

1. Consider Figure 1 from the Hillman et al. paper.

A. What was the experiment that lead to the results? (What did the authors do, what did they measure, what were the results).

- Just look experiment + control: with <sup>non functional</sup> <sup>functional</sup> parental allele **OK**
- measured Ube3a transcript level via quantitative RT-PCR  
protein level via various tissues.  
hippocampus, thymus, liver, heart.

they normalized w/ FPKM values of UBE3A transcripts. **and what did they also look at?**

B. What do the data show?

- in both  $Ube3a^{mfp+} \times mfp^-$ , the Ube3a transcript & protein expression levels are higher in the CNS than N-CNS **i.e....**
- when  $Ube3a$  is normalized via FPKM, only lung is significantly different from the exp. level in the hippocampus
- the  $Ube3a$  FPKM level of maternal is significantly higher than paternal by 20 fold in the hippocampus, but not significantly different in other parts of the body.

C. What can we conclude from the data?

- functional parental allele is not necessary to display WT transcript/protein levels in the CNS & N-CNS in mouse tissues.
- discrepancy in maternal & paternal together makes the level of  $Ube3a$  FPKM in hippocampus overall high FPKM if  $Ube3a$  is similar to other tissues except the lung

D. What do the authors conclude from these data? Do you agree or disagree with their conclusion (e.g. the last sentence of the introduction), and why?

concluded that imprinting does not function to regulate the dosage of  $Ube3a$  /  $UBE3A$  in neurons. I disagree because in Figure 3, the  $Ube3a$  /  $UBE3A$  level is higher in mouse than in opossum for the cortex. Imprinting occurs in mice while opossum biallelically expresses  $Ube3a$ , so imprinting may have a function in increasing dosage in the CNS, but not there must be other mechanisms to result in higher expression in the CNS in opossum remaine.

**very good point! What they assumed was that imprinting would gene product"**

When  $Ube3a$  FPKM is looked separately in CNS vs N-CNS, then the levels are not significant, so not significant even though imprinting would gene product"