Synaptic Vesicles, Mitochondria, and Actin Alterations in SMN-deficient Mice

Margret Neher

Göttingen
Synaptic Vesicles, Mitochondria, and Actin Alterations in SMN-deficient Mice

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Margret Neher
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Dekan: Prof. Dr. rer. nat. H.K. Kroemer

I. Bericherstatter: Prof. Dr. J.H. Weishaupt

II. Berichterstatter: Prof. S. Rizzoli

III. Berichterstatterin: Prof. M. Schön

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Chapter 1

Introduction

1.1 Motor Neuron Diseases

This heterogeneous group of diseases is characterized by a predominant affection of the motor neuron system. The chronic progression of these diseases can still hardly be influenced by therapeutic intervention. The degeneration can affect the first motor neuron (central paralysis), the second motor neuron (peripheral paralysis), or both systems in combination (Poeck and Hacke 2001). An affection of the first motor neuron is characterized by an increase of the muscle tone and enhanced muscle reflexes, while an affection of the second motor neuron is characterized by atonic paralysis and extinction of muscle reflexes (Trepel 2003). Spastic spinal paralysis is an example for a degeneration of the first motor neuron, while spinal muscular atrophy (SMA) belongs to motor neuron diseases with degeneration of the second motor neuron. The prevalent amyotrophic lateral sclerosis (ALS) shows the nearly unique mixed pattern of a central and peripheral paralysis. One well-known ALS patient is the astrophysicist Stephen Hawking, who develops fascinating theories on the origin and evolution of galaxies, while being captured by his own body.

1.2 Spinal Muscular Atrophy (SMA)

There are different forms of SMA. The following study refers to the so-called proximal spinal muscular atrophy, characterized by degeneration of the second motor neurons and skeletal muscular atrophy. The disease is autosomal recessive and caused by a deletion of, or mutations in, a single gene, the survival motor neuron 1 (SMN1) gene. After cystic fibrosis, proximal SMA is the most common autosomal recessive disorder in humans, with a carrier frequency of approximately 1 in 35 and therefore an incidence of 1 in 6000. It is also the most common genetic cause of infant mortality (Monani 2005).

There are three types of proximal SMA ranging in severity from very severe (type I) to intermediate (type II) to mild (type III) (Pearn 1980). Moreover, there are numerous other less frequent forms of SMA with a different genetic background but similar phenotype (Table 1). The leading symptoms of proximal SMA are a progressive predominant proximal
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<table>
<thead>
<tr>
<th>SMA Type</th>
<th>Mode of Inheritance</th>
<th>Gene Chromosome Location</th>
<th>Phenotype Symptoms</th>
<th>Onset of the Disease</th>
</tr>
</thead>
<tbody>
<tr>
<td>Proximal SMA’s Type I (Werdnig- Hoffmann)</td>
<td>Autosomal recessive</td>
<td>SMN1; 5q11.2-13.3</td>
<td>Proximal muscle weakness, patients never sit unaided; death &lt; 2 years</td>
<td>&lt; 6 months</td>
</tr>
<tr>
<td>Type II (intermediate chronic form)</td>
<td>Autosomal recessive</td>
<td>SMN1; 5q11.2-13.3</td>
<td>Proximal muscle weakness, patients sit unaided but become wheelchair bound, develop Scoliosis of spine</td>
<td>6 - 18 months</td>
</tr>
<tr>
<td>Type III (Kugelberg-Welander)</td>
<td>Autosomal recessive</td>
<td>SMN1; 5q11.2-13.3</td>
<td>Proximal muscle weakness, patients walk unaided, normal lifespan</td>
<td>&gt; 18 months</td>
</tr>
<tr>
<td>Distal SMA</td>
<td>Autosomal recessive</td>
<td>11q13</td>
<td>Distal muscle weakness, diaphragmatic involvement</td>
<td>2 months – 20 years</td>
</tr>
<tr>
<td>SMARD (Spinal Muscular Atrophy Respiratory Distress)</td>
<td>Autosomal recessive</td>
<td>IGHMBP2; 11q13.2</td>
<td>Distal lower limb weakness, diaphragmatic weakness, sensory, autonomic neurons also affected</td>
<td>1-6 months</td>
</tr>
<tr>
<td>X-linked infantile SMA</td>
<td>X-linked</td>
<td>Xp11.3-q11.2</td>
<td>Arthrogryposis, respiratory insufficiency, scoliosis, chest deformities, loss of anterior horn cells at birth</td>
<td>at birth</td>
</tr>
<tr>
<td>Distal SMA IV</td>
<td>Autosomal dominant</td>
<td>7p15</td>
<td>Distal muscles affected, bilateral weakness in hands, atrophy of thenar eminencmotor neurone and peritoneal muscle</td>
<td>12-36 years</td>
</tr>
</tbody>
</table>

Table 1: Modes of inheritance, gene location of the underlying mutation, phenotype and onset of the different forms of SMA. Note that the three types of proximal SMA have the same genetic defect but they differ in their phenotypes with lifespan ranging from under 2 years to normal. The other forms of SMA have miscellaneous modes of inheritance and genetic background (from Monani, 2005, p 886; reproduced with kind permission from Elsevier).

Paresis, which appears always symmetric, and displays muscular fasciculation. The muscle reflexes are abolished. Vegetative functions, like bleb and gut evacuations, are not generally affected. Furthermore, no disturbance of the intelligence of patients is apparent. The various forms of SMA are very different in their time course and prognosis (Poeck and Hacke, 2001).

**SMA type I (Werdnig-Hoffmann)** The acute type I form is characterized by severe, generalized muscle weakness and hypotonia at birth or within the next 3 months. The paralysis begins in the pelvic muscles and progresses from the extremities and from cranial towards the facial muscles. The muscular weakness of the inspiratory muscles leads to atelectasis, which promotes the development of infection and inflammatory processes.
1.3 The Genetic Principles of Proximal SMA

Death from respiratory failure or pneumonia occurs within the first two years (Poeck and Hacke 2001).

**SMA type II (intermediate - chronic form)**  The onset of the disease is later than in type I, at an age between 6 and 18 months. The type II patients learn to sit and mostly to crawl while walking is impossible (Kunze, 1999). Although most patients become older than two years pulmonary complications are leading to an early death (Kunze, 1999).

**SMA type III (Kugelberg-Welander)**  The mildest form of proximal SMA can be symptomless until the early adulthood. The onset is mostly within ages 2 and 17 years. A proximal weakness of the legs starts after an initially normal motor development. This might first be noticeable through difficulties in climbing stairs and standing up. In the later disease course most patients become dependent on a wheel chair. The life span is not considerably shortened.

### 1.3 The Genetic Principles of Proximal SMA

All three forms of proximal SMA are autosomal recessive. Regardless of the disease severity, 95% of all patients carry a deletion in a gene containing nine exons that localizes to chromosome 5q11.2-13.3. This gene is termed the telomeric survival of motor neuron gene (SMN1). The remaining 5% carry small mutations in the gene. In humans the SMN1 gene lies within a 500 kb inverted-duplicated region. Through the duplication of this region the centromeric half contains an almost identical copy gene, SMN2 (Lefebvre et al. 1995). SMA patients lack the telomeric SMN1 gene, but they have at least one copy of SMN2. This, however, produces only around 10% functional SMN protein and therefore can not compensate for the lack of the former. The difference in expression between the two genes is due to a translational silent single nucleotide (C to T) transition inside SMN2 exon 7. This critical transition leads to an alternative splicing of the SMN2 gene, resulting in a shortened gene product (SMN ∆Ex7) lacking exon 7. This makes a genetic diagnosis of SMA relatively straightforward, based on a simple PCR reaction (Dreesen et al. 1998). Due to the shorter length of the SMN2 transcript compared to the SMN1 transcript both genes can be distinguish from each other through gel electrophoresis. The SMN ∆7 isoform is unstable and rapidly degraded. While the SMN1- transcript is almost 100% translated to functional full-length protein (FL-SMN), only 10% of SMN2 transcript translated to FL-SMN protein. Patients with a deletion in the SMN1 gene, therefore, express insufficient levels of functional FL-SMN protein. Figure 1 shows the molecular basis of SMA: transcription and translation of SMN1 and SMN2-gene as well as the resulting amounts of protein (Monani 2005).

The number of SMN2 genes varies in the population from 1 to 6 copies. The greater the number of SMN2 genes, the more FL-SMN protein is produced and the milder is the disease phenotype (McAndrew et al. 1997) (Feldkötter et al. 2002). This explains the range in severity among SMA type-I, type-II, and type-III patients. A direct correlation between
The amounts of full length SMN-expression and severity of the phenotype demonstrates that SMA is due to a classical gene dose effect.

In essence, SMA results from low levels of the FL-SMN protein.

### 1.4 The FL-SMN Protein

### 1.4.1 Assembly and Distribution of the FL-SMN Protein

The FL-SMN protein contains 294 amino acids and has a molecular weight of 38 kilo Dalton (kDa). It is ubiquitously expressed and does not exhibit homology to any previously identified protein. The necessity of the SMN for cellular survival and function has been proven in the following organisms and cell types: human, mouse, chicken, DT40 cells,
1.4 The FL-SMN Protein

Figure 2: Schematic diagram of SMN showing coding exons and the relative localization of selected domains with known functions. In SMN2 exon 7 is spliced out resulting in truncated protein lacking the C-terminal most 16 amino acids (Briese et al. 2005), p 947; reproduced with kind permission from John Wiley and Sons).

drosophila melanogaster, caenorhabditis elegans, and schizosaccharomyces pombe, indicating that SMN is essential for viability of all eukaryotic organisms (Miguel-Aliaga et al. 1999; Hannus et al. 2000; Paushkin et al. 2000; Wang and Dreyfuss 2001). One central question around this protein is why defects in a ubiquitously expressed protein have such a selective effect on motor neurons and corresponding muscles.

The C-terminal region of the SMN-protein contains a region for self-oligomerisation encoded by exon 6, which contains numerous tyrosin-glycin pairs (Lorson et al. 1999). This is flanked by a central evolutionary highly conservative tudor domain, encoded by exon 3, which contains 53 amino acids and is important for the interaction between SMN and the Sm-proteins (Bühler et al. 1999). Further interaction domains and a schematic illustration of the SMN-protein are shown in Figure 2.

Within one cell SMN can be found in the cytoplasm as well as in the nucleus (Liu and Dreyfuss 1996). The distribution of SMN in the cytoplasm appears diffuse in all cells studied. Only in muscle cells is SMN existent in large cytoplasmic aggregates (Liu and Dreyfuss 1996) (Burlet et al. 1998). In the nucleus SMN appears as 0.1-1 µm large, dot-like structures. These structures were often found to colocalize with coiled bodies and were therefore termed gems (gemini of coiled bodies). The name coiled bodies has been recently replaced by the name Cajal bodies (Gall et al. 1999). Cajal bodies are known to be rich in factors involved in the transcription and processing of many types of nuclear RNAs. The close interaction of gems with Cajal bodies indicates that gems are also involved in RNA-processing (Liu and Dreyfuss 1996). Beside the localization in gems SMN is also directly localized in Cajal bodies in cultured cells and primary neurons (Carvalho et al. 1999). In HeLa cells the assembly of the coiled bodies is disrupted due to the depletion of SMN (Girard et al. 2006). Furthermore, the group of Carvalho could show that SMN in Cajal bodies colocalize with small nuclear ribonucleoproteins (snRNPs). In this study the majority (>85%) of cells studied contained SMN only in Cajal bodies, while no gems could be identified (Carvalho et al. 1999). Depending on the tissue SMN is expressed in different amounts (Coovert et al. 1997) (Burlet et al. 1998). Particularly high levels of the protein are found in the spinal motor neurons, apparently the most profoundly
affected cells in SMA patients. Furthermore, different studies in human tissue (muscle, heart, kidney and brain) show a clear reduction of SMN-expression after birth (Burlet et al. 1998). Interestingly, it is found that the intracellular distribution changes during the normal CNS-development. There is a shift from SMN being localized more in the nucleus during early stages of development towards an accumulation in the cytoplasm, especially in axons of motor neurons (Giavazzi et al. 2006).

