

Frodyma, et. al. 2020

Data analysis for Frodyma paper

Citation

Coming soon...

RNA-seq analysis methods

RNA sequencing (RNA-seq) analysis was conducted by _____. For RNA sequencing, RNA was purified from _____ by _____. Total RNA samples (_____ ng) were hybridized with _____ (_____) to substantially deplete rRNA from the samples. _____ Stranded/Unstranded _____ RNA sequencing libraries were prepared as using the _____ Kit. Purified libraries were quantified on _____ using _____. Libraries were normalized to _____ nM and equal volumes were pooled in preparation for sequence analysis. _____ etc. _____ Raw sequence data has been deposited as GSE_____

Sequence reads were preprocessed using XPRESSpipe (v0.4.1) [1], with adapter sequences set to AGATCGGAAGAGCACACGTCTGAACTCCAGTCA and AGATCGGAAGAGCGTCGTGTAGGGAAAGAGTGT. Reads were processed using H. sapiens GRCh38.p13 Ensembl release 99. Differential expression analysis was performed using the XPRESSpipe wrapper for DESeq2 (v1.22.1) [2]. Differentially expressed genes were further visualized using XPRESSplot [1]. Isoform abundance analysis was performed using the XPRESSpipe wrapper for Cufflinks (v2.1.1) [3] and IGV (v2.4.19) [4]. Scripts used to perform these analyses can be found at https://github.com/j-berg/froydima_2020.

References

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- [3] Trapnell C, Roberts A, Goff L, Pertea G, Kim D, Kelley D, et al. Differential gene and

transcript expression analysis of RNA-seq experiments with TopHat and Cufflinks. Nat Protoc. 2012;7. <https://doi.org/10.1038/nprot.2012.016>.

[4] Robinson J, Thorvaldsdóttir H, Winckler W, Guttman M, Lander E, Getz G, et al. Integrative Genomics Viewer. Nat Biotechnol. 2011;29:24–26. <https://doi.org/10.1038/nbt.1754>.