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Specific interaction between Sam68 and neuronal mRNAs: implication for the activity-dependent biosynthesis of elongation factor eEF1A.

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Running title: Sam68 and neuronal mRNA translation

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Abstract

In cultured hippocampal neurons and in adult brain, the splicing regulatory protein Sam68 is partially relocated to the somatodendritic domain, and associates with dendritic polysomes. Transfer to the dendrites is activity-dependent. We have investigated the repertoire of neuronal mRNAs to which Sam68 binds *in vivo*. Using co-immunoprecipitation and microarray screening techniques, Sam68 was found to associate with a number of plasticity-related mRNA species, including Eef1a1, an activity-responsive mRNA coding for translation elongation factor eEF1A. In cortical neuronal cultures, translation of the Eef1a1 mRNA was strongly induced by neuronal depolarisation, and correlated with enhanced association of Sam68 with polysomal mRNAs. The possible function of Sam68 in Eef1a1 mRNA utilization was studied by expressing a dominant-negative, cytoplasmic Sam68 mutant (GFP-Sam68 Δ C) in cultured hippocampal neurons. The level of eEF1A was lower in neurons expressing GFP-Sam68 Δ C than in control neurons, supporting the proposal that endogenous Sam68 may contribute to the translational efficiency of the Eef1a1 mRNA. These findings are discussed in the light of the complex, potentially crucial regulation of eEF1A biosynthesis during long-term synaptic change.

Keywords

Dendritic mRNA, activity-dependent translation, RNA-binding protein, plasticity

Introduction

The translation of neuronal mRNAs is regulated by their association with specific RNA-binding proteins, of which an increasing variety have been identified (Bramham and Wells 2007; Kiebler and Bassell 2006). In the well-studied cases of CPEB, FMRP and ZBP-1, the proteins maintain the target mRNAs in a dormant state during transport or storage; this inhibitory function is thought to be suppressed by activity-dependent phosphorylation events, e.g. in the vicinity of synapses (Bagni and Greenough 2005; Huang et al. 2002; Huttenlocher et al. 2005). In addition to these predominantly cytoplasmic proteins, recent evidence from non-neuronal systems indicates that nuclear proteins of the splicing machinery may also contribute to the control of translation. For example, SR splicing factors continuously shuttle between nucleus and cytoplasm, where they can bind to mature mRNAs and promote translation (Sanford et al. 2004). Translational efficiency can also be increased by splicing proteins which are deposited onto RNA substrates during the splicing process itself, and which remain bound to the mRNA after nuclear processing and export have been completed (Kunz et al. 2006; Le Hir et al. 2000; Wiegand et al. 2003). Interestingly, a diversity of splicing factors have been found to exist in neuronal dendrites (Glanzer et al. 2005). One of those is Sam68, a ubiquitous RNA-binding protein implicated in signal-dependent splice site selection, and which can also function as an adaptor and substrate for Src family kinases (Lukong and Richard 2003; Matter et al. 2002; Paronetto et al. 2007). In cultured cell lines, Sam68 is a nuclear, non-shuttling protein; however, it is required to enhance the nuclear export and / or translation of certain retroviral mRNAs, indicating a potential interaction with translation-activating factors (McLaren et al. 2004; Modem et al. 2005). Beside viral mRNAs, Sam68 can bind to a select population of mature cellular mRNAs derived from cultured HeLa cells (Itoh et al. 2002); the functional outcome of these binding interactions has not been determined.

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We previously reported that Sam68 was among the proteins associated with neuronal mRNAs (Ben Fredj et al. 2004; Grange et al. 2004). In adult cortical and hippocampal tissue, biochemical fractionation and immuno-electron microscopy indicated that Sam68 molecules were associated with dendritic polysomes, suggesting a possible positive role in translational regulation (Grange et al. 2004; Khandjian et al. 2004). In cultured hippocampal neurons, a minor but detectable fraction of endogenous or GFP-tagged Sam68 molecules were found to occur in the soma and dendrites, concentrated in generally immobile, RNA-rich granules. Following prolonged membrane depolarisation by elevated KCl, the proportion of Sam68 in the cytoplasm was strongly increased, due to massive migration of the protein to these somatodendritic sites (Ben Fredj et al. 2004). These results suggest that Sam68 may play a part in the regulation of those mRNAs which become translationally activated in response to electrical activity. To test this hypothesis, we have investigated the identity of mRNA species that were bound to Sam68 in cortical neurons in vivo. We found that one major target was Eef1a1, an mRNA known to be activated during long-term synaptic change in the hippocampus. In cultured neurons, depolarisation enhanced the association of both Sam68 and the Eef1a1 mRNA with polyribosomes, and increased the amount of its product, the eEF1A protein. Expression of a dominant-negative Sam68 mutant confined to the cytoplasm was able to lower the amount of eEF1A, supporting the proposal that Sam68 promotes the translation of neuronal Eef1a1.

Materials and Methods

Immunoprecipitation of Sam68-associated RNA and preparation of amplified RNA probes

Cortical Sam68-RNA complexes were prepared as previously described (Grange et al. 2004). Briefly, the cortex from one adult rat was dissected and lysed at 4°C in polysome dissociation buffer [10 mM Tris-HCl, pH 7.5, 50 mM KCl, 1 mM sodium orthovanadate, 2 mM dithiothreitol (DTT), 100 nM microcystine-LR (Sigma), protease inhibitors (Complete; Roche Molecular)] containing 0.5% Triton X-100, 160 mM sucrose, 30 mM EDTA, pH 8.0 and 400 U/ml RNasin (Promega, Inc., Madison, WI). After clarification by centrifugation at 82,000g for 20 min at 4°C, the lysate was precleared with protein G–Sepharose beads (GE Healthcare), then immunoprecipitated with affinity-purified anti-Sam68 IgG (AD-1), or with preimmune protein A-purified IgG from the same rabbit. Immune complexes were collected onto protein G –Sepharose beads, washed, and processed for RNA purification using the “RNEasy” kit (Qiagen France, Courtaboeuf). The immunoprecipitated polyA+ RNAs were linearly amplified by cDNA synthesis and T7 polymerase transcription (Gelder et al. 1990), using the MessageAmp™ aRNA Kit (Ambion). According to whether the template RNA originated in the specific immunoprecipitate or the preimmune control, either Cyanine 3™ (Cy3) or Cyanine 5™ (Cy5)-labeled UTP was incorporated during transcript synthesis, as per manufacturer’s instructions. Transcripts amplified from the specific immunoprecipitate and the corresponding control were mixed in equal amounts and used to probe the 15000 cDNAs arrayed on the NIA micro-array (Tanaka et al. 2000). To control for possible biasing by differential incorporation of Cy3 and Cy5, the experiment was repeated with swapping the fluorophores between “specific” and “background” transcripts.

Quantitation of hybridization

After hybridization and washing, digital images of the micro-array at 635 nm and 532 nm were acquired by laser scanning confocal microscopy. Fluorescent spots were delineated and

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quantitated with GenePix software (Axon Instruments). Raw fluorescence values (F) in each channel were converted into their Log_2 . After Cy3 labelling of background RNAs, immunoprecipitation of any given mRNA was considered specific when the value in the Cy5 channel exceeded that in the Cy3 channel by at least 4 Log_2 units (a 16-fold difference). When background RNAs were labelled with Cy5, the cut-off was brought down to 0.5 Log_2 unit (an 1.4-fold difference). This differential cut-off was applied to compensate for the systematic bias in labelling efficiency in favour of Cy5. Note that the requirement for higher Cy3 labelling of specifically immunoprecipitated RNAs is conservative and likely excludes a number of Sam68-associated mRNAs. Z scores were calculated as $(\text{Log}_2F - \langle \text{Log}_2F \rangle) / \sigma_{\text{Log}_2F}$, where $\langle \text{Log}_2F \rangle$ is the mean and σ_{Log_2F} is the standard deviation, both calculated over the entire population of fluorescence values in the relevant channel (Mayne et al. 2001). Reproducibility of the overall population of z scores was measured by simple linear regression between experiments. 95% confidence intervals were calculated for the z scores of Table Ia using t statistics ($n=3$), and individual mRNAs with lower confidence limit $z < -1.0$ were rejected for inclusion in the Table; more than 80% of these scores had lower limit above zero. Frequencies of GO descriptors were computed from ~4000 functionally described genes of the microarray (available through <http://lgsun.grc.nia.nih.gov/cDNA/NIA-CloneSet-GoTermFinder.html>).

Polymerase chain reactions

Immunoprecipitated or total RNA was reverse-transcribed with the universal primer GGAATTC(T)₁₇, and the resulting cDNA was amplified by using the gene-specific primer pairs listed in Table II. For classical PCR, after 2 min at 95°C, 30 cycles were performed (15 s at 95°C, 30 s annealing, and 30 s at 72°C for each cycle). For quantitative PCR by real-time fluorometry (Light Cycler, Roche Molecular Systems), DNA was denatured at 95°C for 5 min, then subjected to ~35-40 amplification cycles (15 s at 95°C, 5 s annealing, 15 s at 72°C,

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all temperature changes at 20°C/s). Annealing took place at 55°C except for Arc (57°C). The relative amounts of specific DNA template were determined by counting the number of cycles needed to reach the inflection point (“crossing point” at which $d^2[\text{DNA}]/dt^2=0$) of the DNA accumulation curve (typically between 25 and 32), and standardized by relating to a scale generated with known amounts of total cDNA. The specificity of the amplification product was confirmed by checking its melting curve with the Light Cycler, and its size on an agarose gel.

Immunoprecipitation of RNA-mycSam68 complex from HEK cells

HEK-293 cells were grown in Dulbecco’s modified Minimum Essential Medium, containing 10% fetal calf serum (Invitrogen). The myc-Sam68 expressing plasmid was a kind gift from I. Barlat and B. Tocqué (at the time at Aventis Pharma, Vitry, France). One 6 cm dish of cells was transfected with either myc-Sam68 or with control vector (containing a CMV promoter). 48 h later, cells were lysed in polysome dissociation buffer and, after clarifying by centrifugation, the lysate was immunoprecipitated with 2 µg anti-myc monoclonal antibody (12CA5, Santa Cruz) and protein G-Sepharose beads, as described for brain lysates. Total RNA was isolated and amplified by RT-PCR as described above.

Neuronal cultures and transfections

Primary cultures of rat hippocampal neurons were prepared from fetal (E19) hippocampi and seeded at a density of 6000 cells / cm² onto poly-L-lysine-coated glass coverslips (Marienfeld, Germany), adhering to the procedure described by Goslin and Banker (Banker et al. 1998), except that the culture medium was Neurobasal with B27 supplement (Invitrogen). Housing and handling of animals complied with the regulations issued by the European Union and INSERM and were approved by the local veterinary authorities. The GFP-Sam68 (WT) and GFP-Sam68ΔC plasmids were kind gifts from S. Richard (McGill University, Montréal, Canada). The mRFP1 plasmid (expressing monomeric red fluorescent protein) was a kind gift

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from R.Y. Tsien (UCSD, San Diego, CA, USA). Transfection was performed at 9 DIV by calcium phosphate coprecipitation, exactly according to (Goetze et al. 2004). We used a mix of 2.5 μg GFP-Sam68 and 0.5 μg mRFP1 plasmids per coverslip. Expression of the GFP-Sam68 and GFP-Sam68 ΔC plasmid was stable for the next 10 days. Neurons were fixed at 14-19 DIV and processed for immunofluorescent staining.

Primary cortical neurons were prepared from E16 fetal cortices, using essentially the same procedure as for hippocampal neurons. The cortical cells were seeded at a density of 10^5 cells / cm^2 onto poly-L-lysine coated 10-cm dishes (Nunc). Neurons at 9 DIV were treated for 3 h with either depolarizing medium (containing, in mM: KCl, 25; CaCl₂, 1.8; MgCl₂, 0.8; NaCl, 100; NaHCO₃, 26; NaH₂PO₄, 1; glucose, 30; Hepes-NaOH, 15) or control solution (same except for KCl, 5; NaCl, 120).

Sedimentation of polysomes on sucrose gradients

RNA-protein complexes were fractionated as described by (Krichevsky and Kosik 2001). Briefly, cortical neurons were lysed in low salt buffer [20 mM Tris-HCl (pH 7.5), 10 mM NaCl, 3 mM MgCl₂, 1 mM RNasin, 1 mM dithiothreitol, 0.3% Triton X-100, and 0.05 M sucrose] containing protease inhibitors (EDTA-free Complete, Roche Molecular), 1 mM vanadate and 100 nM microcystin LR. Nuclei and the majority of mitochondria were sedimented by centrifugation for 10 min at $10,000 \times g$ at 4°C. The NaCl and MgCl₂ concentrations in the cytoplasmic extracts (supernatants) were adjusted to 170 and 13 mM, respectively. Linear sucrose gradients (15%–45% w/w in 25 mM Tris-HCl [pH 7.5], 25 mM NaCl, 5 mM MgCl₂) were prepared using an AutoDensi Flow apparatus (Büchner). Cytoplasmic extracts (1.2 ml) were overlaid onto 10.6 ml gradients and centrifuged at 32,000 rpm for 2.5 hr at 4°C in a SW41 rotor. Fractions of 1 ml volume were collected with on-line UV absorbance measurement at 254 nm (GE Healthcare Akta Prime). For Western immunoblotting, the protein content of the fractions was concentrated by trichloroacetic acid

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3 precipitation, redissolved in Laemmli buffer, and analysed on 8% polyacrylamide / SDS gels.
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5 For RT-PCR, RNA was extracted by treating the relevant fractions with Trizol reagent
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7 (Invitrogen), precipitated with ethanol, and subjected to RT-PCR with the same primers as
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9 above.
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12 *Detection of eEF1A protein increase by immunoblotting*

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14 Cortical neurons were treated with control or depolarising medium as above, then lysed in ice-
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16 cold RIPA buffer (20 mM Tris-HCl [pH 7.5], 150 mM NaCl, 1 mM EDTA, 1 % Triton X-
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18 100, 0.5 % Na deoxycholate, 0.1 % SDS) containing protease inhibitors (EDTA-free
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20 Complete, Roche Molecular). After clarification by centrifugation (15 min at 12,000 x g),
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22 equal protein amounts of each lysate were analysed by 10 % polyacrylamide / SDS gel
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24 electrophoresis and immunoblotting, first with anti-eEF1A monoclonal antibody (UBI) then
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26 with anti-actin polyclonal antibody (Santa Cruz). Immunoreactive bands were revealed by
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28 chemiluminescence (Pierce Femto) and autoluminography. The films were digitized by
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30 scanning and the integrated intensity of the bands was determined using Metamorph software
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32 (Universal Imaging). Immunoblotting and exposure conditions were calibrated so that the
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34 integrated intensity of the bands was in the linear range with respect to protein amount.
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37 *Immunofluorescence of eEF1A in hippocampal neurons*

