**Identification and functional evaluation of *GRIA1* missense and truncation variants in patients with intellectual disability: An emerging** **neurodevelopmental phenotype**

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**Abstract**

*Background: GRIA1* encodes the GluA1 subunit of α-amino-3-hydroxy-5-methyl-4-isoxazole propionate (AMPA) receptors, which are ligand-gated ion channels that act as excitatory receptors for the neurotransmitter *L*-glutamate (Glu). AMPA receptors (AMPARs) are homo- or heteromeric protein complexes with four subunits, each encoded by different genes, *GRIA1 to GRIA4*. Despite GluA1-containing AMPARshaving a crucial role in brain function, the human phenotype associated with deleterious *GRIA1* sequence variants has not been established. This study shows that mutations in *GRIA1* cause both dominant and recessive neurodevelopmental disorders.

*Methods:* Subjects with *de novo* missense and nonsense *GRIA1* variants were identified through international collaboration. Detailed phenotypic and genetic assessments of the subjects were carried out and the pathogenicity of the variants was evaluated using electrophysiological and biochemical analyses to characterize changes in AMPAR receptor function and expression. In addition, two *Xenopus gria1* CRISPR/Cas9 F0 models were established and successfully used to characterize the *in vivo* consequences.

*Results*: Seven unrelated patients with rare *GRIA1* variants were identified. One patient carried a homozygous nonsense variant (p.Arg377Ter), and six had heterozygous missense mutations (p.Arg345Gln, p.Ala636Thr, p.Ile627Thr, p.Gly745Asp) of which the p.Ala636Thr variant was recurrent in three patients. The cohort revealed subjects to have a recurrent neurodevelopmental disorder mostly affecting cognition and speech. Functional evaluation of major GluA1-containing AMPAR subtypes carrying the *GRIA1* variant mutations showed that three of the four missense variants profoundly perturb receptor function. The homozygous stop-gain variant completely destroys the expression of GluA1-containing AMPARs. In addition, data from the *Xenopus gria1*models shows transient motor deficits, an intermittent seizure phenotype and, by using a newly described method, a significant impairment to working memory in mutants.

*Conclusion*: These data support the first description of a new developmental disorder caused by both heterozygous and homozygous variants in *GRIA1* affecting AMPAR function.

**Keywords**

AMPA receptor, AMPAR, iGluR, Glutamate receptor 1, GLUR1, GLURA, Neurodevelopmental impairment, epilepsy, *Xenopus,* Free movement pattern Y maze, CRISPR.

**Introduction**

AMPARs belong to the ionotropic glutamate receptors (iGluR) superfamily of ligand-gated cation channels that mediate the majority of excitatory synaptic transmission in the central nervous system1. The primary function of AMPARs is to facilitate synaptic transmission by delivering excitatory postsynaptic currents (EPSC), but AMPARs are also involved in [synaptic plasticity](https://en.wikipedia.org/wiki/Synaptic_plasticity) mechanisms thought to underlie learning and memory2-4. AMPARs form as tetrameric assemblies of the four subunits, GluA1-4, encoded by the *GRIA1-4* genes5. The *GRIA1* gene encodes the 907 amino acid GluA1 subunit (Fig. 1A-C). GluA1 can assemble as a homomeric receptor or combine with GluA2-4 subunits into heteromeric AMPARs. Structurally, AMPARs have a four-layer structure with the amino terminal domains (NTDs) from each subunit forming an upper extracellular layer, the agonist-binding domains (ABDs) forming a middle layer containing four Glu binding sites, and the transmembrane domains (TMDs) forming a central, membrane permeating ion channel (Fig. 1B).

Neurodevelopmental disorders (NDDs) encompass a range of phenotypes such as intellectual, behavioral, memory, or motor deficits and are estimated to affect 1-3% of the population in Western countries6-9. A growing body of evidence suggests that a substantial proportion of NDDs have monogenetic causes affecting key proteins in excitatory neurotransmission10-12, including *GRIA* genes. Advances in understanding the genetic architecture of the brain may begin to unravel the genetic causes for NDDs. In particular, appreciating the critical role of AMPARs in excitatory neurotransmission and synaptic plasticity mechanisms is yielding a new perspective for their causal role in NDDs. Previous studies have implicated *GRIA2, GRIA3*, and *GRIA4* genes in NDDs, but *GRIA1* has not been established as a disease-causing gene. Although variants in *GRIA1* have been identified through WES and WGS studies in NDD patient cohorts13-16 and a potential mutation 'hotspot' in *GRIA1* has been postulated15, to date, there has been no detailed phenotypic analysis or functional work completed to classify these variants beyond uncertain clinical significance.

A cohort of unrelated NDD patients with both new and previously-reported GRIA1 missense variants was identified through collaboration. In particular, this included a patient with a homozygous *GRIA1* stop-gain mutation that truncates the GluA1 subunit and appears to disrupt expression of any GluA1-containing AMPAR subtype. Functional evaluation of the impact of the *GRIA1* variants on the function of GluA1-containing AMPAR subtypes showed three out of four to have profound gain- or loss-of-function effects on important functional features of homomeric and heteromeric GluA1 receptor subtypes. In addition, the *in vivo* effects of disruption of GluA1-containing AMPAR expression was assessed in genetically altered *Xenopus* tadpoles using a novel behavioral model for measuring working memory. The results provide evidence that *GRIA1* variants are the cause of a monogenic NDD characterized by ID, speech and language delay, poor sleep, abnormal electroencephalogram (EEG) with or without seizures, normal brain imaging, and endocrine abnormalities, adding to the existing collection of *GRIA* related NDDs.

**Materials and methods**

**Cohort analysis**

Subjects were identified through the authors’ clinical practice, GeneMatcher17 or ClinVar databases18. Medical information including birth parameters, epilepsy, electroencephalograms (EEGs), developmental histories, brain MRIs and physical examinations were collected from the local healthcare providers. The study was conducted in agreement with the Declaration of Helsinki and approved by the local ethics committees. Since all patients had cognitive impairment, their parents or legal guardians gave informed consent.

**Genetic identification and analysis**

Subjects 1 to 5 were investigated by WES or WGS ordered by primary healthcare providers or as part of larger research studies (Supplementary patient information). Patient 6 was investigated by targeted panels (Supplementary patient information). Information on genetic analysis was not available for patient 7.All variants were annotated using the NM\_000827.3 (GRCh37/hg19) transcript of *GRIA1*. The functional consequences of missense variants were predicted using calculation of Combined Annotation-Dependent Depletion (CADD) scores19, and Sorting Intolerant From Tolerant (SIFT)20 and Polymorphism Phenotyping v2 (PolyPhen2)21 analysis. The Genome Aggregation Database (gnomAD v.2.1.1; https://gnomad.broadinstitute.org/) was employed to determine the frequency of the variants in control populations.

**Materials**

All chemicals were from Sigma-Aldrich (St. Louis, MO) unless otherwise stated. Dulbecco's modified Eagle's medium (DMEM), fetal bovine serum, trypsin, and penicillin-streptomycin were from Invitrogen (Carlsbad, CA). DNA modifying enzymes were from New England Biolabs (Ipswich, MA) except PfuUltra II Fusion HS DNA polymerase (Agilent, Carlsbad, CA). Tissue cell culture plasticware was from Sarstedt (Nümbrecht, Germany) unless otherwise stated. Cyclothiazide (CTZ), kainic acid, and NASP were from HelloBio (Bristol, UK). A DNA construct encoding GFP-GluA1 was a gift from Roberto Malinow (University of California, San Diego), and the construct encoding mCardinal-Farnesyl-5 was a gift from Michael Davidson (Addgene plasmid # 56159; http://n2t.net/addgene:56159; RRID:Addgene\_56159).

***Xenopus* *tropicalis* husbandry**

Adult Nigerian strain *Xenopus tropicalis* were housed and maintained within the European *Xenopus* Resource Centre, University of Portsmouth, in recirculating systems at 24 – 25 oC with 15% daily water changes on a 13 – 11-hour light-dark cycle. All work was conducted in accordance with the Home Office Code of Practice under PPL 79/8983 and PP4353452 following approval from the University of Portsmouth's Animal Welfare and Ethical Review Body. For egg recovery, female *Xenopus tropicalis* were primed with 10 IU of Human Chorionic Gonadotropin (Chorulon, Intervet) and boosted with 100 IU the following morning. Egg clutches were fertilized with cryopreserved sperm (EXRC)22-24. Embryos were cultured at 27oC for the first 24 hours and 24oC thereafter in 0.05 X Marc's Modified Ringer's (MMR in mM: 22 NaCl, 0.5 KCl, 0.5 CaCl2, 0.25 MgCl2, 1.25 HEPES, pH 7.4), in complete darkness with a 50% media change every other day, with twice-daily health checks. Once at feeding stages, tadpoles were fed a mixed diet of spirulina and sera micron twice daily, 5 days per week and once daily, 2 days per week.

**Animal strains and genetic alteration**

Experimental data presented in this study were obtained from either wild-type (WT) Nigerian strain *Xenopus tropicalis* or a transgenic line expressing GFP in differentiated neural tissue [Xtr.Tg(tubb2b:GFP)Amaya] RRID:EXRC\_3001 (from here on referred to as tubb2bGFP). In these background strains, three different CRISPR/Cas9-based mosaic models were made and analyzed as detailed below: a *tyrosinase* crispant in which exon 2 was targeted (*Xtr.tyr*em1E*X*RC referred to as “*tyr* crispant” in the text and used as a control); a *gria1* crispant in which exon 2 was targeted (*Xtr.gria1*em1E*X*RC referred to as “*gria1* knockout” in the text and described in Supplemental Fig. S2) and a second *gria1* crispant model in whichexon 8 was targeted (*Xtr.gria1*em2E*X*RC referred to as “*gria 1* crispant” in the text and described in Fig. 2.).

**Generation of knockout animals using CRISPR-Cas9**

The target regions within *gria1* (exon 2 and exon 8) were identified using Xenbase25, and single guide RNAs (sgRNA) were designed using v9.1 of the *X. tropicalis* genome and later blasted against v10 to check for additional off-target sites. Two single-stranded oligonucleotide templates (Supplementary Table S1) for sgRNA synthesis were selected for each region based on the following criteria: high mutagenic activity, minimal predicted off-target events, and a high frameshift frequency using the CRISPRscan26 and inDelphi algorithms27. Following the Taq-based method described by Nakayama *et al.*28*,* single-stranded oligonucleotides (Invitrogen, UK) containing the T7 promoter were annealed and extended with the universal CRISPR oligonucleotide, this template was then transcribed using a T7 Megashortscript kit (Invitrogen, UK). The resulting sgRNAs were purified using SigmaSpin™ Sequencing Reaction Clean-Up columns (Sigma-Aldrich), quantified using a NanoDrop 1000 spectrophotometer (Thermo Fisher Scientific, Loughborough, UK), analyzed by agarose gel electrophoresis, and stored at -80oC as single-use aliquots. Across all experiments, 1000 pg sgRNAs were co-injected with 2.6 ng Cas9 protein (Spy cas9 NLS, New England Biolabs) into single-cell *X. tropicalis* embryos. The efficiency of indel formation was assessed in genomic DNA from crispant embryos. Briefly, lysates were prepared from three batches of embryos collected at Nieuwkoop and Faber (NF) stage 10 and stage 41, by incubation at 56 °Cfor 2 hours in 50 mM Tris, 1 mM EDTA, 0.5% [v/v] Tween-20, 100µg/ml Proteinase K, pH 8.5. Primers for PCR amplification of the target regions of interest were designed using Primer3 software29 (Supplementary Table S1) and blasted against v10 of the *Xenopus tropicalis* genome (Xenbase). Amplicons were column purified (SmartPure PCR Purification Kit, Eurogentec, Belgium), Sanger sequenced (Genewiz, UK), and the resulting trace files were compared using ICE v2 CRISPR Analysis Tool software (Synthego, Redwood City, CA). Indels were confirmed by Sanger sequencing of subcloned PCR amplicons from mutant animals.

**Phenotypic analysis of crispant tadpoles**

To identify gross morphological differences between uninjected tadpoles and crispant tadpoles, animals were anesthetized in 0.025% w/v tricaine mesylate solution and visually inspected using an AxioZoom V16 stereomicroscope (Zeiss, Jena, Germany) with fluorescence for visualizing GFP-expressing animals. To inhibit melanogenesis for the study of neural transgene expression, tadpole medium was supplemented with 75 µM 1-phenyl 2-thiourea (PTU) after hatching at NF26. Analysis of working memory was examined at stage NF50, using the Zantiks MWP unit (Zantiks Ltd, Cambridge, UK) and the free-movement pattern (FMP) Y-maze30. The FMP Y-maze is validated for assessing spatial working memory and cognitive flexibility in zebrafish and has been applied to both mice and humans but has never been used in *Xenopus*. The FMP Y-maze quantifies deviations from randomness in search strategies by analyzing a continuous log of arm entries in the maze in terms of discrete choices (i.e. 'Left' or 'Right'), grouped into a series of four overlapping choices, called 'tetragrams' (e.g. LRLR, LLLR, etc.). Vertebrates such as zebrafish, mice, and humans show a common dominant search strategy (~25 - 30% of turns) of sequentially alternating left/right choices (LRLR or RLRL)30; 31. This pattern is abolished with memory-blocking drugs and reduces in aging 30. The task is also able to quantify behavioral flexibility by analyzing the change across time in alternation strategies30. For example, behavioral flexibility is reduced with dopamine D1 receptor antagonists and in the presence of acute stress30-32.

*Xenopus tropicalis* tadpoles were placed in semi-translucent, acrylic inserts containing two identical Y-mazes (three 10 x 10 x 25mm arms and a 10 x 10 x 10mm central zone). Inserts were filled with 0.05 X MMR and contained a dilute but equally distributed tadpole food mix. The Y-Maze arms were of equal size with no intra-maze cues, and lights were maintained off during each trial to match the rearing conditions of the tadpoles. An infra-red video camera was used for live monitoring and recording of the movement of individual animals. Sample size estimations were calculated *a priori* from pilot studies performed on 16 uninjected control tadpoles and 16 *gria1* knockout tadpoles, using the G\*Power software package version 3.1.9.733. Tadpoles were transferred into the Y-mazes and placed into the Zantiks MWP unit for a pretrial time of 120 seconds to acclimatize. They were then tracked for one hour of free search. Data from each trial were output in two forms: zone entries/exits over time and an AVI video file (with live tracking). Zone entries and exits were converted into left or right turns, grouped into overlapping sequences of four turns (tetragrams) using customized Excel spreadsheets, and normalized against the total number of moves. For experiments with MK-801 ((+)-5-methyl-10,11-dihydroxy-5H-dibenzo(a,d)cyclohepten-5,10-imine]), the concentration and delivery was based on previously published work in Zebrafish30. Specifically, tadpoles were placed in 100 mL beakers containing 50mL 0.05X MMR and 0.75mg/L MK-801 for 2 hours before evaluation in the Y-maze.

