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Physiol Rev 89:1079-1103, 2009. doi:10.1152/physrev.00013.2009

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The Function of Activity-Regulated Genes in the Nervous System

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Loeblich S, Nedivi E. The Function of Activity-Regulated Genes in the Nervous System. *Physiol Rev* 89: 1079–1103, 2009; doi:10.1152/physrev.00013.2009.—The mammalian brain is plastic in the sense that it shows a remarkable capacity for change throughout life. The contribution of neuronal activity to brain plasticity was first recognized in relation to critical periods of development, when manipulating the sensory environment was found to profoundly affect neuronal morphology and receptive field properties. Since then, a growing body of evidence has established that brain plasticity extends beyond development and is an inherent feature of adult brain function, spanning multiple domains, from learning and memory to adaptability of primary sensory maps. Here we discuss evolution of the current view that plasticity of the adult brain derives from dynamic tuning of transcriptional control mechanisms at the neuronal level, in response to external and internal stimuli. We then review the identification of “plasticity genes” regulated by changes in the levels of electrical activity, and how elucidating their cellular functions has revealed the intimate role transcriptional regulation plays in fundamental aspects of synaptic transmission and circuit plasticity that occur in the brain on an every day basis.

I. INTRODUCTION

Groundbreaking work in the 1960s and 1970s demonstrated that brain plasticity is shaped by sensory input during critical periods of development. Experimental manipulation of the sensory environment was shown to profoundly affect neuronal morphology and receptive field properties (120, 160, 274). Later studies demonstrated that brain plasticity is not limited to development but persists in adulthood and seems to be an inherent feature of everyday brain function, critical for learning and memory, and the adaptability of primary sensory maps (reviewed in Refs. 31, 114, 116, 180, 278).

The understanding of neuronal adaptability in response to the external environment is largely owed to the discovery in other cell systems of signal transduction pathways that couple extracellular cues to changes in

transcriptional activity. The study of tumor viruses and their mechanism of action opened the door to our understanding of the cellular pathways that control cell growth and proliferation, and in the mature nervous system mediate neuronal responses to electrical activity.

The first molecules shown to translate extracellular signals to transcriptional programs that alter cell properties were transcription factors activated within minutes of stimulation by growth factors and encoded by the cellular counterparts of tumor virus oncogenes. Early studies of the Rous sarcoma virus (RSV) revealed that its transforming function resided in a single gene called *src* (25, 125, 210). In work for which they received the Nobel prize, Bishop and Varmus (210) showed that the viral *src* gene, *v-src*, was derived from a normal cellular protooncogene *c-src*. Clues to the machinery of intercellular signaling via growth factors and mitogens came when it was discov-

ered that in normal cells protooncogenes function in signaling roles that regulate cell growth and proliferation (189, 256, 272). Cellular oncogenes were found to encode proteins that represent all major components of the growth factor response pathway: from the growth factor *sis* (42, 62, 133, 134) and growth factor receptor *erbB* (64, 218), to the small G protein *ras* (59, 189, 256) or nonreceptor tyrosine kinase *src* (25, 125, 210), and finally nuclear proteins, such as *myc* (3, 71, 105, 146), *fos* (55), or *jun* (26, 88, 202).

It was known that application of growth factors like platelet-derived growth factor (PDGF) leads to rapid activation of gene expression despite the presence of protein synthesis inhibitors such as cycloheximide. These rapid response genes were termed immediate-early genes (IEGs) (46). Nuclear run-off transcription assays after growth factor treatment revealed that the protooncogenes *c-fos* and *c-myc* were among the IEGs (96, 190). Several facts implicated *c-fos* in regulation of gene expression. It was known to encode a nuclear protein (55) associated with chromatin and capable of binding DNA cellulose in vitro (226, 233). It turned out that *c-fos* and *c-jun* are specific members of inducible gene families whose products associate combinatorially to form dimeric complexes that function as transcriptional activators (54). These transcription factors, in turn, help induce expression of second wave genes, termed delayed early genes.

Studies of the protooncogene *c-fos* also pioneered the identification of *cis*- and *trans*-acting regulatory elements that play a role in controlling IEG induction (269). The first operationally defined regulatory *cis*-element, the serum response element (SRE), was initially mapped in the promoter region of the *c-fos* gene (91, 219, 260). The *c-fos* promoter was also found to contain a cAMP responsive element (CRE) (243), common to cAMP-regulated genes (185). The search for *trans*-acting protein factors that bind to CRE identified a bZip transcription factor termed CREB (CRE-binding protein) (111). The CREB protein family was found to comprise more than 10 members (102) able to heterodimerize in multiple combinations and associate with Fos and Jun as heterodimers, further fine-tuning the binding specificity of the complex (101). The *CREB* gene gives rise to three main activating isoforms: α (94), β (22), and Δ (111, 287). In addition, alternative splicing appears to be a common feature of CREB family genes and is used as a mechanism to generate both transcriptional activators and repressors. The gene encoding the CREB homolog CREM (cAMP-responsive element modulator) also shows IEG induction kinetics, is subject to cAMP-dependent activation (183), and is differentially spliced to encode both activators and repressors of cAMP-dependent transcription (reviewed in Ref. 152). The *CREM* products can heterodimerize with each other, as well as with CREB, adding further com-

plexity to the regulation of cAMP-induced transcription (154).

In 1989, Gonzalez and Montminy (94) demonstrated that phosphorylation is a key mechanism in CREB regulation. They showed that elevated cAMP levels led to phosphorylation of CREB at a specific residue, serine-133. Phosphorylation of CREB at this particular residue is a prerequisite for its ability to trigger gene transcription in *trans*, as was first assessed for the *somatostatin* gene (94). The phosphorylated CREB protein associates with CBP (CREB-binding protein) (45) and mediates interaction with TFIIB, a major component of the transcription machinery of the RNA polymerase II complex (150). Termination of CREB signaling can be achieved by dephosphorylation through protein phosphatase 1 (PP-1), or PP-2a (201).

In parallel to in-depth studies of protooncogene IEGs, such as *c-fos*, other studies greatly expanded the pool of known inducible genes by administering different stimuli, such as mitogens and phorbol esters (4, 155, 161, 253), leading to the more general view that gene transcription in the nucleus is part of the cellular response program to alterations in signaling from outside the cell. An integral aspect of this cellular response program is the biphasic nature of transcriptional activation, with IEG induction closely followed by expression of a second wave of delayed early genes. Another important feature is that multiple signals and intracellular pathways can influence IEG transcriptional activation via an assortment of *cis*- and *trans*-acting elements, and nuclear IEGs themselves comprise a diverse array of factors that can potentially act combinatorially to differentially affect distinct second wave gene sets.

II. TRANSCRIPTIONAL ACTIVATION OF IMMEDIATE EARLY GENES IN THE NERVOUS SYSTEM

Generalization of these concepts to neurons started before they were fully understood and articulated, with demonstration that treatment of pheochromocytoma (PC12) cells with nerve growth factor (NGF) stimulates expression of *c-fos* (56, 95, 149). Surprisingly, in this case *c-fos* induction was associated with events related to cell differentiation rather than cell proliferation. This suggested that the role of IEGs and their activation of a late response transcriptional program could differ according to the differentiation and physiological state of the stimulated cell and was not necessarily restricted to proliferative cells. In addition, *c-fos* expression in PC12 cells could be induced by agents other than growth factors and mitogens, including depolarizing stimuli and voltage-gated calcium influx (187). These studies provided motivation for examining expression of *c-fos* and other

IEGs in the brain, and their *in vivo* induction in response to various neural stimuli, most notably seizure paradigms.

Immunostaining revealed Fos-positive neurons throughout the mouse and rat brain (65, 186). Basal levels of *c-fos* expression could be robustly increased by administration of convulsants such as Metrazol (66, 186, 250) or picrotoxin (231), or by electrical kindling (67). The IEGs *c-jun*, *zif/268*, and *junB* were also found to be rapidly and coordinately activated in seizure paradigms (231). This led to the hypothesis that IEGs may play a part in normal neuronal function and that their increased levels were part of a programmed genomic response to intense stimulation analogous to the genomic response of nonneuronal cells to growth factors (231). However, while one could speculate that convulsant-induced transcription was merely an amplification of normal neuronal responses, it had yet to be shown that similar changes in gene expression could be detected after more natural, physiological levels of stimulation.

This critical point was subsequently addressed in the late 1980s. Several studies demonstrated that changes in IEG transcription factor mRNA levels could occur in response to physiological stimuli. For example, water deprivation resulted in Fos protein expression in hypothalamic neurons known to be activated by dehydration (232). Also, activation of small-diameter cutaneous sensory afferents by noxious heat or chemical stimulation rapidly induced *c-Fos* immunoreactivity (123) as well as *zif/268* gene and protein expression (276) in the superficial layers of the spinal cord dorsal horn. Activation of low-threshold cutaneous afferents resulted in Fos-labeled cells with a different laminar distribution. Neither type of stimuli elicited Fos expression in the dorsal root ganglia, nucleus gracilis, or ventral horn (123). Similarly, baseline *zif/268* mRNA levels in the visual cortex of rats was significantly increased by exposure to light after dark-adaptation and decreased after retinal action potential blockade by monocular intravitreal injection of TTX, whereas *c-jun* mRNA levels were unchanged by these treatments (277). Interestingly, *zif/268* mRNA baseline levels were NMDA receptor dependent and dropped significantly throughout cortex and hippocampus upon treatment with the NMDA receptor antagonist MK-801, whereas *c-fos* mRNA levels were unaffected by NMDA receptor blockade.

These findings provided strong support for the idea that endogenous IEG expression in the brain is a result of normal synaptic activity and could therefore be influenced by neuronal stimulation or activity blockade in a pathway and stimulus specific manner. A side product of these developments was the realization that if IEG activation was part of a neuronal response program to natural stimuli, their expression could be used as a metabolic marker for mapping functional pathways at a cellular

level (186, 232). There are excellent examples of successful implementation of this strategy (16, 83, 98, 241, 294), although there are concerns with strictly equating IEG expression with neuronal activity (reviewed in Refs. 100, 141).

