Retinal Neurotransmitters

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Introduction

Fast synaptic signaling in the vertebrate retina encodes presynaptic voltages as time-varying modulations in extracellular neurotransmitter concentrations and decodes neurotransmitter signals with postsynaptic transmembrane ionotropic or heptahelical receptor arrays. Additional heptahelical receptor pathways conditionally modulate synaptic signaling, often on lower temporal and spatial frequency scales. The six major retinal neurotransmitters glutamate, \(\gamma\)-aminobutyrate (GABA), glycine, acetylcholine, dopamine, and serotonin are formed by group transfer reactions and associated with specific cellular sources and targets in precise connective patterns. The sources and targets of less circumscribed small molecule modulators are mentioned but briefly and peptides are treated elsewhere. Vertical channels deploy fast, high-gain glutamatergic synapses between photoreceptors (PRs) and their bipolar (BC) and horizontal cell (HC) targets, and between BCs and their amacrine (AC) and ganglion cell (GC) targets. Lateral channels composed of HCs and ACs primarily use fast, low-gain sign inverting GABAergic and glycinergic synapses, with specialized circuits employing high-gain sign conserving cholinergic signaling. Four modes of fast synaptic transfer dominate neuronal receptive field circuits: high-gain, sign conserving (\(\rightarrow\)) or inverting (i\(\rightarrow\)), and low-gain, sign conserving (\(\Rightarrow\)) or inverting (i\(\Rightarrow\)). The following sections summarize metabolic networks, transporters, receptors, and circuitries for each neurotransmitter species, concluding with a précis of future directions. Tables 1-6 encapsulate metabolic network diagrams, Enzyme (EC) and Transport Commission (TC) codes, localizations of macromolecules associated with a neurotransmitter phenotype and other data.
References are restricted to reviews or recent exemplars of concepts from which original literature may be traced.

**The Vertical Channel Neurotransmitter: L-Glutamate (Table 1)**

*Glutamate Metabolic Networks.* L-glutamate is a net anion at physiological pH and the central amino acid in a vast network of group transfers in all cells. No enzyme exclusively controls intracellular glutamate levels and no enzyme cluster defines a glutamatergic phenotype. Cellular contents of glutamate and other small molecules reflect group transfer, energetic, redox, osmoregulatory and signaling demands that require supramillimolar concentrations $≈10-100\times$ greater than required to charge tRNAs for protein synthesis. Many transport epithelia and other somatic cells maintain 1-10 mM glutamate, levels as high as PRs, BCs and GCs: neurons are not unique in this regard. Neuronal glutamate is predominantly derived from glutamine produced in Müller cells (MC), but the retinal pigmented epithelium (RPE) maintains 2-10 mM glutamine and could be a source for PRs. Normal MCs display avid glutamate transport but contain little glutamate (50-500 $\mu$M) due to robust conversion by glutamine synthetase (glutamate-ammonia ligase, Table 1). Glial glutamine export appears driven by trans-activated System N (SN1) transporters (Chaudhry et al., 1999) and neuronal glutamine import by the $\text{Na}^+$-dependent Glt1 transporter (Varoqui et al., 2000). Glutaminase on the inner mitochondrial membrane exterior deamidates glutamine to glutamate within intermembrane space, from whence glutamate may be transported to the mitochondrial matrix to drive 2-oxoglutarate synthesis or escape to the cytosol via porin channels.

*Glutamate Transporters.* Vesicular glutamate import is mediated by VGlut1 and VGlut2 transporters, members of the $\text{Na}^+$-dependent plasma membrane $\text{PO}_4^{3-}$...
symporter family (e.g. Takamori et al., 2000). VGluT1 is strongly expressed in both the outer and inner plexiform layers (Stobrawa et al., 2001) and VGluT2 has yet to be mapped, though it is differentially distributed in brain (Varoqui et al., 2002). Anionic glutamate import (K\text{m} \approx 1-3 \text{ mM}) is strongly coupled to $\Delta \Psi$ generated by vATPase proton accumulation. The number of VGluTs per vesicle, their transfer numbers and resultant vesicular glutamate concentrations are all undetermined. Electron microscopic immunocytochemical data in lamprey spinal cord estimate vesicular glutamate levels at $\approx 60 \text{ mM}$, but true levels could be up to ten-fold lower.

Plasma membrane high-affinity glutamate transporters (K\text{m} \approx 5-50 \mu\text{M}) are single 10 transmembrane domain (TMD), one re-entrant loop (RL) polypeptides and members of the widely distributed Na\textsuperscript{+}-dependent acidic and neutral amino acid transporter superfamily (Seal and Amara, 1999). Import is coupled to the sodium gradient ($\Delta p\text{Na}$) and $\Delta \Psi$ (increased transport at negative potentials) and experimental collapse of these gradients activates transporter export sufficient to empty a neuron of glutamate in minutes. Physiologically, export is gradually activated by increasingly positive potentials, although the net import/export balance is not known. Import is weakly electrogenic due to Na\textsuperscript{+}/H\textsuperscript{+} symport but can be strongly electrogenic by activating non-stoichiometric Cl\textsuperscript{−} channel conductance increases ($\uparrow g\text{Cl}$). Four glutamate transporters are expressed in the mammalian retina: predominantly EAAT1 in MCs; EAAT2 in cones and BCs; EAAT3 in HCs, ACs and GCs; EAAT5, in non-mammalian MCs and mammalian rods (Pow et al., 2000).

Plasma membrane glutamate transporters likely shape synaptic kinetics, control glutamate overflow and recycle carbon skeletons. HC response kinetics are shaped by
cone glutamate transporters (Gaal et al., 1998; Vandenbranden et al., 1996) and EAAT2 should act similarly at BC → AC and BC → GC synapses. The relative roles of intrasynaptic neuronal and extrasynaptic glial transport are unresolved. Neuronal transport may couple intrasynaptic glutamate levels to cone voltage via transporter $\Delta \Psi$ dependence (Gaal et al., 1998) when Ca$^{2+}$ flux through voltage-independent cone inner segment cation channels drives vesicle fusion at potentials below Ca$^{2+}$ channel activation (Rieke and Schwartz, 1994).

EAATs on BC dendrites could have novel receptor-like activity by activating $\uparrow g\text{Cl}$ but ON-center BC responses are abolished in mGluR6 knockout mice (Masu et al., 1995) and all BC responses can be blocked with glutamate receptor-specific agents. EAAT-like involvement in PR i→ ON-center BC signaling is more compelling in fishes where hyperpolarizations arise from glutamate-activated $\uparrow g\text{Cl}$ (Grant and Dowling, 1996).