**Molecular Biology**

Mutations in *GRIA1* (MIM 138248) were introduced by site-directed mutagenesis into their corresponding positions in rat cDNA expression constructs encoding GluA1. Specifically, the plasmid vector pXOOF34 containing cDNA for the unedited flip isoform of rat *GRIA1* gene was used for site-directed mutagenesis and subsequent heterologous expression in mammalian cells or generation of mRNA for microinjection in *Xenopus laevis* oocytes (XOs). Site-directed mutagenesis was performed using the QuickChange mutagenesis kit (Stratagene, La Jolla, CA). The mutations were verified by Sanger DNA sequencing of the entire GluA1 coding region (GATC Biotech, Constance, Germany). For analysis of cell-surface expression of WT and mutant GluA1 receptors, cDNA encoding a β-lactamase (blac) enzyme was inserted in the GluA1 cDNA in between the segments encoding the *N*-terminal signal sequence and the NTD using In-Fusion cloning (Promega, Mountain View, CA, USA). Specifically, a PCR-amplified DNA fragment encoding blac flanked by two short amino acid linkers (GGSGS and GGSG) was inserted in-frame between the signal sequence and the NTD using an *Xho*I restriction site introduced by site-directed mutagenesis of codon 24 and 25 to create WT and mutant blac-GluA1 constructs. For imaging of the expression patterns of WT and mutant GFP-tagged GluA1 subunit, a GFP-tagged GluA1 construct35 and the red fluorescent membrane-reporter mCardinal-Farnesyl-5 were PCR subcloned into the CMV-IRES cassette of the pmLINK plasmid vector36. Two pmLINK plasmids can be fused by a two-step recombination strategy to form an expression construct for co-expression of cDNA inserts from two identical CMV promoter expression cassettes36. Specifically, pMLink-eGFP-GluA1-WT and pMLink-eGFP-GluA1-R377\* were digested with *Swa*I and combined with *Pac*I digested pMLink-mCardinal-Farnesyl-5 by use of Gibson assembly (Gibson Assembly® Cloning Kit, New England Biolabs) to generate pMLink-eGFP-GluA1-WT-mCardinal-Farnesyl-5 and pMLink-eGFP-GluA1-R377Ter-mCardinal-Farnesyl. For co-expression studies of WT and mutant GluA1 with the AMPAR GluA2 subunit and the auxiliary TARP-class subunit gamma-2, pXOOF plasmid constructs containing the flip isoform of rat GluA2 in the R-edited form (GluA2R) and gamma-2 were used. When used as templates for *in vitro* transcription of mRNA, all pXOOF plasmid constructs were linearized downstream of the 3' untranslated region using *Nhe*I, purified using a NucleoSpin DNA clean-up kit (Macherey-Nagel, Düren, Germany), and stored at a concentration of 1.0 ug/uL at -20 ˚C until use. cRNA transcription was performed using the ARCA mRNA synthesis kit (NEB, Madison, WI, USA). The resulting mRNA was column purified using NucleoSpin RNA Clean-up kit (Macherey-Nagel), diluted to 0.5 ng/nL, and stored at -80 ˚C until use.

**Mammalian Cell Culturing and Expression**

HEK293T cells (American Type Culture Collection, Manassas, VA) were cultured in DMEM medium supplemented with 10% v/v fetal bovine serum, 100 units/ml penicillin, and 100 μg/ml streptomycin at 37 °C in a humidified 5% CO2 environment. For expression of WT and mutant GluA1 in HEK293 cells, TransIT-LT1 DNA transfection reagent (Mirus, Madison, WI) was used as described previously34. Briefly, HEK293 cells in suspension (1e6 cells/mL) were mixed with DNA/transfection complex formed by mixing plasmid DNA, TransIT-LT1 reagent, and DMEM in a 1:3:90 ratio and immediately plated into poly-*D*-lysine coated glass-bottom 96-well plates (MatTek Corporation, Ashland, MA) at 1e6 cells and 1 µg plasmid DNA per well and incubated for two days before experiments.

***Xenopus laevis* oocyte expression**

Defolliculated XOs(stage V to VI) were prepared and injected with mRNA as described previously37. The care and use of *Xenopus laevis* were in strict adherence to a protocol (license 2014−15−0201−00031) approved by the Danish Veterinary and Food Administration. Injected XOs were incubated at 18 °C in Modified Barth’s Solution (MBS) containing (in mM) 88 NaCl, 1 KCl, 0.41 CaCl2, 2.4 NaHCO3, 0.33 Ca(NO3)2, 0.82 MgSO4, 5 Tris (pH 7.4) supplemented with 50 µg/ml gentamycin until use.

**Cell-surface expression levels**

Relative levels of surface-expressed blac-tagged GluA1 were quantified in living cells by measuring the conversion rate of the membrane-impermeable blac substrate nitrocefin by simple absorption spectroscopy38. For analysis in XOs, injected oocytes were placed individually in wells of clear-bottom 96-well plates containing 100 µL MBS followed by the addition of nitrocefin to a final concentration of 50 µM in a total volume of 200 µL per well and incubated at 37 ˚C for 3 hours. During the incubation time, 50 µL samples of medium were removed every 30 minutes, and the absorbance at 486 nm of the nitrocefin conversion product was determined using a microplate reader (Safire2, Tecan, Maennedorf, Switzerland), plotted as a function of time of sampling, and the rate of nitrocefin conversion was determined by linear regression analysis of the slope of the curve in the linear range.

**Western blot analysis**

Transfected HEK293 cells lysate was mixed 1:1 with 2xSDS sample buffer composed of 50 mM Tris-HCl, 2% w/v SDS, 10% w/v glycerol, 1% w/v β-mercaptoethanol, 12.5 mM EDTA, 0.02 w/v % bromophenol blue (pH 6.8) and heated at 65 ˚C for 5 minutes. 20 uL samples were loaded on freshly prepared 10% polyacrylamide gels, resolved based on molecular weight through electrophoresis, and transferred to a polyvinylidene difluoride membrane (Sigma-Aldritch) and incubated with antibodies for GluA1 for 2 h at RT, washed twice for 15 minutes at RT, then incubated with alkaline phosphatase-conjugated secondary antibody for 2 h at RT, and finally rinsed in water for 15 min. The immunoreactive protein content in the membrane was visualized using alkaline phosphatase mediated conversion of 5-bromo-4-chloro-3-indolyl phosphate (BCIP)/nitro blue tetrazolium (NBT) into an insoluble blue-purple product (SIGMAFAST™ BCIP/NBT system, Merck).

**TEVC electrophysiology**

Glass micropipettes (0.69 mm ID/1.2 mm OD, Harvard Apparatus, Holliston, MA) were pulled on a Sutter P-1000 micropipette puller (Sutter Instruments, Novato, CA) to a tip resistance of 0.5-2.5 MΩ and filled with 3 M KCl. Oocytes were clamped using a two-electrode voltage-clamp amplifier (OC-725C, Warner Instruments, Hamden, CT) and continuously perfused with Frog Ringer's solution containing (in mM) 115 NaCl, 2 KCl, 5 HEPES, and 1.8 BaCl2 (pH 7.6 with NaOH) by gravity-assisted perfusion at flow rates of 2 to 4 mL/min into a vertical oocyte flow chamber with a volume of 0.3 mL39. Compounds were dissolved in Frog Ringer's solution and added by bath application. Concentration-response data were generally recorded at holding potentials of -40 mV; otherwise, in the -20 to -80 mV range. Each compound solution was applied for 10 to 60 s depending on time needed to obtain steady-state currents. Data acquisition was accomplished using a CED 1401plus analog-digital converter (Cambridge Electronic Design, Cambridge, UK) interfaced with a PC running WinWCP software (available from Strathclyde Electrophysiology Software, University of Strathclyde, Glasgow, UK). Concentration-response experiments were performed by measuring agonist-evoked current during stepwise application of increasing concentrations of agonist, as illustrated in Fig. 3E. All experiments were performed at RT.

**Confocal Imaging**

A Leica SP2 confocal microscope equipped with an argon laser, a helium/neon laser, and 63X 1.2 Na HCX PL APO water-corrected objective was used. GFP-tagged WT and mutant GluA1 were visualized using 488 laser lines at 25-35% input power as excitation sources and emission measurement in the 500-560 nm spectrum ranges. In addition, the co-expressed plasma membrane marker mCherry-Farnesyl-5 was visualized using the 633-nm helium/neon laser line at 25-35% input power and collection of emission in the 640 to 700-nm spectrum range. Overlay images were produced with Leica LAS AF Lite software (Leica Microsystems GmbH, Wetzlar, Germany).

**Data and statistical analysis**

To construct concentration-response curves from electrophysiological data, agonist-evoked current responses from individual oocytes were determined from TEVC traces using ClampFit 10 software (Molecular Devices, San Jose, CA) and normalized to the current response by maximal agonist concentration. Composite concentration-response plots were constructed from normalized responses from 8 to 30 oocytes and fitted using GraphPad Prism v6.01 (GraphPad Software, San Diego, CA, USA) to a four-variable Hill equation:

In this equation, *bottom* is the fitted minimum response, *top* is the fitted maximum response, *nH* is the Hill slope, and X is the agonist concentration, and EC50 is the half-maximally effective agonist concentration, respectively. All statistical analyses of data from TEVC experiments were performed in GraphPad Prism 9. Unless otherwise stated, summary TEVC data are represented as mean with 95% confidence interval (CI) from n cells. One-way analysis of variance (ANOVA) with Dunn’s *post hoc* multiple comparison test was performed for comparisons of three or more groups in which the data were normally distributed, and where a p-value <0.05 was considered significant

For the FMP Y-maze experiment, data across groups were compared using an ANCOVA with significant effects assessed by Dunn’s *post hoc* multiple comparison test, where a p-value <0.05 was considered significant. Grouped data are shown as the mean ± the standard error of the mean (SEM). Unless otherwise stated, statistical significance was denoted as follows: \*p < 0.05, \*\*p < 0.01, and \*\*\*p < 0.001.

**Results**

**Genetic and clinical findings in NDD patients with homozygous and heterozygous GRIA1 variants.** Seven unrelated NDD patients with rare *GRIA1* variants were identified and included in the study via the author's clinical practice, direct communication, and through the GeneMatcher17 or ClinVar database18. In all cases, the *GRIA1* variants were identified from clinical or research genetic analysis aimed to determine the genetic cause underlying the patient's NDD and were found to include four missense variants (c.1906G>A; p.Ala636Thr, c.1880T>C; p.Ile627Thr, 2234G>A; p.Gly745Asp, and c.1034G>A; p.Arg345Gln) and one truncating variant (c.1129C>T; p.Arg377Ter) (Table 1). All missense variants were heterozygous and arose *de novo*, whereas the truncating stop-gain variant was homozygously inherited from consanguineous parents. Three patients harbored the p.Ala636Thr variant (patients 2-4), which previously has been reported as a recurrent *de novo* *GRIA1* variant15. An overview of the genetic and bioinformatic data is provided in Table 1, while patient clinical features are provided in Table 2. Individual case stories of patients are included in the supplementary information (Supplementary patient information). Patient ages ranged from 7 to 26 years and included an equal number of males and females and one patient for which sex was not reported (Table 2). All patients were diagnosed with ID. In addition, patients were most severely affected in terms of verbal abilities: patients 1, 2, and 3 remained non-verbal while patients 4 and 6 communicated using either simple words or short sentences. Data on speech was not available for patients 5 and 7. All patients could walk independently before 24 months of life (ranging from 12 to 17 months). Patients 1, 2, 3, 4, and 6 had cognitive impairment ranging from moderate to severe based on clinical impression. Due to limited data access, it was impossible to determine the severity of cognitive impairment for patients 5 and 7. Also, data on behavioral phenotype were not available for subjects 5 and 7 but were reported in all remaining patients. These included recurring themes, such as anxiety, ASD, and ADHD phenotypes (Table 2). Patient 6 was described as having a low anger threshold and had challenging behavior such as inconsolable upset, anger tantrums, and occasional aggressive outbursts; however, no self-injurious behavior was described. Patients 3, 4, and 6 had an autism spectrum disorder. Patients 1, 3, and 4 were reported to have a sleeping disorder necessitating treatment with melatonin. Normal brain MRI was reported for all patients except patient 7, for which information on MRI status was not available. None of the patients were reported to have a movement disorder. Epileptic seizures were only reported in patient 1, who carried the homozygous truncating variant and were reported to be treatment-resistant focal seizures (Supplementary patient information). An endocrine disorder was identified in two subjects included premature puberty in patient 1 and hypothyroidism plus polycystic ovarian syndrome in patient 6.

Evaluation of effects of *GRIA1* variants on GluA1 subunit expression

The amino acid residues in the GluA1 subunit protein affected by patient *GRIA1* missense variants are located in different subunit domains (Fig. 1A-B). As shown in the cartoon representation of a single GluA1 subunit protein (Fig. 1A), Arg345 is located in the NTD, Gly745 is located in the ABD that contains the Glu-binding site, and the Ile627 and Ala636 are located within the M3 transmembrane domain of the TMD. The Arg377 residue affected by the stop-gain variant in patient 1 is located in the C-terminal of the NTD close to the linker that connects the NTD to the ABD (Fig. 1A). The stop-gain mutation (R377\*) is therefore predicted to cause expression of the NTD alone. All of the residues are located in GluA1 sequence regions that are highly conserved among species and the other AMPAR subunits GluA2-4 (Fig. 1B). Except for Arg345, no other missense variants affecting the residues exist in the Genome Aggregation Database (gnomAD), which may indicate sensitivity to missense variation. The p.Ile627Thr, p.Ala636Thr, and p.Gly745Asp variants that were not present in gnomAD were classified as potentially damaging by various *in silico* tools for the prediction of deleteriousness of single missense variants (Table 1). Moreover, missense tolerance ratio (MTR) analysis40 of the *GRIA1* coding sequence reveals the codons encoding Ile627 and Ala636 to have MTR scores (0.41 and 0.33, respectively) that are in the lower 5% percentile and close to the global minimum of the entire subunit sequence (Fig. 1D), strongly indicating an unusually high sensitivity of these residues to missense variation. Gly745 also has a low MTR score (0.66) and is located in a local minima region of the MTR (Fig. 1D) and, therefore, is also predicted to have sensitivity to missense variants. In contrast, Arg345 is located in a sequence region without reported unusual sensitivity to missense variants.