III. TRANSCRIPTIONAL ACTIVATION AS THE BASIS FOR LONG-TERM SYNAPTIC PLASTICITY

The realization that synaptic activity induces gene transcription in mature neurons dawned as it was becoming clear that synaptic activity could also drive long-term changes in neuronal structure and function. This led to the proposal that electrical and chemical stimuli that produce long-term change in neurons, such as during learning and memory, act through a mechanism analogous to that of growth factors, namely, through second messenger pathways leading to transcriptional activation (reviewed in Refs. 21, 93).

It had been known for two decades that protein synthesis inhibitors could affect memory retention in mammals, first demonstrated in mice by intracerebral injection of the protein synthesis inhibitor puromycin (82). However, early studies monitoring behavioral performance could not separate the cellular and subcellular effects of the applied drugs from potential effects on aspects of behavior such as attention and motivation. Studies in *Aplysia* were the first to demonstrate that transcriptional activation and subsequent protein translation were critical for learning-related synaptic plasticity (184). In the intact animal, gently touching the siphon leads to its retraction, as well as gill retraction. This defense response could be sensitized by repeated stimulation such that sensitization could last for minutes (short-term sensitization, STS) to hours or days (long-term sensitization, LTS) (33, 39). The circuitry that controls this behavior *in vivo* involves sensory neurons from the siphon that connect to motor neurons in the gill. The sensory-motor-neuron synapses are modulated by facilitating interneurons driven by sensory neurons from the tail (86). Unlike STS, LTS is seen only after repeated stimulation and is mediated by serotonin (137). In this system serotonin acts as a neuromodulator and activates the cAMP-PKA pathway. Cocultured *Aplysia* sensory and motor neurons recapitulate the STS and LTS seen *in vivo*, with the long-term facilitation (LTF) of synaptic connections paralleling behavioral LTS in its requirement for multiple spaced applications of serotonin (184, 224). In an elegant study, Montarolo et al. (184) demonstrated that application of protein or RNA synthesis inhibitors to the *Aplysia* coculture system blocks LTF of sensorimotor connections, but has no effect on short-term facilitation (184). Application of the inhibitors during or immediately

after stimulation with serotonin suppressed LTP, whereas administration before or after the stimulation was ineffective. This is in line with behavioral studies in mammals, where long-term memory is suppressed after administration of RNA or protein synthesis inhibitors during or immediately after learning (82, 182, 223; reviewed in Refs. 58, 108, 179). The *Aplysia* studies clearly demonstrated that long-term plasticity and its dependence on RNA and protein synthesis derives from cellular properties that are independent of the complex architecture or intact circuitry of the living animal.

Hints that transcriptional activation mediated by CREB is essential for synaptic plasticity also came from the *Aplysia* culture system, since bath application of cAMP could be used in place of serotonin to induce LTP (234). Administration of oligonucleotides containing CRE sequences resulted in selective loss of LTP, but not STF in this preparation (57), and repeated serotonin application was shown to induce CREB phosphorylation and activation of CREB-dependent gene transcription (136). Meanwhile, studies in PC12 cells revealed that the CRE element in the *c-fos* promoter was one and the same as the calcium-responsive element (CaRe), and CREB becomes phosphorylated at serine-133 in response to both membrane depolarization and elevation of intracellular Ca^{2+} levels, subsequently leading to transcriptional activation of the *c-fos* gene (243). This study also demonstrated that cAMP and Ca^{2+} signaling had synergistic effects on CREB-dependent *c-fos* transcription in vitro, suggesting that these two signaling pathways converge on the CREB protein to regulate activity-dependent transcription in neurons.

In vertebrates, studies of long-term synaptic plasticity at the cellular and synaptic level mainly focused on the hippocampus, a region of the brain known for its importance in learning and memory. In 1973 Bliss and Lømo (24) reported the phenomenon of long-term potentiation (LTP), whereby high-frequency stimulation of the perforant path produces enhanced transmission in the dentate gyrus lasting hours to weeks. Similar to behavioral memory, LTP in hippocampal slices has distinct temporal phases, an early phase lasting 1–3 h after tetanic stimulation, and a persistent late phase lasting at least 8 h (late LTP, L-LTP) (227). Like hippocampal-dependent learning, L-LTP in vitro and in vivo requires repetitive stimulation and is dependent on coincidence detection by the NMDA receptor (49, 188). These attributes and the fact that it can be induced in brain slices in vitro have contributed to the emergence of LTP as the premier synaptic model of memory in the hippocampus, and more generally as a model for activity-dependent synaptic plasticity in the vertebrate brain (23).

Two groups that first monitored the expression of IEGs in response to LTP (48, 276) found that high-frequency stimulation of the perforant path in rat hippocampus at an intensity and frequency sufficient for LTP induc-

tion also markedly increased *zif/268* mRNA in the dentate gyrus. Other IEG mRNAs, such as *c-fos*, *c-jun*, and *junB*, were less consistently increased under the same experimental conditions, in line with earlier findings (63). While Cole et al. (48) found that stimulation of convergent inhibitory synaptic inputs known to block LTP also blocked IEG induction, Wisden et al. (276) using a similar protocol but slightly different placement of the stimulating electrode, saw a bilateral increase in mRNA levels of *zif/268*, *c-fos*, and *c-jun* alike. This finding indicated that upregulation of IEG mRNA is not always indicative of LTP (276). However, induction of LTP was always accompanied by concomitant *zif/268* mRNA expression. The fact that *zif/268* mRNA upregulation and LTP induction by high-frequency stimulation both require similar stimulus conditions, including activation of the NMDA receptor (48), argued that they were regulated by the same synaptic mechanisms. Consistent with this, induction of L-LTP in hippocampal slices produced either by tetanic stimulation or application of a membrane-permeable cAMP analog could be blocked by inhibitors of protein synthesis or transcriptional blockers (85, 118, 200). The similarity to the *Aplysia* system was further established with the demonstration that stimuli that induced L-LTP in CA1 of the hippocampus could also induce CRE-mediated gene expression (126) and CREB phosphorylation (60). L-LTP-induced gene expression could be blocked by PKA inhibitors or L-type Ca^{2+} channel blockers (117), again implicating CREB as a convergence point for both cAMP and Ca^{2+} signaling.

The first conclusive test of whether CREB was required for memory was performed in *Drosophila*, where genetic screens for learning and memory mutants identified several mutants with defects in genes encoding key enzymes that regulate intracellular cAMP levels (reviewed in Ref. 266). Inducible overexpression of a dominant negative truncated CREB protein containing the CRE-binding domain but lacking transactivation activity interfered with the formation of long-term memory if induced prior to training, without affecting short-term memory (290). Attempts to show the requirement for CREB in long-term memory using a knockdown approach in mice was complicated by the existence of multiple CREB isoforms (reviewed in Ref. 246). Initial results from analysis of *CREB α* ^Δ knockout mice were consistent with the *Aplysia* and *Drosophila* studies in that short-term memory was unimpaired whereas long-term memory was severely compromised (28), despite upregulation of the CREB β -isoform in these mutants. In contrast, Gass et al. (90) found that *CREB α* ^Δ knockout mice with a different genetic background did not show deficits in social transmission of food preference, nor in dentate gyrus or CA1 LTP in slice preparations. A strain in which all CREB isoforms were reduced showed clear behavioral deficits in the Morris water maze (90), although concerns

remain that the mutant mice may be impaired in behavioral flexibility, rather than spatial memory. Another study used conditional CREB knockouts and found both hippocampal LTP and long-term depression (LTD) to be normal (12). Also, learning and memory in the water maze task were not grossly affected. Surprisingly, the mutant mice showed impairment in a conditioned taste aversion learning task, thought to be hippocampus independent. These studies cast a controversial light on the findings of CREB deletion work in mammals and call to mind that gene dosage, genetic background, and compensatory mechanisms (22, 122) may confound the interpretation of such findings.

Mouse knockout studies of other IEG transcription factors linked to activity-dependent gene transcription and synaptic plasticity further validated the view that the cellular underpinnings of activity-dependent synaptic plasticity and learning and memory were one and the same. *Zif/268* mutant mice (259) that lack the *Zif/268* protein were found to be defective in L-LTP, indicating that *zif/268* mRNA expression is not just correlated with LTP, but is in fact required for its expression (135). Long-term memory was also impaired in *zif/268* mutant mice when assessed by a variety of behavioral tests, including object recognition and spatial memory tasks. Genetic evidence for a role of c-Fos in memory formation came from *c-fos* knockout mice that displayed impaired spatial memory in a water maze task (213), although it could not be ruled out that this was partially due to developmental abnormalities in the *c-fos* mice (132).

Thus, over the course of 15 years, three model systems, *Aplysia*, *Drosophila*, and mouse, have contributed to our understanding of how behavioral plasticity, such as is manifested during learning and memory, derives from activity-dependent plasticity at the synaptic level and is mediated by transcriptional activation of IEG transcription factors. The transcription factor CREB was key to establishing this sequence and emerged as a potential convergence point for two major signaling pathways communicating extracellular signals to the neuronal nucleus: depolarization-induced Ca^{2+} signaling by ionotropic glutamate receptors, and cAMP linked catecholamine or growth factor signaling. The importance of these pathways for activity-dependent neuronal function and malfunction cannot be overstated and has recently been comprehensively reviewed by Greenberg and colleagues (47, 80).

IV. SEARCH FOR ACTIVITY-REGULATED GENES

The above studies lent credence to the idea that in postmitotic neurons IEG activation is a normal response to synaptic activity and the first step towards long-term activity-dependent plasticity. However, they all focused

on IEGs, such as *c-fos*, *c-jun*, or *zif/268*, identified in screens for rapid response to growth factors. It was clear that to fully understand the activity-dependent genetic programs utilized by neurons in the mammalian brain, one had to screen for genes that were induced by neuronal activity, rather than by growth factor stimulation.