The amount of SMN is reduced in all reviewed tissues of SMA patients (Burlet et al. 1998), with patients with SMA type I having smaller amounts of SMN protein than type II or type III patients (Coovert et al. 1997; Lefebvre et al. 1997; Burlet et al. 1998). The spinal cord, with an almost complete absence of FL-SMN protein in type I patients, is especially affected, while the amount of SMN in lymphocytes and muscle tissue is only moderately reduced. Although in SMA mostly lower motor neurons are affected, a reduction in all reviewed tissues is detectable (Jablonska et al. 2007). However SMA patients develop in the first line motor neuron degeneration and muscular atrophy (Jablonska et al. 2007). In order to shed light on the selective effect of SMN on the nervous system several groups studied in particular the distribution of SMN in motor neurons. Béchade et al. (1999) could for the first time demonstrate that SMN is located in the proximal part of dendrites, where it is associated with microtubules. One year later, another research group also came to the conclusion that the SMN is associated with elements of the cytoskeleton in spinal dendrites and axons, in particular during the early postnatal development (Pagliardini et al. 2000). The discovery of SMN in axonal compartments led to the assumption that SMN is actively transported. Recent studies showed granular accumulations of SMN in axons of motor neurons. They demonstrated that the transport of SMN over longer distances is microtubules-dependent, while for shorter distances microfilaments are required (Zhang et al. 2003). SMN is also found in the growth cones of motor neurons (Rossoll et al. 2003).

1.4.2 The SMN-Complex

SMN is the central component of a large oligomeric complex, the so-called SMN-complex (Meister et al. 2001; Gubitz et al. 2004). Already in 1996 Liu and Dreyfuss showed that SMN binds amongst others to itself, to the RGG-rich region of hnRNP U, and to fibrillarin. One year later this group identified the SMN interacting Protein1 (SIP-1), which was later termed gemin2. Until now seven more gemins have been identified. Figure 1 shows a scheme of all seven interacting proteins called gemin 2-7.

This complex has been postulated to function as an “assemblyosome” important for snRNP biogenesis and pre-mRNA splicing, which will be described in the next section. The binding partners of the SMN-complex can also appear independently from each other. A recent study showed that gemin2 forms a complex with SMN including no other gemin proteins. This leads to the assumption that the pair SMN-gemin2 is the smallest available subunit. Overall a multitude of other proteins has been identified, which directly interact with SMN and/or gemins, although they are not integral components of the SMN-complex. To these proteins belong among others, the Sm- and Lsm-proteins of the small ribonucleoprotein family (snRNPs) (Liu et al. 1997; Bühler et al. 1999; Friesen and Dreyfuss 2000).
1.4 The FL-SMN Protein

Figure 3: Schematic illustration of the SMN-complex with known components (taken from Carissimi et al. 2006). SMN is interacting directly with gemin2, gemin3, gemin4 and gemin5 and gemin7 (Liu et al. 1997); Charroux et al. 2000; Gubitz et al. 2004), while gemin4 binds over gemin3 (Charroux et al. 2000). For the interaction of gemin6 with SMN-complex is Gemin7 required. Gemin8 binds to Gemin6 and Gemin7 on the multiprotein complex (Carissimi et al. 2006). p. 8134; reproduced with kind permission from the author).

the helicase A (Pellizzoni et al. 2001) and hnRNP Q (Mourelatos et al. 2001).

The exact implication of these interactions remains to be determined. However the knowledge of these interactions may shed light on additional neuron specific functions of SMN, which may explain why reduced SMN levels have relatively little impact on cells other than motor neurons.

1.4.3 The Role of SMN in SnRNP Biogenesis and pre-mRNA Splicing

The most extensively documented function of the SMN has been the so called housekeeping role in providing small ribonucleoproteins, which are the main components of the spliceosome. One essential step of posttranscriptional processing of the pre-mRNA to mature mRNA is the splicing of pre-mRNA in the nucleus before being exported to the cytoplasm. During splicing the noncoding introns are removed from the pre-mRNA and the protein-encoding exons are linked to each other. This process is catalyzed by the spliceosome. The main components of the spliceosomes are U snRNPs (uridine-rich small nuclear ribonucleoproteins) (Will and Lührmann 2001). Each snRNP contains seven Sm-proteins plus one or two small nuclear RNAs (snRNA U1, U2, U4/U6 and U5). The role of the SMN-complex in the assembly of spliceosomal U snRNPs U1, U2, U4/U6 and U5 in the cytoplasm has been studied in detail (Fischer et al. 1997); Selenko et al. 2001); Jablonka et al. 2002); Winkler et al. 2005); Kolb et al. 2007). The SMN-complex builds the U snRNPs by loading the Sm-proteins in a well defined order onto the U snRNA (Pellizzoni, 2007). In this
Introduction

Figure 4: The mode of action of the SMN and PRMT complexes. The Sm-proteins are newly synthesized by the cell and bind to the PRMT5-complex (1). This adds methyl groups to the Sm-proteins, which cause their handing-over to the SMN-complex (2). The SMN-complex subsequently transfers the Sm-proteins to a nucleic acid, called U snRNA (3), thereby forming U snPNP, a subunit of the spliceosome. SMN- and PRMT5-complexes may then enter into a new cycle (Fischer et al. 1997, p. 750; reproduced with kind permission from the author).

function the SMN-complex is strictly controlled by a further protein assembly called the PRMT5-complex. Figure 4 shows the sequential course of this clustering reaction towards U snRNPs.

This process takes place in the cytoplasm. The snRNPs are then imported to the nucleus, where they are often found together with SMN-complexes in cajal bodies, where they undergo further maturation processes before they take up their function in pre-mRNA splicing.

The role of SMN in snRNP biogenesis and pre-mRNA splicing has been most extensively documented (Meister et al. 2000; Hannus et al. 2000). However, it still remains unknown whether SMA and the motor neuron phenotype are a direct consequence of a disruption in snRNP biogenesis and pre-mRNA splicing.

A defect in a function as universal as snRNP biogenesis and pre-mRNA splicing is likely to affect all tissues equally. There are basically three schools of thought to explain why defects in a ubiquitously expressed protein may have such a selective effect on motor neurons.
(i) Considering that low levels of SMN lead to a general defect of mRNA splicing, the tissue-specific nature of SMA could be explained by the fact that motor neurons simply have lower tolerance for depleted SMN levels, being large, high-energy requiring cells. However, a lack of an adverse effect on other large, high-energy requiring cells, such as cortical motor neurons and sensory neurons makes this a questionable hypothesis.

(ii) Another possibility would be that the unique sensitivity of affected motor neurons in SMA is a result of aberrant splicing of one or more RNAs, which are critical for the appropriate functioning of these cells. However, such RNAs have yet to be identified.

(iii) The SMN may have additional motor neuron specific functions besides its role in snRNP biogenesis and pre-mRNA splicing. Studies confirming this school of thought will be discussed in the next section.

### 1.4.4 Motor Neuron Specific Function of SMN

The main feature distinguishing neurons from other cell types is that they possess dendrites and axons that convey messages from one neuron to another or target muscle cells by chemical and electrical processes. This feature implies that cargo has to be transported over long distances in order to secure sustainability and growth of neurons. Several lines of evidence support additional neuron-specific functions of SMN. The hypothesis that SMN has a neuron-specific function firstly emerged from the observations demonstrating accumulation of SMN protein in the axon and growth cones of neurons specifically in cells in vitro (Fan and Simard 2002) and anterior horn cells in vivo (Tizzano et al 1998). These observations have been fostered by experiments identifying the presence of SMN-containing granules within the neurites of chick cortical neurons and rat spinal motor neurons (Figure 5), that associate with microtubules and exhibit bidirectional movement between the cell body and the growth cone (Zhang et al 2003).

Moreover, it is well established that SMN can bind RNA (Lorson and Androphy 1998) (Bertrandt et al 1999) and ribonucleoprotein particles (Jones et al 2001); (Liu and Dreyfuss 1996); (Mourelatos et al 2001).

These findings lead to the assumption that SMN may be involved in the transport of specific mRNAs to the growth cone in response to local cues during development. Rosoll and colleagues reported that SMN and its binding partner hn-RNP R modulate axon growth cones. They finally discovered that SMN and its hnRNP are involved in the processing and localization of β-actin mRNA to the growth cones of developing motor neurons (Rossoll et al 2003). Reduced growth cone size, lower β-actin levels and shorter neurites in primary motor neurons from SMA mice add further weight to this line of thought. The role of actin in the SMA disease will be discussed in a following section.

Recent studies could show that SMN directly binds to the alpha-subunit of the vesicle coat protein ‘coat protein I’ (COPI). This subunit, alpha-COP, is Golgi associated and co-immunoprecipitates with SMN, small nuclear ribonucleoprotein-associated assembly factors and β-actin mRNA. This observation leads to the proposal that neurons utilize
Figure 5: A chick cortical neuron stained with an antibody against SMN showing abundant nucleocytoplasmic staining as well as the presence of granules (arrows) within neurites and the growth cone. The granules, RNP particles, are found associated with microtubules, indicating movement along the axon. This supports the idea of a motor neuron specific function of the SMN protein ([Monani 2005], p. 889; reproduced with kind permission from Elsevier).

the Golgi associated COPI vesicles to deliver cargos necessary for motor neuron integrity and function ([Peter et al 2011]). Thus, another study adds further weight to the hypothesis that mutated SMN interferes with the transport of cargo in the axon, which might provide an explanation of the neuron-specific nature of SMA.

1.5 Mouse Models of SMA

Animal models can shed considerable light on the pathogenesis and mechanisms of their respective human disease. Since the SMN mutations were first identified as responsible of SMA, numerous organisms have been manipulated in order to generate animal models of SMA. In the following, the approach leading to the mouse model we used for our experiments will be explained. Several problems had to be circumvented before obtaining a suitable mouse model.

The identification of the murine homolog of the SMN gene (Smn) ([DiDonato et al 1997]) made it possible to knock it out ([Schrank et al 1997]). However, humans are the only species with an SMN2 gene and all SMA patients carry at least one intact SMN2 gene. A complete knock out in mice, in other words a complete lack of the SMN protein, is embryonically lethal and therefore did not result in a useful mouse model, but provided evidence that Smn is an essential gene ([Schrank et al 1997]). This is not surprising, given the fact that in SMA patients the SMN1 gene is also deleted or mutated but the SMN2 gene is still producing small amounts of FL-SMN protein.

One approach taken to overcome the embryonic lethality in Smn -/- mice was to introduce the SMN2 gene into this genetic background ([Monani et al 2000]). This strategy showed that the phenotype depends on the SMN2 transgene copy number. The introduction of eight copies of SMN2 completely rescued the SMA phenotype, two copies resulted in SMA mice that die at 5 days of age ([Hsieh-Li et al 2000]; Monani et al. 2000) and one
1.6 Neuro Muscular Junction (NMJ) Pathology in SMA

1.6.1 Synaptic Vesicles

Synaptic vesicles allow the communication between neurons and other cells through releasing neurotransmitter, at the NMJ acetylcholine, into the synaptic gap, as chemical response to the electrical impulse arriving at the nerve terminal. Vesicles are constantly recreated and grouped into three pools: the readily releasable pool, the recycling pool, and the reserve pool. The vesicles of the readily releasable pool are docked to the cell membrane, making them the first group of vesicles to be released on stimulation. The recycling pool is defined as those vesicles that maintain release upon (physiological) moderate stimulation. The majority of vesicles (typically ~ 80-90%) belong to the reserve pool from, which release neurotransmitter only during intense stimulation (Rizzoli and Betz 2005). Figure 6 illustrates this three-pool model.

Several functional studies at the NMJ in SMA mice lead to the question whether there is a disruption regarding synaptic vesicles in SMA mice. Kong and colleagues measured a two-fold reduction of the evoked endplate currents (EPCs). The EPCs depend on the number of fused vesicles and the muscle response to the transmitter released from a single vesicle. The spontaneous miniature EPC (MEPC), defined as muscle response to one single vesicle, was found to be normal, indicating that a decreased number of fused vesicle leads to the reduction of the EPCs (Kong et al 2009).

In agreement with this study it was found that the kinetics of the postsynaptic potentials are slowed and evoked neurotransmitter release is decreased by approximately 55% in Transversus Abdominus (TVA) muscle (Ruiz et al 2010). Indeed electron microscopy images of the NMJ in TVA showed that individual synaptic vesicles within presynaptic terminals had normal diameters and morphology, but overall density of synaptic vesicles within the presynaptic terminal area was reduced by 56%, while the number of docked
Figure 6: “The classic three-pool model. The reserve pool makes up \(\sim 80-90\%\) of the total pool, and the recycling pool is significantly smaller (\(\sim 10-15\%\)). The readily releasable pool (RRP) consists of a few vesicles (\(\sim 1\%\)) that seem to be docked and primed for release”. Figure and caption from (Rizzoli and Betz 2005), p. 58; reproduced with kind permission from Nature Publishing Group.

vesicles, defined as those within 20 nm of the membrane, was only reduced by 32\% (Kong et al 2009). In contrast, in diaphragm no significant difference in vesicle number was found (Kariya et al 2009).