For evaluation of the effect of the *GRIA1* variants on expression and function of GluA1-containing AMPARs subtypes, the mutations reflecting the *GRIA1* variants p.Arg345Gln (R345Q), p.Arg377Ter (R377\*), p.Ile627Thr (I627T), p.Ala636Thr (A636T), and p.Gly745Asp (G745D) were generated in the GluA1 subunit (Materials & methods). Mutational effects on the ability of the GluA1 subunit protein to fold correctly, assemble into receptors, and traffic to the cell surface membrane were first characterized using a β‐Lactamase (β‐lac) enzyme reporter assay34; 41; 42. Specifically, β‐lac was fused to the extracellular N-terminal of WT and mutant GluA1. The resulting β‐lac tagged subunit constructs were expressed in *Xenopus* *laevis* oocytes (*Materials & methods*). Following two days of expression, the relative levels of WT and mutant receptor present at the oocyte cell surface were determined by measuring the rate of β‐lac catalyzed cleavage of the cell‐impermeable chromogenic substrate nitrocefin added to the cells38; 42. All mutants except R377\* displayed β-lac activity similar to WT GluA1, which indicates that the mutations do not affect normal efficiency of subunit expression, folding, assembly, and surface trafficking (Fig. 1E). In contrast, R377\* expressing oocytes showed a similar level of β‐lac activity to un-injected oocytes (0.002 OD/h versus 0.001 OD/h; *p* = 0.72; Fig. 1E), supporting that the R377\* mutation truncates the GluA1 subunit in the NTD such that the subunit lacks the ABD and TMD that is essential for subunit assembly into functional receptors in the membrane (Fig. 1). This was further tested by expressing WT GluA1 and the R377\* mutant tagged with green fluorescent protein (GFP) in the N-terminal of the NTD in HEK293 cells together with the red-fluorescent cell-surface membrane reporter protein construct mCardinal-farnesyl (Materials and Methods). Confocal imaging of GFP fluorescence in the transfected cells showed clear membrane localization of WT GFP-GluA1 protein that overlapped with mCardinal-farnesyl fluorescence. In contrast, GFP fluorescence for GFP-GluA1-R377\* cells was confined to the intracellular compartments (Fig. 1F). Furthermore, Western blot analysis of GluA1 protein size in protein extracts from WT and R377\* transfected HEK293 cells were performed (Fig. 1G). Analysis with an antibody directed against an NTD epitope showed GluA1 protein with a size corresponding to full-length GluA1 (100 kDa) in WT transfected cells, whereas R377\* transfected cells showed a protein product with a size corresponding to truncation at Arg377 (43 kDa). A similar analysis using an antibody directed at the GluA1 C-terminal showed a 100 kDa size for the WT protein and detected no GluA1 protein in extracts from R377\* transfected cells (Fig. 1G), indicating that translational read-through of the stop codon generated by the p.Arg377Ter variant does not occur. Together, these data show that the homozygous *GRIA1* variant p.Arg377Ter identified in patient 1 is a stop-gain variant that truncates GluA1 at Arg377 and prevents any expression of functional GluA1 subunit in human and amphibian cells.

Indels in exon 8 cause working memory deficits in tadpoles.

The homozygous stop-gain variant p.Arg377\* was modeled in *Xenopus tropicalis* tadpoles using CRISPR-Cas9 to investigate how the disrupted expression of the GluA1 subunit contributes to neurological development and behavior.Gene editing in *Xenopus* speciesis now so efficient that analysis is routinely performed in founder animals, enabling rapid testing to support the causality of genetic disruption across a range of genes43-46. To support the use of *X. tropicalis* to model variants in *gria1,* both human and *Xenopus tropicalis* share identical gene structures and produce proteins that are >87% conserved (Supplementary Fig. S1A and B). Additionally, the exon 8 target region corresponding to the homozygous nonsense variant identified in Patient 1 (p.Arg377\*) is well conserved in *Xenopus* (Supplementary Fig. S1C). CRISPR/Cas9 editing (Materials and Methods) was used to disrupt exon 2 (*Xtr.gria1*em1E*X*RC, *gria1* knockout) or the genomic DNA encoding 18 amino acids upstream of Arg377 in exon 8 (*Xtr.gria1*em2E*X*RC, *gria1* crispant) (Supplementary Table S1 and Supplementary Fig. S1). Arg377 is the penultimate amino acid of exon 8, and targeting upstream of the variant was chosen to avoid altered splicing outcomes in the crispant model. Sanger sequencing of the target region within exon 8 in genomic DNA samples collected from crispant tadpoles demonstrated a good penetrance of indels. Further, sequencing of subcloned genomic amplicons revealed none of the sequenced clones represented the WT allele and demonstrated that half of the sequenced clones truncated the protein (denoted by the asterisk, Fig. 2A). The genotype of the *gria1* knockout model targeting exon 2 is predominantly a 7bp deletion(>75% indels,Supplementary Fig. S2A).

All phenotyping experiments were replicated in embryos from at least three different females, and no consistent early developmental abnormalities were noted across these experiments. Transient motor differences were seen in almost all post-hatching knockout (27 of 30 animals) and crispant tadpoles (23 of 30 animals). Both types of *gria1* mutantswere slower to hatch than controls and subsequently displayed an abnormal escape response to tactile stimulation. Post-hatching control animals were observed to move away from all tactile stimuli (trunk and head), whereas both types of mutant animals responded either by moving in tight circles or not at all. At later stages (NF42 onwards), all tadpoles were seen to develop normally, adopting an appropriate filter-feeding (head down, tail up) posture with the ability to navigate their environment freely. Imaging revealed no obvious or consistent craniofacial abnormalities in either model (Fig. 2B; Supplemental Fig. 2B). Similarly, no significant gross structural differences were noted in the forebrain, midbrain, or hindbrain regions when *gria1* knockout (Supplemental Fig. S2C) or *gria1* crispant tadpoles (Fig. 2C) were made in a tubb2bGFP background to enable visualization of the brain. A small number of tadpoles in each batch were observed to demonstrate episodic periods of abnormal behavior (5 of 50 animals), which included 'C-shaped' alternating axial contractions of the tail coupled with rapid changes in direction. These 'manic' bouts were followed by a prolonged and unusual period of immobility. Interestingly, this behavior is consistent with descriptions in the literature of seizures in *Xenopus* tadpoles47; 48.

All patients in this study have varying degrees of cognitive impairment (Table 2 and Supplementary patient information). Until now, quantitative measures of higher executive functions in *Xenopus* have not been described. Therefore, a novel free-movement pattern behavioral model was developed in *Xenopus tropicalis* tadpoles to assess the impact of *gria1* knockout on cognitive functions (Materials and Methods). Previous FMP Y-maze studies show a dominant vertebrate search strategy largely consisting of alternating left/right choice patterns (LRLR or RLRL). Importantly, this strategy can be impaired by pharmacological agents that disrupt working memory and cognitive flexibility30; 31. The results presented here show that control tadpoles demonstrate a predominant search strategy consisting largely of alternations (black bars, Fig. 2D, Supplemental Fig. 2D and 3A). The alternation strategy observed in control tadpoles was abolished following administration of the NMDA receptor antagonist MK-801 (green bars, Supplemental Fig. 3). Specifically, MK-801 treated tadpoles showed "primitive" search patterns consisting of repetitions (Supplemental Fig. 3A and D), similar to those observed in invertebrate models 30. These findings are consistent with those reported in rodent and zebrafish FMP Y-maze studies30; 49; 50, which show that glutamatergic disruption impairs spatial working memory and provides a further demonstration of the suitability of *Xenopus* tadpoles to study genetic disruption within *gria1*.

Next, the 1-hour FMP Y-maze assay was used to compare WT and *gria1* crispant (Fig. 2D-G) or *gria1* knockout tadpoles (Supplemental Fig. S2D-G). Both the *gria1* crispant (Fig. 2D) and *gria1* knockout tadpoles (Supplemental Fig. S2D) showed a significant decrease in alternations compared to WT tadpoles. This decrease in alternations was present across the entire 1-hour trial (Fig. 2E and Supplemental Fig. S2E) and is consistent with a working memory deficit. These results agree with the finding of short-term working memory deficits in the *gria1*-/- mouse model51-55, Unlike in the *Gria1-/-* mouse, there was no evidence to support hyperactivity in either type of mutant tadpole (Fig. 2F, Supplemental Fig. S2F). However, caution must be applied to this conclusion since these assessments were made on the movement into and out of zones rather than the total distance covered. Finally, the tadpoles show a relatively static search strategy and appear to perform a decreasing number of turns across the trial, suggesting that the 1-hour trial period could be reduced in future studies (Fig. 2G, Supplemental Fig. 2G).

Impact of variants on homomeric GluA1 receptor function

The GluA1 subunit can assemble as a homomeric receptor and as heteromeric receptor subtypes with GluA2-4 subunits56-58. First, the effect of the *GRIA1* variants on ligand-gated ion channel function of homomeric GluA1 receptors was evaluated by measuring current responses to Glu application in *Xenopus* *laevis* oocytes expressing WT and the mutant GluA1 subunits (Fig. 3). Like all AMPAR subtypes, homomeric GluA1 receptors display fast and profound desensitization1; 59; resulting in current responses to Glu that within milliseconds decline by more than 95% from a peak response level to a steady-state level that represents the majority of the receptor population to reside in the desensitized receptor state60; 61. This response waveform cannot be resolved in *Xenopus* oocytes. Therefore, to enable measurement of both desensitized and non-desensitized response levels, recordings of Glu-evoked currents were performed in the presence and absence of cyclothiazide (CTZ), a compound that blocks AMPAR desensitization (Materials & methods) (Fig. 3A-B). Un-injected oocytes did not show any responses to Glu in absence or presence of CTZ (Fig. 3A), confirming that *Xenopus laevis* oocytes do not express endogenous AMPA receptors at functional detectable levels. All mutants except R345Q showed current responses that were significantly different from WT (Fig. 3C; Table 4). As expected from the biochemical analysis of surface expression, oocytes expressing R377\* did not produce current response to Glu in the absence or presence of CTZ (Fig. 3A; Table 4); further confirming that the p.Arg377Ter variant prevents the expression of a functional GluA1 subunit. Also, no detectable desensitized or non-desensitized currents were observed for the G745D mutant (Fig. 3A; Table 4). The I627T mutant displayed detectable currents during desensitizing and non-desensitizing conditions, however, with amplitudes more than 10-fold lower than WT (Fig. 3A; Table 4). These results indicate that the p.Ile627Thr and p.Gly745Asp variants identified in patients 5 and 6, respectively, have loss-of-function effects on the function of the GluA1 subunit. In contrast, oocytes expressing the A636T mutant on average displayed 10-fold increased currents compared to oocytes expressing WT GluA1 (Fig. 3A and 3E; Table 4). Importantly, in individual A636T expressing oocytes, the Glu-evoked currents recorded sequentially in the absence and presence of CTZ had near-identical amplitudes (Fig. 3A), whereas currents in WT expressing oocytes increased 61-fold when desensitization was blocked by CTZ (Fig. 3D; Table 4); a factor that corresponds well to previously reported ratios between non-desensitized peak current and desensitized steady-state current amplitudes for homomeric GluA1 receptors recorded in HEK293 cells using fast-application protocols60; 61. This result strongly indicates that the A636T mutation disrupts the ability of homomeric GluA1 receptors to desensitize and explains the dramatic increase in current amplitude during non-desensitizing conditions. A similar analysis of the factor by which CTZ increased Glu-evoked current for R345Q and I627T showed increases that were not significantly different from WT (Fig. 3D and Table 4) to indicate that these mutations do not change receptor desensitization properties.

In addition to changing receptor desensitization properties, mutations can also change the current response to Glu by perturbing the activation properties of the receptor, e.g., the ability of the receptor to open the channel when the agonist is bound. A measure of the activation properties of AMPARs is to determine the efficacy of the weak partial agonist kainic acid (KA) relative to the full agonist Glu for activating receptor currents62; 63 (Fig. 3H). When desensitization was blocked, KA evoked current at WT GluA1 that was 21% of the Glu-evoked current (Fig. 3G; Table 4). The R345Q mutant showed similar KA efficacy (26%), indicating the mutation does not change receptor activation properties. In contrast, I627T showed a significantly lower KA efficacy of 5% compared to WT, indicating a decreased ability of the GluA1 subunit to translate agonist binding to channel opening (Fig. 3G and Table 4). This result explains the dramatic decrease in desensitized and non-desensitized current for I627T (Fig. 3B). In contrast, A636T showed a significantly increased KA efficacy (52%) (Fig. 3G and Table 4), indicating increased channel-opening ability. Notably, the effect of the A636T mutation has previously been studied for GluA164-66. Ala636 is the third Ala residue in the SYTANLAAF motif that is completely conserved in all eukaryotic iGluR subunits (Fig. 1D). This motif forms the upper M3 helix that lines the extracellular entrance to the channel and acts as a gate during channel opening67-69. Profound effects of mutation of this conserved Ala to Thr were first identified in the GluD2 iGluR in the mutant *lurcher* mice strain70; 71 that causes widespread neuronal cell death in homozygous animals. The same Ala-Thr mutation has subsequently been applied in several iGluR subunits to study the role of SYTANLAAF motif for, in particular, channel activation, including in GluA164; 66. The results presented here corroborate previous findings that the Ala-Thr mutation profoundly increases the ability of GluA1 to activate channel-opening and further shows that desensitization is abolished.

Dose-response curves for Glu were generated for WT and the R345Q, I627T, and A636T mutants and used to determine the half‐maximally effective concentration (EC50) of Glu (Materials and Methods) (Fig. 3E; Table 4). Due to the minimal current responses of the I627T mutant under desensitizing conditions, the dose-response experiments for this mutant were performed in the presence of CTZ. The R345Q and I627T showed EC50 values identical or very close to WT (Table 4), indicating that these mutations overall do not change receptor sensitivity to Glu. In contrast, the A636T mutant showed greatly increased sensitivity towards Glu compared to WT, leading to a 25-fold reduced EC50 (Fig. 3F; Table 4), which agrees with previous reports64-66.

Impact of variants on heteromeric GluA1 receptor function and in presence of the auxiliary subunit γ-2

Homomeric GluA1 AMPARs are thought to exist *in vivo*72; however, heteromeric GluA1/A2 and GluA1/A3 receptors are considered to constitute the main population of GluA1-containing receptors in most CNS regions56-58; 73; 74. Furthermore, most native AMPARs form a complex with auxiliary subunits that regulate their function, such as the transmembrane AMPA receptor regulatory proteins (TARPs)75. To evaluate the effects of the *GRIA1* variants in the context of heteromeric receptors, WT and mutant GluA1 were expressed together with the GluA2 subunit and the electrophysiological studies of receptor function were repeated (Fig. 4). Specifically, WT and mutant GluA1 were expressed with GluA2 subunit in the R-edited (GluA2R) form in a 1:2 ratio which previously has been shown to result in the majority of GluA1 subunits to assemble with GluA2R to form a receptor population mainly composed of GluA1/A2R receptors (Material and methods)37; 76. Heteromeric assembly was conﬁrmed by measuring the current-voltage (IV) relationship of Glu-evoked currents (Fig. 4B). The IV relationship changes from inwardly rectifying for homomeric GluA1 to linear when GluA1 assembles with GluA2R subunits (Fig. 4B). Except for R377\*, all of the mutants displayed a near-complete shift from inwardly rectifying IV curves for homomeric expression to linear IV curves when co-expressed with GluA2R, showing that the mutations do not affect the ability of GluA1 to preferentially assemble with GluA2R to form heteromeric receptors (Fig. 4B). Also, mutant IV curves were identical to the equivalent WT, confirming that the mutations do not change rectification properties of homo- or heteromeric receptors. Although the GluA2R subunit can form homomeric receptors, these have very low channel conductance and traffic poorly to the cell surface77; 78. Consequently, homomeric GluA2R rarely produce detectable currents in *Xenopus* oocytes or other cells and are therefore not a concern in functional studies. In addition, to further mimic native AMPARs, GluA1/A2 receptors were also expressed with the prototypical AMPAR auxiliary TARP-class subunit γ-2 (also known as stargazin) (Materials and Methods)79. Successful incorporation of γ-2 subunits into the receptor complex was determined by measuring the ratio of sequential currents evoked by GLU and KA in the absence of CTZ (Fig. 4A). Previous work has shown that γ-2 increases steady-state current to KA relatively more than the Glu current, and determination of the KA/GLU current ratio provides a robust test for functional expression of AMPARs in complex with γ-237; 79-83. For WT and all mutants except A636T, γ-2 co-expression increased the KA/Glu ratio approximately 4- to 6-fold and maintained linear IV curves, verifying the formation of heteromeric GluA1/A2R receptors in complex with γ-2 (Fig. 4B and Supplementary Fig. S4). Notably, as the GluA1/A2-A636T receptor showed a profoundly increased KA/Glu ratio, it is likely that further increase of this ratio as measure for γ-2 incorporation is not possible.