In the early 1990s, several groups set out to identify genes that are regulated by changes in activity and could therefore be relevant to activity-dependent neuronal plasticity (196, 222, 283). While the differential screening and subtractive hybridization methods that constituted the state of the art at the time may seem outdated, these screens laid the conceptual groundwork for later screen design using more up-to-date methods. In addition, their combined output provided a significant fraction of the activity-regulated genes whose cellular function has been characterized in the context of synaptic plasticity, therefore affording us a first glimpse of the cellular machinery activated by IEGs in response to neuronal activity.

In screening for activity-regulated genes, as in any differential screen, a critical aspect of screen design is selection of the tissues for comparison. The greater the number of differences between two tissue sources, the more difficult it is to select from the cloned genes those that are relevant to the specific difference of interest. The more complex a tissue source, containing multiple cell types or multiple functional regions, the more differences will likely be detected. For these reasons it is best to compare cell populations that are as similar and homogeneous as possible. This consideration independently led all three groups to the hippocampus, a region associated with learning and memory, known to undergo cellular plasticity, and relatively uniform regarding cell type, compared with cortex or whole brain. The choice of adult rather than developing tissue circumvented the confounding influence of developmental change. They all used seizure to induce strong, synchronized neural activity with protocols previously shown to result in IEG induction in the brain. Many of the activity-regulated genes identified using the seizure protocol were later demonstrated to be induced by less severe stimulation (see Table 1), such as during LTP (110, 166, 196, 222, 263, 283, 285, 286), and by physiological stimuli, for example, visual input (30, 51, 157, 170, 195, 263, 283, 285), suggesting their induction was not specifically seizure related, but was likely to occur in the brain with day to day levels of activity.

The Kuhl and Worley screens focused on IEGs by applying protein synthesis inhibitors prior to stimulation (222, 283). Kuhl and colleagues (222) identified, in addition to *c-fos* and *zif/268*, five independent genes upregulated by metrazole-induced seizures, that they termed brain activity dependent (BAD) 1 through 5. BAD1, BAD2, and BAD3 were novel genes not further examined. BAD4 encoded a secreted protein of unknown function, TIS21,

TABLE 1. *IEG summary*

Protein	IEG	Stimulus Protocol	Plasticity Paradigm	Cellular Function
tPA	Yes	Seizure (222), LTP (222), learning (239)	L-LTP (13)	Extracellular matrix degradation (262), synaptic growth (13)
MHC I	No	Seizure (51), sensory (51)	LTP (121), LTD (121)	Competitive structural refinement (121)
<i>Intracellular signaling</i>				
SNK	Yes	Seizure (142), LTP (142), electroconvulsive therapy (199), epileptiform activity (238)	Synaptic scaling (236, 238)	Kinase, synapse weakening (236, 238)
RGS2	Yes	Seizure (128), LTP (128), drugs (32, 128)	ND	G protein signaling (20, 128)
CPG16	No	Seizure (196)	ND	S/T-kinase (110, 247), cortical architecture?, neuronal migration?
Rheb	Yes	Seizure (286), LTP (110, 286), sensory (195)	ND	Ras-Raf signaling (289)
<i>Extracellular signaling</i>				
BDNF	No	Seizure (5, 74, 110, 131, 199, 292), LTP (212), sensory (41)	LTP (78, 138, 148, 211)	Neurite outgrowth (15), neuronal survival (113), competitive structural refinement (34), axonal and dendritic arborization (129, 176-178)
COX-2	Yes	Seizure (5, 110, 199, 283), LTP (283), neuronal activity (283), stress (283)	ND	Arachidonic acid metabolism (248), transcellular signaling (109)
Narp	Yes	Seizure (263), LTP (263), sensory (263)	ND	Pentraxin (263), neurite outgrowth (263), AMPAR clustering (203), excitatory synaptogenesis (181, 203, 245)
CPG15	Yes	Seizure (196), diabetes (139), ischemia (104), axotomy (258), sensory (107, 157, 195)	ND	Dendritic outgrowth (198), axonal outgrowth (37), synapse maturation (37, 87), neuronal survival (221)
Arcadlin	Yes	Seizure (284), LTP (284)	ND	Cadherin (284, 288), endocytosis of N-cadherin (288), spine turnover (288)
<i>Synaptic machinery</i>				
Arc	Yes	Seizure (166, 170), LTP (110, 166, 170), sensory (170)	Synaptic scaling (244), L-LTP (215), LTD (215), mGluR-LTD (209, 273), memory consolidation (215, 244)	AMPA receptor internalization (44, 215, 228)
Homer-1a	Yes	Seizure (30, 196), sensory (30, 195)	ND	mGluR clustering (30, 280), signaling (9, 77), and surface expression (8, 10, 229); coupling of mGluRs to the PSD (264), IP ₃ R (265); TRPC1 gating (291); AMPAR cycling (169)
CPG2	No	Seizure (196), sensory (195)	ND	Glutamate receptor endocytosis (53)

ND, not determined. Reference numbers are given in parentheses. See text for definitions.

also induced by NGF (29). BAD5 turned out to be the tissue-type plasminogen activator (tPA), an extracellular protease known to activate the protease plasmin by proteolytic cleavage of plasminogen (262). Further analysis showed that tPA mRNA is upregulated in electrophysiological paradigms that induce LTP in hippocampus and that tPA induction relies on the NMDA receptor (222). Consistent with a potential role in synaptic plasticity, tPA is induced in the cerebellum in response to learning (239), and tPA inhibitors suppress the expression of late LTP (13). Follow-up screens by the Kuhl group identified *arg3.1*, coding for a dendritically enriched protein (166) also identified by the Worley (170) and Nedivi groups (110).

The screen performed by Worley and co-workers proved even more fruitful thanks to the use of subtractive hybridization, yielding 15 novel IEGs (153), approximately half of which encoded transcription factors, including a novel zinc finger protein *Egr3*, a *Zif/268* homolog (285). The MECS paradigm also identified a number of specifically upregulated IEGs with predicted cellular functions unrelated to transcriptional regulation, including COX-2, an inducible form of prostaglandin H synthase (283); Rheb, a novel Ras-related small GTP-binding protein (286, 289); Arc, a novel cytoskeleton-associated protein enriched in dendrites (170) (also termed Arg3.1 by Link et al., see above); Narp, a novel member of the pentraxin family of secreted lectins (263); Homer, a PDZ

protein that binds metabotropic glutamate receptors (30) (also termed CPG22 by the Nedivi group, Ref. 195), and Arcadlin, a novel cadherin-like molecule (284). These are further discussed below.

The Nedivi screen also benefited from enrichment by subtractive hybridization and was further enhanced by a highly sensitive differential screening procedure (110, 196). In contrast to the above-mentioned screens, it was performed without protein synthesis inhibitors. As a result, this screen identified a large number of candidate plasticity genes (CPGs) induced in the hippocampal dentate gyrus in response to kainic acid seizure, including both IEGs and late response genes. The authors initially reported 52 genes, listing 17 with known cellular function, among them, the IEG transcription factors *c-Jun*, *Zif/268*, *c-Fos*, and *CREM* (196). Continued screening eventually expanded this number to 362 upregulated CPGs, 66 of them known, and 7 novel CPGs with motifs predicting their putative function (110). Among the candidate genes there was a clear representation of membrane-, vesicle-, and synapse-related proteins.

Even before characterization of the novel "plasticity" genes identified in these screens, perusing the cellular functions of known genes newly recognized in the context of a seizure response, provided some strong clues regarding the cellular response of neurons to synaptic activity. It became clear that the nervous system employs a wide array of genes in response to activity, with the potential of affecting multiple cellular properties, ranging from process outgrowth and rearrangement of the cytoskeleton and the extracellular matrix, to changes in neuronal excitability and modulation of synaptic strength.

The Nedivi and Worley screens also allowed a preliminary estimate of the number of genes that may be activity-regulated in the brain. Considering that CPGs comprised 5% of the screened clones, and a complexity of ~20–30,000 brain RNA species (158), the total number of activity-regulated genes was estimated at 500–1,000 (196). Exhaustive screening of the libraries prepared after MECS stimulation in the presence of protein synthesis inhibitors generated an estimate of 30–40 independent activity-regulated IEGs, of which perhaps 10–15 were transcription factors (153). These estimates likely represent lower bounds, limited by the finite sensitivity of the screens.

A conceptually different screen for activity-regulated genes in this early generation was not based on seizure activity, but rather on genes downregulated by activity blockade during development (51). This screen identified the immune molecule class I major histocompatibility complex (MHC) antigen as an activity-regulated gene and showed for the first time that the mRNAs for CD3 ζ , a component of the MHC I receptor complex, and for β 2-microglobulin, a cosubunit of the MHC I complex, are

present in the central nervous system (CNS) (51). Hence, molecules formerly described as mediating cell-cell recognition in the immune system were shown to be regulated by activity in the central nervous system. In mutant mice lacking functional MHC I signaling, developmental refinement of connections between the retina and central targets is incomplete, hippocampal LTP is enhanced, and LTD is absent (121).

Approximately 10 years after these first screens, with the advent of microarray chip technology, a second wave of studies attempted to provide a more comprehensive view of transcriptional changes associated with enhanced activity. Since the microarray approach does not require the amount of tissue necessary for conventional differential screening or subtractive hybridization, it allows the use of less efficient, but more specific stimulation protocols, resulting in a greater diversity of approaches. Some chip screens were similar to the earlier differential screens, using hippocampal tissue after an *in vivo* seizure-like stimulus (5, 72, 84, 199), or hippocampal slices after an LTP-inducing stimulus (208). Other screens made use of cultured cortical neurons treated with different pharmacological agents (115, 147, 279). Whether the stimulus was seizure, LTP, NMDA application, or other pharmacological manipulations that affect activity, consensus groups were identified for many of these screens. Two common groups were those related to signal transduction and transcriptional activation. Effector gene categories were more varied between the different screens but did contain genes seen in earlier screens representing neuritogenesis, synaptic transmission, and cytoskeletal regulation. Interestingly, a genome-wide screen for seizure-induced transcriptional changes in the *Drosophila* nervous system also yielded a crop of transcripts, suggesting activity-dependent modulation of membrane excitability and synaptic transmission, as well as cytoskeletal architecture and synaptic growth (97).

