<u>University of Michigan DNA Sequencing Core</u> Services, Pricing and Information for UMich internal Clients

The University of Michigan DNA Sequencing Core ("UMDSC") provides analytical services to UMich-affiliated researchers ("internal clients") and non-UMich-affiliated researchers ("external clients"). This document lists the services provided to internal clients of the UMDSC, including the prices of those services as of the above revision date. Revisions to this document will be made periodically with approval of the UM Office of Financial Analysis (OFA), and may include changes in prices or services. Revisions become effective upon the date of OFA approval, and at that time supersede all previous price lists. Actual prices charged will be those approved by the OFA. In event of errors in this document, the OFA approved prices supersede those stated here.

The UMDSC offers nucleic acid analysis services as listed in the following table. For definitions or for detailed information on the services, please consult the Core's web site (http://seqcore.brcf.med.umich.edu) or with Core staff members. The actual prices that a client will pay generally requires a complicated mixture of the below charges. All prospective clients should contact the Sequencing Core (seqcore@umich.edu) for discussion and clarification of costs for *their* project.

Discounts are available for members of certain Centers. In most cases, these are provided by the Center administration in the form of rebates, after the service is rendered by the Core and after payment has been made by the Investigator. Please contact your Center administrators to discuss potential discount opportunities.

Sanger Sequencing:	
Sanger Sequencing, standard	\$3.00/rxn
Sanger Sequencing, Rush	\$5.00/rxn
Sanger Sequencing, Same Day	\$8.00/rxn
(Same Day Rush is available only if enabled on the Sample Submission form.)	
Sanger Seq, full 96-well plates	\$2.40/rxn
(Will be charged for any empty wells)	
Sanger Seq, Reduced Service	\$2.25/rxn
(Full 96-well plate only. Please consult Core staff before attempting to use this sea	
Sanger Seq, Sec Structure DNA	\$6.00/rxn
(siRNA, hairpin)	
Sanger Sequencing, Large DNA	\$12.00/rxn
Sanger Seq, large projects	contact us
Fragment Analysis:	Φ 50.00 /
FragAnalysis, Instrument Use	\$58.00/run
FragAnalysis, CE Loading Plate	\$36.00/plate \$35.00/rxn
FragAnalysis, IdentifilerPlus	\$35.00/rxn
qPCR Instrument Usage:	
qPCR Instrument Usage	\$20.00/run
(Run one SIBR Green or Tagman 96- or 384-well plate, client-prepared)	+
(
Sequenom MassARRAY Services:	
Sequenom Pinset, reagents only	\$316.00/pinset
(Pinset is 24 reactions, each of which is 1 sample tested against 1 multiplex)	
Sequenom Quadrant reagnts only	\$800.00/quadrant
(Quadrant is 96 reactions, each of which is 1 sample tested against 1 multiplex)	
Sequenom HalfChip reagnts only	\$1,420.00/half chip
(HalfChip is 192 reactions, each of which is 1 sample tested against 1 multiplex)	
Sequenom FullChip reagnts only	\$2,580.00/full chip
(FullChip is 384 reactions, each of which is 1 sample tested against 1 multiplex)	
EpiTyper reagents only	\$750.00/chip
(Priced as fraction of a chip)	
Sequenom technician time	\$90.00/hour
(e.g. assay design, sample processing, reaction spotting, data analysis)	

PyroMark pyrosequencing services:

FullPlate reagents, std labor \$600.00/plate
PartialPlate reagnts, std labor \$290.00/plate

(24 wells)

PyroMark technician time \$90.00/hour

(Extra effort beyond standard processing labor)

Illumina BeadArray services:

Illumina kits at market price

BeadArray technician time \$61.00/hour BeadArray reagnts and supplies \$257.00/plate

(Per 96-well plate)

BeadArray Scanning \$12.00/hour

(Instrument usage)

Illumina Sequencing services, Library generation:

Standard Genomic libraries \$102.00/library 96-well Genomic libraries \$34.00/library

