



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

## Experiential contributions to social dominance in a rat model of Fragile-X syndrome

**Citation for published version:**

Saxena, K, Webster, J, Hallas-Potts, AM, MacKenzie, R, Spooner, PA, Thomson, D, Kind, P, Chattarji, S & Morris, R 2018, 'Experiential contributions to social dominance in a rat model of Fragile-X syndrome', *Proceedings of the Royal Society B-Biological Sciences*, vol. 285, no. 1880.  
<https://doi.org/10.1098/rspb.2018.0294>

**Digital Object Identifier (DOI):**

[10.1098/rspb.2018.0294](https://doi.org/10.1098/rspb.2018.0294)

**Link:**

[Link to publication record in Edinburgh Research Explorer](#)

**Document Version:**

Peer reviewed version

**Published In:**

Proceedings of the Royal Society B-Biological Sciences

**General rights**

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

**Take down policy**

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact [openaccess@ed.ac.uk](mailto:openaccess@ed.ac.uk) providing details, and we will remove access to the work immediately and investigate your claim.



**Experiential contributions to social dominance in a  
rat model of Fragile-X syndrome**

Journal:	<i>Proceedings B</i>
Manuscript ID	RSPB-2018-0294
Article Type:	Research
Date Submitted by the Author:	06-Feb-2018
Complete List of Authors:	Saxena, Kapil; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience Webster, Jemma; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience A.M.Hallas-Potts@sms.ed.ac.uk, Amelia; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience MacKenzie, Ruairi; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience Spooner, Patrick; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience Thomson, Derek; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience Kind, Peter; University of Edinburgh, Simons Center for the Developing Brain Chattarji, Sumantra; National Centre for Biological Sciences, Centre for Brain Development and Repair MORRIS, Richard; The University of Edinburgh, Centre for Cognitive and Neural Systems, Edinburgh Neuroscience
Subject:	Behaviour < BIOLOGY, Neuroscience < BIOLOGY
Keywords:	Fragile-X, Social Dominance, Hierarchy, Gene-environment interactions, Fmr1, Tube test
Proceedings B category:	Neuroscience & Cognition

Dr FNL - Jan 2018

## **Experiential contributions to social dominance in a rat model of Fragile-X syndrome**

K. Saxena, J. Webster, A. Hallas-Potts, R. Mackenzie,  
P.A.Spooner, D. Thomson, P. Kind, S. Chatterji  
and R.G.M. Morris

Simons Initiative for the Developing Brain and Centre for Cognitive and Neural Systems,  
Edinburgh Neuroscience, The University of Edinburgh, 1 George Square, EH8 9JZ  
*and*  
Centre for Brain Developmental and Repair, NCBS-InSTEM, Bangalore

### **Corresponding Author:**

R.G.M. Morris ([r.g.m.morris@ed.ac.uk](mailto:r.g.m.morris@ed.ac.uk)) - Edinburgh Neuroscience

### **Key Words:**

Fragile-X, Social Dominance, Hierarchy, *Fmr1*, Autism spectrum disorders, gene-environment interactions

## Abstract

Social withdrawal is one phenotypic feature of the monogenic neurodevelopmental disorder Fragile-X. Using a ‘knock-out’ rat model of Fragile-X, we examined whether deletion of the *Fmr1* gene that causes this condition would affect the ability to form and express a social hierarchy as measured in a tube-test. Male Fragile-X ‘knock-out’ rats living together could successfully form a social dominance hierarchy, but were significantly subordinate to wild-type animals in mixed group cages. Over 10 days of repeated testing, the Fragile-X mutant rats gradually showed greater variance and instability of rank during their tube-test encounters. This affected the outcome of future encounters with stranger animals from other cages, with the initial phenotype of wild-type dominance lost to a more complex picture that reflected, regardless of genotype, the prior experience of winning or losing. Our findings offer a novel insight into the complex dynamics of social interactions between laboratory living groups of Fragile X and wild-type rats. Even though this is a monogenic condition, experience has an impact upon future interactions with other animals. Gene/environment interactions should therefore be considered in the development of therapeutics.

(1252 characters)

*Key words:* Social Hierarchy, social dominance, Tube-Test, Fragile-X, Rat.

## Introduction

The aim of this study was to investigate whether deletion of the Fragile-X mental retardation protein (FMRP) would impact the expression of a social hierarchy in group-living laboratory rats. Fragile-X typically arises via CGG trinucleotide repeats in its promotor region of *Fmr1*, the X-linked gene that encodes FMRP, and a leading monogenic cause of inherited learning disability and autism (1, 2). Heterogeneous clinical symptoms are observed, including macro-orchidism (enlarged testes), abnormal facial features and a high prevalence of epilepsy; other phenotypic features can include alterations in aggressive behaviour, attentional and cognitive deficits and severe mental retardation (3, 4). Social withdrawal is also a prominent characteristic. New treatments are urgently sought for this autism-spectrum disorder.

Animal models of Fragile-X began with studies using mutant mice (5) and only more recently turned to the use of rats. One focus has been on the hypothesis that deletion of *Fmr1* affects signalling downstream of metabotropic glutamate receptors (mGluR). Specifically, synaptic changes have been observed in mice that mimic the human syndrome including alterations in dendritic spines (6), an exaggeration of the capacity to induce or sustain activity-dependent long-term depression (LTD) of synaptic efficacy in the hippocampus (7, 8), and alterations in long-term potentiation (LTP) in the amygdala (9, 10). Behavioural changes are seen when the mice are tested singly, including deficits in 'flexibility' such as repetitive behaviours (11, 12), a reversal deficit in the watermaze (5), and reactivity to novelty when objects are experienced in changed places and testing contexts (13). These behavioural changes may be linked to the underlying changes in activity-dependent synaptic plasticity that mediates the capacity to learn, remember, and update stored information in response to change.

Some animal studies have also directly examined the social phenotypes of human Fragile-X such as social withdrawal, social anxiety and difficulty in forming peer relationships. Impairments in social communication and social interactions have been reported (12, 14-17), reflecting construct validity, as well as rodent-characteristic social behaviour such as ultrasound communication (18). However, a key issue in examining any aspect of social behaviour is establishing the relative influence of genetic and environmental factors in the determination of phenotype because social interactions offer the opportunity for

transitivity of dominance relationships, and stability or change over time (12). It is known that a rat's age is a significant correlate of success in dominance interactions by virtue of an animal's memory of previous social interactions with another animal (19). This memory may be the basis of both 'transitivity' ( $A > B$ ,  $B > C$ , and  $A > C$ ) and the stability of such inter-animal dominance relationships.

