Behavioral flexibility in a mouse model for obsessive-compulsive disorder: Impaired Pavlovian reversal learning in SAPAP3 mutants

Bastijn J.G. van den Boom1,2 | Adriana H. Mooij1 | Ieva Misevičiūtė1 | Damiaan Denys1,2 | Ingo Willuhn1,2

1Netherlands Institute for Neuroscience, Royal Netherlands Academy of Arts and Sciences, Amsterdam, The Netherlands
2Department of Psychiatry, Amsterdam UMC, University of Amsterdam, Amsterdam, The Netherlands

Correspondence
Ingo Willuhn, Netherlands Institute for Neuroscience, Meibergdreef 47, 1105 BA, Amsterdam, The Netherlands.
Email: i.willuhn@nin.knaw.nl

Funding Information
H2020 European Research Council, Grant/Award Number: ERC-2014-STG 638013; Nederlandse Organisatie voor Wetenschappelijk Onderzoek, Grant/Award Number: 864.14.010, 2015/06367/ALW

Obsessive-compulsive disorder (OCD) is characterized by obsessive thinking, compulsive behavior and anxiety, and is often accompanied by cognitive deficits. The neuropathology of OCD involves dysregulation of cortical-striatal circuits. Similar to OCD patients, SAPAP3 knockout mice (SAPAP3−/−) exhibit compulsive behavior (grooming), anxiety and dysregulated cortical-striatal function. However, it is unknown whether SAPAP3−/− display cognitive deficits and how these different behavioral traits relate to one another. SAPAP3−/− and wild-type (WT) littermates were trained in a Pavlovian conditioning task pairing visual cues with the delivery of sucrose solution. After mice learned to discriminate between a reward-predicting conditioned stimulus (CS+) and a non-reward stimulus (CS−), contingencies were reversed (CS+ became CS− and vice versa). Additionally, we assessed grooming, anxiety and general activity. SAPAP3−/− acquired Pavlovian approach behavior similarly to WT, albeit less vigorously and with a different strategy. However, unlike WT, SAPAP3−/− were unable to adapt their behavior after contingency reversal, exemplified by a lack of re-establishing CS+ approach behavior (sign tracking). Surprisingly, such behavioral inflexibility, decreased vigor, compulsive grooming and anxiety were unrelated. This study shows that SAPAP3−/− are capable of Pavlovian learning, but lack flexibility to adapt associated conditioned approach behavior. Thus, SAPAP3−/− not only display compulsive-like behavior and anxiety, but also cognitive deficits, confirming and extending the validity of SAPAP3−/− as a suitable model for the study of OCD. The observation that compulsive-like behavior, anxiety and behavioral inflexibility were unrelated suggests a non-causal relationship between these traits and may be of clinical relevance for the treatment of OCD.

KEYWORDS
anxiety, behavioral flexibility, cognitive flexibility, compulsive behavior, general activity, obsessive-compulsive disorder, Pavlovian conditioning, reversal learning, SAPAP3 mutant mice, symptoms relationship

INTRODUCTION

Obsessive-compulsive disorder (OCD) is a psychiatric disorder that is characterized by recurrent unwanted thoughts, anxiety and compulsive behavior, but is also often associated with cognitive deficits.1–5 The persistence of maladaptive patterns of inflexible thoughts and behaviors suggest a lack of cognitive flexibility, the ability to adapt behavior in response to changing situational requirements.

Preclinical animal models are a valuable tool to elucidate neurobiological mechanisms of OCD, but also enable us to investigate how...
different symptoms relate to one another. Mice with genetic deletion of Synapse-associated protein 90/postsynaptic density protein 95 associated protein 3 (SAPAP3−/−), a postsynaptic scaffolding protein predominantly expressed in cortico-striatal circuits,6–8 have been used for the study of OCD. Although certain genetic variants of the human homolog of SAPAP3 occur more frequently in OCD patients9 and variation in the SAPAP3 gene was found to be associated with grooming-related disorders in humans (without direct association to OCD),10 the genetic link of SAPAP3 to OCD needs further study. However, the previously showed virtue of the SAPAP3−/− model lies outside of genetics: The phenotype of these mice maps remarkably well onto symptoms of human OCD patients. For example, both OCD patients11–14 and SAPAP3−/−7,8 exhibit dysregulation of projections from cortex to striatum. Similar to subtypes of OCD patients, SAPAP3−/− display compulsive-like grooming that can be decreased by deep-brain stimulation.15 Furthermore, optogenetic stimulation of cortico-striatal projections can restore normal grooming,16 whereas stimulation of cortico-striatal projections in wild-type (WT) mice evokes increased grooming.17 In addition to excessive grooming, SAPAP3−/− mice show increased anxiety, both of which can be reduced by viral rescue of striatal SAPAP3.7 Similarly, administration of selective serotonin reuptake inhibitors, the primary pharmacotherapy for OCD, normalizes self-grooming and anxiety-like behavior in SAPAP3−/−. Despite this promising validation of the model, cognitive deficits have not been assessed in SAPAP3−/− until now (this manuscript and Ref. 18). To study cognitive flexibility in both humans and animals, reversal learning paradigms are often used.19,20 Previous studies examining behavioral deficits in reversal learning in OCD patients yielded mixed outcomes, with some studies observing deficits,21–27 whereas others did not.28–30 Notably, deficits in reversal learning associated with altered recruitment of fronto-striatal circuitry have been observed more consistently.31–35 During reversal learning, previously acquired contingencies of stimulus-reward associations are reversed, and the subjects’ adaptation to this is assessed. Pavlovian conditioning is the most basic type of associative learning, during which a conditioned stimulus (CS) can trigger approach behavior, a procedure called “autoshaping”.36–38 Autoshaping enables differentiation between approach towards the predictive CS itself (so-called sign tracking), thought to be driven by model-free strategies, and the reward location (goal tracking), presumably driven by model-based strategies.39–41 thereby probing cognitive mechanisms underlying the behavior. To investigate cognitive flexibility in SAPAP3−/−, we trained mice in an autoshaping paradigm in touchscreen boxes. Upon task acquisition, reward contingencies were reversed. In addition, we investigated the relationship between behavioral flexibility, compulsive-like behavior, and anxiety. Such a multi-faceted behavioral investigation of SAPAP3−/− may contribute to the understanding of behavioral deficits in OCD patients.

