



REPLY TO HARTL AND GAO:

Lack of between-batch difference in the distributions of measured extracellular RNA levels

Zixu Zhou^a, Qiuyang Wu^a, Zhangming Yan^b, Haizi Zheng^a, Chien-Ju Chen^b, Yuan Liu^a, Zhijie Qi^b, Riccardo Calandrelli^b, Zhen Chen^c, Shu Chien^{a,d,1}, H. Irene Su^{e,f,1}, and Sheng Zhong^{a,b,d,1} 

We thank Drs. Hartl and Gao for pointing out the difference in read lengths between the cancer and normal samples (1). Indeed, Illumina discontinued their 50-base pair (bp) sequencing kits after we finished sequencing the cancer samples, forcing us to switch to 75-bp sequencing kits on the normal samples. The different sequencing kits did incur small changes of sequence alignment metrics. While acknowledging the possibility that this may bias the assessment of cancer vs. normal discrimination performance of our method, we believe such a possibility is substantially ameliorated by the following considerations.

First, a recent study reported that “with the exception of 25 bp reads, there is little difference for the detection of differential expression regardless of the read length” (2). In line with this report, the absolute difference of mismatch rates (0.5% vs. 0.8%) between the 50-bp and 75-bp reads is much smaller than the typically used threshold (4%) of the mapping software (STAR). Furthermore, among the sequence alignment metrics Hartl and Gao list (1), only the rates of uniquely mapped reads directly affect the quantification of extracellular RNA (exRNA) levels. However, the rates of uniquely mapped sequences contributed to sample-to-sample variation in PC1, which is independent of the observed cancer–normal difference (PC2) (second row in figure 1B of ref. 1).

Second, we cannot identify a batch difference in the distributions of the measured exRNA levels: “The distributions of TPM exhibited little difference between any 2 cancer samples or between a cancer

sample and a normal sample (Fig. 4A and *SI Appendix*, Fig. S9). Thus, every sample contains a similar proportion of highly expressed exRNAs, regardless of the threshold for calling highly expressed exRNAs” (3). Furthermore, the pairwise differences of the normal samples in exRNA levels nearly completely overlapped with the between-batch pairwise differences (Fig. 1). Taken together, read length is unlikely to affect the quantification of exRNAs or bias our cancer–normal comparison, which was completely based on exRNA quantification.

However, the aforementioned considerations cannot dwarf our caution that “this study did not rule out all possible confounding factors that may contribute [to] the separation of cancer and normal samples” and “future studies . . . are needed, preferably as double-blind prospective trials” (3).

Finally, the read-length difference does not affect other major findings in our paper. Specifically, Small Input Liquid Volume Extracellular RNA Sequencing (SILVER-seq) can reproducibly produce exRNA sequencing libraries from different aliquots of the same serum sample. exRNA levels measured by SILVER-seq correlated with those measured by standard exRNA sequencing methods. SILVER-seq detected fragments of tissue-specific RNAs in human sera. SILVER-seq–derived exRNA levels reflected chronological age. SILVER-seq–based classifiers exhibited good discrimination power between recurrent and nonrecurrent cancer samples.

^aGenemo Inc., San Diego, CA 92121; ^bDepartment of Bioengineering, University of California San Diego, La Jolla, CA 92093; ^cDepartment of Diabetes Complications and Metabolism, Beckman Research Institute, City of Hope, Duarte, CA 91010; ^dInstitute of Engineering in Medicine, University of California San Diego, La Jolla, CA 92093; ^eMoores Cancer Center, University of California San Diego, La Jolla, CA 92093; and ^fDepartment of Obstetrics, Gynecology and Reproductive Sciences, University of California San Diego, La Jolla, CA 92093

Author contributions: H.Z., C.-J.C., Y.L., Z.Q., and R.C. analyzed data; and Z.Z., Q.W., Z.Y., Z.C., S.C., H.I.S., and S.Z. wrote the paper.

The authors declare no competing interest.

Published under the [PNAS license](#).

¹To whom correspondence may be addressed. Email: shuchien@ucsd.edu, hisu@ucsd.edu, or szhong@ucsd.edu.

First published January 21, 2020.

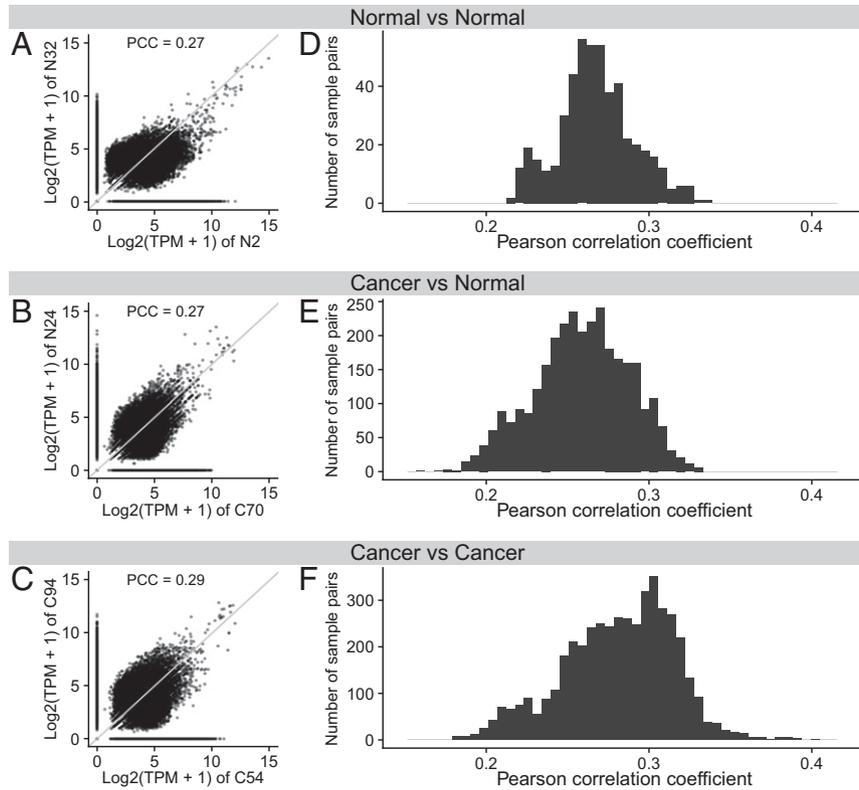


Fig. 1. Genome-wide comparisons of exRNA levels within and between the 2 batches. Pairwise difference is measured by 1-correlation. (A–C) Scatter plots of log₂-transformed transcripts per million (TPMs) of all genes between 2 normal samples (A), a cancer (x axis) and a normal sample (y axis) (B), and 2 cancer samples (C). (D–F) Histograms of pairwise Pearson correlation coefficients (PCCs) between every pair of normal samples (D), every cancer–normal pair (E), and every pair of cancer samples (F), based on the log₂-transformed TPMs of all genes.

- 1 C. Hartl, Y. Gao, Clarifying the effect of library batch on extracellular RNA sequencing. *Proc. Natl. Acad. Sci. U.S.A.* **117**, 1849–1850 (2020).
- 2 S. Chhangawala, G. Rudy, C. E. Mason, J. A. Rosenfeld, The impact of read length on quantification of differentially expressed genes and splice junction detection. *Genome Biol.* **16**, 131 (2015).
- 3 Z. Zhou et al., Extracellular RNA in a single droplet of human serum reflects physiologic and disease states. *Proc. Natl. Acad. Sci. U.S.A.* **116**, 19200–19208 (2019).