Inspired by a prominent study of causal mechanisms contributing to social dominance in mice by (20), we chose to study 'social dominance' and 'social hierarchy' as relatively tractable indices of social interactions using the 'tube test' (21). We sought to contrast *direct* effects of the FMRP mutation on the expression of a social hierarchy, with *indirect* effects mediated by an individual animal's experience of its rank within its social group across repeated testing sessions. The simplest question is: does a cage of male  $Fmr1^{-/y}$  rats living together develop a social hierarchy? Second, would wild-type rats be dominant in a social group consisting of both lines living together with  $Fmr1^{-/y}$  rats? Third, beyond measurement of an individual's 'rank' in such hierarchies, would  $Fmr1^{-/y}$  rats show differential variability in rank over time? That is, might normal rats display stability by virtue of remembering previous encounters, but  $Fmr1^{-/y}$  rats display a more variable pattern? Fourth, what would be the impact of an individual animal's experience of his rank upon his subsequent social interactions with a stranger animal (i.e. from a different cage)? That is, what would be the impact on future contests with strangers of a rat being highly- or lowly ranked in his home cage? Would genotype or relative rank in the home-cage dominate in determining the outcome? Pertinent to these questions is that a new study in mice suggests that synaptic plasticity on a thalamic-prefrontal pathway in the brain of mice, reflecting the history of winning or losing social contests, can be causally involved in determining an animal's rank (22).

We tested a new line of  $Fmr1$  knock-out rats on a Long-Evans background originally developed in conjunction with Sage/Horizon (LEH rats (23)). This line was made using Zinc finger nuclease (ZFN)-mediated disruption of  $Fmr1$ . In a first report, Asiminas et al (24) describe in detail: 1) the target site for ZFN cleavage, 2) the donor sequence including eGFP and a nuclear localisation signal (NLS) flanked by 5' and 3' homology recombination arms for homology directed repair (HDR), and 3) the resulting targeted locus. Immunohistochemical localisation of FMRP in P15 WT and  $Fmr1$  KO rats shows the absence of FMRP in transgenic rats. This line has been calibrated by our colleagues as showing

several of the same aspects of phenotype as previously observed in a different line of rat (13). The switch from mouse to the rat is because rats display complex patterns of social behaviour that have been extensively described (25). Our hope was that, by using this more social species, we might thereby create an alternative and potentially useful rodent model of the social anxiety phenotype of Fragile-X humans.

## Methods

### Subjects

The subjects were adult male Long-Evans hooded rats (n=56), weighing between 320 and 680 gm across the three separate cohorts (2 tested at 3 months, 1 at 6 months). Rats of similar weights were group housed, 4 rats in a cage since weaning, with *ad libitum* access to food and water. A 12h light and dark cycle was maintained with all the experiments carried out in the light phase between 9 am to 5 pm. *Fmr1*<sup>-y</sup> male and wildtype littermate control rats were generated by mating female *Fmr1* heterozygotes crossed to wild-type Long-Evans hooded rats (Charles River Labs). The wild-type (WT) rats were littermate matched.

The experiment was conducted in 3 successive cohorts over 2 years by KS, JW, RB and AH-P. In each cohort, the rats were housed in 3 major groups: WT only, KO only and mixed WT and KO. In each case, there were 4 rats together in a cage, with the mixed group composed of 2 WT and 2 KO animals. In total, there were 56 rats divided between 3 WT cages, 4 KO cages and 7 Mixed cages. In all the cases, the experimenters were blind to the genotype of the subjects, the animals in each cage being colour-coded on their tails, with the code made and retained by independent person (DT).

### Apparatus

The tube-test was adapted for rats from the study of Wang et al (20). A transparent plastic tube 1 m in length (small tube: 6 cm internal diameter; large tube: 7 cm-internal diameter) was placed between two holding boxes (42x26x18cm; Figure 1A). The tube size was selected so that rats could easily run down the tube, but were unable to pass each other. Removable barriers were placed at the entrances of the tube from each holding box, and at a central location. Bedding from the rats' home cage was placed in each holding box. The tube and boxes were placed on a table in a laboratory room with subdued lighting, the

experimenter at a table facing the equipment. A video system was used to record selected trials for illustrative purposes only.

*Insert Figure 1 about here*

## **Protocols**

*Habituation:* Each cage was handled daily for a minimum of 3 days prior to testing. During a single day of habituation, all animals in a cage were placed singly in the apparatus and allowed to run freely in the tube between the holding boxes for 30 min. All barriers were removed. Small sections of tubing had also been placed in all home cages 2 weeks prior to testing to help familiarise the animals.

*The tube-test of social dominance:* Two rats were tested by placing one rat at either end of the tube in their respective holding boxes. The entrance barriers were then lifted, and the rats introduced into the tube whereupon they generally moved to the centre. Once both rats touched the central barrier, it was raised to allow the dominance test to begin. The rats competed for dominance in the tube in various ways ranging from submission and retreat by one of the animals through to direct physical pushing of one opponent backwards out of the tube. “Winning” was defined as reaching the entrance barrier slit at either end of the tube. There was never any ambiguity about which rat had “won”. Each pair of rats were run 5 times, with the winning rat recorded each time, with the side on which each rat started swapped between successive runs (to counterbalance left and right). The time between lifting the middle barrier and the submissive animal leaving the tube was recorded as the latency (s). The apparatus was cleaned with an amphoteric disinfectant between cages but not between all encounters. Cage order was varied so that start times per cage changed between trials. A random order generator assigned pair orders within cages, which changed between sessions. In each session, fodder from the home cage was placed in the holding boxes to mirror the home-cage environment.

*Phase I - Dominance hierarchies:* The tube test was initially used to establish within-cage dominance hierarchies (Fig 1B Phase I). In each session, every animal competed against all 3 of its cage-mates. Each pair was run in the tube 5 consecutive times. A rat winning at least 3 times in a 5-trial block was declared a clear winner, but testing always continued for 5 trials. A winning rat was allocated 2 points and the loser rat given 0 points per trial. Tallying



up these points across all 5 trials allowed dominance ranking of the 4 cage-mate rats. In case of equal points, the ranking was decided based on the outcome of the specific competition between the two equal scoring individuals, with the winner ranked higher. This test of position in the hierarchy was repeated 10 times across successive sessions.