2 | MATERIALS AND METHODS

2.1 | Subjects

Animal procedures were in accordance with European and Dutch laws and approved by the Animal Experimentation Committee of the Royal Netherlands Academy of Arts and Sciences. SAPAP3−/− mice were donated by the Feng laboratory.7 Male and female SAPAP3−/− (mean age 9 months, n = 20, two excluded because of skin lesions) and WT littermates (mean age 8 months, n = 14) were housed individually on a reversed light-dark cycle (lights on from 19:00 to 07:00) and food-restricted to 85% of their free-feeding body weight. All behavioral procedures were performed during the dark phase.

2.2 | Procedure

First, grooming was assessed in an open field (OF; preautoshaping) for 60 minutes. Next, animals were tested in the autoshaping task, followed by a second 60-minute OF test (postautoshaping), and 10 minutes on the elevated plus maze (EPM) to probe anxiety. A Janelia Automatic Animal Behavior Annotator classifier42 was used to quantify self-grooming (see Ref. 43). More methodological details can be found in the supplemental information.
2.3 | Pavlovian conditioning (autoshaping)

2.3.1 | Apparatus
Training was performed in trapezoid-shaped Bussey-Saksida touchscreen chambers (Campden Instruments, Leics, UK)44 (Figure 1A). CS (white rectangles presented for 10 seconds) appeared in two different positions on the touchscreen (12.1 in., screen resolution 600 x 800), but were otherwise indistinguishable. Chambers were equipped with infrared beam detectors—one for trial initiation (opposite to touchscreen), two measuring stimuli approaches and one to count reward magazine entries. The experiment was based upon the autoshaping protocol presented in Hore et al.44

2.3.2 | Training
Mice were trained twice a day in the autoshaping for 30 minutes per session, for a total of 72 sessions (36 before and 36 after reversal). A trial started when the mouse interrupted the infrared beam at the back of the chamber (invisible to the mouse) after a variable intertrial interval of 25 seconds (low contiguity between interrupting beam and trial initiation). Therefore, trial initiation is a consequence of exploration and general activity. Immediately after trial initiation, an auditory cue was presented (0.5 seconds) and a visual stimulus displayed on either the left (rewarded CS; CS+) or right (nonrewarded CS; CS−) side of the screen (position counterbalanced between animals) for 10 seconds. Upon CS+ offset, reward was delivered to the magazine (7 µL, liquid strawberry milkshake, Melkunie) (Figure 1B). During CS+ trials, we expected mice to approach the CS+ location or the reward magazine (both measured with infrared beams) (Figure 1A). During CS− trials, we expected animals to refrain from making such approaches.

2.3.3 | Reversal training
Reward contingencies were reversed after 36 sessions (spatial reversal of CS), whereby the previous CS+ became the CS− and the previous CS− became the CS+.

2.3.4 | Exclusion criteria
We excluded animals that did not associate the CS+ with reward.45−47 Thus, animals that failed to approach screen or reward magazine during CS+ in over 70% of trials (in the last 10 sessions before reversal) and/or failed to avoid screen or reward magazine during CS− less than 70% of trials in the same sessions, were excluded from the analysis. This resulted in three excluded WT and six excluded SAPAP3−/−. Additionally, on a session-by-session basis, individual sessions in which animals initiated less than 10 trials were excluded.

