Dissertation

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Role of plectin in microtubule dynamics

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“Caminante, son tu huellas el camino, y nada más; caminante, no hay camino, se hace camino al andar.”

“Wanderer, your footsteps are the path, and nothing more; wanderer, there is no path, the path is made by walking.”

- Antonio Machado
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SUMMARY

Plectin, a polypeptide of very large size (>500 kDa) and member of the plakin cytolinker protein family, is one of the most abundant and versatile cytolinkers expressed in mammalian cells. One of its outstanding features is its functional diversity that is mainly based on the alternative splicing of a series of different first coding exons resulting in different isoforms.

A number of observations made in previous studies, including the targeting of overexpressed plectin isoform 1c (P1c) to microtubules (MTs), the partial colocalization of P1c with MTs in cultured cells, and the in vitro binding of neural plectin to MT-associated proteins (MAPs), pointed towards a potential interaction of plectin with MTs. Moreover, ACF7 and BPAG1, other members of the plakin cytolinker protein family, have been shown to be involved in the regulation of MT dynamics, both acting as MT stabilizers. Based on these previous observations and additional unpublished data from this laboratory one of the major goals of my thesis was to identify the role of plectin in MT dynamics. Of particular interest in this context was the question whether plectin isoform P1c antagonizes MAP-mediated MT stabilization, and if so, what was the underlying molecular mechanism. I report here that contrary to other cytolinker proteins, P1c acts as a destabilizer of MTs and that this destabilization relies on IF-dependent P1c localization.

Since P1c is a major isoform in keratinocytes and neurons, I searched for a mechanistic link between plectin-related changes in MT dynamics and alterations in MT-dependent processes of physiological significance in keratinocytes and neurons isolated from P1c-deficient mice. Functioning as a promoter of MT dynamics, I found plectin to affect vital MT-dependent functions and properties of keratinocytes, including glucose uptake, cell division and growth, focal adhesion turnover, shape and polarized migration of cells. P1c-deficient primary DRG neuronal cultures were found to be a suitable system to study the consequences
of altered MT dynamics in neurons. The data revealed alterations in the processes of neuritogenesis, such as outgrowth and branching of neurites, distorted synaptic vesicle transport along neurites, and ability of neuronal membrane to depolarize.

Finally, I describe a model where P1c destabilizes MTs by antagonizing MAP-mediated MT stabilization. The reported findings unmask P1c as a MT destabilizer and show that cytolinker-mediated destabilization is required for the correct performance of multiple physiological processes. These findings point towards a fascinating new feature of cytoskeletal filament cross-talk, namely the potential of IFs to destabilize MTs via an associated cytolinker protein, thereby stimulating MT dynamics.
1 INTRODUCTION

The Cytoskeleton

The cytoskeleton is a system of intracellular filaments responsible for organizing the cell shape and the cell interaction with the environment. It provides mechanical strength as well as enables the cells to rearrange their components to grow, migrate, divide and adapt to changing environmental conditions. Although long time considered as a unique feature of eukaryote cells, prokaryotes also present an active and dynamic cytoskeleton. Prokaryotes show a surprisingly complex cytoskeletal composition that share with the eukaryotic cytoskeleton the main functions and properties, providing resistance to external forces, acting as scaffold or enabling cell division (Wickstead & Gull, 2011).

The cytoskeleton of eukaryotic cells is composed of three major types of filaments. Actin filaments provide protrusive and contractile forces. Microtubules (MTs) form a polarized network allowing vesicle and organelle movement along the cell. Finally, intermediate filaments (IFs) endow cells with mechanical strength and resistance to shear stress. All of them are essential to the spatial organization of the cells, are dynamic and adaptable. Each type of filament is constructed from smaller protein subunits that can be repetitively assembled and disassembled, facilitating rapid structural reorganization. Despite these common features, each cytoskeletal filament type has unique mechanical properties and stability that result in their individual roles. In addition, there are a large number of cytoskeleton-associated proteins that regulate the dynamics of the filaments and coordinate their functions, by regulating their polymerization, providing energy, or anchoring them to cellular structures. Among these proteins we find the MT- and actin-associated proteins and proteins belonging to the cytolinker family.
1.1.1 Actin