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39 Neurons were fixed with 4% PFA containing 4% sucrose (30 min, 4°C); permeabilized with
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41 0.2% Triton X-100; blocked with 10% preimmune goat serum in PBS (25°C, 1 h); and stained
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43 with 1 µg/ml anti-eEF1A monoclonal antibody (UBI) in PBS containing 3% goat serum
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45 (25°C, 2 h), then with Alexa350TM-conjugated secondary antibody (Molecular Probes). After
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47 washing, coverslips were mounted with Mowiol and observed on a Zeiss microscope fitted
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49 with an epifluorescence setup and a cooled CCD camera (Zeiss Axiocam), or with a confocal
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51 laser scanning microscope (Zeiss LSM 510).
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For quantitation of eEF1A fluorescence, fields containing transfected neurons were imaged at low magnification (20x). Imaging conditions were identical for all fields. Images were processed with Metamorph (Universal Imaging). For each image, a background image was produced with a low-pass filter and subtracted so as to remove low-frequency fluorescence noise due e.g. to local unevenness in illumination. Next, a lower fluorescence threshold was set, which allowed automatic delineation and analysis of the fluorescent objects (cells), using Integrated Morphometric Analysis. The same threshold was used throughout the entire analysis. The relative eEF1A fluorescence of each transfected neuron i was calculated according to the equation: $IF_{\text{transf},i} = e\text{EF1A}_{\text{transf},i} / \langle e\text{EF1A} \rangle_{\text{nontransf}}$

where $e\text{EF1A}_{\text{transf},i}$ is the mean grey level per pixel within the transfected neuron, and $\langle e\text{EF1A} \rangle_{\text{nontransf}}$ is the average (over 15 surrounding neurons) of the mean grey level per pixel within non-transfected neurons. These $IF_{\text{transf},i}$ values were averaged over the entire set of transfected neurons, to yield the values of Fig. 4E.

Immunoprecipitation of Sam68-NMDA receptor complexes

Rat cortical synaptic membranes were prepared by differential centrifugation as described by Grange et al. (2004), and resuspended in ice-cold buffer containing 4mM HEPES (pH 7.4), 1mM preboiled sodium orthovanadate, 1mM dithiothreitol, 100 nM microcystin (Sigma), and a mix of protease inhibitors ("Complete", Roche). An aliquot of suspension (530 μl , 4.5 $\mu\text{g}/\mu\text{l}$ total protein) was solubilized by adding 1:9 vol. of detergent solution (10% sodium deoxycholate in 0.5M Tris, pH 9.0) and gently agitating for 20 min. Triton X-100 (0.1% final concn.) was then added, and the extract was dialyzed overnight against immunoprecipitation buffer (10 mM Tris-HCl, pH 7.4, 100 mM NaCl, 0.1 mM sodium orthovanadate, 5 mM EDTA, 1 mM dithiothreitol, 0.1 % Triton X-100, 1 nM microcystin, and protease inhibitors). The dialyzed extract was centrifuged for 30 min at 10^5 g in a TL-100 tabletop ultracentrifuge (Beckman), and the supernatant (300 μg protein diluted to 500 μl buffer per IP) was used for

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3 immunoprecipitation for 3 h with 10 μ g affinity-purified anti-Sam68 polyclonal antibody, or
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6 preimmune IgG purified from the same rabbit, or with anti-NR1 monoclonal antibody
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8 (Pharmingen). Immunoprecipitates were collected onto protein G-Sepharose beads and
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10 analysed by SDS-PAGE (8% polyacrylamide gel) and Western immunoblotting. Antibodies
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12 used for immunoblotting were purchased from UBI (anti-PSD95 monoclonal antibody) and
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14 Chemicon (anti-GluR2/3 rabbit polyclonal antibody). The bands were revealed by
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16 chemiluminescence (Pierce Femto).
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For Peer Review

Results

Identification of Eef1a1 in the repertoire of Sam68-associated mRNAs

Following an approach initiated by Tenenbaum et al. (2000), we decided to use a combination of co-immunoprecipitation and array screening to probe the repertoire of Sam68-associated mRNAs in cortical neurons. Sam68-mRNA complexes were immunoprecipitated from a rat cortical cytoplasmic extract with an anti-Sam68 antibody, as previously reported (Ben Fredj et al. 2004). A parallel immunoprecipitation was performed with preimmune antibody. The polyA⁺ RNAs associated with the Sam68 and control immunoprecipitates were linearly amplified by cDNA synthesis and T7 polymerase-mediated transcription, as initially described by Eberwine and coworkers (Gelder et al. 1990), and the amplified transcripts were fluorescently labeled with Cy3 or Cy5, respectively. The two transcript populations were then mixed and hybridized with the ~15,000 cDNAs of the NIA “15K” clone set arrayed onto a glass slide (Tanaka et al. 2000), and for each cDNA the fluorescence values at both wavelengths were measured and compared (Fig. 1 A, B). A given mRNA was considered to be specifically immunoprecipitated when the fluorescence level was consistently higher in the channel corresponding to the immunospecific than in the background precipitation. To compare mRNA abundance measurements across several experiments, normalized measures of the relative abundance of each mRNA in the immunoprecipitated population were calculated by converting the fluorescence values into z scores (Lopez de Silanes et al. 2004; Mayne et al. 2001; see Materials and Methods). Duplicate assays of the same RNA population showed that z scores were highly reproducible through the entire set of ~15,000 genes, except for a limited number of low abundance mRNAs, indicating that abundance measurements were not grossly perturbed by fluctuations in the hybridization and detection procedure (Fig. 1C). When distinct experiments with different brain samples were compared, fluctuations in individual z scores were larger, likely due to random differences in the hydrolysis or

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dissociation rates of individual mRNAs during immunoprecipitation. The correlation between samples remained sufficient to ensure that the species scoring highest ($z > 1.5$) in one trial retained a 85% probability of scoring positive ($z > 0$) in a second trial (Fig. 1D). To obtain reliable identification of the most abundant mRNAs in the immunoprecipitate, z scores were measured in three independent experiments, mRNAs were ranked by decreasing mean z score, and some species with unacceptably large standard deviations were excluded. We also discarded a few arrayed sequences for which no cognate mRNAs had been identified. Table Ia shows those specifically immunoprecipitated mRNA species for which the mean relative abundance was highest after applying this screen .

The dual-channel procedure was then used to compare the relative abundances of mRNA species in the immunoprecipitated vs. total polyA+ populations. The total population of polyA+ RNAs present in the input extract (before immunoprecipitation) was amplified and the resulting Cy5-labeled transcripts were mixed with an aliquot of Cy3-labeled transcripts derived from the Sam68 immunoprecipitate. After hybridization to the array, z scores were calculated for each channel. The species of Table Ia scored 0.6 to 1.0 units higher in the immunoprecipitated than in the total population, consistent with their enrichment.

There was a discernible pattern in the biological functions of these mRNAs (Fig. 1E). When assessed by Gene Ontology annotation for generic cellular process or component, more than 20% of the mRNAs (9 transcripts) were involved in intracellular or receptor-associated signal transduction, as they coded for e.g. receptor subunits, protein kinases, protein phosphatases, and associated regulatory factors. Another 20% of the mRNAs coded for proteins involved in protein transport, either as regulators of vesicular traffic, or of microtubule-dependent motion. These proportions are significantly higher than what would be expected based on the overall distribution of biological functions among the genes represented on the micro-array (Fig. 1E). By comparison, transcription regulators appeared to be under-represented among the major

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Sam68-associated mRNAs. Assessment of the literature further indicates that some of the proteins of Table I contribute to exocytic and endosomal traffic (Arf-1, BRI-3, vacuolar ATPase) or to cytoskeletal regulation (stathmin) even though this is not apparent from their GO function label. All of these processes are heavily involved in the local response to stimuli which affect neuronal homeostasis and plasticity. Indeed, more than one third of the genes of Table Ia code for proteins directly involved in synaptic plasticity, connectivity development, or neurodegenerative disease (Table Ib).

The β -actin mRNA, which was detected among the most abundant species in our screen, had formerly been detected by RT-PCR in immunoprecipitates of brain Sam68 (Ben Fredj et al. 2004), and was known as a specific target for recombinant Sam68 (Itoh et al. 2002). This provided a positive control for the immunoprecipitation procedure. The Gephyrin mRNA provided a negative control, as it was found neither by RT-PCR, nor in the micro-array analysis (data not shown and (Grange et al. 2004)).

The most enriched mRNA, Eef1a1, coded for the ubiquitous isoform of translation elongation factor eEF1A, an important player in both protein synthesis and long-term synaptic change (see Discussion). The z score value of the Eef1a1 fluorescence indicates that the representation of Eef1a1 among immunoprecipitated mRNAs is significantly higher than average ($p=0.004$).

RT-PCR with gene-specific primer pairs (see Table II) was used to confirm the specific coimmunoprecipitation of a range of mRNAs detected in the micro-array analysis, either among the more enriched species listed in Table Ia, or among mRNAs with lower relative abundance. In addition to β -actin, confirmed species included Eef1a1, Calmodulin I and Uchl1 (Fig. 2). Interestingly, the Arc / Arg 3.1 mRNA was present among the less abundant mRNAs in the micro-array analysis, and could also be detected by RT-PCR (see Fig. 2A). Since both Arc and Eef1a1 are dendritic, activity-responsive mRNAs, this raises the

possibility that upon exit to the somatodendritic compartment Sam68 may play a role in activity-induced translation of neuronal mRNAs.

As Eef1a1 was the most enriched of Sam68-associated mRNAs in our assay, and since its translational regulation by synaptic inputs is well-documented (see below), we decided to investigate further its relationship with Sam68. To reconstitute the interaction between Eef1a1 mRNA and Sam68 in a heterologous system, myc-tagged Sam68 or a control plasmid were transiently expressed in HEK 293 cells; cytoplasmic extracts prepared from the transfected cells were immunoprecipitated with anti-myc antibody, and the immunoprecipitates were probed by RT-PCR using the Eef1a1-specific primer pair. Endogenous Eef1a1 transcripts were detected in anti-myc immunoprecipitates from myc-Sam68-expressing cells but not from control cells (Fig. 2C, lanes 1-6). This result was reproduced by quantitative RT-PCR (Fig. 2C). Thus, the Eef1a1 mRNA associates with *bona fide*, overexpressed Sam68. By contrast, the β -tubulin mRNA was absent from myc-Sam68 immunoprecipitates (Fig. 2B, lanes 7-12), supporting the mRNA sequence specificity of binding.

Identification of Eef1a1 as a translationally activated mRNA in depolarized neurons

The Eef1a1 mRNA contains a 5' oligopyrimidine tract and belongs to the general group of signal-responsive mRNAs encoding components of the translation apparatus (Petroulakis and Wang 2002). Moreover, translation of Eef1a1 is stimulated during synaptic change ((Tsokas et al. 2005). To determine whether Eef1a1 might belong to the population of translationally activated mRNAs in our system of in vitro depolarised neurons, primary cultures of cortical neurons were subjected to control treatment or prolonged depolarisation, lysed under non-denaturing conditions, and the post-nuclear supernatants were fractionated on sucrose density gradients, as described by (Krichevsky and Kosik 2001). Fractions were analysed for their RNA content by continuous monitoring of UV absorbance during collection. Fractions corresponding to the free mRNPs, 80S monosomes, polyribosomes, and RNA granules

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(migrating down to the bottom of the gradient) were analysed by RT-PCR with Eef1a1-specific primers. In non-stimulated neurons, the Eef1a1 mRNA was associated to the same extent with free mRNPs and with polysomes. Upon depolarisation, Eef1a1 became nearly undetectable in the free mRNP fraction, and strongly accumulated in the polysome fraction (Fig.3A). These data indicate that Eef1a1 mRNA translation is activated by depolarisation. Consistent with this result, depolarisation of cultured cortical neurons induced a significant increase in the total level of eEF1A protein, as detected by Western immunoblotting (Fig. 3B).

Depolarisation-inducible association of Sam68 with polysomes

We previously reported that Sam68 was associated with polyribosomes in cytoplasmic extracts derived from adult rat cortical tissue (Grange et al. 2004). To confirm that a similar association occurred in cultured cortical neurons, and to determine whether the interaction could be regulated by depolarisation, extracts were prepared and fractionated exactly as in Fig. 3A. Fractions were then analysed by Western immunoblotting with anti-Sam68 antibody. In extracts from control, quiescent neurons, Sam68 mostly comigrated with free mRNPs; a minor fraction sedimented in the RNA granule fraction (Fig. 3C, top). Following depolarisation, Sam68 accumulated in fractions containing polysomes and granules, even though the total amount of RNA in those fractions was slightly lower than in the control condition (Fig. 3, bottom). This result is consistent with the depolarisation-induced accumulation of Sam68 fluorescence in RNA-enriched dendritic clusters, which we have previously reported (Ben Fredj et al. 2004). Thus, in depolarised neurons, the increase in Eef1a1 translation correlates with the loading of Sam68 onto translated mRNAs.