2.3.5 | Performance measures
We measured the following variables based on infrared beam interruptions (Figure 1A): (a) number of trials initiated, (b) number of trials with an approach towards screen (sign tracking), (c) number of trials with an approach towards reward magazine (goal tracking), (d) combined number of trials with an approach towards screen and/or reward magazine [CS screen approach OR CS reward magazine approach] (in some cases both elements are approached in the same trial, which is then accounted for as a single trial with approach behavior in this calculation), (e) difference (score) of combined number of trials [combined CS+ approaches-combined CS− approaches], and (f) general activity during autoshaping measured as total number of infrared beam breaks outside of CS presentation. Combined number of trials with an approach (during CS presentation) consisted of the animals’ approach to the cue, the reward magazine, or both.36

2.4 | Data analysis
OF grooming data were analyzed using two-way mixed ANOVA with within-factor time between OF testing and between-factor genotype, followed by post-hoc analyses using independent t tests (except Mann-Whitney U test to compare grooming duration after autoshaping). EPM data were analyzed using independent t tests (except Mann-Whitney U test to compare entries to open arms). OF activity data were analyzed using Kruskal Wallis test (within-factor time between OF testing and between-factor genotype), followed by Wilcoxon signed-rank post-hoc test. For trial-initiation analysis, two-way repeated ANOVA was used (within-factors session and reversal). Comparison within genotype employed two-way repeated ANOVA (between-factor CS and reversal), followed by post-hoc paired t tests (except Wilcoxon signed-ranks tests to compare CS+ vs CS− prerereversal for both genotypes). Direct comparison of genotypes using difference score was analyzed using two-way mixed ANOVA (within-factor acquisition or maintenance and between-factor genotype), followed by post-hoc independent t tests. Correlation coefficients are expressed as R squared and estimates of 95% confidence intervals are reported. Statistical analyses were carried out using SPSS (version 23.0) and Graphpad Prism (version 6). P values were adjusted for multiple comparisons using the Holm-Bonferroni correction.48 Statistical significance was defined as P < 0.05.

3 | RESULTS

3.1 | SAPAP3−/− mice phenotyping
Grooming was assessed in the OF before (preautoshaping) and after (postautoshaping) Pavlovian conditioning. A main effect of genotype on grooming duration (F(1, 29)=8.69, P = 0.006), but no main effect of session (F(1, 29)=0.50, NS), nor an interaction effect (F(1, 29)=0.26, NS) was found. SAPAP3−/− showed increased grooming duration both preautoshaping (Figure 2A, mean ± SEM: 159.43 ± 19.67 seconds WT vs 320.53 ± 58.67 seconds SAPAP3−/−, t(19.50) = −2.60, P = 0.017) and postautoshaping (Figure 2A, 114 ± 15.71 seconds WT vs 312.29 ± 66.62 seconds SAPAP3−/−, U = 57, z = −2.46, P = 0.026). Grooming bouts showed a similar effect (supplementary Figure 1A).

A significant correlation was found between preautoshaping and postautoshaping grooming for SAPAP3−/− (Pearson R2 of 0.243; P = 0.04; 95% CI 0.02-0.79), indicating relatively consistent grooming over time (3 months) (Figure 2D). These grooming durations were averaged to compute a grooming trait value per animal.

On the EPM, SAPAP3−/− spent less time in open arms (Figure 2B: 152.25 ± 19.52 seconds WT vs 97.38 ± 8.51 seconds SAPAP3−/−).
3.2 | General activity during behavioral tasks

Throughout different behavioral tasks, we assessed general activity. Visual inspection of trajectories of SAPAP3−/− in the OF showed similar movement patterns compared to WT (Figure 3A, B). During OF testing, movement was quantified during periods when animals did not groom (to exclude effects of grooming on activity). A main effect of genotype on movement was found (F(1, 29) = 14.47, P = 0.001), but no main effect of time between OF sessions (F(1, 29) = 0.62, NS), nor an interaction effect (F(1, 29) = 0.12, NS). Post-hoc analyses showed that SAPAP3−/− showed decreased movement during preautoshaping OF (Figure 3C: 21081.5 ± 4001 cm WT vs 10 644.6 ± 769.1 cm SAPAP3−/−, U = 38, z = −3.22, P = 0.001) and postautoshaping OF (Figure 3C: 21772.6 ± 2083.3 cm WT vs 12 445.2 ± 961 cm SAPAP3−/−, U = 18, z = −4.01, P < 0.0001). In addition, analyzing movement in the inner (data not shown, t(29)=2.15, P = 0.04) and outer diameter (data not shown, t(29)=2.24, P = 0.03) of the OF also showed that SAPAP3−/− showed decreased movement. To test if decreased general activity in SAPAP3−/− was because of excessive grooming, movement was divided by time not spent grooming. Similar to movement, a main effect of genotype on movement per minute was found (F(1, 29)=12.32, P < 0.0001), but no main effect of time in between OF testing (F(1, 29)=0.51, NS), nor an interaction effect (F(1, 29)=0.16, NS). Post-hoc tests showed that SAPAP3−/− showed decreased movement per minute preautoshaping (Figure 3D: 368.45 ± 71.42 cm/min WT vs 194.29 ± 13.1 cm/min SAPAP3−/−, U = 46, z = −2.9, P = 0.003) and postautoshaping (Figure 3D: 377.05 ± 37.63 cm/min WT vs 224.51 ± 15.13 cm/min SAPAP3−/−, U = 19, z = −3.97, P < 0.0001). Movement during OF was not related to grooming (Pearson R² of 0.03; NS; 95% CI −0.60 to 0.34) (Figure 3E).