Actin filaments or microfilaments are long filamentous polymers of ~7 nm diameter formed by globular actin subunits. They are composed by two parallel protofilaments that twist around each other in a right-handed helix. As a result of the regular and parallel orientation of actin subunits, the ends of the polymer are different presenting distinct kinetic rate constants for association and dissociation. The plus end or barbed end of actin filaments polymerizes and depolymerizes faster than the minus end or pointed end. During this process known as treadmilling, actin subunits are recruited at the plus end and shed from the minus end. When the subunits change rapidly from the free to the polymerized state, but keeping the length of the filament unchanged, the process is known as “steady-state treadmilling”. Actin dynamics can be altered by drugs like phalloidin (that binds to and stabilizes filaments, causing net polymerization), or latrunculin (that binds to and stabilizes actin monomers, causing net depolymerization).

Actin filaments determine the cell shape and are necessary for cell movement. There are many types of actin-based superstructures (Chhabra & Higgs, 2007; Doherty & McMahon, 2008). Actin filaments can be organized as a mesh underlying and tightly in contact with the plasma membrane. They can also form a dense network of highly branched (lammellipodia and ruffles) or long unbranched (filopodia) actin filaments at the leading edge of migratory cells. Long bundles of actin cables (stress fibers) are usually anchored at adhesion sites, and actin-rich structures can be associated with invaginations in endocytic and phagocytic structures.

1.1.2 Intermediate filaments

IFs are a superfamily of 10 nm-fibers. They are named after their diameter, between thin actin filaments (~7 nm) and myosin filaments (~15 nm) (Ishikawa et al, 1968). There is a large
variety of IFs that are differentially expressed in different tissues and were originally classified into five categories (Fuchs & Weber, 1994). (i) Type I IFs correspond to acid keratins and include eleven epithelial keratins, K9-K20, and four hair keratins, Ha1-Ha4. (ii) Type II keratins are basic or neutral, and include eight epithelial keratins, K1-K8, and four hair keratins, Hb1-Hb4. Both types of keratins assemble as obligatory heteropolymers and are expressed differentially at various stages of development and differentiation. (iii) Type III IF proteins encompass vimentin (mesenchymal cell types and transformed cell lines), desmin (smooth, skeletal, and cardiac muscle), glial fibrillary acidic protein (GFAP) (glial cells and astrocytes) and peripherin (peripheral nervous system). Type III IFs form homopolymers, but they can also assemble as a heteropolymers with other type III IF proteins or with neurofilaments (NFs). (iv) The type IV group comprehends NF proteins and α-internexin. NFs are divided into NF-L (light, 62 kDa), NF-M (medium, 102 kDa), and NF-H (heavy, 110 kDa). NF proteins are expressed in axons, dendrites, and perikarya, while α-internexin is also expressed in neurons, but playing a more important role during embryonic development. (v) Type V group contains the proteins composing the nuclear lamina, lamins. Lamins form a meshwork on the inner surface of the nuclear membrane, providing a framework for the nucleus and participating in chromatin organization. There are three types of lamins, lamins A, B, and C, presented in different combinations depending on the cell type and differing in their functions. (vi) There are also a so-called type VI group of IFs that can not be included in any of the previous groups, but they are considered part of the IF superfamily according to their sequence and structure. Within this group we can find nestin and filensin, proteins expressed in the developing central nervous system and during differentiation of the vertebrate lens epithelia, respectively.

All IF proteins share a common structure. Unlike actin and tubulin, they form non-polarized filaments. IFs have a central α-helical domain, the rod, flanked by nonhelical head (N terminus) and tail (C terminus) domains. The assembly takes places after a first formation of
parallel and in-register dimers, followed by their association into stable tetramers. Contrary to other cytoskeletal filaments, IFs do not require auxiliary proteins or factors to assemble. However, their function and reorganization can be regulated by phosphorylation. Different kinases, like protein kinase A, protein kinase B, cAMP-dependent kinase, and cdc2 kinase phosphorylate vimentin (Chou et al., 1990; Eriksson et al., 2004), desmin (Huang et al., 2002), NFs (Shea & Lee, 2011; Sunil et al., 2012), and lamins (Fields & Thompson, 1995).