(High throughput automated process, in development)

Ultra-low input genomic lib
PCR-free genomic libraries
\$120.00/library
10x Genomics Long-Contig lib
Exome captures, pools
Non-Stranded mRNA polyA lib
\$123.00/library
\$120.00/library
\$1027.00/library
\$531.00/pool

(Discontinued by Illumina 12/31/2017)

Stranded mRNA polyA libraries\$121.00/libraryStranded mRNA Ribo-reduced lib\$285.00/librarySMARTer stranded low input\$209.00/library

(Client supplies Ribo-Gone kit if needed)

smRNA-seq libraries \$194.00/library 10x Genomics Single-cell RNA \$1721.00/library

(Must be scheduled in advance!)

Library QC/Normalize/Pooling \$22.00/library
Pool assessment for loading \$48.00/pool
Illumina Library Tech time \$97.00/hour

(Extra services for custom projects)

Illumina Sequencing Services on the HiSeq 2500 - 2 lanes per flow cell:

HS 2500v2 Single-end, 50 RAPID \$1047.00/lane HS 2500v2 Paired-end, 50 RAPID \$1651.00/lane HS 2500v2 Paired-end 100 RAPID \$2185.00/lane HS 2500v2 Add 50nt RAPID run \$433.00/lane

(Custom projects only)

Illumina Sequencing Services on the HiSeq 2500 and HiSeq 4000 - 8 lanes per flow cell:

 HS 2500 v4 Single-end, 50
 \$980.00/lane

 HS 2500 v4 Paired-end, 50
 \$1551.00/lane

 HS 2500 v4 Paired-end, 125
 \$2347.00/lane

 HS 2500 v4 Add 50nt to v4 run
 \$402.00/lane

HS 4000 Single-end, 50 \$973.00/lane
HS 4000 Paired-end, 50 \$1558.00/lane
HS 4000 Paired-end, 75 \$1667.00/lane
HS 4000 Paired-end, 150 \$2296.00/lane
HS 4000 Add 50nt to HS4000 run \$404.00/lane

(Custom projects only)

Illumina Sequencing Services on the NextSeq 500 and MiSeq- 1 lane per flow cell:

NextSeq Mid-Output 150 cycle \$1328.00/lane
NextSeq Mid-Output 300 cycle \$1975.00/lane
NextSeq High-Output 75 cycle \$1612.00/lane
NextSeq High-Output 150 cycle \$2923.00/lane
NextSeq High-Output 300 cycle \$4525.00/lane

 MiSeq 300 cycle Micro
 \$706.00/lane

 MiSeq 300 cycle Nano
 \$592.00/lane

 MiSeq 500 cycle Nano
 \$713.00/lane

 MiSeq 150 cycle v3
 \$1214.00/lane

 MiSeq 500 cycle v2
 \$1601.00/lane

Duo cBot RAPID kit\$212.00/eachDrive, 1T\$75.00/eachIllumina Lanes Tech time\$88.00/hour

(Extra services for custom projects)

Pacific Biosciences Sequencing services:

PacBio shearing and QC
PacBio DNA library preparation
\$468.00/sample

(Additional reagents may be required, in certain cases. Discuss with Core staff.)

BluePippin Isolation \$111.00/sample
PacBio Technician time \$75.00/hour
PacBio SMRT Cell sequencing \$425.00/SMRT cell

MicroArray Services:

There are two cost components to a MicroArray project, the array and the processing time. Arrays are purchased by the Core and billed with processing, after project completion. Except where noted, prices below are for processing ONLY.