During EPM testing, SAPAP3−/− exhibited decreased movement (Figure 3F: 4008.17 ± 156.34 cm WT vs 2292.15 ± 259.57 cm SAPAP3−/−, t(29)=5.36, P < 0.0001) that was not because of increased grooming (grooming duration EPM vs movement EPM; Pearson R² of 0.03; NS; 95% CI −0.61 to 0.34), nor because of increased anxiety-like behavior (ratio of time spent in closed vs open arms vs movement EPM; Pearson R² of 0.22; NS; 95% CI −0.78 to 0.01).

During autoshaping, average total beam breaks during intertrial intervals (as a proxy for activity) were averaged across sessions. SAPAP3−/− showed decreased activity compared to WT (Figure 3G:...
FIGURE 3  SAPAP3−/− display decreased general activity during behavioral testing that did not affect the overall number of initiated trials. Representative open field (OF) trajectories of a wild-type littermate (WT) (A) and a SAPAP3−/− (B) during OF testing. Activity in the OF measured during periods when animals were not grooming for WT (n = 14, blue, circles are individual animals) and SAPAP3−/− (n = 17, orange, triangles are individual animals) (C). Average activity in the OF per minute of time not spent grooming (D). No correlation between averaged grooming duration and average activity in the OF per minute (E). General activity on the elevated plus maze (F). Beam breaks measured during autoshaping intertrial intervals (no CS) as a proxy for activity for WT (n = 11, circles) and SAPAP3−/− (n = 12, triangles) (G). Percentage of WT autoshaping sessions excluded based on criterion of a minimum of 10 initiated trials per session (H). Percentage of SAPAP3−/− autoshaping sessions excluded (I). The average number of excluded WT and SAPAP3−/− sessions is significantly different (J). Average number of initiated trials per session for WT (excluded sessions removed) (K). Average number of initiated trials per session for SAPAP3−/− (excluded sessions removed) (L). Average number of trials initiated by WT and SAPAP3−/− before and after reversal is not significantly different (M). Data are mean ± SEM; *P < 0.05. EPM, elevated plus maze; NS, not significant; OF, open field

438.35 ± 57.82 breaks WT vs 206.44 ± 28.13 breaks SAPAP3−/−, t(23)=3.71, P = 0.001.

Mice had to initiate trials by interrupting an infrared beam located opposite to the screen. Some animals initiated only approximately five trials per session, whereas the majority of the animals initiated about 30. Sessions with less than 10 initiated trials were excluded from analyses (Figure 3H I J). Analyses on the number of excluded sessions showed a significant main effect of genotype (F(1, 23)=4.26, P = 0.05) and a main effect of reversal (F(1, 23)=7.94, P = 0.01), but no interaction (F(1, 23)=0.19, NS), suggesting that although reversal had an effect on trial initiation in both genotypes, SAPAP3−/− were more inactive in general.

After exclusion of inactive sessions, the number of initiated trials for WT showed no significant effect of reversal (F(1, 569.59) = 2.14, NS), nor an effect of session (F(35,30.3) = 0.73, NS) nor an interaction effect (F(35,30.3) = 1.01, NS) (Figure 3K). SAPAP3−/− displayed a minor decrease in initiated trials after reversal (F(1,481.69) = 28.31, P < 0.001), but no effect of session (F(35,27.49) = 0.45, NS) nor an interaction effect (F(35, 27.49) = 1.01, NS) (Figure 3L). Direct comparison of initiated trials between WT and SAPAP3−/− showed no difference (Figure 3M: 33.82 ± 2.39 trials WT vs 27.83 ± 2.01 trials SAPAP3−/−, t(23)=1.93, NS).

3.3 | Autos shaping performance

During CS+ presentation before reversal, WT interacted with the reward magazine (Figure 4A) as well as the CS itself (Figure 4C) with no systematic preference. After reversal, WT re-acquired the new reward contingencies, showed by increased CS approaches, but refrained from magazine approaches.

Similar to WT, SAPAP3−/− learned to discriminate between CS+ and CS−, but mainly only approached the reward magazine (Figure 4B) and not the CS (Figure 4D). After reversal, SAPAP3−/− showed diminished discrimination between the CS+ and CS−, but still retrieved rewards.

Because mice interacted with both screen and magazine, we calculated combined approaches towards screen and magazine during either CS+ or CS− presentation. This allowed direct comparison of autoshaping performance within and between genotypes, independent of applied behavioral strategy (Figure 4E, F).

Statistics were performed on the 10 sessions before reversal and on the last 10 sessions after reversal averaged over animals, as performance became asymptotic. In WT, a main effect of CS on autoshaping performance (F(1, 10) = 142.66, P < 0.0001), no main effect of reversal (F(1, 10) = 0.21, NS), nor an interaction effect (F(1, 10) = 0.11, NS) was found. Post-hoc analyses showed a significant difference between approach behavior towards CS+ and CS− before (Figure 4E, bar graphs prereversal: 16.64 ± 1.46 trials with approaches CS+ vs 8.07 ± 0.99 trials with approaches CS−, P = 0.009) and after reversal (postreversal: 15.02 ± 1.31 trials with approaches CS+ vs 8.71 ± 0.99 trials with approaches CS−, t(10) = 9.1, P < 0.001), accompanied by no difference between CS+ before and after reversal (t(10) = 1.29, NS) nor between CS− before and after reversal
suggesting that mice reached similar performance after reversal.

