Parameter optimization for automated behavior assessment: plug-and-play or trial-and-error?

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Introduction

Over the years, fear conditioning research in rodents has moved from purely “manual” scoring of freezing behavior by human observers to mainly automated measurements. Since a few years, conditioning researchers have (re)focused on generalization, including generalization of context conditioning, because of its relevance to the study of learning and memory, and to the development and maintenance of anxiety disorders (Wang et al., 2009; Wiltgen et al., 2010; Hermans et al., 2013). Contextual generalization/discrimination research necessitates that freezing in several contexts is compared directly (e.g., Wang et al., 2009; Wiltgen et al., 2010; Yu et al., 2010). Counterbalancing of contexts is not always possible, e.g., in case of a generalization gradient with some contexts having a grid and others a plastic floor (Luyten et al., 2013; Poulos et al., 2013). Furthermore, the differences in freezing between the originally trained and a similar context are often modest, and not as clear-cut as between the original and a dissimilar context (unpublished data), resulting in limited effect sizes. Concurrently, there is an increasing interest in transgenic animals, to investigate the role of certain genes in discrimination and generalization (e.g., Yu et al., 2010; Tayler et al., 2011; Cushman et al., 2012). Genetic modifications may, however, result in phenotypes with only small behavioral deficits.

Taken together, this growing domain is confronted with the challenge to distinguish subtle behavioral effects in different contexts. Our data show that automated behavioral measurements may not always be appropriate for these purposes, or that, at least, they should be implemented and interpreted with great care.

To our knowledge, there is only one paper that has systematically validated settings for a widely used system to detect freezing in rodents (i.e., Med Associates, Inc. Video Fear Conditioning System, with VideoFreeze® software). The optimized software parameters for freezing mice [motion index threshold 18 and minimum freeze duration 30 frames (1 s)], were published in this journal (Anagnostaras et al., 2010). One of the challenges of parameter optimization is finding a balance between the detection of “non-freezing” (e.g., tail) movements, while at the same time ignoring respiratory and cardiac motion during freezing. VideoFreeze is also frequently used to assess rat freezing behavior and others have reported their validated settings for rats (motion threshold 50) (Zelikowsky et al., 2012a). It makes sense to use a higher activity threshold for rats, as they are bigger animals and respiratory movements—which do not preclude freezing—will result in more pixels changing than for mice. Those are also the settings that we used in our studies.

Materials and methods

Experiments were conducted on 48 male Wistar rats (±275 g), in three replications of 16 rats each (eight rats per group). All sessions were meticulously scheduled using free ExpTimer software (Luyten and Van Cappellen, 2013). All experiments were approved by the KU Leuven animal ethics committee, in accordance with the Belgian Royal Decree of 29/05/2013 and European Directive 2010/63/EU.

On the first day, rats were trained in context A (Figure 1A). Four minutes after the start of the session, rats received five unsignaled footshocks (0.8 mA, 1 s), separated by 90 s. One minute after the last shock, animals were returned to their home cage. Twenty-four hours later, half of the rats were tested in context A and the other half in similar context B (Figure 1B). During this test, rats were exposed to the context for 8 min, without shocks. We hypothesized that there would be significantly less freezing (unpaired t-test) in context B than in A on day 2, because of a generalization decrement. The results presented here are part of an ongoing study, including other control groups, which will be discussed as a whole elsewhere. Freezing...
RESULTS AND DISCUSSION

Three consecutive series of rats (16 each, eight per group) were compared using software measurements and showed significant contextual discrimination between contexts A and B [A > B; series 1: \( t_{(14)} = 2.65, p = 0.02 \); series 2: \( t_{(14)} = 3.79, p < 0.01 \); series 3: \( t_{(14)} = 2.79, p = 0.01 \)].

