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## Functional impairment of cortical AMPA receptors in schizophrenia

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### Abstract

Clinical and preclinical studies suggest that some of the behavioral alterations observed in schizophrenia (SZ) may be mechanistically linked to synaptic dysfunction of glutamatergic signaling. Recent genetic and proteomic studies suggest alterations of cortical glutamate receptors of the AMPA-type (AMPA receptors), which are the predominant ligand-gated ionic channels of fast transmission at excitatory synapses. The impact of gene and protein alterations on the electrophysiological activity of AMPA receptors is not known in SZ. In this proof of principle work, using human postmortem brain synaptic membranes isolated from the dorsolateral prefrontal cortex (DLPFC), we combined electrophysiological analysis from microtransplanted synaptic membranes (MSM) with transcriptomic (RNA-Seq) and label-free proteomics data in 10 control and 10 subjects diagnosed with SZ. We observed in SZ a reduction in the amplitude of AMPA receptor currents elicited by kainate, an agonist of AMPA receptors that blocks the desensitization of the receptor. This reduction was not associated with protein abundance but with a reduction in kainate's potency to activate AMPA receptors. Electrophysiologically-anchored dataset analysis (EDA)

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was used to identify synaptosomal proteins that linearly correlate with the amplitude of the AMPARs responses, gene ontology functional annotations were then used to determine protein-protein interactions. Protein modules associated with positive AMPARs current increases were down-regulated in SZ, while protein modules that were up-regulated in SZ were associated with decreased AMPARs currents. Our results indicate that transcriptomic and proteomic alterations, frequently observed in the DLPFC in SZ, converge at the synaptic level producing a functional electrophysiological impairment of AMPARs.

## Keywords

AMPA receptors; microtransplantation of synaptic membranes; schizophrenia; synaptic dysfunction

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## 1. Introduction

Alterations of excitatory synaptic function have been found in transcriptomic, genetic, and proteomic studies of schizophrenia (SZ) (Akbarian et al., 1996; Banerjee et al., 2015; Catts et al., 2015; Catts et al., 2016; Gluck et al., 2002; Hall et al., 2020; Harrison, 1999; Harrison and Weinberger, 2005; Hu et al., 2015; Mirnic et al., 2000; Pinner et al., 2016; Sequeira et al., 2012; Vawter et al., 2002). Clinical and preclinical studies suggest that abnormal synaptic transmission and several behavioral abnormalities in SZ may be mechanistically linked to abnormal glutamatergic neurotransmission, probably through hypofunction of N-methyl-D-aspartate (NMDA) receptors (NMDARs), in brain regions affected in SZ (Coyle et al., 2003; Gilmour et al., 2012; Hardingham and Do, 2016; Jadi et al., 2016; Krystal et al., 2002; Krystal et al., 2003; Krystal et al., 1994). However, the limited clinical efficacy of pharmacological modulation of NMDARs in clinical and preclinical trials indicate that synaptic dysfunction in SZ is a far more complex process (Fleischhacker and Miyamoto, 2016; Miyamoto et al., 2012). For instance, the function of NMDARs at the synapse is interdependent with the activity of  $\alpha$ -amino-3-hydroxy-5-methyl-4-isoxazole propionic acid (AMPA) receptors (AMPARs) (Cull-Candy and Leszkiewicz, 2004). At rest, NMDARs are blocked by extracellular  $Mg^{2+}$ ; activation of postsynaptic AMPARs removes the  $Mg^{2+}$  blockade allowing NMDARs to be permeable and electrically active (Mayer and Westbrook, 1987; Mayer et al., 1984; Nowak et al., 1984). Thus, abnormal responses of AMPARs may lead to hypofunction of NMDARs. Indeed, a recent genetic study found that mutations in GRIA2, the gene for the most expressed cortical AMPARs subunit GluA2, is strongly associated with SZ (Gulsuner et al., 2020). Similarly, synaptic GluA2 and GluA3 expression were mostly, but not uniformly, reduced in the auditory cortex of people with SZ (MacDonald et al., 2019). Taken together these recent studies strongly suggests a primary role of AMPARs in SZ.

Although AMPARs are constructed as heterotetramers from a pool of only four subunits (GluA1-GluA4), a large functional heterogeneity arises from diverse heteromeric assemblies, alternative splicing (Sommer et al., 1990), RNA editing (Wright and Vissel, 2012), post-translational modifications (Lussier et al., 2015), and transmembrane accessory proteins (Carbone and Plested, 2016; Diaz, 2010). Notably, transcriptional and proteomic

changes of AMPARs have been demonstrated in the dorsolateral prefrontal cortex (DLPFC) of brains from SZ patients; although some studies have also found no changes (Corti et al., 2011; Hammond et al., 2010; Pinner et al., 2016; Rubio et al., 2012). Importantly, the direct measurement of ion currents through synaptic receptors from fresh-frozen brains from subjects with SZ has never been undertaken, and the impact of reported transcriptional and proteomic alterations that can affect the physiology of synaptic receptors in SZ is not known. In this proof of principle work using human postmortem brain synaptic membranes, we show that the function of synaptic AMPARs is significantly altered in SZ. Our results indicate that transcriptomic and proteomic alterations observed in the dorsolateral prefrontal cortex (DLPFC) from SZ brains, converge at the synaptic level producing functional impairment of AMPARs currents.

## 2. Materials and Methods

This preliminary research was performed using gray matter from the DLPFC in a cohort of 10 SZ cases and 10 controls from which we first analysed expression using RNA-Seq; second, label-free proteomics was performed using liquid chromatography-mass spectrometry (LC-MS/MS) in enriched-synaptosome preparations, and finally, we studied two-electrode voltage clamp electrophysiological (TEVC) profiles using microtransplantation of synaptic membranes (MSM) (Fig. 1A). Postmortem brain specimens of the DLPFC (BA9 plus 46, without selection for the left or right hemisphere) were obtained from the University of California Irvine Brain Bank (UCIBB) following the university's Institutional Review Board and after obtaining verbal and written consent from next of kin. A psychological autopsy was completed based on family informant interview, medical and psychiatric records, coroner's toxicology reports, and the subject's medication history. The UCIBB autopsy protocol is based largely on procedures validated by Kelly and Mann (Kelly and Mann, 1996) and includes questions concerning the decedents' demographics, medical history, psychiatric symptoms, medication use, hospitalizations, substance use, and physical health. The human brain dissection and freezing protocol are described in detail elsewhere (Jones et al., 1992). Briefly, after collection, frontal cortex samples were frozen in isopentane at  $-40^{\circ}\text{C}$  and then stored at  $-80^{\circ}\text{C}$  for downstream studies. The subjects' demographic comparison for age, gender, postmortem interval (PMI), and RNA integrity number (RIN) showed that pH and RIN were significantly different in subjects with SZ compared to controls (Table 1 and supplementary data 1 for demographics, use of drugs, and prescribed medications use). Correction for confounding variables is described in the transcription profile section. This cohort was free of known disorders that could potentially cause synaptic alterations due to neurodegeneration, such as Alzheimer's disease, epilepsy, AIDS, or Parkinson's disease. The DLPFC was selected because the altered activation of this region in functional imaging studies have shown that 'hypofrontality' is one of the best-replicated findings in patients with SZ (Barch et al., 2001; Bolkan et al., 2016; Perlstein et al., 2001; Weinberger and Berman, 1996; Weinberger et al., 1988).