To expand the perspective of the amount and distribution of synaptic vesicles (SVs) in SMA mice, we explored both total area of the terminal covered by SVs and the SVs spatial organization through fluorescent microscopy. First we undertook this study in TVA, a postural muscle of the anterior wall innervated by lower intercostal nerves. This muscle is known to be severely affected in the disease.

Additionally, we studied SVs in Levator Auris Longus muscle (LAL) from the dorsal surface of the head, innervated by the facial nerve. These studies allowed us to quantify pre- and postsynaptic pathology in muscle groups from two anatomical regions of the mouse and to compare pathologies in predominately slow-twitch (TVA) versus fast-twitch (LAL). Two further characteristics render the LAL muscle an interesting object of study: First, the LAL has two constituent muscle bands, among which the caudal band is known to be more affected in the disease than the rostral band (Murray et al 2008). After a comprehensive comparison between the rostral and the caudal bands, it was shown that the two bands differ in their “Delayed Synapsing” (DeSyn) and “Fast Synapsing” (FaSyn) characterictis (Murray et al 2008). It is therefore likely that selective vulnerability is due to the FaSyn characterictics, dominant in the caudal band. These two classes, DeSyn and FaSyn, differ in the focal AChRs clustering, the alignment of presynaptic nerve with AChRs and the alignment of schwann cells. In FaSyn muscles, this focal organization process is achieved in less than 1 day, whereas in DeSyn muscles up to 5 days are required (Murray et al 2008). Experimentally these two classes can be distinguished by their reaction to paralysis with botulinum toxin A. While NMJs conforming to a DeSyn phenotype have been shown
to undergo dramatic collateral sprouting, FaSyn NMJ remains largely unaffected (Murray et al. 2008).

Second, postsynaptic terminals have been described to mature almost at the same pace as in control mice (Murray et al. 2008). This allows us to assess the question to what extent pre-and postsynaptic phenotypes are interrelated by proving whether the pathology regarding SVs found in TVA is also found in LAL.

The question how deficiency of SMN protein can contribute to the observed abnormalities of synaptic vesicles will be addressed in the discussion part.

### 1.6.2 Mitochondria

Mitochondria are organelles of elaborate structure, that in addition to supplying cellular energy and producing reactive oxygen species (ROS) have significant roles in calcium homeostasis and apoptosis. At the presynaptic side they regulate intraterminal Ca$^{2+}$ levels, providing an especially important buffering action during trains of action potentials (David and Barrett 2000). Exploring the neurotransmission of severe SMA revealed that asynchronous neurotransmitter release is increased by $\sim 300\%$ compared to the wild type (Ruiz et al. 2010).

Presynaptic terminals release neurotransmitter at rest (spontaneous release), upon a single action potential (synchronous evoked release), and during intraterminal Ca$^{2+}$ accumulation resulting from prolonged electrical stimulation (asynchronous release). Therefore, the mentioned increase of asynchronous release indicates an anomalous augmentation of intraterminal bulk Ca$^{2+}$ in SMA mice. A possible explanation for this augmentation is a decreased Ca$^{2+}$ reuptake by mitochondria during trains of action potentials (Ruiz et al. 2010).

Mitochondria dysfunction is known to be involved in many neurodegenerative diseases, however little is known about the role of mitochondria in SMA pathogenesis. Mitochondria are briefly mentioned in two electron microscopy (EM) studies on P14 SMA mice. The study in diaphragmatic muscle showed that presynaptic mitochondria are decreased in size but not in numbers (Kariya et al. 2008). In contrast, another research group demonstrated in Tibiales Anterior muscle a reduced density of mitochondria, while mitochondrial morphology was normal (Kong et al. 2009).

To study the density, reliability, and spatial distribution of mitochondria in Transversus Abdominal muscle (TVA) at P14 under the confocal microscope, we used Mito Tracker, a cell-permanent fluorescent dye that is sequestered by functioning mitochondria. Moreover we explored the spatial relationship between Mito Tracker spots and SV clusters; reflecting the two main organelles of the presynaptic compartment, for which actin serves as a track for short-range transport (Langford 2002).

### 1.6.3 F-Actin

Actin is one of the most abundant proteins in eukaryotic cells. Actin is the monomeric subunit of microfilaments known as globular actin (G-actin). G-actin subunits assemble
into long filamentous polymers called F-actin.

In neurons, actin-filament plays an important role in multiple functions such as axon initiation, growth, guidance, and branching during neuron development. In order to secure sustainability in neurons, cargo has to be transported over long distances. The “dual transport model” was proposed in which microtubule-based motors ensure long-range axonal transport, whereas short-range movement of organelles such as vesicle and mitochondria at nerve terminals and subcortical plasma membrane regions depends primarily on actin-based myosin (Langford 2002). In addition to these functions, the F-actin-based network (Hirokawa and Takeda 1998) may participate in creating a scaffold for SV clustering, and/or in supporting ordered vesicle mobility. Moreover, it has been suggested that F-actin anchors synaptic vesicles to AZs by a labile link formed with synapsin, a vesicle protein (De Camilli et al 1990; Greengard et al. 1993)

Several lines of evidence suggest that a deregulation of actin plays an important role in SMA pathogenesis. SMN with its binding partner hnRNP R has been shown to interact with β-actin mRNA and mediates the transport along the axon. A decrease in actin protein content in growth cones of SMN-deficient motor neurons in culture and reduced axon growth adds further weight to this line of thoughts (Rossoll et al 2003). Additional SMN-deficient PC12 cells have increased levels of neuronal profilin IIa protein, accompanied by a decrease in levels of plastin3 (Bowerman et al 2009). This is leading to an inappropriate activation of the Rho/ROCK pathway. RhoA is a small GTPase that plays an important role in the regulation of actin cytoskeletal dynamics, which is essential for a balanced equilibrium between actin depolymerization and filament stabilization (Luo et al 1997). In neurons, by signaling through various pathways, RhoA mediates neuronal growth, formation, polarization, regeneration, branching, pathfinding, guidance, and retraction (Govek et al 2005).

Given that actin might play a key role in SMA pathogenesis, we wanted to explore further its content and distribution relative to SVs in TVA motor terminals of SMA mutant

---

**Figure 7:** Simplified model of how SMN depletion impacts the regulators of actin cytoskeletal dynamics and thereby causes in motor neuron degeneration (Bowerman et al 2009 p. 72; reproduced with kind permission from author).
1.6 Neuro Muscular Junction (NMJ) Pathology in SMA

Previous research about spinal muscular atrophy using animal models provided important insights regarding the pathomechanisms of this disease.

Rosoll and colleagues (2003) could show in motor neurons in culture that SMN together with its binding partner heterogeneous nuclear ribonucleoprotein (hnRNP) R interacts with the β-actin mRNA and mediates its transport along the axon. Among others, this study provided evidence for a motor neuron specific function of SMN and indicated that actin plays an important role in SMA pathogenesis.

Short-range movements of organelles, such as vesicles and mitochondria, belong, among others, to the processes mediated by actin in motor neurons (Langford 2002). Moreover the F-actin-based network may participate in creating a scaffold for synaptic vesicle clustering (Hirokawa et al. 1989) and it has been suggested that F-actin anchors synaptic vesicles to AZs by a labile link formed together with synapsin, a vesicle-associated protein (De Camilli et al. 1990; Greengard et al. 1993).

Functional studies at the NMJ of SMA mice revealed that evoked neurotransmitter release is decreased by approximately 55% in TVA muscle, indicating a decreased number of fused vesicles. On the other hand asynchronous release is increased by ~300% due to an anomalous augmentation of intraterminal bulk Ca$^{2+}$ in SMA mice. A possible explanation of this augmentation is a decreased Ca$^{2+}$ reuptake by mitochondria during trains of action potentials (Ruiz et al 2010).

The work described here has the aim to explore the amount and organization of Synaptic Vesicles (SVs), mitochondria, and actin in nerve terminals of SMA mice compared to controls, in order to provide morphological evidence for the observed disturbed neurotransmission at the NMJ of SMA mice. Moreover, it is my aim to gain insight into the question whether a deregulation in actin dynamics provides a possible link to abnormalities observed in SMA pathogenesis.

All experiments regarding SVs, mitochondria and actin were done with TVA muscle of 14 days (P14) old mice in a mouse model of a severe form of this disease.

The study about SVs has been extended in the following aspect:

To get insight into the maturation process in SMA disease, the distribution of SVs were studied additionally in seven days (P7) old mice, both in TVA and levator auris longus (LAL) muscle. The LAL consists of a rostral and a caudal band. The caudal band is known to be more affected than the rostral band in the disease (Murray et al. 2008). This allowed me to study SVs for three different muscle conditions at P7 and P14. Another characteristic of the LAL is that the postsynaptic terminals have been described to mature almost at the same pace as in control mice, while the presynaptic morphology shows abnormalities in P5-P6 SMN deficient mice (Murray et al 2008). It was my aim to test in LAL muscle whether SVs abnormalities occur also in the absence of a postsynaptic pathology to gain
insight into the question how much pre- and the postsynaptic phenotypes are interrelated in SMN deficient mice.

In particular the following aspects will be quantified:

1. Synaptic Vesicles
   
   (a) Total area of SV clusters
   (b) Mean area of a single SV cluster
   (c) Number of SVs clusters

2. Mitochondria
   
   (a) Total area of *Mito Tracker* spots
   (b) Mean area of a single *Mito Tracker* spots
   (c) Number of *Mito Tracker* spots

3. Organization of Synaptic Vesicles and Mitochondria within the nerve terminal

4. Amount and distribution of F-Actin related to SVs clusters.
Chapter 2

Materials and Methods

2.1 Materials

2.1.1 Mouse Model

SMA mouse lines were kindly provided by Dr. A. Burghes (Ohio State University, Ohio, USA) and are presently available in the Animal House in Seville. Experimental mice were obtained by breeding pairs of SMA carrier mice (Smm+/--;SMN2+/++;SMNΔ7+/+) on a FVB/N background. Mice were housed, handled, and bred in accordance with the regulations of the E.U. for animal care and use.

2.1.2 Equipment

<table>
<thead>
<tr>
<th>Genotyping</th>
<th>Equipment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Centrifuge</td>
<td>Eppendorf 5415 D</td>
</tr>
<tr>
<td>Multipipette</td>
<td>Eppendorf plus</td>
</tr>
<tr>
<td>Thermocycler</td>
<td>Biometra TGradient</td>
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<td>UV Transilluminator</td>
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<table>
<thead>
<tr>
<th>Surgery</th>
<th>Equipment</th>
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<td>Mayor scissors</td>
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<tr>
<td>Forceps</td>
<td>F·S·T stainless (Germany) 14022-12</td>
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<td>Microsurgical scissors</td>
<td>Nopa CE Germany Stainless AC778/02</td>
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<td>Surgical Blade</td>
<td>Steriler CE 0434</td>
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## 2. Materials and Methods

### Immunocytochemistry

<table>
<thead>
<tr>
<th>Materials</th>
<th>Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Microscope Slides</td>
<td>Menzel Glaeser</td>
</tr>
<tr>
<td>Coverslip</td>
<td>Menzel Glaeser</td>
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<tr>
<td>Orbital Shaker</td>
<td>Biosan 0S-20</td>
</tr>
<tr>
<td>Plates</td>
<td>Cellstar</td>
</tr>
<tr>
<td>Vapor Pressure Osmometer</td>
<td>Wescor 5500</td>
</tr>
<tr>
<td>pH Meter</td>
<td>Crison Basicco</td>
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<tr>
<td>Scale</td>
<td>Mettler PJ360</td>
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### Microscopy

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<td>Nikon SMZ1500</td>
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<tr>
<td>Confocal Laser Scanning Microscope</td>
<td>Olympus FV 1000</td>
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### Image Analysis

<table>
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<th>Website</th>
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<tbody>
<tr>
<td>ImageJ</td>
<td><a href="http://www.macbiophotonics.ca/downloads.htm">www.macbiophotonics.ca/downloads.htm</a></td>
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### 2.1.3 Genotyping

#### Genotyping

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<th>Reagent</th>
<th>Supplier</th>
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<tr>
<td>Alkaline Lysis Reagent</td>
<td>(25mM NaOH; 0.2 mM EDTA)</td>
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<tr>
<td>Neutralization Reagent</td>
<td>40mM Tris-HCL adjusted to pH)</td>
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<td>Ethidium Bromide</td>
<td>Sigma 160535</td>
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#### Primers

<table>
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<th>Supplier</th>
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<tbody>
<tr>
<td>Neo B Primer</td>
<td>Sigma VC00021</td>
</tr>
<tr>
<td>A2R Primer</td>
<td>Sigma 700023</td>
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<tr>
<td>A2F Primer</td>
<td>Sigma 700025</td>
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#### Oligonucleotides

| dNTPs                | Invitrogen 10043 |

#### Enzymes

<table>
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<th>Enzyme</th>
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<tr>
<td>Taq 5U</td>
<td>Biotools 10043</td>
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#### Solution and Buffers