*Phase II - Inter-cage tournaments:* The second phase was to assess dominance in novel social conflicts (Figure 1B Phase II). Because the full experiment was conducted as 3 separate cohorts, we could only test a subset of the inter-cage competitions. Four combinations were tested as follows: WT vs KO (2 cages), Mixed vs Mixed (3 cages), WT vs Mixed (2 cages), and KO vs Mixed (2 cages). Each rat competed against every animal from the paired cage (total 16 pairs), with 5 runs per pair per day. A random order generator established pair-order as well as cage-order. Each tournament was repeated 3 times across days. The tube was changed between pairs and fresh bedding was laid down in the holding cages for each tournament. Before testing, one cage was allowed to freely explore the tube for 5 min, followed by the second for 5 min and the first again for 2 min, ensuring no particular animals/cages smell dominated the environment. The cages were renumbered by a third party to keep the experimenter blind.

### **Statistical Analyses**

Test for the statistical significance were used. A Chi-square test was used to compare the expected and observed values of number of wins between genotypes. A t-test was used to compare average rank and wins per pair between WT and KO in mixed cages. Where the statistical variance of rank and/or a simple change/no-change measure of rank stability is presented, a t-test was used to test the level of significance. For the variance scores, we adjusted the values to a percentage score in which 100% was the maximal variance score possible. Due to presence of two different genotypes in the mixed cages, their analysis was processed separately to that of the single-line WT and KO cages. The Pearson correlation-coefficient was also computed to examine stability across sessions. The significance level is quoted for all tests. All the plots are represented as mean  $\pm$  SEM.

## **Results**

### *Qualitative observations*

The rats behaved appropriately in the tube test, readily walking down the tube to the centre, confronting their 'opponent', investigating him and interacting socially. Upon opening

the partition separating the two sides, the two animals investigated each other and then either advanced or retreated as appropriate (Figure 1). Very little aggressive behaviour was observed. Latencies to complete individual tests typically ranged from 3 to 30 s (rarely longer), with considerable within and between animal variability (such that latency did not prove to be a useful measure). As described in the Methods, the contests between two individuals were repeated 5 times each day to ascertain a clear “winner” of each session.

### *Phase 1: Dominance hierarchies*

The first result, secured over 10 successive sessions of testing, was that a clear social hierarchy was apparent in both the single-line cage groups containing only wild-type (WT) or only *Fmr1*<sup>-y</sup> KO animals (Figure 2). Panels A and B show the stable and shifting rank of animals in two representative single-line cages over the full 10 days of testing; Panel A displays an example of a wild-type cage that was particularly stable, whereas Panel B shows a KO cage displaying apparently greater day-to-day variability of rank. From the full set of 7 single-line cages, the average rank over 10 days collectively revealed a clear hierarchy in both lines; the data is plotted to show the highest (top), lowest (bottom), and the two mid-ranked animals across cages (Panels C,D).

*Insert Figure 2 about here*

Second, we then sought to measure the apparent variability in rank across sessions. Two separate measures were used: (a) *Variance* - the statistical measure of variability of absolute rank; (b) *Stability* - measured as a count of the number of times an animal changed its rank between one session and the next, without regard to extent of the change (as used by the Hu laboratory (20)). Panel E shows that percent variance was significantly lower in the WT single-line cages than in the KO cages ( $t=2.99$ ,  $df=26$ ,  $p=0.006$ ). Stability showed a similar trend (Panel H), although the change did not reach statistical significance ( $t=2.05$ ,  $df=26$ ,  $p=0.054$ ). We then examined whether variance and stability changed across the 10 sessions. With each measure calculated separately for sessions 1-5 and 6-10 (Panels F,G,I,J), the WT rats showed a significant correlation of rank stability across the two sub-sets of 5 sessions (single-line cages,  $r=0.69$ ,  $p=0.01$ ), whereas the KO rats did not ( $r=0.39$ ,  $p=0.13$ ). The corresponding data for variance is difficult to interpret as the ‘clumping’ of the very low variance scores for the WT rats differs from the much more dispersed scores of the KO rats

(Panels F,G). In both cases, the trend was for the slope of the regression line to be lower than  $45^\circ$ , pointing to a gradual assumption of stability across repeated tube-test sessions.

*Insert Figure 3 about here*

We then, third, examined the mixed-line cages in each of which 2 WT and 2 KO rats had been living together for several weeks. The key finding was that WT rats won significantly more contests than KO animals (Figure 3). This is shown for a representative cage (Panel A), in the analyses of the overall number of contests (chi-square=14.63, df 1,  $p=0.0001$ ; Panel B), and for the average rank of individual WT (2.27) and KO (2.72) animals calculated across all 280 contests ( $t=3.48$ , df 278,  $p=0.0006$ ; Panel C). We wondered whether there was any relationship between rank and weight of the animal, surmising that larger animals might be more inclined to win. The concern was that, in comparing WTs and KOs in animals living together, the comparison of genotypes might be secondary to a difference in weight. However, not only was there no relationship between weight and rank ( $r=0.057$ , NS; data not shown), the mean weight of the WT rats (468.4 gm) was, if anything, slightly lower than that of the KO animals (488.8 gm; NS). Thus, the apparent dominance of the WT over the KO animals is not an artefact of greater weight.

As in the single-line cages, both variance and stability were computed across the ten sessions, and separately for sessions 1-5 and 6-10. No overall mean differences were observed (Panels D, G). However, in examining change across the two sub-sets of 5 sessions, only the WT animals displayed significant correlations of variance ( $r=0.55$ ,  $p=0.04$ ) and stability ( $r=0.56$ ,  $p=0.03$ ). The KO rats showed non-significant regression coefficients ( $r=0.17$  and  $r=0.11$ , N.S.), reflecting instability of rank, but interestingly, with correlation slopes that were well below  $45^\circ$ . Strikingly, the KO rats subdivided across sessions into 'hyper-stable' ( $n=8$ ) and 'hyper-unstable' ( $n=6$ ) sub-groups (note that such stability and instability may reflect either a high or low absolute rank, Panel I).

Thus, KO rats have a phenotype reflecting lower dominance than cage-mate wild-type rats, reflecting social withdrawal, but apparently greater variability in their experience of tube-test contests that may affect their future interactions across sessions.