## 2.1 Transcriptomic Profile

We characterized the transcriptome from the gray matter of 20 DLPFC samples by RNA-Seq. RNA was prepared from 80 to 100 mg of frozen tissue samples, with Omni Prep Multi-Sample Homogenizer (Omni International, Kennesaw, GA). Total RNA isolation was performed with TRIZOL™ reagent (Invitrogen, Carlsbad, CA), following the manufacturer's instructions. RNA was quantified by OD260/280 with a UV spectrophotometer and treated with RNase-free DNase using the RNeasy MinElute columns (Qiagen, Valencia, CA). The quality of the total RNA was finally evaluated using the Agilent 2100 Bioanalyzer RNA Chip (Santa Clara, CA). Libraries were prepared using Illumina TruSeq RNA Sample Prep kit v2., following manufacturer's instructions. Pooled libraries (8–12 samples per lane) were then sequenced on a HiSeq 2000 platform, generating 100 bp paired-end reads, yielding an average of 299,831,258 reads per lane. For data analysis, first, TruSeq adapter sequences were trimmed using cutadapt (v2.5; with settings -n 3 --trim-n -m 40). Reads mapping to the human transcriptome (Gencode version 32) were then quantified using Salmon (Patro et al., 2017) (v0.14.1; indexed with -k 31 and quantified with settings --validateMappings --mimicBT2 --gcBias --seqBias). Mapping rates were  $83.63\% \pm 1.05\%$  (mean  $\pm$  SD). Read counts were then summarized at the gene level, and genes with counts per million (CPM)  $>5$  in at least two samples were considered expressed, yielding a total of 13,849 expressed genes used for all downstream analyses. Latent variables were estimated using surrogate variable analysis (SVA; svaseq approach (Leek, 2014; Leek et al., 2012). Differential gene expression was analyzed using the edgeR Bioconductor package (Robinson et al., 2010); gene-level counts were fit to a negative binomial model containing diagnosis and four surrogate variables representing latent variables. Two of the four surrogate variables (SVs) were highly correlated with RIN and pH respectively (supplementary figure 1). Differential gene expression between controls and SZ was assessed using Likelihood Ratio Tests. To adjust for multiple testing false-discovery rates were obtained using the Benjamini-Hochberg method.

## 2.2 Synaptosomes preparation

Human synaptoneuroosomes were isolated from  $\approx 50$  mg frozen gray matter samples of the DLPFC using the Syn-PER method (Thermo Fisher Scientific) using the manufacturer instructions and as previously reported (Limon et al., 2019). The resultant pellet, enriched in synaptoneuroosomes (P2 fraction), was resuspended in sterile distilled to break up the synaptosomes and to create small proteoliposomes containing synaptic membranes and their synaptic proteins. The protein concentration of the P2 fraction was determined by using Qubit protein assay reagent kit (Thermo Fisher Scientific). No differences in the total amount of protein in synaptic preparations was found between groups (2.756 mg/mL in controls vs 2.76 mg/mL in SZ (n= 10 ctrl, 10 SZ; t-test, double tailed, p-value = 0.920). These P2 membrane preparations were stored at  $-80^{\circ}\text{C}$  until needed for downstream studies.

## 2.3 Proteomic Profile

One aliquote containing 50  $\mu\text{g}$  of protein from each P2 membrane preparation of each subject was analyzed by LC-MS/MS to measure relative protein abundance in the synapto-proteome (English et al., 2015; Focking et al., 2015). The analysis was done on a Thermo

Scientific Q Exactive mass spectrometer connected to a Dionex Ultimate 3000 (RSLCnano) chromatography system as has been previously described (English et al., 2015; Focking et al., 2015). All data were acquired with the mass spectrometer operating in positive ion and automatic data-dependent switching mode. A high resolution (70,000) MS scan (300–1600 m/z) was performed using the Q Exactive to select the 12 most intense ions prior to MS/MS analysis using high energy collision-induced dissociation (HCD). We collected two types of mass spectrometry data in the experiments. It is formally a ‘shotgun’ experiment and the collected tandem mass spectra were used to identify the proteins. The parent ion signal (MS1 scan) on high-resolution instruments such as the Q Exactive is widely used to calculate protein abundance in so-called ‘label-free’ proteomics experiments (Cox and Mann, 2011). The signal for each peptide derived from a given protein is integrated over time (the time it takes to enter the instrument via HPLC), and the summed signals are used to estimate the relative abundance of each protein. To analyze the data, we used the MaxQuant program specifically for label-free experiments using high-resolution instruments supported by Andromeda as a database search engine for peptide identification (Cox et al., 2011). This program has several statistical control steps to ensure that only high-quality reliable ion signals are accepted. Label-free quantitation was performed as previously described (Luber et al., 2010). Carbamidomethylation was defined as a fixed modification, while oxidation and acetylation of the protein N-terminus were defined as variable modifications. Only peptides with seven or more amino acid residues were allowed for identification. Additionally, at least one unique peptide was required to identify a protein. The cut off for false discovery rate for peptide and protein identification was set to 0.01 (1%). To avoid bias associated with protein under-representation between groups, proteins were excluded in cases where there was less than 50% availability of the LFQ intensities in each biological group. The label-free algorithm takes the maximum number of identified peptides between any two samples and compares the intensity of these peptides to determine peptide ratios. Label-free quantitation (LFQ) intensity values were used for protein quantification across the groups. Raw LFQ intensities were extracted from the MaxQuant software and log base 2 transformed prior to analysis to eliminate distributional skew and to give approximate normality. Regression normalization was performed to remove technical variation between samples (Callister et al., 2006). Significance testing was performed at the 5% level between SZ and control samples. Fold changes were obtained from an exponentiation (power of 2) of the difference in group means. Fold changes <1 were inverted (–1/fold change) to present a consistent scale of measurement. Two control samples and one SZ sample that had poor chromatographic profiles as compared to the rest of the samples LC-MS/MS were excluded from the proteomic analyses. All 20 subjects were included in electrophysiological experiments using microtransplantation of synaptic membranes (MSM) procedures.

#### 2.4 Two Electrode Voltage Clamp (TEVC) electrophysiology Profile

Aliquots from the original P2 preparation were used in MSM experiments to record ion currents elicited by native AMPARs embedded in synaptic membranes (Limon, 2019; Mazzo et al., 2016). Briefly, synaptosome-enriched preparations were sonicated, 3 times for 5 seconds at 1 min intervals in ice, to break the synaptosomes and create small proteoliposomes. These sonicated preparations were injected into stage V-VI *Xenopus*

*laevis* oocytes using protocols previously published for cellular membranes (Conti et al., 2013; Limon et al., 2008, 2011, 2012). Each oocyte was injected with 50 nL of synaptic proteoliposomes (2 mg/mL protein concentration) and recorded after 18–36 hrs of the injection. This is the period of time when responses are most stable. For electrophysiological recordings, oocytes were placed in a recording chamber (volume  $\approx 0.1$  ml) and perfused continuously (5–10 ml/min) with Ringer's solution [115 mM NaCl, 2 mM KCl, 1.8 mM CaCl<sub>2</sub>, 5 mM HEPES (pH 7.4)] at room temperature (19–21°C). Data acquisition and analyses were performed by using WinEDR v2.3.8 Strathclyde Electrophysiology software (John Dempster, Glasgow, United Kingdom). To determine the amplitude of AMPA receptors the maximum amplitude of ion currents activated by 100  $\mu$ M kainate, which elicits non-desensitizing currents through AMPARs (Limon et al., 2010; Stern-Bach et al., 1998) was used. The use of kainate allows consistent recordings of ion currents via AMPARs and produces less intra-subject variation compared to currents elicited by the combination of AMPA and cyclothiazide, which are coapplied to remove the strong desensitization of AMPARs (Limon et al., 2007). Other chemicals were from Sigma-Aldrich. Electrophysiological results are expressed as mean  $\pm$  SEM. Statistical comparisons to identify the effect of diagnosis and the contribution of demographic variables (age, RIN, PMI, and pH) used the mean of kainate current for each subject (n = 10–14 injected oocytes were technical replicates per subject) in an analysis of covariance (ANCOVA) model. Fourteen oocytes per subject were recorded to measure current amplitude, but only oocytes that showed a stable baseline and signal to noise ratio of at least 2 were included in the analysis. To determine differences in current amplitude an ANCOVA (between-subject factor: diagnosis; covariates: pH and RIN; interaction: pH\*Diagnosis; (JMP, version 14) was used. The EC<sub>50</sub> for kainate for microtransplanted oocytes was determined as previously reported (Limon, 2019). In all cases, p<0.05 was considered significant. Pearson product-moment was used for linear correlations using JMP version 14, Adjusted p-values are shown when multivariate analyses were implemented. Previously we have shown stability of the electrical recording with long PMIs (Limon et al., 2008).