*Ionotrophic Glutamate Receptors (iGluRs).* Vertebrate iGluRs are a diverse group (Dingledine et al., 1999), decoding glutamate signals as cation currents via ionotropic AMPA and KA receptors: apparently non-ordered tetramers of 3 TMD-1 RL subunits with mid-to-high micromolar glutamate thresholds, separable pharmacologies and transient channel opening properties. AMPA receptor GluR1-4 subunits are produced by a single gene family, with a large number of alternatively spliced and RNA edited forms, while KA receptor GluR5-7 and KA1, 2 subunits arise from two gene families, also with post-translational modifications. The *orphan* receptor subunits $\delta 1,2$ are yet another family of likely iGluR subunits. This diversity potentially yields iGluR assemblies with varied unitary conductances, ionic selectivities, kinetics and affinities, as well as
protein kinase A and C (PKA, PKC) modulation, and post-synaptic aggregation control through specialized domains. Receptor diversity is further enhanced by mixed iGluR expression (e.g. Ghosh et al., 2001) and the coexistence of different functional AMPA assemblies in one cell (Zhang et al., 1995). AMPA and KA receptors tend to activate brief conductances in the 1-20 pS range, though some KA receptors can activate larger conductances, while AMPA receptors containing GluR2(R) subunits have substantially smaller unitary conductances.

Functional NMDA receptors are obligate heteromeric 3TMD-1RL tetramers or pentamers of NR1 and NR2 subunits, requiring concurrent glutamate and glycine binding and depolarization relief of uncompetitive Mg$^{++}$ channel block, and gating unitary conductances 2-3× larger than AMPA/KA receptors that activate/deactivate more slowly, with less desensitization that AMPA/KA receptor currents, and are more glutamate-sensitive. Co-activated with AMPA receptors in retina (Diamond and Copenhagen, 1995), functional NMDA-activated pathways appear restricted to AC and GC subsets (Fletcher et al., 2000; Marc, 1999).

_Heptahelical Glutamate Receptors._ Heptahelical glutamate receptors are GPCR Group C members requiring homodimer formation for signaling. The only heptahelical receptor that completely regulates a synaptic pathway is the mGluR Group III mGluR6 localized to ON-center BC dendrites (Vardi et al., 2000a). How mGluR6 activation effects closure of ON-center BC cation channels is unknown and as all mammalian ON-center BCs express a single isoform of mGluR6, any filtering differences must arise within or after in the G-protein signaling path, where $G_{\alpha_o}$ appears to be the coupler
(Dhingra et al., 2000). ON-center BCs in tetrapods, but perhaps not fishes, appear to be under the complete control of mGluR6 signaling.

Other mGluRs have been localized to the retina, but specific pathway functions have not been delineated. Group I-like systems generally mediate intracellular Ca\(^{2+}\) modulations and isoforms of both mGluR1 and 5 have been localized to the inner retina (Koulen et al., 1997b). Group II mGluR2 isoforms may inhibit synaptic release and the most distinctive, but likely not exclusive, localization is the expression of mGluR2 by starburst ACs (Koulen et al., 1996). Group III mGluRs 4,7,8 are widely expressed in the inner retina but their functional roles await analysis.

**Glutamatergic Signal Processing.** AMPA receptors dominate physiological PR \(\rightarrow\) HC signaling, (Blanco and de la Villa, 1999) even though HCs may co-express AMPA GluR2,3,4 (Haverkamp et al., 2001; Morigiwa and Vardi, 1999) and KA GluR6,7 receptors (Brändstatter et al., 1997). Cone \(\rightarrow\) OFF BC signaling in mammals is decoded by KA receptors in two classes of OFF-center BCs and AMPA receptors in a third (DeVries, 2000), generating fast AMPA-driven and slower KA-driven BC classes, perhaps initiating transient and sustained OFF channels, analogous to ON-center BC shaping of transient and sustained ON channels described by Awatramani and Slaughter (2000). OFF-center BCs of non-mammalians may be dominated by AMPA receptors. Signaling via mGluR6 appears as the sole process controlling mammalian ON-center BCs. Mammalian rod BCs have been shown to expresses GluR2 AMPA subunits, but their role is yet unknown

BC \(\rightarrow\) AC and GC signaling in all vertebrates is dominated by AMPA receptors (Marc, 1999), though KA receptors may shape some responses. ACs and GCs display
cell-specific responses to glutamate agonists and responsivity fractionation suggests that physiological attributes such as “sluggish” or “brisk” responses may be determined in part by different AMPA receptor assemblies (Marc and Jones, 2002). NMDA receptor expression varies across ACs and GCs and both immunocytochemical and physiological mappings reveal that specific inner plexiform layer strata driven by cone BCs are enriched with AC and GC dendrites bearing NMDA receptors (Fletcher et al., 2000; Marc, 1999) while rod BC sublayer AC targets lack NMDA receptors (Ghosh et al., 2001; Marc, 1999).

**Glutamate: Future Directions.** Is there a macromolecular signature for a glutamatergic neuron phenotype? VGluT is currently the sole identifier of a glutamatergic phenotype (Eiden, 2000), but other gene products such as EAAT2 may be coordinately regulated with VGluT and transcription factor clusters may ultimately define the glutamatergic phenotype. Detailed spatial mapping and biophysics will be required to resolve the contributions of glial and neuronal EAAT transporters to synaptic kinetics. The mGluR6 transduction pathway and its target channel remains a major puzzle. Do adapter proteins modify its properties and does receptor oligomerization influence signaling? What are the consequences expressing of mixed AMPA, KA, and δ receptors. Do mGluRs shape specific networks or are they diffuse subtle adaptive elements? Why are NMDA receptors relegated to subsets of ACs and GCs and absent from rod BC targets?

**The Dominant Lateral Channel Fast Inhibitory Neurotransmitter: GABA (Table 2)**

**GABA Metabolic Networks.** GABA, an achiral zwitterionic non-protein amino acid, evolved into a signal from its ancestral metabolic role. Vertebrate GABA synthesis
glutamate is driven by two glutamic acid decarboxylases (GAD1/GAD67 and GAD2/GAD65, Table 2), differing in targeting and regulation. GABA catabolism is aerobic, largely driven by mitochondrial matrix GABA transaminase serial oxidation of GABA to succinate semialdehyde and succinate, driving the citrate cycle (Kalloniatis and Tomisich, 1999). Most GAD-containing retinal neurons are likely GABAergic, although some somatic tissues synthesize GAD and GABA, and developing glutamatergic motor neurons in *Drosophila* require GAD expression (Featherstone et al., 2000). Supramillimolar GABA content remains strong marker of GABAergic function and neuronal GABA signals span the inner plexiform layer in all species.

Subsets of HCs and ACs have been shown to express GABA signals, GABA transport and GAD content (reviewed by Marc, 1992). Most ACs are GABA+ and GAD+, though a complete match has yet to be achieved. GABA signal strengths vary widely across AC classes (≈ 1-10+ mM), forming characteristic quantitative GABAergic signatures. All cholinergic, serotonergic and many peptidergic ACs GABAergic signatures and are thus multi-neurotransmitter neurons. Interplexiform cells are AC variants whose processes target the outer retina and several IPC classes may contain GABA (Marc, 1995).