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<tr>
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<th>Composition</th>
</tr>
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<tbody>
<tr>
<td>PBS Buffer 10X</td>
<td>Biotools 10044</td>
</tr>
<tr>
<td>Electrophoresis Buffer</td>
<td>192 mM Glycine, 0.1% SDS, 25mM Tris-HCl, pH 8.3</td>
</tr>
<tr>
<td>SDS-Sample Buffer</td>
<td>0.125 M Tris-HCl, pH 6.8, 4% SDS, 0.15 M DTT, 20% Glycerol, 0.01% Bromphenol blue</td>
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### 2.1.4 Immunocytochemistry

#### Chemicals

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<th>Supplier/EC</th>
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<tr>
<td>Paraformaldehyde (PFA)</td>
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<tr>
<td>Phosphate Buffered Saline (PBS)</td>
<td>Sigma-Aldrich EC 231-834-5</td>
</tr>
<tr>
<td>Albumin from Bovine Serum (BSA)</td>
<td>Sigma-Aldrich EC 232-936-2</td>
</tr>
<tr>
<td>Glycine</td>
<td>Sigma-Aldrich EC 200-2722</td>
</tr>
<tr>
<td>Slowfade Gold Antifade Reagent</td>
<td>Invitrogen 586012</td>
</tr>
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</table>
2.1 Materials

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Concentration</th>
<th>Type</th>
<th>Reference</th>
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</thead>
<tbody>
<tr>
<td>Primary antibody Anti-VAChT</td>
<td>1:500</td>
<td>polyclonal rabbit</td>
<td>Invitrogen 139 103</td>
</tr>
<tr>
<td>Primary antibody Anti-SV2</td>
<td>1:500</td>
<td>monoclonal mouse</td>
<td>DSHB 149-81791</td>
</tr>
<tr>
<td>Secondary antibody Alexa 488</td>
<td>1:500</td>
<td>polyclonal donkey</td>
<td>Invitrogen A11029</td>
</tr>
<tr>
<td>Secondary antibody Alexa 647</td>
<td>1:500</td>
<td>polyclonal donkey</td>
<td>Invitrogen A21236</td>
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<table>
<thead>
<tr>
<th>Dye</th>
<th>Concentration</th>
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<th>Reference</th>
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<tr>
<td><em>Mito Tracker</em> Orange CMX Ros</td>
<td>400 nM</td>
<td>thiol-reactive chlormethyl moiety</td>
<td>Invitrogen M7510</td>
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<tr>
<td>α-Bungarotoxin Alexa Fluor 647</td>
<td>1.25 µl/ml</td>
<td>snake toxin</td>
<td>Invitrogen L2911</td>
</tr>
<tr>
<td>α-Bungarotoxin Rhodamine Alexa flour 647 phalloidin</td>
<td>1.25 µl/ml</td>
<td>snake toxin</td>
<td>Sigma T0195</td>
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<tr>
<td>Alexa Fluor 647 phalloidin</td>
<td>25 µl/ml</td>
<td>mushroom toxin</td>
<td>Invitrogen A22287</td>
</tr>
</tbody>
</table>

Table 2: Summary of reagents used for immunostaining.

Solutions and Buffers

Hepes solution:

- 145 mM NaCl
- 5 mM KCl
- 2 mM CaCl₂
- 1 mM MgCl₂
- 15 mM glucose
- 10 mM Hepes.

NaOH was used to adjust the pH between 7.35 and 7.4. The osmolarity was adjusted between 300 and 310 mmol/kg by using an osmometer.
2.2 Methods

2.2.1 Genotyping

SMA mice could be distinguished from wild type mice through clinical observation; however, all mice were genotyped for confirming their genetic characteristic.

**Visual and clinical Identification of SMA**  The SMA mice (Δ7) are smaller in size and appear weak and less active compared to WT mice. Further identification was done through the righting reflex test: when turned onto there back SMA mice showed some latency in righting themselves.

**Marking and caudal biopsy**  Five days after birth each mouse received an individual identification number by cutting a fingertip. Through this mark the mice could be identified before the experiment. The PCR genotyping was done using tail DNA.

**DNA Extraction**  To lyse the cells, the tissue samples from caudal biopsy were centrifuged for 30 sec at 13,000 rpm with 50 µl added alkaline lysis reagent and then incubated, first for 30 min at 65°C, followed by 15 min incubation at 98°C. After lysis, the samples were placed on ice for 1 min and treated with 50 µl Neutralizing Reagent. To separate the DNA from the rest, the samples were centrifuged for 1 min at 13,000 rpm. After centrifugation 50 µl of the supernatant, containing the DNA, were pipetted and put into a clean tube.

**PCR**  The polymerase chain reaction (PCR) is a technique to amplify a particular DNA sequence. With this technique we could amplify a DNA sequence identifying either the wild type gene or SMA variant. The method relies on thermal cycling, consisting of cycles of repeated heating and cooling of the reaction for DNA melting and enzymatic replication of the DNA. During the first step the DNA physically separate into two strands through a high temperature around 90°C. At a lower temperature, around 50°C, the specific primers can complementarily bind to the DNA. Starting from the primer, at a temperature around 65°C, each strand is then used as the template in DNA synthesis by the heat stable DNA polymerase (Taq Polymerase) to selectively amplify the target DNA. Building blocks for the synthesis are the deoxyribonucleoside triphosphates (dNTPs). Repeating this cycle, in our case 35 times, leads to an exponential amplification of the target DNA. The protocol we used is based on a single tube reaction with 3 primers, two located in the WT gene (A2F and A2R) region and one in the neomycin cassette present in KO mice.
2.2 Methods

<table>
<thead>
<tr>
<th>PRIMER</th>
<th>MER</th>
<th>Sequence 5’-3’</th>
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</thead>
<tbody>
<tr>
<td>NeoB</td>
<td>21</td>
<td>GCA GCT GTG CTC GAC GTT GTC</td>
</tr>
<tr>
<td>A2R</td>
<td>25</td>
<td>CCT TAA AGG AAG CCA CAG CTT TAT C</td>
</tr>
<tr>
<td>A2F</td>
<td>21</td>
<td>GCT CTT GGC AAC CCT ACT GTC</td>
</tr>
</tbody>
</table>

Table 3: Primer Sequences

The reaction was done with the following components:

- 9.3 µl MQ water
- 4 µl MgCl2 (25mM)
- 2.5 µl buffer 10X (Biotools)
- 0.5 µl dNTPs 25mM (Invitrogen)
- 0.5 µl Neo B primer 10 µM (Sigma)
- 0.5 µl A2R primer 10 µM (Sigma)
- 0.5 µl A2F primer 10 µM (Sigma)
- 0.2 µl Taq 5U/ µl (Biotools)
- 1 µl DNA

Using the following PCR program: (1) 95 °C / 4 min, (2) 95 °C/ 1:30 min, (3) 62 °C / 1:30 min, (4) 72 °C / 1:30 min; 35 x steps 2 to 4, (5) 72 °C / 4 min, (6) 4 °C / 1 h.

Electrophoresis

The amplified target DNA could be detected by gel electrophoresis, showing two different bands, one at 900 base pairs (bp) corresponding to the KO and another at 600 bp corresponding to the WT gene. Both bands were amplified in heterozygous mice whereas in homozygote mice only one band was detectable (Fig.8)

Figure 8: Electrophoresis showing null (-/-), heterozygous (+/-) and WT (+/+) genotypes with 900 bp corresponding to absence of the gene and 600 bp to the presence of the gene.
2.2.2 Muscle Dissection

Mice were sacrificed by means of \( CO_2 \) and immediately exsanguinated.

**Transversus Abdominis (TVA)**

The TVA is the inner muscle of the abdominal wall. For an optimal penetration during immunochemistry all tissues above the TVA need to be removed carefully (Figure 9). As a first step the mouse was positioned in dorsal decubitus and the abdominal skin was opened through a midline incision and pulled to the sides. Second, the whole left hemi-abdominal muscle wall, up to the sternum, was cut out and pinned to the bottom of a 2 ml chamber, over a bed of cured silicone rubber (Sylgard, Dow Corning). Third, the Obliquus Externus muscle and the fat tissue were removed carefully to obtain a free area of intact TVA muscle fibers with their corresponding nerve branches.

![Figure 9: Abdominal muscle wall from mice. In red Transversus Abdominis Muscle (TVA) used for experiments (self-made drawing).](image)

**Levator Auris Longus (LAL)**

The LAL is located directly under the neck skin reaching from the cranial midline towards the ears (Fig. 10). For the dissection, the hair was removed by shaving the area between the left ear and the cranial midline. Second, the left neck muscle block with skin was removed and pinned upside down to the bottom of a 2 ml chamber. Before removing several other muscles (M.Spleniue Capitis, M. Capitis Post Mayor, M. Cervicoscularis) located over the LAL, the skin underneath was pulled to the side. Finally, the LAL was cleaned from fat and connective tissue. Figure 10 shows the LAL with the two constituent muscle bands (rostral and caudal).
2.2 Methods

2.2.3 Immunocytochemistry and Toxin-Conjugated Labeling

Visualization of the different investigation targets

**Postsynaptic side.** The endplates were labeled with α-Bungarotoxin (α-BTX), a snake toxin that binds to the muscle-type nicotinic acetylcholine receptor (nAChR). α-BTX was conjugated to either rhodamine or Alexa Fluor 647.

**Synaptic Vesicles** were marked by using a primary antibody, anti-VACHT, against the vesicular acetylcholine transporter. This primary antibody was visualized using a secondary antibody containing a fluorescent dye. The vesicular acetylcholine transporter VACHT is an integral membrane protein with 12 putative trans-membrane domains. VACHT translocates acetylcholine from the cytoplasm into synaptic vesicles, where it stays until release.

A second type of vesicle staining was done using a primary antibody against the synaptic vesicular transmembrane protein (SV2). This primary antibody was again visualized using a secondary antibody containing a fluorescent dye.

**Mitochondria** were labeled with *Mito Tracker*, a fluorescent dye containing a mildly thiol-reactive chlormethyl moiety. This cell-permeant probe is oxidized when it enters an actively respiring cell. Depending on the mitochondrial membrane potential *Mito Tracker* diffuses passively across the membrane and accumulates in active mitochondria (Poot et al 1996). Specific staining of mitochondria depends critically on the viability of the preparation. Therefore, the dissection was done as quickly as possible with a minimum of cell damage. **Actin** was labeled with phalloidin, a high-affinity probe for filamentous-actin.
(F-actin). The toxin phalloidin is extracted from a mushroom, amanita phalloides and conjugated to a bright, far-red-fluorescent dye, *Alexa Fluor 647*.

**The protocols**

The different protocols differ only slightly from each other nevertheless they are described completely to facilitate replication of the experiment.

**Synaptic Vesicles and mitochondria staining**

The muscles were dissected in hepes solution at pH between 7.35 and 7.4 (see section 2.3.4.3. Solutions and Buffers), followed by 40 minutes incubation with 400 nM *Mito Tracker* diluted in hepes solution. After washing with hepes solution, the preparations were fixed in 4 % paraformaldehyde, washed with PBS and incubated at 4 °C over night. Subsequently they were incubated with glycine 0.1 M in PBS for 30 min at room temperature to bind and removes aldehydes left over from the fixation step. Subsequently muscles were permeabilized with 1% Triton X-100 in PBS. Unspecific binding sites were blocked by incubation with 5% BSA in PBS-T for 3 hours, followed by incubation with primary antibody (anti-VAChT) 1:500 (2.5% BSA in PBS-T 1%) for 30 minutes at room temperature. Muscles were stored at 4 °C over night. The next day they were incubated at room temperature for 30 min; they were washed using 0.05 % Triton in PBS, then incubated with the secondary antibody (goat anti-rabbit Alexa 488 diluted 1:500) and the postsynaptic dye (1.25 µl/ml BTX Alexa Fluor 647) in 5% BSA in PBS-T 0.05%. After washing with PBS-T 0.05% the muscles were finally mounted under the stereoscopic microscope with slowfade medium.

**Synaptic Vesicles staining with SV2 and AChT labeling**

The muscles were dissected in hepes solution (see section solutions and buffers). Subsequently fixed in 4% paraformaldehyde and washed with PBS, followed by 30 min incubation with glycine 0, 1 M in PBS at room temperature and permeabilization with 1% Triton X-100 in PBS. Unspecific binding sites were blocked by incubation with 5 % BSA in PBS-T for 2 hours at room temperature and then at 4 °C over night. After incubation at room temperature for 30 min, the preparations were incubated with primary antibodies (anti-SV2) and (anti- VAChT) 1: 500 (2.5% BSA in PBS-T 1%) for 2 hours at room temperature, followed by washing with PBS-T 1%. Together with the postsynaptic dye (1 µl /ml BTX Rhodamine), they were incubated for 1 hour with secondary antibodies goat anti-rabbit Alexa 488 and goat anti-mouse Alexa 647 1:500 diluted in 5% BSA in PBS-T 0.05 %. Before mounting the preparation under the stereoscopic microscope they were washed with PBS-T 0, 05 %.