## 2.5 Functional enrichment and protein-protein interaction analyses.

Functional enrichment analysis was done with Metascape web interface which incorporates access to 40 knowledgebases (Zhou et al., 2019). Transcriptomic analysis of differentially expressed genes was done against the whole genome. For pathway and protein enrichment a minimum overlap of 3, p-value cutoff of 0.01, minimum enrichment of 1.5, and gene prioritization by evidence counting (GPEC) was used (Zhou et al., 2019). For protein-protein interaction enrichment, the minimum network size was 3 using the databases BioGrid, InWeb\_IM (human) and OmniPath (human). Proteomics data was analyzed by obtaining dysregulated proteins using GPEC and the total list of identified proteins by LC-MS/MS as background. For the integrated analysis of transcriptomic, proteomic and subset of proteins that correlated with ion currents from AMPARs, we used the multi-list feature of Metascape using pathway and protein enrichment with a minimum overlap of 3, p-value cutoff of 0.00001, minimum enrichment of 3 and GPEC; the proteomics dataset was used as background.



### 3. Results

#### 3.1 Whole transcriptomic and synaptosome-proteomic profiles

Gene expression levels measured by RNA-Seq were significantly upregulated in SZ for 151 genes at the  $p < 0.01$  level (supplementary data 2). Functional and protein-protein interaction analyses of these genes showed significant enrichment for chaperone-mediated folding, positive regulation of inflammation, stress response, negative regulation of phosphorylation and ATP biosynthesis (Fig. 1, supplementary data 3). In contrast, 186 genes were downregulated in SZ. These downregulated genes were found to be significantly involved in synaptic signaling, signal release and transmission across chemical synapses (supplementary data 4). These results are in agreement with a reduction of synaptic communication and loss of dendritic spines that have been observed in the DLPFC of SZ subjects (Glantz and Lewis, 2000). However, the expression of AMPARs subunits (GRIA1, GRIA2, GRIA3 and GRIA4), the principal gates for synaptic input at synaptic spines, was not different between control and SZ subjects (Fig. 1F).

To explore the possible effect of medications we reanalyzed the non-human primate CommonMind Consortium (CMC) Knowledge Portal macaque collection consisting of clozapine ( $n=9$ ), and high haloperidol dosage ( $n=7$ ) and low dosage ( $n=10$ ) compared with placebo ( $n=10$ ). The collection was obtained from Synapse (MP Vawter, PI) and analyzed by using 3 surrogate variables in an SVA analysis, and drug group as a factor. The details of the CMC Macaque collection can be found on the CMC website (<https://www.synapse.org/#!Synapse:syn2759792/wiki/69613>). The analysis of the macaque frontal cortex showed that none of the GRIA1, -2, -3, -4 genes were differentially expressed in the low Haldol or Clozapine drug group compared to the control group (supplementary data). GRIA2 and GRIA3 were nominally decreased ( $p = 0.04$ , not significant following FDR correction) in expression in the high Haldol dose group compared to controls by 7%.

Comparative studies between transcriptome and proteome in animal models have shown that levels of transcripts and proteins do not necessarily correlate (Ghazalpour et al., 2011); therefore, we determined the relative abundance of AMPARs subunits within the context of the synapto-proteome. In agreement with transcriptomic analysis, the abundance of AMPARs subunits in the proteomics analysis was not different between controls and SZ subjects (Fig. 2). However, GluA4 was not found in our synaptic preparations. This was probably due to its very low abundance in the cortex (Schwenk et al., 2014; Schwenk et al., 2012) and the detection limits of label-free proteomics (Sandberg et al., 2014). Interestingly, only mRNA and protein levels for GRIA3 and GluA3 correlated ( $r=0.59$   $p=0.013$ ; supplementary figure 2).

In both groups the protein levels for GluA2 was the most abundant subunit, accounting for  $75.3 \pm 1.7$  % of the total protein available to assemble in AMPARs (mean  $\pm$  SD;  $n=17$  subjects), followed by GluA3 ( $22.6 \pm 1.6$  %) and GluA1 ( $2.1 \pm 0.3$  %). The relative proportion of GluA2 > GluA3 > GluA1 > GluA4 has the same order of that reported in the cortex of rodents, although the proportion in rodent for those subunits are approximately 45%, 27%, 21%, and 6%, respectively (Schwenk et al., 2014).



No differences in the relative proportions of AMPARs subunits were found between controls and SZ subjects. The proportions and high correlation between GluA2 and GluA3 ( $r(17) = 0.737$ ;  $p=0.0007$ ; Pearson's correlation) compared to that of GluA2 and GluA1 ( $r(17) = 0.381$ ;  $p=0.13$ ) suggest the presence of heteromeric GluA2/3 and homomeric GluA2 receptors as the major drivers of synaptic inputs in the DLPFC and a significantly lower proportion of GluA1/2 receptors (Fig. 2D).

Further analysis of the 2487 identified proteins in the P2 fractions identified 128 proteins that were differentially expressed between SZ and control subjects ( $P<0.05$ ; supplementary data 5), 54 and 74 proteins were upregulated and downregulated in SZ, respectively. Protein to protein interaction (PPI) and functional annotation analyses found that upregulated proteins participate in ERBB2/ERBB3 signaling, stress response, and autophosphorylation pathways (supplementary data 6). Downregulated proteins were found to be involved in translational silencing of gene expression, particularly LI3a-mediated (e.g. *EIF4B*, *RPL28*), tissue regeneration (*GAP43*, *GJA1*), and neurotransmitter uptake and release (*SLC6A1*, *SNAP25*) (supplementary data 7). These results also indicate convergence between RNA-Seq and synaptic-proteomic data on the downregulation of synaptic function, although the total abundance of AMPARs subunits is maintained in the DLPFC of SZ subjects.

### 3.2 Electrophysiology of native AMPA receptors profile.

AMPA receptors are a complex of proteins which in addition to the four principal subunits (GluA1-GluA4), they are co-assembled with a large number of auxiliary proteins (Schwenk et al., 2012; Schwenk et al., 2009) that modulate the gating, permeability and pharmacology of the channel (Cho et al., 2007; Gill et al., 2011; Herring et al., 2013; Kato et al., 2010a; Kato et al., 2010b; Milstein et al., 2007). Similarly, AMPA receptors activity is modulated by many intracellular pathways (Wang et al., 2005). To determine whether proteomic changes in SZ impact the electrophysiological activity of AMPARs, aliquots of the original P2 samples used for proteomics were microtransplanted into *Xenopus* oocytes. All samples from the cohort exhibited functional agonist-induced responses when tested by TEVC in MSM, confirming the viability of AMPARs. Figure 3A shows ion currents elicited by 100  $\mu$ M kainate from DLPFC synaptosome membranes injected into oocytes. Kainate is a low-affinity agonist of AMPARs that keeps the channel open in a non-desensitized state, allowing the measurement of kainate-induced AMPARs responses in the steady-state (Limon et al., 2010). Because *Xenopus* oocytes do not express endogenous AMPARs (Limon et al., 2008; Limon, 2019), electrophysiological responses elicited by AMPARs agonists in microtransplanted oocytes are from functional and successfully incorporated, receptor protein complexes docked into the surface of the oocyte. This is consistent with previous work indicating that synaptic function and intracellular signaling is preserved in fresh-frozen and thawed human postmortem brain slices (Hahn et al., 2006).