In SAPAP3−/−, we found a significant main effect of CS on autoshaping performance ($F_{(1, 10)} = 33.96, P < 0.0001$), no main effect of reversal ($F_{(1, 10)} = 1.66, NS$), but accompanied by a significant interaction effect ($F_{(1, 10)} = 9.81, P = 0.01$), suggesting that SAPAP3−/− approached CS+ and CS− differently before and after reversal. Indeed, post-hoc analyses showed that SAPAP3−/− differentiated between CS+ and CS− before reversal (Figure 4F bar graphs: 13.99 ± 1.1 trials with approaches CS+ vs 8 ± 0.88 trials with approaches CS−, $t_{(10)} = 7.96, P < 0.0001$), but not after reversal (10.69 ± 1.4 trials with approaches CS+ vs 8.11 ± 1.11 trials with approaches CS−, $t_{(10)} = 2.53, P = NS$). A significant decrease in CS+ approach behavior was found after reversal ($t_{(10)} = 2.91, P = 0.045$), but not in CS− approach behavior ($t_{(10)} = −0.21, NS$).

### 3.4 Direct performance comparison between genotypes

We computed a difference score of combined approach behavior for both genotypes (Figure 5A) and performed statistics on the first 10 sessions before and after reversal (preacquisition and
postacquisition, respectively) and the last 10 sessions before and after reversal (premaintenance and postmaintenance, respectively). A main effect of reversal on acquisition \((F(1, 21) = 96.37, P < 0.0001)\) was found, but no main effect of genotype \((F(1, 21) = 0.04, NS)\), nor an interaction effect \((F(1, 21) = 0.001, NS)\), suggesting similar acquisition rate between WT and SAPAP3\(^{−/−}\).

A significant main effect of reversal on maintenance was found \((F(1, 20) = 11.38, P = 0.003)\), accompanied by a significant main effect of genotype \((F(1, 20) = 10.47, P = 0.004)\), without an interaction effect \((F(1, 20) = 0.51, NS)\). Interestingly, post-hoc analyses showed no significant difference between WT and SAPAP3\(^{−/−}\) before reversal (Figure 5A bar graphs premaintenance: 8.58 ± 1.06 difference score WT vs 5.98 ± 0.75 difference score SAPAP3\(^{−/−}\), \(t(21) = 2.02, NS\)); however, a significant difference after reversal (Figure 5A bar graphs postmaintenance: 6.31 ± 0.69 difference score WT vs 2.58 ± 1.02 difference score SAPAP3\(^{−/−}\), \(t(20) = 3.03, P = 0.028\)) was found.

To examine the relation between grooming and autoshaping performance in SAPAP3\(^{−/−}\), correlation analyses between the grooming trait measure and the difference score were performed. No correlation between grooming and premaintenance difference score was found (Pearson \(R^2\) of 0.004; NS; 95% CI −0.53 to 0.62), nor between grooming and postmaintenance difference score (Pearson \(R^2\) of 0.05; NS; 95% CI −0.44 to 0.73) (Figure 5B).

We explored the relation between anxiety and autoshaping performance in SAPAP3\(^{−/−}\). No correlation was found between anxiety and premaintenance difference score (Pearson \(R^2\) of 0.08; NS; 95% CI −0.34 to 0.74), or between anxiety and postmaintenance difference score Pearson \(R^2\) of 0.04; NS; 95% CI −0.72 to 0.45) (Figure 5C).

Finally, successful reversal in WT was accompanied by re-emerging of cue approach behavior (Figure 4C). Thus, we tested the correlation between cue approach behavior during CS+ and difference score after reversal and found a strong relationship between these two measures (Pearson \(R^2\) of 0.75; \(P < 0.0001\); 95% CI 0.69-0.94) (Figure 5D), suggesting that CS+ approach behavior is involved in successful reversal learning. No correlation was found between reward magazine approach behavior during CS+ and difference score after reversal (Pearson \(R^2\) of 0.02; NS; 95% CI −0.53 to 0.30) (Figure 5E).

4 | DISCUSSION

The first aim of this study was to investigate the ability of SAPAP3\(^{−/−}\), a transgenic mouse model for compulsive behavior in psychiatric disorders such as OCD, to acquire Pavlovian conditioned responding and their ability to flexibly adjust acquired behavior to reversed reward contingencies. We found that SAPAP3\(^{−/−}\), although less vigorous in their responses compared to WT, acquired responses to Pavlovian CS,
but unlike WT, were unable to adapt their conditioned approach behavior upon contingency reversal. Both genotypes developed Pavlovian “goal-tracking” approaches during the CS+ (reward-magazine approaches), but ceased to goal-track after reversal. In contrast to SAPAP3−/−, WT exhibited “sign tracking” (CS+ approaches), which emerged during task acquisition and re-emerged towards the new CS+ after reversal, suggesting that this behavioral strategy contributed to successful reversal learning. Our second aim was to assess how behavioral flexibility is related to other OCD-like symptoms such as compulsive behavior and anxiety. Surprisingly, both grooming and behavioral flexibility is related to other OCD-like symptoms such as compulsive behavior and anxiety. Surprisingly, both grooming and behavioral flexibility were unrelated to Pavlovian behavioral flexibility in SAPAP3−/−, suggesting that these traits have independent etiologies. Together, our results refine the SAPAP3−/− mouse model for OCD by identifying another OCD-like trait and its relationship to other cardinal OCD-like symptoms.