Series 3 (Figure 1C) was also scored manually, to allow comparison with yet another context (not described here) which was not located in a Med Associates box. Manual scoring was done by two independent observers, blind to the software scores. Surprisingly, hand-scored freezing (average of observers 1 and 2) did not yield significant differences between contexts A and B \( [t_{(14)} = 1.78, p = 0.10] \). Moreover, software scores were significantly higher than manual scores in context A [74 vs. 66%, i.e., a 8% difference, paired \( t \)-test \( t_{(7)} = 3.93, p < 0.01 \)], while there was virtually no difference between the software and manual scores in context B [48 vs. 49%, paired \( t \)-test \( t_{(7)} = -1.28, p = 0.24 \)].

Because of this finding, we decided to examine inter-rater agreement and to retrospectively reevaluate freezing, also manually, in the two previous series of rats.

First, we investigated our findings in series 3 more thoroughly. The agreement between both human observers was substantial (Landis and Koch, 1977) [Cohen’s kappa, a statistic to assess rater concordance, here using 20 ordered categories (0–5% freezing, 6–10%, etc.) = 0.65], with an average difference of 2.6% freezing. Therefore, we combined both ratings and used the average as the manual score. Correlations between software and hand-scored measurements were high in both contexts (93% in A and 99% in B), but while agreement in context B was substantial (kappa 0.71), it was only poor in context A (kappa 0.05). Additional analyses comparing software scores with those of a human observer for the two previous series of rats (16 rats per context) led to similar conclusions: only slight agreement in context A (kappa 0.18), but moderate agreement in context B (kappa 0.45).

To conclude, we find good agreement between software and manual scores in context B, but not in context A, while using identical software settings.

We therefore decided to reevaluate our camera calibration. Before the start of our studies, both cameras were calibrated in the base setup (without inserts) as described in the manual and discussed with experienced VideoFreeze users, and both cameras showed crisp and well-contrasted images (Figure 1). In addition, both cameras were calibrated using the “Calibrate-Lock” function before each rat. However, it is possible that differences in camera white balance (because of different plastic inserts used in both contexts) caused the observed discrepancies. To investigate this, we adapted the camera white balance in context A until it matched the other context (average grayscale intensity of 119, higher values led to very overexposed images), resulting in a slightly whiter image.

We trained and tested seven more naïve rats in context A and compared software scores with those of two human observers. Two additional rats were excluded from further analyses due to extremely low freezing (\( \leq 5\% \)) on day 2. Unfortunately, adapting the white balance did not resolve the discrepancy between the software and observers’ scores. Software scores were on average still 3% higher than human scores and when removing one
outlying case (Grubb’s test, \(p < 0.05\)) with an average observer score that was 23% higher than the software measure, this difference was even 8%. Analyses indicated that there was still poor agreement between software and observer scores (kappa 0.06).

In conclusion, changing the white balance did not improve agreement between software and manual scores in context A.

Given these findings, we decided to probe into other researchers’ experiences and contacted seven research groups who recently (2011–2013) used the Med Associates software and setup for rat research and who implemented different context configurations in their papers. We asked which software settings they used and how they calibrated their systems. It turns out that there are considerable differences between various labs, as to how they define what the software should consider as freezing (Table 1).

All studies used several floors (different grids and/or plastic floors) and inserts (e.g., black A-frame and/or white curved back wall), as in our own experiments. The applied software settings were quite variable, with motion thresholds ranging from 18 to 150, and a minimum freeze duration of less than 1 s up to 3 s. Some authors used settings that were previously optimized for mice (Anagnostaras et al., 2010; Halladay et al., 2012; Beeman et al., 2013; Broadwater and Spear, 2013a,b), while others performed their own validations for rats (Zelikowsky et al., 2012b). Although not mentioned in their papers, several researchers reported to us that they optimized their parameters as well. J. Long used a 50 or 100 motion threshold depending on the context and in K. Goosens’ lab, the motion threshold usually ranges from 100 to 150, depending on the context configuration and size and strain of the animal (but kept constant for all animals in a certain context on a given test day), and is determined by the experimenter (personal communication).