The mean of kainate-induced AMPARs responses (AMPARs currents) for each subject was not significantly correlated with age or PMI ( $r(20) = 0.0245$ ,  $p = 0.918$  for age, and  $r(20) = 0.3$ ,  $p = 0.19$  for PMI), but found significant correlations with pH and RIN in the controls, but not in SZ. A screening effect for sources of variation confirmed the presence of an interaction between diagnosis and pH for the amplitude of AMPARs currents. Therefore,

an ANCOVA model (between-subject factor: diagnosis; covariates: pH and RIN; interaction: pH\*Diagnosis; N=20,  $F(4,15) = 5.85$ ,  $p = 0.0048$ ) was used to test for diagnosis effects. This analysis found a significant effect of diagnosis for the reduction of AMPARs currents in SZ (Diagnosis LogWorth = 3.1;  $p = 0.0008$ ). Corrected AMPARs current was found to correlate with the protein levels of GluA2 and GluA3 in the cohort ( $r(17) = 0.61$ ,  $p = 0.038$  for GluA2 and ( $r(17) = 0.61$ ,  $p = 0.01$  for GluA3) and the sum of all AMPARs subunits ( $\Sigma$ GluA; ( $r(17) = 0.54$ ,  $p = 0.03$ ). Notably, the correlation between AMPARs currents and protein levels was higher in SZ than in the control group e.g. AMPA responses vs GluA3 in control ( $r(8) = 0.583$ ,  $p = 0.129$ , and AMPA responses vs GluA3 in SZ ( $r(9) = 0.708$ ,  $p = 0.033$ ). Similarly, GluA1 was correlated with AMPARs currents in SZ ( $r(9) = 0.77$ ,  $p = 0.0149$ ) but not in the control ( $r(8) = 0.126$ ,  $p = 0.75$ ), suggesting a higher participation of GluR1 in AMPARs responses in SZ.

Because gene and proteomic analysis did not find changes in the abundance of AMPARs subunits, we hypothesized that lower amplitude of AMPARs currents in SZ was caused by a reduction in the  $EC_{50}$  for kainate. In agreement with this hypothesis, AMPARs receptors in SZ had lower  $EC_{50}$  for kainate than controls ( $p = 0.03$ , one-sided tail testing a priori for lower sensitivity, Fig 3E). The lower amplitude of AMPARs currents correlated with lower kainate potency (Fig. 3F). These results are consistent with a reported hypofunction of AMPARs (Coyle et al., 2003; Hammond et al., 2010; Pinner et al., 2016) and a loss of excitatory connectivity in the DLPFC of SZ patients (Glantz and Lewis, 2000; Harrison, 1999; Penzes et al., 2011).

### 3.3 Electrophysiologically-anchored dataset analysis (EDA) in SZ

Transcriptomic and proteomic datasets provide converging evidence indicating synaptic dysfunction in SZ; however, loss of synaptic communication in schizophrenia and compensatory rearrangements are cellular processes that involve a large number of structural and accessory proteins at the synapse. As a first examination of the utility in using MSM data to unravel this complexity, and to better understand the relationship between protein expression and physiological activity of AMPARs, we implemented what we called an EDA approach, to generate lists of proteins that correlate with the physiological activity of AMPARs (supplementary data 8 for all correlations). We found that in addition to the positive correlation with GluA subunit protein counts and AMPARs current, the AMPARs currents were also positively correlated with 118 proteins representing the excitatory postsynaptic density, and mitochondrial and ribosomal complexes (Fig. 4; supplementary data 9 shows all the modules assessed). A molecular complex detection algorithm MCODE, that identifies densely connected regions in large PPI networks (Bader and Hogue, 2003), identified 5 molecular modules: 1) Nonsense mediated decay (NMD) complex, which is an mRNA surveillance system that maintains synaptic architecture and synaptic vesicle efficacy (Long et al., 2010); 2)  $Ca^{2+}$  signaling pathway, which is fundamental for buffering of synaptic  $Ca^{2+}$ , neurotransmitter release and activity-dependent regulation of synaptic function (Greer and Greenberg, 2008); 3) the mitochondria electron transport respiratory chain complex I, 4) the mitochondria F1F0-ATP synthase of Complex V which is fundamental to supply the high metabolic demand of active synapses, and 5) activation of AMPARs and synaptic plasticity, which are the proteins responsible

for generating the AMPARs currents. In contrast, AMPARs currents were negatively correlated with 107 proteins involved in intracellular signaling, phosphorylation, and DNA biosynthetic processes (Fig. 5, supplementary data 10 shows all the modules assessed). Three MCODE modules that correlated negatively with AMPARs were representative of complexes involved in 1) formation of tubulin folding intermediates by CCT/TRiC, which are chaperonin mechanisms important for synaptic proteostasis (Gorenberg and Chandra, 2017), 2) glucagon signaling pathway, and 3) AURKA activation by TPX2 pathway, which is important for neurite elongation and dendrites differentiation (Kahn et al., 2015; Mori et al., 2009).

A metanalysis including transcriptomic, proteomic, and EDA-generated lists of proteins shows that 15.3% of proteins that positively correlate with the amplitude of AMPARs currents are downregulated in SZ (Fig. 6); in contrast, 10% of proteins with negative correlation with AMPARs currents are upregulated in SZ (supplementary data 11). Interestingly, functional convergence of upregulated transcriptome and synapto-proteome indicated that the MAPK pathway (Table 2), which has been consistently implicated in SZ studies, had a negative association with AMPARs currents (Figure 6).

#### 4. Discussion

In this preliminary study, we used a variant of the method of microtransplantation of plasma membranes from postmortem tissue (Miledi et al., 2004) to record, for the first time, the activity of synaptic AMPARs from the DLPFC of people diagnosed with SZ. It was previously shown that microtransplanting synaptic membranes from autopsy tissue allows the pharmacological and electrophysiological characterization of synaptic receptor complexes in human brain regions; a prerequisite for drug discovery (Limon, 2019; Mazzo et al., 2016; Zwart et al., 2019). MSM opens the door to use biophysical parameters of ion currents as functional endpoints for more sophisticated analysis in neurological and neuropsychiatric disorders. In this study, by integrating multi-dimensional data with physiological responses of postsynaptic receptors our results suggest a reduced function of AMPARs in SZ. This reduction may be at least partially explained by changes in potency and by alterations in protein modules that impact the trafficking and function of AMPARs. However, we cannot rule out reverse causation. It may be possible that altered AMPAR responses cause changes in protein subunit levels since previous studies have shown reductions (Beneyto and Meador-Woodruff, 2006; Vawter et al., 2002), no changes (Healy et al., 1998), or elevations (Chinopoulos et al., 2007; Dracheva et al., 2005) in the expression levels of AMPARs in postmortem studies of SZ.

Our transcriptomic and proteomic analysis of this cohort found no changes in the DLPFC levels of transcripts or proteins for any of the AMPARs subunits in SZ DLPFC; however, we observed functional deficits in the receptors' electrophysiological activity ( $EC_{50}$ ) that is supportive of AMPARs deficits in SZ. Normal abundance of mRNA and protein of AMPARs subunits in SZ may be a compensatory attempt to normalize synaptic function by homeostatic mechanisms (Turrigiano, 2017). Taken together, the data shows that alterations in cellular pathways and altered AMPARs function in the DLPFC are part of a molecular

alteration of synaptic signaling which may lead to a decreased excitatory input and functional dysconnectivity in the DLPFC in SZ.

The convergence of different lines of evidence suggests that synaptic functional alterations of AMPARs may underlie the hypofunction and decreased activation that has been consistently observed in the DLPFC of SZ patients (Berman, 2002). Reduced excitation and the subsequent potential shunt of NMDARs activation may underlie the generation of psychoses in individuals with SZ due to the complementary causes for NMDA receptor hypofunction that our data suggest. While this hypothesis is far from new, it does represent a functional measure that augments previous investigations that relied solely on singular methods that counted abundance of genes or proteins.