SAPAP3−/− have been shown to groom excessively to the point of removing fur and occasionally producing skin lesions. Because of these negative consequences, this behavior is considered compulsive. Consistently, we confirm that SAPAP3−/− display increased grooming compared to WT, reflected in both number of grooming bouts and duration of grooming. Increased grooming was detected both before and after the Pavlovian conditioning and on the EPM, suggestive of a stable phenotype that is not affected by behavioral testing. Furthermore, grooming before and after autoshaping was correlated significantly, indicating that individual mice display a relatively reliable degree of grooming, even over a period of months. Our results are consistent with previous reports, demonstrating robustness of the SAPAP3−/− grooming phenotype and further validate this behavioral readout as a proxy for compulsivity.

In addition to grooming, we measured other behavioral traits that are central to OCD symptomology. We assessed anxiety on the EPM and confirmed previously reported augmentation of anxiety in SAPAP3−/−. Previous studies measured anxiety in the OF, in the light-dark box, and on the elevated zero-maze. The light-dark box test and the elevated mazes are widely used assays for anxiety-like behavior, thought to assess different forms of anxiety, bright-space and open-space anxiety, respectively. Thus, SAPAP3−/− show increased anxiety-like behavior on different paradigms, indicating a broad anxiety phenotype.

We then asked whether increased grooming and anxiety in SAPAP3−/− were related, but found no correlation between these two variables. More specifically, the degree of anxiety measured on the EPM did not correlate with grooming on the EPM itself (grooming state during paradigm), nor with grooming repeatedly assessed in the OF (grooming trait over time). Therefore, our findings imply that compulsive behavior and anxiety are not causally related to one another in SAPAP3−/−, a question of clinical relevance, where some hypothesized that anxiety causes compulsion in OCD, and others hypothesized that compulsivity causes anxiety.

Unexpectedly, we discovered another SAPAP3−/− trait that is not commonly reported as an OCD symptom: General activity (ie, locomotion plus overall movement) was diminished compared to WT. This decreased activity was not an indirect consequence of SAPAP3−/− spending more time grooming instead of being active otherwise, because decreased activity remained, even after grooming periods were excluded from the analysis (ie, activity relative to time spent not grooming). In addition, this relative inactivity was not correlated with grooming itself. However, although no correlation was found between movement and anxiety-like behavior assessed on the EPM, we cannot exclude that anxiety may play a role in decreased overall activity. To ensure that this differential activity did not confound Pavlovian learning, we took several measures: (a) Animals were required to initiate trials in the autoshaping task, which enabled the exclusion of low-activity sessions and caused the total number of initiated trials not to differ between genotypes. (b) Rather than analyzing the total number of approaches during CS presentation, we analyzed the number of trials in which mice completed at least one response during the CS. (c) In order to not bias towards exclusive approaches to either cue or magazine, a measure of “combined approach” responding during CS presentation was computed (ie, counting whether an animal approached either CS or reward magazine during CS presentation). Together, these methods precluded general activity differences between genotypes from penetrating learning variables (instead of assessing how vigorous a mouse responded) and enabled direct comparison of SAPAP3−/− and WT.

Although we focused on minimizing the potentially confounding effects of decreased SAPAP3−/− general activity on reversal learning, it cannot be excluded as a trait of potential OCD relevance. For instance, OCD shows high comorbidity with depression and anhedonia, pathologies that produce decreased activity marked by loss of motivation and inability to experience pleasure. Furthermore, patients with severe OCD tend to exhibit depressive symptoms, elaborate avoidance behavior, and high levels of anhedonia, all of which are consistent with decreased general activity. Finally, it has been reported that OCD patients move around less in their homes during everyday life compared to healthy controls. However, whether diminished general activity is an undereexplored symptom of OCD that could potentially be studied in SAPAP3−/− will have to be evaluated in future studies.