Vertebrate HCs remain neurochemically recalcitrant despite a long history of physiological analysis. GABA content, GABA transport and GAD are expressed by subsets of HCs in many species, but many HCs express no known neurotransmitter signature. A neurotransmitter-free model of feedback signaling via connexin currents has been proposed by Kamermans et al. (2001) though not all problems are thereby solved. Why do any HCs contain GABA if they use connexins for feedback? Some non-
mammalian HCs form conventional synapses onto glycinergic interplexiform cells (IPCs) and other targets in fishes (reviewed in Marc, 1992; Marc, 1995), but may use connexin signaling for cone feedback. Mammalian HCs are equally challenging. Rodent HCs are GABA- but express GABA in development; macular primate HCs are GABA+ but become GABA- in the periphery; rabbit type B HCs are all GABA- while type A HCs are GABA+ in the streak and GABA- in the periphery. All rabbit HCs are GAD67 immunoreactive at their dendritic tips, suggesting fine regulation of signaling (Johnson and Vardi, 1998). All feline, canine and porcine HCs express GABA (Kalloniatis and Tomisich, 1999; Marc, 1992).

Many species display weak GABA signals in small subsets of the BC cohort and some have been shown to contain GAD. Some may acquire GABA through heterocellular coupling while others may be true sign inverting elements. The ganglion cell layer in many species contains many GABA+ cells, mostly displaced starburst ACs, while the remainder are GCs coupled to GABAergic ACs (Marc and Jones, 2002).

**GABA Transporters.** The vesicular transporter VGAT is a 10 TMD polypeptide related to the plant amino acid permease family, is 2-3 fold selective for GABA (Km \( \approx 5 \) mM) over glycine (McIntire et al., 1997) and widely distributed in the inner plexiform layer and in mammalian HCs (Haverkamp et al., 2000).

Most non-mammalians express only neuronal AC and HC GABA transport while mammals express neuronal AC and glial MC transport. Three mammalian, single polypeptide12 TMD, Na\(^+\)-coupled GABA transporters (GAT1, 2,3) are present in the mammalian retina (Johnson et al., 1996), though GAT2 is more closely related to the epithelial osmoregulatory GABA/betaine transporter and is expressed in the RPE, which
contains no detectable GABA. GAT1 is widely expressed in some but not all GABAergic ACs, while MCs preferentially display GAT3. No HC has been found to express these proteins, even those with robust GABA transport.

GATs may regulate synaptic kinetics: they are localized on presynaptic GABAergic neurons, and neuronal transporter blockade can dramatically slow IPSP kinetics (Cherubini and Conti, 2001). GAT3-mediated GABA transport by MCs is avid in mammals, buffering any additional synaptic overflow, though non-mammalians do not require glial support.

Is GABA export a surrogate for or adjunct to vesicular GABAergic synaptic transmission (O'Malley et al., 1992; Schwartz, 1999; Yazulla, 1995)? GABA import is \( \Delta p_{Na} \) and \( \Delta \Psi \)-coupled, with greater import at negative potentials. Collapse of \( \Delta p_{Na} \) or decreasing \( \Delta \Psi \) activates ligand export, but is it physiologically significant? Isolated fish HCs can export physiologically detectable GABA in vitro upon depolarization (Schwartz, 1999), suggesting transport can mediate feedback or feedforward. However, neither GABA transport nor GATs have been detected in mammalian HCs and reconciliation of these observations awaits new work.

**Ionotropic GABA Receptors.** GABA\(_A\) and GABA\(_C\) receptors, members of the ligand-gated channel superfamily, are partially ordered pentameric assemblies of 4 TMD \( \alpha(1-6) \), 2 \( \beta(1-3) \), \( \gamma(1-4) \), \( \delta, \epsilon, \theta \), and \( \pi \) subunits, some with splice variants. Co-assembly of 2\( \alpha \) and 2\( \beta \) subunits, at least, is required to form functional surface-expressed GABA\(_A\) receptors (Connor et al., 1998). GABA\(_A\) receptors gate large, rapidly desensitizing \( \hat{g}_{Cl} \) with GABA thresholds in the 10 \( \mu \)M range. GABA\(_C\) receptors are homomeric assemblies of \( \rho(1-3) \) subunits, are more GABA sensitive and activate
weaker but relatively non-desensitizing $\uparrow g_{Cl}$. Cones, BCs and GCs are known to concurrently express functional $\text{GABA}_A$ and $\text{GABA}_C$ receptors (Feigenspan and Bormann, 1994; Picaud et al., 1998; Zhang et al., 1997b).

*Heptahelical GABA Receptors.* Functional $\text{GABA}_B$ receptors (Group C GPCRs) are R1-R2 isoform heterodimers (Jones et al., 1998; Sullivan et al., 2000). R1a and R1b versions have been mapped to HCs, ACs and GCs (Koulen et al., 1998; Zhang et al., 1998a) and physiological data have shown GCs and BCs to bear functional, pharmacologically defined $\text{GABA}_B$ receptors, in some cases exhibiting complex switching behavior (Zhang et al., 1998b). GPCR coupling of $\text{GABA}_B$ receptors can activate $\uparrow g_K$, hyperpolarizing target neurons, whereas others gate increases in $[\text{Ca}^{2+}]_i$ (Zhang et al., 1997b) or inhibit presynaptic $\text{Ca}^{2+}$ currents (Matthews et al., 1994).

*GABAergic Signal Processing.* GABAergic signaling is primarily inhibitory, effected through ionotropic hyperpolarizations/shunts or GPCR pathways. The HC $\rightarrow$ cone GABAergic pathway has been difficult to validate due to its varied expression. HC $\rightarrow$ cone feedback in amphibians clearly involves ionotropic GABA receptors (Wu, 1994) and most of the components of the pathway are expressed in most vertebrates. Unconventional feedback mechanisms such as connexin-based feedback currents and $\Delta \Psi$-coupled GABA transporter export enrich signaling possibilities but do not reduce uncertainty. Mammalian HCs lack GABA transport while the presence of VGAT provokes the idea that vesicular transmission can occur from HC dendrites. Mammalian HCs display presynaptic vesicle clusters within photoreceptor synaptic terminals but are documented infrequently (e.g. Linberg and Fisher, 1988).
The HC to BC path has argued from anatomical data but the relative HC- and AC-driven surround strengths remain unclear. BCs express dendritic GABA$_A$ and GABA$_C$ receptors (Greferath et al., 1994; Haverkamp et al., 2000; Koulen et al., 1997a) consistent with GABAergic HC $\Rightarrow$ OFF BC signaling through $\uparrow$gCl, forming a proper surround polarity. GABAergic HC $\Rightarrow$ ON BC signaling through $\uparrow$gCl must be sign conserving for proper surround polarity and specialized chloride importers likely shift ON-center BC dendritic E$_{Cl}$ to positive levels to achieve polarity reversal, while preserving a proper negative E$_{Cl}$ at the axon terminal (Vardi et al., 2000b).

The AC $\Rightarrow$ BC signaling is supported by abundant evidence in all vertebrates. How GABA$_A$ and GABA$_C$ receptors differentially shape BC responses is just now emerging. GABA-sensitive, slower GABA$_C$ receptors may initiate feedback control at low contrasts, with less sensitive, faster GABA$_A$ receptors dominating at high contrast. The involvements of GABA$_B$ receptors appear complex and may vary over BC types. GABAergic AC $\Rightarrow$ AC signaling numerically dominates retinal circuitry, is largely GABA$_A$ mediated (Lukasiewicz and Shields, 1998) with concatenated inhibitory chains enriching network assembly (Marc and Liu, 2000). Alls GCs show significant GABAergic inputs, much of it driven by receptors with GABA$_A$–like pharmacologies (Akopian et al., 1998), though some clearly use concurrent GABA$_A$ / GABA$_B$ / GABA$_C$ signaling (Zhang et al., 1997b).