**Synaptic Vesicles and Actin staining**

The muscles were dissected in Hepes solution (see section 2.3.4.3. solutions and buffers) and then fixed in 4 % paraformaldehyde, washed with PBS and set at 4 °C over night. Subsequently they were incubated with glycine 0, 1 M in PBS at room temperature, then permeabilized with 1 % Triton X-100 in PBS. Unspecific binding sites were blocked by incubation with 5% BSA in PBS-T for 3 hours. Followed by incubation with primary antibody (anti-VAChT) 1:500 (2.5% BSA in PBS-T 1%) for 30 minutes at room temperature. They were stored at 4 °C over night. The next day they were incubated at room temperature for 30 min; they were washed using 0.05% Triton in PBS, then incubated with
the secondary antibody (goat anti-rabbit Alexa 488 diluted 1:500), the actin dye (25 µl/ml Alexa fluors 647 phalloidin) and the postsynaptic dye (1.25 µl/ml BTX-Rhodamine red) in 5% BSA in PBS-T 0,05 %. After washing with PBS-T 0,05 % the muscles were finally mounted under the stereoscopic microscope with slowfade medium.

2.2.4 Confocal Microscopy

Muscles were imaged with an upright Olympus FV1000 confocal laser scanning microscope, equipped with three excitation laser lines (488, 561 and 633 nm). A 10x objective was used for coarse localization of nerve terminals within the preparation. Images were taken using a 63x oil-immersion objective with a numerical aperture of 1.4. Fluorescence emission was detected sequentially by selecting the appropriate filter settings, as given in the Table 4.

<table>
<thead>
<tr>
<th>Label</th>
<th>Exc Max (nm)</th>
<th>Em Max nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>α-Bungarotoxin Alexa Fluor 647</td>
<td>650</td>
<td>668</td>
</tr>
<tr>
<td>α-Bungarotoxin Rhodamine</td>
<td>552</td>
<td>579</td>
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<td>Secondary Antibody Alexa488</td>
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<tr>
<td>Alexa flour 647 phalloidin</td>
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<td>668</td>
</tr>
<tr>
<td>Mito Tracker Orange CMX Ros</td>
<td>554</td>
<td>576</td>
</tr>
</tbody>
</table>

**Table 4:** Filter settings for detection of specific fluorescent labels.

2.2.5 Image Analysis with ImageJ

Transferring the image from the Olympus Fluoview program to the Image J program

The Olympus Fluoview program was used to acquire the emission of each dye, for example BTX far red, Mito Tracker and Alexa Flour 488 in individual channels in three (X-Y-Z) dimensions. The Z dimension was represented as a series of slices, showing the emission from different depth levels at intervals of 0.5 µm. The images were imported from the Olympus Fluoview program to the Image J program in a RGB red green and blue color format as a sequence containing images at about 8 z-positions per fluorescence channel. The image sequence from different depth levels of each channel was transformed to a Z-projection, when required. After setting the scale in µm corresponding to one pixel, Z-projections were used for imaging analysis. In some cases single slices were used for analysis, if mentioned in the text.

Measuring the postsynaptic area

The Z-projection showing the postsynaptic area labeled by BTX-Rhodamine or BTX-Far red, was converted from RGB color image to an 8 bit grayscale image. The image was segmented into regions of interest and background by setting interactively lower and upper
threshold values. The area of a selected region was then measured in square micrometers using image J routines.

**Measuring the area of synaptic vesicle clusters**

In order to measure the synaptic vesicle clusters, lower and upper threshold values were set as for the postsynaptic side, with the difference that the areas of interest were analyzed in “particles analyzing mode” by manually setting the threshold until edges of each vesicle cluster were defined. Then the individual areas of each cluster were measured.

**Measuring the area of the mitochondria clusters**

In contrast to synaptic vesicles, mitochondria are present in all cells. To measure only the mitochondria inside the terminal we used the postsynaptic area as a mask and cleared all signals outside this region. The mitochondria inside the synaptic terminal were then measured in “particles analyzing mode”, similar to the analysis of vesicle clusters.

**Actin measurements**

In transverse single confocal slices the diameters of F-actin rings surrounding SVs could be estimated from their respective line-intensity profiles. From maximum projected confocal images actin areas were determined automatically, by defining masks based on brightness thresholding. To measure only the actin inside the terminal we used the postsynaptic area as a mask and cleared outside signals before measuring actin.

**Quantification of the nearest-neighbor distances between SV clusters and mitochondria spots**

Nearest-neighbor distances between mitochondria and SV clusters were determined by fitting each of the red and green spots to a 2D gaussian distribution, and using the distances between the peaks (x,y) of the distributions.

**2.2.6 Statistics**

Unless otherwise stated all values indicate mean ± standard error of the mean (SEM) and numbers \( (n) \) refer to the number of terminals. Differences between groups were tested using the \( t \)-test (2 tailed).
Chapter 3

Results

The results of this work have been incorporated into our publication “SMN Requirement for Synaptic Vesicle, Active Zone and Microtubule Postnatal Organisation in Motor Nerve Terminals” (Torres-Benito et al 2011).

3.1 Synaptic Vesicles

3.1.1 Deficits in Synaptic Vesicle Maturation

Synaptic Vesicles (SV) together with mitochondria are the main organelles in the nerve terminal. A disturbed distribution or reduced amount of SVs could provide an explanation for the observed 55% decrease of evoked neurotransmitter release in TVA muscle of SMA mice (Ruiz et al 2010).

We used an antibody against the vesicular ACh transporter (VAChT) in combination with a secondary antibody, conjugated with a green fluorophore to explore the SVs spatial organization. In particular the total area of the terminal covered by SVs was determined. To better localize the NMJ we labeled, like in all the following experiments, also the postsynaptic side with α-Bungarotoxin (α-BTX), which binds to the nicotinic acetylcholine receptor (nAChR), in this case conjugated with rhodamine.

Spinal Muscular Atrophy progresses during development. To get insight into the presynaptic maturation process, we examined synaptic structures in mice at postnatal day seven (P7) and 14 (P14). We used TVA muscle, one of the most affected ones in this disease (Murray et al 2008) (Ruiz et al 2010).

The confocal images show that SVs are organized in clusters (Fig. 11 for P7 and Fig. 12 for P14). In wild-type (WT) mice, at the beginning of the NMJ postnatal maturation period (first week of life), SV clusters were observed to be small (Fig. 11A), but by the second week, as WT NMJs matured, SVs covered larger areas of the terminal (Fig. 12A). In SMN deficient terminals, however, SV clusters remained small (compare Fig. 12A, upper and lower panels).

Also, at the postsynaptic side, images taken at P14 showed signs of immaturity.
Figure 11: Representative en face views of NMJs from the TVA muscle at P7 stained with BTX-Rho (red), which binds specifically to postsynaptic AChRs, and anti-VAChT (green), which labels synaptic vesicles. Images are Z-stack projections. A. Terminals of WT (upper panels) and SMN\textDelta7 mice (lower panels) respectively, at P7. Panels B; C & D show mean values (±sem) of BTX area, total synaptic vesicle area and their ratio respectively both for WT (white bars) and mutants (grey, filled bars). Scale bars: 10 μm *:\(P < 0.05\); **:\(P < 0.005\); ***:\(P < 0.0005\).
3.1 Synaptic Vesicles

Figure 12: Representative en face views of NMJs from the TVA muscle at P14 stained with BTX-Rho (red), which binds specifically to postsynaptic AChRs, and anti-VACHT (green), which labels synaptic vesicles. A. Terminals of WT (upper panels) and SMNΔ7 mice (lower panels) respectively, at P14. Note the signs of immaturity in the SMNΔ7 terminals: no clear folders or perforations at the postsynaptic side in SMNΔ7 (left, lower panel) compared to WT (left, upper panel). SVs at the presynaptic terminal remain small in SMNΔ7 mice (right, lower panel). B. Mean postsynaptic area in SMNΔ7 (grey, filled bars) and in WT terminals (white bars) are significantly different. C. The total area of vesicles stained with VACHT is smaller in Δ7 mice than in WT motor terminals. D. The relative area covered by vesicles in relation to the area covered by postsynaptic receptors is less than half in SMNΔ7 terminals. Scale bars: 10 μm.

*: P < 0.05; **: P < 0.005; ***: P < 0.0005.
While in control littermates band structures were apparent, representing BTX, which resides at the lips of infoldings, no such features were obvious in images from mutants (compare Fig. 12A upper and lower panels). A quantitative analysis of the images is shown in Figure 11B-D and 12B-D.

Measuring the area of the postsynaptic side (see methods) showed that NMJs are smaller in mutants both at P7 and P14 (Fig. 11B & 12B, respectively). Therefore, the areas covered by SVs were normalized to those of the postsynaptic terminal, in order to compare the fractional area covered by SVs between mutants and control. The total surface of the terminal covered by SVs was $\sim 30\%$ smaller in SMN deficient terminals ($n = 37$ NMJs from 3 mice) than in WTs ($n = 38$ NMJs from 3 mice) already at P7 (Fig. 11C & D $P < 0.0001$). At P14, this difference became even larger, near 50%, (Fig. 12C & D; $P < 0.0001$; WT: $n = 52$; $\Delta 7$: $n = 58$ terminals). In addition, the size of SV clusters was smaller in mutants than in WTs, both at P7 ($\sim 30\%$; $P = 0.02$) and, much more so, at P14 ($75\%$; $P < 0.0001$). This analysis shows the phenomena of maturation in a remarkable way: in littermate controls the clusters size increased and the number of clusters diminished (see Table 5) from P7 to P14. In contrast, in mutants this process was not observed, no change in cluster size, nor in the number of clusters per terminal took place during the second week of life.
3.1.2 VACHT and SV2 colocalize in SMA Motor Terminals

To test whether the clustering of SVs observed in the TVA muscle of SMA mutants with anti-VACHT antibodies could also be seen with another vesicle marker, a double staining of SVs using antibodies against VACHT, and the synaptic vesicle protein 2 (SV2) was performed. The vesicular fluorescence patterns observed with anti-VACHT (red) were also seen with anti-SV2 (green), both in WT (Fig. 13A), and in mutant terminals (Fig. 13B). Quantitative analysis of the immunofluorescence signals indicated that the colocalization indexes (see Methods) of these two markers were \( \sim 0.9 \) in both types of mice. These data corroborate the smaller size of the clusters in SMA mutant terminals, and rule out the existence of an atypical vesicle population devoid of the vesicular transporter.

Figure 13: VACHT and SV2 colocalize in SMA synaptic vesicles. A & B. Examples of WT and SMA \( \Delta 7 \) terminals at P14 from TVA muscles. Images are Z-stack projections. Scale bar: 5 \( \mu \text{m} \).
3. Results

3.1.3 A reduced SV pool can coexist with a mature Postsynaptic Terminal in LAL Muscle.

The data presented above show that SMA deficiency produces pre- and postsynaptic defects in the TVA muscle, and raise the question of how much the pre- and the postsynaptic phenotypes are interrelated in SMN deficient mice. Therefore, it was explored whether the vesicle content was also anomalous in the LAL muscle of SMA mutants, a muscle, in which postsynaptic terminals have been described to mature almost at the same pace as in control mice Murray et al. (2008). The analysis, as described for the TVA, therefore, was repeated in the LAL.

Because previous work has identified a selective vulnerability of the nerve terminals in the caudal band in comparison to the rostral band of the LAL muscle (Murray et al., 2008; Ruiz et al., 2010), the size of the postsynaptic side and the organization of SVs at NMJs in both bands was compared.

In the rostral band there was no significant difference in the size of the postsynaptic side at P7 (Table 5), and P14 (Fig. 14 E and Table 5). Also in the caudal band there was no difference in size of the postsynaptic side at P7 (Table 5). Only in the caudal band at P14 the size of the postsynaptic side was ~23% reduced in mutants (Fig. 14B). In comparison: in TVA muscle at P14 the size of the postsynaptic side was ~33% smaller in mutant relative to WT.