Most of the above-mentioned screens partially validate their positives at the transcript level, but the significance of each individual hit to neuronal plasticity can only be assessed after a characterization of its cellular function in this context. This holds true for a second type of recent microarray screens that did not target the general category of activity-regulated genes, but focused on gene expression changes during developmental activity-dependent plasticity in the visual system (151, 172, 261). Interpretation of these data is even more complex due to the confounding effects of age-specific gene regulation, multiple developmental events occurring concurrently during the period examined, and the difference in outcome of the deprivation protocols used to manipulate activity. While there is some overlap with genes previously identified in response to more general seizure manipulations, especially in the transcription factor category, for the most part, the visual system development screens have yielded a pool of genes distinct from each

other and from other screens for activity-regulated genes. It is too early to interpret the mechanistic significance of these differences.

Microarray screens have some major limitations: they are only capable of interrogating a predefined set of sequences, and they report relative, rather than absolute, abundance of a given transcript. In addition, because they are based on sequence-specific probe hybridization, they are particularly vulnerable to false positives due to probe cross-reactivity. Lately, newer techniques that are less susceptible to these problems are replacing gene expression microarray screens (for overview, see Ref. 50). Serial analyses of gene expression (SAGE) has been used to identify genes regulated in response to NGF in PC12 cells (7) and to identify candidate early target genes of JNK or AP-1 in *Drosophila* (76). SAGE technique has also been combined with chromatin immunoprecipitation (ChIP) for serial analysis of chromatin occupancy (SACO) in a genome-wide screen for CREB target sites (127). Comparing the SACO results with those obtained from affymetrix chips with and without cAMP stimulation, the authors validated their results and greatly expanded our view of the CREB regulon. Deep sequencing, a new, unbiased approach for identifying active transcripts (254), allows the detailed detection of multiple splice variants from a single gene, a goal notoriously difficult to achieve with conventional hybridization techniques. Although their application with respect to the activity-regulated gene program in the nervous system is yet to come, the advent of these powerful screening technologies holds promise for more comprehensive approaches to identification of activity-regulated genes.

In summary, forward genetic screens for activity-regulated genes have advanced our thinking regarding the cellular changes set in motion by synaptic activity. While signaling from the synapse to the nucleus utilizes traditional second messenger pathways similar to those activated by growth factors, inducing many of the same transcription factor IEGs, the cellular response to these two types of stimuli clearly diverges at the transcriptional level. Transcriptional programs initiated by synaptic activity are dominated by gene products that directly impinge on neuronal structure and function, with particular emphasis on proteins that promote process outgrowth and differentiation, or interface with the synaptic machinery.

V. IMPLEMENTATION OF ACTIVITY-DEPENDENT NEURONAL TRANSCRIPTIONAL PROGRAMS

Forward genetic screens have yielded a large crop of activity-regulated genes with cellular functions described in nonneuronal contexts or in contexts of early neural

developmental. While these gave a feel for what kind of cellular processes may be recruited in response to synaptic activation, few were followed up and studied mechanistically in the context of cellular or behavioral plasticity. Below we focus on activity-regulated genes that have been characterized sufficiently to yield insight into the cellular mechanisms recruited in response to altered levels of neuronal activity.

A. Intracellular Signaling Molecules

As might have been anticipated by analogy to the growth factor response pathway, two major classes of activity-regulated genes encode transcription factors and signaling proteins. In both classes there are “generic” members in common with the response pathways to extracellular stimuli unrelated to synaptic activity. Also identified were novel transcription factors and signaling proteins, predominantly kinases (168, 247), that seem unique to synaptic stimulation. We will not elaborate further on the role of activity-dependent transcription factors in synaptic plasticity, as this topic is comprehensively discussed in a number of excellent recent reviews (2, 47, 80). We discuss several candidate plasticity genes in the signaling molecule category newly investigated in the context of activity-dependent neuronal response pathways.

1. SNK

The serum-induced kinase SNK, also called Plk2, belongs to the family of pololike kinases, first identified in *Drosophila* (168, 255). In mammals, three genes encode pololike kinases that have been described in the context of proliferating cells, and their function in oncogenic transformation is well characterized (for review, see Ref. 69). In postmitotic neurons, SNK was found to be upregulated in response to various stimuli, including pharmacologically induced seizures, LTP-producing high-frequency stimulation (142), electroconvulsive seizure therapy (199), and epileptiform activity (238). Early studies showed that SNK is activity-regulated both at the mRNA and protein levels, with the most robust induction in hippocampus, amygdala, and the cerebral cortex. SNK protein is localized to soma and dendrites (142) and in particular to dendritic spines (6, 206).

While an early study demonstrated a direct *in vitro* interaction of SNK with the Ca²⁺-binding protein Cib (142), an integrin α IIB binding factor (192), the relevance of this interaction to activity-dependent plastic changes in the nervous system remains unclear. Evidence for a role in synaptic plasticity came from a study demonstrating SNK interaction with spine-associated Rap GTPase activating protein (SPAR) (206). SPAR is part of the postsynaptic scaffolding associated with PSD95, as well as the NMDA receptor (207), and was found to regulate den-

driftic spine morphology by reorganizing the F-actin cytoskeleton (207). SNK inhibits SPAR-mediated cytoskeletal reorganization, inducing loss of SPAR, as well as PSD95 and even presynaptic Bassoon clusters, suggesting it may play a role in weakening or possibly eliminating synaptic contacts (206). The molecular mechanism by which SNK phosphorylation of SPAR leads to SPAR loss is through association with the multisubunit E3 ubiquitin ligase complex SCF (β -TRCP), which directs SPAR towards proteasome-mediated degradation (6).

Two recent studies suggest a requirement for SNK in synaptic scaling. Disruption of SNK function using a dominant-negative approach abolished synaptic scaling in cultured hippocampal neurons (236). Eliminating CDK5 function has a similar outcome, since CDK5-mediated phosphorylation of SPAR is a prerequisite for its recognition by SNK (6, 236). Other experiments in organotypic slice cultures showed that SNK is required for the downregulation of membrane excitability in pyramidal neurons. Thus SNK loss of function results in a persistent potentiation of synaptic strength which precludes the expression of LTP (238). Interestingly, SNK seems to regulate itself negatively, likely through autophosphorylation, and is therefore capable of limiting its own actions over time (144). Taken together, these data demonstrate that SNK is an activity-induced molecule that plays a role in homeostatic synaptic plasticity. The potential substrate specificity of the SNK kinase and the fact that not every spine head contains SPAR in detectable quantities (207) warrants consideration of a synaptic tagging-like mechanism for SNK action (237).

2. *RGS2*

The RGS gene family (regulators of G protein signaling) encodes proteins that accelerate the intrinsic GTPase activity of heterotrimeric G protein α -subunits, thus functioning to dampen G protein-mediated signal transduction (20). In contrast to other members of the mammalian RGS family, *RGS2* is specifically induced by neuronal activity, with induction characteristics of an IEG (32, 128). Early studies using single turnover assays performed in solution indicated that *RGS2* functions as a GTPase-activating protein (GAP) selectively for $G_q\alpha$ subunits. However, *RGS2* also stimulates $G_i\alpha$ when G protein is reconstituted in vesicles together with the muscarinic acetylcholine receptor (mAChR) (128). The same study showed that *RGS2* inhibits both M1- and M2-dependent mAChR-mediated activation of mitogen-activated protein (MAP) kinase, consistent with its GAP activity for both $G_q\alpha$ and $G_i\alpha$ subunits, and indicating how *RGS2*'s function may be integrated in activity-induced downstream signaling pathways.

Behavioral analyses of *RGS2* knockout (KO) mice revealed increased anxiety and reduced male (but not

female) aggression, although no other cognitive or motor deficits were observed (205). These findings were partially supported by a genetic trait analysis of outbred mice that identified the *RGS2* locus as a quantitative trait gene influencing anxiety (282). Anatomically, *RGS2* KO mice display reduced apical and basilar spine density in Golgi-Cox stained hippocampal slices and impaired basal electrical activity in hippocampus CA1 neurons seen as a reduction in the input-output relationship. Surprisingly, LTP in CA1 neurons of these mice was normal (205). A recent study looking more closely for circuit deficits that potentially underlie the behavioral anxiety seen in the *RGS2* KO mice examined *RGS2* KO neurons in culture and found that in the absence of *RGS2* paired-pulse ratios were increased, while expression of *RGS2* in either wild-type or *RGS2*^{-/-} background led to reduced paired-pulse ratios (103). Pharmacological analyses demonstrated that this *RGS2* action is specific to $G_{i/o}$ trimeric α -subunits and that G_q pathways are not involved. Consistent with its effects on paired-pulse facilitation, the authors show that *RGS2* increases presynaptic release probability by increasing local intracellular Ca^{2+} concentrations, likely by downregulating the $G_{i/o}$ -mediated inhibition of presynaptic Ca^{2+} channel activity (103). Taken together, these studies establish a role for *RGS2* in short-term synaptic plasticity, that may underlie its contribution to anxiety- and aggression-related behaviors.

B. Extracellular Signaling Molecules

Activity-dependent signaling is not confined within the cell. A significant fraction of the novel activity regulated genes examined in detail encode extracellular molecules that mediate intercellular communication. We are accustomed to thinking of extracellular signaling as mediated by diffusible molecules that act systemically or in controlled gradients across long distances. Some of the extracellular activity-dependent gene products, such as brain-derived neurotrophic factor (BDNF), clearly act in this manner too. However, an unusual aspect of this category as a whole is that their signaling can also be distinctly localized to the synaptic vicinity.

1. *BDNF*

BDNF was first isolated on the basis of its ability to support the survival of, and process outgrowth from, CNS neurons (15, 113). It was thought to act with other neurotrophins to promote neuronal survival and differentiation during nervous system development (reviewed in Ref. 249). However, expression of neurotrophins, including BDNF, in the adult brain, particularly in the hippocampus (11, 75, 112, 171), suggested they may play additional roles in regulating neuronal function (reviewed in Ref. 225). Shortly thereafter it was shown that transcription of

the *BDNF* gene is strongly regulated by experimentally induced seizure (74, 131, 292), as well as by LTP (212) and physiological levels of activity such as exposure to light (41). Not surprisingly, BDNF also surfaced in screens for activity-regulated genes (5, 110, 199).