IFs provide cells with resistance to mechanical stress. Up to date, 93 distinct diseases related to IFs have been identified. Among them we find Alexander disease, Charcot-Marie-Tooth disease and amyotrophic lateral sclerosis, caused by mutations in IF of glial and neural cells (Liem & Messing, 2009). Mutations in desmin cause skeletal muscle dystrophies and cardiopathies (Carlsson & Thornell, 2001). Keratin mutations lead to diseases classified as epidermolysis bullosa simplex (EBS), a heterogeneous group of skin blistering disorders (Intong & Murrell, 2012).

### 1.1.3 Microtubules

MTs are hollow polarized cylinders. They are cytoskeletal polymers composed of α- and β-tubulin monomers. Tubulin monomers assemble linearly into protofilaments that associate laterally to form ~24 nm-wide cylindrical structures. The initial structures forming are small sheets of protofilaments that later close to form MTs containing 13 protofilaments. MT polymerization occurs in three steps. First, MT nucleation requires γ-tubulin that acts as a scaffold for α- and β-tubulin aggregation. The nucleation rate depends on the initial concentration of soluble tubulin. Second, during aggregation or elongation, tubulin dimers are incorporated as a complex with GTP. Finally at steady state, MT length remains constant but the two polymer ends behave different. The plus end incorporates new tubulin dimers, while the minus end looses them. As occurs in actin filaments, this process is named treadmilling.
However, it is very rare that MT length remains constant. Usually individual MTs present long phases of assembly (rescue) and rapid phases of disassembly (catastrophe) (Figure 1). Rapid transitions from one state to the other are known as dynamic instability (Mitchison & Kirschner, 1984; Schulze & Kirschner, 1986). MT polymerization can be altered using drugs.

![Figure 1. Schematic representation of MT dynamics. Tubulin dimers are incorporated as a complex with GTP at the plus end of the MT. At the minus end, tubulin dimers are bound to GDP. Dynamic MTs undergo length fluctuations due to spontaneous phases of catastrophe and rescue (Adapted from Conde & Cáceres, 2009).](image)

Colchicine and vinblastine bind to MTs and induce their depolymerization. By sequestering tubulin dimers, also nocodazole induces MT depolymerization. In contrast, taxol binds to MTs and stabilizes them, inhibiting their dynamics. MTs are required for maintaining cell shape and polarization, cell division, intracellular transport and they participate together with actin
filaments in cell locomotion. In addition, MTs have been shown to play an essential role in organization and dynamics of axons and dendrites. Their dynamics and functions are fine-tuned and regulated by other proteins binding to them. There are proteins that facilitate MT stabilization, like MT-associated proteins (MAPs), and cytolinkers like ACF7/MACF or BPAG1. Proteins leading to MT destabilization are Op18/stathmin, SCG10 (Conde & Cáceres, 2009). MT or MT-related protein dysfunction have been identified in a variety of neurological disorders such as Alzheimer and Parkinson’s disease (Tischfield et al, 2011; Lee et al, 2001).

1.1.4 Microtubule associated proteins (MAPs)

MAPs form a diverse protein family characterized by their interaction with MTs (Olmsted, 1986; Wiche, 1989). In most of the cases MAP-tubulin interaction is responsible for MT stabilization and promotion of polymerization. Although the first attempts to isolate MAPs were done by purifying them from brain due to the high content in MTs of this tissue, MAPs are expressed ubiquitously. MAP proteins have been divided in different categories according to their structure and expression pattern. Among them, tau presents a highly complex variety of isoforms as a result of distinct splicing of transcripts. These isoforms differ in the number of MT-binding repeats and in the presence of exons 2 and 3 (Figure 2). MAP1, MAP2, and tau are expressed primarily in neurons, however MAP2 has been found also in keratinocytes and in other non-neuronal tissues (Wiche et al, 1984; Liu et al, 2007). Tau was identified in other non-neuronal cells like muscle as well (Gu et al, 1996; Janué et al, 2010). E-MAP-115 is another MAP restricted to epithelia (Masson & Kreis, 1993). In contrast, MAP4 is expressed ubiquitously (Parysek et al, 1984). XMAP215 shows also a MT-polymerizing activity in addition to its MAP-stabilizing activity (Brouhard et al, 2008).