Effect of dominant-negative Sam68 on neuronal eEF1A expression

The association of Sam68 to actively translated Eef1a1 mRNA suggests that Sam68 might regulate the synthesis of eEF1A. To investigate this point, we assayed the level of eEF1A

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3 immunoreactivity in cultured hippocampal neurons that expressed GFP-tagged derivatives of
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5 either wild-type Sam68 or Sam68 Δ C, a C-terminally truncated Sam68 mutant. The latter
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7 mutant retains a significant affinity for specific RNA sequences, but is deprived from
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9 tyrosine-rich and proline-rich motifs involved in protein-protein interactions with SH2 and
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11 SH3-containing ligands, and also from a C-terminal (nonclassical) nuclear localisation signal
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13 (Fig. 4F). In cultured cell lines, Sam68 Δ C is constitutively cytoplasmic and displays
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15 transdominant-negative activity towards endogenous Sam68, when coexpressed with
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17 retroviral (HIV-1) mRNAs that require Sam68 for efficient utilization (Reddy et al. 1999;
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19 Soros et al. 2001). Sam68 Δ C is specifically recruited to these viral mRNAs once they have
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21 reached the cytoplasm, and appears to block their normal interaction with the translation
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23 apparatus (Soros et al. 2001). We reasoned that expression of Sam68 Δ C in neurons should
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25 similarly allow us to probe the contribution of endogenous Sam68 to the translational activity
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27 of target neuronal mRNAs, as distinct from its role in nuclear mRNA processing. In
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29 transfected neurons, the GFP-Sam68 Δ C protein was mostly excluded from the neuronal
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31 nucleus and concentrated in clusters within the cell body; it was also detectable in dendritic
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33 shafts (Fig. 4, A2, B). The relative expression level of endogenous eEF1A in transfected and
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35 non-transfected neurons was evaluated by immunofluorescent staining with an anti-eEF1A
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37 monoclonal antibody. Among the neurons of a field, those expressing GFP-Sam68 Δ C
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39 typically had a lower level of eEF1A immunoreactivity than surrounding, nontransfected
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41 neurons (Fig. 4, A1, C1). This was not the case in neurons that had been transfected with
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43 GFP-tagged wild-type Sam68 (Fig. 4, D1). Neurons expressing GFP-Sam68 Δ C maintained a
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45 normal morphology (Fig. 4, A1, C3), indicating that the decrease in eEF1A did not arise from
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47 some general toxicity of the construct. The effect of GFP-Sam68 Δ C expression on
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49 endogenous eEF1A immunofluorescence was quantitated by using as internal reference value
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51 the average immunofluorescence of nontransfected neurons surrounding the transfected cells
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(see Materials and Methods). This quantitation procedure indicated that GFP-Sam68 Δ C reproducibly lowered the level of eEF1A immunofluorescence, whereas GFP-Sam68 did not (Fig. 4E). As a control, GFP-Sam68 Δ C did not affect the abundance of coexpressed mRFP. Since GFP-Sam68 Δ C functions in the cytoplasm, and presumably competes with endogenous Sam68 for binding sites in mRNA-protein complexes, this result suggests that endogenous Sam68 positively regulates the translation of eEF1A-encoding mRNAs.

Possible post-synaptic anchoring by Sam68

In non-neuronal cells, Sam68 is known to bind to membrane-associated signalling proteins (Lukong and Richard 2003). In neurons, given its association with dendritic mRNAs that become actively translated in response to local synaptic stimuli (Tsokas et al. 2005), we wondered whether Sam68 might physically interact with synaptic receptors. In synaptic membranes prepared from rat cortex, both the NMDA receptor and the post-synaptic density protein PSD95 could be co-immunoprecipitated with Sam68 (Fig. 5, lanes 1-6). By contrast, AMPA receptors were essentially absent from these immunoprecipitates (lanes 7-9), consistent with the notion that Sam68 was specifically recruited by the NMDA receptor complex. We could also detect Sam68 in NMDA receptor immunoprecipitates (lanes 10-11). Thus Sam68 might conceivably provide a platform for docking specific neuronal mRNAs and / or for recruiting specific translation factors at subsynaptic sites.

Discussion

The present study indicates that brain Sam68 is associated with a wide but selective range of cytoplasmic mRNAs, among which transcripts encoding long-term plasticity related proteins are prominent. Sam68 might collectively regulate the synthesis of an entire group of such proteins, in an activity-dependent fashion. The set of Sam68-associated mRNAs we found in brain differs from those reported in non-neuronal cell lines or lineages (Itoh et al. 2002; Paronetto et al. 2006; Tremblay and Richard 2006). Extensive comparison of the different sets is difficult, since the screening methods were different. However, two of the mRNAs, β -actin and dynein, were found in both the study by (Itoh et al. 2002) and our screen, providing a reliable positive control. Taken together, the various screens suggest that mRNA recognition by Sam68 is highly dependent on cellular context.

One of most enriched among the Sam68-associated cortical transcripts was the Eef1a1 mRNA. Although Eef1a1 is ubiquitous, the Sam68-Eef1a1 mRNA interaction was apparently not detected in the previous studies. This may be due to the sensitivity of the micro-array sampling procedure; alternatively, the interaction may be favoured by neuron-specific factors. However, it was possible to reconstitute the Sam68-Eef1a1 mRNA interaction in transfected HEK cells by using overexpressed, myc-tagged Sam68. The interaction is phylogenetically conserved since HEK 293 cells are of human origin; this is consistent with the >90% identity of human and rat Eef1a1 nucleotide sequences, including in the 3' untranslated region. Interestingly, based on free energy minimization, the entire 3'UTR of the Eef1a1 mRNA is predicted to fold into a single, long hairpin displaying at its extremity a U-rich terminal loop (Fig. 6). This is reminiscent of the U6 motif-containing stem-loop structure that mediates recognition of the β -actin mRNA by Sam68 *in vitro* (Itoh et al. 2002). However, our experiments do not rule out the possibility that the Sam68 – Eef1a1 mRNA association may

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be mediated by a third molecule. Further studies will be required to establish whether the 3' UTR structure forms a direct binding site for Sam68 on the Eef1a1 mRNA.

In dissociated cortical neurons, chemically induced depolarisation is known to generate bursts of action potentials and to recruit some of the pathways that underlie synaptic plasticity. While causing the translational repression of multiple mRNA species, depolarisation activates the translation of a select group of transcripts (Krichevsky and Kosik 2001). Our data indicate that Eef1a1 belongs to this group. Eef1a1 encodes the ubiquitous isoform of translation elongation factor eEF1A, which transports amino-acyl-tRNA to the A site of the ribosome during protein synthesis. In addition, eEF1A also anchors mRNAs to the actin cytoskeleton, and can bundle actin filaments according to a unique spatial arrangement (Liu et al. 2002a). Local provision of eEF1A may play a role in both biosynthetic and structural aspects of synaptic plasticity, and perhaps establish a link between these two facets of long-term synaptic change. In *Aplysia*, synthesis of the eEF1A homologue is instrumental to maintenance of synapse-specific long-term facilitation (Giustetto et al. 2003). In the rodent hippocampus, translation of the Eef1a1 mRNA is regulated by synaptic signals in complex ways, which appear to differ between preparations. In hippocampal slices from young rats, Eef1a1 translation has been shown to be rapidly induced during tetanus-induced LTP of CA3-CA1 synapses, subject to activation of the rapamycin-sensitive mTOR / S6 Kinase pathway (Tsokas et al. 2005). This finding is consistent with the fact that the Eef1a1 mRNA contains a 5' oligopyrimidine tract, which is a signature for mTOR-inducible mRNAs. Interestingly, in adult rat hippocampus *in vivo*, LTP of dentate granule synapses induced redistribution rather than neosynthesis of eEF1A protein. In contrast, long-term depression elicited by pharmacological stimulation of metabotropic glutamate receptors (mGluRs) was accompanied by a large increase in eEF1A protein, without change in Eef1a1 mRNA level (Huang et al. 2005). Thus, the induction of eEF1A synthesis seems to be exquisitely sensitive

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to the spatio-temporal and cell-specific details of synaptic signalling, suggesting that multiple effectors regulate the translation of Eef1a1 in neurons. One of these effectors may be the RNA-binding protein FMRP, since in vitro and in cell lines, FMRP has been reported to bind and repress the Eef1a1 mRNA (Sung et al. 2003); furthermore it is tempting to speculate that removal of this putative inhibition underlies the effect of mGluRs on Eef1a1 translation (Bear et al. 2004). The present data suggest that Sam68 is also part of the regulatory machinery that controls neuronal Eef1a1 translation, with an action distinct from that of FMRP. Since depolarisation increased the recruitment of Sam68 to polysomal mRNAs, it is likely that Eef1a1 transcripts bound to Sam68 were engaged in translation. The finding that the dominant-negative GFP-Sam68 Δ C, but not GFP-Sam68(WT) inhibited eEF1A protein accumulation suggests that endogenous Sam68 may be required for the translational activation of Eef1a1 mRNA. Since the dominant-negative effect of GFP-Sam68 Δ C was already detected in unstimulated neurons, the low amount of Sam68 existing in the cytoplasm prior to depolarisation may contribute to generating the proportion of actively translated Eef1a1 mRNA seen under basal conditions. In line with this possibility, Sam68 has been reported to interact with RNA helicase A and with the mRNA export factor Tap (Reddy et al. 2000), two proteins known to promote translation of specific viral and cellular mRNAs (Hartman et al. 2006; Li et al. 2006). Kinases recruited by the C terminus of Sam68 might also be required to phosphorylate and release mRNA-bound translation inhibitory factors, similar to the case of hnRNP-K phosphorylation by Src (Ostareck-Lederer et al. 2002). GFP-Sam68 Δ C, which is defective for certain protein-protein interactions, may displace the endogenous Sam68 and associated translation factors out of the target mRNA; indeed, in the case of viral mRNAs, GFP-Sam68 Δ C functionally antagonizes RNA helicase A and Tap (Reddy et al. 2000). Upon depolarisation, the largely enhanced export of Sam68 to the cytoplasm may cooperate with the action of signalling cascades such as the mTOR pathway

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3 for translational stimulation of activity-responsive mRNAs exemplified by Eef1a1. A similar
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5 scheme may apply to certain non-neuronal systems; in the case of meiotic male gametes,
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8 Paronetto et al. (2006) recently confirmed the occurrence of signal-induced transfer of Sam68
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10 to the cytoplasm and its binding to polysomes.
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For Peer Review

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For Peer Review

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For Peer Review

Legends to Tables and Figures

Table Ia

ID: Genbank identifier of cloned sequence on the NIA 15 K micro-array. Name: name of transcript homologous to the sequence on the array (available through <http://lgsun.grc.nia.nih.gov/cDNA/15k.html>). Mean z score and its standard deviation (n=3) were calculated for each transcript as described in the text. P values were calculated from z scores, assuming normal distribution of scores. Biological process or component associated with transcripts were identified by reporting Gene Ontology (GO) descriptors used to annotate the transcripts in the NIA database (available through <http://lgsun.grc.nia.nih.gov/cDNA/NIA-CloneSet-GoTermFinder.html>). Rightmost column: identification of the GO descriptors by their numbering in the Gene Ontology database.

Fig. 1

A. Scheme of the RNA-protein coimmunoprecipitation procedure, and image of a representative hybridization experiment. Green channel: Cy5-labelled probes obtained by amplification of polyA⁺ mRNA immunoprecipitated with the anti-Sam68 antibody. Red channel: Cy3-labelled probes obtained after immunoprecipitation with control (preimmune) antibody. **B.** Distribution of paired fluorescence values obtained in A. X-axis: specifically immunoprecipitated transcripts. Y-axis: adventitious transcripts. A large number of transcripts are more abundant in the specific immunoprecipitate than in the control. **C.** Within-sample reproducibility of z score measurements. The results of two different hybridization experiments performed with the same sample are compared. The regression line is shown in red. **D.** Between-sample reproducibility of z scores. The results of hybridization experiments performed on two different brain extracts are compared. The regression line is shown in red.

Note that mRNAs with highest values in either sample (to the right of the vertical green line and above the horizontal green line) are mostly confined to the top right quadrant. **E.** Functional profile of the 43 mRNAs displaying the highest mean z scores in the immunoprecipitates (see Table Ia). For each transcript, a generic function was registered (identified by Gene Ontology descriptors of the mRNA in the NIH database; see Table Ia). Black bars: number of occurrences of the indicated functional descriptors in the 43 mRNA sample. Empty bars: expected number of occurrences of the descriptors, based on their frequencies among mRNAs of the micro-array. The difference is highly significant ($p < 0.0001$, chi-square test), indicating enrichment for specific functional categories.

Fig. 2

A. Validation of specific micro-array hybridization results by RT-PCR. Cortical tissue extract was subjected to immunoprecipitation with anti-Sam68 (“IP”) or preimmune (“Pre”) IgG as shown; total mRNA was purified from the immunoprecipitates, and subjected to reverse transcription and amplification by PCR, using the indicated gene-specific primers (Table II). The PCR products were visualized by fluorescent staining with ethidium bromide. The expected sizes of PCR products are indicated (Eef1a1: 336 bp; Uch-L1: 187 bp; β -actin: 178 bp; Calm-1: 229 bp; Arc: 287 bp). Positive controls for the amplification reactions were obtained by performing PCR with total mRNA (“Tot”) directly derived from the input extract. No amplification took place without cDNA (“None” or “0”).

B. Reconstitution of the association between Sam68 and the Eef1a1 mRNA in cultured cells. HEK 293 cells were transfected with a myc-tagged Sam68 expression construct, or with control vector, as indicated. Cytoplasmic extracts were prepared and immunoprecipitated with anti-myc antibody, and subjected to RT-PCR with Eef1a1 (lanes 1, 3) or with β -tubulin (lanes

7,9) primers. As positive controls for amplification, RT-PCR products were generated with total mRNA from the transfected cells (lanes 2, 4, 8, 10) or from rat cortex (lane 5, 11). Lanes 6, 12: negative control for PCR.

C. The amounts of endogenous Eef1a1 mRNA in anti-myc immunoprecipitates from the indicated transfections were determined by quantitative (“real-time”) PCR (n=3 for each transfection). *a.u.*, arbitrary units.

Fig. 3

A. Depolarisation induces an increase in Eef1a1 mRNA translation. Cultured cortical neurons were incubated for 6 h in 5mM or 25 mM KCl, then cytoplasmic extracts were prepared and equal protein amounts were subjected to fractionation on sucrose density gradients. For each gradient, representative fractions corresponding to translationally inactive mRNPs, 80S monosomes, polysomes, and RNA granules were analysed by RT-PCR with specific primers to amplify the Eef1a1 mRNA. The PCR products were analysed side by side on an agarose gel (“-” and “+” lanes : gradient fractions from control and depolarised neurons; respectively) and visualized by fluorescent staining with ethidium bromide. Greyscale was inverted to improve contrast.

B. Depolarisation increases the amount of eEF1A protein. Cortical neurons were treated as in A, then lysed, and equal amounts of total protein were analysed by SDS-polyacrylamide gel electrophoresis and immunoblotting. Left: the same blot was probed first with anti-eEF1A monoclonal antibody, then with anti-actin antibody. Right: the relative amount of eEF1A in each lane was measured by densitometric scanning, and normalized to the amount of actin in the same lane to correct for gel loading error. The histogram shows the average (+/- S.D.) of normalized eEF1A amount in each condition.

