

Publication

An intrinsic distinction in neuromuscular junction assembly and maintenance in different skeletal muscles

JournalArticle (Originalarbeit in einer wissenschaftlichen Zeitschrift)

ID 155369

Author(s) Pun, S.; Sigrist, M.; Santos, A. F.; Ruegg, M. A.; Sanes, J. R.; Jessell, T. M.; Arber, S.; Caroni, P.

Author(s) at UniBasel [Rüegg, Markus A.](#) ; [Arber, Silvia](#) ;

Year 2002

Title An intrinsic distinction in neuromuscular junction assembly and maintenance in different skeletal muscles

Journal Neuron

Volume 34

Number 3

Pages / Article-Number 357-370

Keywords Agrin/genetics/metabolism; Animals; Botulinum Toxin Type A/pharmacology; Bungarotoxins/pharmacology; Hindlimb; Immunohistochemistry; Mice; Knockout; Models; Neurological; Muscle Denervation; Muscle; Skeletal/cytology/drug effects/*embryology/*innervation/physiology; Neuromuscular Agents/pharmacology; Neuromuscular Junction/*embryology/growth & development/*physiology; Receptor Aggregation; Receptors; Cholinergic/*metabolism; Schwann Cells/metabolism

We analyzed the formation of neuromuscular junctions (NMJs) in individual muscles of the mouse embryo. Skeletal muscles can be assigned to one of two distinct classes of muscles, termed "Fast Synapsing" (FaSyn) and "Delayed Synapsing" (DeSyn) muscles, which differ significantly with respect to the initial focal clustering of postsynaptic AChRs, the timing of presynaptic maturation, and the maintenance of NMJs in young adult mice. Differences between classes were intrinsic to the muscles and manifested in the absence of innervation or agrin. Paralysis or denervation of young adult muscles resulted in disassembly of AChR clusters on DeSyn muscles, whereas those on FaSyn muscles were preserved. Our results show that postsynaptic differentiation processes intrinsic to FaSyn and DeSyn muscles influence the formation of NMJs during development and their maintenance in the adult.

Publisher Cell Press

ISSN/ISBN 0896-6273

edoc-URL <http://edoc.unibas.ch/dok/A5259090>

Full Text on edoc Restricted;

Digital Object Identifier DOI 10.1016/S0896-6273(02)00670-0

PubMed ID <http://www.ncbi.nlm.nih.gov/pubmed/11988168>

ISI-Number WOS:000175214700008

Document type (ISI) Article