Particularly noteworthy is the MAP2/tau family of MAPs (Dehmelt & Halpain, 2004).
The MAP2/tau family includes MAP2, MAP4 and tau. All of them share a C-terminal domain harboring MT-binding repeats and have alternative splice isoforms (Figure 2). MAP2 and tau increase MT stability and rigidity (Felgner et al., 1997) and are responsible for MT-bundling (Lewis et al., 1989). MAP2 is mainly found in dendrites of differentiated neurons (Sanchez et al., 2000) and MAP2c has been related to Alzheimer’s disease due to its specific interaction with apolipoprotein E (Huang et al., 1994). Tau is normally described as an axonal protein but an additional dendritic function of tau has been identified (Ittner et al., 2010). In addition, tau has been shown to inhibit kinesin-dependent traffic of organelles (Stamer et al., 2002; Dixit et al., 2008) and to participate in axonal branching by protecting MTs against severing proteins like katanin (Qiang et al., 2006). Tau has also been implicated in neurofibrillary tangle formation and amyloid-β toxicity in Alzheimer’s disease (Kampers et al., 1999; Ballatore et al., 2007; Ittner...
et al., 2010) and in frontotemporal dementias like parkinsonism, frontotemporal lobar degeneration and motor neuron disease (Fu et al, 2010).

1.1.5 Cytoskeletal linker proteins

The plakin family of proteins (also known as cytolinkers) are responsible for strengthening cells against mechanical stress and regulating cytomatrix plasticity by networking and anchoring cytoskeletal filament systems to organelles and junctional complexes (Wiche, 1998; Fuchs & Karakesisoglou, 2001; Leung et al, 2002; Sonnenberg & Liem, 2007). They are a family of large, multi-domain proteins that link cytoskeletal networks to each other, to organelles and to membranes (Figure 3). They are built up by a combination of modular domains, but not all of

![Figure 3. Mammalian cytolinker proteins. Functional domains of mammalian plakins including actin-binding domain, plakin domain, coiled-coil rod domain, spectrin repeat rod, plectin repeat domains are specified. (R. Spurny, PhD thesis).]
them are shared by all cytolinkers: (i) a calponin-type actin-binding domain (ABD) consisting of
two calponin-homology (CH) domains presented in alternatively spliced forms; (ii) a plakin
domain, containing six α-helical segments, that is important for interaction with different
proteins; (iii) a coiled-coil α-helical rod domain (with 10.4 residue charge periodicity) that
mediates the dimerization of molecules, as it occurs in plectin (Wiche et al., 1991); (iv) a plectin
repeat domain (PRD) composed by four complete and one partial 38-residue motif. PRDs can
be categorized in A, B, and C repeats; the IF-binding domain is located in the PRD domain; (v)
a spectrin repeat (SR) domain characteristic of the spectrin family made up of three α-helices
connected by two loop regions; (vi) EF-hand calcium-binding motifs; (vii) Gas2-homology
region named GAR domain or a GSR domain functioning as MT-binding sites in some of the
cytolinker proteins (Leung et al., 2002).