GABA: Future directions. Diverse data support GABAergic signaling by some HCs, but inconsistencies persist and many HCs lack detectable GABA. How do these cells function? Where are the synapses and GABA transporters in mammalian HCs? The mysteries involving AC $\rightarrow$ BC, AC, GC signaling events are more straightforward.
We need to understand the functional consequences GABA receptor types and subtype mixtures. What are the contributions of glial and neuronal GATs to synaptic kinetics and do ACs use GABA export to supplant or augment vesicular release?

3 The Minor Lateral Channel Fast Inhibitory Neurotransmitter: Glycine (Table 3)

_Glycine Metabolic Networks_. Glycine is an achiral zwitterion at physiological pH with limited conformers. Glycine content is elevated in specific ACs with sparse, varicose dendrites: e.g. mammalian AII and DAPI-3 ACs. Lower levels are found in mammalian ON-center BCs that acquire glycine by coupling leakage from AII ACs. ACs with high glycine levels may also contain low GABA levels in many species, perhaps from heterocellular coupling. Retinal glycine synthesis is still unresolved. The glycine hydroxymethyltransferase is reportedly elevated in retina and spinal cord and converts precursor serine to glycine, glycinergic ACs contain no significant precursor serine (Kalloniatis and Tomisich, 1999). Of course, many GABAergic ACs contain little glutamate, so precursors are not proven indices of phenotype. Conversely, somatic cells also use alanine-glyoxylate transferase to produce glycine and precursor L-alanine is elevated in retina and perhaps in glycinergic ACs. Glycine transport has been proposed as a novel mechanism for elevating AC glycine levels based on the depletion of AC glycine by sarcosine (methylglycine), a glycine transport agonist (Pow, 1998), though this effect may have been complicated by trans-activated glycine export.

_Glycine Transporters_. No glycine-selective vesicular transporter has been identified and the nominal inhibitory amino acid vesicle transporter VIAAT transports
GABA and glycine with similar efficacy, while VGAT is only 2-3 fold selective for GABA over glycine. Either might serve a neuron with elevated levels of glycine and no GABA.

Plasma membrane AC glycine transport is mediated by GlyT1, a member of the sodium-coupled solute symporter family and the signature macromolecule of the retinal glycinergetic phenotype. As with EAATs and GATs, collapse of ΔΨ or ΔpNa can evoke complete glycine export in minutes but under normal conditions, GlyT1 likely regulates synaptic kinetics.

**Ionotropic Glycine Receptors.** Glycinergic signaling is exclusively mediated by ionotropic glycine receptors (GlyRs), apparently non-ordered multimers of α1-4 and β subunits. Subunit α1 is abundant in the mammalian inner plexiform layer and is expressed on BCs and GCs (Wässle et al., 1998), though ACs with well-documented anatomical glycinergetic inputs must certainly express GlyR as well. Glycine activates large, rapidly desensitizing ↑gCl, especially in GCs. Since glycine activates gCl in parallel with GABA_A and GABA_C receptors, often on adjacent synapses (Marc and Liu, 2000), glycinergetic signaling may avoid inhibitory occlusion: subadditivity due to GABA spillover at adjacent synapses.

**Glycinergic Signal Processing.** The best-known mammalian glycinerergic circuit is the rod i→ rod BC → Aβ AC i⇒ cone OFF BC pathway, where OFF BC GlyRs render the pathway net sign conserving, as is appropriate for OFF-center channels. This arcane evolutionary capture of cone pathways to serve scotopic signaling is absent in advanced non-mammalian retinas expressing complete, separable rod and cone ON and OFF pathways to GCs. Most GCs receive glycinergetic input and the distributions of GlyRs on identified GCs match relative GABAergic and glycinergetic presynaptic process
densities in specific inner plexiform layer strata. Though their processes are sparse, glycineric ACs are potent and mediate complex behavior via local and wide-field systems, intercalating in sign inverting chains with GABAergic ACs ((Cook et al., 2000; Marc and Liu, 2000; Zhang et al., 1997a).

The glycineric IPC of non-mammalian species is best characterized in teleosts (Marc, 1995). It is presynaptic and postsynaptic in both plexiform layers and part of its role may be to transfer H1 signals from the outer to the inner plexiform layer, bypassing the BCs spatial filter.

**Glycine: Future Directions.** The vesicle transporter of glycineric ACs and the mechanism that elevates glycine content remain unknown. Does glycine transport shape synaptic kinetics in the retina? Does glycineric transmission prevent synaptic occlusion? Why are glycineric ACs of mammals and non-mammalian species structurally similar but involved in such different networks?

**The Lateral Channel Fast Excitatory Neurotransmitter: Acetylcholine (Table 4)**

**Acetylcholine Metabolic Networks.** Acetylcholine is a small quaternary cation synthesized by cholineacetyl transferase (ChAT), a cholinergic phenotype signature macromolecule. Attempts to localize retinal acetylcholine by immunocytochemistry have not succeeded. Conversely, [³H] choline uptake autoradiography and ChAT immunocytochemistry label the same ACs. Extracellular retinal acetylcholinesterase (AChE) terminates synaptic acetylcholine signaling.

**Acetylcholine and Choline Transporters.** Acetylcholine is transported into AC vesicles by VAChTs (Koulen, 1997), a member of the toxin-extruding proton-translocating antiporter family, and under coordinate regulation with ChAT expression.
The ligands for these transporters are cations, so proton antiport and $\Delta p$H dominates vesicle loading.

Synaptic or overflow acetylcholine is cleared by AChE with transfer rate of $\approx 10^4$, effecting rapid hydrolysis of acetylcholine into choline and acetate. Choline is a significant agonist at some acetylcholine receptors and Na$^+$-coupled choline transport via ChT1 may be essential to prevent adventitious receptor desensitization as well for choline recovery.

*Ionotropic Acetylcholine Receptors.* Ionotropic cholinergic transmission is mediated by nicotinic acetylcholine receptors (nAChRs), ordered pentameric assemblies of 4 TMD $\alpha$, $\beta$, and $\gamma$ subunits. Nine neuronal $\alpha$ subunits are known, imparting distinctive properties to channels: e.g., $\alpha_7$ subunits are thought to form homomers with high Ca$^{2+}$ permeability and are expressed widely in the inner plexiform layer. The $\beta_2$ subunit is also abundant in ACs and GCs (Keyser et al., 2000) and $\alpha_3\beta_2$ assemblies are apparently involved in developmental excitatory periodicity linked to retinothalamic patterning (Bansal et al., 2000).

*Heptahelical Acetylcholine Receptors.* The muscarinic acetylcholine receptors (mAChRs) are Group A GPCRs. Subtypes M$_2$, M$_3$ and M$_4$ have been immunolocalized in avian retinas and are expressed by GCs, ACs and BCs (Fischer et al., 1998). Cholinergic starburst ACs express type M$_2$ receptors consistent with autoreceptor regulation of acetylcholine release.

