Running the DGE pipeline on LSF:

Prerequisites:

- 1) Download the DGE package and unpack it with tar
- 2) You'll need both python (I usually run it in 2.7) and bwa installed. bwa should be runnable by simply typing the command "bwa".

Usage is as follows:

usage: python /path/to/dge-prod/Scripts/run_DGE_analysis.py [-h]

[--short_lsf_queue SHORT_LSF_QUEUE]
[--long_lsf_queue_LONG_LSF_QUEUE]

[--loose barcodes] [--cleanup]

sample_map reference barcodes alignment_dir

analysis dir

positional arguments:

sample_map location of sample map file

reference reference genome: Human|Mouse|Rat|Chicken barcodes barcode plate: P1|P2||P3|P1P2|Trugrade_384_set1|

Trugrade_96_set1|Trugrade_96_set2|Trugrade_96_set3|

Trugrade_96_set4

alignment_dir directory to process alignments analysis_dir directory to calculate gene expression

optional arguments:

-h, --help show this help message and exit

--cleanup removes resulting fq and sam files upon successful completion

Most arguments should be self explanatory. The sample_map should be a tab-delimited file where there's a line for each pair of fastqs like:

sample id subsample id /path/to/read1.fg /path/to/read2.fg

*Please note: the pipeline expects read1 to have at least 16 called bases, where bases 1-6 represent the well barcode and bases 7-16 represent the UMI.

An example usage of the pipeline is:

python /path/to/dge-prod/Scripts/run_DGE_analysis.py --short_lsf_queue queue --long_lsf_queue queue SampleMap Human Trugrade 384 set1 Alignment DGE

The word "queue" should replaced with the name of your queue in LSF. At the Broad we have different queues for short and long-running jobs. If you don't, you can use the same queue for both settings.