We show that SAPAP3−/− were able to learn to discriminate between environmental stimuli predicting reward (CS+) and no reward (CS−) similar to WT and displayed Pavlovian conditioned approach responses during presentation of these stimuli, indicating no overall Pavlovian learning deficit. However, already during initial acquisition (prior to reversal), SAPAP3−/− employed a different approach strategy than WT. In anticipation of reward, WT approached both the CS location and the reward magazine equally during CS+ presentation, whereas SAPAP3−/− only approached the magazine. These two approach strategies are thought to differ in the amount of incentive salience assigned to the CS. Approach towards the CS+ itself (sign tracking) is thought to be rooted in the CS gaining incentive salience, a process consistent with model-free learning. In contrast, approach towards the reward location (goal tracking) suggests underlying model-based learning independent of incentive motivation. Surprisingly, after reversal, both genotypes refrained from reward magazine approaches. SAPAP3−/− did not recover responding, whereas WT re-acquired approach behavior under the reversed reward contingencies, although exclusively towards the CS+, suggesting model-free mechanisms to enable this flexible behavior. To take this speculation one step further: The lack of model-free learning-
based approaches in SAPAP3−/− may explain their inability to adapt to the reversal. However, future studies are necessary to test these ideas in more depth.

Previous studies indicate crucial involvement of the prefrontal cortex (PFC) in reversal learning. One PFC region, the orbitofrontal cortex (OFC), is thought to be particularly important, as OFC lesions consistently result in impaired reversal learning.60–67 SAPAP3−/− display altered OFC-striatal activity,7,8 and deficits in behavioral response inhibition that can be rescued by optogenetic stimulation of the OFC-striatal network.64 We report that once SAPAP3−/− learned CS contingencies, they were unable to update their behavioral response upon reversal. One explanation for this finding is that SAPAP3−/− were not able to “disinhibit” responding for previously unrewarded cues, despite successful inhibition of responding to the previously rewarded cue. This is consistent with the reported intact acquisition of Pavlovian responses, but impaired reversal learning in OFC-lesioned animals.65–67 Thus, a compromised PFC-striatal network present in SAPAP3−/−, which was shown to be involved in their excessive grooming,68,15,16 is possibly responsible for the lack of adaptation to changing situational requirements. Furthermore, striatal regions that receive PFC input are thought to be critical for model-free learning,68 suggesting that both the lack of model-free response strategies in SAPAP3−/− and their behavioral inflexibility may be a consequence of SAPAP3−/−-inherent PFC-striatal dysfunction.

The persistent, compulsive behavior of OCD patients can be conceptualized as inflexible behavior. However, previous studies examining symptom-unrelated cognitive flexibility in OCD patients yielded mixed outcomes, with some studies observing behavioral deficits in reversal learning,21–27 whereas others did not.28–30,69 As discussed above, deficits in reversal learning are associated with altered recruitment of fronto-striatal circuitry (suggestive of altered cognitive processing), which has been observed more consistently in OCD patients during cognitively-flexibility demanding tasks.31–34 Moreover, a recent neuroimaging study employing Pavlovian fear conditioning found that OCD patients failed to flexibly update fear responses after reversal, despite normal acquisition of fear conditioning.70 Similarly, we found that SAPAP3−/− acquire Pavlovian conditioning but fail to flexibly update their responses after contingency reversal. Thus, dysfunctional cortical-striatal circuitry in both OCD patients and SAPAP3−/− may be responsible for behavioral deficits in flexibly updating conditioned responses, further validating the SAPAP3−/− model for OCD.

SAPAP3−/− acquired Pavlovian conditioned responding, similar to WT, but failed to flexibly update their behavior upon reversal of reward contingencies. This lack of behavioral adaptation was robust and persisted for an extended period of training after contingency reversal. Such inflexibility could potentially contribute to the persistence of compulsive behavior despite negative consequences. However, individual traits of SAPAP3−/− measured here (anxiety, compulsivity, flexibility, vigor) were not linearly related to one another, suggesting at least partial independence, which may prove to be of relevance for the treatment of OCD patients. In summary, we report that in addition to compulsive behavior and augmented anxiety, SAPAP3−/− display decreased vigor and cognitive deficits, thereby mapping well onto OCD symptomatology. Thus, our work provides further support for the use of the SAPAP3−/− model to study OCD-like behavior and its underlying neurobiology.

ACKNOWLEDGMENTS
We thank Dr. Matthijs Feenstra (Netherlands Institute for Neuroscience) and Dr. Nienke Vulink (Amsterdam UMC) for their insightful comments on the manuscript, Dr. Nicole Yee (Netherlands Institute for Neuroscience) for her technical assistance and input on the manuscript, Ralph Hamelink (Netherlands Institute for Neuroscience) for his practical contributions to this study, and Dr. Guoping Feng (Massachusetts Institute of Technology) for providing us with SAPAP3−/−.

This work was in part supported by the NWO VIDI grant to I.W. (864.14.010, 2015/06367/ALW) and by the ERC Starting Grant to I.W. (ERC-2014-STG 638013). This research did not receive any funding from agencies in the for-profit sector.

CONFLICTS OF INTEREST
None of the authors have conflicts of interest associated with this study.

