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Titolo	The Glutamate/GABA-Glutamine Cycle : Amino Acid Neurotransmitter Homeostasis // edited by Arne Schousboe, Ursula Sonnewald
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Nota di contenuto	Introduction to the Glutamate-Glutamine cycle -- Glucose, lactate, - hydroxybutyrate, acetate, GABA, and succinate as substrates for synthesis of glutamate and GABA in the glutamine-glutamate/GABA cycle -- Anaplerosis for glutamate synthesis in the neonate and in adulthood -- Enzyme complexes important for the glutamate-glutamine cycle -- BCAA metabolism and NH <sub>3</sub> homeostasis -- Glutaminases Vesicular Glutamate Uptake -- The glutamine transporters and their role in the Glutamate/GABA-Glutamine Cycle -- Glutamine Metabolism in Gliomas -- Oligodendrocytes: development, physiology and glucose metabolism -- Dysregulation of glutamate cycling mediates methylmercury-induced neurotoxicity -- Astroglia, glutamatergic transmission and psychiatric diseases -- Glutamine Synthetase: Role in Neurological Disorders -- The Glutamate – Glutamine Cycle in Epilepsy -- Index.
Sommario/riassunto	Fundamental biochemical studies of basic brain metabolism focusing on the neuroactive amino acids glutamate and GABA combined with the seminal observation that one of the key enzymes, glutamine synthetase is localized in astroglial cells but not in neurons resulted in the formulation of the term “The Glutamate-Glutamine Cycle.” In this cycle glutamate released from neurons is taken up by surrounding astrocytes, amidated by the action of glutamine synthetase to glutamine which can be transferred back to the neurons. The conversion of glutamate to glutamine is like a stealth technology,

hiding the glutamate molecule which would be highly toxic to neurons due to its excitotoxic action. This series of reactions require the concerted and precise interaction of a number of enzymes and plasma membrane transporters, and this volume provides in-depth descriptions of these processes. Obviously such a series of complicated reactions may well be prone to malfunction and therefore neurological diseases are likely to be associated with such malfunction of the enzymes and transporters involved in the cycle. These aspects are also discussed in several chapters of the book. A number of leading experts in neuroscience including intermediary metabolism, enzymology and transporter physiology have contributed to this book which provides comprehensive discussions of these different aspects of the functional importance of the glutamate-glutamine cycle coupling homeostasis of glutamatergic, excitatory neurotransmission to basic aspects of brain energy metabolism. This book will be of particular importance for students as well as professionals interested in these fundamental processes involved in brain function and dysfunction.

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