The analysis of the electrophysiological activity and relative abundances of native receptors from human brain tissue provides a glimpse of functional information that warrants further studies. For example, we observed that postmortem pH was negatively correlated with the amplitude of AMPARs currents in the control but not the SZ group. While it has been reported that patients with SZ may exhibit extracellular acidification in magnetic resonance spectroscopy (MRS) studies and postmortem brains associated with impaired oxidative phosphorylation and mitochondria dysfunction (Dogan et al., 2018; Hagihara et al., 2018; Sullivan et al., 2019b), and also that AMPARs are negatively modulated by extracellular H<sup>+</sup> (Ihle and Patneau, 2000; Traynelis and Cull-Candy, 1991), it is not known whether AMPARs in a chronically acidified milieu have compensatory modifications that make them less sensitive to agonal changes of pH. However, the experimental setting in which we studied native receptors was constant in terms of buffer and pH. This does not rule out the possibility that there are post-translational modifications or alternative splicing changes to the AMPARs that we have not accounted for in our proteomic analysis.

Further analyses studying the role of auxiliary subunits, which are known to modify the kinetics, pharmacology and polyamine block of AMPA receptors depending on the cellular needs (Kato et al., 2010b), in this process are needed. Auxiliary subunits may also be involved in the pharmacological differences that we observed in MSM experiments. Previous work by Meador-Woodruff's lab had shown a reduction in the transmembrane AMPA regulator protein TARP- $\gamma$ 8 in the anterior cingulate in SZ (Drummond et al., 2013); although we did not observe group differences in TARP- $\gamma$ 8 in the DLPFC in our cohort, in physiological conditions TARP- $\gamma$ 8 is known to increase glutamate and kainate affinity (Kato et al., 2010b; Tomita et al., 2007), and a reduction in this protein may explain the reduced potency of AMPARs that we observed in the DLPFC of some SZ subjects. In support of this hypothesis, the subject marked with an arrow in MSM experiments (Figure 3F) had the lowest gene expression for TARP- $\gamma$ 8 (CACGN8) in the SZ group and had no detectable protein quantities in LC-MS/MS experiments. Future studies in larger cohorts should aid in the categorization of SZ subtypes based on the specific alterations of genes or proteins that correlate with the excitatory electrical activity of the synapses.

Our analysis indicates that additional mechanisms converge into producing deficits of AMPARs. For instance, both, RNA-Seq and LC-MS/MS data implicate MAPK3 signaling in SZ, which is in agreement with *de novo* mutations and alterations in copy number



## Supplementary Material

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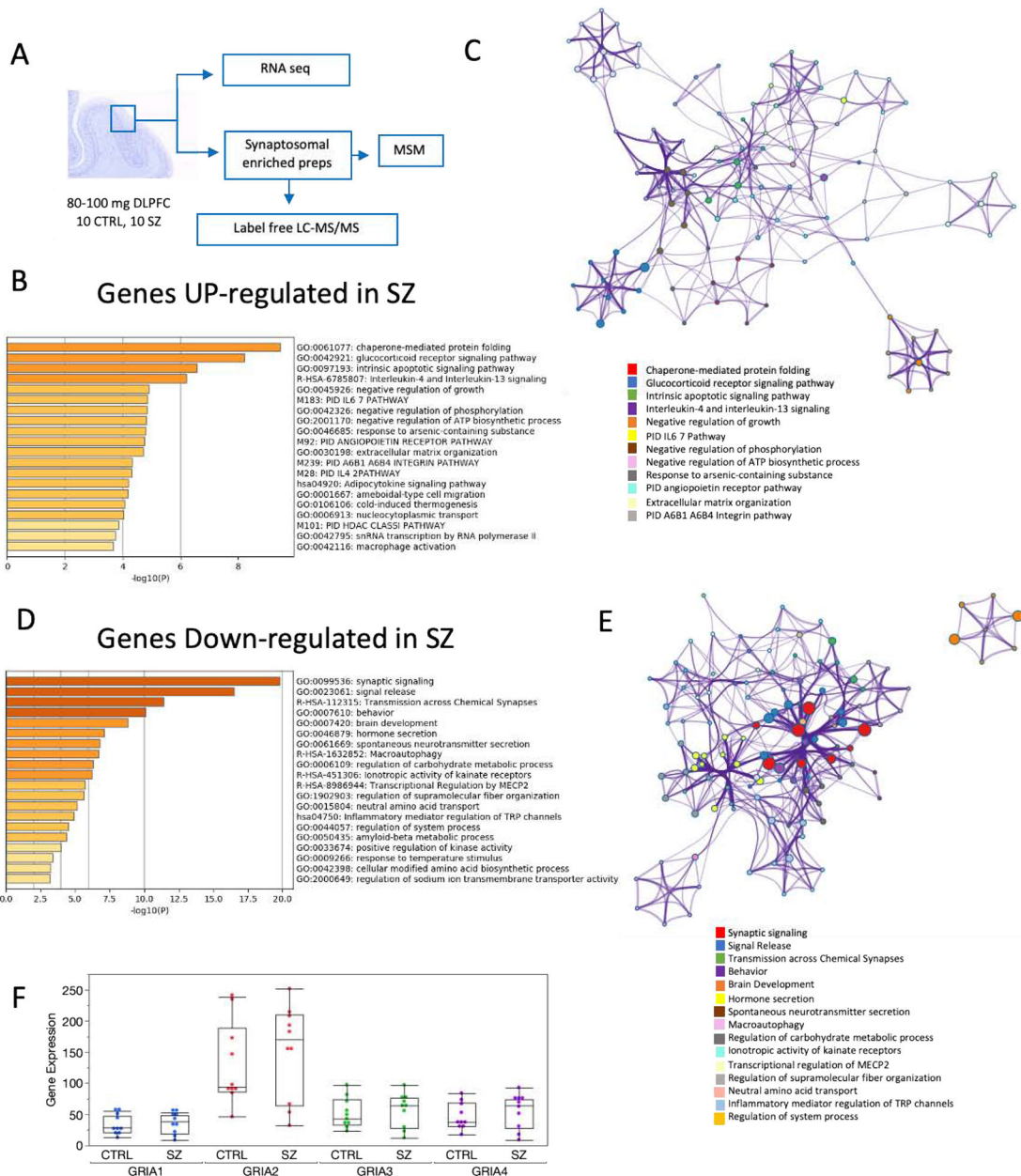
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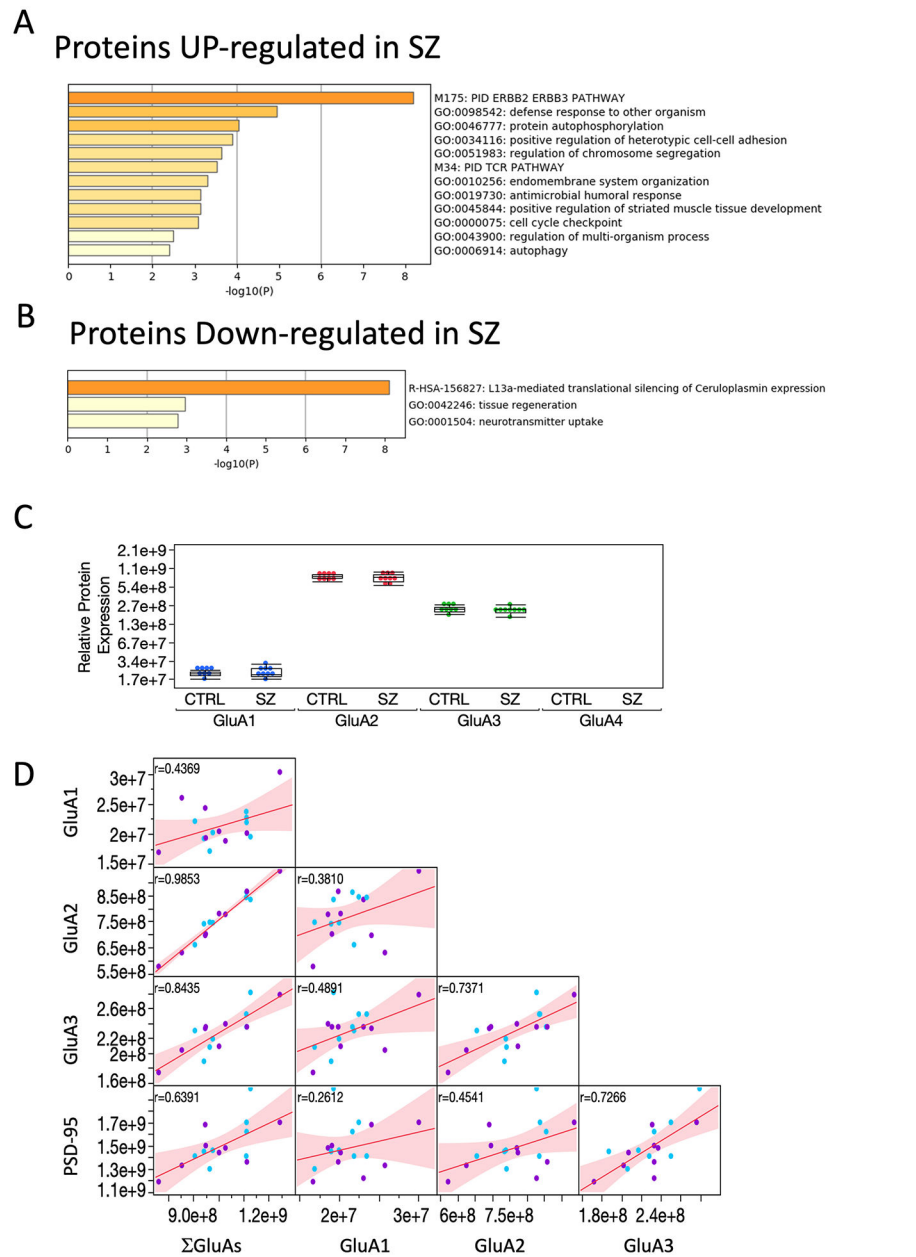
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**Figure 1. Functional enrichment and network interactions of differentially regulated genes in RNA-Seq datasets.**