### *Phase 2: Inter-cage Tournaments*

The second phase of experimentation involved ‘tournaments’ in which all animals of a given cage were, over three sessions, pitted against all animals of another cage, with 5 trials per pair per session (Figure 4). These contests were therefore always with ‘stranger’ animals (note that this could not be done for all 14 cages of Phase 1 as the study was conducted in 3 replications; during any one replication, the subset of cages that could be tested was against others of that same replication). There were, nonetheless, a total of 1320 contests (sufficient for statistical reliability). This phase allowed us to ask, over all permutations, whether the dominance rank an animal had assumed by the end of Phase 1 was predictive of his behaviour in the tournament tests of Phase 2.

*Insert Figure 4 about here*

Surprisingly, comparison of WT and KO rats now revealed no difference as a function of genotype (Figure 4A; chi-square=0.11, df=1,  $p=0.74$ , NS); the phenotype displayed in Phase 1 appeared to be lost. Comparison of the cage tournaments between only the single-line cages revealed, paradoxically, that KO rats won *more* contests against WT rats (chi-square=7.00, df=1,  $p=0.0081$ ; Panel B). The situation for the mixed cages, for which the KO rats would have had more experience of losing than WT rats, was more evenly matched (chi-square=0.54, df=1,  $p=0.46$ , NS; Panel C). In keeping with an emerging pattern, when single-line WT rats were pitched against mixed-cage KO rats, there was a significant dominance of the WT animals (chi-square=27, df=1,  $p<0.0001$ ; Panel D). This might have been expected to continue for the comparison of WT rats of mixed-line cages vs KO single-line cages (Panel E), but this was not observed - possibly because WT rats that had experience of KO rats were for some reason wary of stranger KO rats (chi-square=0.06, df=1,  $p=0.79$ ; NS; Panel E).

The impact of experience gained from the contests of first phase was most striking when the same data of Figure 4 were re-examined with regard to Phase 1 rank as well as genetic status (Figure 5). Panel A shows a cartoon of the rank (1-4) of the 4 rats in a given cage (Cages 1 and 2) with each animal of Cage 1 tested against each ‘stranger’ animal of Cage 2 (graphically representative example shown for the rank 3 animal of Cage 1). The Phase 1 rank was partially predictive of such contests because, when animals of both

genotypes were pooled together regardless of genotype into a sub-group that had first or second rank in Phase 1 (high ranked, Panel A) and had gone through the contest with the sub-group of third or fourth ranked animals (low ranked) (i.e. excluding higher vs. higher and lower vs. lower), the higher ranked individuals won significantly more contests (Chi-square=102, df=1,  $p<0.0001$ ; Figure 4B). While such a result may seem trivial, this suggests that the experience of winning or losing over the 10 days of testing in Phase 1 has consequences for future interactions with the strangers.

It then became of interest to ask, the outcome with respect to genetic status of stranger-contests between higher ranked and separately between lower ranked animals. Strikingly, KO high-rank rats won significantly more often against WT high-rank animals (Chi-square= 39, df= 1,  $p=0.0001$ ; Panel C), but this pattern was reversed in the comparison of the WT and KO low-ranked animals (Chi-square= 5.3, df= 1,  $p=0.02$ ; Panel D). Note the number of contests in panels C and D are necessarily half those of panels B, which is half of overall contest number of 1320 (Panel A, Figure 4) as it (Panel B; Figure 5) excludes the competition between the Higher vs Higher and Lower vs Lower.

## Discussion

Previous studies of social dominance in animal models of autism spectrum disorders have focused on mice due the relative ease of making deletions of murine genes by homologous recombination (12, 26). One novel feature of this study is the use of the rat following the development of this ‘knock-out’ rat-model of Fragile-X syndrome (24). Accordingly, we sought to examine whether deletion of FMRP would have an impact on (a) the ability to form and express a social hierarchy as measured in a tube-test, (b) social dominance behaviour when animals are confronted by another rat living in the same cage, and (c) social dominance when confronted by a stranger rat from another cage.

### *The tube-test measure of hierarchy*

Our data indicate that male *Fmr1*<sup>-y</sup> rats living together in cage groups of 4 animals readily form a social hierarchy that can be measured reliably using the tube test (which is known to correlate well with several other measures of social dominance in mice (20). However, in groups of animals consisting of 2 WT and 2 *Fmr1*<sup>-y</sup> rats, the WT rats were

significantly more dominant, although not exclusively so on a rat-by-rat basis. Thus, there is a clear social dominance phenotype associated with this rat model of Fragile-X that may correspond to aspects of the social anxiety and/or social withdrawal of affected individuals when confronted by others.

A single tube test encounter has only one output - one animal is a winner (dominant) and the other a loser (subordinate). By testing every member of a cage with all others, and doing so several times within a session (to ensure within-day reproducibility), we could decipher at least one measure of the social hierarchy in that cage and how it evolved over successive days of testing. Other tests examined by Wang et al. (20) include whisker trimming also known as “Dalila effect” but this phenomenon is observed in mice but not in rats (27). A urine marking assay, in which the so-called dominant animals are observed to mark a larger territory than subordinates, was a further possibility; however, one study observed that male rats spend more time adjacent to the urine of other male rats than that of an odourless substance, pointing to a lack of avoidance of urine marking under some conditions (28). It has also been observed that sniffing frequency decreases in the subordinate animal of a pair (29), and we could examine this in a follow-up study (see below). The extent of access to food or water in a group cage is potentially another measure, but was not suitable for us as it requires the food and/or water deprivation which we did not use; resource competition can, however, be well correlated with the tube-test in rats (30). As these measures, where they could be measured, correlated well for mice (see Wang et al [18]), we believe that our focus on the tube-test is justified.

#### *Interaction of genetic and experiential factors*

This study revealed the impact of both genetic and, to a limited extent, experiential factors. One aspect of the latter was repeated testing over several days, and this revealed greater dominance stability in WT rats than *Fmr1* KO animals. Moreover, the rank held by a WT rat during the first 5 days of testing was predictive of its rank during the second 5 days of testing in both single-line and mixed-line cages. This is not surprising as rats readily form a stable hierarchy - in laboratory conditions (31) and in the (semi) wild (19). Stability is also promoted under laboratory conditions when group-living animals must compete for food resources from the time of weaning (32). In contrast, *Fmr1* KO animals displayed changes in stability over the course of the 10 days of testing. The second experiential aspect we

examined was the impact on social dominance of prior experience of testing with familiar home-cage animals upon later tournaments with stranger rats. This revealed an apparent loss of the earlier identified genetic phenotype, with social dominance behaviour now dominated by the animals' prior experience of having won or lost earlier contests. That is, rats of either genotype ranking relatively highly in Phase 1 won more Phase 2 contests against animals ranked relatively lowly (although again not exclusively). This is consistent with the memory of 'previous outcome' concept with respect to social dominance (19). However, when this expected pattern was broken down into sub-groups reflecting both genotype and relative rank, a more complex picture emerged that, if anything, constituted a partial reversal of the earlier phenotype.