ORCID
Bastijn J.G. van den Boom https://orcid.org/0000-0002-0853-3763
Ingo Willuhn https://orcid.org/0000-0001-6540-6894

REFERENCES
1. Abramowitz JS, Taylor S, McKay D. Obsessive-compulsive disorder. Lancet. 2009;374:491-499.
2. Denys D. Pharmacotherapy of obsessive-compulsive disorder and obsessive-compulsive spectrum disorders. Psychiatr Clin North Am. 2006;29:553-584.
3. Bakor G, Anderson PD. Obsessive-compulsive disorder. J Pharm Pract. 2014;27:116-130.
4. Gruner P, Pittenger C. Cognitive inflexibility in obsessive-compulsive disorder. Neuroscience. 2017;345:243-255.
5. Benzina N, Mallet L, Burguière E, N'Diaye K, Pelissolo A. Cognitive dysfunction in obsessive-compulsive disorder. Curr Psychiatry Rep. 2016;18:80. Retrieved April 5, 2018, from https://c9nlhk4bmfrp5ifldfkis07rb.sec.amc.nl/content/pdf/10.1007/s11920-016-0720-3.pdf.
6. Welch JM, Wang D, Feng G. Differential mRNA expression and protein localization of the SAP90/PSD-95-associated proteins (SAPAPs) in the nervous system of the mouse. J Comp Neurol. 2004;472:24-39.
7. Welch JM, Lu J, Rodriguez RM, et al. Cortico-striatal synaptic defects and OCD-like behaviours in Sapap3-mutant mice. Nature. 2007;448:894-900.
8. Han Y, Ade K, Caffall Z, et al. Circuit-selective striatal synaptic dysfunction in the sapap3 knockout mouse model of obsessive-compulsive disorder. Biol Psychiatry. 2014;75:623-630.
9. Züchner S, Wendland JR, Ashley-Koch AE, et al. Multiple rare SAPAP3 missense variants in trichotillomania and OCD. Mol Psychiatry. 2009;14:6-9.
10. Bienvenu OJ, Wang Y, Shugart YY, et al. Sapap3 and pathological grooming in humans: results from the OCD collaborative genetics study. Am J Med Genet Part B Neuropsychiatr Genet. 2009;150:710-720.
11. Ting JT, Feng G. Neurobiology of obsessive-compulsive disorder: insights into neural circuitry dysfunction through mouse genetics. Curr Opin Neurobiol. 2011;21:842-848.
12. Milad MR, Rauch SL. Obsessive-compulsive disorder: beyond segregated cortico-striatal pathways. Trends Cogn Sci. 2012;16:43-51.
59. Dayan P, Berridge KC. Model-based and model-free Pavlovian reward learning: revaluation, revision, and revelation. Cogn Affect Behav Neurosci. 2014;14:473-492.

60. Dias R, Robbins TW, Roberts AC. Dissociation in prefrontal cortex of affective and attentional shifts. Nature. 1996;380:69-72.

61. McAlonan K, Brown VJ. Orbital prefrontal cortex mediates reversal learning and not attentional set shifting in the rat. Behav Brain Res. 2003;146:97-103.

62. Boulougouris V, Dalley JW, Robbins TW. Effects of orbitofrontal, infralimbic and prelimbic cortical lesions on serial spatial reversal learning in the rat. Behav Brain Res. 2007;179:219-228.

63. Chudasama Y, Robbins TW. Dissociable contributions of the orbitofrontal and infralimbic cortex to pavlovian autoshaping and discrimination reversal learning: further evidence for the functional heterogeneity of the rodent frontal cortex. J Neurosci. 2003;23:8771-8780.

64. Chang SE. Effects of orbitofrontal cortex lesions on autoshaped lever pressing and reversal learning. Behav Brain Res. 2014;273:52-56.

65. Burke KA, Takahashi YK, Correll J, Leon Brown P, Schoenbaum G. Orbitofrontal inactivation impairs reversal of Pavlovian learning by interfering with "disinhibition" of responding for previously unrewarded cues. Eur J Neurosci. 2009;30:1941-1946.

66. Panayi MC, Killcross S. Functional heterogeneity within the rodent lateral orbitofrontal cortex dissociates outcome devaluation and reversal learning deficits. Elife. 2018;7:1-27.

67. Tait DS, Brown VJ. Difficulty overcoming learned non-reward during reversal learning in rats with ibotenic acid lesions of orbital prefrontal cortex. Ann N Y Acad Sci. 2007;1121:407-420.

68. Yin HH, Knowlton BJ, Balleine BW. Lesions of dorsolateral striatum preserve outcome expectancy but disrupt habit formation in instrumental learning. Eur J Neurosci. 2004;19:181-189.

69. Cavedini P, Ferri S, Scarone S, Bellodi L. Frontal lobe dysfunction in obsessive-compulsive disorder and major depression: a clinical-neuropsychological study. Psychiatry Res. 1998;78:21-28.

70. Apergis-Schoute AM, Gillan CM, Fineberg NA, Fernandez-Egea E, Sahakian BJ, Robbins TW. Neural basis of impaired safety signaling in obsessive compulsive disorder. Proc Natl Acad Sci. 2017;114:3216-3221.

SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section at the end of the article.

How to cite this article: van den Boom B, Mooij AH, Misevičiūtė I, Denys D, Willuhn I. Behavioral flexibility in a mouse model for obsessive-compulsive disorder: Impaired Pavlovian reversal learning in SAPAP3 mutants. Genes, Brain and Behavior. 2019;18:e12557. https://doi.org/10.1111/gbb.12557