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C. Depolarisation enhances the association of Sam68 with polysomal RNA and RNA granules. Cytoplasmic extracts from control and depolarised neurons were prepared and fractionated as in **A**. The fractions were analysed by immunoblotting with anti-Sam68 antibody (left). The distribution of RNA along the gradient was determined by continuous monitoring of UV absorbance at 254 nm during collection of the fractions (right). The various types of ribonucleoproteic assemblies are indicated.

Fig. 4

Effect of dominant-negative Sam68 on eEF1A expression in cultured hippocampal neurons.

A-C: The GFP-Sam68 (WT) and GFP-Sam68 Δ C proteins were expressed in hippocampal neurons, together with monomeric Red Fluorescent Protein (mRFP) as a morphological marker. Five days later (at 14 DIV), the neurons were fixed and stained by immunofluorescence with anti-eEF1A antibody; mRFP, GFP-Sam68 and endogenous eEF1A were simultaneously visualized by triple-colour fluorescence microscopy.

A1-A3: close-up view of the three fluorescent markers in a GFP-Sam68 Δ C-transfected neuron and a nearby, untransfected neuron (wide-field microscopy). **A1:** Note the lack of eEF1A immunostaining in the transfected cell (arrow), compared to the neighbouring cell. Scalebar: 13 μ m. **B:** clustered distribution of the GFP-Sam68 Δ C protein within the dendrite of a hippocampal neuron (single plane of a confocal z series). Scalebar: 4.7 μ m.

C1-C3: representative field showing GFP-Sam68 Δ C transfected neurons and neighbouring untransfected cells at low magnification. Scalebar: 65 μ m. **C1:** the relative intensity of eEF1A immunofluorescence is represented by pseudocolour coding. Perinuclear staining is brighter in untransfected neurons (arrowheads) than in transfected (arrows). **C3:** normal neuronal morphology of the transfected cells.

D1-D3: same as C1-C3 with GFP-Sam68WT transfected neuron.

E: the eEF1A immunofluorescence was measured (as mean grey level per pixel) in transfected neurons and in surrounding, nontransfected neurons of the same field. The ratio between the fluorescence of transfected neurons and the average fluorescence of surrounding nontransfected cells was then calculated for each transfected neuron. The histogram bars show the average (\pm S.D.) of these ratios for neurons transfected with GFP-Sam68 Δ C and with GFP-Sam68WT. Neurons expressing GFP-Sam68 Δ C had a ratio lower than 1, indicating a decrease in endogenous eEF1A. Neurons expressing GFP-Sam68WT had a ratio close to 1. The difference between GFP-Sam68 Δ C and GFP-Sam68WT was significant ($p=0.03$, two-sided Mann-Whitney U test) ($n=3$ experiments).

F: schematic depiction of the wild-type and C-terminally deleted, dominant-negative Sam68 proteins. The major protein domains are shown as boxes. KH: core RNA-binding module (hnRNP-K homology domain). P1 to P5: proline-rich motifs. Y: tyrosine-rich region. NLS: nuclear localisation signal.

Fig. 5

Binding of Sam68 to the post-synaptic NMDA receptor complex. Purified synaptic membranes were solubilized under non-denaturing conditions, and subjected to immunoprecipitation with the indicated antibodies. Lanes 1, 4, 7: aliquot (10%) of input lysate. After detecting PSD-95, the blot of lanes 4-6 was stripped and reprobed to detect GluR2/3 subunits of AMPA receptors, shown in lanes 7-9. Lane 10: control immunoprecipitation with a mouse monoclonal IgG, class-matched to the anti-NR1 antibody.

Fig. 6

The 3' UTR sequences of human EEF1A1 and rat Eef1a1 mRNAs have been aligned using Clustal W (v. 1.83). The first nucleotide of each sequence is that immediately following the

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2
3 stop codon, and the last nucleotide corresponds to the major polyadenylation site. Stars
4 indicate identical residues. Using the Rnaalifold programme of the Vienna RNA package (Ivo
5 L Hofacker, <http://www.tbi.univie.ac.at/~ivo/RNA/>), the predicted folding of the aligned
6 sequences displays a large stem structure with a U-rich terminal loop (overlined).
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Specific interaction between Sam68 and neuronal mRNAs: implication for the activity-dependent biosynthesis of elongation factor eEF1A.

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Abstract

In cultured hippocampal neurons and in adult brain, the splicing regulatory protein Sam68 is partially relocated to the somatodendritic domain, and associates with dendritic polysomes. Transfer to the dendrites is activity-dependent. We have investigated the repertoire of neuronal mRNAs to which Sam68 binds *in vivo*. Using co-immunoprecipitation and microarray screening techniques, Sam68 was found to associate with a number of plasticity-related mRNA species, including Eef1a1, an activity-responsive mRNA coding for translation elongation factor eEF1A. In cortical neuronal cultures, translation of the Eef1a1 mRNA was strongly induced by neuronal depolarisation, and correlated with enhanced association of Sam68 with polysomal mRNAs. The possible function of Sam68 in Eef1a1 mRNA utilization was studied by expressing a dominant-negative, cytoplasmic Sam68 mutant (GFP-Sam68 Δ C) in cultured hippocampal neurons. The level of eEF1A was lower in neurons expressing GFP-Sam68 Δ C than in control neurons, supporting the proposal that endogenous Sam68 may contribute to the translational efficiency of the Eef1a1 mRNA. These findings are discussed in the light of the complex, potentially crucial regulation of eEF1A biosynthesis during long-term synaptic change.

Keywords

Dendritic mRNA, activity-dependent translation, RNA-binding protein, plasticity

Introduction

The translation of neuronal mRNAs is regulated by their association with specific RNA-binding proteins, of which an increasing variety have been identified (Bramham and Wells 2007; Kiebler and Bassell 2006). In the well-studied cases of CPEB, FMRP and ZBP-1, the proteins maintain the target mRNAs in a dormant state during transport or storage; this inhibitory function is thought to be suppressed by activity-dependent phosphorylation events, e.g. in the vicinity of synapses (Bagni and Greenough 2005; Huang et al. 2002; Huttenlocher et al. 2005). In addition to these predominantly cytoplasmic proteins, recent evidence from non-neuronal systems indicates that nuclear proteins of the splicing machinery may also contribute to the control of translation. For example, SR splicing factors continuously shuttle between nucleus and cytoplasm, where they can bind to mature mRNAs and promote translation (Sanford et al. 2004). Translational efficiency can also be increased by splicing proteins which are deposited onto RNA substrates during the splicing process itself, and which remain bound to the mRNA after nuclear processing and export have been completed (Kunz et al. 2006; Le Hir et al. 2000; Wiegand et al. 2003). Interestingly, a diversity of splicing factors have been found to exist in neuronal dendrites (Glanzer et al. 2005). One of those is Sam68, a ubiquitous RNA-binding protein implicated in signal-dependent splice site selection, and which can also function as an adaptor and substrate for Src family kinases (Lukong and Richard 2003; Matter et al. 2002; Paronetto et al. 2007). In cultured cell lines, Sam68 is a nuclear, non-shuttling protein; however, it is required to enhance the nuclear export and / or translation of certain retroviral mRNAs, indicating a potential interaction with translation-activating factors (McLaren et al. 2004; Modem et al. 2005). Beside viral mRNAs, Sam68 can bind to a select population of mature cellular mRNAs derived from cultured HeLa cells (Itoh et al. 2002); the functional outcome of these binding interactions has not been determined.

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We previously reported that Sam68 was among the proteins associated with neuronal mRNAs (Ben Fredj et al. 2004; Grange et al. 2004). In adult cortical and hippocampal tissue, biochemical fractionation and immuno-electron microscopy indicated that Sam68 molecules were associated with dendritic polysomes, suggesting a possible positive role in translational regulation (Grange et al. 2004; Khandjian et al. 2004). In cultured hippocampal neurons, a minor but detectable fraction of endogenous or GFP-tagged Sam68 molecules were found to occur in the soma and dendrites, concentrated in generally immobile, RNA-rich granules. Following prolonged membrane depolarisation by elevated KCl, the proportion of Sam68 in the cytoplasm was strongly increased, due to massive migration of the protein to these somatodendritic sites (Ben Fredj et al. 2004). These results suggest that Sam68 may play a part in the regulation of those mRNAs which become translationally activated in response to electrical activity. To test this hypothesis, we have investigated the identity of mRNA species that were bound to Sam68 in cortical neurons in vivo. We found that one major target was Eef1a1, an mRNA known to be activated during long-term synaptic change in the hippocampus. In cultured neurons, depolarisation enhanced the association of both Sam68 and the Eef1a1 mRNA with polyribosomes, and increased the amount of its product, the eEF1A protein. Expression of a dominant-negative Sam68 mutant confined to the cytoplasm was able to lower the amount of eEF1A, supporting the proposal that Sam68 promotes the translation of neuronal Eef1a1.

Materials and Methods

Immunoprecipitation of Sam68-associated RNA and preparation of amplified RNA probes

Cortical Sam68-RNA complexes were prepared as previously described (Grange et al. 2004). Briefly, the cortex from one adult rat was dissected and lysed at 4°C in polysome dissociation buffer [10 mM Tris-HCl, pH 7.5, 50 mM KCl, 1 mM sodium orthovanadate, 2 mM dithiothreitol (DTT), 100 nM microcystine-LR (Sigma), protease inhibitors (Complete; Roche Molecular)] containing 0.5% Triton X-100, 160 mM sucrose, 30 mM EDTA, pH 8.0 and 400 U/ml RNasin (Promega, Inc., Madison, WI). After clarification by centrifugation at 82,000g for 20 min at 4°C, the lysate was precleared with protein G–Sepharose beads (GE Healthcare), then immunoprecipitated with affinity-purified anti-Sam68 IgG (AD-1), or with preimmune protein A-purified IgG from the same rabbit. Immune complexes were collected onto protein G –Sepharose beads, washed, and processed for RNA purification using the “RNEasy” kit (Qiagen France, Courtaboeuf). The immunoprecipitated polyA+ RNAs were linearly amplified by cDNA synthesis and T7 polymerase transcription (Gelder et al. 1990), using the MessageAmp™ aRNA Kit (Ambion). According to whether the template RNA originated in the specific immunoprecipitate or the preimmune control, either Cyanine 3™ (Cy3) or Cyanine 5™ (Cy5)-labeled UTP was incorporated during transcript synthesis, as per manufacturer’s instructions. Transcripts amplified from the specific immunoprecipitate and the corresponding control were mixed in equal amounts and used to probe the 15000 cDNAs arrayed on the NIA micro-array (Tanaka et al. 2000). To control for possible biasing by differential incorporation of Cy3 and Cy5, the experiment was repeated with swapping the fluorophores between “specific” and “background” transcripts.

Quantitation of hybridization

After hybridization and washing, digital images of the micro-array at 635 nm and 532 nm were acquired by laser scanning confocal microscopy. Fluorescent spots were delineated and

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quantitated with GenePix software (Axon Instruments). Raw fluorescence values (F) in each channel were converted into their Log_2 . After Cy3 labelling of background RNAs, immunoprecipitation of any given mRNA was considered specific when the value in the Cy5 channel exceeded that in the Cy3 channel by at least 4 Log_2 units (a 16-fold difference). When background RNAs were labelled with Cy5, the cut-off was brought down to 0.5 Log_2 unit (an 1.4-fold difference). This differential cut-off was applied to compensate for the systematic bias in labelling efficiency in favour of Cy5. Note that the requirement for higher Cy3 labelling of specifically immunoprecipitated RNAs is conservative and likely excludes a number of Sam68-associated mRNAs. Z scores were calculated as $(\text{Log}_2F - \langle \text{Log}_2F \rangle) / \sigma_{\text{Log}_2F}$, where $\langle \text{Log}_2F \rangle$ is the mean and σ_{Log_2F} is the standard deviation, both calculated over the entire population of fluorescence values in the relevant channel (Mayne et al. 2001). Reproducibility of the overall population of z scores was measured by simple linear regression between experiments. 95% confidence intervals were calculated for the z scores of Table Ia using t statistics ($n=3$), and individual mRNAs with lower confidence limit $z < -1.0$ were rejected for inclusion in the Table; more than 80% of these scores had lower limit above zero. Frequencies of GO descriptors were computed from ~4000 functionally described genes of the microarray (available through <http://lgsun.grc.nia.nih.gov/cDNA/NIA-CloneSet-GoTermFinder.html>).

Polymerase chain reactions

Immunoprecipitated or total RNA was reverse-transcribed with the universal primer GGAATTC(T)₁₇, and the resulting cDNA was amplified by using the gene-specific primer pairs listed in Table II. For classical PCR, after 2 min at 95°C, 30 cycles were performed (15 s at 95°C, 30 s annealing, and 30 s at 72°C for each cycle). For quantitative PCR by real-time fluorometry (Light Cycler, Roche Molecular Systems), DNA was denatured at 95°C for 5 min, then subjected to ~35-40 amplification cycles (15 s at 95°C, 5 s annealing, 15 s at 72°C,

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all temperature changes at 20°C/s). Annealing took place at 55°C except for Arc (57°C). The relative amounts of specific DNA template were determined by counting the number of cycles needed to reach the inflection point (“crossing point” at which $d^2[\text{DNA}]/dt^2=0$) of the DNA accumulation curve (typically between 25 and 32), and standardized by relating to a scale generated with known amounts of total cDNA. The specificity of the amplification product was confirmed by checking its melting curve with the Light Cycler, and its size on an agarose gel.

Immunoprecipitation of RNA-mycSam68 complex from HEK cells

HEK-293 cells were grown in Dulbecco’s modified Minimum Essential Medium, containing 10% fetal calf serum (Invitrogen). The myc-Sam68 expressing plasmid was a kind gift from I. Barlat and B. Tocqué (at the time at Aventis Pharma, Vitry, France). One 6 cm dish of cells was transfected with either myc-Sam68 or with control vector (containing a CMV promoter). 48 h later, cells were lysed in polysome dissociation buffer and, after clarifying by centrifugation, the lysate was immunoprecipitated with 2 µg anti-myc monoclonal antibody (12CA5, Santa Cruz) and protein G-Sepharose beads, as described for brain lysates. **Total RNA was isolated and amplified by RT-PCR as described above.**

Neuronal cultures and transfections

Primary cultures of rat hippocampal neurons were prepared from fetal (E19) hippocampi and seeded at a density of 6000 cells / cm² onto poly-L-lysine-coated glass coverslips (Marienfeld, Germany), adhering to the procedure described by Goslin and Banker (Banker et al. 1998), except that the culture medium was Neurobasal with B27 supplement (Invitrogen). Housing and handling of animals complied with the regulations issued by the European Union and INSERM and were approved by the local veterinary authorities. The GFP-Sam68 (WT) and GFP-Sam68ΔC plasmids were kind gifts from S. Richard (McGill University, Montréal, Canada). The mRFP1 plasmid (expressing monomeric red fluorescent protein) was a kind gift

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3 from R.Y. Tsien (UCSD, San Diego, CA, USA). Transfection was performed at 9 DIV by
4 calcium phosphate coprecipitation, exactly according to (Goetze et al. 2004). We used a mix
5 of 2.5 μg GFP-Sam68 and 0.5 μg mRFP1 plasmids per coverslip. Expression of the GFP-
6 Sam68 and GFP-Sam68 ΔC plasmid was stable for the next 10 days. Neurons were fixed at
7 14-19 DIV and processed for immunofluorescent staining.