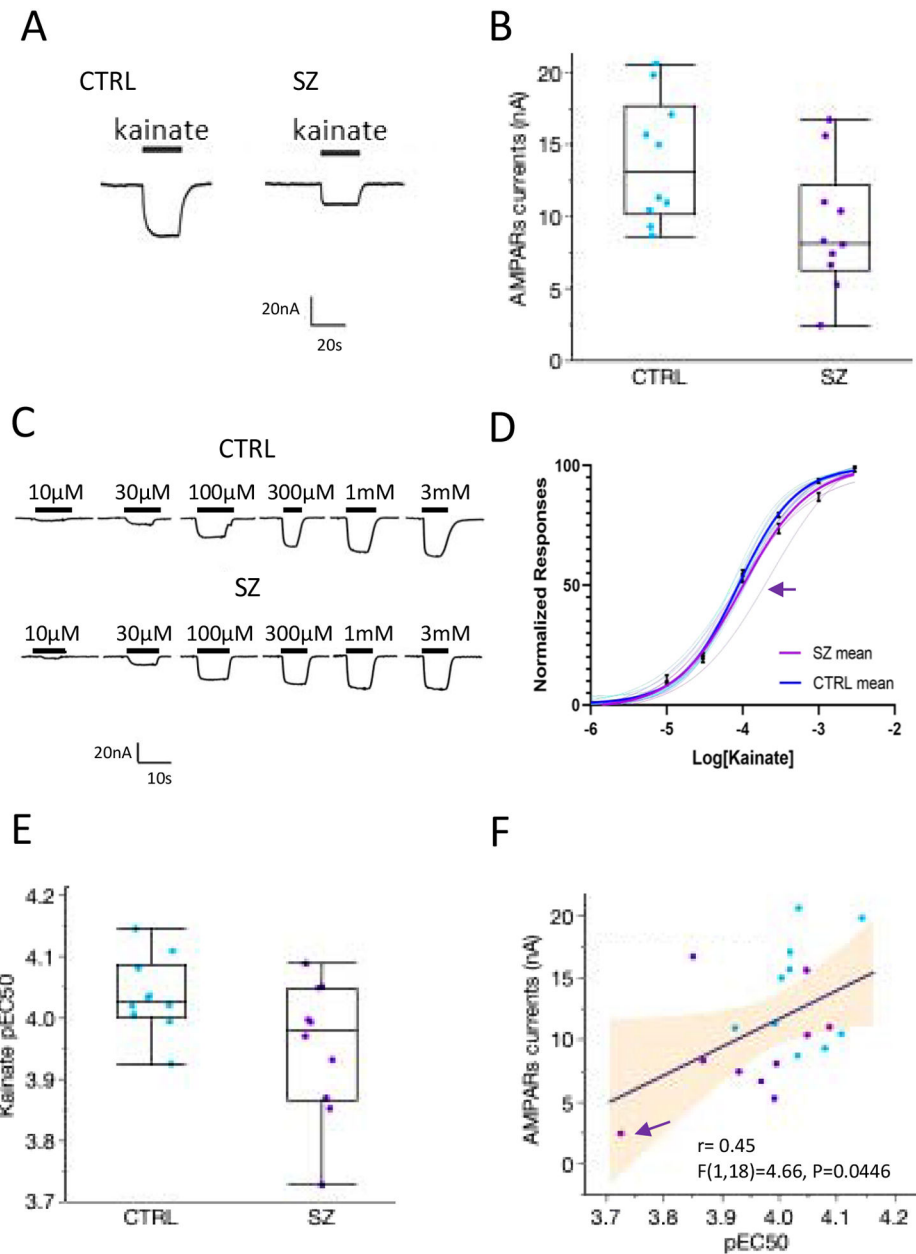
**A.** Diagram illustrating the methodological approach for processing fresh-frozen DLPFC for the different levels of analyses presented in this study. **B. D.** Enriched ontology clusters for genes found to be differentially expressed in SZ at the  $p < 0.01$  level of significance using Metascape (Zhou et al., 2019). **C, E.** Network layout of gene interactions wherein each gene is a circle node. The node size is proportional to the number of input genes that fall in that gene and color is the enrichment identity, as per the color insert at the bottom of the network. **F.** Gene expression levels transcripts per million (TPM) for AMPARs subunits in control and schizophrenia (mean  $\pm$  SEM).



**Figure 2. Functional enrichment of differentially regulated proteins in synaptosomal-enriched preparations.**

**A. B.** Enriched ontology clusters for proteins differentially expressed in SZ at the  $P < 0.05$  level of statistical significance. **C.** Label-free quantitation (LFQ) intensity of expression levels for GluA subunits in control and schizophrenia (mean  $\pm$  SEM). **D.** Multivariate correlation between GluA, their sum that represents the combination of all available AMPARs heteromers, and the postsynaptic density protein PSD-95 that anchors AMPARs to the synaptic cell membrane.