On this background, mutant WT and mutant GluA1/A2R receptors with and without γ-2 were characterized for changes in the average current response, desensitization, and activation properties, and Glu EC50 using similar electrophysiological recording protocols as for homomeric GluA1 (Fig. 4A and Supplementary Fig. S4). Co-expression of R377\* with GluA2R did not produce currents, showing that the NTD-truncated GluA1 subunit cannot form a functional receptor with GluA2R. However, upon co-expression with γ-2, very small currents (10-20 nA) were detected in some oocytes (Fig. 4A; Table 4), but the amplitude of the currents was more than 300-fold smaller than the mean currents in oocytes expressing WT GluA1/A2R with γ-2 and, therefore, likely originate from homomeric GluA2R receptors, which previously have been shown to produce detectable currents when co-expressed with γ-237.

Similar to the results from homomeric receptors, GluA1-I627T/GluA2 receptors showed currents that were significantly smaller than WT both during desensitizing conditions (40 nA for I627T versus 511 nA for WT; Fig. Table 4) as well as non-desensitizing conditions (2728 nA for I627T versus 7051 nA for WT; Table 4). Also similar to homomeric I627T, the desensitization ratio was not significantly different from WT (Fig. 4D and Table 4), and the KA/Glu efficacy was decreased (9% for G745D versus 30% for WT; Fig. 4D and Table 4). These results show that the effect of the I627T mutation on GluA1 activation properties lead to heteromeric GluA1/A2 receptors with reduce current amplitudes. However, when WT and I627T was co-expressed with the γ-2 auxiliary subunit, the mean current amplitudes were not significantly different (Fig. 4A and C; Table 4). Generally, inclusion of the γ-2 auxiliary subunit in AMPAR subtypes enhances receptor activation by increasing the efficiency of receptor subunits to translate agonist binding to channel opening79; 84; 85, and this effect is manifest as a marked increase in apparent KA efficacy79; 86. Indeed, when γ-2 was co-expressed with the heteromeric receptors, the KA/Glu response ratio increased from 30% to 74% for WT (Fig. 4A and D; Table 4). This effect was also observed for the I627T mutant where KA/Glu increased from 6% to 66% (Fig. 4A and D; Table 4). Therefore, this result suggests that inclusion of γ-2 into the GluA1/A2 receptor partly rescues the detrimental effect of the I627T mutation on receptor activation properties. Other TARP subtypes as well as non-TARP auxiliary subunits vary greatly in their influence on AMPAR function75; 86; 87. Further work examining functional effects of AMPAR variants in the context of different auxiliary subunits may therefore be warranted.

For the A636T mutation, the effects observed in homomeric GluA1 were maintained in heteromeric GluA1/A2R overall; these showed significantly increased Glu-evoked currents, decreased desensitization, decreased Glu EC50, and increased KA efficacy (Fig. 4A, C and D; Table 4). This effect pattern was maintained upon γ-2 co-expression, except for the KA efficacy found not to be significantly different from WT (Fig. 4D). However, as previously mentioned, γ-2 in general enhances activation, and it might be speculated that this masks any effect of the A636T mutation on activation in heteromeric GluA1/A2 receptors with γ-2.

The G745D mutation that yielded homomeric GluA1 receptors that are completely inactive, displayed desensitized and non-desensitized Glu-evoked currents when co-expressed with GluA2R (Fig. 4A); however, the current amplitudes were 6-fold and 19-fold decreased, respectively, compared to the WT currents (Fig. 4C; Table 4). Surprisingly, analysis of the CTZ potentiation ratio and KA/Glu efficacy ratio showed G745D to have decreased desensitization and increased KA/Glu efficacy ratio, which would suggest steady-state currents to be increased (Fig. 4D; Table 4). The EC50 value for Glu was not changed (Supplementary Fig. S2). These effects of G745D were maintained when co-expressed with γ-2, except for the CTZ potentiation ratio, which was not different from WT GluA1/A2R with γ-2 (Fig. 4D; Table 4). These results show that the G745D changes functional properties of the GluA1 subunit to overall lower heteromeric GluA1/A2R receptor currents; however, they do not unequivocally reveal how the mutation disrupts function. Finally, the R345Q mutant displayed similar properties to WT GluA1/A2R with and without γ-2 with comparable current amplitudes, desensitization, and KA/Glu ratios that were not different from WT (Fig. 4D; Table 4).

In summary, the electrophysiological evaluation of the effect of the *GRIA1* variants on AMPAR function confirmed the prediction that the stop-gain variant p.Arg345Ter completely prevents the formation of functional GluA1-containing AMPARs. Notably, this finding supports the results from the *Xenopus* tadpole behavioral experiments that showed the knockout and the exon 8 deletion crispant tadpoles to have indistinguishable phenotypes. Together, this suggests that the homozygous p.Arg345Ter variant leads to complete loss of GluA1 in patient 1. For the missense variants in patients 2 to 7, the electrophysiological results strongly suggest that the p.Ile627Thr and p.Gly745Asp variants overall lead to severe loss-of-function phenotypes for homomeric GluA1 receptors and significantly decrease heteromeric GluA1-containing receptor function. In contrast, the p.Ala636Thr variant produces a clear gain-of-function phenotype in homomeric and heteromeric receptors characterized by loss of desensitization and increased Glu sensitivity. The p.Arg345Gln variant was found not to change functional parameters of both homomeric and heteromeric receptors, and therefore may be considered benign in terms of function.

Structural modeling of GluA1 shows that variants affect key receptor domains

Based on X-ray and cryo-EM structures representing the main functional states of the AMPARs, the structural mechanisms that underlie receptor function is becoming increasingly well-understood67-69; 88-90. Overall, the mechanism can be described by a four-state mechanistic model as illustrated in Fig. 5A. The Ile627, Ala636, and Gly745 residues where the *GRIA1* variants lead to altered receptor function are located in regions of the receptor structure that hold critical roles in this model. Specifically, Ile627 and Ala636 are located in the upper part of the M3 helix, which lines the ion channel and contains the channel gate (Fig. 5A). Asp745 is located in the clam-shell shaped ABD containing the Glu binding site and which undergoes the initial conformational changes leading to channel opening and receptor desensitization (Fig. 5A). To understand how mutation of these residues can influence the stability of the key receptor states, the structural role of these residues was analyzed using homology models of homomeric GluA1 created from AMPAR structures that represent the resting, active, and desensitized receptor states (Supplementary methods) (Fig. 5B-D).

Gly745 is located in the center of a short beta-strand that acts as a hinge between the upper and lower domains (denoted D1 and D2) that form the clamshell-shaped ABD (Fig. 5A–B). Upon agonist binding, D1 and D2 close around the agonist, leading to a pre-active, closed state from which the receptor can transition to an active, open-channel state or a desensitized, closed-channel state (Fig. 5A). In general, glycine adds flexibility to the peptide backbone that is lost upon substitution with any residue. Therefore, it can be speculated that the G745D mutation changes hinge properties to destabilize the closed-cleft ABD conformation in the pre-active, active, and desensitized states. Also, in the tetrameric AMPAR complex, the four ABDs are arranged pairwise into two identical dimers. In each dimer, the two ABDs are in a back-to-back orientation with the agonist binding clefts facing outwards (Fig. 5A and B). Gly745 is part of the dimer interface that is formed mainly between the D1 subdomains (Fig. 5B). In the resting and active state GluA1 models, the side chain introduced by any mutation of Gly745 will face into a hydrophobic motif formed by side chains of Ile495, Pro508, and Leu765 that contributes to the D1/D1 interface (Fig. 5B, *lower panels*). The G745D mutation introduces an acidic side chain towards this motif, thereby possibly destabilizing the interface (Fig. 5B). Notably, the strength of the D1/D1 interface is important for the stability of the pre-active and active states, and perturbation is known to be determinant for the equilibrium between receptor states67; 91-93(Fig. 5A). Specifically, mutations that destabilize the interface promote entry into the desensitized, closed-channel state and prevent entry into the active, open-channel state94. Therefore, the loss-of-function effects of the G745D mutation may also be caused by the destabilization of the D1/D1 interface. Notably, in heteromeric receptors such as GluA1/A2R receptors, the ABD dimers form between GluA1 and GluA2R subunits. Thus, the ABD interface in both dimers will be affected by the GluA1-G745D mutation, explaining the dominant effect of the mutation in heteromeric GluA1/A2R receptors.

Ile627 and Ala636 are located in the transmembrane M3 helix that forms the ion channel pore and the channel gate (Fig. 5C-D). Ala636 is at the extracellular-facing tip of M3 that contains the channel gate and undergoes conformational changes during channel opening and closing68; 69; 90. The GluA1 models show Ala636 in the four subunits to be arranged in nearly identical conformations in the resting and desensitized states, where the channel gate is closed (Fig. 5D). In these states, the Ala636 resdiues pack closely against the side chains of Leu638 and Thr639 on the adjacent subunit and form a hydrophobic interaction network that likely contributes to the stability of the closed-channel states (*lower panels*, Fig. 5D). The substitution of alanine with threonine introduces additional bulk and polarity into this network which is predicted to destabilize the closed-channel configuration of the upper M3. In contrast, in the active, open-channel model, the tips of the M3 helices have moved away from the channel center axis. As a result, the four Ala636 side chains do not interact with any channel residues, facing a direction allowing the extra methyl and hydroxyl groups introduced by the A636T mutation (Fig. 5D). Therefore, the models suggest that A636T selectively destabilize the closed-channel conformations observed in the resting and non-desensitized states, thereby shifting receptor equilibrium towards the active state. This analysis is in good agreement with the functional results that show the A636T mutant to have enhanced activation and decreased desensitization (Fig. 5A).

Ile637 is located down the M3 helix below the channel gate and above the selectivity filter at the tip of the M2 re-entry loop. In the resting and desensitized states, the models show the isoleucine side chain projecting towards the channel center (indicated with a blue arrow in the top panel, Fig. 5C) and not forming any interactions with other residues. In these conformations, the substitution of isoleucine with threonine, which has a similar-sized, but polar side chain, can be expected to be tolerated without changing the stability of the closed-channel states. In contrast, in the open-channel conformation of the active state, the Ile627 side chain is close to Phe598 and Gln600 at the tip of the re-entry M2 helix and forms hydrophobic interactions that stabilize the open-channel state. The polar side chain of threonine may disrupt these interactions to destabilize the open-channel conformation and explain the effect of decreased activation of the I627T mutation (Fig. 5A). In summary, except for p.Arg345Gln, the *GRIA1* variants implicated in NDDs affect residues that are positioned in critical structural domains in the GluA1 subunit protein, and analysis of the potential structural effect of the variants agree well with their observed functional effects to further substantiate their pathogenic status.

**Discussion**

The *GRIA1-4* genes are emerging as candidate disease-causing genes in NDDs; particularly in forms with severe intellectual disability (ID), but also in autism spectrum disorder (ASD) and attention-deficit disorder (ADD). Indeed, multiple studies employing *GRIA* targeted or whole-exome sequencing (WES) of individual NDD patients or cohorts have reported variants in all four *GRIA* genes as potential or verified pathogenic13; 15; 62; 95-102. These include studies utilizing electrophysiological and biochemical analysis of potential variant effects on the function or expression of recombinant AMPARs containing the subunit variant. Functionally validated variants are so far best described for *GRIA2* and *GRIA3* for which more than 20 missense, insertion/deletion (indel), or stop-gain variants have been reported to change normal receptor function or disrupt or truncate subunit structure; strongly suggesting a linkage between specific *GRIA2* and *GRIA3* variants and NDD phenotypes7; 62; 97; 102; 103. For example, Salpietro *et al.*103 performed functional evaluation of 11 *GRIA2* variants identified in NDD patients with severe ID and found the majority to impact function or expression of GluA2-containing AMPAR subtypes. Similar evaluation of five *GRIA3* missense variants identified in a cohort of 400 unrelated males with X-linked mental retardation (XLMR) found all to drastically alter or destroy the function of GluA3-containing AMPARs97. *GRIA4* missense variants have also been associated with NDD phenotypes with severe ID95, but functional evaluation has not yet established whether these *GRIA4* variants change the function of GluA4-containing AMPAR subtypes.

In contrast to *GRIA2*, *GRIA3,* and *GRIA4*, the identity of *GRIA1* as an NDD-causing gene has yet to be established, with only three *GRIA1* missense variants so far reported in NDD patients (p.Ile627Thr, p.Ala636Thr, and p.Gly745Asp)13-15. Interestingly, the p.Ala636Thr variant has been recurrently identified in six unrelated patients via WES or targeted *GRIA1* sequencing in two large NDD cohorts13; 15. However, none of these *GRIA1* variants reported in the literature have been functionally evaluated for the potential impact on the expression and function of GluA1-containing AMPARs and, therefore, the pathogenic significance of *GRIA1* in NDDs has remained unclear. Consequently, the diagnostic interpretation and reporting of *GRIA1* variants are at present challenging, with *GRIA1* variants classified as variants of uncertain significance as per the Association for Clinical Genomic Science (ACGS) guidelines for variant classification104. Furthermore, *GRIA1* is not part of most commercial or custom-made gene panels used for genetic diagnosis of NDD patients and is not included in most clinical genetic knowledgebases such as, for example, the widely used UK-based PanelApp web resource, where virtual gene panels related to human disease are curated. Together, these uncertainties regarding the pathophysiological role of *GRIA1* variants have limited patient diagnosis and potential AMPAR-targeted drug treatment options.