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10 Primary cortical neurons were prepared from E16 fetal cortices, using essentially the same
11 procedure as for hippocampal neurons. The cortical cells were seeded at a density of 10^5 cells
12 / cm^2 onto poly-L-lysine coated 10-cm dishes (Nunc). Neurons at 9 DIV were treated for 3 h
13 with either depolarizing medium (containing, in mM: KCl, 25; CaCl₂, 1.8; MgCl₂, 0.8; NaCl,
14 100; NaHCO₃, 26; NaH₂PO₄, 1; glucose, 30; Hepes-NaOH, 15) or control solution (same
15 except for KCl, 5; NaCl, 120).

16 *Sedimentation of polysomes on sucrose gradients*

17 RNA-protein complexes were fractionated as described by (Krichevsky and Kosik 2001).
18 Briefly, cortical neurons were lysed in low salt buffer [20 mM Tris-HCl (pH 7.5), 10 mM
19 NaCl, 3 mM MgCl₂, 1 mM RNasin, 1 mM dithiothreitol, 0.3% Triton X-100, and 0.05 M
20 sucrose] containing protease inhibitors (EDTA-free Complete, Roche Molecular), 1 mM
21 vanadate and 100 nM microcystin LR. Nuclei and the majority of mitochondria were
22 sedimented by centrifugation for 10 min at $10,000 \times g$ at 4°C. The NaCl and MgCl₂
23 concentrations in the cytoplasmic extracts (supernatants) were adjusted to 170 and 13 mM,
24 respectively. Linear sucrose gradients (15%–45% w/w in 25 mM Tris-HCl [pH 7.5], 25 mM
25 NaCl, 5 mM MgCl₂) were prepared using an AutoDensi Flow apparatus (Büchner).
26 Cytoplasmic extracts (1.2 ml) were overlaid onto 10.6 ml gradients and centrifuged at 32,000
27 rpm for 2.5 hr at 4°C in a SW41 rotor. Fractions of 1 ml volume were collected with on-line
28 UV absorbance measurement at 254 nm (GE Healthcare Akta Prime). For Western
29 immunoblotting, the protein content of the fractions was concentrated by trichloroacetic acid
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3 precipitation, redissolved in Laemmli buffer, and analysed on 8% polyacrylamide / SDS gels.
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5 For RT-PCR, RNA was extracted by treating the relevant fractions with Trizol reagent
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7 (Invitrogen), precipitated with ethanol, and subjected to RT-PCR with the same primers as
8
9 above.
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12 *Detection of eEF1A protein increase by immunoblotting*

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14 Cortical neurons were treated with control or depolarising medium as above, then lysed in ice-
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16 cold RIPA buffer (20 mM Tris-HCl [pH 7.5], 150 mM NaCl, 1 mM EDTA, 1 % Triton X-
17
18 100, 0.5 % Na deoxycholate, 0.1 % SDS) containing protease inhibitors (EDTA-free
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20 Complete, Roche Molecular). After clarification by centrifugation (15 min at 12,000 x g),
21
22 equal protein amounts of each lysate were analysed by 10 % polyacrylamide / SDS gel
23
24 electrophoresis and immunoblotting, first with anti-eEF1A monoclonal antibody (UBI) then
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26 with anti-actin polyclonal antibody (Santa Cruz). Immunoreactive bands were revealed by
27
28 chemiluminescence (Pierce Femto) and autoluminography. The films were digitized by
29
30 scanning and the integrated intensity of the bands was determined using Metamorph software
31
32 (Universal Imaging). Immunoblotting and exposure conditions were calibrated so that the
33
34 integrated intensity of the bands was in the linear range with respect to protein amount.
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37 *Immunofluorescence of eEF1A in hippocampal neurons*

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39 Neurons were fixed with 4% PFA containing 4% sucrose (30 min, 4°C); permeabilized with
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41 0.2% Triton X-100; blocked with 10% preimmune goat serum in PBS (25°C, 1 h); and stained
42
43 with 1 µg/ml anti-eEF1A monoclonal antibody (UBI) in PBS containing 3% goat serum
44
45 (25°C, 2 h), then with Alexa350TM-conjugated secondary antibody (Molecular Probes). After
46
47 washing, coverslips were mounted with Mowiol and observed on a Zeiss microscope fitted
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49 with an epifluorescence setup and a cooled CCD camera (Zeiss Axiocam), or with a confocal
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51 laser scanning microscope (Zeiss LSM 510).
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For quantitation of eEF1A fluorescence, fields containing transfected neurons were imaged at low magnification (20x). Imaging conditions were identical for all fields. Images were processed with Metamorph (Universal Imaging). For each image, a background image was produced with a low-pass filter and subtracted so as to remove low-frequency fluorescence noise due e.g. to local unevenness in illumination. Next, a lower fluorescence threshold was set, which allowed automatic delineation and analysis of the fluorescent objects (cells), using Integrated Morphometric Analysis. The same threshold was used throughout the entire analysis. The relative eEF1A fluorescence of each transfected neuron i was calculated according to the equation: $IF_{\text{transf},i} = eEF1A_{\text{transf},i} / \langle eEF1A \rangle_{\text{nontransf}}$ where $eEF1A_{\text{transf},i}$ is the mean grey level per pixel within the transfected neuron, and $\langle eEF1A \rangle_{\text{nontransf}}$ is the average (over 15 surrounding neurons) of the mean grey level per pixel within non-transfected neurons. These $IF_{\text{transf},i}$ values were averaged over the entire set of transfected neurons, to yield the values of Fig. 4E.

Immunoprecipitation of Sam68-NMDA receptor complexes

Rat cortical synaptic membranes were prepared by differential centrifugation as described by Grange et al. (2004), and resuspended in ice-cold buffer containing 4mM HEPES (pH 7.4), 1mM preboiled sodium orthovanadate, 1mM dithiothreitol, 100 nM microcystin (Sigma), and a mix of protease inhibitors ("Complete", Roche). An aliquot of suspension (530 μ l, 4.5 μ g/ μ l total protein) was solubilized by adding 1:9 vol. of detergent solution (10% sodium deoxycholate in 0.5M Tris, pH 9.0) and gently agitating for 20 min. Triton X-100 (0.1% final concn.) was then added, and the extract was dialyzed overnight against immunoprecipitation buffer (10 mM Tris-HCl, pH 7.4, 100 mM NaCl, 0.1 mM sodium orthovanadate, 5 mM EDTA, 1 mM dithiothreitol, 0.1 % Triton X-100, 1 nM microcystin, and protease inhibitors). The dialyzed extract was centrifuged for 30 min at 10^5 g in a TL-100 tabletop ultracentrifuge (Beckman), and the supernatant (300 μ g protein diluted to 500 μ l buffer per IP) was used for

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3 immunoprecipitation for 3 h with 10 μ g affinity-purified anti-Sam68 polyclonal antibody, or
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6 preimmune IgG purified from the same rabbit, or with anti-NR1 monoclonal antibody
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8 (Pharmingen). Immunoprecipitates were collected onto protein G-Sepharose beads and
9
10 analysed by SDS-PAGE (8% polyacrylamide gel) and Western immunoblotting. Antibodies
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12 used for immunoblotting were purchased from UBI (anti-PSD95 monoclonal antibody) and
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14 Chemicon (anti-GluR2/3 rabbit polyclonal antibody). The bands were revealed by
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16 chemiluminescence (Pierce Femto).
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Results

Identification of Eef1a1 in the repertoire of Sam68-associated mRNAs

Following an approach initiated by Tenenbaum et al. (2000), we decided to use a combination of co-immunoprecipitation and array screening to probe the repertoire of Sam68-associated mRNAs in cortical neurons. Sam68-mRNA complexes were immunoprecipitated from a rat cortical cytoplasmic extract with an anti-Sam68 antibody, as previously reported (Ben Fredj et al. 2004). A parallel immunoprecipitation was performed with preimmune antibody. The polyA⁺ RNAs associated with the Sam68 and control immunoprecipitates were linearly amplified by cDNA synthesis and T7 polymerase-mediated transcription, as initially described by Eberwine and coworkers (Gelder et al. 1990), and the amplified transcripts were fluorescently labeled with Cy3 or Cy5, respectively. The two transcript populations were then mixed and hybridized with the ~15,000 cDNAs of the NIA “15K” clone set arrayed onto a glass slide (Tanaka et al. 2000), and for each cDNA the fluorescence values at both wavelengths were measured and compared (Fig. 1 A, B). A given mRNA was considered to be specifically immunoprecipitated when the fluorescence level was consistently higher in the channel corresponding to the immunospecific than in the background precipitation. To compare mRNA abundance measurements across several experiments, normalized measures of the relative abundance of each mRNA in the immunoprecipitated population were calculated by converting the fluorescence values into z scores (Lopez de Silanes et al. 2004; Mayne et al. 2001; see Materials and Methods). Duplicate assays of the same RNA population showed that z scores were highly reproducible through the entire set of ~15,000 genes, except for a limited number of low abundance mRNAs, indicating that abundance measurements were not grossly perturbed by fluctuations in the hybridization and detection procedure (Fig. 1C). When distinct experiments with different brain samples were compared, fluctuations in individual z scores were larger, likely due to random differences in the hydrolysis or

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3 dissociation rates of individual mRNAs during immunoprecipitation. The correlation between
4 samples remained sufficient to ensure that the species scoring highest ($z > 1.5$) in one trial
5 retained a 85% probability of scoring positive ($z > 0$) in a second trial (Fig. 1D). To obtain
6 reliable identification of the most abundant mRNAs in the immunoprecipitate, z scores were
7 measured in three independent experiments, mRNAs were ranked by decreasing mean z
8 score, and some species with unacceptably large standard deviations were excluded. We also
9 discarded a few arrayed sequences for which no cognate mRNAs had been identified. Table Ia
10 shows those specifically immunoprecipitated mRNA species for which the mean relative
11 abundance was highest after applying this screen .
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24 The dual-channel procedure was then used to compare the relative abundances of mRNA
25 species in the immunoprecipitated vs. total polyA+ populations. The total population of
26 polyA+ RNAs present in the input extract (before immunoprecipitation) was amplified and
27 the resulting Cy5-labeled transcripts were mixed with an aliquot of Cy3-labeled transcripts
28 derived from the Sam68 immunoprecipitate. After hybridization to the array, z scores were
29 calculated for each channel. The species of Table Ia scored 0.6 to 1.0 units higher in the
30 immunoprecipitated than in the total population, consistent with their enrichment.
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44 There was a discernible pattern in the biological functions of these mRNAs (Fig. 1E). When
45 assessed by Gene Ontology annotation for generic cellular process or component, more than
46 20% of the mRNAs (9 transcripts) were involved in intracellular or receptor-associated signal
47 transduction, as they coded for e.g. receptor subunits, protein kinases, protein phosphatases,
48 and associated regulatory factors. Another 20% of the mRNAs coded for proteins involved in
49 protein transport, either as regulators of vesicular traffic, or of microtubule-dependent motion.
50 These proportions are significantly higher than what would be expected based on the overall
51 distribution of biological functions among the genes represented on the micro-array (Fig. 1E).
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60 By comparison, transcription regulators appeared to be under-represented among the major

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Sam68-associated mRNAs. Assessment of the literature further indicates that some of the proteins of Table I contribute to exocytic and endosomal traffic (Arf-1, BRI-3, vacuolar ATPase) or to cytoskeletal regulation (stathmin) even though this is not apparent from their GO function label. All of these processes are heavily involved in the local response to stimuli which affect neuronal homeostasis and plasticity. Indeed, more than one third of the genes of Table Ia code for proteins directly involved in synaptic plasticity, connectivity development, or neurodegenerative disease (Table Ib).

The β -actin mRNA, which was detected among the most abundant species in our screen, had formerly been detected by RT-PCR in immunoprecipitates of brain Sam68 (Ben Fredj et al. 2004), and was known as a specific target for recombinant Sam68 (Itoh et al. 2002). This provided a positive control for the immunoprecipitation procedure. The Gephyrin mRNA provided a negative control, as it was found neither by RT-PCR, nor in the micro-array analysis (data not shown and (Grange et al. 2004)).

The most enriched mRNA, Eef1a1, coded for the ubiquitous isoform of translation elongation factor eEF1A, an important player in both protein synthesis and long-term synaptic change (see Discussion). The z score value of the Eef1a1 fluorescence indicates that the representation of Eef1a1 among immunoprecipitated mRNAs is significantly higher than average ($p=0.004$).

RT-PCR with gene-specific primer pairs (see Table II) was used to confirm the specific coimmunoprecipitation of a range of mRNAs detected in the micro-array analysis, either among the more enriched species listed in Table Ia, or among mRNAs with lower relative abundance. In addition to β -actin, confirmed species included Eef1a1, Calmodulin I and Uch-L1 (Fig. 2). Interestingly, the Arc / Arg 3.1 mRNA was present among the less abundant mRNAs in the micro-array analysis, and could also be detected by RT-PCR (see Fig. 2A). Since both Arc and Eef1a1 are dendritic, activity-responsive mRNAs, this raises the

possibility that upon exit to the somatodendritic compartment Sam68 may play a role in activity-induced translation of neuronal mRNAs.

