPCR, and customer charged a per-sample QC fee. Results will be returned to the customer at their request.

Shipping

- 1. A filled out submission form must accompany all shipments.
- 2. Samples <12 should be submitted in either a 0.5 mL or 1.5 mL nuclease-free microfuge tube with the sample name **clearly** written on the cap—NO more than 6-8 letters/numbers—**and the PI's name on the side**. If >12 samples submitted, the samples should be in a nuclease-free PCR plate sealed appropriately with the PI name on the side of the plate skirt (spreadsheet with plate layout should be e-mailed to <u>genomics@vai.org</u> and should also accompany plate). All items shipped should be sealed in a plastic bag.
- 3. Sample should be shipped on dry ice overnight to

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