At P7 in the rostral band, SV area and mean cluster area (Table 5) were not different in WTs (n = 21 terminals) and mutants (n = 22 terminals), while both parameters were significantly reduced in mutants at P14 (Table 5). More specifically, at P14 the mean SV area was reduced ~52% and mean size of SVs clusters was reduced ~66% in mutant (Fig. 14F & G), (P = 0.0003 and P = 0.006, respectively). In the caudal part, however, both parameters were already decreased at P7 in mutants. At P7 the SV area was ~49% and the SV cluster size was ~57% smaller (Table 5). At P14 this difference even increased, the SV area was ~73% and the SVs cluster size ~ 83% reduced (Fig.14C & D, respectively).

These results demonstrate that the observed abnormalities are not a property of the TVA muscle alone. Moreover, they confirm the selective vulnerability of the caudal band of the LAL muscle. The finding that the maturation of the postsynaptic end-plates takes place normally while there are early presynaptic defects, shows that pre- and postsynaptic defects can occur independently from each other.
Figure 14: Synaptic vesicle reduction and anomalous distribution in SMA motor terminals from LAL muscles despite mature postsynaptic side. A & B. Representative images of pre- and postsynaptic characteristics of NMJs in the LAL muscles of WT (upper panels) and SMNΔ7 (lower panels) mice at P14. Postsynaptic terminals were labeled with BTX-Rho (red) and presynaptic vesicles with antibodies against VAChT (green). Note the clustering of SVs despite the mature appearance of the postsynaptic side in the SMNΔ7 terminal. Scale bar: 5μm. B-G. Mean postsynaptic area (B & E), SV area (C, F), and cluster area (D, G) in SMNΔ7 and WT terminals, from rostral (B-D) and caudal (E-G) divisions of the muscle at P14. Note that the area of vesicles and the cluster sizes were significantly reduced at P14. However, in the caudal part SV deficiencies in SMNΔ7 terminals were evident since P7 (n = 16 WT terminals and n = 15 SMNΔ7 terminals). Scale bars: 10 μm *:P < 0.05; **:P < 0.005; ***:P < 0.0005 (modified from our publication Torres-Benito L, Neher MF, Cano R, Ruiz R, Tabares L. 2011).
Table 5: Summary of synaptic vesicle and postsynaptic side properties in TVA and LAL muscle. Comparison between WT (wild type) and D7 (SMA mutant) in TVA (Transversus Abdominis Muscle) and LAL (Levator Auris Longus muscle) at P7 (seven days old mice) and P14 (fourteen days old mice), regarding BTX Area ($\mu m^2$) (area of postsynaptic side), SV Area/BTX Area (Area of the synaptic vesicles divided by area of postsynaptic side), SV cluster size ($\mu m^2$) (mean size of one single vesicle synaptic vesicle cluster) and number of clusters/BTX Area. Data in mean, sem (standard error of the mean) and n (number of observations).

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3.2 Mitochondria

3.2.1 Mitochondria Clusters are smaller in SMA Motor Terminals

Mitochondria provide ATP for synaptic transmission, ion pumps, and the assembly of the actin cytoskeleton involved in the clustering of SVs and mitochondria themselves ([Hirokawa and Takeda 1998, Langford 2002]). Furthermore they regulate intraterminal Ca$^{2+}$ levels, providing an especially important buffering action during trains of action potentials ([David...])
3.2 Mitochondria

<table>
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**Table 6: Summary of mitochondria properties in TVA muscle.** Comparison between WT (wild type) and D7 (SMA mutant) in TVA (Transversus Abdominus Muscle) at P14 (fourteen days old mice), regarding Number of Mito Tracker spots per terminal. Total area of Mito Tracker Spots /BTX Area (µm²) and Mito Tracker spot size (µm²). Data in mean, sem (standard error of the mean) and n (number of observations).

Therefore, a reduction in the number of mitochondria, alterations in their spatial distribution, or impairment of their functional capacities may decrease their Ca²⁺ buffering capability and produce toxic Ca²⁺ overloads. To assess whether the spatial distribution of mitochondria or their density was altered in SMN deficient terminals, the TVA muscle was incubated with Mito Tracker (400 nM), a cell-permeant probe that is sequestered by intact mitochondria and retained during cell fixation (Poot et al 1996).

Figure 16 A-D (central panels) shows representative examples of the distribution of Mito Tracker labeled mitochondria in motor terminals at P14. The total area stained by Mito Tracker per terminal was about half in mutants (19.4 ± 2.7 µm²; n = 20) as compared to WT terminals (38.8 ± 4.9 µm²; n = 22; P = 0.002). Normalization of these areas to postsynaptic surfaces also gave significantly smaller values in mutants (WT: 0.29 ± 0.03; mutant: 0.19 ± 0.02; P = 0.01) (Figure 15A). This difference was mainly due to the smaller size of the Mito Tracker spots (WT: 1.72 ± 0.36 µm², n = 360 spots; mutant: 0.89 ± 0.1 µm², n = 214 spots. P = 0.03) in mutants (Figure 15 C). Quantification of the number of Mito Tracker microregions per terminal showed no significant differences between WT and mutants (WT: 22.9± 2.1 spots/terminal, n= 18 terminals; mutant 23.5 ± 3.1, n = 19 terminals) (P = 0.9) (Figure 15C).

These properties of mitochondria in TVA muscle of SMA mice compared to wild type are also shown in Table 6.

3.2.2 Co-clustering of Mitochondria and SV clusters

Next, we explored the spatial relationship between Mito Tracker spots and SV clusters. In most cases, as in the illustrated examples, mitochondria (red) appeared near SV clusters (green) in both WT (Fig. 16A & B) and mutant terminals (Fig. 16C & D). Moreover, in many cases a core of mitochondria was clearly surrounded by a SV rim (Fig. 16A-D, right panels), reminiscent of the donut-like structures described in the calyx of Held, Wimmer et al (2006) and Perkins et al (2010).
Figure 15: A-D. Staining of mitochondria with *Mito Tracker* in WT (wild type) and D7 (mutant mice) in TVA muscle at P14.  A. The total area stained by *Mito Tracker* normalized to the postsynaptic surfaces was significantly reduced in mutant.  B. Quantification of numbers of *Mito Tracker* spots per terminal showed no significant difference between WT and mutant.  C. The size of *Mito Tracker* spots is smaller in mutant.  D. Relationship between mean areas covered by SV and mitochondria in WT (blue) and SMA Δ7 (red) terminals. *: $P < 0.05$. 
To quantify this observation we measured the nearest-neighbor distances between SV clusters and mitochondria spots. Nearest-neighbor distances were determined by fitting each of the red and green spots to a 2D Gaussian distribution, and using the peak (x,y) of each distribution. The results in nine mutant terminals (202 measurements) and 13 WT terminals (282 measurements), showed that in mutants SV and mitochondria were closely associated, as in controls (Fig. 16 E). Even more, the percent of red-green spots nearest distances within 1 µm were larger in mutants (41%) than in WT (31%) (P = 0.0006), as expected for the smaller sizes of SV and mitochondria cluster in mutants. Taken together, these results show that in nerve terminals mitochondria and SVs clusters are localized in pairs. This aspect of the spatial organization is not altered in mutant terminals, whereas the fractional area covered by mitochondria is reduced.

### 3.3 F-actin

#### 3.3.1 F-actin forms ring-like Structures around SV Clusters

Filamentous actin (F-actin) is a prominent cytoskeletal element in nerve terminals. The F-actin-based network may participate in creating a scaffold for SV clustering, and/or in supporting ordered vesicle mobility ([Hirokawa et al][]{ref}) In addition, it has been suggested that F-actin anchors synaptic vesicles to AZs by a labile link formed with synapsin, a vesicle protein ([De Camilli et al][]{ref} and [Greengard et al][]{ref}).

SMN deficiency produces defects in β-actin mRNA axonal transport, and a decrease in actin protein content in growth cones of motoneurons in culture ([Rossoll et al][]{ref}). We explored F-actin content and distribution in presynaptic TVA motor terminals of SMA mutant mice. F-actin was revealed by binding to fluorescent Phalloidin-Alexa 647, which binds to all isoforms of F-actin but not to monomeric actin ([Wulf et al][]{ref}). In addition, to gain insight into the role of F-actin in the organization of SV, the distributions of F-actin relative to SV were also examined.

F-actin was localized in the sub-plasmalemmal region, extending into the cytoplasm and forming a thin network. Figure 17A shows a WT terminal with F-actin labeled with phalloidin (upper panel), SV labeled with VACHT antibodies (green), and postsynaptic receptors labeled with BTX-Rho (red, lower panel). In Figure 17B a typical terminal from a mutant mouse is shown (left panel, phalloidin image; central panel, anti-VACHT and BTX-Rho merged). In some cases, F-actin was also clearly visualized in the axon (Fig. 17B, arrow in right panel). In WT and mutant mice, F-actin filaments run parallel to the major axis of the terminal and SV clusters are positioned along them.

The association between SV clusters and actin could be better visualized in transverse single confocal slices. Figure 17C (WT) and 17D (mutant) present two representative examples showing SV clusters surrounded by F-actin, forming ring-like structures, similar to actin rings described in the lamprey reticulospinal giant synapse ([Shupliakov et al][]{ref}). Interestingly, the diameter of F-actin rings surrounding SV were apparently smaller in mutant than in WT terminals, as could be appreciated by their respective line intensity
Figure 16: Mitochondrial density in SMA presynaptic terminals is reduced. Data are from the TVA muscle (P14). A - D. The upper panels (A & C) show Z-stack projections while the lower panels (B & D) show single confocal sections. The staining from left to right are: BTX-A647 (grey), SVs (green), Mito Tracker (red), merge of SV and mitochondria. E. Nearest neighbor distributions of SV clusters and mitochondrial regions (calculated from their respective center of mass), in WT and mutants terminals. Scale bars A-D: 5 µm; insets: 600 nm (from our publication Torres-Benito L, Neher MF, Cano R, Ruiz R, Tabares L., 2011)
profiles across SVs and phalloidin loops (Fig. 17E). However, these structures were not visualized frequently enough for statistical analysis. Thus, the percentage of the terminal area covered by phalloidin, normalized to the postsynaptic area, was quantified (Fig. 17F). In WT mice, although the area of the terminal covered by phalloidin was apparently larger than in mutants (WT: $41 \pm 9\%$, $n = 6$ terminals; mutants: $34 \pm 4\%$, $n = 6$ terminals), this difference did not reach statistical significance ($P = 0.5$).

Taken together, these results show that, although a slight reduction of the F-actin based network is difficult to quantify, F-actin is still present in high amounts in the mutant and, similarly to the wild type case, closely associated with SVs in the form of ring-like structures.
Figure 17: F-actin and SV are closely associated in SMA motor terminals. Data are from the TVA muscle (P14). A - B. Representative examples of the distribution of F-actin (gray) in a WT (A) and a mutant (B) terminal. The same terminals are also shown stained for SV (green) and postsynaptic receptors (red). Arrow in the right panel of B points to actin in the axon. C - D. Phalloidin (blue), SV (green), and AChRs stained with BTX-Rho (red), showing the organization of actin around SV clusters in a WT and a mutant (D) terminal. E. Intensity profiles across actin loops and SV clusters showing the reduced diameter and thickness of the loop in mutants. F. Phalloidin areas in WT and mutant terminals were not significantly different (P < 0.05). Scale bars: A & B: 5 µm; C & D: 2 µm (from our publication Torres-Benito L, Neher MF, Cano R, Ruiz R, Tabares L., 2011).
Spinal muscular atrophy (SMA), a frequent neurodegenerative disease, is caused by reduced levels of functional SMN. SMN is a ubiquitously expressed protein, however reduced levels of SMN predominantly affect motor neurons and muscles.

The following studies revealed important insights to understand this phenomena:

- In search of a neuron-specific function of SMN it was shown that SMN interacts with β-actin mRNA and mediates its transport along the axon [Rossoll et al. 2003].

- Functional studies at neuromuscular junctions (NMJ) showed that the evoked neurotransmitter release was decreased by approximately 55% in most affected muscles, indicating a decreased number of fused vesicles, while asynchronous release is increased by ~300% due to an anomalous accumulation of intraterminal bulk Ca$^{2+}$ in SMA mice [Ruiz et al. 2010]. A possible explanation of this augmentation is a decreased Ca$^{2+}$ reuptake by mitochondria during trains of action potential.

Starting from these results, here, we undertook a manifold morphological study regarding synaptic vesicles (SVs), mitochondria and actin at the NMJ of SMN deficient mice in order to better understand the disturbed neurotransmission mentioned above and to learn more about actin as a possible key player in SMA pathogenesis.

### 4.1 Synaptic Vesicle Clusters remain small in SMA mice during Maturation

Our fluorescence microscopy images using an antibody against the vesicular acetylcholine transporter (vAChT) show clearly, that synaptic vesicles are organized in clusters in the presynaptic motor terminal.

