

**UMass Medical School  
Illumina Sequencing Library Construction Service  
Molecular Biology Core Labs**



**Client Information:** Name: \_\_\_\_\_ Phone: \_\_\_\_\_  
 Email: \_\_\_\_\_ Date: \_\_\_\_\_  
 PI/Lab: \_\_\_\_\_ Account #: \_\_\_\_\_  
 PI signature: \_\_\_\_\_

**Library Construction Information (Please submit one ticket for each set of libraries to be mixed):**

Illumina Library	Minimum Requirements:
Indicate starting material: ___ Genomic DNA ___ Amplicon(s) ___ cDNA (double stranded) ___ Total RNA for cDNA library* ___ Small RNA* ___ NanoString collection <sup>‡</sup> , # of samples: _____ ___ Cells for 10X Chromium single-cell library ___ Other cells ___ Other _____ (describe)	genomic DNA: 2-3 µg amplicon: 1 µg (2 µg if shearing required) cDNA: 1 µg (2 µg if shearing required) total RNA: 2-5 µg small RNA: 50-100 ng NanoString: dried + Seq Code plates and enzyme Master Mix cells for Chromium: 10,000 viable cells other cells: ask before submission
What is the preferred insert size? _____ (Optimal library inserts are ≤ 500 bases)	
Instrument: _____	
Desired read length: _____	
<i>* Must be stored in water</i>	
<i>‡ Include the _SeqCodeIndices.csv file from the NanoString instrument</i>	

Species: \_\_\_\_\_ Source: \_\_\_\_\_

How was the material prepared? \_\_\_\_\_

What buffer is the material in? \_\_\_\_\_

List the name and part # of any kits used in preparation: \_\_\_\_\_

How was the material quantified? \_\_\_\_\_

**Is the material from a human pathogen, or does it contain any restricted, potentially hazardous, or infectious material?** If yes, please detail: \_\_\_\_\_

*Your signature confirming this statement:* \_\_\_\_\_

Sample Name	Concentration	Volume (µl)

(If additional samples, list on supplemental sheet.)

Please attach any available information about the sample material (e.g. gel photos, Fragment Analyzer/Bioanalyzer traces, etc).