*Cholinergic Signal Processing.* Every vertebrate displays displaced ON and conventional OFF cholinergic starburst AC homologues and bistratification of cholinergic signatures in the inner plexiform layer. Non-mammalians express two or three additional
cholinergic ACs, though the functions of the additional cells are unknown. All cholinergic ACs are also GABAergic, confounding simple circuitry analysis. Only starburst AC circuits have been properly analyzed (Famiglietti, 1991) and they are driven by cone BCs through a high-sensitivity AMPA receptor (Marc, 1999), primarily targeting GCs. Though few starburst AC > AC contacts have been validated, unclassified GABAergic ACs may be excited by starburst ACs through nAChRs (Dmitrieva et al., 2001), perhaps amplifying GC surround inhibition GCs in dim photopic conditions. Any direct signaling between starburst ACs is likely GABAergic and sign inverting, since starburst AC receptive fields are small. Starburst ACs drive directionally selective (DS) GCs in the rabbit retina through nAChRs, but are not needed for directional selectivity per se (He and Masland, 1997; Kittila and Massey, 1997).

_Acetylcholine: Future Directions._ Understanding cholinergic function requires more physiological data in light-driven preparations, discrimination of cholinergic and GABAergic synapses, and pharmacological dissection of nAChRs and mAChRs. Though starburst ACs can be driven to release GABA by export (O'Malley et al., 1992), they lack the neuronal GABA transporter GAT-1 (Dmitrieva et al., 2001): which do they use? Does ChT1 play any role in signal termination?

**The Global Modulator: Dopamine (Table 5)**

_Dopamine Metabolic Networks._ Dopaminergic retinal neurons are ACs or IPCs (reviewed by Witkovsky and Dearry, 1991) that express tyrosine 3-mono-oxygenase (tyrosine hydroxylase, TH) and lack conversion of dopamine to norepinephrine and epinephrine. Their sparse processes facilitate global signaling and dopamine is not spatially buffered, effectively reaching sites tens of microns from the inner plexiform
Tyrosine is an essential amino acid acquired exogenously and accumulated through TAT1 aromatic amino acid transport. In dopaminergic neurons, it is converted to DOPA by TH and DOPA to dopamine by aromatic L-amino acid decarboxylase. Aromatic amines are highly oxidizable and rapid turnover is common in aminergic neurons. Mitochondrial monoamine oxidase converts dopamine to DOPAL, a highly toxic intermediate, then converted to the acetate form for export, apparently by diffusion. As TH is the first stage in tyrosine conversion to neuroactive monoamines, it is present in rare additional neurons that may synthesize norepinephrine or epinephrine, though little is known of their dispositions and roles.

**Dopamine Transporters.** Dopamine is loaded into synaptic vesicles by VMAT2, the neuronal form of the vesicle amine transporter family ((Erickson and Varoqui, 2000). As with other cationic amines, loading is strongly coupled to ∆pH.

The 12 TMD dopamine transporter DAT is similar to most other Na⁺-coupled transporters and is susceptible to trans-activation of dopamine export via transporter agonists such as amphetamines. Its involvement in spatial buffering is somewhat unclear since diffusing dopamine is a potent signal. However, dopaminergic neurons form many synapse-like contacts and highly targeted axonal fields, suggesting that specific connective zones are under higher regulation than others.

**Heptahelical Dopamine Receptors.** All known dopamine receptors are heptahelical Group A (rhodopsin-like) GPCRs coupled through Gs (subtypes D1,D5) or Gi/o (D2,D3,D4), and grouped as pharmacological D1/D2 adenyl cyclase activating/suppressing cohorts, respectively. No retinal cell, including MCs and the RPE, lacks some form of dopamine receptor, and many express both.
Dopaminergic Signal Processing. Dopaminergic neurons apparently signal the onset of photopic epochs through vesicular and dopaminergic effects emerge in seconds to minutes, rather than milliseconds. In teleosts, dopamine activates cone contraction, uncouples HCs and renders GCs more transient through D1 mechanisms, mimicking light adaptation (Vaquero et al., 2001; Witkovsky and Darry, 1991). Coupling control between HCs in teleosts is effected by the axonal fields of dopaminergic IPCs, while that between mammalian AII ACs is effected by axonal fields of dopaminergic ACs in the distal inner plexiform layer (Hampson et al., 1992). In amphibians, dopamine shifts the balance of PR → HC signaling in favor of cones, in part by reducing rod I_h currents via D2 receptors (Akopian and Witkovsky, 1996). The actual patterning and control of dopamine release remains uncertain, but dopaminergic ACs/IPCs appear to be under massive GABA_A receptor-gated suppression and relief from inhibition uncovers spontaneous DA release, perhaps generated by constitutive repetitive spiking ((Feigenspan et al., 1998). Nevertheless, many dopaminergic neurons also receive explicit BC inputs and express iGluRs, so the situation is far from clear. Mammalian, reptilian and avian dopaminergic ACs have also been reported to contain GABA, complicating interpretations further.

Dopamine: Future Directions. A tremendous amount of analysis of dopamine receptor pharmacology has already been done, but how those signaling pathways are themselves regulated and how adaptation state controls and is controlled by dopaminergic neurons demands further exploration.
The mystery neurotransmitter: Serotonin

*Serotonin Metabolic Networks.* Serotonin is present at high levels in specific non-mammalian AC subsets. The mammalian retina contains at most 10-fold less serotonin than dopamine, much of that attributable to platelets and photoreceptor synthesis of melatonin. In the CNS, the phenotype-defining enzyme tryptophan hydroxylase (TrpH) converts TAT1-imported tryptophan to 5-hydroxytryptophan, but thereafter the same enzymes expressed in all other aminergic neurons control serotonin production. There is yet no evidence that any mammalian AC expresses TrpH.

*Serotonin Transporters.* As for dopaminergic neurons in brain, VMAT2 is the obligatory vesicle transporter, though its intraretinal distribution is yet unknown. The content of VMAT2-expressing vesicles thus tracks the substrate amine content of cytosol and VMAT2 expression is not a phenotype signature. All known and suspected serotoninergic ACs are also GABAergic neurons, perhaps expressing both VMAT2 and VGAT.

High-affinity serotonin transport is mediated by SERT, a classic Na⁺-coupled single polypeptide, 12 TMD transporter susceptible to inhibition by numerous agents such as fluoxetine. In many vertebrates, more neurons express serotonin transport (PRs, BC subsets, AC subsets) than are immunoreactive for serotonin, including the GABAergic mammalian A17/S1/S2: No satisfactory explanation has emerged: some of these cells may truly express SERT while others (e.g. some BCs) may have a coupling leak with a *bona fide* serotoninergic AC.

*Ionotropic Serotonin Receptors.* The 5HT3 serotonin receptor is an assembly of unknown stoichiometry of presumed 4 TMD subunits. 5HT3A and B variants have been
described and both must apparently be expressed to mimic conductances and pharmacological profiles of native 5HT3 receptors (Davies et al., 1999). 5HT3A subunits have been localized to mammalian rods, suggesting potential presynaptic control of rod signaling by endogenous serotonin of unknown provenance (Pootanakit and Brunken, 2001).

