

The Art of Grant Packaging or How Your Grant Looks Does Matter

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How your grant looks does matter

Common features of NIH study section member

Tired eyes

Busy brain

Tired brain

No one comes to study section saying: “I am so rested and relaxed that I had lots of time to read my grants”



Text

1. Left justified vs. both sides justified?
2. Create space between paragraphs to provide visual breaks
3. Check spelling, check grammar; it matters
4. Make sure text flow around figures is clear

Figures are key

1. Need to balance # figures and volume of text: more is not necessarily better
2. Figures need to be necessary for points being made
3. Stand alone – point made is visually obvious: my criteria is that I can easily see all aspects of the figure at 125%; If I have to expand to 200% or greater that is not good
4. Quality of figures; use of Adobe Illustrate to great figure in .eps format
5. Figure legend text; succinct

One Example: note size of text in figure relative to size of text in the document

2. Generation of muscle specific *Bmal1* KO mice. Our observations that muscles *Bmal1* KO mouse were exciting and provided evidence to support links between circadian rhythms, the molecular clock and skeletal muscle homeostasis. However, those mice do not allow us to discern the relative contribution of the skeletal muscle to address the function of the molecular clock in skeletal muscle we generated an inducible *Bmal1*^{fl/fl} mouse [iMS*Bmal1*^{+/+}; iMS*Bmal1*^{-/-}] for the targeted disruption of the molecular clock in skeletal muscle. The tamoxifen inducible Cre recombinase mouse (HSA-MCM) was provided by Dr. David M. Nathans at the University of Kentucky and the efficacy of this mouse for studies of skeletal muscle recombination in adult mice has been demonstrated.

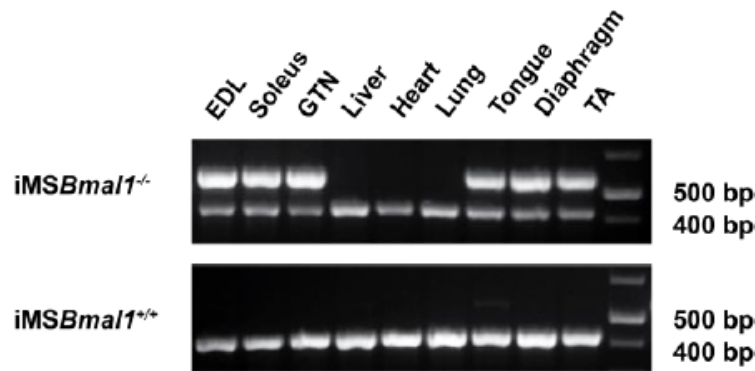


Figure 1. PCR results showing recombination of *Bmal1* in muscle

recombination in adult mice has been demonstrated. The *Bmal1* mouse is available through Jackson Laboratories [B6.129S4(Cg)-Arntl^{tm1}Weit/J] and has been used to target *Bmal1* function in selected cell types. We generated the iMS*Bmal1* mouse and treated mice at 8-12 weeks of age with either vehicle or tamoxifen for three weeks following the treatment with tamoxifen. We isolated skeletal muscles, heart and other non-muscle tissues, extracted DNA and performed PCR to evaluate *Bmal1* recombination. As seen from the results provided in Figure 1, *Bmal1* (band = 571bp) only occurs in skeletal muscle in tamoxifen treatment and not in heart, liver, lung, tongue, diaphragm, and tail.