One intriguing possibility is that KO rats fail to process the social cues from the opposing animal as effectively as do WT rats. Thus, in considering the pattern shown during the later contests with stranger rats, we suspect that an important aspect of the animals' experience was not just rank but the relative variance or stability of the rank, with a phenotype clearly emerging to the effect that the WT rats showed greater stability (66%) than KO rats (49%). These absolute levels are commensurate to those reported for one study in mice that measured stability across testing sessions (59% stability, see (20)). However, as the testing sessions of Phase 1 continued, the KO rats showed a bimodal pattern of becoming either 'hyper-stable' or 'hyper-unstable'. As noted in the Methods, our observations and scoring of the animals were conducted blind to genotype, but we sensed that some animals behaved as though sensitive to cues from the other animal whereas other animals behaved as if they were not. If there is a genotype-associated difference in the processing of social cues, KO rats may develop the behavioural patterns of either pushing forward regardless ('bully' strategy) or retreating the moment another animal is experienced ('submissive' strategy). Both of these patterns would then populate the sub-group showing 'hyper-stability' of high-rank and low-rank respectively. If this analysis is correct, it would be expected that high-rank KO rats displaying hyper-stability would win over stranger WT rats, even highly-ranked WT rats whereas low-ranked KO rats displaying hyper-stability would lose in contests with stranger WT rats, even lowly-ranked WT rats. This is likely to have contributed to the pattern that was observed. In future research, it would therefore be valuable to video carefully all encounters between animals in the tube-test, as done with mice (22, 33), with a view to objectively identifying patterns of behaviour that may reflect differential sensory processing (e.g. whisker movements), motor movements related to advance or retreat, and social

interactions (e.g. head contact, overt aggression). This would be demanding to do because it would likely require a replication and video capture of all conditions described here, again conducted and filmed ‘blind’ over hundreds of encounters, but then decoded with respect to genetic status and previously established rank.

Studies of social dominance in mice were the first to establish that *Fmr1* KO mice displayed abnormalities in the tube-test (14). Paylor’s pioneering study also showed that experience matters when studying encounters between familiar (cage-mate) and unfamiliar animals, and between mice that had experience of tube-test encounters compared to mice on the first day of testing. Spencer et al (14) observed either dominance by WT animals or equivalence of the two genotypes as a function of these experiential factors, a finding replicated by Goebel-Goody et al. (34). More recently, however, a study by de Esch and colleagues (35) reported a variety of tube-test outcomes, as a function of differential housing and experience. Unlike our cage-mate results (Phase 1), they observed greater dominance by *Fmr1* KO over WT mice that was highly significant. Dominance increased and stabilised over 5 days of testing, but is largely the opposite of our findings with rats. They went on to establish that knocking out the mGLUR5 sub-unit of metabotropic glutamate receptors in *Fmr1* KO mice had a dramatic effect. Specifically, *Fmr1* KO animals won tube-test encounters against animals with the double knock-out, the double knock-out animals won against WTs, but the WTs won against animals with only the mGLUR5 knock-out. We are uncertain what to conclude from this pattern of results except to agree with Spencer et al’s (14) assertion that: “*in future investigations....*” of *Fmr1* KO animals, it will be necessary “*...to incorporate experimental designs to clearly assess the influence of test experience on behaviour...*” (page 428). Whether the use of rats (as here) rather than mice will help in this task remains to be established, but a strong case has been made for their use now that technologies for rat genetic interventions are becoming available (36). From the perspective of brain mechanisms, (22) have recently provided that neural activation of the dorso-medial prefrontal cortex (dmPFC) is both necessary and sufficient to induce dominance in tube-test social competitions. The causal evidence derives from both chemogenetic inhibition studies and optogenetic activation experiments directed at a thalamic-prefrontal pathway. In a further demonstration of the validity of measures from the tube-test, transfer from it to a “warm-spot” test in a cage with an otherwise ice-cold floor was shown to display a significant correlation of these two measures of rank. They note that “*an important parameter for the*



*cost-benefit computation in a social confrontation is the history of winning .... our results might shed light on the treatment of .... psychiatric diseases.”*

There is growing interest in the interaction of genetic and environmental influences on the cognitive phenotype of Fragile-X children (37), and wider interest in psychiatry (17). It would seem to be of value to continue our analysis of gene-environment interactions in this rat model of Fragile-X, diversifying into a wider set of tests of social interaction followed by exploring the impact of pharmacological treatments that target abnormalities in mGLUR signalling.

### **Acknowledgements**

This work was supported by an Erasmus Mundus EXPERTS - SUSTAIN (2016-2017) scholarship to KS, a Royal Society Commonwealth Conference Travel Award to RM and KS, the Patrick Wild Centre (University of Edinburgh) and the Centre for Brain Development and Repair (NCBS, InStem, Bangalore).

## Figure Legends

**Figure 1: *Experimental apparatus and design.*** **A.** An acrylic (Perspex) tube 1 m in length connected two holding boxes containing ‘bedding’ from the home cages of the animals being tested. Three separators limited access to the tube from the cages (1 and 2), and at the centre (3), contact between the two animals. A trial commenced when the central separator was lifted. **B.** The design involved two phases. Phase 1 involved measurement of the relative rank and variability in rank within each of the cages. Phase 2 involved competitions between each animal of one cage against each animal of another cage (e.g. rat 3 of cage 1 against each of the 4 animals of Cage 2). Note colour-coding of the animals with a tail mark.