We demonstrated that in wild type muscles, as a pattern of maturation, the SV clusters increased highly in size while their number diminished between P7 and P14. In mutant mice, however, the clusters remained small and the number of clusters did not change. Hence at P7 the total surface of the terminal covered by SV and the average size of any
given cluster was equally reduced by $\sim 30\%$ in mutant mice. At P14 these differences became even larger, while in littermate controls the cluster size increased, and the number of clusters diminished. In mutants no change in cluster size, nor in the number of clusters per terminal, took place. In numbers, at P14 the total surface of the terminal covered by SV was near 50\%, and the size of SV clusters was $\sim 75\%$ reduced in mutant mice. This is in agreement with an electron microscopy study of NMJ in tibialis anterior muscle at P13 demonstrating that individual synaptic vesicles within the presynaptic terminal had normal diameter and morphology, but their overall density within the presynaptic terminal area was reduced by 56\% (Kong et al 2009). Accordingly, another ultrastructural study showed a decrease in SV density in SMA mice (Lee et al 2011). In contrast in diaphragm at P14 no significant difference in vesicle number was found (Kariya et al 2009). The finding that the size of SV clusters increases during maturation leads to the question whether single vesicles increase in size, their density within clusters decreases, or the number of vesicles increases. The amplitude of spontaneous miniature endplate currents (mEPC), representing the muscle response to neurotransmitter release from a single vesicle, was found to be normal in SMA mice. This indicates that the size of single vesicles is normal and suggests that a reduced number of vesicles or increased vesicle density leads to the smaller SV cluster size (Kong et al 2009).

However our images provide only information about the area of maximum projections of SV clusters. Also, light-microscopic images of clusters do not provide numbers on the density of vesicles within clusters. Disregarding such problems and assuming tightest possible vesicle packing a very rough estimation on how many SV might form one SV cluster can be obtained by dividing the volume of a sphere with the diameter of one SV cluster (2.6 $\mu$m, in wild type TVA at P14) by the volume of one vesicle (0.0395 $\mu$m in diameter). Doing so we arrive at a very high number - 285000. This is orders of magnitude larger than estimates from electron microscopic studies on the number of vesicles per active zone for mammalian glutamatergic synapses (200 to 250). It is, however, compatible with estimates in ribbon synapses (10000 to 13000), if a volume fraction of about 0.05 is assumed (see (Fernandez-Alfonso and Ryan 2006), 2006, for a review on vesicle numbers). Under the same assumption the total vesicle pool in frog neuromuscular junction contains $\sim 500,000$ vesicles ((Ceccarelli et al 1973); (Heuser and Reese 1973); (Molgo and Pecot-Dechavassine 1988); (Rizzoli and Betz 2005)). We could show that the area of the cluster size increases three times from P7 to P14 in wild type TVA, which would mean a 5.2 times increase in the number of vesicles at constant packing density. In contrast in SMA mice the cluster size remains almost the same from P7 to P14.

However, repeating similar experiments using a fluorescent microscope with a resolution high enough to identify single vesicles or else an electronmicroscopic analysis would certainly be an interesting extension of our study.
4.2 Selective Vulnerability of Motor Neurons innervating the Caudal Band of Levator Auris Longus muscle (LAL) despite only modest Abnormalities at the Postsynaptic Side

To study SV in a different anatomical region we performed the same analysis we did in TVA muscle in LAL muscle. In LAL muscle we showed that SV pathology is more severe in the caudal band than in the rostral band. There was no differences between mutant and wild type regarding the SV area and mean size of SV clusters in the rostral band at P7, while both parameters were significantly reduced in mutants at P14. On the contrary, in the caudal band both parameters were already decreased at P7. The reduction in the caudal band of the area covered by SV and mean size of SV clusters was even more profound than in TVA muscle.

These results, identifying the caudal band to be more affected in the disease than the rostral band, is in agreement with a study showing that presynaptic nerve terminal loss was widespread in the caudal band but almost absent in the rostral band (Murray et al., 2008). After excluding other possibilities, it is likely that this selective vulnerability in the caudal band is due to the Fast Synapsing (FaSyn) characteristic, while in rostral band the Delayed Synapsing (DeSyn) characteristic is dominant (Murray et al., 2008). However this does not demonstrate a direct link between FaSyn characteristics and selective vulnerability since genetic tools to experimentally manipulate FaSyn and DeSyn characteristics are not available yet. To get more insight into the question what makes some muscles more vulnerable than others, it would be of interest to proof whether also the TVA belongs to the class of Fast Synapsing muscles.

The comparison of postsynaptic sizes revealed that the reduction at the postsynaptic side is much more profound in TVA than in LAL. This is in agreement with a study showing that the reduction of muscle fiber diameter and postsynaptic endplate shrinkage was profound in TVA, but only modest in LAL (Murray et al., 2008). For a better understanding of the disease it is essential to define whether postsynaptic pathology at the NMJ is a consequence of pathology of the nerve or occurs independently from presynaptic pathology.

In case postsynaptic pathology is a consequence of presynaptic pathology we would expect that abnormalities at the postsynaptic side are correlated to pre-synaptic abnormalities. Here we could show, however, that in the caudal band of LAL a marked reduction of the pool size of SV is present in the presynapse, despite only modest abnormalities of its postsynaptic side. This finding suggests that postsynaptic pathology occurs largely independently from presynaptic pathology. According to that it was shown that presynaptic nerve terminal loss can occur independently from postsynaptic endplate shrinkage at the neuromuscular junction (Murray et al., 2008). Thus, it is unlikely that postsynaptic pathology is a plain consequence of disturbed neuronal signaling. One possible explanation for this is that perturbed SMN gene activity may be required in both neuronal and muscle tissue to result in the pathological phenotype (Chan et al., 2003). Neuron-specific functions
4. Discussion

of SMN are well documented with SMN being shown to be important for events including axon outgrowth, pathfinding, and neuronal maintenance (McWhorter et al. 2003; Rossoll et al. 2003; Ferri et al. 2004).

The muscle-specific role is still not comprehensively explored. Deletion of SMN exon 7 in mouse muscle is known to cause severe muscular dystrophy (Cifuentes-Diaz et al. 2001) and a recent study, utilizing a drosophila model of SMA, has identified an important role for the SMN protein in specifying form and function of muscle sarcomeres (Rajendra et al. 2007). Accordingly, a study in SMA mice shows that the molecular profile of muscle pathology in SMA, with an upregulation of voltage-dependent anion-selective protein 2 (Vdac2) and downregulation of parvalbumin, was distinct from that of denervated muscle. This adds further to the hypothesis that SMN-induced changes in muscle cannot solely be attributed to motor neuron degeneration (Mutsaers et al. 2011).

One possibility to better clarify to what extent pre- and postpathology are interrelated would be to repeat the current set of experiments in a mouse model, in which SMN expression has been reduced specifically either in muscle or nerve tissue.

4.3 Mitochondrial Clusters are smaller in SMA Motor Terminals

We could show that mitochondria are organized in clusters and that the number of clusters per terminal is the same in mutant and wild type at P14. However the mean size of clusters was reduced by about one half in mutants and accordingly the total area of mitochondrial clusters per terminal was $\sim 50\%$ reduced in mutants.

Little is known about the role of mitochondria in SMA pathogenesis. In one electron microscopy (EM) study on P14 SMA mice presynaptic mitochondria in diaphragmatic muscle are decreased in size but not in numbers (Kariya et al. 2008). In contrast, another electron microscopy study on P14 SMA mice showed in Tibiales Anterior muscle a reduced density of mitochondria, while mitochondrial morphology was normal (Kong et al. 2009). Therefore it remains unclear, whether our finding of a reduced size of mitochondrial clusters is due to smaller mitochondria or reduced number of mitochondria in SMA mice. The staining of mitochondria with the fluorescent dye Mito Tracker is problematic: the uptake of Mito Tracker into mitochondria depends on the negative resting membrane potential. Therefore mitochondria had to be in a fairly viable condition to achieve specific staining. We had to exclude many preparations from further analysis because Mito Tracker did not enter sufficiently into the mitochondria.

On the other hand, although the Mito Tracker signal does not provide information about the functionality of mitochondria in all aspects, one advantage of this probe is that it does confirm that mitochondria are still active to maintain a negative resting membrane potential. Hence we could compare the amount of active mitochondria in mutant and wild type.

Mitochondrial dysfunction is known to play a central role in the pathogenesis of ma-
4.3 Mitochondrial Clusters are smaller in SMA Motor Terminals

Mito- nor neurodegenerative diseases such as Alzheimer’s, Parkinson’s, Huntington’s disease, and amyotrophic lateral sclerosis. This is not surprising regarding the two faces of mitochondria. Besides being sources of ATP, mitochondria are also the main producers of reactive oxygen species (ROS). If ROS levels overwhelm the defense mechanisms of the cells, oxidative damage of proteins, lipids, and DNA occurs ((Valko et al 2007); (Carvalho et al 2009)). Another consequence of mitochondrial impairment is the release of pro-apoptotic factors and a reduced Ca\(^{2+}\) buffering capacity. At the presynaptic side rapid recovery of Ca\(^{2+}\) transients is essential for nerve signaling. Interestingly at NMJ of SMA mice the asynchronous neurotransmitter release (see introduction) was found to be increased by $\sim$ 300 % indicating an anomalous augmentation of intraterminal bulk Ca\(^{2+}\) during repetitive stimulation (Ruiz et al 2010).

Permanent Ca\(^{2+}\) overload is toxic for the cell as a result of several cross-amplifying cascades, which will be briefly mentioned here. First, Ca\(^{2+}\) activates the cysteine proteases of the calpain and caspase families, that degrade a variety of substrates and trigger apoptotic cascades ((Chan and Mattson 1999); (Nixon 2003)). Second, Ca\(^{2+}\) induces oxidative stress, among others through the activation of oxygenases (Lipton, 1993; Mattson and Pedersen, 1998). Third, Ca\(^{2+}\) triggers apoptosis through activation of pro-apoptotic proteins such as Bax, Par-4, and p53 leading to mitochondrial membrane permeability changes, release of cytochrome C, and caspase activation ((Duan and Cooke 1999); (Dargusch et al 2001); (Culmsee and Mattson 2005)).

Regarding the finding of Ca\(^{2+}\) accumulation at the presynaptic side of SMA mice and the obliterative effect of Ca\(^{2+}\) overload on the cell in general, it is obvious that answering the question how Ca\(^{2+}\) accumulation comes about in SMA might provide an important step towards a possible therapy of the disease. At the presynaptic side, the influx of Ca\(^{2+}\) through voltage-dependent and ligand-gated channels in the plasma is a critical signal for the release of neurotransmitters ((Yuste et al 2000); (Burnashev and Rozov 2005)).

Normally, Ca\(^{2+}\) is then removed from the cytoplasm by plasma membrane Na\(^+\)/Ca\(^{2+}\) exchanger, plasma membrane and ER Ca\(^{2+}\)-ATPase, as well as Ca\(^{2+}\)-binding proteins such as calbindin and parvalbumin (Hof et al 1999). Moreover, Ca\(^{2+}\) can be transported into and released from mitochondria ((Werth et al 1994); (Duchen 2000); (Nicholls 2003)). This raises the question to what extent mitochondrial dysfunction contributes to the altered Ca\(^{2+}\) homeostasis in SMA. Here we could show that the amount of active mitochondria is about half in mutants. Further work is required to find out whether the remaining mitochondria still function in their role to regulate intraterminal Ca\(^{2+}\) levels. This could be addressed through intramitochondrial Ca\(^{2+}\) measurements upon stimulation using Ca\(^{2+}\) -sensitive fluorescent indicator dyes (Vila et al 2003).
4. Discussion

4.4 SV Clusters and Mitochondria are localized in Pairs

Surprisingly our confocal images with a double staining of SV and Mitochondria revealed that SV clusters are localized right next to mitochondrial clusters (Fig. 16). In many cases SV were scattered around a core of mitochondria forming a ring-like structure. Quantification of the nearest neighbor distances confirmed this observation and showed that this aspect of spatial organization is not altered in mutant terminals. Similar to our finding, SV has been described to form donut-like assemblies around a central cluster of mitochondria in the calyx of Held synapse. Further estimation assume that each of these units consist out of ∼ 800 synaptic vesicles and six to nine mitochondria (Wimmer et al. 2006).

In the calyx of Held this structural specialization appeared with the onset of hearing which was accompanied by distinct changes in the properties of synaptic transmission. After the onset of hearing, mature synapses show much less short-term synaptic depression, low release probabilities, and narrow action potentials (Taschenberger and von Gersdorff, 2000; Taschenberger et al., 2002). It is therefore likely that the pairing from SV clusters and mitochondria is tightly linked to enhanced synaptic function both in the calyx of Held and probably also in the NMJ. As a consequence the ATP, produced by mitochondria, could be directly and more efficiently used for priming, transport, and filling of SVs. Moreover mitochondria might be perfectly located to fulfill their role as Ca\(^{2+}\) buffer, which is essential for sustaining high-frequency synaptic transmission. Taken together, we here show, to our knowledge for the first time at the NMJ, that SV clusters are colocalized with mitochondria. This provides evidence that the structural principle described in the calyx of Held might be of more general relevance for pre-synaptic terminals.