The potential relevance of BDNF function to synaptic plasticity was bolstered by studies manipulating its levels in vivo. Intracerebral infusion of BDNF or NT4/5, which share TrkB as their common receptor (130), inhibited formation of ocular dominance columns in the developing visual cortex of cats (34). Other known neurotrophins such as NGF and NT3 did not show these effects, hinting that the TrkB receptor specifically is involved in mediating competition-driven segregation of thalamocortical afferents into eye-specific domains. In a later complementary study, Cabelli et al. (35) reduced endogenous BDNF levels by infusing visual cortex at the peak of ocular dominance segregation with soluble fusion proteins that combine the TrkB receptor extracellular domain with IgG (TrkB-IgG receptor bodies) and act as scavengers of TrkB ligands (242). After intraocular injection of ³H-labeled proline and subsequent visualization by autoradiography, it became clear that ocular dominance stripes were absent near the infusion cannula (35).

Involvement of BDNF in the synaptic plasticity paradigm of LTP was first demonstrated in mice with targeted disruption of the *BDNF* gene. Hippocampal slices from *BDNF* knockout mice showed significantly reduced LTP in the CA1 region of the hippocampus (148, 211). Treatment of these slices with recombinant BDNF entirely rescued deficits in LTP as well as in basal synaptic transmission. Thus synaptic impairments in the knockout mice are not secondary to developmental defects but due to a direct requirement for BDNF during synaptic function and plasticity (211). These findings are consistent with studies showing that TrkB-IgG receptor bodies reduce the synaptic response to tetanic stimulation and diminish the magnitude of LTP in adult hippocampal slices, while exogenous BDNF promotes LTP induction in young slices (78). Additional studies showed that application of BDNF to hippocampal slices from young adults elicited a sustained enhancement of synaptic transmission that mimicked, but did not occlude, high-frequency induced LTP (138). These combined studies strongly indicate an acute role for BDNF in hippocampal synaptic plasticity.

Another aspect of neuronal function influenced by BDNF is structural plasticity of axonal and dendritic arbors. Application of BDNF to organotypic slice cultures of developing neocortex resulted in a rapid increase in length and complexity of pyramidal neuron dendrites (178). Conversely, removing endogenous BDNF (and NT4–5) in this system using TrkB-IgG receptor bodies showed the requirement for BDNF in the growth and maintenance of pyramidal cell dendritic arbors (177). Although evidence for its effect on axons is less extensive

than for dendritic growth, application of BDNF has also been shown to enhance arborization of retinal axons in vitro (129). Inhibition of spontaneous neuronal activity, synaptic transmission, or L-type calcium channels all prevents the positive effect of BDNF on dendritic arborization (176), suggesting that BDNF may serve to link levels of synaptic activity with process outgrowth and activity-dependent structural plasticity.

2. *Narp*

Narp, neuronal activity-regulated pentraxin, is a member of the pentraxin family of secreted proteins. Consistent with the conservation of its Ca²⁺ and carbohydrate binding domains, Narp has the biochemical properties of a calcium-dependent lectin (263). Since pentraxins are structurally conserved across species (73), and the plant-derived lectin concanavalin A promotes neurite outgrowth in mammalian neurons (163), Narp was tested in a heterologous expression system where COS-1 cells transfected with a Narp expression construct were cocultured with cortical explants. Neurons, but not glia, from the explant migrated out to the surrounding area and exhibited exuberant process outgrowth (263). These effects of Narp were not apparent when explants were cultured on collagen, but were most obvious on a less supportive substrate, such as poly-L-ornithine. While it is possible that Narp is simply a better substrate for growth than poly-L-ornithine, its potency in promoting outgrowth is in the nanomolar range, unlike substrates such as NCAM or laminin, and more comparable to neurotrophins and growth factors (159). The growth-promoting effects of plant lectins, also effective at nanomolar concentrations, are attributed to binding of specific receptors on the neuronal surface (164). Like other lectins, Narp is multivalent, and its binding efficacy as well as physiological activity may be dependent on this multivalency (263).

Heterologous expression in HEK 293 cells revealed that secreted Narp binds to itself and forms large aggregates that can trap cotransfected GluR1 subunits (203). A closer look at Narp distribution in the mammalian nervous system shows it is predominantly localized to aspiny neurons, where it resides almost exclusively at GluR1-positive excitatory synapses on the shaft of dendrites, whereas it is largely absent from glutamatergic synapses on spiny neurons, such as pyramidal cells. Ultrastructural analyses detected Narp at both pre- and postsynaptic terminals and in the synaptic cleft as well (203). Native Narp in brain is part of a pentraxin complex with neuronal pentraxin 1 (NP1, described in Ref. 235). Narp and NP1 form quarternary complexes covalently linked by disulfide bonds. Whereas Narp is readily induced in neuronal activity paradigms (263), NP1 is constitutively expressed and is nonresponsive to elevated activity (281). Compared with Narp, NP1 has only minor synaptogenic activity

(281). Thus Narp/NP1 ratios affect the synaptogenic efficacy of pentraxin complexes. Since Narp levels are directly coupled to synaptic activity, excitatory synaptogenesis is dynamically regulated by Narp through its contribution to pentraxin complex composition. A third member of the neuronal pentraxin family is the neuronal pentraxin receptor (NPR). NPR is physically associated with Narp and NP1, and like Narp and NP1 it binds AMPA receptors and contributes to synaptogenesis (61, 245). NPR possesses a transmembrane domain that likely tethers a pentraxin complex containing all three neuronal pentraxins to the cell membrane. Interestingly, both NP1 and NPR are axonally derived and are secreted exclusively by excitatory cell axons at their contact points with interneurons (181, 245). Overexpression of Narp in neurons leads to an increase in the number of excitatory but not inhibitory synapses (203). These data taken together support a model in which Narp facilitates excitatory synaptogenesis by acting as an extracellular aggregating factor for AMPA receptors, specifically on dendritic shafts (181, 203, 245).

However, in response to activation of group 1 metabotropic glutamate receptors, the extracellular metalloprotease TACE releases a soluble form of NPR and the other neuronal pentraxins from the transmembrane tether, allowing their trafficking to endosomes together with attached AMPA receptors, thereby contributing to the downregulation of synaptic AMPA receptors (43). It is possible that the role of nonactivity regulated neuronal pentraxins is to mediate complex turnover, allowing real time incorporation of activity-regulated Narp, thus linking synaptogenic efficacy with neuronal activity.

3. *CPG15 (neurtin)*

cpg15 (candidate plasticity gene 15 also known as *Nrn1*) was initially identified in the Nedivi screen as seizure regulated. *cpg15* has since been shown to be induced by physiological stimuli such as light (157, 195) and whisker stimulation (107), as well as in response to chemically induced diabetes (139), ischemia (104), and axotomy (258). Interestingly, while *cpg15* is an IEG regulated by the same signal transduction pathways, transcription factors, and promoter elements that regulate expression of transcription factors such as Fos (87), *cpg15* encodes a small extracellular protein (191, 195, 198). The CPG15 protein contains both a signal sequence and a consensus sequence for a glycosylphosphatidylinositol (GPI) anchor, and in its processed form is attached to the cell surface (191, 198). In vitro functional characterization of CPG15 suggested that it promotes neurite outgrowth in cultured hippocampal neurons (191). In vivo studies, using virally mediated overexpression in the optical tectum of *Xenopus laevis* tadpoles followed by live imaging, revealed that CPG15 promotes

dendritic outgrowth in a non-cell-autonomous manner (198). Using the same system, Cantalops et al. (37) showed that CPG15 expression significantly increased the growth rate of retinal axons and promoted synaptic maturation by recruitment of functional AMPA receptors to synapses. Electrophysiological recording showed that while NMDA receptor-mediated currents remained unchanged in response to increased CPG15 levels, AMPA receptor currents were significantly increased and failure rates at hyperpolarizing potentials were decreased, potentially due to AMPA receptor recruitment to otherwise silent, immature synapses (37).

CPG15 protein is concentrated in axon tracts (197) and is distributed inside the axon as well as on the axon surface (36). Depolarization by KCl and kainate rapidly increases surface expression of CPG15, suggesting that its delivery to the axon membrane can be rapidly regulated by neural activity (36). These data support a model in which activity plays a dual role in CPG15 action, increasing its expression levels, and promoting delivery of axonally trafficked protein to the surface where it acts to locally modulate dendritic and axonal elaboration, and synaptic maturation.

CPG15 has no sequence homology with traditional neurotrophin ligands and is unusual as a growth factor in its action as a membrane-bound ligand. It does share with neurotrophins an ability to act both as a neuronal growth and differentiation factor and as a survival factor. Consistent with its identification as an activity-regulated gene, *cpg15* expression is contemporaneous with critical periods for activity-dependent plasticity (52, 195). However, *cpg15* is also expressed during early brain development before neuronal differentiation and circuit formation, suggesting a different role at this stage (221). Indeed, soluble CPG15 purified from the culture media in a heterologous expression system can rescue both cortical progenitors and differentiated cortical neurons from starvation-induced apoptosis in vitro (221). CPG15 overexpression in vivo during early development expands the cortical progenitor pool by preventing apoptosis, resulting in an expanded cortical plate and heterotypic cell masses in the ventricular zone. Conversely, RNAi-mediated CPG15 knock-down leads to a shrunken cortical plate with fewer neurons as a result of decreased progenitor survival (221). It is still unclear whether the soluble form of CPG15 harvested from culture media in vitro exists in vivo as a biologically active molecule, and whether CPG15's distinct roles as a survival or differentiation factor are mediated by the soluble versus membrane-bound form.