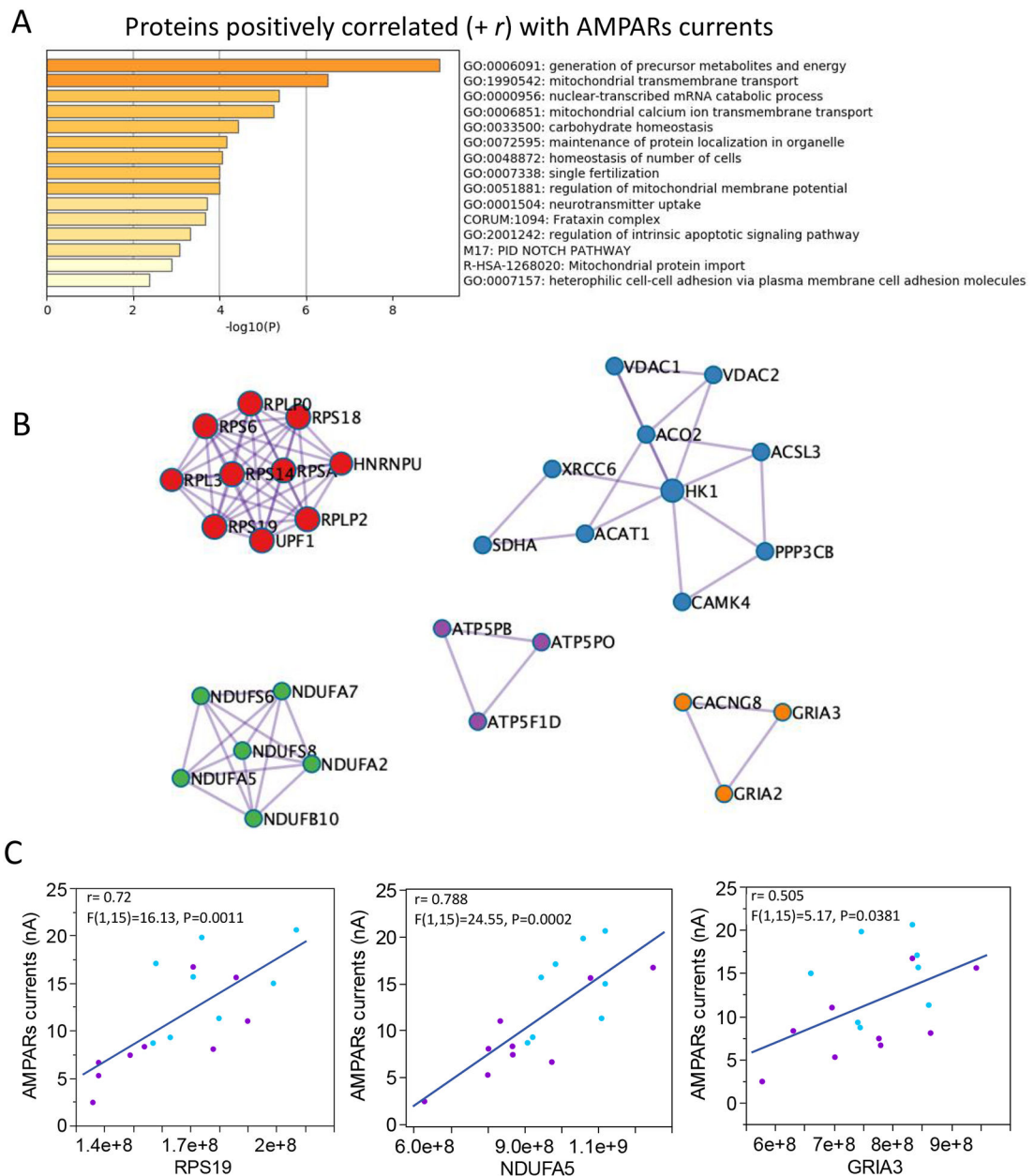




**Figure 3. Altered AMPARs responses in SZ.**

**A.** Functional responses of AMPARs elicited by 100  $\mu$ M kainate of oocytes microtransplanted with DLPFC synaptic membranes from a control and an SZ brain. **B.** AMPARs responses in SZ cases were reduced by 34% ( $n = 10$  CTRL, 10 SZ;  $F(4,15) = 5.85, p = 0.0048$ ; ANCOVA correcting for pH, RIN and the interaction between diagnosis and pH). Each point in the graph represents the mean for each subject ( $n = 10$ –14 oocytes measured per each subject). **C.** Representative concentration-dependent responses to kainate in oocytes microtransplanted with SZ and control synaptic membranes. **D.** Hill equation fits to normalized concentration-response data for each subject (thin lines) and for the group average (thick lines). Each curve for a single subject was done with a minimum of 3 oocytes per subject. Symbols indicate the mean  $\pm$  SEM of responses of each group,

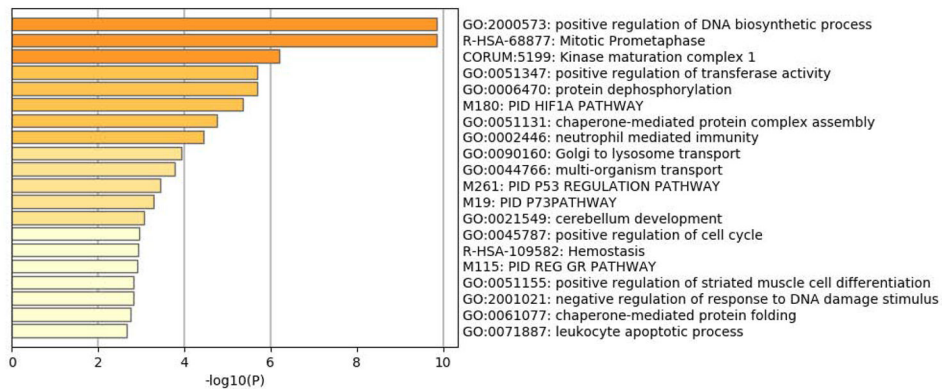
the arrow indicates a subject with reduced affinity respect to the whole cohort. **E. F.** The affinity AMPARs was reduced in SZ controls ( $P = 0.03$ , one sided tail), and the differences in affinity were positively correlated with the amplitude of AMPARs.  $pEC_{50}$  indicates the negative logarithm of the  $EC_{50}$  for each subject. Arrow shows a subject with no detectable CACGN8 protein levels.



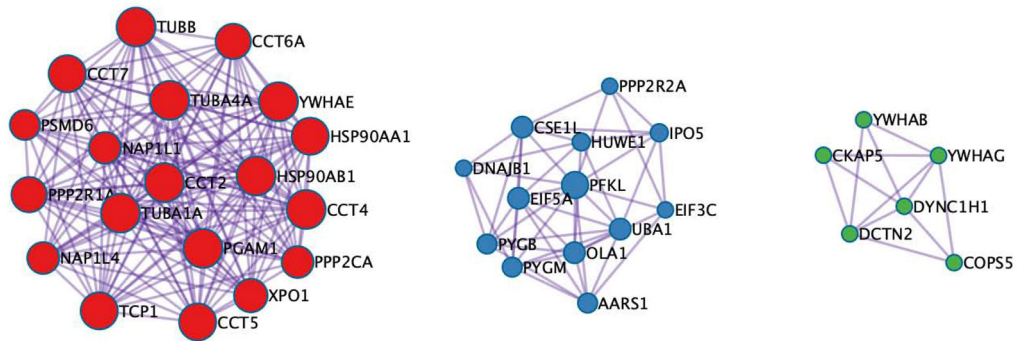
**Figure 4. Functional enrichment and MCODE modules of proteins positively correlated with AMPARs responses.**

**A.** Enriched ontology clusters for proteins differentially expressed in SZ at the  $P < 0.05$  level. **B.** Proteins densely connected identified by molecular complex detection algorithm (MCODE). Protein complexes involved with postsynaptic densities and AMPARs subunits, energy production in mitochondria, and protein translation in ribosomes were positively correlated with AMPARs currents. **C.** Linear correlations between AMPARs currents and proteins representative of MCODE modules shown in B.