**Heptahelical Serotonin Receptors.** The GPCR Group A serotonin receptors comprise a complex array of response element receptors, with 15 subtypes in seven families. Among this cohort, 5HT2A receptors are known to be expressed on PRs and rod BCs in the mammalian retina (Pootanakit et al., 1999). Some controversy exists whether phospholipase C coupling is activated through Gq or Gi signaling, but the potential for generating IP3 or diacylglycerol signals near the membrane raises the possibility that serotonin could activate transient receptor potential (TRP) non-selective cation channels, some of which (TRP7) are expressed in retina.

**Serotoninergic Signal Processing.** Little is known of the functions of serotonin in the retina, due to the pharmacological complexities of serotonin receptors, the fact that many GPCRs have constitutive activity and that many agents act as inverse agonists, capable of generating effects in the absence of signaling. Serotoninergic ACs may be a central switch in controlling retinal function, though these ACs must also serve GABAergic roles. Non-mammalian serotoninergic ACs are directly driven by mixed rod-cone OFF-center BCs, form feedback synapses to BCs and target GCs and ACs with feedforward synapses (Marc et al., 1988). However, no data exist to discriminate GABAergic versus serotoninergic signaling at these sites and serotonin may act globally through diffusion. Serotonin reciprocally modulates ON and OFF channels: 5HT3R
activation suppresses scotopic mammalian OFF-center GC responses while 5HT3R antagonism inhibits ON-center GC responses, sparing cone-driven responses (Jin and Brunken, 1998). These complex reciprocal effects could act at PRs or BCs and much remains to be resolved.

**Serotonin. Future directions.** The mammalian serotonin-transporting AC lacks histochemically and immunochemically detectable serotonin. Is an undiscovered monoamine involved or are serotoninergic synapses and vesicles rare, and serotonin synthesis is restricted to small dendritic volumes? Is TrpH or VMAT2 expressed in mammalian ACs? Does serotoninergic signaling involve PR-derived serotonin and are TRP receptors involved?

**Other neuroactive molecules.**

Many non-peptide species can target ionotropic receptors, GPCRs, tyrosine kinase receptors and intracellular response elements and more will likely be found. These additional signals emanate from multifarious sources and modulate signaling within diverse neuronal, glial and epithelial targets, though none is known to be a primary fast signal.

**Nitric oxide** (NO) derived from Ca\(^{2+}\)-coupled NO synthase arginine-citrulline cycling potentially arises from numerous vertical and lateral channel sources and can target an array of cells through guanyl cyclase activation (Eldred, 2001). The effects are potent and include cGMP modulation of cone synaptic Ca\(^{2+}\) channels and HC coupling. The involvement of carbon monoxide signaling in retina is only now being explored.

**Melatonin** signaling, somewhat the inverse of dopamine signaling, initiates with melatonin production in scotophase PRs, diffusion to target melatonin receptors
Retinal Neurotransmitters (Wiechmann and Smith, 2001), and activates dark-adapted and suppresses light-adapted states. Melatonin signaling is coupled to intrinsic circadian oscillator pathways in PRs, but such pathways are complex and simple assignment of photic states to melatonin-versus-dopamine signaling is certainly inaccurate.

Retina expresses a variety of ATP-activated ionotropic P2X receptors on neurons and MCs (Pannicke et al., 2000; Taschenberger et al., 1999) and P2Y GPCRs are expressed in retina (Deng et al., 1998), though the ATP sources and magnitudes of receptor-gated signaling remain to be resolved.

One of the newest signal candidates in retina is anandamide (N-arachidonylethanolamine) or a related molecule that activates cannabinoid CB1 and CB2 GPCRs expressed on retinal neurons (Straiker et al., 1999; Yazulla et al., 1999). Retinal distributions of CNS anandamide transporters and potential interactions with vanilloid receptors are terra incognita.

Fin
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  168.


Table 1: L-glutamate (2S)-2-Aminopentanedioic acid
MW: 147.13
Zwitterion at pH 6-8

Localization:
rod & cones (≈ 1 mM)
BCs, GCs (5-10 mM)
ACs (0.04-1 mM)
MCs (0.04-0.3 mM)

Metabolism: enzyme | EC | map | site | major reactants | → | products
--- | --- | --- | --- | --- | --- | ---
1 glutaminase | 3.5.1.2 | G7 | mim | Q | → | E
2 aspartate transaminase | 2.6.1.1 | F4-5 | mx | C + 2-oxoglutarate | ↔ | E + oxaloacetate
3 multiple transaminases | 2.6.1.x | mx | aa + 2-oxoglutarate | ↔ | E + 2-oxo acid
4 GMP synthase | 6.3.4.1 | G1 | c | Q + xanthonosine | → | E + 5’ GMP
5 CTP synthase | 6.3.4.2 | J7 | c | Q + UTP | → | E + CTP
6 glutamate-F6P transaminase | 2.6.1.16 | D4 | c | Q + F6P | → | E + 2-oxo acid
7 glutamate dehydrogenase | 1.4.1.2 | H6-7G7 | mx | E | → | Q
8 glutamate-ammonia ligase | 6.3.1.2 | G7 H7 | c | E | → | Q
9 glutamate-cysteine ligase | 6.3.2.2 | H7 | c | E | → | γ glutamylcysteine

Transporters
plasma membrane | TC | Localization
--- | --- | ---
a EAAT1 (GLAST) | 2.A.23.2.1 | \( ∀ \) MCs
b EAAT2 (GLT1) | 2.A.23.2.2 | \( ∀ \) PRs, \( ∀ \) cBCs
c EAAT3 (EEAC1) | 2.A.23.2.3 | \( ∀ \) ACs
d EAAT4 | 2.A.23.2.n | \( ∀ \) not in retina
e EAAT5 | 2.A.23.2.n | \( ∀ \) rods
f GluT (Q import) | 2.A.18.6.1 | \( ∀ \) neurons

Receptors: Ionotropic
- KA | 1.A.10.1.1 | KA12, GluR567
- AMPA | 1.A.10.1.2 | GluR1234
- Orphan δ | 1.A.10.1.n | δ1,2
- NMDA | 1.A.10.1.3 | NR1abcd, NR2bc

Receptors: Heptahelical
- GPCR Class C mGluR Group I Gs | mGluR1,5
- GPCR Class C mGluR Group II Gi | mGluR2
- GPCR Class C mGluR Group III Gi | mGluR4,7,8

Abbreviations
- aa: amino acid
- c: cytosol
- cs: carboxylic acid
- F6P: fructose-6-P
- mim: mitochondrial inner membrane
- mx: mitochondrial matrix
- pm: plasma membrane
- px: peroxisome
- vm: vesicular membrane
- vx: vesicle matrix

Neurons expressing glutamatergic phenotypes include all of the cells using ribbon synapses for signaling: rods, cones, and all BCs (grey). All GCs also display a glutamatergic phenotype but with a different metabolic signature.

