

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 12 μ L, 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 A₂₆₀/A₂₈₀ 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, which 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 impedes downstream enzymatic reactions.
- 5. Upon receipt, all libraries will be QC'd by Agilent Bioanalyzer and qPCR; the customer will be charged a persample QC fee. Results will be returned to the customer at their request.

Shipping

- 1. A completed 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 genomics@vai.org and should also accompany plate). All items shipped should be sealed in a plastic bag.
- Sample should be shipped on dry ice overnight to: Van Andel Institute Attention: Marie Adams
 333 Bostwick Ave. NE, Room 1211 Grand Rapids, MI 49503