**Figure 2: *Cage hierarchies - single-line cages.*** **A,B.** Representative session rankings of individual colour tail-marked rats in a single line WT cage (blue) and a single line KO cage (red) across 10 days of testing. Note greater instability of rank of the KO rats. **C,D.** Mean rank of rats as a function of their ordinal position in a cage hierarchy averaged across ten days. Note clear hierarchy of WT single line (3 cages) and KO single line (4 cages). **E-G.** Mean percentage variance of rank for WT and KO rats across 10 sessions of testing, and correlation between variance observed during sessions 1-5 and 6-10 for WT and KO rats respectively. **H-J.** Mean stability for WT and KO rats across the same sessions as E-G. \*\*  $p < 0.01$ ; Means  $\pm$  1SEM.

**Figure 3: *Cage hierarchies - mixed-line cages.*** **A.** Session rankings of individual colour tail-marked rats in a representative mixed-line cage across 10 days of testing (colour-coding as in Figure 2). Note relative but not absolute dominance of WT rats. **B,C.** Number of wins of tube-test contests by genotype and the distinct but non-independent measure of mean rank. See text for chi-square statistics. **D-F.** Mean percent variance of rank for WT and KO rats across 10 sessions of testing, and correlation between variance observed during sessions 1-5 and 6-10 for WT and KO rats respectively. **G-I.** Mean stability for WT and KO rats across the same sessions as D-F. Note that WT rats show inter-session predictability between s1-5 and s6-10, whereas KO rats in these mixed cages divide bi-modally into two sub-groups with low and high stability over s6-10. \*\*\*  $p < 0.001$ ; Means  $\pm$  1SEM.

**Figure 4:** *Cage tournaments - organised by genotype.* **A.** When all WT and KO rats from different cages met in contests in each of the three cohorts (resulting in 1320 contests), there was a loss of the phenotype that in Phase 1 had reflected dominance by WTs. Hatched colour coding reflects pooled data from both single-line and mixed-line cages. **B.** Significant dominance of KO rats in single-line cages in contests with equivalent single-line WT rats. **C.** No dominance pattern when the mixed-line cage animals competed against each other. **D.** WT rats from single-line cages were dominant over KO mixed-line rats. **E.** No dominance pattern when the WT mixed-line cage animals competed against KO single-line rats. \*\*\*\*  $p < 0.0001$ ; \*\*  $p < 0.01$ ; Means  $\pm$  1SEM.

**Figure 5:** *Cage tournaments - organised by experience of winning.* **A.** The data of Figure 4 were re-examined taking into account the rank assumed in phase 1 of the study (contests with cage-mates). All pairwise contests between cages were tested (3 times) and the cartoon depicts a rank 3 animal in cage 1 competing against all rats of cage 2. **B.** Rats that had previously been higher ranked (Ranks 1 and 2) were dominant over stranger rats that had been lower ranked (Ranks 3 and 4), when scores were computed without regard to genotype. **C,D.** Sub-sets of the cage tournament data analysed only for the contests between Higher ranked WT vs. Higher ranked KO animals, and separately for Low-ranked WT vs. Low-ranked KO animals. When previous ranking and genetic line were considered together, the dominance by KO rats in Phase 1 was specific to the higher-ranked animals. For lower ranked animals, WT rats were dominant were marginally but significantly dominant. \*\*\*\*  $p < 0.0001$ ; \*  $p < 0.05$ ; Means  $\pm$  1SEM.

## References

1. Turner G, Webb T, Wake S, Robinson H. Prevalence of fragile X syndrome. *Am J Med Genet.* 1996;64(1):196-7.
2. Sherman SL, Marsteller F, Abramowitz AJ, Scott E, Leslie M, Bregman J. Cognitive and behavioral performance among FMR1 high-repeat allele carriers surveyed from special education classes. *Am J Med Genet.* 2002;114(4):458-65.
3. Hagerman RJ, Hagerman PJ. The fragile X premutation: into the phenotypic fold. *Curr Opin Genet Dev.* 2002;12(3):278-83.
4. Hagerman RJ, Berry-Kravis E, Kaufmann WE, Ono MY, Tartaglia N, Lachiewicz A, et al. Advances in the treatment of fragile X syndrome. *Pediatrics.* 2009;123(1):378-90.
5. Bakker CE, Verheij C, Willemsen R, van der Helm R, Oerlemans F, Vermey M, et al. Fmr1 knockout mice: a model to study fragile X mental retardation. *Cell.* 1994;78(1):23-33.
6. Comery TA, Harris JB, Willems PJ, Oostra BA, Irwin SA, Weiler IJ, et al. Abnormal dendritic spines in fragile X knockout mice: maturation and pruning deficits. *Proceedings of the National Academy of Sciences.* 1997;94(10):5401-4.
7. Bear MF, Huber KM, Warren ST. The mGluR theory of fragile X mental retardation. *Trends Neurosci.* 2004;27(7):370-7.
8. Dolen G, Bear MF. Role for metabotropic glutamate receptor 5 (mGluR5) in the pathogenesis of fragile X syndrome. *J Physiol.* 2008;586(6):1503-8.
9. Suvrathan A, Hoeffler CA, Wong H, Klann E, Chattarji S. Characterization and reversal of synaptic defects in the amygdala in a mouse model of fragile X syndrome. *Proc Natl Acad Sci U S A.* 2010;107(25):11591-6.
10. Suvrathan A, Chattarji S. Fragile X syndrome and the amygdala. *Curr Opin Neurobiol.* 2011;21(3):509-15.
11. Spencer CM, Alekseyenko O, Hamilton SM, Thomas AM, Serysheva E, Yuva-Paylor LA, et al. Modifying behavioral phenotypes in Fmr1KO mice: genetic background differences reveal autistic-like responses. *Autism Res.* 2011;4(1):40-56.
12. Crawley J. What's wrong with my mouse? Behavioral phenotyping of transgenic and knockout mice. New York: Wiley; 2007. 544 p.
13. Till SM, Asiminas A, Jackson AD, Katsanevaki D, Barnes SA, Osterweil EK, et al. Conserved hippocampal cellular pathophysiology but distinct behavioural deficits in a new rat model of FXS. *Hum Mol Genet.* 2015;24(21):5977-84.
14. Spencer C, Alekseyenko O, Serysheva E, Yuva-Paylor L, Paylor R. Altered anxiety-related and social behaviors in the Fmr1 knockout mouse model of fragile X syndrome. *Genes, Brain and Behavior.* 2005;4(7):420-30.
15. McNaughton CH, Moon J, Strawderman MS, Maclean KN, Evans J, Strupp BJ. Evidence for social anxiety and impaired social cognition in a mouse model of fragile X syndrome. *Behav Neurosci.* 2008;122(2):293-300.
16. Gantois I, Pop AS, de Esch CE, Buijsen RA, Pooters T, Gomez-Mancilla B, et al. Chronic administration of AFQ056/Mavoglurant restores social behaviour in Fmr1 knockout mice. *Behav Brain Res.* 2013;239:72-9.
17. Caspi A. Gene-environment interactions in psychiatry: joining forces with neuroscience. *Nature Reviews Neurosci.* 2006;7:583-90.