4. *Arcadlin*

Arcadlin (activity-regulated cadherin-like protein), found by the Worley group to be upregulated in rat hippocampus after MECS or tetanic stimulation of the per-

forant path, was the first activity-regulated gene identified from the cadherin superfamily (284). Northern blot analyses showed expression of the *arcadlin* gene to be brain specific, with highest levels in hippocampus, cortex, and the striatum. The deduced amino acid sequence predicts a single pass transmembrane domain, an extracellular domain with six identified cadherin repeats, and an intracellular COOH terminus. The protein is localized at the soma, and at synapses it colocalizes with the marker synaptophysin. The extracellular domain of Arcadlin is most homologous to the human protocadherin 8, and similar to other cadherins, Arcadlin possesses Ca^{2+} -dependent homophilic binding mediated by this domain. The intracellular portion of Arcadlin, however, seems to be unique among the cadherin family.

Anti-Arcadlin antibody application to acute hippocampal slice preparations suppressed synaptic transmission and impaired LTP induction by tetanic stimulation (284). While these initial results indicated a role for Arcadlin in synaptic transmission, the molecular mechanisms were not elucidated until recently, when Yasuda et al. (288) demonstrated that Arcadlin can regulate dendritic spine numbers by inducing endocytosis of N-cadherin. Using a variety of techniques, their study elegantly demonstrated that Arcadlin and N-cadherin colocalize at synapses and that Arcadlin diminishes the adhesive properties of N-cadherin. Furthermore, the authors show that the molecular pathway for activity-induced internalization of N-cadherin is regulated by Arcadlin through the following sequence of events: synaptic levels of Arcadlin are increased by activity, increasing homophilic interaction of extracellular Arcadlin domains. These interactions activate TAO2 β kinase, which is constitutively bound to the intracellular domain of Arcadlin. TAO2 β is a MAPKKK which is upstream of p38 MAPK. p38 MAPK then feeds back to phosphorylate a key residue of TAO2 β that is required for N-cadherin endocytosis. Hippocampal neurons from Arcadlin-deficient mice show a significantly larger number of spines, perhaps due to increased adhesivity as a result of reduced N-cadherin endocytosis (288). These findings identify the synaptic adhesive apparatus as a target of activity-dependent synaptic remodeling and suggest that Arcadlin may be a negative regulator of synapse and spine numbers through its effect on N-cadherin internalization.

5. Cyclooxygenase 2

Cyclooxygenase 2 (COX-2) first identified by the Worley lab as an inducible immediate-early gene (283) also surfaced in other seizure screens (5, 110, 199). Cyclooxygenases catalyze the first step in prostaglandin synthesis (248) and are the pharmacological target of nonsteroidal anti-inflammatory drugs (reviewed in Ref. 173). The COX-2 substrate arachidonic acid is generated in the stri-

atum in response to NMDA receptor activation (68) and in the hippocampus after synaptic stimulation that induces LTP (275). Hence, increased synaptic activity not only upregulates COX2 mRNA, but also increases availability of the enzyme's substrate. Prostaglandin H₂, the product of COX-2 catalysis, serves as the precursor for further prostaglandin synthesis, but also for the production of prostacyclins and thromboxanes (38, 173, 248), all of which are diffusible molecules that can cross the plasma membrane and therefore spread a local signal to influence neighboring cells (109). It is of particular interest that COX-2 immunostaining is seen in dendritic spines (140), suggesting it can produce diffusible signaling molecules that act locally at the synapse.

C. Regulation of the Synaptic Machinery

While both intra- and extracellular activity-regulated signaling molecules impinge on the synapse, the third category of activity-regulated genes encode proteins that are integral parts of the synaptic machinery.

1. Arc

Arc (activity-regulated cytoskeleton-associated protein) was first described in 1995 as a nontranscription factor IEG upregulated in response to seizure activity (170). At the same time, another group characterized an activity-regulated gene with a 3.1-kb transcript (*arg3.1*) that was later found to be identical to *arc* (166). Robust *arc/arg3.1* mRNA upregulation is seen as early as 30 min after induction (166, 170), and recent work has identified a synaptic activity-responsive element (SARE) through study of the *arc/arg3.1* promoter region (143). It is represented by a ~100-bp element located 5 kb upstream of the transcription initiation site and harbors four boxes that differentially respond to CREB, MEF2, SRF, and TCF to regulate SARE activation (143).

Both initial studies identified homologies between Arc and α -spectrin, characterized its rapid induction in response to neuronal activity, and demonstrated the dendritic localization of *arc/arg3.1* mRNA in the hippocampus (166, 170). While dendritic mRNA localization had been seen for other transcripts, e.g., MAP-2 (89), this feature of *arc* expression is unique among the IEGs (251). Taking advantage of the precise laminar targeting of perforant path projections in the middle molecular layer of the hippocampal dentate gyrus, Steward et al. (251) showed that direct activation of this pathway induces *arc* expression, and accumulation of newly synthesized *arc* mRNA and Arc protein, selectively in the synaptically activated dendritic laminae (251). The local accumulation of both transcript and protein gave rise to the idea that *arc/arg3.1* mRNA might be locally translated in dendrites, a speculation that was confirmed in hippocampal slices

(273). Later, it became evident that *arc/arg3.1* mRNA targeting to specific sites of synaptic activation requires NMDA receptor activation (252). Importantly, this study demonstrated that the induction of *arc/arg3.1* mRNA and its targeting to regions of synaptic activity are separable, since MECS induced *arc/arg3.1* mRNA was evenly distributed in the molecular layer, but when MECS was followed by region-specific synaptic input, the induced transcript accumulated specifically in the region of elevated activity.

The *arc/arg3.1* transcript is one of few naturally occurring mRNAs where the STOP codon of the open reading frame is not located in the last exon. Hence, upon splicing of the transcript, an exon-junction-complex (EJC) tag labels the mRNA at a position 3' to the STOP codon, triggering nonsense-mediated decay of the message (92). This suggests that not only transcriptional activation of *arc/arg3.1* but also its translation is tightly regulated, possibly leading to a single pulse of Arc protein translation in activated terminals and enabling the system to react sensitively to sudden changes in local neuronal activity.

The consequences of Arc protein loss of function were first studied by administration of *arc/arg3.1* antisense oligo-deoxynucleotides via a chronically implanted infusion cannula in the hilus of the fascia dentate (99). These and other experiments revealed that Arc is required for maintenance of LTP and memory consolidation, but not short-term memory (for review, see Ref. 268). These findings were reinforced by studies of an *arc/arg3.1* knockout mouse. In these mice, LTP is compromised: while the early phase of LTP is enhanced, the late phase is essentially absent. The mice show disrupted memory consolidation during spatial learning, fear conditioning, conditioned taste aversion (which is hippocampus independent), and object recognition, although short-term memory was intact (215). A more recent study established a role for Arc in Pavlovian fear conditioning in the lateral amygdala (216). Together, these findings indicate that Arc is required for consolidation of enduring synaptic plasticity and memory storage.

The finding that Arc is necessary for expression of L-LTP and establishment of certain forms of memory led to examination of its role in another form of neuronal plasticity, synaptic scaling (244). In wild-type neurons, surface expression of postsynaptic AMPA-type glutamate receptors is upregulated upon chronic blockade of activity and downregulated under conditions of elevated activity to globally and homeostatically compensate for changes in general activity levels, a process termed synaptic scaling (for review, see Ref. 267). In *arc/arg3.1* knockout neurons, synaptic scaling is absent (244). The effect on surface glutamate receptors seems to be specific for GluR1, since surface GluR2 levels are unchanged in *arc/arg3.1* knockout neurons (244).

Screening for Arc interacting proteins with the yeast-2-hybrid system identified two critical components of the endocytic machinery, dynamin 2 and endophilin 3 (44). Both proteins coimmunoprecipitate with Arc from transfected HEK cells and from forebrain preparations. In HeLa cells, they colocalize on functional early endosomes. Fine-mapping of the binding sites showed that Arc binds to dynamin 2 and endophilin 3 via distinct domains. Amino acids 89–100 in Arc are necessary and sufficient for binding to endophilin 3 (44), and these same amino acids are also required for mediating the decrease in AMPA receptor currents (228), as is the motif for dynamin binding (44), further arguing for an endocytosis-based mechanism of Arc function. Analysis in neurons showed that Arc associates with endophilin and dynamin on early endosomes as well. *arc/arg3.1* knockout neurons show increased AMPA receptor surface expression and reduced endocytosis (44) in line with findings in the *arc/arg3.1* knockout mouse (215).

Since AMPA receptors have been demonstrated to undergo constitutive internalization that is regulated by synaptic activity (70, 162), glutamate receptor internalization has been suggested as a fundamental molecular mechanism to reduce synaptic strength during synaptic plasticity, leading to LTD (174). Two studies presented evidence for a direct role for Arc in AMPA receptor internalization (44, 228). Virally mediated overexpression of green fluorescent protein (GFP)-tagged Arc in CA1 pyramidal neurons in hippocampal slices reduced the amplitude of AMPA receptor-mediated synaptic currents. This was a result of GluR2/3 containing AMPA receptor removal from the synaptic membrane (228). siRNA directed against *arc/arg3.1* could block the Arc-mediated reduction in AMPA currents, as did the deletion of the Arc region known to interact with endophilin and required for clathrin-mediated endocytosis. The interaction between the tail domain of GluR2 and the general endocytic adaptor AP2 can be suppressed using a specific peptide which competes for receptor tail binding (156). When delivered intracellularly through the recording pipette, this peptide abolishes the Arc-induced reduction of AMPA currents. Finally, Arc expression occludes NMDA-dependent LTD. These data suggest that Arc activates synaptic mechanisms similar to LTD, namely, the removal of functional AMPA-type glutamate receptors from the synapse (156, 228).