In the present work, a phenotype has been identified in NDD patients with *GRIA1* variants that includes ID, speech and language delay, poor sleep, abnormal electroencephalogram (EEG) with or without seizures, normal brain imaging, and endocrine abnormalities. Additionally, only missense heterozygous variants have been reported within the current literature, with our index case as the first homozygous nonsense variant in *GRIA1* causing a neurodevelopmental delay phenotype. Functional evaluation of the *GRIA1* variants identified in the patients was performed using electrophysiological and biochemical analyses, which characterized variant-induced changes in receptor function and expression. Crucially, three of the four missense variants caused significant changes in the function of homomeric GluA1 and heteromeric GluA1/A2 subtypes, whether this was in the current response amplitudes, degree of desensitization, or receptor activation. The most pronounced abnormalities were demonstrated by the p.Gly745Asp variant, which had a minimal current response, and the homozygous nonsense variant (p.Arg377ter), which demonstrated no current response to Glu. In addition to the lack of current response, shown in the p.Arg377ter variant, there was no cell surface expression. These findings support the notion that homozygous nonsense variants result in no functioning *GRIA1* gene, leading to increased severity in phenotype, and is internally consistent with the results of mutants in the *Xenopus* model (below). Interestingly, the p.Ala636 variant showed an increased current response which was thought to be secondary to increased sensitivity to glutamate and a loss of desensitization. Our findings demonstrate that *GRIA1* contributes to two syndromes with an autosomal recessive and autosomal dominant inheritance pattern and increased severity demonstrated by early-onset seizures and profound speech and language delay in the autosomal recessive index case.

Fundamental insight into the physiological role of GluA1 has come from gene deletion studies in mice where knockout models of the GluA1-encoding gene *Gria1* have been established and report normal development and life expectancy in *Gria1-/-* animals54; 105-109. However, behavioral inconsistencies are often reported in studies employing the *Gria1-/-* model, including hyperactivity54; 106-110, impaired spatial working memory51-54, and abnormalities in prepulse inhibition111 and sleep EEG in keeping with those found in schizophrenia112. Here, *GRIA1* gene function analysis is extended to a second model utilizing a CRISPR-based loss-of-function analysis in crispant *X. tropicalis* tadpoles and used to test explicitly the genotype-phenotype link caused by truncation of *GRIA1*. *Xenopus* have an extensive track record for cost-effective, high-throughput gene function analysis113-117 and high evolutionary similarity to mammals, but broadly lack robust assays to measure higher executive functions. To date, most behavioral studies in *Xenopus* have focused on understanding behavior in the wild, with some reports detailing laboratory schooling118; 119, swim and search patterns120-123, color differentiation124-126, seizure induction47; 127, and learned behaviours126; 128. In contrast, sophisticated quantitative behavioral analysis has so far been limited in tadpoles. Against this background, the present work demonstrates the successful adaptation of an established method from other vertebrate models with high translational relevance to humans30 to test the working memory of *gria1* mutant tadpoles. The initial characterization of the nonsense variant using *Xenopus* tadpoles was undertaken by creating homozygous indels within exon 8 and a separate *gria1* knockout model. Our model demonstrates the quantitative behavioral analysis of higher cognitive functions in *Xenopus* tadpoles for the first time, using a test with the potential for future direct comparison between the animal model and the patient cohort. Homozygous indels created within exon 8 to mimic the homozygous nonsense variant (p.Arg377ter) functionally support the description of an NDD phenotype by showing working memory deficits without detectable structural changes to the brain. Overall, these findings are in keeping with those reported in the *gria1*-/- mouse, providing a second, cost-effective model organism to investigate further the functional role of GluA1-containing AMPARs in the brain.

In summary, this study establishes *GRIA1* as a human NDD causing gene that merits being part of the existing collection of *GRIA*-related NDDs.

### **Supplemental Data**

Supplemental Data include seven figures and one tables and a supplementary methods section*.*

### **Declaration of Interests**

*The authors declare no competing interests.*

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### **Web Resources**

GeneMatcher: https://genematcher.org

OMIM: <https://www.omim.org>

Decipher: https://decipher.sanger.ac.uk

ClinVar: https://www.ncbi.nlm.nih.gov/clinvar

gnomAD: https://gnomad.broadinstitute.org

Addgene: <http://n2t.net/addgene>

Xenbase: http://www.xenbase.org/entry/

PanelApp: <https://panelapp.genomicsengland.co.uk>

MTR-viewer: http://biosig.unimelb.edu.au/mtr-viewer/

**Data and Code Availability**

This study did not generate datasets or code.

**References**

1. Traynelis, S.F., Wollmuth, L.P., McBain, C.J., Menniti, F.S., Vance, K.M., Ogden, K.K., Hansen, K.B., Yuan, H., Myers, S.J., and Dingledine, R. (2010). Glutamate receptor ion channels: structure, regulation, and function. Pharmacological reviews 62, 405-496.

2. Bliss, T.V., and Collingridge, G.L. (2013). Expression of NMDA receptor-dependent LTP in the hippocampus: bridging the divide. Molecular brain 6, 5.

3. Huganir, R.L., and Nicoll, R.A. (2013). AMPARs and synaptic plasticity: the last 25 years. Neuron 80, 704-717.

4. Malinow, R., and Malenka, R.C. (2002). AMPA receptor trafficking and synaptic plasticity. Annual review of neuroscience 25, 103-126.

5. Hollmann, M., and Heinemann, S. (1994). Cloned glutamate receptors. Annual review of neuroscience 17, 31-108.

6. Leonard, H., and Wen, X. (2002). The epidemiology of mental retardation: challenges and opportunities in the new millennium. Mental retardation and developmental disabilities research reviews 8, 117-134.

7. Chelly, J., and Mandel, J.L. (2001). Monogenic causes of X-linked mental retardation. Nature reviews Genetics 2, 669-680.

8. Ropers, H.H. (2010). Genetics of early onset cognitive impairment. Annual review of genomics and human genetics 11, 161-187.

9. Sheridan, E., Wright, J., Small, N., Corry, P.C., Oddie, S., Whibley, C., Petherick, E.S., Malik, T., Pawson, N., McKinney, P.A., et al. (2013). Risk factors for congenital anomaly in a multiethnic birth cohort: an analysis of the Born in Bradford study. Lancet 382, 1350-1359.

10. Hamdan, F.F., Gauthier, J., Araki, Y., Lin, D.T., Yoshizawa, Y., Higashi, K., Park, A.R., Spiegelman, D., Dobrzeniecka, S., Piton, A., et al. (2011). Excess of de novo deleterious mutations in genes associated with glutamatergic systems in nonsyndromic intellectual disability. American journal of human genetics 88, 306-316.

11. Kaufman, L., Ayub, M., and Vincent, J.B. (2010). The genetic basis of non-syndromic intellectual disability: a review. Journal of neurodevelopmental disorders 2, 182-209.

12. Yuan, H., Low, C.M., Moody, O.A., Jenkins, A., and Traynelis, S.F. (2015). Ionotropic GABA and Glutamate Receptor Mutations and Human Neurologic Diseases. Molecular pharmacology 88, 203-217.

13. de Ligt, J., Willemsen, M.H., van Bon, B.W., Kleefstra, T., Yntema, H.G., Kroes, T., Vulto-van Silfhout, A.T., Koolen, D.A., de Vries, P., Gilissen, C., et al. (2012). Diagnostic exome sequencing in persons with severe intellectual disability. The New England journal of medicine 367, 1921-1929.

14. Guo, H., Duyzend, M.H., Coe, B.P., Baker, C., Hoekzema, K., Gerdts, J., Turner, T.N., Zody, M.C., Beighley, J.S., Murali, S.C., et al. (2019). Genome sequencing identifies multiple deleterious variants in autism patients with more severe phenotypes. Genetics in medicine : official journal of the American College of Medical Genetics 21, 1611-1620.

15. Geisheker, M.R., Heymann, G., Wang, T., Coe, B.P., Turner, T.N., Stessman, H.A.F., Hoekzema, K., Kvarnung, M., Shaw, M., Friend, K., et al. (2017). Hotspots of missense mutation identify neurodevelopmental disorder genes and functional domains. Nat Neurosci 20, 1043-1051.

16. Turner, T.N., Hormozdiari, F., Duyzend, M.H., McClymont, S.A., Hook, P.W., Iossifov, I., Raja, A., Baker, C., Hoekzema, K., Stessman, H.A., et al. (2016). Genome Sequencing of Autism-Affected Families Reveals Disruption of Putative Noncoding Regulatory DNA. American journal of human genetics 98, 58-74.

17. Sobreira, N., Schiettecatte, F., Valle, D., and Hamosh, A. (2015). GeneMatcher: a matching tool for connecting investigators with an interest in the same gene. Hum Mutat 36, 928-930.

18. Landrum, M.J., Lee, J.M., Benson, M., Brown, G.R., Chao, C., Chitipiralla, S., Gu, B., Hart, J., Hoffman, D., Jang, W., et al. (2018). ClinVar: improving access to variant interpretations and supporting evidence. Nucleic Acids Res 46, D1062-D1067.

19. Rentzsch, P., Witten, D., Cooper, G.M., Shendure, J., and Kircher, M. (2019). CADD: predicting the deleteriousness of variants throughout the human genome. Nucleic Acids Res 47, D886-D894.

20. Ng, P.C., and Henikoff, S. (2003). SIFT: Predicting amino acid changes that affect protein function. Nucleic Acids Res 31, 3812-3814.

21. Adzhubei, I.A., Schmidt, S., Peshkin, L., Ramensky, V.E., Gerasimova, A., Bork, P., Kondrashov, A.S., and Sunyaev, S.R. (2010). A method and server for predicting damaging missense mutations. Nat Methods 7, 248-249.

22. Sargent, M.G., and Mohun, T.J. (2005). Cryopreservation of sperm of Xenopus laevis and Xenopus tropicalis. Genesis 41, 41-46.

23. Mansour, N., Lahnsteiner, F., and Patzner, R.A. (2009). Optimization of the cryopreservation of African clawed frog (Xenopus laevis) sperm. Theriogenology 72, 1221-1228.

24. Xenopus Online Resource | Xenopus Resource Centre (EXRC). In. (

25. Karimi, K., Fortriede, J.D., Lotay, V.S., Burns, K.A., Wang, D.Z., Fisher, M.E., Pells, T.J., James-Zorn, C., Wang, Y., Ponferrada, V.G., et al. (2018). Xenbase: a genomic, epigenomic and transcriptomic model organism database. Nucleic Acids Res 46, D861-D868.

26. Moreno-Mateos, M.A., Vejnar, C.E., Beaudoin, J.D., Fernandez, J.P., Mis, E.K., Khokha, M.K., and Giraldez, A.J. (2015). CRISPRscan: designing highly efficient sgRNAs for CRISPR-Cas9 targeting in vivo. Nat Methods 12, 982-988.

27. Shen, M.W., Arbab, M., Hsu, J.Y., Worstell, D., Culbertson, S.J., Krabbe, O., Cassa, C.A., Liu, D.R., Gifford, D.K., and Sherwood, R.I. (2018). Predictable and precise template-free CRISPR editing of pathogenic variants. Nature 563, 646-651.

28. Nakayama, T., Blitz, I.L., Fish, M.B., Odeleye, A.O., Manohar, S., Cho, K.W., and Grainger, R.M. (2014). Cas9-based genome editing in Xenopus tropicalis. Methods Enzymol 546, 355-375.

29. Untergasser, A., Cutcutache, I., Koressaar, T., Ye, J., Faircloth, B.C., Remm, M., and Rozen, S.G. (2012). Primer3--new capabilities and interfaces. Nucleic Acids Res 40, e115.

30. Cleal, M., Fontana, B.D., Ranson, D.C., McBride, S.D., Swinny, J.D., Redhead, E.S., and Parker, M.O. (2021). The Free-movement pattern Y-maze: A cross-species measure of working memory and executive function. Behav Res Methods 53, 536-557.

31. Fontana, B.D., Cleal, M., Clay, J.M., and Parker, M.O. (2019). Zebrafish (Danio rerio) behavioral laterality predicts increased short-term avoidance memory but not stress-reactivity responses. Anim Cogn 22, 1051-1061.

32. Fontana, B.D., Cleal, M., Gibbon, A.J., McBride, S.D., and Parker, M.O. (2021). The effects of two stressors on working memory and cognitive flexibility in zebrafish (Danio rerio): The protective role of D1/D5 agonist on stress responses. Neuropharmacology 196, 108681.

33. Faul, F., Erdfelder, E., Lang, A.G., and Buchner, A. (2007). G\*Power 3: a flexible statistical power analysis program for the social, behavioral, and biomedical sciences. Behav Res Methods 39, 175-191.

34. Stenum-Berg, C., Musgaard, M., Chavez-Abiega, S., Thisted, C.L., Barrella, L., Biggin, P.C., and Kristensen, A.S. (2019). Mutational Analysis and Modeling of Negative Allosteric Modulator Binding Sites in AMPA Receptors. Molecular pharmacology 96, 835-850.

35. Hayashi, Y., Shi, S.H., Esteban, J.A., Piccini, A., Poncer, J.C., and Malinow, R. (2000). Driving AMPA receptors into synapses by LTP and CaMKII: requirement for GluR1 and PDZ domain interaction. Science 287, 2262-2267.

36. Lu, P., Bai, X.C., Ma, D., Xie, T., Yan, C., Sun, L., Yang, G., Zhao, Y., Zhou, R., Scheres, S.H.W., et al. (2014). Three-dimensional structure of human gamma-secretase. Nature 512, 166-170.

37. Poulsen, M.H., Lucas, S., Stromgaard, K., and Kristensen, A.S. (2014). Evaluation of PhTX-74 as subtype-selective inhibitor of GluA2-containing AMPA receptors. Molecular pharmacology 85, 261-268.

38. Lam, V.M., Beerepoot, P., Angers, S., and Salahpour, A. (2013). A novel assay for measurement of membrane-protein surface expression using a beta-lactamase. Traffic 14, 778-784.

39. Joshi, P.R., Suryanarayanan, A., and Schulte, M.K. (2004). A vertical flow chamber for Xenopus oocyte electrophysiology and automated drug screening. J Neurosci Methods 132, 69-79.

40. Traynelis, J., Silk, M., Wang, Q., Berkovic, S.F., Liu, L., Ascher, D.B., Balding, D.J., and Petrovski, S. (2017). Optimizing genomic medicine in epilepsy through a gene-customized approach to missense variant interpretation. Genome Res 27, 1715-1729.

41. Li, J., Zhang, J., Tang, W., Mizu, R.K., Kusumoto, H., XiangWei, W., Xu, Y., Chen, W., Amin, J.B., Hu, C., et al. (2019). De novo GRIN variants in NMDA receptor M2 channel pore-forming loop are associated with neurological diseases. Hum Mutat 40, 2393-2413.

42. Swanger, S.A., Chen, W., Wells, G., Burger, P.B., Tankovic, A., Bhattacharya, S., Strong, K.L., Hu, C., Kusumoto, H., Zhang, J., et al. (2016). Mechanistic Insight into NMDA Receptor Dysregulation by Rare Variants in the GluN2A and GluN2B Agonist Binding Domains. American journal of human genetics 99, 1261-1280.

43. Barbosa, S., Greville-Heygate, S., Bonnet, M., Godwin, A., Fagotto-Kaufmann, C., Kajava, A.V., Laouteouet, D., Mawby, R., Wai, H.A., Dingemans, A.J.M., et al. (2020). Opposite Modulation of RAC1 by Mutations in TRIO Is Associated with Distinct, Domain-Specific Neurodevelopmental Disorders. American journal of human genetics 106, 338-355.