18. Rotschafer SE, Trujillo MS, Dansie LE, Ethell IM, Razak KA. Minocycline treatment reverses ultrasonic vocalization production deficit in a mouse model of Fragile X Syndrome. *Brain research*. 2012;1439:7-14.
19. Berdoy M, Smith P, Macdonald DW. Stability of social status in wild rats: age and the role of settled dominance. *Behaviour*. 1995;132(3):193-212.
20. Wang F, Zhu J, Zhu H, Zhang Q, Lin Z, Hu H. Bidirectional control of social hierarchy by synaptic efficacy in medial prefrontal cortex. *Science*. 2011;334(6056):693-7.
21. Lindzey G, Winston H, Manosevitz M. Social Dominance in Inbred Mouse Strains. *Nature*. 1961;191(4787):474-6.
22. Zhou T, Zhu H, Fan Z, Wang F, Chen Y, Liang H, et al. History of winning remodels thalamo-PFC circuit to reinforce social dominance. *Science*. 2017;357(6347):162-8.
23. Hamilton SM, Green JR, Veeraragavan S, Yuva L, McCoy A, Wu Y, et al. Fmr1 and Nlgn3 knockout rats: novel tools for investigating autism spectrum disorders. *Behavioral neuroscience*. 2014;128(2):103.
24. Asiminas A, Jackson A, Louros S, Till S, Dando O, Bear M, et al. Sustained correction of associative learning deficits following brief, early treatment in a rat model of Fragile X Syndrome. Submitted for publication. 2018.
25. Lore R, Flannelly K. Rat societies. *Sci Am*. 1977;236(5):106-11, 13-6.
26. Thomas KR, Capecchi MR. Site-directed mutagenesis by gene targeting in mouse embryo-derived stem cells. *Cell*. 1987;51(3):503-12.
27. Long SY. The Dalila effect: C57BL6 mice barber whiskers by plucking. *Behav Brain Res* 1972;108:39-45.
28. D.B. R, D.A. S. Evidence for marking with urine by rats. *Behavioral Biology*. 1974;12:517-23.
29. Wesson DW. Sniffing behavior communicates social hierarchy. *Current Biology*. 2013;23(7):575-80.
30. Jupp B, Murray J, Jordan E, Xia J, Fluharty M, Shrestha S, et al. Social dominance in rats: effects on cocaine self-administration, novelty reactivity and dopamine receptor binding and content in the striatum. *Psychopharmacology*. 2016;233:579-89.
31. Blanchard RJ, Flannelly KJ, Blanchard DC. Life-span studies of dominance and aggression in established colonies of laboratory rats. *Physiology & behavior*. 1988;43(1):1-7.
32. Craig J. Measuring social behavior: social dominance. *Journal of animal science*. 1986;62(4):1120-9.
33. Rodriguiz RM, Chu R, Caron MG, Wetsel WC. Aberrant responses in social interaction of dopamine transporter knockout mice. *Behavioural brain research*. 2004;148(1):185-98.
34. Goebel-Goody SM, Wilson-Wallis ED, Royston S, Tagliatela S, Naegele JR, Lombroso PJ. Genetic manipulation of STEP reverses behavioral abnormalities in a fragile X syndrome mouse model. *Genes, Brain and Behavior*. 2012;11(5):586-600.
35. de Esch CE, van den Berg WE, Buijsen RA, Jaafar IA, Nieuwenhuizen-Bakker IM, Gasparini F, et al. Fragile X mice have robust mGluR5-dependent alterations of social behaviour in the Automated Tube Test. *Neurobiol Dis*. 2014;75:31-9.
36. Iannaccone PM, Jacob HJ. *Rats! : The Company of Biologists Ltd*; 2009.
37. Dyer-Friedman J, Glaser B, Hessl D, Johnston C, Huffman LC, Taylor A, et al. Genetic and environmental influences on the cognitive outcomes of children with fragile X syndrome. *Journal of the American Academy of Child & Adolescent Psychiatry*. 2002;41(3):237-44.

Saxena et al - Figure 1

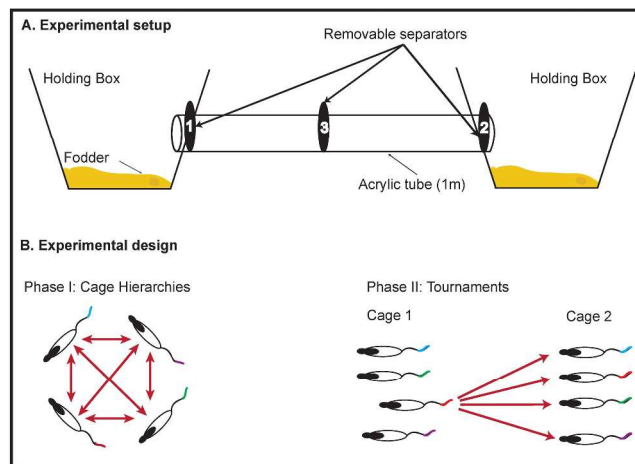


Figure 1

297x420mm (300 x 300 DPI)

Saxena et al - Figure 2

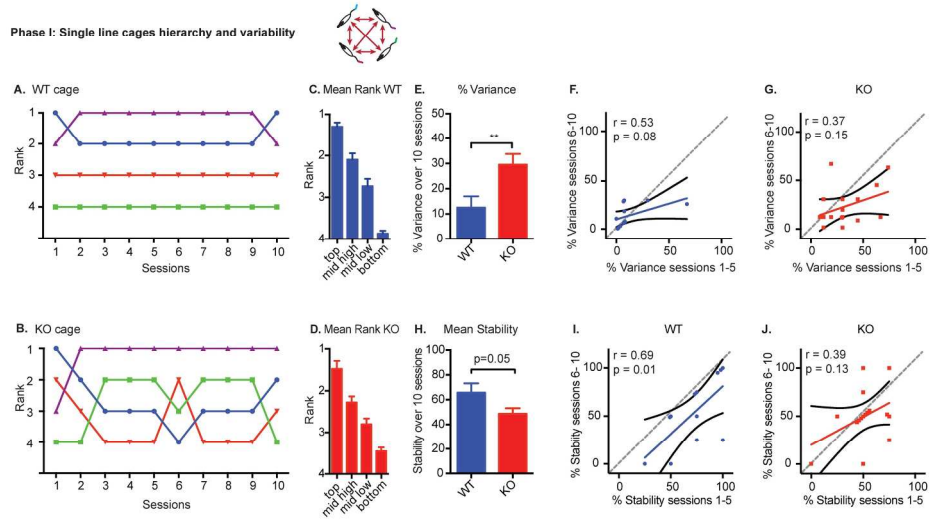


Figure 2

209x148mm (300 x 300 DPI)