As Eef1a1 was the most enriched of Sam68-associated mRNAs in our assay, and since its translational regulation by synaptic inputs is well-documented (see below), we decided to investigate further its relationship with Sam68. To reconstitute the interaction between Eef1a1 mRNA and Sam68 in a heterologous system, myc-tagged Sam68 or a control plasmid were transiently expressed in HEK 293 cells; cytoplasmic extracts prepared from the transfected cells were immunoprecipitated with anti-myc antibody, and the immunoprecipitates were probed by RT-PCR using the Eef1a1-specific primer pair. Endogenous Eef1a1 transcripts were detected in anti-myc immunoprecipitates from myc-Sam68-expressing cells but not from control cells (Fig. 2C, lanes 1-6). This result was reproduced by quantitative RT-PCR (Fig. 2C). Thus, the Eef1a1 mRNA associates with *bona fide*, overexpressed Sam68. **By contrast, the β -tubulin mRNA was absent from myc-Sam68 immunoprecipitates (Fig. 2B, lanes 7-12), supporting the mRNA sequence specificity of binding.**

Identification of Eef1a1 as a translationally activated mRNA in depolarized neurons

The Eef1a1 mRNA contains a 5' oligopyrimidine tract and belongs to the general group of signal-responsive mRNAs encoding components of the translation apparatus (Petroulakis and Wang 2002). Moreover, translation of Eef1a1 is stimulated during synaptic change ((Tsokas et al. 2005). To determine whether Eef1a1 might belong to the population of translationally activated mRNAs in our system of in vitro depolarised neurons, primary cultures of cortical neurons were subjected to control treatment or prolonged depolarisation, lysed under non-denaturing conditions, and the post-nuclear supernatants were fractionated on sucrose density gradients, as described by (Krichevsky and Kosik 2001). Fractions were analysed for their RNA content by continuous monitoring of UV absorbance during collection. Fractions corresponding to the free mRNPs, 80S monosomes, polyribosomes, and RNA granules

(migrating down to the bottom of the gradient) were analysed by RT-PCR with Eef1a1-specific primers. In non-stimulated neurons, the Eef1a1 mRNA was associated to the same extent with free mRNPs and with polysomes. Upon depolarisation, Eef1a1 became nearly undetectable in the free mRNP fraction, and strongly accumulated in the polysome fraction (Fig.3A). These data indicate that Eef1a1 mRNA translation is activated by depolarisation. Consistent with this result, depolarisation of cultured cortical neurons induced a significant increase in the total level of eEF1A protein, as detected by Western immunoblotting (Fig. 3B).

Depolarisation-inducible association of Sam68 with polysomes

We previously reported that Sam68 was associated with polyribosomes in cytoplasmic extracts derived from adult rat cortical tissue (Grange et al. 2004). To confirm that a similar association occurred in cultured cortical neurons, and to determine whether the interaction could be regulated by depolarisation, extracts were prepared and fractionated exactly as in Fig. 3A. Fractions were then analysed by Western immunoblotting with anti-Sam68 antibody. In extracts from control, quiescent neurons, Sam68 mostly comigrated with free mRNPs; a minor fraction sedimented in the RNA granule fraction (Fig. 3C, top). Following depolarisation, Sam68 accumulated in fractions containing polysomes and granules, even though the total amount of RNA in those fractions was slightly lower than in the control condition (Fig. 3, bottom). This result is consistent with the depolarisation-induced accumulation of Sam68 fluorescence in RNA-enriched dendritic clusters, which we have previously reported (Ben Fredj et al. 2004). Thus, in depolarised neurons, the increase in Eef1a1 translation correlates with the loading of Sam68 onto translated mRNAs.

Effect of dominant-negative Sam68 on neuronal eEF1A expression

The association of Sam68 to actively translated Eef1a1 mRNA suggests that Sam68 might regulate the synthesis of eEF1A. To investigate this point, we assayed the level of eEF1A

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immunoreactivity in cultured hippocampal neurons that expressed GFP-tagged derivatives of either wild-type Sam68 or Sam68 Δ C, a C-terminally truncated Sam68 mutant. The latter mutant retains a significant affinity for specific RNA sequences, but is deprived from tyrosine-rich and proline-rich motifs involved in protein-protein interactions with SH2 and SH3-containing ligands, and also from a C-terminal (nonclassical) nuclear localisation signal (Fig. 4F). In cultured cell lines, Sam68 Δ C is constitutively cytoplasmic and displays transdominant-negative activity towards endogenous Sam68, when coexpressed with retroviral (HIV-1) mRNAs that require Sam68 for efficient utilization (Reddy et al. 1999; Soros et al. 2001). Sam68 Δ C is specifically recruited to these viral mRNAs once they have reached the cytoplasm, and appears to block their normal interaction with the translation apparatus (Soros et al. 2001). We reasoned that expression of Sam68 Δ C in neurons should similarly allow us to probe the contribution of endogenous Sam68 to the translational activity of target neuronal mRNAs, as distinct from its role in nuclear mRNA processing. In transfected neurons, the GFP-Sam68 Δ C protein was mostly excluded from the neuronal nucleus and concentrated in clusters within the cell body; it was also detectable in dendritic shafts (Fig. 4, A2, B). The relative expression level of endogenous eEF1A in transfected and non-transfected neurons was evaluated by immunofluorescent staining with an anti-eEF1A monoclonal antibody. Among the neurons of a field, those expressing GFP-Sam68 Δ C typically had a lower level of eEF1A immunoreactivity than surrounding, nontransfected neurons (Fig. 4, A1, C1). This was not the case in neurons that had been transfected with GFP-tagged wild-type Sam68 (Fig. 4, D1). Neurons expressing GFP-Sam68 Δ C maintained a normal morphology (Fig. 4, A1, C3), indicating that the decrease in eEF1A did not arise from some general toxicity of the construct. The effect of GFP-Sam68 Δ C expression on endogenous eEF1A immunofluorescence was quantitated by using as internal reference value the average immunofluorescence of nontransfected neurons surrounding the transfected cells

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(see Materials and Methods). This quantitation procedure indicated that GFP-Sam68 Δ C reproducibly lowered the level of eEF1A immunofluorescence, whereas GFP-Sam68 did not (Fig. 4E). As a control, GFP-Sam68 Δ C did not affect the abundance of coexpressed mRFP. Since GFP-Sam68 Δ C functions in the cytoplasm, and presumably competes with endogenous Sam68 for binding sites in mRNA-protein complexes, this result suggests that endogenous Sam68 positively regulates the translation of eEF1A-encoding mRNAs.

Possible post-synaptic anchoring by Sam68

In non-neuronal cells, Sam68 is known to bind to membrane-associated signalling proteins (Lukong and Richard 2003). In neurons, given its association with dendritic mRNAs that become actively translated in response to local synaptic stimuli (Tsokas et al. 2005), we wondered whether Sam68 might physically interact with synaptic receptors. In synaptic membranes prepared from rat cortex, both the NMDA receptor and the post-synaptic density protein PSD95 could be co-immunoprecipitated with Sam68 (Fig. 5, lanes 1-6). By contrast, AMPA receptors were essentially absent from these immunoprecipitates (lanes 7-9), consistent with the notion that Sam68 was specifically recruited by the NMDA receptor complex. We could also detect Sam68 in NMDA receptor immunoprecipitates (lanes 10-11). Thus Sam68 might conceivably provide a platform for docking specific neuronal mRNAs and / or for recruiting specific translation factors at subsynaptic sites.

Discussion

The present study indicates that brain Sam68 is associated with a wide but selective range of cytoplasmic mRNAs, among which transcripts encoding long-term plasticity related proteins are prominent. Sam68 might collectively regulate the synthesis of an entire group of such proteins, in an activity-dependent fashion. The set of Sam68-associated mRNAs we found in brain differs from those reported in non-neuronal cell lines or lineages (Itoh et al. 2002; Paronetto et al. 2006; Tremblay and Richard 2006). Extensive comparison of the different sets is difficult, since the screening methods were different. However, two of the mRNAs, β -actin and dynein, were found in both the study by (Itoh et al. 2002) and our screen, providing a reliable positive control. Taken together, the various screens suggest that mRNA recognition by Sam68 is highly dependent on cellular context.

One of most enriched among the Sam68-associated cortical transcripts was the Eef1a1 mRNA. Although Eef1a1 is ubiquitous, the Sam68-Eef1a1 mRNA interaction was apparently not detected in the previous studies. This may be due to the sensitivity of the micro-array sampling procedure; alternatively, the interaction may be favoured by neuron-specific factors. However, it was possible to reconstitute the Sam68-Eef1a1 mRNA interaction in transfected HEK cells by using overexpressed, myc-tagged Sam68. The interaction is phylogenetically conserved since HEK 293 cells are of human origin; this is consistent with the >90% identity of human and rat Eef1a1 nucleotide sequences, including in the 3' untranslated region. Interestingly, based on free energy minimization, the entire 3'UTR of the Eef1a1 mRNA is predicted to fold into a single, long hairpin displaying at its extremity a U-rich terminal loop (Fig. 6). This is reminiscent of the U6 motif-containing stem-loop structure that mediates recognition of the β -actin mRNA by Sam68 *in vitro* (Itoh et al. 2002). However, our experiments do not rule out the possibility that the Sam68 – Eef1a1 mRNA association may

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3 be mediated by a third molecule. Further studies will be required to establish whether the 3'
4 UTR structure forms a direct binding site for Sam68 on the Eef1a1 mRNA.
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8 In dissociated cortical neurons, chemically induced depolarisation is known to generate bursts
9 of action potentials and to recruit some of the pathways that underlie synaptic plasticity.
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11 While causing the translational repression of multiple mRNA species, depolarisation activates
12 the translation of a select group of transcripts (Krichevsky and Kosik 2001). Our data indicate
13 that Eef1a1 belongs to this group. Eef1a1 encodes the ubiquitous isoform of translation
14 elongation factor eEF1A, which transports amino-acyl-tRNA to the A site of the ribosome
15 during protein synthesis. In addition, eEF1A also anchors mRNAs to the actin cytoskeleton,
16 and can bundle actin filaments according to a unique spatial arrangement (Liu et al. 2002a).
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18 Local provision of eEF1A may play a role in both biosynthetic and structural aspects of
19 synaptic plasticity, and perhaps establish a link between these two facets of long-term
20 synaptic change. In *Aplysia*, synthesis of the eEF1A homologue is instrumental to
21 maintenance of synapse-specific long-term facilitation (Giustetto et al. 2003). In the rodent
22 hippocampus, translation of the Eef1a1 mRNA is regulated by synaptic signals in complex
23 ways, which appear to differ between preparations. In hippocampal slices from young rats,
24 Eef1a1 translation has been shown to be rapidly induced during tetanus-induced LTP of CA3-
25 CA1 synapses, subject to activation of the rapamycin-sensitive mTOR / S6 Kinase pathway
26 (Tsokas et al. 2005). This finding is consistent with the fact that the Eef1a1 mRNA contains a
27 5' oligopyrimidine tract, which is a signature for mTOR-inducible mRNAs. Interestingly, in
28 adult rat hippocampus *in vivo*, LTP of dentate granule synapses induced redistribution rather
29 than neosynthesis of eEF1A protein. In contrast, long-term depression elicited by
30 pharmacological stimulation of metabotropic glutamate receptors (mGluRs) was
31 accompanied by a large increase in eEF1A protein, without change in Eef1a1 mRNA level
32 (Huang et al. 2005). Thus, the induction of eEF1A synthesis seems to be exquisitely sensitive

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to the spatio-temporal and cell-specific details of synaptic signalling, suggesting that multiple effectors regulate the translation of Eef1a1 in neurons. One of these effectors may be the RNA-binding protein FMRP, since in vitro and in cell lines, FMRP has been reported to bind and repress the Eef1a1 mRNA (Sung et al. 2003); furthermore it is tempting to speculate that removal of this putative inhibition underlies the effect of mGluRs on Eef1a1 translation (Bear et al. 2004). The present data suggest that Sam68 is also part of the regulatory machinery that controls neuronal Eef1a1 translation, with an action distinct from that of FMRP. Since depolarisation increased the recruitment of Sam68 to polysomal mRNAs, it is likely that Eef1a1 transcripts bound to Sam68 were engaged in translation. The finding that the dominant-negative GFP-Sam68 Δ C, but not GFP-Sam68(WT) inhibited eEF1A protein accumulation suggests that endogenous Sam68 may be required for the translational activation of Eef1a1 mRNA. Since the dominant-negative effect of GFP-Sam68 Δ C was already detected in unstimulated neurons, the low amount of Sam68 existing in the cytoplasm prior to depolarisation may contribute to generating the proportion of actively translated Eef1a1 mRNA seen under basal conditions. In line with this possibility, Sam68 has been reported to interact with RNA helicase A and with the mRNA export factor Tap (Reddy et al. 2000), two proteins known to promote translation of specific viral and cellular mRNAs (Hartman et al. 2006; Li et al. 2006). Kinases recruited by the C terminus of Sam68 might also be required to phosphorylate and release mRNA-bound translation inhibitory factors, similar to the case of hnRNP-K phosphorylation by Src (Ostareck-Lederer et al. 2002). GFP-Sam68 Δ C, which is defective for certain protein-protein interactions, may displace the endogenous Sam68 and associated translation factors out of the target mRNA; indeed, in the case of viral mRNAs, GFP-Sam68 Δ C functionally antagonizes RNA helicase A and Tap (Reddy et al. 2000). Upon depolarisation, the largely enhanced export of Sam68 to the cytoplasm may cooperate with the action of signalling cascades such as the mTOR pathway

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3 for translational stimulation of activity-responsive mRNAs exemplified by Eef1a1. A similar
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5 scheme may apply to certain non-neuronal systems; in the case of meiotic male gametes,
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7 Paronetto et al. (2006) recently confirmed the occurrence of signal-induced transfer of Sam68
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9 to the cytoplasm and its binding to polysomes.
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Legends to Tables and Figures

Table Ia

ID: Genbank identifier of cloned sequence on the NIA 15 K micro-array. Name: name of transcript homologous to the sequence on the array (available through <http://lgsun.grc.nia.nih.gov/cDNA/15k.html>). Mean z score and its standard deviation (n=3) were calculated for each transcript as described in the text. P values were calculated from z scores, assuming normal distribution of scores. Biological process or component associated with transcripts were identified by reporting Gene Ontology (GO) descriptors used to annotate the transcripts in the NIA database (available through <http://lgsun.grc.nia.nih.gov/cDNA/NIA-CloneSet-GoTermFinder.html>). Rightmost column: identification of the GO descriptors by their numbering in the Gene Ontology database.