Seven plakin protein family members have been identified up to date: desmoplakin,
bullous pemphigoid antigen 1 (BPAG1), MT-actin cross-linking factor (ACF7), periplakin,
enoplakin, epiplakin, and plectin; they are expressed in a large variety of tissues.
Desmoplakin is responsible for anchoring IFs to desmosomes, structures that mediate
intercellular adhesion in tissues subjected to mechanical stress (for review see Getsios et al.,
2004). As a result of alternative splicing in the central rod domain, there are two isoforms of
desmoplakin, DPI (322 kDa) and DPII (259 kDa) (Virata et al., 1992). Both isoforms are
expressed in all types of epithelia, but DPI is also found in cardiac cells and in follicular
dendritic cells associated with desmin and vimentin, respectively (Angst et al., 1990; Franke &
Moll, 1987). Desmoplakin is involved in autoimmune diseases such as paraneoplastic
pemphigus and genetic disease like striate palmoplantar keratoderma. In mice, desmoplakin
deletion leads to embryonic death (Leung et al., 2002). The autoimmune skin blistering disease
bullous pemphigoid (Moll & Moll, 1998) is associated with autoantibodies against BPAG1 and
BPAG2. In addition, mutations in the Bpag1 gene lead to sensory neuron degeneration
observed in mice with dystonia musculorum (Brown et al, 1995). Dystonia musculorum is a recessive neuropathy affecting sensory neurons and including disorganization of the cytoskeleton, accumulation of neuronal IFs and muscle weakness (Dalpe et al, 1998; Dalpe et al, 1999). ACF7 also known as MACF1, trabeculin or macrophin, is a cytolinker protein that is highly expressed in epidermis, and responsible for interconnecting MTs and actin filaments (Leung et al, 1999). Albeit no disease involving ACF7 has been reported yet, ACF7-deficient mice die at the gastrulation stage of the embryogenesis (Kodama et al, 2003). Periplakin and envoplakin are closely related and represent the smallest proteins of the plakin family (~200 kDa) (Ruhrberg & Watt, 1997; Ruhrberg et al, 1996). Both proteins are components of the cornified envelope and localized at desmosomes and IFs in differentiated keratinocytes. They are involved in paraneoplastic pemphigus and pemphigus foliaceus autoimmune diseases. Epiplakin is a very large protein (725 kDa) composed solely of PRDs, 13 in humans and 16 in mice. It was originally isolated as an autoantigen in a serum obtained from a patient with subepidermal blistering disease and its expression is restricted to epithelial tissues (Fujiwara et al, 1992; Fujiwara et al, 1996; Spazierer et al, 2003).

1.1.1 MT regulation

MT dynamics and functions are fine-tuned and regulated by other proteins binding to the polymer. Some MT-binding proteins are MT stabilizers or growth promotors, while others promote MT depolymerization (Figure 4). Post-translational modifications (PTMs) of MTs seem to be crucial to spatially and temporally control the activity of MT regulatory proteins.

Post-translational modification (PTMs)

PTMs are chemical modifications that occur to the proteins after their translation. Many PTMs, such as phosphorylation or ubiquitinylation, directly affect the function and activity of the
proteins. Albeit, there is no apparent direct effect of PTMs on MTs, PMTs are emerging as control elements in the regulation of the interaction of tubulin with MT-binding proteins (for a review see Janke & Bulinski, 2011). The best studied PTMs of MTs are acetylation, detyrosination, Δ2-tubulin generation (see below), polyglutamylation, and polyglycylation.

Acetylation of Lys40 on α-tubulin takes place once the MT is polymerized, functioning as a marker of relatively old and stable MTs (Piperno et al., 1987; Bulinski et al., 1988). Acetylated MTs increase the MT-binding affinity of molecular motors like KIF5A, KIF5B/kinesin,

**Figure 4. Model of MT-regulating proteins and PTMs.** MT dynamics are regulated by proteins that bind along the MTs. MT-stabilizing proteins are depicted in green and destabilizing proteins in red.

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and dynein (Reed et al, 2006), stimulating vesicle and membrane bound organelle transport (Friedman et al, 2010). Acetylation of tubulin has also been proposed to induce enzymatic MT severing (Sudo & Baas, 2010).

Tyrosination occurs on dimerized α-tubulin (Raybin & Flavin, 1977) while detyrosination is mediated by a cytosolic carboxypeptidase that is active only on polymerized MTs, leaving a Glu residue exposed at the C terminus. Detyrosinated tubulin can be further converted to Δ2-tubulin (by removal of the Glu residue exposed at the C terminus after tubulin detyrosination) that irreversible locks MTs in a stable state, since tubulin tyrosin ligase is unable to again tyrosinate Δ2-tubulin (Paturle-Lafanèche et al, 1991). Similar to acetylated MTs, detyrosination increases the binding of kinesin and IFs to MT (Liao & Gundersen, 1998; Kreitzer et al, 1999).

Polyglutamylation consists in the progressive addition of one or more glutamate residues near the C terminus of polymerized α- or β-tubulin (Edde et al, 1990). MT-stabilizing proteins like MAPs or MT-severing proteins such as katanin or spastin might be attracted to MTs by long poly-glutamate tails (Bonnet et al, 2001; Lacroix et al, 2010).