44. Macken, W.L., Godwin, A., Wheway, G., Stals, K., Nazlamova, L., Ellard, S., Alfares, A., Aloraini, T., AlSubaie, L., Alfadhel, M., et al. (2021). Biallelic variants in COPB1 cause a novel, severe intellectual disability syndrome with cataracts and variable microcephaly. Genome Med 13, 34.

45. Deniz, E., Mis, E.K., Lane, M., and Khokha, M.K. (2018). CRISPR/Cas9 F0 Screening of Congenital Heart Disease Genes in Xenopus tropicalis. Methods Mol Biol 1865, 163-174.

46. Sega, A.G., Mis, E.K., Lindstrom, K., Mercimek-Andrews, S., Ji, W., Cho, M.T., Juusola, J., Konstantino, M., Jeffries, L., Khokha, M.K., et al. (2019). De novo pathogenic variants in neuronal differentiation factor 2 (NEUROD2) cause a form of early infantile epileptic encephalopathy. J Med Genet 56, 113-122.

47. Hewapathirane, D.S., Dunfield, D., Yen, W., Chen, S., and Haas, K. (2008). In vivo imaging of seizure activity in a novel developmental seizure model. Exp Neurol 211, 480-488.

48. Pratt, K.G., and Khakhalin, A.S. (2013). Modeling human neurodevelopmental disorders in the Xenopus tadpole: from mechanisms to therapeutic targets. Dis Model Mech 6, 1057-1065.

49. Shapiro, M.L., and O'Connor, C. (1992). N-methyl-D-aspartate receptor antagonist MK-801 and spatial memory representation: working memory is impaired in an unfamiliar environment but not in a familiar environment. Behav Neurosci 106, 604-612.

50. Brosnan-Watters, G., Wozniak, D.F., Nardi, A., and Olney, J.W. (1996). Acute behavioral effects of MK-801 in the mouse. Pharmacol Biochem Behav 53, 701-711.

51. Sanderson, D.J., Good, M.A., Skelton, K., Sprengel, R., Seeburg, P.H., Rawlins, J.N., and Bannerman, D.M. (2009). Enhanced long-term and impaired short-term spatial memory in GluA1 AMPA receptor subunit knockout mice: evidence for a dual-process memory model. Learn Mem 16, 379-386.

52. Reisel, D., Bannerman, D.M., Schmitt, W.B., Deacon, R.M., Flint, J., Borchardt, T., Seeburg, P.H., and Rawlins, J.N. (2002). Spatial memory dissociations in mice lacking GluR1. Nat Neurosci 5, 868-873.

53. Schmitt, W.B., Deacon, R.M., Seeburg, P.H., Rawlins, J.N., and Bannerman, D.M. (2003). A within-subjects, within-task demonstration of intact spatial reference memory and impaired spatial working memory in glutamate receptor-A-deficient mice. J Neurosci 23, 3953-3959.

54. Zamanillo, D., Sprengel, R., Hvalby, O., Jensen, V., Burnashev, N., Rozov, A., Kaiser, K.M., Koster, H.J., Borchardt, T., Worley, P., et al. (1999). Importance of AMPA receptors for hippocampal synaptic plasticity but not for spatial learning. Science 284, 1805-1811.

55. Sanderson, D.J., Sprengel, R., Seeburg, P.H., and Bannerman, D.M. (2011). Deletion of the GluA1 AMPA receptor subunit alters the expression of short-term memory. Learn Mem 18, 128-131.

56. Greger, I.H., Watson, J.F., and Cull-Candy, S.G. (2017). Structural and Functional Architecture of AMPA-Type Glutamate Receptors and Their Auxiliary Proteins. Neuron 94, 713-730.

57. Herguedas, B., Garcia-Nafria, J., Cais, O., Fernandez-Leiro, R., Krieger, J., Ho, H., and Greger, I.H. (2016). Structure and organization of heteromeric AMPA-type glutamate receptors. Science 352, aad3873.

58. Herguedas, B., Krieger, J., and Greger, I.H. (2013). Receptor heteromeric assembly-how it works and why it matters: the case of ionotropic glutamate receptors. Progress in molecular biology and translational science 117, 361-386.

59. Sommer, B., Keinanen, K., Verdoorn, T.A., Wisden, W., Burnashev, N., Herb, A., Kohler, M., Takagi, T., Sakmann, B., and Seeburg, P.H. (1990). Flip and flop: a cell-specific functional switch in glutamate-operated channels of the CNS. Science 249, 1580-1585.

60. Plested, A.J., Wildman, S.S., Lieb, W.R., and Franks, N.P. (2004). Determinants of the sensitivity of AMPA receptors to xenon. Anesthesiology 100, 347-358.

61. Robert, A., Irizarry, S.N., Hughes, T.E., and Howe, J.R. (2001). Subunit interactions and AMPA receptor desensitization. J Neurosci 21, 5574-5586.

62. Davies, B., Brown, L.A., Cais, O., Watson, J., Clayton, A.J., Chang, V.T., Biggs, D., Preece, C., Hernandez-Pliego, P., Krohn, J., et al. (2017). A point mutation in the ion conduction pore of AMPA receptor GRIA3 causes dramatically perturbed sleep patterns as well as intellectual disability. Human molecular genetics 26, 3869-3882.

63. Schmid, S.M., Korber, C., Herrmann, S., Werner, M., and Hollmann, M. (2007). A domain linking the AMPA receptor agonist binding site to the ion pore controls gating and causes lurcher properties when mutated. J Neurosci 27, 12230-12241.

64. Klein, R.M., and Howe, J.R. (2004). Effects of the lurcher mutation on GluR1 desensitization and activation kinetics. J Neurosci 24, 4941-4951.

65. Taverna, F., Xiong, Z.G., Brandes, L., Roder, J.C., Salter, M.W., and MacDonald, J.F. (2000). The Lurcher mutation of an alpha-amino-3-hydroxy-5-methyl- 4-isoxazolepropionic acid receptor subunit enhances potency of glutamate and converts an antagonist to an agonist. J Biol Chem 275, 8475-8479.

66. Kohda, K., Wang, Y., and Yuzaki, M. (2000). Mutation of a glutamate receptor motif reveals its role in gating and delta2 receptor channel properties. Nat Neurosci 3, 315-322.

67. Twomey, E.C., and Sobolevsky, A.I. (2018). Structural Mechanisms of Gating in Ionotropic Glutamate Receptors. Biochemistry 57, 267-276.

68. Twomey, E.C., Yelshanskaya, M.V., Grassucci, R.A., Frank, J., and Sobolevsky, A.I. (2017). Channel opening and gating mechanism in AMPA-subtype glutamate receptors. Nature 549, 60-65.

69. Chen, S., Zhao, Y., Wang, Y., Shekhar, M., Tajkhorshid, E., and Gouaux, E. (2017). Activation and Desensitization Mechanism of AMPA Receptor-TARP Complex by Cryo-EM. Cell 170, 1234-1246 e1214.

70. Phillips, R.J.S. (1960). ‘Lurcher’, a new gene in linkage group XI of the house mouse. Journal of Genetics 57, 35.

71. Zuo, J., De Jager, P.L., Takahashi, K.A., Jiang, W., Linden, D.J., and Heintz, N. (1997). Neurodegeneration in Lurcher mice caused by mutation in delta2 glutamate receptor gene. Nature 388, 769-773.

72. Ge, Y., and Wang, Y.T. (2021). GluA1-homomeric AMPA receptor in synaptic plasticity and neurological diseases. Neuropharmacology 197, 108708.

73. Schwenk, J., Baehrens, D., Haupt, A., Bildl, W., Boudkkazi, S., Roeper, J., Fakler, B., and Schulte, U. (2014). Regional diversity and developmental dynamics of the AMPA-receptor proteome in the mammalian brain. Neuron 84, 41-54.

74. Schwenk, J., Harmel, N., Brechet, A., Zolles, G., Berkefeld, H., Muller, C.S., Bildl, W., Baehrens, D., Huber, B., Kulik, A., et al. (2012). High-resolution proteomics unravel architecture and molecular diversity of native AMPA receptor complexes. Neuron 74, 621-633.

75. Jackson, A.C., and Nicoll, R.A. (2011). The expanding social network of ionotropic glutamate receptors: TARPs and other transmembrane auxiliary subunits. Neuron 70, 178-199.

76. Kristensen, A.S., Jenkins, M.A., Banke, T.G., Schousboe, A., Makino, Y., Johnson, R.C., Huganir, R., and Traynelis, S.F. (2011). Mechanism of Ca2+/calmodulin-dependent kinase II regulation of AMPA receptor gating. Nat Neurosci 14, 727-735.

77. Greger, I.H., Khatri, L., and Ziff, E.B. (2002). RNA editing at arg607 controls AMPA receptor exit from the endoplasmic reticulum. Neuron 34, 759-772.

78. Swanson, G.T., Kamboj, S.K., and Cull-Candy, S.G. (1997). Single-channel properties of recombinant AMPA receptors depend on RNA editing, splice variation, and subunit composition. J Neurosci 17, 58-69.

79. Tomita, S., Adesnik, H., Sekiguchi, M., Zhang, W., Wada, K., Howe, J.R., Nicoll, R.A., and Bredt, D.S. (2005). Stargazin modulates AMPA receptor gating and trafficking by distinct domains. Nature 435, 1052-1058.

80. Korber, C., Werner, M., Kott, S., Ma, Z.L., and Hollmann, M. (2007). The transmembrane AMPA receptor regulatory protein gamma 4 is a more effective modulator of AMPA receptor function than stargazin (gamma 2). J Neurosci 27, 8442-8447.

81. Kott, S., Sager, C., Tapken, D., Werner, M., and Hollmann, M. (2009). Comparative analysis of the pharmacology of GluR1 in complex with transmembrane AMPA receptor regulatory proteins gamma2, gamma3, gamma4, and gamma8. Neuroscience 158, 78-88.

82. Kott, S., Werner, M., Korber, C., and Hollmann, M. (2007). Electrophysiological properties of AMPA receptors are differentially modulated depending on the associated member of the TARP family. J Neurosci 27, 3780-3789.

83. Sager, C., Tapken, D., Kott, S., and Hollmann, M. (2009). Functional modulation of AMPA receptors by transmembrane AMPA receptor regulatory proteins. Neuroscience 158, 45-54.

84. Priel, A., Kolleker, A., Ayalon, G., Gillor, M., Osten, P., and Stern-Bach, Y. (2005). Stargazin reduces desensitization and slows deactivation of the AMPA-type glutamate receptors. J Neurosci 25, 2682-2686.

85. Soto, D., Coombs, I.D., Gratacos-Batlle, E., Farrant, M., and Cull-Candy, S.G. (2014). Molecular mechanisms contributing to TARP regulation of channel conductance and polyamine block of calcium-permeable AMPA receptors. J Neurosci 34, 11673-11683.

86. Shelley, C., Farrant, M., and Cull-Candy, S.G. (2012). TARP-associated AMPA receptors display an increased maximum channel conductance and multiple kinetically distinct open states. J Physiol 590, 5723-5738.

87. Kamalova, A., and Nakagawa, T. (2021). AMPA receptor structure and auxiliary subunits. J Physiol 599, 453-469.

88. Twomey, E.C., Yelshanskaya, M.V., Grassucci, R.A., Frank, J., and Sobolevsky, A.I. (2017). Structural Bases of Desensitization in AMPA Receptor-Auxiliary Subunit Complexes. Neuron 94, 569-580 e565.

89. Chen, S., and Gouaux, E. (2019). Structure and mechanism of AMPA receptor - auxiliary protein complexes. Curr Opin Struct Biol 54, 104-111.

90. Durr, K.L., Chen, L., Stein, R.A., De Zorzi, R., Folea, I.M., Walz, T., McHaourab, H.S., and Gouaux, E. (2014). Structure and dynamics of AMPA receptor GluA2 in resting, pre-open, and desensitized states. Cell 158, 778-792.

91. Sun, Y., Olson, R., Horning, M., Armstrong, N., Mayer, M., and Gouaux, E. (2002). Mechanism of glutamate receptor desensitization. Nature 417, 245-253.

92. Armstrong, N., and Gouaux, E. (2000). Mechanisms for activation and antagonism of an AMPA-sensitive glutamate receptor: crystal structures of the GluR2 ligand binding core. Neuron 28, 165-181.

93. Mayer, M.L., Olson, R., and Gouaux, E. (2001). Mechanisms for ligand binding to GluR0 ion channels: crystal structures of the glutamate and serine complexes and a closed apo state. J Mol Biol 311, 815-836.

94. Horning, M.S., and Mayer, M.L. (2004). Regulation of AMPA receptor gating by ligand binding core dimers. Neuron 41, 379-388.

95. Martin, S., Chamberlin, A., Shinde, D.N., Hempel, M., Strom, T.M., Schreiber, A., Johannsen, J., Ousager, L.B., Larsen, M.J., Hansen, L.K., et al. (2017). De Novo Variants in GRIA4 Lead to Intellectual Disability with or without Seizures and Gait Abnormalities. American journal of human genetics 101, 1013-1020.

96. Cherot, E., Keren, B., Dubourg, C., Carre, W., Fradin, M., Lavillaureix, A., Afenjar, A., Burglen, L., Whalen, S., Charles, P., et al. (2018). Using medical exome sequencing to identify the causes of neurodevelopmental disorders: Experience of 2 clinical units and 216 patients. Clinical genetics 93, 567-576.

97. Wu, Y., Arai, A.C., Rumbaugh, G., Srivastava, A.K., Turner, G., Hayashi, T., Suzuki, E., Jiang, Y., Zhang, L., Rodriguez, J., et al. (2007). Mutations in ionotropic AMPA receptor 3 alter channel properties and are associated with moderate cognitive impairment in humans. Proc Natl Acad Sci U S A 104, 18163-18168.

98. Philips, A.K., Siren, A., Avela, K., Somer, M., Peippo, M., Ahvenainen, M., Doagu, F., Arvio, M., Kaariainen, H., Van Esch, H., et al. (2014). X-exome sequencing in Finnish families with intellectual disability--four novel mutations and two novel syndromic phenotypes. Orphanet journal of rare diseases 9, 49.

99. Allen, N.M., Conroy, J., Shahwan, A., Lynch, B., Correa, R.G., Pena, S.D., McCreary, D., Magalhaes, T.R., Ennis, S., Lynch, S.A., et al. (2016). Unexplained early onset epileptic encephalopathy: Exome screening and phenotype expansion. Epilepsia 57, e12-17.

100. Alkelai, A., Shohat, S., Greenbaum, L., Schechter, T., Draiman, B., Chitrit-Raveh, E., Rienstein, S., Dagaonkar, N., Hughes, D., Aggarwal, V.S., et al. (2021). Expansion of the GRIA2 phenotypic representation: a novel de novo loss of function mutation in a case with childhood onset schizophrenia. Journal of human genetics 66, 339-343.