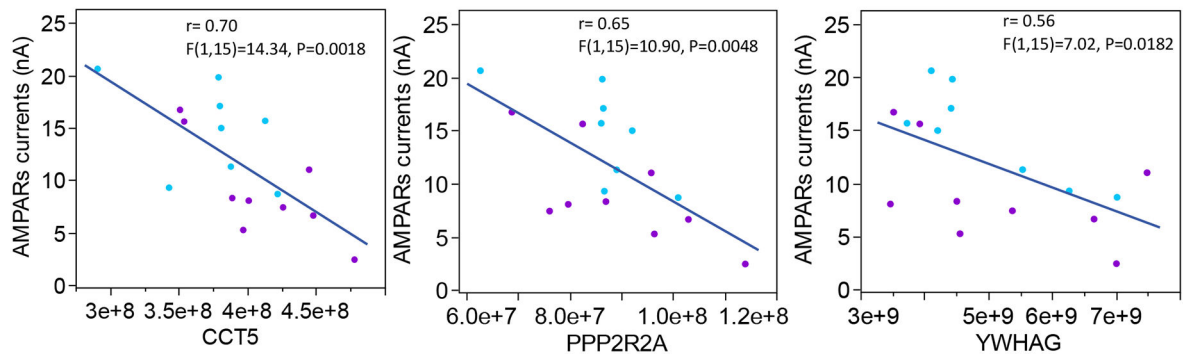
**A** Proteins negatively correlated ( $-r$ ) with AMPARs currents



**B**

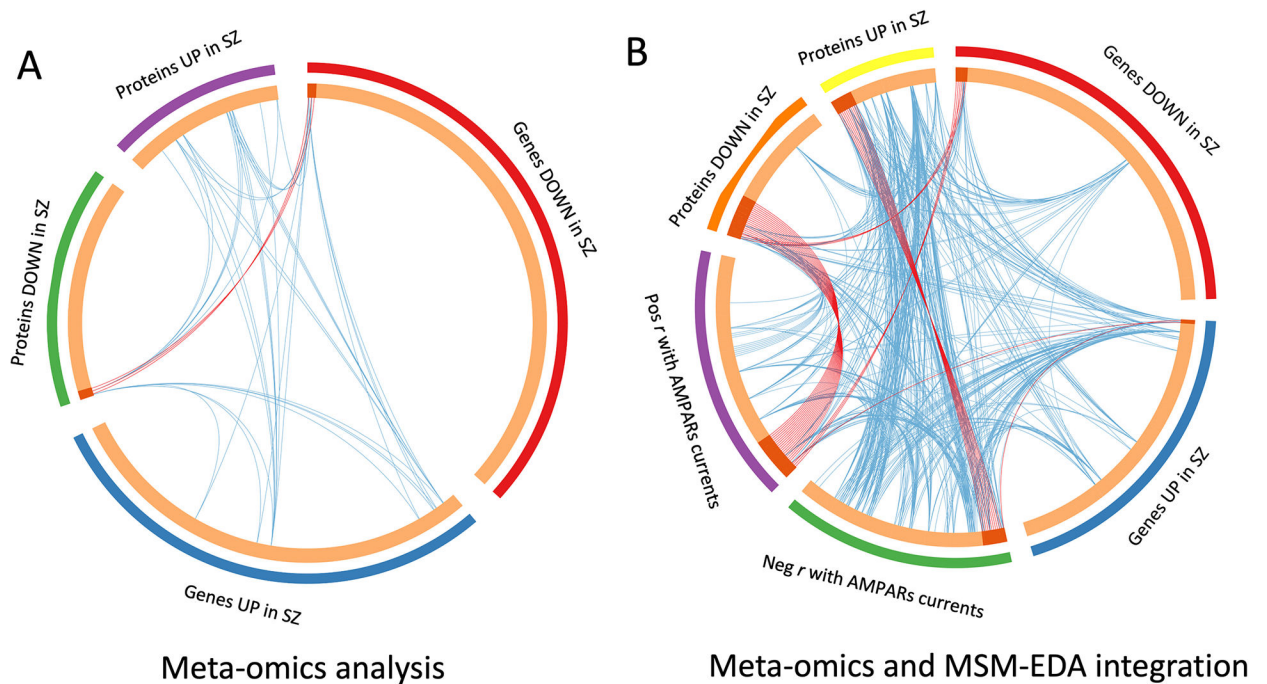


**C**



**Figure 5. Functional enrichment and MCODE modules of proteins negatively correlated with AMPARs responses.**

**A.** Enriched ontology clusters for proteins differentially expressed in SZ at the  $p < 0.05$  level. **B.** Proteins densely connected identified by molecular complex detection algorithm (MCODE). Proteins complexes typically involved in intracellular signaling and DNA biosynthesis were negatively correlated with AMPARs currents. **C.** Linear correlations between AMPARs currents and proteins representative of MCODE modules shown in B.



**Fig. 6. Transcriptomic and synaptic-proteomic convergence on the amplitude of AMPARs currents in SZ.**

A. Circos plot showing gene/protein overlap across different levels of analysis. Outside arches represent the different gene/protein lists generated during this study. Inside arches contain genes/proteins. Dark orange arches indicate the genes/proteins that appear in multiple lists. Purple lines connect the same gene that is shared by multiple lists. Blue links indicate the amount of functional overlap among the input gene lists. B. Similar analysis as in A, including proteins that correlate with the amplitude of AMPARs currents. Notice that genes and proteins that are downregulated in SZ overlap with proteins that are involved in the generation of the amplitude of AMPARs currents measured in microtransplanted membranes. In contrast, genes and proteins upregulated in SZ overlap with proteins that negatively impact the amplitude of the responses measured. Functional overlap is also stronger between lists of genes that negatively impact the amplitude of the responses in SZ, indicating a convergence of mechanisms of synaptic dysfunction.

**Table 1.**Cohort demographics showing mean  $\pm$  standard deviation.

Diagnosis	Gender (F/M)	Age (years)	PMI (hr)	pH	RIN
SZ	3/7	44 $\pm$ 10	22.8 $\pm$ 10	6.6 $\pm$ 0.4	5.4 $\pm$ 1.2
CTRL	3/7	48 $\pm$ 13	20.5 $\pm$ 8	6.2 $\pm$ 0.2*	6.6 $\pm$ 1.3*

\*P<0.05 (Student *t* test), N = 10 each group. PMI, postmortem interval.

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**Table 2.**

Functional convergence between upregulation of MAPK-involved signaling pathways and lower AMPARs responses in SZ

SOURCE	GENE/PROTEIN	NAME
RNA-Seq	CDKN1A	Cyclin Dependent Kinase Inhibitor 1A
RNA-Seq	GADD45B	Growth Arrest and DNA Damage Inducible Beta
RNA-Seq	MAPK9	Mitogen-Activated Protein Kinase 9
MSM-EDA	PLCB1	Phospholipase C Beta 1
LC-MS/MC	PLCD1	Phospholipase C Delta 1
MSM-EDA	PPP3CB	Protein Phosphatase 3 Catalytic Subunit Beta
LC-MS/MC;RNA-Seq	PRKCB	Protein Kinase C Beta
RNA-Seq	PRKCE	Protein Kinase C Epsilon
RNA-Seq	SHC1	SHC Adaptor Protein 1
LC-MS/MC	CAPN1	Calpain 1
LC-MS/MC	CDC42	Cell Division Cycle 42
LC-MS/MC	COL4A2	collagen type IV alpha 2 chain
LC-MS/MC	EIF2S1	Eukaryotic Translation Initiation Factor 2 Subunit Alpha
LC-MS/MC	HRAS	HRas proto-oncogene, GTPase
LC-MS/MC	KRAS	KRAS proto-oncogene, GTPase
LC-MS/MC	MAPK1	mitogen-activated protein kinase 1
LC-MS/MC	MAPK3	mitogen-activated protein kinase 3
RNA-Seq; MSM-EDA	HSPB1	Heat Shock Protein Family B (Small) Member 1
MSM-EDA	MAPK10	mitogen-activated protein kinase 10
MSM-EDA	PPP2CA	Protein Phosphatase 2 Catalytic Subunit Alpha
MSM-EDA	PPP2R1A	Protein Phosphatase 2 Scaffold Subunit Aalpha
MSM-EDA	PTPN11	Protein Tyrosine Phosphatase Non-Receptor Type 11
MSM-EDA	TUBA1A	Tubulin Alpha 1a
MSM-EDA	TUBA4A	Tubulin Alpha 4a
MSM-EDA	YWHAB	tyrosine 3-monooxygenase/tryptophan 5-monooxygenase activation protein beta
LC-MS/MC; MSM-EDA	FGA	Fibrinogen Alpha Chain
LC-MS/MC; MSM-EDA	MAP2K2	mitogen-activated protein kinase kinase 2