More recent studies shed light on Arc's role in AMPA-receptor endocytosis during mGluR-mediated LTD. Two major mechanisms of long-term synaptic depression are known: one mediated through the ionotropic NMDA receptors, the other by metabotropic G protein-coupled mGluRs. Both scenarios lead to a loss of AMPA-type glutamate receptors in the postsynaptic membrane, but employ different molecular mechanisms (204). In an elegant study, Waung et al. (273) reported that brief

activation of mGluRs led to local dendritic Arc synthesis and to a persistent increase of GluR1 endocytosis rate. Furthermore, overexpression of GFP-Arc occludes further mGluR-induced decrease of surface GluR1, arguing that Arc and mGluR-LTD employ the same molecular mechanism. Finally, the authors show that endogenous Arc is required for mGluR-mediated AMPA receptor endocytosis and the expression of mGluR-LTD (273).

Another study examining the same pathway showed that *arc/arg3.1* translation is induced within minutes of mGluR activation and that this induction is essential for mGluR-LTD (209). Arc translation in response to mGluR activity requires the translation elongation factor eEF2 kinase, a CaM kinase that binds mGluR and dissociates upon its activation, whereupon it phosphorylates eEF2. Phosphorylation of eEF2 by eEF2K leads to a decrease in global translation, but an increase in specific Arc translation. Thus Park et al. (209) link local Arc translation to a key regulatory component of the translational machinery. Indeed, *eEF2K* knockouts show impaired mGluR-LTD, showing that eEF2K and Arc act in the same molecular pathway. Interestingly, synaptic scaling is normal in *eEF2K* knockouts, suggesting that Arc's function in this plasticity paradigm (244) involves a molecular mechanism distinct from eEF2K function. The fragile X mental retardation protein (FMRP) has also been shown to bind to *arc/arg3.1* mRNA (124, 293) and is hypothesized to inhibit its translation (17). In concordance with this model, the authors observe a dependence of mGluR-LTD upon Arc in *fmr1* knockout slices, arguing that the otherwise distinct signaling pathways for eEF2K and FMRP converge on Arc (209).

The product of an activity-regulated gene, Arc, has been shown as essential for the formation of long-term memories and the expression of both L-LTP and mGluR-LTD. It regulates AMPA receptor surface expression both in an acute way, and in homeostatic forms of plasticity such as synaptic scaling. While its molecular mode of action is being unraveled, we are also beginning to understand how *arc/arg3.1* mRNA localization and translation are themselves tightly regulated.

2. Homer

Homer-1a was first described as an IEG regulated by synaptic activity (30) that binds to group 1 metabotropic glutamate receptors (mGluRs). It was subsequently shown to be a member of a family of Homer-related proteins derived from three distinct genes (Homer 1–3) that each generate several splice variants (280). Like Homer-1a, all Homer proteins bind group 1 mGluRs, yet *Homer-1a* is the only IEG among them. Homer-1a is also distinct from other Homer proteins in that it does not contain a COOH-terminal coiled-coil domain that allows for leucine zipper-mediated dimerization and is hence

unable to oligomerize (27, 280). Homer proteins are synaptically localized (264, 280) and contain a proline-rich NH₂-terminal EVH1 domain that is critical for their specific binding to G protein-linked metabotropic glutamate receptors mGluR1 α and mGluR5 (19, 30, 264) that are coupled to phospholipase C and thereby regulate phosphoinositide hydrolysis (194, 214). The same EVH domain apparently binds other Homer ligands. Binding partners of particular interest for synaptic function are the inositol trisphosphate receptor (IP₃R) (265), the TRPC1 cation channel (291), and the PSD scaffolding molecule Shank (264).

Consistent with their multimeric nature, Homer complexes can be coimmunoprecipitated from brain with both mGluR1 α and the IP₃R, suggesting that Homer couples these receptors in a higher order signaling complex (265). Similarly, Homer facilitates physical association between the TRPC1 channel and the IP₃R (291). Shank binds Homer through a single EVH binding site that is distinct from its GKAP binding domain, thus bridging between these two proteins (264). Since GKAP had been described as part of the NMDA receptor anchoring and signaling complex in the postsynaptic density (PSD) (193), the Homer-Shank interaction potentially links NMDA receptors to both mGluRs and IP₃Rs. Using heterologous expression in COS-7 cells, Tu et al. (265) showed that Shank mediates attachment of Homer to PSD95/GKAP clusters and that mGluR5 clustering relies on the presence of both Homer and Shank. Importantly, Homer self-association and Shank binding are mediated by different portions of Homer, potentially allowing for assembly of a multivalent adapter complex (265).

The IP₃R is located in the ER specialization of the spine apparatus in close proximity to the PSD (270) and functions as an intracellular store calcium channel that can be activated by IP₃ (which can be produced through mGluR signaling and phospholipase C activation; reviewed in Ref. 230). In turn, release of intracellular Ca²⁺ stores by ER IP₃Rs regulates entry of extracellular Ca²⁺ through plasma membrane TRPC channels (145, 291). Homer proteins thus physically couple intracellular IP₃R signaling to the surface mGluRs that activate them, and to surface TRPC channels that they activate themselves. In addition, Homer proteins in this large signaling complex also link mGluR signaling to NMDA receptor signaling cascades via their association with Shank and GKAP, potentially linking IP₃ receptor-mediated Ca²⁺ emission to general synaptic transmission. Elegantly, the Homer-linked signaling complex can be dynamically regulated by Homer-1a which can bind either mGluRs or IP₃R, but does not contain a cross-linking domain that would allow their association within the same complex. Thus Homer 1a effectively functions as an activity-regulated dominant negative that can terminate mGluR/IP₃R interactions (265, 280) as well as IP₃R/TRPC1 interactions (291) and per-

haps disassociate them from the NMDA signaling complex at the PSD.

While the above functions may imply that Homer is a "mere" scaffolding molecule, there is also evidence that Homer can act as an intracellular agonist of G protein-coupled mGluR1 α and mGluR5 (77). Apparently these receptors can shift from an inactive to an active conformation even in the absence of agonist (217). Ango et al. (9) showed that constitutive (agonist-independent) activity of these receptors is controlled by Homer proteins and that disruption of the Homer/mGluR interaction leads to constitutively active mGluR1 and mGluR5 receptors (9). Using a pharmacological approach, they further demonstrated that Homer-1a directly competes with Homer-3 for mGluR1 α and triggers constitutive activity of the receptor. Thus, in this case, too, Homer-1a functions as an endogenous, activity-regulated dominant negative that dynamically competes with constitutive Homer isoforms to regulate mGluR function.

Additionally, Homer proteins may play a role in trafficking and surface expression of group I mGluRs, although this role has not been entirely clarified. In cerebellar neurons where endogenous Homer-1b and -1c are absent, mGluR5 receptors display restricted somatic localization. Cotransfection of Homer-1b and -1c results in dendritic targeting of Homer-mGluR5 complexes (8). Furthermore, Homer-1b expression prevents trafficking of mGluRs from the ER to the cell surface in heterologous systems (229) and suppresses their surface expression in dendrites (10). In contrast, mGluR surface expression is positively regulated by expression of Homer-1a (10, 229).

A recent paper also shows a role for Homer in mediating a physical link between the PSD and the endocytic zone adjacent to it, perhaps facilitating capture and maintenance of a proximal pool of cycling AMPA receptors (169). Although understanding of all Homer functions is still preliminary, clearly, the identification of Homer-1a as an activity-regulated gene product has opened a window into the inner workings of the basic synaptic machinery and its regulation by synaptic activity.

3. CPG2

CPG2, candidate plasticity gene 2, was identified as the product of an activity-regulated transcript (195, 196). Similar to other activity-regulated transcripts, it seems to be the only activity-regulated product of a multitranscript gene. However, as a splice variant of the massive *syne-1* gene, it gives rise to a protein that is unrelated structurally or functionally to its siblings transcribed from the same gene (53). CPG2 is expressed exclusively in brain, predominantly in hippocampus, neocortex, striatum, and cerebellum. In mature cultured hippocampal neurons, CPG2 immunoreactivity is punctate and mainly localizes adja-

cent to PSD95 clusters, a marker for excitatory postsynaptic sites (53).

Immunoelectron microscopic studies have shown that CPG2 resides in the postsynaptic compartment, adjacent to the PSD and in close vicinity of clathrin-coated pits and clathrin-coated vesicles. Clathrin-mediated endocytosis is believed to underlie glutamate receptor internalization (18, 40, 175, 271). The finding that CPG2 colocalizes with clathrin light chain in the postsynaptic endocytic zone lateral to the PSD prompted Cottrell et al. (53) to investigate the role of CPG2 in glutamate receptor internalization. RNAi-mediated knockdown of CPG2 showed that both constitutive and activity-induced AMPA-type glutamate receptor internalization were impaired in the absence of CPG2. Under the same conditions, a fourfold increase of clathrin-coated vesicles was observed, some of which carried the NR1 subunit of the NMDA receptor (53). Spine head size was also affected by CPG2 levels. Under conditions where CPG2 was overexpressed, spine head areas were increased by 7%, upon CPG2 knockdown they were decreased by 18%.

Given that CPG2 is located adjacent to F-actin in spine heads and features a spectrin repeat and coiled-coil domains, it has been speculated that CPG2 may act as a nexus connecting components of the endocytic machinery to the actin cytoskeleton (53). However, it is equally possible that CPG2 fulfills multiple functions, and that variations in spine head size seen upon altered CPG2 levels are not secondary to altered membrane internalization resulting from defective endocytosis, but are direct effects of CPG2 on the spine cytoskeleton.

Neurons may express CPG2 after elevated synaptic activity to prompt removal of AMPA- and NMDA-type glutamate receptors from the synapse as a homeostatic mechanism to counteract the effect of elevated synaptic activity. It is not clear whether CPG2 acts globally as a mechanism of synaptic scaling on all synapses of a given neuron, or whether synaptic tagging directs it to specific synapses that undergo subsequent glutamate receptor internalization.

D. Activity-Regulated Genes Coordinating Inhibitory Synapse Formation

In addition to orchestrating functional and structural changes at the excitatory synapse, recent work shows that the activity-dependent transcription program also coordinates the formation of inhibitory synapses, and likely underlies the maturation of inhibitory circuits.