101. Trivisano, M., Santarone, M.E., Micalizzi, A., Ferretti, A., Dentici, M.L., Novelli, A., Vigevano, F., and Specchio, N. (2020). GRIA3 missense mutation is cause of an x-linked developmental and epileptic encephalopathy. Seizure 82, 1-6.

102. Sun, J.H., Chen, J., Ayala Valenzuela, F.E., Brown, C., Masser-Frye, D., Jones, M., Romero, L.P., Rinaldi, B., Li, W.L., Li, Q.Q., et al. (2021). X-linked neonatal-onset epileptic encephalopathy associated with a gain-of-function variant p.R660T in GRIA3. PLoS Genet 17, e1009608.

103. Salpietro, V., Dixon, C.L., Guo, H., Bello, O.D., Vandrovcova, J., Efthymiou, S., Maroofian, R., Heimer, G., Burglen, L., Valence, S., et al. (2019). AMPA receptor GluA2 subunit defects are a cause of neurodevelopmental disorders. Nature communications 10, 3094.

104. Richards, S., Aziz, N., Bale, S., Bick, D., Das, S., Gastier-Foster, J., Grody, W.W., Hegde, M., Lyon, E., Spector, E., et al. (2015). Standards and guidelines for the interpretation of sequence variants: a joint consensus recommendation of the American College of Medical Genetics and Genomics and the Association for Molecular Pathology. Genetics in medicine : official journal of the American College of Medical Genetics 17, 405-424.

105. Procaccini, C., Aitta-aho, T., Jaako-Movits, K., Zharkovsky, A., Panhelainen, A., Sprengel, R., Linden, A.M., and Korpi, E.R. (2011). Excessive novelty-induced c-Fos expression and altered neurogenesis in the hippocampus of GluA1 knockout mice. Eur J Neurosci 33, 161-174.

106. Vekovischeva, O.Y., Zamanillo, D., Echenko, O., Seppala, T., Uusi-Oukari, M., Honkanen, A., Seeburg, P.H., Sprengel, R., and Korpi, E.R. (2001). Morphine-induced dependence and sensitization are altered in mice deficient in AMPA-type glutamate receptor-A subunits. J Neurosci 21, 4451-4459.

107. Chourbaji, S., Vogt, M.A., Fumagalli, F., Sohr, R., Frasca, A., Brandwein, C., Hortnagl, H., Riva, M.A., Sprengel, R., and Gass, P. (2008). AMPA receptor subunit 1 (GluR-A) knockout mice model the glutamate hypothesis of depression. FASEB J 22, 3129-3134.

108. Fitzgerald, P.J., Barkus, C., Feyder, M., Wiedholz, L.M., Chen, Y.C., Karlsson, R.M., Machado-Vieira, R., Graybeal, C., Sharp, T., Zarate, C., et al. (2010). Does gene deletion of AMPA GluA1 phenocopy features of schizoaffective disorder? Neurobiol Dis 40, 608-621.

109. Barkus, C., Feyder, M., Graybeal, C., Wright, T., Wiedholz, L., Izquierdo, A., Kiselycznyk, C., Schmitt, W., Sanderson, D.J., Rawlins, J.N., et al. (2012). Do GluA1 knockout mice exhibit behavioral abnormalities relevant to the negative or cognitive symptoms of schizophrenia and schizoaffective disorder? Neuropharmacology 62, 1263-1272.

110. Bannerman, D.M., Deacon, R.M., Brady, S., Bruce, A., Sprengel, R., Seeburg, P.H., and Rawlins, J.N. (2004). A comparison of GluR-A-deficient and wild-type mice on a test battery assessing sensorimotor, affective, and cognitive behaviors. Behav Neurosci 118, 643-647.

111. Wiedholz, L.M., Owens, W.A., Horton, R.E., Feyder, M., Karlsson, R.M., Hefner, K., Sprengel, R., Celikel, T., Daws, L.C., and Holmes, A. (2008). Mice lacking the AMPA GluR1 receptor exhibit striatal hyperdopaminergia and 'schizophrenia-related' behaviors. Molecular psychiatry 13, 631-640.

112. Ang, G., McKillop, L.E., Purple, R., Blanco-Duque, C., Peirson, S.N., Foster, R.G., Harrison, P.J., Sprengel, R., Davies, K.E., Oliver, P.L., et al. (2018). Absent sleep EEG spindle activity in GluA1 (Gria1) knockout mice: relevance to neuropsychiatric disorders. Transl Psychiatry 8, 154.

113. DeLay, B.D., Corkins, M.E., Hanania, H.L., Salanga, M., Deng, J.M., Sudou, N., Taira, M., Horb, M.E., and Miller, R.K. (2018). Tissue-Specific Gene Inactivation in Xenopus laevis: Knockout of lhx1 in the Kidney with CRISPR/Cas9. Genetics 208, 673-686.

114. Hwang, W.Y., Marquez, J., and Khokha, M.K. (2019). Xenopus: Driving the Discovery of Novel Genes in Patient Disease and Their Underlying Pathological Mechanisms Relevant for Organogenesis. Front Physiol 10, 953.

115. Kariminejad, A., Szenker-Ravi, E., Lekszas, C., Tajsharghi, H., Moslemi, A.R., Naert, T., Tran, H.T., Ahangari, F., Rajaei, M., Nasseri, M., et al. (2019). Homozygous Null TBX4 Mutations Lead to Posterior Amelia with Pelvic and Pulmonary Hypoplasia. American journal of human genetics 105, 1294-1301.

116. Naert, T., Colpaert, R., Van Nieuwenhuysen, T., Dimitrakopoulou, D., Leoen, J., Haustraete, J., Boel, A., Steyaert, W., Lepez, T., Deforce, D., et al. (2016). CRISPR/Cas9 mediated knockout of rb1 and rbl1 leads to rapid and penetrant retinoblastoma development in Xenopus tropicalis. Scientific reports 6, 35264.

117. Naert, T., and Vleminckx, K. (2018). CRISPR/Cas9 disease models in zebrafish and Xenopus: The genetic renaissance of fish and frogs. Drug Discov Today Technol 28, 41-52.

118. Katz, L.C., Potel, M.J., and Wassersug, R.J. (1981). Structure and mechanisms of schooling intadpoles of the clawed frog, Xenopus laevis. Animal Behaviour 29, 20-33.

119. Lopez, V., 3rd, Khakhalin, A.S., and Aizenman, C. (2021). Schooling in Xenopus laevis Tadpoles as a Way to Assess Their Neural Development. Cold Spring Harb Protoc 2021.

120. Truszkowski, T.L., James, E.J., Hasan, M., Wishard, T.J., Liu, Z., Pratt, K.G., Cline, H.T., and Aizenman, C.D. (2016). Fragile X mental retardation protein knockdown in the developing Xenopus tadpole optic tectum results in enhanced feedforward inhibition and behavioral deficits. Neural Dev 11, 14.

121. Roberts, A., Hill, N.A., and Hicks, R. (2000). Simple mechanisms organise orientation of escape swimming in embryos and hatchling tadpoles of Xenopus laevis. J Exp Biol 203, 1869-1885.

122. Hanzi, S., and Straka, H. (2018). Wall following in Xenopus laevis is barrier-driven. J Comp Physiol A Neuroethol Sens Neural Behav Physiol 204, 183-195.

123. Videlier, M., Cornette, R., Bonneaud, C., and Herrel, A. (2015). Sexual differences in exploration behavior in Xenopus tropicalis? J Exp Biol 218, 1733-1739.

124. Moriya, T., Kito, K., Miyashita, Y., and Asami, K. (1996). Preference for background color of the Xenopus laevis tadpole. J Exp Zool 276, 335-344.

125. Viczian, A.S., and Zuber, M.E. (2014). A simple behavioral assay for testing visual function in Xenopus laevis. J Vis Exp.

126. Blackiston, D.J., and Levin, M. (2012). Aversive training methods in Xenopus laevis: general principles. Cold Spring Harb Protoc 2012.

127. Bell, M.R., Belarde, J.A., Johnson, H.F., and Aizenman, C.D. (2011). A neuroprotective role for polyamines in a Xenopus tadpole model of epilepsy. Nat Neurosci 14, 505-512.

128. Blackiston, D.J., and Levin, M. (2013). Ectopic eyes outside the head in Xenopus tadpoles provide sensory data for light-mediated learning. J Exp Biol 216, 1031-1040.

**Figure legends**

**Figure 1. Location of residues in GluA1 affected by *GRIA1* variants and effect on receptor expression**.

**(A)** Cartoon representation of the structure and domain organization of the human GluA1 subunit protein (NP\_000817.1) encoded by the *GRIA1* gene. Residues affected by the GRIA1 variants evaluated in this study are shown in spheres (carbon and nitrogen atoms in purple and blue, respectively) and indicated with arrows. The GluA1 subunit structure was modelled from the structure of the rat GluA2 receptor (*Suplementary materials & methods*).

**(B)** Structure of the homotetrameric GluA1 AMPA receptor. The locations of the residues affected by the *GRIA1* missense variants are indicated by red spheres in the tetrameric receptor structure (*left*) and further shown in cross-sectional top views (*right*).

**(C)** Multiple alignment of the amino acid sequences that surround the residues affected by the *GRIA1* variants (highlighted in red) in GluA1 from human, rat, chicken, frog (*Xenopus*), and fish, and in human GluA2, GluA3, and GluA4 subunits. Residues with diverging physiochemical properties are shown in bold.

**(D)** Missense tolerance ratio (MTR)40 analysis of the population-level variation in the coding regions in *GRIA1* predict the tolerance of missense variation along the GluA1 primary structure to indicate residues or regions that are functionally sensitive to mutation. The plot was created using the MTR-Viewer online service. Horizontal lines show gene-specific MTR percentiles 5th, 25th, 50th, and neutrality (MTR = 1.0). Red domains in the plot indicate regions in the primary structure of GluA1 that are highly sensitive to missense mutation. The positions of the residues affected by the *GRIA1* variants in this study are shown as red circles. For reference, a linear representation of the GluA1 domain structure is shown above the plot.

**(E)** Biochemical assessment of WT and mutant GluA1 cell-surface expression. Summary of blac enzyme activity levels from the surface of oocytes expressing WT and mutant GluA1 receptors tagged in the N-terminal of the NTD for measurement of receptor cell surface level. The enzyme blac catalyzes hydrolysis of the β-lactam ring of the membrane-impermeable substrate nitrocefin to cause a color change from yellow to red, which is measured as increase in optical density (OD) at 486 nm. As nitrocefin does not permeate the cell membrane, only extracellular blac activity is measured. The data shown are the mean ± SEM of parallel measurements from 10 to 16 live oocytes expressing the indicated receptor subunits at 48 hours post RNA injection.

**(F)** Confocal imaging of HEK293 cells expressing the membrane reporter mCardinal-farnesyl (*left images*) and GFP-tagged WT and R377\* mutant GluA1 (*middle images*) to visualize cellular GluA1 distribution patterns. Co-localization of green and red fluorescence indicates cell surface localization of GFP-tagged GluA1 and is visualized by merging left and middle images (*right images*), thereby appearing yellow.

**(G)** Western blot analysis of WT and R377\* GluA1 protein expression in HEK293 cells using antibodies to the N – and C-termimi of the protein (methods).

**Figure 2. Tadpoles bearing CRISPR/Cas9 mediated insertion and deletion changes to exon 8, *gria1* (*Xtr.gria1*em2E*X*RC)** **have significant deficits in working memory.**

**(A)** Targeted disruption within *gria1* exon 8 generates a range of insertion-deletion changes *in vivo*. Sanger sequencing of eight subcloned genomic amplicons (exon 8) from *Xtr.gria1*em2E*X*RC tadpoles revealed a range of indels that occurred in samples at the CRISPR cut site (indicated by the blue line), including the re-occurring 1bp deletion and 3bp deletion. Further, four of the identified clones were found to cause a frameshift that truncated the protein (denoted by the asterisk).

**(B)** Representative micrograph images of the head region of uninjected control and *gria1* crispant tadpoles under bright-field conditions reveal no gross difference in craniofacial morphology.

**(C)** Representative bright-field (*left*) and fluorescence micrograph images (right, *in green*) of the head regions of transgenic [Xtr.Tg(tubb2b:GFP)Amaya, RRID: EXRC\_3001] tadpoles reveal gross morphology of the forebrain, midbrain, and hindbrain regions. Crispant (*Xtr.gria1*em2E*X*RC) tadpoles examined across three batches were generally indistinguishable from age-matched control tadpoles both in their craniofacial appearance and brain morphology. Specifically, brain length measured as the distance from the forebrain to the hindbrain (mean ± SD) was not significantly different between control and *gria1* crispant tadpoles (control tadpoles: 1.33 ± 0.1 mm, n=36; crispant tadpoles: 1.31 ± 0.1 mm, n=36, *p* = 0.537).

**(D-E)** The relative frequency distribution plots of the 1-hour global search strategy of wild-type (*black bars*) and *gria1* crispant (*blue bars*) tadpoles in the FMP Y-maze. Shown is the summative (D: *left,* mean ± SEM) and individual (D: *right*) tadpole performances from 60 wild-type and 60 *gria1* crispant tadpoles. The *gria1* crispant tadpoles were observed to perform significantly fewer alternations than stage-matched control animals (ANCOVA: F(2, 176) = 10.3, p<0.001 (n=60), D), and this difference in the overall proportion of alternations performed was observed throughout the trial (E).

**(F-G)** Overall, there was no significant difference in the number of turns performed by the uninjected control and *gria1* crispant tadpoles (t116.364 = -0.564, p=0.574), and all tadpoles were observed to perform fewer turns as the length of the trial increased (mean ± SEM).

**Figure 3. Functional characterization of homomeric mutant GluA1 receptors.**

**(A-B)** Representative steady-state currents evoked by sequential 10-20 s applications of Glu (1 mM, *red bar*) and Glu in the presence of CTZ (100 μM, *green bar*) from un-injected oocytes and oocytes expressing WT and mutant GluA1. The holding potential was -40 mV in all shown recordings. Note that the R377\*, I627T, and G745D mutants (B) show no or very small currents relative to WT, R345Q, and A636T.

**(C)** Scatter plot with bars of individual and mean amplitude of Glu-evoked currents in oocytes expressing WT and mutant GluA1 in the absence (*red bars and symbols*) and presence (*green bars and symbols*) of CTZ block of desensitization. Error bars indicate the 95% confidence interval of the mean amplitude. Note the semi-log y-axis. The stipulated line indicates the mean amplitude level for WT GluA1.

**(D)** Summary of fold desensitization for Glu-evoked currents calculated from the amplitudes of currents evoked by sequential application of Glu in the absence and presence of CTZ. Data represent the mean of 10 to 50 oocytes for each WT and mutant GluA1. Error bars indicate the 95% confidence interval of the means.

**(E)** Representative steady-state currents evoked by sequential applications (*black bars*) of increasing concentrations of Glu at oocytes expressing WT and the A636T mutant GluA1.