With regard to the calibration procedure, there was no uniformity either. While half of the research groups (including ours) calibrated the camera before the start of the experiments using a base context, the other half readjusted brightness, gain, and shutter in each context. A quick survey among two labs using VideoFreeze for mice gave a similar picture, with one of both calibrating in each context (Tayler et al., 2011) and the other at initial setup (McDermott et al., 2012). Although some authors mentioned to us that they found that camera calibration can greatly influence the measurements, we did not find meaningful improvements when adapting white balance in our experiments.

Taken together, there is considerable variability in the settings that are being applied by various research groups using the same software and equipment. It is somewhat surprising that rather divergent parameters are being put forward as optimal settings, although this might partially depend on the hand-scoring technique that was used for validation. A recent paper (Shoji et al., 2014) describing a new software package provides hints on how to determine motion thresholds and calibrate in each context. Nevertheless, the question remains whether studies and contexts become more comparable when using different, “optimized” settings in each context or if, conversely, this results in less commensurable measurements. Our finding that the agreement between manual and software measures varied substantially across contexts when using identical settings, is quite alarming. Thus, researchers should be aware of a potential divergence of agreement in different contexts when using a fixed motion threshold and minimum freeze duration. On the one hand, direct comparison of different contexts, e.g., in contextual generalization research, may lead to biased conclusions when using software measurements (in our case artificially increasing differences between contexts A and B). On the other hand, it is self-evident that manual scores by human observers also have drawbacks compared to the objectivity and time-efficiency of automated measurements. We feel that researchers should carefully compare both measurements and decide on the best choice for their research question.

Finally, we would like to stress that in the papers mentioned in Table 1, we see no interpretation problems, as most of these studies did not directly compare freezing between several contexts, and in case they did (Zelikowsky et al., 2012b), contexts were counterbalanced. Note however that, theoretically, even limited measurement deviations between contexts may induce heightened variability when counterbalancing contexts, thereby decreasing the chance of finding significant effects.

| Motion threshold | Minimum freeze duration (seconds) | Camera calibration | References |
|------------------|-----------------------------------|--------------------|------------|
| 18               | 1                                 | In each context    | Beeman et al., 2013 |
| 18               | 1                                 | In standard context | Broadwater and Spear, 2013a,b |
| 20               | 1.07                              | In standard context | Moffett et al., 2011 |
| 50\#             | 1\#                               | In standard context | Zelikowsky et al., 2012b |
| 50 or 100        | 0.77                              | In each context    | Long et al., 2011 |
| 120              | 1                                 | In each context    | Vander Weeke et al., 2013 |
| 150\#            | 3\#                               | In each context    | Sticht et al., 2012 |

*Most information was obtained through personal communication with the authors. Settings indicated with # were mentioned in the corresponding publications.*

**Table 1 | Software settings and calibration procedures (adjustment of brightness, gain, and shutter) in seven research groups who use VideoFreeze software and several context configurations for rat conditioning studies.**

**CONCLUSION**

While each researcher should balance the (dis)advantages of automated and manual scoring against one another, we believe that caution is required when using software measurements, particularly when comparing different context configurations or in case of subtle behavioral effects.

**AUTHOR CONTRIBUTIONS**

Laura Luyten designed the experiments, partially conducted them, analyzed and interpreted all data and wrote the manuscript. Natalie Schroyens carried out part of the studies and contributed to the analyses and manuscript draft. Laura Luyten and Natalie Schroyens manually scored the freezing videos. Dirk Hermans and Tom Beckers contributed to the conceptual design and
manuscript draft. All authors approved the final version of the manuscript.

ACKNOWLEDGMENTS

This work was supported by the Research Foundation—Flanders (FWO) Project G072909N, Research Grant (to Laura Luyten) 1504614N and KU Leuven grant PF/10/005. Laura Luyten is a postdoctoral fellow of the FWO. The authors would like to thank all researchers who provided information about their methods.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Received: 28 November 2013; accepted: 20 January 2014; published online: 04 February 2014.

Citation: Luyten, L., Schroyens, N., Hermans, D. and Beckers, T. (2014) Parameter optimization for automated behavior assessment: plug-and-play or trial-and-error? Front. Behav. Neurosci. 8:28. doi: 10.3389/fnbeh.2014.00028

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