In an elegant study, Lin et al. (165) used microarrays to identify activity-regulated genes that respond to membrane depolarization. Out of more than 300 potential hits, one candidate gene was responsive only to

Ca²⁺ influx, showed neuron-specific expression, and is expressed coincidentally with inhibitory synapse development. The study further shows that this protein, the bHLH-PAS transcription factor Npas4, regulates the development of inhibitory synapses by orchestrating the expression of a large set of genes that controls formation and/or maintenance of GABAergic synapses, among them *BDNF*. Notably, Npas4 binds to sites in the *BDNF* promoter region (165) responsible for its activity-dependent transcription (1, 167, 257). Furthermore, *BDNF* knockdown partially attenuates Npas4-induced elevation of GABAergic synapses, suggesting that Npas4 regulation of *BDNF* transcription is at least partially responsible for its effect on inhibitory synapse numbers.

Another study from the same lab analyzed the targets of the transcriptional activator MEF2 on a genome-wide scale (81). In neurons, MEF2 responds to neurotrophins and synaptic Ca²⁺ influx (79) and has been shown to restrict excitatory synapse numbers in certain brain regions (14, 79, 220, 240). Using a microarray approach, the new study by Flavell et al. (81) identifies 182 activity-dependent MEF2 target genes that are implicated in various aspects of synapse function, among them several genes that encode proteins with a structural or functional role at the inhibitory synapse. Together with the findings related to Npas4, which also seems to be a direct target of MEF2 transcription (81), these data demonstrate that the neuronal activity-regulated gene program orchestrates inhibitory synapse formation and function, and likely underlies earlier observations of activity-dependent maturation of inhibitory circuits in the brain (106, 119).

VI. SUMMARY

It has been commonly accepted that during development environmental cues act through specific receptors to influence a cell's transcriptional profile and hence its fate. The idea that the environment similarly affects transcription as a means to alter the response properties of mature differentiated cells is newly emerging. In the brain in particular, perhaps because of the paucity of cell proliferation after birth, the view of a hard-wired structure has long dominated the field.

Studies of protooncogene IEGs such as *c-fos* introduced the general concept that gene transcription in the nucleus is part of the cellular response program to alterations in signaling from outside the cell. Integral to this concept was the biphasic nature of transcriptional activation, with IEG induction closely followed by a second wave of delayed-early gene expression. Stimulation of IEG expression by NGF in nonproliferating PC12 cells suggested that this biphasic response could differ according to the differentiation and physiological

state of the stimulated cell and was not necessarily restricted to proliferative cells. Furthermore, IEG expression in culture could be induced by agents other than growth factors and mitogens, including depolarizing stimuli and voltage-gated calcium influx. In vivo, IEGs were shown to be responsive to physiological stimuli, providing strong support for the idea that endogenous IEG expression in the brain is a result of normal synaptic activity and can therefore be influenced by neuronal stimulation or activity blockade in a pathway and stimulus specific manner. At the same time it was becoming clear that synaptic activity can also drive long-term changes in neuronal structure and function. These concepts were unified with the proposal that electrical and chemical stimuli that produce long-term change in neurons, such as during learning and memory, act through mechanisms analogous to those of growth factors, namely, through second messenger pathways that lead to transcriptional activation.

Forward genetic screens aimed at identifying genes induced by neuronal activity rather than by growth factor stimulation proved essential to understanding the activity-dependent genetic programs utilized by neurons in the mammalian brain on an everyday basis. Characterization of activity-regulated genes identified in these screens has advanced our knowledge of the cellular changes set in motion by synaptic activity. While signaling from the synapse to the nucleus utilizes traditional second messenger pathways, similar to those activated by growth factors and inducing many of the same transcription factor IEGs, the cellular response program activated by these two types of stimuli diverges at the transcriptional level. Transcriptional programs initiated by synaptic activity are dominated by gene products that directly impinge on neuronal structure and function, with particular emphasis on proteins that promote process outgrowth and differentiation or interface with synaptic machinery (see Fig. 1).

Given the large number of candidates identified in screens for activity-regulated genes, many of which remain uncharacterized in a neuronal context, we are only beginning to fathom the complexity of processes regulated by synaptic activity in the brain. Yet, despite our incomplete understanding, several themes are beginning to emerge surrounding the role of activity-regulated genes in brain function. Neurons react to changes in levels of synaptic activity with a complex response that takes place on multiple levels and at different time points. Neuronal plasticity, Hebbian or homeostatic, is a consequence of potentially overlapping and interacting actions of a wide number of genes. Furthermore, multiple signals and intracellular pathways can influence transcriptional activation via an assortment of *cis*- and *trans*-acting elements, and nuclear IEGs themselves make up a diverse array of factors that can potentially act in a multitude of combinations to differentially affect distinct second-wave gene sets.

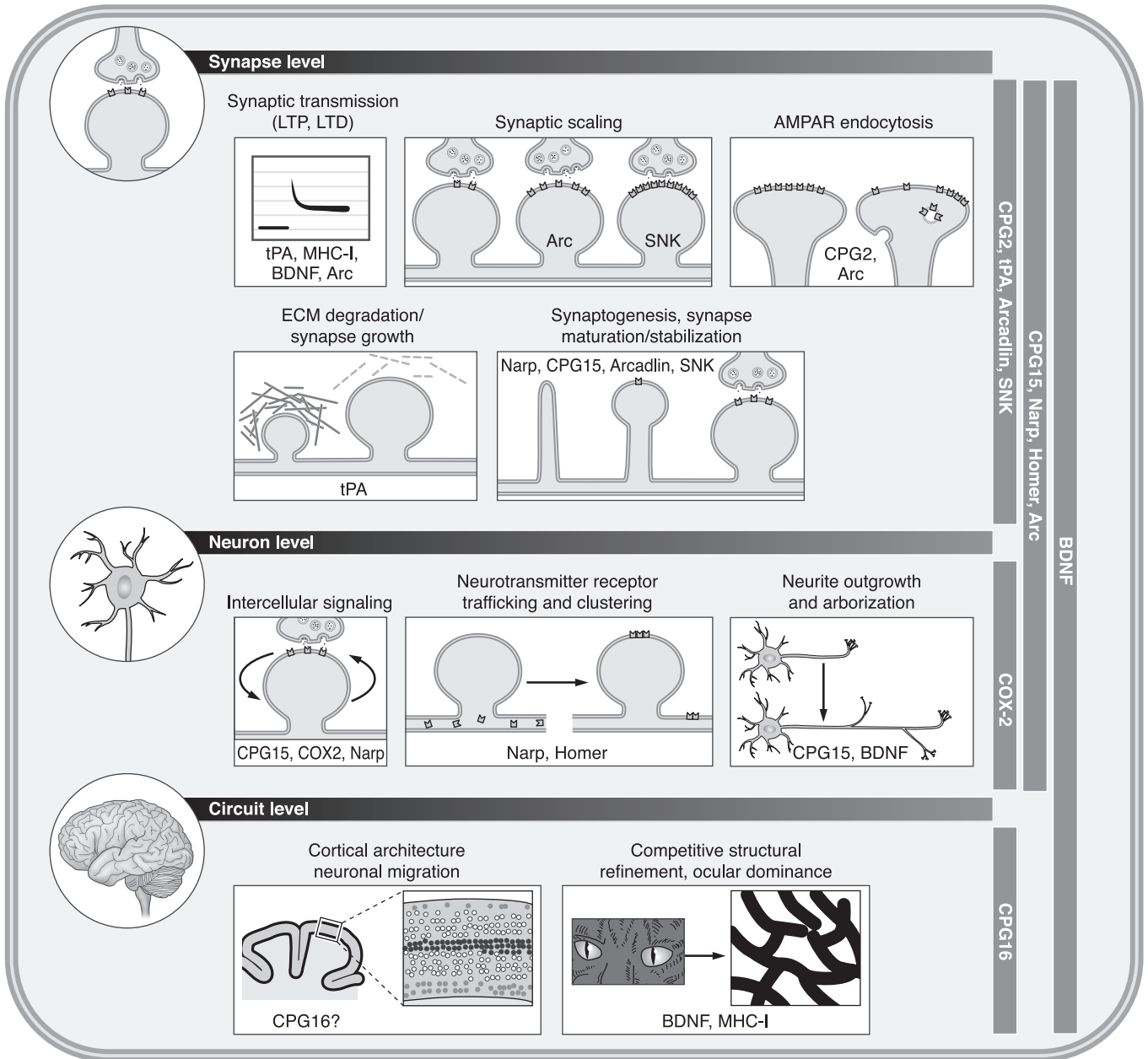


FIG. 1. Activity-regulated genes affect multiple cellular functions at different organizational levels. Boxed sketches depict specific biological processes and show the activity-regulated gene products involved. The boxes are arranged according to the level of organization they represent: *top*, synaptic level; *middle*, cellular level; *bottom*, systems level. Vertical bars on the right indicate which levels each activity-regulated gene product spans with respect to its mechanism of operation.

The large number of genes induced by neuronal activity, and their individual functions suggest that “plasticity genes” are not a unique regulatory gene set whose action is layered on top of constitutive cellular and synaptic genes; rather, they constitute the backbone of the synaptic and cellular machinery and their regulated expression is a means to dynamically tune everyday neuronal function. In addition, regulation of activity-dependent genes is tightly controlled not only at the transcriptional level, but also at the levels of translation and localization,

and perhaps at the levels of mRNA and protein half-life, allowing fine spatial and temporal control of the machinery they oversee.

C-fos, *c-jun*, CREB, CREM, *zif/268*, tPA, Rheb, RGS2, CPG16, COX-2, Narp, BDNF, CPG15, Arcadlin, Homer-1a, CPG2, and Arc likely constitute only the tip of the iceberg in the fascinating array of cellular mechanisms our brains use to bring about remarkable functions such as learning and memory, as well as cognitive abilities like thought and reasoning.

ACKNOWLEDGMENTS

We thank Jeff Hoch, Robert Weinberg, and members of the Nedivi lab for helpful comments on the manuscript.

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GRANTS

We acknowledge funding by the National Eye Institute, the Picower Institute Innovation Fund, the American Federation for Aging Research, and the Deutsche Forschungsgemeinschaft.

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