Saxena et al - Figure 3

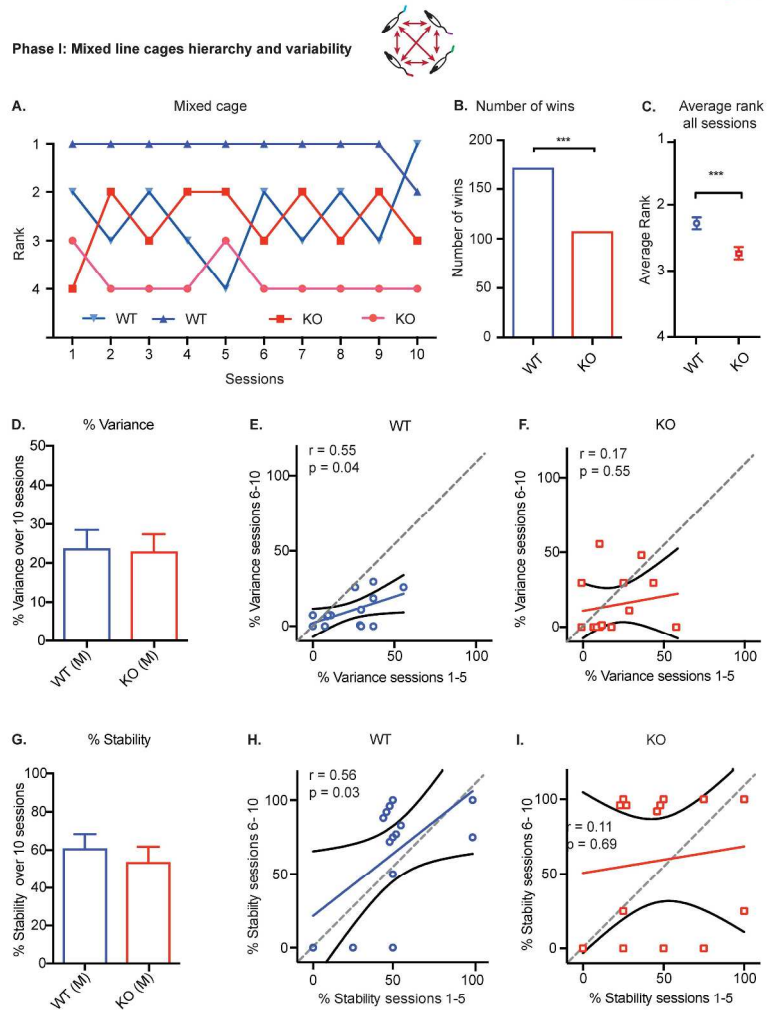


Figure 3

297x420mm (300 x 300 DPI)



Saxena et al - Figure 4

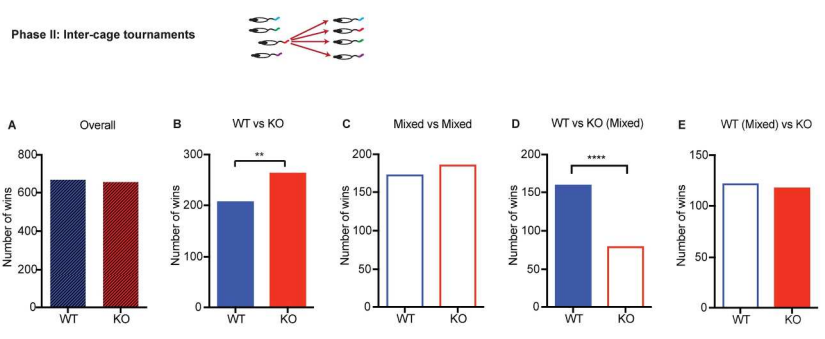


Figure 4

209x148mm (300 x 300 DPI)

Saxena et al - Figure 5

## Phase II: Inter-cage tournaments considering the rank from Phase I

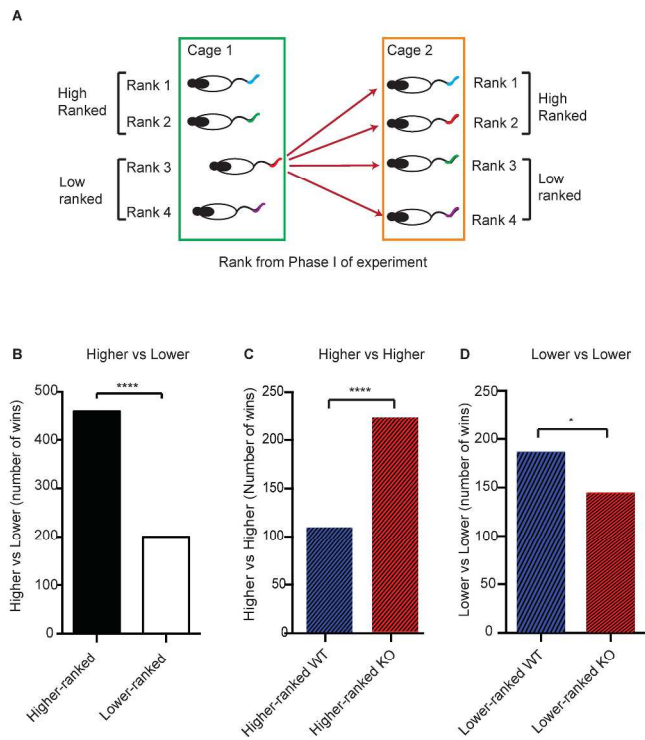


Figure 5

297x420mm (300 x 300 DPI)