The extension of glycine side chains from glutamate residues near the accessible C terminus of α- or β-tubulin is known as polyglycylation, and in contrast to other PTMs, it is confined to cilia and flagella (Redeker et al, 1994).

**MT-stabilizing proteins**

Among the MT-stabilizing proteins, there is a complex and diverse set of MT-interacting proteins that can be classified according to their localization on MTs. According to this, there are MT-stabilizing proteins that bind all along the MT surfaces, plus end- (+TIP), and minus end-binding proteins.
The best known MT-stabilizing proteins are those that belong to the MAP2/tau family. They bind along the sides of acetylated MTs and contribute to their stabilization by crosslinking adjacent MTs (Felgner et al., 1997). However, MAPs are not the only MT-stabilizing proteins that bind along MTs. Septins form a novel family of filamentous GTPases that have been shown to inhibit MT catastrophes and participate in MT guidance during polymerization (Spiliotis, 2010; Bowen et al., 2011). The septin family consists of multiple genes and protein isoforms that can associate with cell membranes or the cytoskeleton. They decorate fragments of the MTs near the nuclear envelope and the cell periphery. Septins have been suggested to function as spatial localizers within the MTs, interacting with MAPs, motors and tubulin. Indeed, SEPT2 was found to compete with MAP4 for MT-binding at poly-glutamylated sites (Spiliotis et al., 2008). In addition, motor proteins, such as kinesins, participate in transporting tubulin heterodimers and oligomers promoting the assembly of tubulin (Kimura et al., 2005).

MT plus ends are subjected to a highly complex regulation. A delay between polymerization and GTP hydrolysis of GTP-β-tubulin creates a GTP-cap. The loss of this cap induces rapid depolymerization of the MTs, however some GTP remnants have been identified in older parts of MTs (Dimitrov et al., 2008). Nevertheless, the GTP-cap is not the only way how the plus end is regulated. The +TIP protein family has been extensively studied. It is composed of structurally unrelated proteins that often colocalize and share common functions (Lansbergen & Akhmanova, 2006; Akhmanova & Steinmetz, 2008; Jiang & Akhmanova, 2011). Albeit there are some +TIP proteins with a MT-destabilizing role, most of them are involved in promoting MT growth and stabilization. Although proteins within the +TIP family can be quite different from each other in structure, they contain serine-rich sequences that have been shown to interact with EB1, and have autoinhibitory domains. The end-binding (EB) protein family consists of proteins of very similar structure and sequence. The members of this family are expressed depending on the differentiation stage of different cell types. EB1 is a
ubiquitously expressed EB protein that dimerizes and promotes MT polymerization by increasing the rescue frequencies and decreasing the rate of depolymerization and the time of MTs spend in pausing. It has also been shown that EB1 promotes persistent growth by suppressing catastrophes in vitro (Komarova \textit{et al}, 2009). EB3 promotes MT growth and it is important during neurite formation by coordinating filamentous actin-MT interaction (Geraldo \textit{et al}, 2008) and for the organization of MTs during myotube differentiation (Straube & Merdes, 2007). EB3 binding to MTs is regulated via aurora A-mediated phosphorylation that leads to EB3-SIAH1 binding. SIAH1 is ubiquitin E3-ligase that targets EB3 for degradation (Ban \textit{et al}, 2009).

Another essential component of the +TIP family is cytoplasmic linker protein 170 (CLIP170). Deletion of CLIP170 results in very low MT rescue frequency (Komarova \textit{et al}, 2002). It is part of a subgroup of +TIP proteins named CAP-Gly proteins (cytoskeleton-associated protein Gly-rich) that interacts with the EEY/F domain of EB1 and tyrosinated α-tubulin (Peris \textit{et al}, 2006; Mishima \textit{et al}, 2007). CLIP170 was described as a phospho-sensitive MAP (Rickard & Kreis, 1991), and indeed, its binding to MTs is regulated by mTOR and AMP kinase PKA (Choi \textit{et al}, 2002; Nakano \textit{et al}, 2010; Lee \textit{et al}, 2010). CLASPs are CLIP-associated proteins that contribute to promote rescue events by recruiting tubulin dimers at the tips of MTs and by their interaction with EB1 (Akhmanova \textit{et al}, 2001; Al-Bassam \textit{et al}, 2010). The binding affinity of CLASPs to EB1 is decreased by GSK3β phosphorylation (Kumar \textit{et al}, 2009).

