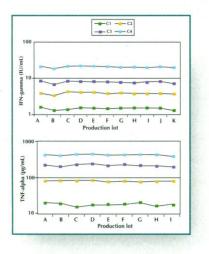
# BioSource EASIA Kits

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**Show Me!** Individual production lots were analyzed using 4 levels of control specimens according to standard protocol. Interlot CV for all controls ranged from 5.1-6.6%



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#### SCIENCE'S COMPASS

standard protocol in most mouse laboratories. Group-housed and isolated male mice differ in the dynamics of the patterns of their dominance hierarchies, as well as in their aggressive behavior, and these differences are strain-dependent (1). Moreover, individual housing can lead to increased anxiety-like behavior in an elevated plus maze (2). Just how housing practices might interact with laboratory site to affect strain differences is not readily predictable from literature of which we are aware, nor for our data were there differences in housing—all mice were grouped.

Dawson et al. agree that multiple tests using different approaches should be used to solidify inferences about the genetic structure of behavior, although they are hardly unique in adopting this practice. They imply without directly asserting that the fact that our measures were unconditioned, as well as ill defined, may have led us to be unable to isolate small genetic differences reliably. Only some behavioral domains are best tapped with conditioned responses, and we avoided these in our study for practical reasons. We do not agree that our responses were ill defined. For example, on our water escape task, group differences were not particularly large (multiple  $R^2 = 0.18$ ), but the intertrial consistency of behavior, as indicated by Cronbach's coefficient alpha, was reasonably high ( $\alpha = 0.81$  over the first four trials). Such consistency is not the hallmark of ill-defined tasks. They also err in their assertion that "the main source of variation...in one laboratory was where the experimenter was highly allergic to mice." We offered this as an example of a laboratory difference, but there are no data suggesting it was the "main source of variation"-this appears to be Dawson et al.'s opinion. There was probably ultrasound emitted from the motor of the Racal Airmate 1 device strapped to the small of the back of the Edmonton experimenter at waist level. However, this was a constant: the experimenter wore the unit whenever working with the mice, from the day they arrived in the colony until the end of testing, and there was ample time for habituation. Whether wearing the Airmate apparatus had any effect on mouse behavior in standard tests can only be addressed with a controlled study using people not allergic to mice who wear or do not wear the filter unit. Data relevant to this question are needed before the effects can be called "profound."

Hen notes that his knockout mouse colony has been maintained on a genetic background involving multiple 129 substrains. This, we suspect, is true for many other knockout colonies as well. To explain the loss of alcohol drinking phenotype in the 5-HT<sub>1B</sub> knockouts over time, he proposes that an increasing influence of modifier genes from the 129/SvEvTac strain reduces alcohol preference. This hypothesis can be tested definitively by rederiving cryopreserved embryos from the original population. The stability of reduced alcohol-induced ataxia in the knockouts suggests that the effects of such modifier genes are trait-specific, which is consistent with our other findings. Hen elaborates a breeding *strategy* that can protect against such modifier gene effects; maintaining knockouts on fully inbred rather than segregating populations also will accomplish this.

Tordoff et al. suggest our results may have been influenced by differences among labs in the composition of Purina diets. This is quite feasible because there were modest but statistically significant differences among our three labs in mouse body and brain weights. We agree that it would be interesting to run further experiments of this nature using rigorously defined semisynthetic diets. Our study rigorously equated the behavioral test apparatus and testing protocols, and we sought to restrict variation in many aspects of the lab environment. We did not seek to equate the lab environment, however. We wanted to know whether commonplace variations in lab environments would modify the pattern of genetic effects, and we found that for certain behaviors they did, whereas other behavioral tests yielded substantially the same results in all three labs, despite the differences among diets and drinking water. It is doubtful that differences between labs can be explained by a single environmental factor; instead, both the environmental and genetic contributions are probably multifactorial and complex

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## CORRECTIONS AND CLARIFICATIONS

It should have been made clear in George B. Dyson's letter "Darwin in Kansas" (*Science*'s Compass, 27 Aug., p. 1355) that Erasmus Darwin's warning was one that could have been given to today's Kansas Board of Education. It was not literally given to the Board in 1794, as the state of Kansas did not then exist.