PRL photoreceptor layer
OPL outer plexiform layer
HCL horizontal cell layer
BCL bipolar cell layer
ACL amacrine cell layer
IPL inner plexiform layer
GCL ganglion cell layer

All Metabolism map loci index the Boehringer Mannheim Biochemical Pathway diagram accessed through www.expasy.ch and its mirrors.
Table 2: γ-aminobutyric acid

| 4-Aminobutanoic acid | MW:103.12 | Zwitterion at pH 7 |

**Localization:**
- HC subsets (0.5-2 mM)
- γ+ AC subsets (1-10 mM)
- γ− AC subsets (0.04-0.5 mM, possible AC → AC coupling leaks)
- GC subsets (0.04-0.5 mM, AC → GC coupling leaks)
- MCs (0.08-0.5 mM → rises in anoxia)

**Transmitters:**
- Metabolism: **enzyme**
- 10 4-aminobutyrate decarboxylase
- 11 4-aminobutyrate transaminase
- 12 aminobutyraldehyde dehydrogenase

**Transporters**
- plasma membrane
- TC
- Localization
- mitochondria
- TC
- Localization

**Receptors:**
- Ionotropic
  - GABA_A
  - GABA_C
- Heptahelical
  - GPCR Class C GABA_B

**Abbreviations:**
- pa: polyamines
- *: links to Table 1

Neurons expressing GABAergic phenotypes are lateral interneurons, including a subset of or all HCs, depending on species, and most ACs, including displaced starburst ACs. GABA signals fill the inner plexiform layer with no interruption due to overlapping AC arborizations. Some GABAergic ACs are also IPCs, though there specific connectivities are unknown.
Table 3: Glycine

Aminoethanoic acid
MW:75.02
Zwitterion at pH 7

Localization:
G+ ACs (≈ 1-5 mM)
G- ACs (0.1-0.3 mM)
ON BCs (≈ 0.5-1 mM, AC → BC coupling leak in mammals)
class 7 GCs (≈ 0.5 mM, AC → GC coupling leak)
all other cells <0.3 mM

Metabolism: enzyme EC map site reactants → products
13 alanine-glyoxylate transaminase 2.6.1.44 B7 px mx A + glyoxalate → G + pyruvate
14 glycine N-methyltransferase 2.1.1.20 ... c SAH + S ↔ G+ SAM
15 sarcosine dehydrogenase 1.5.99.1 B7 mx Sar + acc → G + H2C=O + acc-H
16 glycine hydroxymethyltransferase 2.1.2.1 B6 c mx TH4 + S ↔ 5,10-CH2TH4 + G
17 PRamine-glycine ligase 6.3.4.13 D2 c PRamine + G → PRG

Transporters
plasma membrane TC Localization mitochondria TC Localization
p GlyT1 2.A.22.2.2 ∀ ACs s unidentified G porter mim
q GlyT2 2.A.22.2.2 not in retina synaptic vesicles
r Low affinity n.n.n.n.n ? t VIAAT 2.A.1.14.n Φ brain vm

Receptors: ionotropic
GlyR TC Subtypes Dominant Localization
1.A.9.3.1 α1, α2, α2*, α3, β ∀Φ BCs, ACs, GCs Φ HCs

Abbreviations: see Tables 1,2
PRG: PR-glycinamide SAH: S-adenosyl-L-homocysteine
px: peroxisome matrix SAM: S-adenosyl-L-methionine

Neurons expressing the glycinergic phenotype (high-affinity glycine transporter) in all species are ACs of several narrow-field and diffuse types, with sparse arbors forming broad distal and proximal bands in the inner plexiform layer. The circuitries of some mammalian glycinergic ACs differ markedly from non-mammals yet are similar in shape. Many non-mammals also have a glycinergic IPC that receives input from GABAergic HCs and targets many neuronal types in the inner plexiform layer.
Table 4: Acetylcholine

2-(acetyloxy)-N,N,N-trimethylethanaminium
MW: 147
Cation at physiological pH

Localization:
ACs (enzyme immunocytochemistry)

<table>
<thead>
<tr>
<th>Metabolism: enzyme</th>
<th>EC</th>
<th>map site</th>
<th>reactants</th>
<th>products</th>
</tr>
</thead>
<tbody>
<tr>
<td>18 cholineacetyl transferase</td>
<td>2.3.1.6</td>
<td>B8 c pm</td>
<td>acetyl-CoA + ch</td>
<td>CoA + ACh</td>
</tr>
<tr>
<td>19 acetylcholinesterase</td>
<td>3.1.1.7</td>
<td>B8 c pm ecm</td>
<td>ACh</td>
<td>ch + acetate</td>
</tr>
<tr>
<td>20 choline dehydrogenase</td>
<td>1.1.99.1</td>
<td>B8 mim</td>
<td>ch</td>
<td>BetA</td>
</tr>
<tr>
<td>21 BetA dehydrogenase</td>
<td>1.2.1.8</td>
<td>B8 mx</td>
<td></td>
<td>Bet</td>
</tr>
</tbody>
</table>

Transmitters

<table>
<thead>
<tr>
<th>plasma membrane</th>
<th>TC Localization</th>
<th>TC Localization</th>
<th>TC Localization</th>
</tr>
</thead>
<tbody>
<tr>
<td>u CHT1</td>
<td>2.A.21.8.1</td>
<td>^ ACs, PRs</td>
<td>na</td>
</tr>
<tr>
<td>v MCT1</td>
<td>2.A.1.13.1</td>
<td></td>
<td>synapt ic vesicles</td>
</tr>
<tr>
<td></td>
<td></td>
<td>w ch transporter</td>
<td>x VACHT</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>2.A.1.2.13</td>
</tr>
</tbody>
</table>

Receptors: Ionotropic
nAChR
TC 1.A.9.1.1
Subunits α2-10, β2-4
Dominant Localization ∀Φ ACs, GCs

Receptors: Heptaheical
GPCR Class A mAChR Gs
Subtypes M1, M3, M5
Dominant Localization ∀ BCs, ACs (M3)

GPCR Class A mAChR Gi/o
Subtypes M2,4
Dominant Localization ∀ ACs, GCs

Abbreviations: see Tables 1-3
Bet: betaine
BetA: betaine aldehyde
ch: choline
ecm: extracellular matrix

Neurons expressing the cholinergic phenotype include conventional and displaced starburst ACs in all species and, in many non-mammalian, additional conventional pyriform ON, pyriform OFF, and bistratified cells ACs (grey) that also arborize in the two cholinergic strata.