Fig. 1

A. Scheme of the RNA-protein coimmunoprecipitation procedure, and image of a representative hybridization experiment. Green channel: Cy5-labelled probes obtained by amplification of polyA⁺ mRNA immunoprecipitated with the anti-Sam68 antibody. Red channel: Cy3-labelled probes obtained after immunoprecipitation with control (preimmune) antibody. **B.** Distribution of paired fluorescence values obtained in A. X-axis: specifically immunoprecipitated transcripts. Y-axis: adventitious transcripts. A large number of transcripts are more abundant in the specific immunoprecipitate than in the control. **C.** Within-sample reproducibility of z score measurements. The results of two different hybridization experiments performed with the same sample are compared. The regression line is shown in red. **D.** Between-sample reproducibility of z scores. The results of hybridization experiments performed on two different brain extracts are compared. The regression line is shown in red.

Note that mRNAs with highest values in either sample (to the right of the vertical green line and above the horizontal green line) are mostly confined to the top right quadrant. **E.** Functional profile of the 43 mRNAs displaying the highest mean z scores in the immunoprecipitates (see Table Ia). For each transcript, a generic function was registered (identified by Gene Ontology descriptors of the mRNA in the NIH database; see Table Ia). Black bars: number of occurrences of the indicated functional descriptors in the 43 mRNA sample. Empty bars: expected number of occurrences of the descriptors, based on their frequencies among mRNAs of the micro-array. The difference is highly significant ($p < 0.0001$, chi-square test), indicating enrichment for specific functional categories.

Fig. 2

A. Validation of specific micro-array hybridization results by RT-PCR. Cortical tissue extract was subjected to immunoprecipitation with anti-Sam68 (“IP”) or preimmune (“Pre”) IgG as shown; total mRNA was purified from the immunoprecipitates, and subjected to reverse transcription and amplification by PCR, using the indicated gene-specific primers (Table II). The PCR products were visualized by fluorescent staining with ethidium bromide. The expected sizes of PCR products are indicated (Eef1a1: 336 bp; Uch-L1: 187 bp; β -actin: 178 bp; Calm-1: 229 bp; Arc: 287 bp). Positive controls for the amplification reactions were obtained by performing PCR with total mRNA (“Tot”) directly derived from the input extract. No amplification took place without cDNA (“None” or “0”).

B. Reconstitution of the association between Sam68 and the Eef1a1 mRNA in cultured cells. HEK 293 cells were transfected with a myc-tagged Sam68 expression construct, or with control vector, as indicated. Cytoplasmic extracts were prepared and immunoprecipitated with anti-myc antibody, and subjected to RT-PCR with Eef1a1 (lanes 1, 3) or with β -tubulin (lanes

7,9) primers. As positive controls for amplification, RT-PCR products were generated with total mRNA from the transfected cells (lanes 2, 4, 8, 10) or from rat cortex (lane 5, 11). Lanes 6, 12: negative control for PCR.

C. The amounts of endogenous Eef1a1 mRNA in anti-myc immunoprecipitates from the indicated transfections were determined by quantitative (“real-time”) PCR (n=3 for each transfection). *a.u.*, arbitrary units.

Fig. 3

A. Depolarisation induces an increase in Eef1a1 mRNA translation. Cultured cortical neurons were incubated for 6 h in 5mM or 25 mM KCl, then cytoplasmic extracts were prepared and equal protein amounts were subjected to fractionation on sucrose density gradients. For each gradient, representative fractions corresponding to translationally inactive mRNPs, 80S monosomes, polysomes, and RNA granules were analysed by RT-PCR with specific primers to amplify the Eef1a1 mRNA. The PCR products were analysed side by side on an agarose gel (“-” and “+” lanes : gradient fractions from control and depolarised neurons; respectively) and visualized by fluorescent staining with ethidium bromide. Greyscale was inverted to improve contrast.

B. Depolarisation increases the amount of eEF1A protein. Cortical neurons were treated as in A, then lysed, and equal amounts of total protein were analysed by SDS-polyacrylamide gel electrophoresis and immunoblotting. Left: the same blot was probed first with anti-eEF1A monoclonal antibody, then with anti-actin antibody. Right: the relative amount of eEF1A in each lane was measured by densitometric scanning, and normalized to the amount of actin in the same lane to correct for gel loading error. The histogram shows the average (+/- S.D.) of normalized eEF1A amount in each condition.

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C. Depolarisation enhances the association of Sam68 with polysomal RNA and RNA granules. Cytoplasmic extracts from control and depolarised neurons were prepared and fractionated as in **A**. The fractions were analysed by immunoblotting with anti-Sam68 antibody (left). The distribution of RNA along the gradient was determined by continuous monitoring of UV absorbance at 254 nm during collection of the fractions (right). The various types of ribonucleoproteic assemblies are indicated.

Fig. 4

Effect of dominant-negative Sam68 on eEF1A expression in cultured hippocampal neurons.

A-C: The GFP-Sam68 (WT) and GFP-Sam68 Δ C proteins were expressed in hippocampal neurons, together with monomeric Red Fluorescent Protein (mRFP) as a morphological marker. Five days later (at 14 DIV), the neurons were fixed and stained by immunofluorescence with anti-eEF1A antibody; mRFP, GFP-Sam68 and endogenous eEF1A were simultaneously visualized by triple-colour fluorescence microscopy.

A1-A3: close-up view of the three fluorescent markers in a GFP-Sam68 Δ C-transfected neuron and a nearby, untransfected neuron (wide-field microscopy). **A1:** Note the lack of eEF1A immunostaining in the transfected cell (arrow), compared to the neighbouring cell. Scalebar: 13 μ m. **B:** clustered distribution of the GFP-Sam68 Δ C protein within the dendrite of a hippocampal neuron (single plane of a confocal z series). Scalebar: 4.7 μ m.

C1-C3: representative field showing GFP-Sam68 Δ C transfected neurons and neighbouring untransfected cells at low magnification. Scalebar: 65 μ m. **C1:** the relative intensity of eEF1A immunofluorescence is represented by pseudocolour coding. Perinuclear staining is brighter in untransfected neurons (arrowheads) than in transfected (arrows). **C3:** normal neuronal morphology of the transfected cells.

D1-D3: same as C1-C3 with GFP-Sam68WT transfected neuron.

E: the eEF1A immunofluorescence was measured (as mean grey level per pixel) in transfected neurons and in surrounding, nontransfected neurons of the same field. The ratio between the fluorescence of transfected neurons and the average fluorescence of surrounding nontransfected cells was then calculated for each transfected neuron. The histogram bars show the average (\pm S.D.) of these ratios for neurons transfected with GFP-Sam68 Δ C and with GFP-Sam68WT. Neurons expressing GFP-Sam68 Δ C had a ratio lower than 1, indicating a decrease in endogenous eEF1A. Neurons expressing GFP-Sam68WT had a ratio close to 1. The difference between GFP-Sam68 Δ C and GFP-Sam68WT was significant ($p=0.03$, two-sided Mann-Whitney U test) ($n=3$ experiments).

F: schematic depiction of the wild-type and C-terminally deleted, dominant-negative Sam68 proteins. The major protein domains are shown as boxes. KH: core RNA-binding module (hnRNP-K homology domain). P1 to P5: proline-rich motifs. Y: tyrosine-rich region. NLS: nuclear localisation signal.

Fig. 5

Binding of Sam68 to the post-synaptic NMDA receptor complex. Purified synaptic membranes were solubilized under non-denaturing conditions, and subjected to immunoprecipitation with the indicated antibodies. Lanes 1, 4, 7: aliquot (10%) of input lysate. After detecting PSD-95, the blot of lanes 4-6 was stripped and reprobed to detect GluR2/3 subunits of AMPA receptors, shown in lanes 7-9. Lane 10: control immunoprecipitation with a mouse monoclonal IgG, class-matched to the anti-NR1 antibody.

Fig. 6

The 3' UTR sequences of human EEF1A1 and rat Eef1a1 mRNAs have been aligned using Clustal W (v. 1.83). The first nucleotide of each sequence is that immediately following the

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2
3 stop codon, and the last nucleotide corresponds to the major polyadenylation site. Stars
4 indicate identical residues. Using the Rnaalifold programme of the Vienna RNA package (Ivo
5 L Hofacker, <http://www.tbi.univie.ac.at/~ivo/RNA/>), the predicted folding of the aligned
6 sequences displays a large stem structure with a U-rich terminal loop (overlined).
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Table Ia Identity and generic function of major Sam68-associated mRNAs.

ID	Name	Mean		p value	Process or component	GO
		z	± S.D.			
AW557547	Translation elongation factor 1 alpha 1, Eef1a1	2.66	± 0.97	0.004	protein biosynthesis	06412
BG086025	ATPase, H ⁺ transporting, V1 subunit B, isoform 2, Atp6v1b2	2.55	± 1.07	0.005	proton transport	15992
BG076172	Calmodulin-1, Calm1	2.36	± 1.11	0.009	signal transduction	07165
BG088462	Zfhx1a, C2-H2 zinc finger protein homeobox 1a	2.35	± 1.29	0.009	regulation of transcription	06355
AW558451	Reticulon 3, Rtn3	2.28	± 0.60	0.011	protein transport	15031
BG088310	Prosaposin	2.1	± 0.88	0.018	metabolism	08152
BG071424	BRI3	2.04	± 0.28	0.021	integral to membrane	16021
BG073409	Stathmin 1, Stmn1	1.98	± 0.48	0.024	signal transduction	07165
BG065012	14-3-3 eta, Ywhah	1.86	± 0.82	0.031	signal transduction	07165
BG075901	Gamma-aminobutyric acid receptor subunit, Gabra3	1.86	± 0.36	0.031	synaptic transmission	07268
BG076067	Inositol hexaphosphate kinase 1	1.86	± 0.50	0.032	metabolism	08152
BG085717	14-3-3 epsilon, Ywhae	1.85	± 0.84	0.032	signal transduction	07165
BG082675	ADP-ribosylation factor 1, Arf-1	1.85	± 0.90	0.032	signal transduction	07165
BG075782	Protein phosphatase 1, catalytic subunit, gamma, Ppp1cc	1.83	± 0.76	0.034	signal transduction	07165
BG075243	Zinc transporter ZnT-3	1.8	± 0.27	0.036	cation transport	06812
BG077677	Beta-actin	1.78	± 0.53	0.037	cytoskeleton	07010
BG065113	Branched chain aminotransferase, bcat-1	1.78	± 1.51	0.038	metabolism	08152
BG064914	Malate dehydrogenase, Mor2	1.73	± 0.61	0.042	metabolism	08152
BG087365	Neuroplastin	1.69	± 0.86	0.046	surface receptor signal transduction	07166
BG073370	Tetratricopeptide repeat domain 3, Ttc3	1.69	± 1.29	0.046	protein binding	05515
BG074109	Heat shock protein 1, alpha, Hspca	1.67	± 1.15	0.047	chaperone activity	03754
BG088451	Tissue inhibitor of metalloproteinase 2, Timp2	1.61	± 0.77	0.053	extracellular matrix	05578
BG075319	Mitogen activated protein kinase 3, pp44 MAPK, Erk1	1.61	± 0.57	0.054	signal transduction	07165
BG074398	Matrix glycoprotein Sparc1 / hevin	1.54	± 0.64	0.062	extracellular matrix	05578
BG073255	Tetraspanin TM4-A	1.5	± 0.86	0.066	integral to membrane	16021
BG082737	Proline-arginine-rich end leucine-rich repeat protein, Prelp	1.46	± 0.64	0.073	extracellular matrix	05578
BG073185	BTB (POZ) domain containing 4, KBTBD4	1.41	± 1.05	0.079	regulation of transcription	06355
BG088334	Vacuolar protein sorting 41	1.4	± 0.05	0.081	protein transport	15031

BG087861	Dynein, cytoplasmic, intermediate chain 1, Dncic1	1.39 ± 0.39	0.082	microtubule-based movement	07018
BG085988	Heterogeneous nuclear ribonucleoprotein H2, Hnrph2	1.35 ± 0.91	0.089	mRNA processing	06397
BG073339	Splicing factor, arginine/serine-rich 5, Srp40 / HRS	1.31 ± 0.72	0.096	mRNA processing	06397
AW553322	G protein-coupled receptor associated sorting protein 1	1.27 ± 0.86	0.102	protein transport	15031
BG073253	Rab11B	1.23 ± 1.25	0.109	protein transport	15031
BG086823	Myeloid leukemia factor 2, parathymosin	1.23 ± 0.78	0.109	immune response	06955
BG072963	Dynein heavy chain	1.23 ± 0.21	0.110	microtubule-based movement	07018
BG086722	Ubiquitin carboxyl-terminal esterase L1, UCH-L1	1.16 ± 0.98	0.122	ubiquitin-dependent protein catabolism	06511
BG086087	Fanconi anemia-associated protein of 100 kDa, Faap100	1.11 ± 1.31	0.134	response to DNA damage	06974
BG086010	Dynein intermediate chain 2, Dync1i2	1.11 ± 0.12	0.134	microtubule-based movement	07018
BG088587	Beclin 1 (coiled-coil, myosin-like BCL2-interacting protein)	1.09 ± 0.43	0.137	autophagy	06914
BG080018	Nuclear protein Nulp1/ bHLH Tcf25	0.99 ± 0.72	0.161	regulation of transcription	06355
BG088909	Transducin-like enhancer of split 6, TLE6 / Grg6	0.99 ± 1.01	0.162	regulation of transcription	06355
BG076556	Farnesyltransferase beta subunit	0.98 ± 1.10	0.162	regulation of cell proliferation	42127
BG069542	hnRNP-E1, poly(rC)-binding protein 1	0.98 ± 1.38	0.163	mRNA processing	06397

Table Ib Sam68-associated transcripts with documented neurobiological function.**Transcripts involved in synaptic plasticity**

Translation elongation factor 1 alpha 1, Eef1a1
 ATPase, H⁺ transporting, V1 subunit B2, Atp6v1b2
 Calmodulin-1, Calm1
 14-3-3 eta, Ywhah
 Protein phosphatase 1, catalytic subunit, gamma, Ppp1cc
 Beta-actin
 Mitogen activated protein kinase 3, pp44 MAPK, Erk1

Published evidence

Tsokas et al. 2005, Huang et al. 2005
 Munton et al. 2007
 Xia and Storm 2005
 Simsek-Duran et al. 2004
 Hu et al. 2006
 Eom et al. 2003
 Thomas and Huganir 2004