Life Technologies/Affymetrtix:

Std, GeneChip cartridge	\$257.00/sample
Std, GeneAtlas 4-array strip	\$230.00/sample
Standard, GeneTitan plate	\$156.00/sample
(Plates are available in 16-, 24-and 96-array formats)	
Low-input, GeneChin cartridge	\$222.00/sample

Low-input, GeneChip cartridge Low-input, GeneAtlas strip \$206.00/sample Low-input, GeneTitan plate \$166.00/sample GeneChip, hyb scan only \$81.00/sample GeneAtlas, hyb scan only \$30.41/sample GeneTitan, hyb scan only \$41.73/sample GeneChip SNP typing \$290.00/sample GeneChip SNP typing, hyb only \$43.00/sample Axiom (GeneTitan) SNP typing \$27.00/sample OncoScan CNV and SNP typing \$765.00/sample

(Includes the cost of the arrays)

Targeted qPCR Arrays:

Qiagen RT2 Profiler prep \$244.00/plate

(Qiagen 96- or 384-well plates are billed at market prices.)

Life Tech OpenArray prep\$128.00/arrayLife Tech OpenArray miRNA prep\$150.00/array

(Life Tech arrays and accessories are billed at market prices.)

DNA Isolation Services:

\$40.50/sample
\$28.50/sample
\$49.00/sample
\$33.00/sample
\$19.50/sample

Subaliquot, transfer, dilution \$1.70/sample Quantify, Nanodrop or Qubit \$1.70/sample DNA and RNA Quality Control Service: Agilent Bioanalyzer 2100 - Submission requests are per chip. We cannot bill for partial chips. Bioan DNA 1000, 7500 or 12000 \$70.00/chip **Bioan DNA High Sensitivity** \$92.00/chip \$68.00/chip Bioan RNA 6000 Nano Bioan RNA 6000 Pico \$72.00/chip \$72.00/chip **Bioan Small RNA** Agilent TapeStation 2200 **TapeStation Standard DNA 1000** \$18.50/sample \$20.40/sample

TapeStation Standard DNA 1000\$18.50/sampleTapeStation High-Sens DNA 1000\$20.40/sampleTapeStation Genomic DNA\$20.40/sampleTapeStation Standard RNA 6000\$18.60/sampleTapeStation High-Sens RNA 6000\$19.00/sample

Please note the following for all services:

The UM DNA Sequencing Core will process samples on a first-come, first-served basis. Samples from external clients will be processed at lower priority than internal clients. For the Pacific Biosciences sequencer, HHMI Investigators have priority above that of UM researchers, up to certain limits. Clients must be registered with the UM DNA Sequencing Core via our computer system (see the "PI Administration" section of Core's web site). All samples must be entered into our computer system, and must be properly labeled with our tracking number(s) before they are delivered to this Core. We reserve the right to discard any samples that are not properly labeled. The Core does not return unused samples or portions of samples after completion of work; they are discarded.

Prices listed in this document are those charged to internal (University of Michigan) clients, as determined by the method of billing employed by the University of Michigan Financial Operations. If a client pays using funds from a "shortcode" within the UM Financial Operations, that client will be charged the rate for internal clients, and the prices listed in this document are applicable. If a client pays by *any* other method, then the prices listed in this document are not applicable. No exception to this policy will be made for any actual or perceived affiliation the client may have with the University of Michigan, nor for any collaborative agreement the client may have with a University of Michigan employee, faculty, student, officer or agent.

An analysis performed by the UMDSC can fail for many reasons, including failures on the Core's part (e.g. instrument malfunction, Core technician error) and failures on the Customer's part (e.g. misquantitated sample, misdesigned experiment, impure sample). We will, to the best of our ability, assess whether a failed analysis is due to a failure on the part of the Core, and we may in such cases repeat an analysis at our cost. When, in our sole judgment, a failure is not due to error on the part of the Core, we reserve the right to refuse repeating the analysis. We will not be held liable for analytical failures arising due to errors or problems in the Client's laboratory. We will not bear the cost of repeat runs unless we explicitly state that we are doing so of our own accord. If, on completion of any repeat, the results prove that the Core was at fault in the original failure, we may at that time choose to absorb the cost of the repeat analysis. Determination of the cause of failure is based solely on the judgment of the Core staff members, and ultimate decision making in such cases resides with the Director of the UM DNA Sequencing Core.