

## UMGC's DNA Sample Submission Instructions for Illumina Genotyping BeadChips

**Overview:** The following are Sample Preparation and DNA Submission Form Filling Instructions.

**Audience:** Researchers who want to submit samples for various Illumina Genotyping Beadchips.

**SYNOPSIS:** PLEASE submit 500 ng of DNA at 25 ng/ul in a 96-WELL PLATE in a COLUMN-WISE fashion RANDOMIZING samples and keeping two unique blank wells.

Protocol:

### \_\_\_ 1) Prepare DNA in TE or water.

- a. Extract DNA using a kit of your choice.

### \_\_\_ 2) Quantitate and Normalize DNA samples to 25 ng/ul

- a. Quantitative DNA using your method of choice (UV absorption, PicoGreen, etc).
- b. Normalize your DNA to 25 ng/ul (a range of about 15-35 ng/ul is acceptable).
- c. **There will be extra charges if samples are not normalized.**
- d. We require a minimum of 500 ng of DNA. *\*Certain Genotyping BeadChips may require more DNA, please contact Service Manager for the DNA requirements for your specific BeadChip.*

### \_\_\_ 3) Plate Samples Column Wise in a 96-well plate

- a. We only accept DNA in a 96-well plate NOT tubes!
- b. Please use a Bio-Rad 96-well fully skirted plate (Cat # MSP9601) or ABgene 96-well plate (part # AB-0800) – we will transfer samples to a Bio-Rad plate if you do not have one. **Sample transfer charges will apply.**
- c. Since the processing for Illumina processes is performed in a column-format, please submit your samples in a COLUMN-ORDERED FASHION ONLY!!

### \_\_\_ 4) Randomize Samples

- a. If your samples are from different tissue types (and/or paired samples like case controls or duplicates), please RANDOMIZE them when plating. (This will effectively eliminate chip-to-chip bias – In general Illumina's BeadChips have little variation but randomizing samples will help reduce it further).
- b. Again, BeadChips are hybridized by a liquid handling robot in a column fashion. The hybridization pattern varies between the different beadchips. Please contact the service manager for the layout for your requested BeadChips in order to arrange your samples accordingly.

## \_\_\_\_ 5) Blank wells

- a. Please have 2 uniquely located blank wells (EMPTY wells) for every 96-well sample plate to which nothing has been added. Blanks serve a dual purpose – they serve as a fingerprint to identify the plate and its orientation. And we will run CEPH DNA controls in place of the blanks that serve as controls.
- b. The blank wells must be asymmetrical, such that a 180-degree rotation of the plate results in a shift of the fingerprint. For example, wells A01 and D07 are acceptable but A01 and H12 are not.
- c. If you have less than 48 samples, please keep 1 well blank instead of 2.
- d. **Extra charges will apply if the UMGC has to re-format the plate to create/modify blank wells.**

## \_\_\_\_ 6) Please fill the DNA submission form

- a. Please refer to detailed instruction on the DNA submission form and fill it accordingly.

## \_\_\_\_ 7) Email us the filled submission form

- a. University of Minnesota researchers can attach it to their online order under SNP Genotyping within the online ordering system.
- b. External users can email it to us at [genotype@umn.edu](mailto:genotype@umn.edu).