**(F)** Composite concentration-response curves for WT and mutant GluA1 homomeric receptors. Data points represent the mean of 10 to 31 oocytes. Error bars are the SEM and are shown when larger than symbol size. The current responses are normalized to the maximal response evoked by Glu (1 mM).

**(G)** Overlay of representative steady-state currents evoked by sequential applications of Glu (1 mM, *green bars*) and KA (300 uM, *purple bars*) in the presence of CTZ (100 uM) to show the efficacy of KA relative to Glu for evoking current. The KA currents are shown normalized to the Glu current.

**(H)** Summary of the average KA/Glu current response ratios for WT and mutant GluA1. Error bars indicate the 95% confidence interval of the means.

**Figure 4. Functional characterization of heteromeric mutant GluA1 receptors.**

**(A)** Representative steady-state currents evoked by sequential 10-20 s applications of 1 mM Glu (red bars), 1 mM Glu in the presence of 100 μM CTZ (*green bars*), 300 uM KA (blue bars), and 300 uM KA in the presence of 100 uM CTZ (300 uM, *purple bars*), from WT and mutant GluA1/A2R receptors in the absence (*upper traces*) and presence (*lower traces*) of the TARP auxiliary subunit γ-2. The holding potential was -40 mV in all shown recordings. Note different amplitude scale for traces with and without γ-2.

**(B)** IV relationships of Glu-evoked currents from oocytes expressing WT and mutant GluA1 subunits alone (*white circles*), with the GluA2R subunit (black circles), and with the GluA2R subunit and γ-2 (*grey circles*). The current amplitude at the different holding potentials is shown normalized to the current at -40 mV. Data points represent the mean from 6 to 10 oocytes. Error bars indicate the SEM and are shown when larger than symbol size.

**(C)** Summaries of the amplitudes of Glu-evoked currents in individual oocytes expressed recorded at -40 mV in absence (*red* *symbols*) and presence (*green symbols*) of CTZ.

**(D)** Summaries of the average desensitization and KA/Glu ratios for WT and mutant GluA1/A2R with (*right panels*) and without γ-2 (*left panels*). Error bars indicate the 95% confidence interval of the mean.

**Figure 5. Structural role of GluA1 residues affected by the *GRIA1* variants.**

**(A)** Cartoon illustration of a four-state model for the structural mechanism underlying AMPAR function and summary of the effects of the I627T, A636T, and G745D mutations. For simplicity, only two subunits with the ABD and M3 segments (*green and gray*, respectively) are shown, organized in a dimer complex. In the model, agonist (*black spheres*) binding to the clamshell-shaped ABD promotes the transition from the resting state to the pre-active state where the D1 and D2 subdomains of the ABD adopt a closed conformation around the agonist. From the pre-active state, the receptor can transition into the active state, which involves conformational changes in the upper region of the M3 helices that open the channel, or to a desensitized state, which involves conformational changes around the ABD dimer interface. The D1/D2 hinge region is highlighted in orange. The table shows a qualitative summary of the mutational effects on receptor function with upward and downward arrows indicating increase and decrease, respectively, at homomeric GluA1 and heteromeric GluA1/A2 receptors.

**(B)** Upper panels show side-on views of the ABD region in a GluA1 subunit dimer (*green and gray, respectively*) in the resting (*left*), active (*middle*), and desensitized (*right*) states with the Gly745 alpha carbon shown as red spheres. Lower panels show zoomed views of the side chains of Ile495 (*blue*), Pro508 (*cyan*), and Leu765 (*black*) as stick representations with the atomic surface indicated by dots. These residues form a hydrophobic interaction network across the D1/D1 interface in the resting and active states. The G745D mutation will project the negatively charged aspartate side chain into this interaction network (indicated by red arrow).

**(C)** Upper panels show side-on views of the channel region formed by the M3 helices in a GluA1 subunit dimer in the resting (*left*), active (*middle*), and desensitized (*right*) states. The lower panels show zoomed views of the side chains of Ile627 (*red*), Phe598 (*cyan*), and Gln600 (*blue*) as sticks with atomic surfaces indicated by dots. In the active state conformation, but not in the resting and desensitized states, these residues form interactions that might stabilize the open-channel conformation of the M3 helices. The channel center axis is indicated by the blue arrow.

**(D)** Upper panels show extracellular top views of the channel gate with the side chains of Ala636 (*red*), Thr639 (*cyan*), and Leu638 (*blue*) are shown as sticks with atomic surfaces indicated by dots. The Ala636 side chains contribute to stabilizing the closed-gate conformation by forming hydrophobic interactions with Thr639 and Leu636 in the resting and desensitized conformations (*lower panels*).

**Tables**

**Table 1. Variant information**

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| Patient | Variant | GRCh38 | cDNA | CADD | SIFT | PolyPhen2 | gnomAD frequency |
| 1 | p.Arg377Ter | g.153686324C>T | c.1129C>T homozygous | 36.0 | - | - | 0 |
| 2 - 4 | p.Ala636T | g.153764516G>A | c.1906G>A heterozygous | 29.3 | deleterious | probably damaging | 0 |
| 5 | p.Gly745Asp | g.153770379:G>A | c.2234G>A heterozygous | 28.1 | deleterious | probably damaging | 0 |
| 6 | p.Ile627Thr | g.153764490T>C | c.1880T>C heterozygous | 28.5 | deleterious | possibly damaging | 0 |
| 7 | p.Arg345Gln | g:153686229:G>A | c.1034G>A heterozygous | 21.8 | tolerated | benign | 1.24 × 10e-5 |

The table shows the five *GRIA1* variants identified in subjects, the resultant change in amino acid, the genomic DNA nucleotide change in *GRIA1*, and the site of the variants in cDNA encoding the GluA1 subunit protein. Combined annotation-dependent depletion (CADD) scores 19 predicted that variants p.Arg377Ter, p.Ala636T, p.Gly745Asp, p.Ile627Thr are highly likely to deleterious variants. Sorting Intolerant From Tolerant (SIFT)20) and Polymorphism Phenotyping v2 (PolyPhen2)21 analysis predicted all variants to be deleterious or damaging, except for p.Arg345Gln, which is predicted as tolerated or benign. Note that the p.Ala636Thr, p.Ile627Thr, and p.Gly745Asp variants (in patients 2, 5, and 6, respectively) were reported previously 13-15.

**Table 2. Clinical features of patients harboring GRIA1 variants**

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **Patient** | **Patient 1** | **Patient 2** | **Patient 3** | **Patient 4** | **Patient 5** | **Patient 6** | **Patient 7** |
| **Transcript number** | NM\_000827.3 | NM\_000827.3 | NM\_000827.3 | NM\_000827.3 | NM\_000827.3 | NM\_000827.3 | NM\_000827.3 |
| ***GRIA1* variant** | c.1129C>T  p.Arg377Ter | c.1906G>A  p.Ala636Thr | c.1906G>A  p.Ala636Thr | c.1906G>A  p.Ala636Thr | c.1880T>C p.Ile627Thr | c.2234G>A  p.Gly745Asp | c.1034G>A p.Arg345Gln |
| **Current age** | 10 | 7 | 13 | 26 | N.R. | 21 | N.R. |
| **Sex** | Female | Female | Male | Female | Male | Female | N.R. |
| **Intellectual disability (ID)** | Yes | Yes | Yes | Yes | Yes | Yes | Yes |
| **Level of cognitive impairment based on the clinical impression** | Severe | Severe | Severe | Moderate | Not classified | Moderate | Not classified |
| **Level of speech impairment** | Non verbal | Non verbal | Non verbal | Language difficult to understand | N.R. | Simple verbal language | N.R. |
| **Motor development** | Walked at 14 months. | Walked at 12 months  Sat at 12 months | Walked at 18 months Sat at 5 months. | Walked at 17 months | N.R. | Walked at 13 months  Delayed fine motor skills and coordination issues | N.R. |
| Epilepsy diagnosis | Yes | N.R. | No | No | No | No | No |
| Electroencephalogram (EEG) | Frequent interictal epileptiform discharges with spikes/spike over  posterior regions | N.R. | N.R. | N.R. | N.R. | Normal | N.R. |
| **Other movement disorder** | Tip-toe walking | N.R. | N.R. | No | N.R. | Dystonia  Catatonia | N.R. |
| **Behavioral issues** | Self-injurious behavior | Unspecified behavioral problems | ADHD  ASD | ADHD | Unspecified behavioral problems | Anxiety Anger tantrums  ASD | N.R. |
| Brain MRI | Normal | Normal | Normal | Normal | Normal | Normal | N.R. |
| Sleep | Poor sleep requiring melatonin | N.R. | Poor sleep requiring melatonin | Normal | Normal | Poor sleep | N.R. |
| **Vision** | Left intermittent divergent squint | N.R. | Possible squint, not diagnosed | Hypermetropia Astigmatism Glasses at 11 months | N.R. | Normal | N.R. |
| Dysmorphic features | No | Mild upslanting palpebral fissures | Broad forehead and telecanthus | No | Flushed cheeks and ears High arched palate | Normal | N.R. |
| Endocrine/Metabolic disease | Precocious puberty from 6 years of life | N.R. | No | No | No | Hypothyroidism Polycystic ovarian syndrome | N.R. |
| Cardiovascular disease | N.R. | N.R. | N.R. | No | Bicuspid aortic valve | No | N.R. |
| **Head circumference (cm)** | 49 | 50 | 53 | N.R. | N.R. | N.R. | N.R. |
| **Age at physical assessment (years)** | 9 ½ | 5 ½ | 4 | 26 | N.R. | 19 ½ | N.R. |
| **Weight (kg)** | 31 | 26 | 21 | 60 | N.R. | 54 | N.R. |
| **Height (cm)** | 140.5 | 118 | 107.3 | 149 | N.R. | 157.5 | N.R. |
| **Other genetic findings** | Normal *SLC2A1* and Epilepsy and Severe delay gene panel | Normal 250k SNP array and *FMR1* analysis | N.R. | normal array CGH | N.R. | N.R. | N.R. |

ASD, autism spectrum disorder; ADHD, Attention deficit hyperactivity disorder; ID, intellectual disability; N.R., not reported. Some information for Patient 2 and 5 were reported previously13; 15.

**Table 3. Functional parameters for WT and mutant GluA1-containing AMPAR subtypes**

|  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Receptor** | **Mean current (nA)** | ***n*** | **Mean current with CTZ (nA)** | ***n*** | **Glu EC50**  **(uM)** | ***n*** | **Glu EC50 with CTZ**  **(uM)** | ***n*** | **Fold desensitization** | ***N*** | **KA/Glu**  **(%)** | ***n*** |
| **WT** | 104  (81-127) | 53 | 4074  (2316-5832) | 45 | 23 (21-25) | 25 | 46 (40-51) | 18 | 62 (51-72) | 21 | 21 (18-25) | 36 |
| + A2R | 511  (372-651) | 54 | 7051 (4631-9472) | 26 | 18 (17-18) | 31 | n.d. | - | 57 (46-66) | 19 | 30 (27-32) | 25 |
| + A2R  + γ-2 | 1474 (1028-1919) | 53 | 7930 (4350-11509) | 23 | 14 (12-17) | 11 | n.d. | - | 7.2  (5.6-8.7) | 46 | 74 (69-78) | 27 |
| **R354Q** | 114  (63-164) | 20 | 2081  (524-1959) | 14 | 14 (12-16) | 10 | n.d. | - | 71 (48-94) | 15 | 24 (17-32) | 9 |
| + A2R | 255  (179-331) | 26 | 3587  (2187-4987) | 10 | 9.4 (8.4-11) | 13 | n.d. | - | 38 (26-48) | 9 | 30 (27-35) | 16 |
| + A2R  + γ-2 | 1001  (720-1282) | 43 | 4433  (2637-6230) | 16 | 5.9  (4.7-7.2) | 12 | n.d. | - | 5.9  (4.3-7.5) | 16 | 80 (73-87) | 16 |
| **R377\*** | 0\*\*\* (0-0) | 10 | 0\*\*\* | 10 | n.d. | - | n.d. | - | n.d. | - | n.d. | - |
| + A2R | 0\*\*\* (0-0) | 10 | 0\*\*\* | 10 | n.d. | - | n.d. | - | n.d. | - | n.d. | - |
| + A2R  + γ-2 | 0\*\*\* (0-0) | 10 | 22\*\*\* (2-40) | 10 | n.d. | - | n.d. | - | n.d. | - | n.d. | - |
| **I627T** | 1\*\*\* (0-1) | 26 | 12\*\*\* (8-17) | 31 | n.d. | - | 28\* (26-30) | 20 | 52 (35-69) | 10 | 5\*\*\* (4-7) | 10 |
| + A2R | 40\*\*\* (28-52) | 25 | 2728\*\*  (1821-3635) | 16 | 15 (13-17) | 10 | n.d. | - | 72 (47-97) | 12 | 9\*\*\* (8-11) | 17 |
| + A2R  + γ-2 | 825 (610-1041) | 41 | 13505\* (7835-19174) | 9 | 11 (9-13) | 17 | n.d. | - | 14 (12-18) | 13 | 66 (61-71) | 11 |
| **A636T** | 993\*\*\*  (746-1246) | 43 | 1105\*\*  (729-1482) | 22 | 0.9 (0.8-1.0) | 26 | n.d. | - | 1.9\*\*\* (1.6-2.4) | 24 | 52\*\*\* (46-56) | 25 |
| + A2R | 2064\*\*  (1437-2690) | 27 | 2474\*\* (1636-3313) | 16 | 4.2 (3.9-4.6) | 10 | n.d. | - | 1.1\*\*\*  (1.0-1.3) | 16 | 81\*\*\* (70-90) | 15 |
| + A2R  + γ-2 | 2504  (1151-3857) | 20 | 1961\*\*  (1650-3758) | 12 | 5.8 (4.8-7.1) | 13 | n.d. | - | 1.3\*\*\* (1.1-1.4) | 14 | 65 (55-75) | 11 |
| **G745D** | 0\*\*\* (0-1) | 34 | 0\*\*\* (0-1) | 22 | n.d. | - | n.d. | - | n.d. | n.d. | n.d. | - |
| + A2R | 83\*\*\*  (57-109) | 29 | 347\*\*\* (191-504) | 11 | 14 (13-16) | 11 | n.d. | - | 6.7\*\*\*  (6.0-7.0) | 11 | 58\*\*\* (52-63) | 11 |
| + A2R  + γ-2 | 219\*\*\*  (134-303) | 31 | 767\*\*\* (354-1179) | 22 | 5.8 (4.3-8.0) | 10 | n.d. | - | 7.3 (5.8-8.9) | 26 | 94\*\*\* (89-99) | 11 |

Values are given for mean currents, Glu EC50, fold-desensitization, and KA/Glu response ratios from the electrophysiological experiments as described in the text. The data represent means ± 95% confidence intervals. *n*, number of individual experiments or oocytes. Statistical information in the form of probability value (*p*) level is given where values are significantly different from WT as: \*, *p* < 0.05; \*\*, *p* < 0.01; \*\*\*, *p* < 0.001. CTZ, cyclothiazide; Glu, glutamate; KA, kainic acid; N.D, not determined; WT, wild-type.