- **ACs** (enriched with choline, betaine, and acetate)
- **Sublamina a**
- **Sublamina b**
- **IPL**
- **GCL**
- **ACL**
- **MNT**
- **VACHT**

- **Localization**: ACs (enzyme immunocytochemistry)

- **Metabolism**: enzyme
  - **18 cholineacetyl transferase**
    - EC 2.3.1.6
    - map site B8 c pm
    - reactants acetyl-CoA + ch
    - products CoA + ACh
  - **19 acetylcholinesterase**
    - EC 3.1.1.7
    - map site B8 c pm ecm
    - reactants ACh
    - products ch + acetate
  - **20 choline dehydrogenase**
    - EC 1.1.99.1
    - map site B8 mim
    - reactants ch
    - products BetA
  - **21 BetA dehydrogenase**
    - EC 1.2.1.8
    - map site B8 mx
    - reactants BetA
    - products Bet

- **Transporters**
  - **plasma membrane**
    - TC u CHT1 2.A.21.8.1
    - TC v MCT1 2.A.1.13.1
  - **Localization**
    - u CHT1: φ ACs, PRs
    - v MCT1: synapt ic vesicles x VACHT
  - **Transporters**
    - TC 2.A.1.2.13
    - Localization φ brain vm

- **Receptors**
  - **Ionotropic**
    - nAChR
      - TC 1.A.9.1.1
      - Subunits α2-10, β2-4
      - Dominant Localization ∀Φ ACs, GCs
  - **Heptaheical**
    - GPCR Class A mAChR Gs
      - Subtypes M1, M3, M5
      - Dominant Localization ∀ BCs, ACs (M3)
    - GPCR Class A mAChR Gi/o
      - Subtypes M2,4
      - Dominant Localization ∀ ACs, GCs

- **Abbreviations**
  - Bet: betaine
  - BetA: betaine aldehyde
  - ch: choline
  - ecm: extracellular matrix

- **Localization**
  - ACs (enzyme immunocytochemistry)
  - Neurons expressing the cholinergic phenotype include conventional and displaced starburst ACs in all species and, in many non-mammalian, additional conventional pyriform ON, pyriform OFF, and bistratified cells ACs (grey) that also arborize in the two cholinergic strata.
Table 5: Dopamine

3,4 dihydroxyphenethylamine
MW: 153.18
cation at neutral pH

Localization:
ACs, IPCs

Metabolism: enzyme

<table>
<thead>
<tr>
<th>Enzyme</th>
<th>EC</th>
<th>map</th>
<th>site</th>
<th>reactants</th>
<th>products</th>
</tr>
</thead>
<tbody>
<tr>
<td>22 phenylalanine 4-monooxygenase</td>
<td>1.14.16.1</td>
<td>F3</td>
<td>c</td>
<td>F + BH4</td>
<td>Y + BH2</td>
</tr>
<tr>
<td>23 tyrosine 3-monooxygenase</td>
<td>1.14.16.2</td>
<td>G2</td>
<td>c</td>
<td>Y + BH4</td>
<td>DOPA + BH2</td>
</tr>
<tr>
<td>24 aromatic L-aa decarboxylase</td>
<td>4.1.1.28</td>
<td>G2</td>
<td>c</td>
<td>DOPA</td>
<td>DA</td>
</tr>
<tr>
<td>25 amine oxidase (flavin containing)</td>
<td>1.4.3.4</td>
<td>G2</td>
<td>mom</td>
<td>DA</td>
<td>DOPAL</td>
</tr>
<tr>
<td>26 aldehyde dehydrogenase (DHA5)</td>
<td>1.2.1.3</td>
<td>G2</td>
<td>mx</td>
<td>DOPAL</td>
<td>DOPAC</td>
</tr>
</tbody>
</table>

Transporters

<table>
<thead>
<tr>
<th>plasma membrane</th>
<th>TC</th>
<th>Localization</th>
<th>synaptic vesicles</th>
<th>TC</th>
<th>Localization</th>
</tr>
</thead>
<tbody>
<tr>
<td>plasma membrane</td>
<td>TC</td>
<td>Localization</td>
<td>synaptic vesicles</td>
<td>TC</td>
<td>Localization</td>
</tr>
<tr>
<td>y TAT1 Na⁺-coupled</td>
<td>2.A.1.13.2</td>
<td>?</td>
<td>aa VMAT2</td>
<td>2.A.1.22.1</td>
<td>?</td>
</tr>
<tr>
<td>z DAT Na⁺-coupled</td>
<td>2.A.22.1.3</td>
<td>?</td>
<td>aa VMAT2</td>
<td>2.A.1.22.1</td>
<td>?</td>
</tr>
</tbody>
</table>

Receptors: Heptahelial

<table>
<thead>
<tr>
<th>GPCR Class A DAR Gs</th>
<th>Subtypes</th>
<th>Dominant Localization</th>
</tr>
</thead>
<tbody>
<tr>
<td>GPCR Class A DAR Gs</td>
<td>D1,D5</td>
<td>PRs, ACs, HCs, BCs</td>
</tr>
<tr>
<td></td>
<td>D2,3,4</td>
<td>PRs, (D4) ACs (D2)</td>
</tr>
</tbody>
</table>

Abbreviations: see Tables 1-4

BH2: dihydrobiopterin
BH4: tetrahydrobiopterin

Neurons expressing a dopaminergic phenotype are conventional ACs in most species (black) and arborize in the distal inner plexiform layer with dendritic (heavy lines) and dense axonal fields (fine mesh). Many species exhibit very sparse IPC-like processes targeting the outer plexiform layer. Advanced fishes (teleosts) possess a dopaminergic IPC (grey) with a unique, dense axonal field in the outer plexiform layer and multistratified inner plexiform layer arborizations.
Table 6: Serotonin

5-hydroxytryptamine
MW: 176.21
Cation at physiological pH

Localization:
ACs in non-mammalians
PRs (biochemistry)

Metabolism: enzyme

<table>
<thead>
<tr>
<th>Metabolism</th>
<th>EC</th>
<th>map</th>
<th>site</th>
<th>reactants</th>
<th>products</th>
</tr>
</thead>
<tbody>
<tr>
<td>28 tryptophan hydroxylase</td>
<td>1.14.16.4</td>
<td>H2 H3</td>
<td>c</td>
<td>W + BH4</td>
<td>5HTP + BH2</td>
</tr>
<tr>
<td>24 aromatic L-aa decarboxylase</td>
<td>4.1.1.28</td>
<td>H2 I2</td>
<td>c</td>
<td>5HTP</td>
<td>5HT</td>
</tr>
<tr>
<td>25 amine oxidase (flavin containing)</td>
<td>1.4.3.4</td>
<td>I2</td>
<td>mom</td>
<td>5HT</td>
<td>5HIAA</td>
</tr>
<tr>
<td>27 aldehyde dehydrogenase (DHA5)</td>
<td>1.2.1.3</td>
<td>I2J2</td>
<td>mx</td>
<td>5HIAA</td>
<td>5HIAC</td>
</tr>
</tbody>
</table>

Transporters
plasma membrane
y TAT1 Na+ -coupled 2.A.1.13.2
bb SERT Na+ -coupled 2.A.22.1.1

Receptors: Ionotropic
5HT3 1.A.9.2.1

Receptors: Heptahelical
GPCR Class A 5HTR G/o
GPCR Class A 5HTR Gq
GPCR Class A 5HTR Gs

Abbreviations: see Tables 1-5
OPL: 5HIAA: 5-hydroxyindole acetaldehyde
5HIAAC: 5-hydroxyindole acetate

5HT: 5-hydroxytryptamine, serotonin
5HTP: 5-hydroxytryptophan

Only non-mammalians express a complete serotoninergic phenotype in ACs (grey) with heavily varicose dendrites arborizing heavily in distal and sparsely in proximal strata of the inner plexiform layer. Many of these cells possess fine, sparse axons targeting the outer plexiform layer. Mammalian A17 (S1,2) ACs (dotted) lack a complete phenotype, expressing only SERT-like transport. In all species, ACs with serotonin transport are GABAergic.