4.5 Actin Dynamics in SMA Pathogenesis

The finding, that the SMN protein is involved in the transport of β-actin mRNA along the axon, raises the question to what extent a disturbance of the actin cytoskeleton influences the pathogenesis in SMA (Rossoll et al. 2003).

For our fluorescent microscopy study we labeled β-actin with phalloidin and the postsynaptic side with BTX-Rhodamine in TVA muscle at P14. In addition, to gain insight into the role of β-actin in the organization of SV, we labeled SV with VACHT antibodies. Our images showed that β-actin is localized in the sub-plasmalemmal region and extends deep into the cytoplasm forming a thin network. Interestingly, single confocal slices revealed that β-actin surrounds SV clusters forming ring-like structures. Similarly, in the lamprey giant synapse and hippocampal synaptic terminals actin was described to form ring-like structures around SV (Shupliakov et al. 2002; Sankaranarayanan et al. 2003).

To our knowledge it is the first time that this organization of actin forming ring-like structures around SV is described at the NMJ of WT mice and we could show that this organization pattern was equally found in SMA mice. However the diameter of β-actin rings surrounding SV appeared smaller in mutant than in WT terminals, as could be estimated
by respective line intensity profiles across SV and phalloidin loops. Moreover, we showed that in WT mice the area of the terminal covered by phalloidin was apparently larger than in mutants (WT: $41 \pm 9\%$, $n = 6$ terminals; mutants: $34 \pm 4\%$, $n = 6$ terminals). However this difference did not reach statistical significance ($P = 0.5$).

A decreased actin content in SMA nerve terminals would be in agreement with the reduced $\beta$-actin protein and mRNA staining in growth cones in culture (Rossoll et al. 2003). Compatible to the proposed function of SMN in the transport of $\beta$-actin mRNA along the axon, it was shown that there is a significant reduction of actin signals in the distal axon, and growth cone of SMN deficient motor neurons compared to the wild type, whereas in the soma no significant difference to the wild type has been found (Mayer 2009). In addition it was shown that treatment of Smn -/-; SMN2 motor neurons in culture by cyclic adenosinmonophosphate (cAMP) induces increases of SMN protein in the soma, axon, and growth cones. This could be explained by the presence of a CREII binding element in the SMN promoter region, which mediates effects of cAMP and therefore stimulates the SMN-transcription of the examined cells. Interestingly, along with the increase of SMN protein in cAMP treated SMN -/-; SMN2 motor neuron an increase of the $\beta$-actin signal in the distal axon and growth cone compared to untreated cells was found (Mayer 2009). This adds further weight to the supposition that a deficit in SMN protein, which underlies SMA, leads to a disruption of the actin cytoskeleton. Synaptic efficacy depends in multiple ways on the actin cytoskeleton: Using an actin depolymerizing agent, latrunculin A, it could be shown that actin is essential for the development and maintenance of young synapses, defined by synaptophysin-labeled vesicle clusters, synaptic vesicle recycling, and ultrastructure. Also, it is proposed that actin, because it is so highly regulated, is a principal target for stabilizing or destabilizing signals that ultimately determine synapse maintenance or elimination (Zhang and Benson 2001). Moreover actin seems to serve as a molecular scaffold to increase the local concentration of an important regulatory molecule, synapsin, in the vicinity of the presynaptic vesicle cluster (Sankaranarayanan et al. 2003). Synapsin, a soluble protein of the synaptic terminal, is regulated through phosphorylation and dephosphorylation. Synapsin forms a link between synaptic vesicles and actin (Greengard et al. 1993). After vesicle fusion, synapsin disassociates from synaptic vesicles due to its curvature-sensing ALS-motif (Krabben et al. 2011). Synapsin then accumulates in the endocytic region and interacts with a soluble pool of G-actin, where the actin-rich cytomatrix is formed. Then synapsin reassociates with vesicle membranes (Bloom et al. 2003). Together with synapsin actin has been implicated at multiple steps of the synaptic vesicle cycle, including endocytosis, transport of recycled vesicles to the synaptic vesicle cluster, and exocytosis (Shupliakov et al. 2002).

Regarding this multiple functions of actin in the vesicle cycle it appears plausible that disorganization in the actin cytoskeleton can affect SV and further neurotransmission.

In the present work we demonstrate that SV clusters remain small during maturation and the total area of the terminal covered by vesicles is reduced by about 50\% in mutant mice at P14. This is accompanied by a decrease of evoked neurotransmission by 55\% and a two fold reduction of the evoked endplate signals (Kong et al. 2009; Ruiz et al. 2010). Besides the reduction of SV vesicles we could show that mitochondria are organized in clusters
next to SV clusters at the NMJ. The total area covered by mitochondria per terminal was
∼ 50% reduced in mutants. Axonal transport of mitochondria is ensured by microtubule-
based motors whereas short range movement of vesicles and organelles at growth cones
and nerve terminals depends primarily on actin-myosin-based transport (Langford 2002).
The interaction between microtubule-and actin-based motors suggests that mitochondrial
transport can be coordinated along both cytoskeletal systems (Bridgman 2004). These
tight interactions seem to be controlled through complex phosphorylation processes of mo-
tor proteins, cytoskeletal structures, or mitochondrial membrane proteins (Müller et al
2005).

In conclusion, actin strongly interacts with SV and mitochondria. Also other charac-
teristics of the disease phenotype, like for example reduced axon growth and branching
(Rossoll et al 2003; Kariya et al 2009), are highly actin-dependent mechanisms. Interes-
tingly, in order to explain the tissue-specific nature of SMA, the only neuron specific
function of the SMN protein which could be identified is that SMN mediates the transport
of β-actin mRNA along the axon. It remains to be determined, whether SMN also medi-
ates the transport of other mRNA species. Above all, further work is required to clarify
whether the characteristics of the disease phenotype described here, such as abnormal SV
clustering and reduction of mitochondria, occur independently or are the consequence of
a dysfunction of the cytoskeleton, which would assign a key role in SMA pathogenesis to
actin.
Chapter 5

Abstract

Proximal spinal muscular atrophy (SMA) is an autosomal recessive disorder characterized by degeneration of the second motor neurons and progressive paralysis with atrophy of the proximal muscles. After cystic fibrosis, SMA is the most common autosomal recessive disorder in humans and the most common genetic cause for infant mortality. SMA, monogenic in origin, is caused by a mutation in a single gene, the Survival of Motor Neuron 1 (SMN1) gene, which leads to reduced levels of Survival Motor Neuron (SMN) protein. SMN is a ubiquitously expressed protein with house-keeping roles in snRNP biogenesis and pre-mRNA splicing. However, reduced levels of SMN predominantly affect motor neurons and muscles for reasons that are poorly understood. It was shown that SMN interacts with β-actin mRNA and mediates its transport along the axon. Functional studies at neuromuscular junctions (NMJ) revealed that the evoked neurotransmitter release was decreased by approximately 55% in most affected muscles, indicating a decreased number of fused vesicles, while asynchronous release is increased by ∼300% due to an anomalous accumulation of intraterminal bulk Ca$^{2+}$ in SMA mice. A possible explanation of this augmentation is a decreased Ca$^{2+}$ reuptake by mitochondria during trains of action potential.

This study presents a comprehensive fluorescent confocal microscopy analysis of the organization and abundance of synaptic vesicles (SVs), mitochondria, and actin in nerve terminals of SMA mice (Smn-/-; SMN2; SMNΔ). The results are also part of our joint publication of Torres-Benito L, Neher MF, Cano R, Ruiz R, Tabares L (2011) SMN Requirement for Synaptic Vesicle, Active Zone and Microtubule Postnatal Organization in Motor Nerve Terminals. PlosOne 6(10):e26164.doi:10.1371/journal.pone.0026164. We visualized SV clusters with an antibody against the vesicular acetylcholine transporter (VChT) and could show that in Transversus abdominis (TVA) muscle SV clusters remain small during maturation with the total area covered by SVs reduced by ∼50%. This severe reduction of SVs was also found in the caudal band of the Levator auris longus muscle (LAL). However, only modest abnormalities at the postsynaptic side were detectable in this muscle. This finding of presynaptic pathology, despite an almost normal postsynaptic status, supports the hypothesis that SMN-induced changes in muscle cannot be solely attributed to motor neuron degeneration.

Next, we stained mitochondria with Mito Tracker and found that the area covered by
mitochondria in mutants was about half compared to wild type. Surprisingly SVs and mitochondria were strongly colocalized. In many cases a core of mitochondria was clearly surrounded by a SV rim. This spatial organization was not altered in mutants and might be of general relevance for the functional integrity in nerve terminals.

Phalloidin stains of F-actin showed that F-actin forms ring-like structures around SV clusters. These structures and the percentage of the terminal area covered by actin, normalized to the postsynaptic area, were smaller in mutant mice.

Actin is implicated at multiple steps of the synaptic vesicle cycle. Short range movement of vesicles and organelles, such as mitochondria at growth cones and nerve terminals, depends primarily on actin-myosin-based transport. Further work is required to clarify whether the characteristics of the disease phenotype described here, such as abnormal SV clustering and reduction of mitochondria, occur independently or are the common consequence of dysfunction of the actin cytoskeleton, which would assign a key role in SMA pathogenesis to actin.
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List of abbreviations

α-BTX  α-Bungarotoxin
ALS  amyotrophic lateral sclerosis
cAMP  cyclic adenosinmonophosphate
DeSyn  delayed synapsing
EM  electron microscopy
EPCs  evoked endplate currents
FaSyn  fast synapsing
KO  knockout
LAL  levator auris longus
MEPC  spontaneous miniature EPC
nAChR  nicotinic acetylcholine receptor
NMJ  neuro muscular junction
P7  postnatal day seven
P14  postnatal day fourteen
PCR  polymerase chain reaction
RGG  arginine glycine
ROS  reactive oxygen species
RRP  readily releasable pool
SMA  spinal muscular atrophy
SMN  survival motor neuron 1
SV  synaptic vesicles
SV2  synaptic vesicular transmembrane
Vdac2  voltage-dependent anion-selective protein 2
WT  wild-type
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I would like to thank the Department of Medical Physiology and Biophysics at the School of Medicine in Sevilla and everyone, who contributed to turn my stay in Sevilla into a time, in which I learned the wonderful Spanish language and culture, met inspiring people and had a great scientific experience through my studies about a motor neuron disease, such as SMA.
Curriculum Vitae

I, Margret Neher, was born on the 18th of April 1987 in Göttingen and grew up as the fifth child of Eva-Maria Neher and Erwin Neher in Eddigehausen, a village near Göttingen. From 1994 to 2006 I visited the Rudolf Steiner School in Weende (Göttingen). During High School I went on a student exchange from October 2003 until April 2004 in Napier, New Zealand.

In April 2007 I started my medical studies at University of Göttingen and took my first state exam successfully in 2009. I completed my medical clerkships in Neurology (Tübingen, Germany), Psychiatrics (Göttingen, Germany), Cardiology (Oxford, Great Britain) and Dermatology (Rabat, Morocco). I obtained teaching experiences as an assistant at anatomy preparations in 2008 and physiology tutor to third semester students in 2009.

From 2000 to present I have been an active canoeist. My best achievements in Canoe Polo have been the gold medalist in the European Champion ships in 2003 and 2007, silver medalist in the World Champion ships 2004, 2008 and 2010 and again gold medalist of the World Games 2005. In flat water kayak racing, an olympic sport, I won 2006 the European Champion Under 23 Championship in K2 over 500m, and the German Championship senior in K1 over 200m and 500m. With these achievements, I qualified among the top 10 preselection candidates for the 2008 Olympics in Peking.

From February to October 2010 I lived in Sevilla, Spain to perform all my experiments for my doctoral thesis in the lab of Prof. Lucia Taberes. The results are published in the journal PlosOne as indicated below.

After the clinical semesters I spent my practical year partly abroad. For the obligatory subjects Surgery and Internal Medicine I went to the Waid Spital in Zürich (Switzerland) from August to October 2012, to the Hospital Italiano, Buenos Aires (Argentina) from October 2012 to February 2013, and from February to April 2013 to Neu-Mariahilf in Goettingen. The optional subject, Neurology, I fulfilled at the Klinikum of the University of Goettingen. In spring 2014 I will prospectively conclude my medical studies.

Publications:

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