Transcripts involved in neurodegeneration

14-3-3 epsilon, Ywhae
 Tetratricopeptide repeat domain 3, Ttc3
 Reticulon 3, Rtn3
 Prosaposin
 Bri3
 Ubiquitin carboxyl-terminal esterase L1, Uch-L1
 Beclin-1

Published evidence

Toyo-oka et al. 2003
 Berto et al. 2007
 He et al. 2004
 Matsuda et al. 2004
 Wickham et al. 2005
 Liu et al. 2002, Gong et al. 2006
 Pacheco et al. 2007

Transcript involved in psychiatric disease

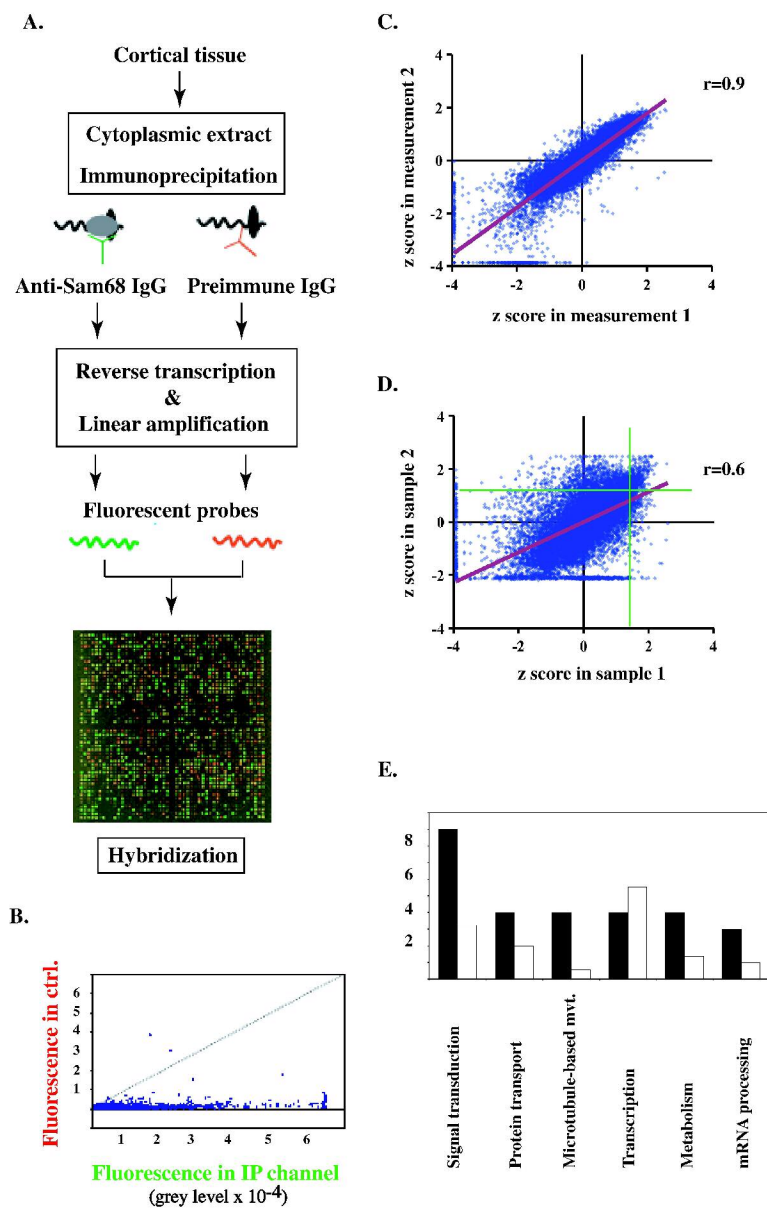
Gamma-aminobutyric acid receptor, subunit Gabra3

Published evidence

Massat et al. 2002

Table II: PCR primers used in this study.

Transcript	Primers
Arc	Forward: 5'-TGTTGGTAACTGCTCGTGTCTGTAG-3' Reverse: 5'-TTTGAGGTAAGATGGTGTGGGC-3'
β -Actin	Forward: 5'-TGGCACCACACTTTCTACAATGAG-3' Reverse: 5'-AGGCATACAGGGACAACACAGC-3'
Calm-1	Forward: 5'-TCTTCAAGTGCCCCAATCCC-3' Reverse: 5'-TCAAGTCCACAGACACAGCCTACTC-3'
Eef1a1	Forward: 5'-GGACAGCAAAAATGACCCACC-3' Reverse: 5'-TTGATGACACCCACAGCAACTG-3'
Gephyrin	Forward: 5'-AAAGAGGGGAGTGTGTTTTGGC-3' Reverse: 5'-AAGGCATTGAGTAAGTCATCTGGG-3'
NR1-1	Forward: 5'-CGTATCCTAGGCATGGTGTGG-3' Reverse: 5'-CCCGGTGCTCGTGTCTTTGG-3'
β -Tubulin	Forward: 5'-CAGACCAGACAACCTTCGTTTTTCG-3' Reverse: 5'-GTATCAGACACTTTGGGCGAGG-3'
Uch-L1	Forward: 5'-TGCTGCTGCTGTTTCCCCTC-3' Reverse: 5'-CCATCCTCAAATCCCAGGTTG-3'

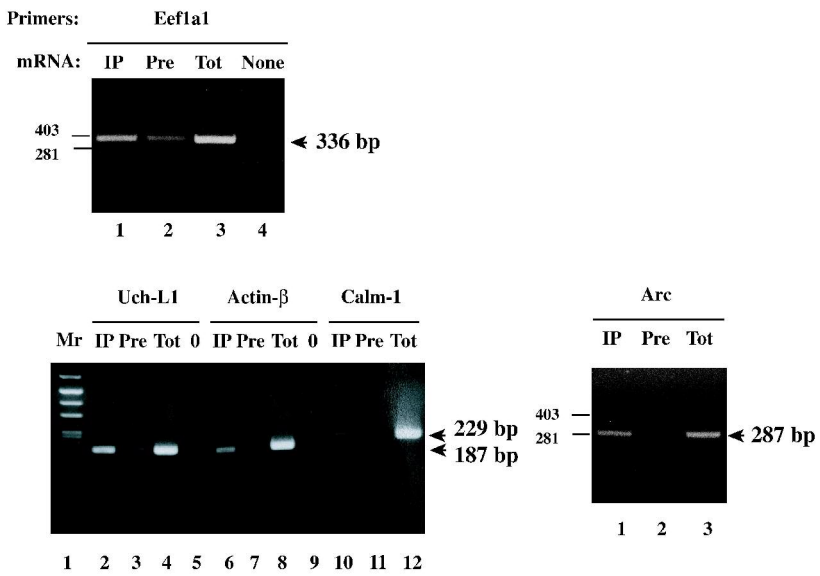


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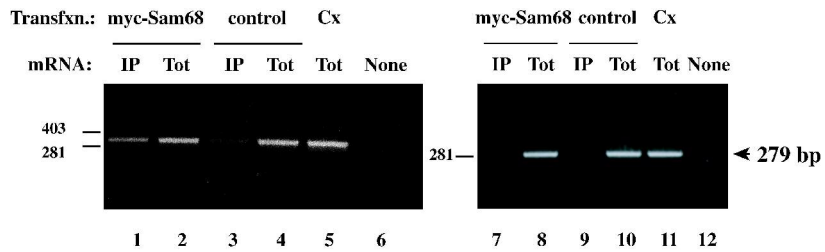
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inserm-00451270, version 1, 28 Jan 2010

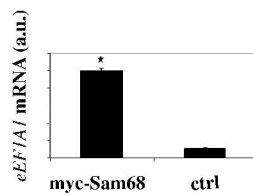
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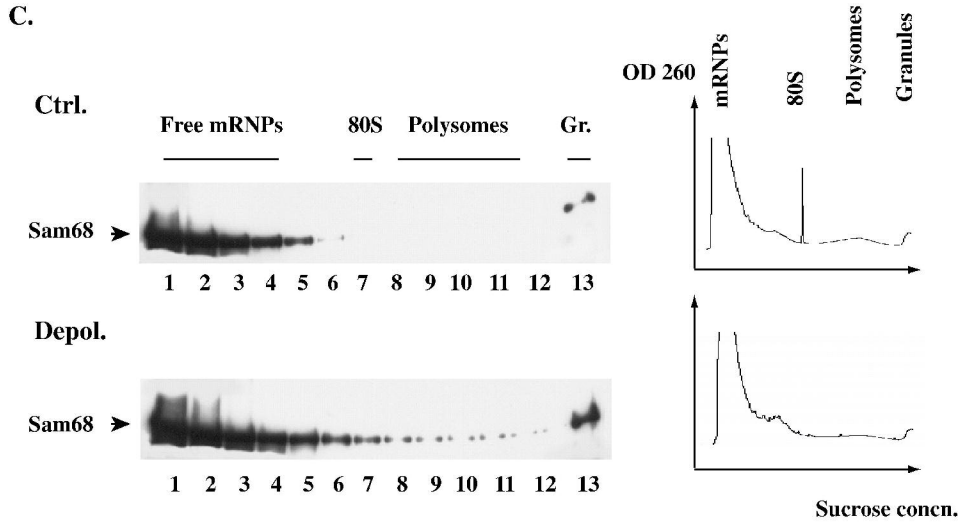
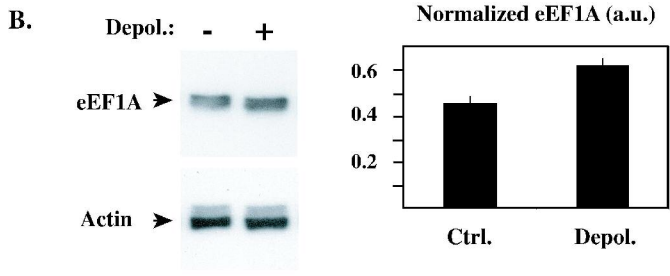
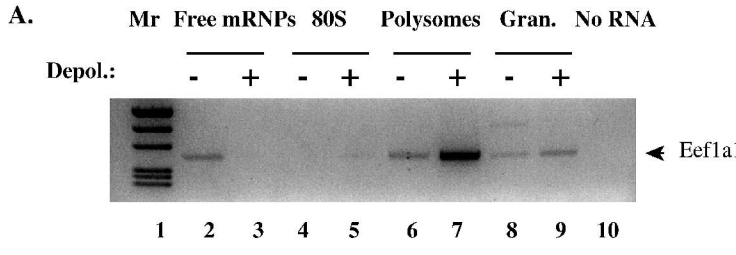
C.



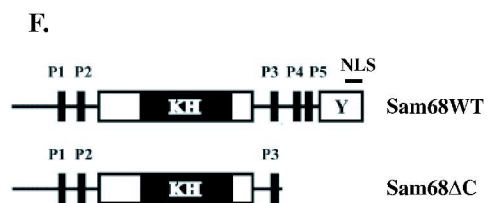
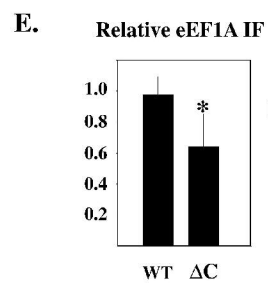
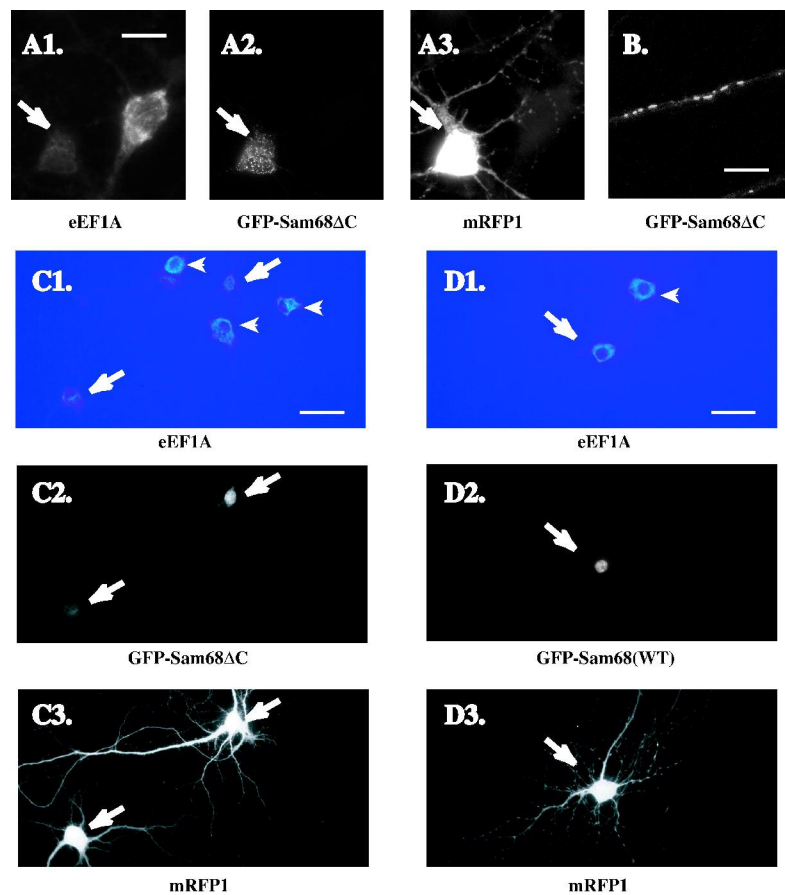
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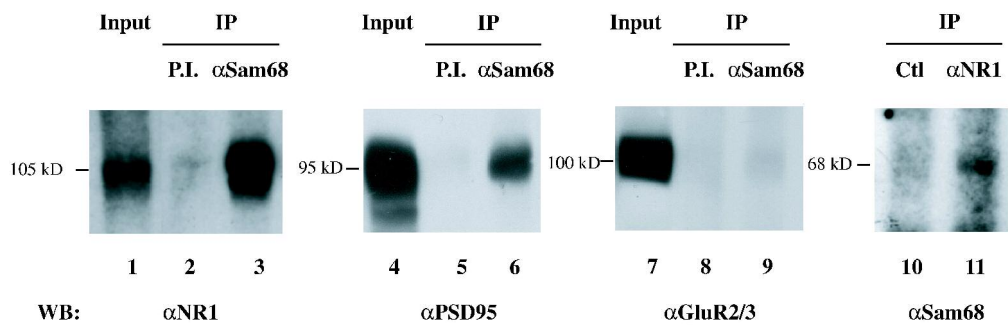
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194x276mm (600 x 600 DPI)

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194x62mm (600 x 600 DPI)

Peer Review