Another +TIP protein recruiting tubulin dimers at the plus-end and promoting MT growht is TOG protein, a MAP with MT-polymerizing activity (Brouhard \textit{et al}, 2008). There are some +TIP proteins specifically expressed in a cell-type dependent manner, such as Lis1 and its binding partner doublecortin in neurons. Absence of any of them leads to type I lissencephaly, a disease characterized by a smooth cerebral surface. Lis1 is responsible for targeting
CLIP170, dynein and dynactin (Tai et al, 2002) and reduces the catastrophe events of MTs (Sapir et al, 1997). Among the +TIP protein family, other tissue specific proteins have been described, such as neuron-navigator 1 in neurons (Martínez-López et al, 2005) or melanophilin in melanocytes (Wu et al, 2005).

Finally, there are proteins responsible for MT stabilization at the minus end of MTs. In interphase cells, MTs are organized in a MT-organizing center like centrosomes (MTOC), or loosely organized with a large number of free minus ends generated by release from MTOC, cytoplasmic assembly, or severing of pre-existing MTs (Dammerman et al, 2003). MT minus ends are stabilized due to the association of \( \gamma \)-tubulin with other proteins to form the \( \gamma \)-tubulin ring complex (\( \gamma \)-TuRC) that functions as a nucleation seed for MTs but also as stabilizing cap to avoid depolymerization. Pericentrin and ninein are \( \gamma \)-TuRC proteins that are localized at the centriole and anchor MTs to the centrosome (Abal et al, 2002; Lin et al, 2006). Pericentrin and ninein, relocate \( \gamma \)-TuRCs from the centrosome to the nuclear envelope during skeletal muscle differentiation (Bugnard et al, 2009). Ninein also anchors MT at epithelial cell junctions through an interaction with desmoplakin (Lechler & Fuchs, 2007). Nehzah is another MT stabilizing protein that anchors MT minus ends at junctions in epithelia cells independently of the \( \gamma \)-TuRC (Meng et al, 2008).

**MT-destabilizing proteins**

Several types of MT-binding proteins have been identified as MT destabilizers and are crucial for mitotic spindle organization. The MT-severing proteins katanin, spastin and fidgetin form a family of closely related enzymes that regulate the number and length of MTs through their severing activity. PTMs and other MT-binding proteins regulate MT activity and function (Zhang et al, 2007; Roll-Mecak & McNally, 2010). Members of the kinesin-13 family, such as KIF2A and MACK, have been shown to induce a conformational change of tubulin dimers that
triggers catastrophe events. They are thought to function by bending and peeling off individual protofilaments to generate rings (Ems-McClung & Walczak, 2010). Removal of Y from $\alpha$-tubulin leads to decreased binding of MCAK and KIF2A to MTs (Peris et al, 2009). In addition, +TIP protein EB1 recruits MCAK to the plus end of MTs and this interaction is regulated by aurora B-mediated phosphorylation (Andrews et al, 2004; Lan et al, 2004). Stathmin is a MT-destabilizing protein that, distinctly to MACK, binds to tubulin heterodimers, and curves them into a complex that cannot be incorporated into polymerized MTs (Belmont et al, 1996).

**Actin-MT crosstalk**

Cell motility, growth cone guidance, cell division, and wound healing require coordination between MT and actin filament dynamics (Rodríguez et al, 2003; de Forges et al, 2012). Different MT- and actin-binding proteins, cytolinkers and signaling cascades mediate the coordination between the two systems. Different +TIP proteins are responsible for anchoring MTs to the actin cell cortex. Adenomatous polyposis coli (APC) protein is transported by kinesin to the MT plus end where it stimulates their polymerization by binding to EB1 and anchors them at the cell cortex via mDia (a RhoA effector) and IQGAP1 (an Rac/Cdc42 effector) (Wen et al, 2004; Reilein & Nelson, 2005). CLASP proteins attach distal MTs ends to cortical sites and act independently of APC (Mimori-Kiyosue et al, 2004). Dynein and p150$^{\text{glued}}$ are the biggest subunits of the dynactin complex that is recruited by EB1 and CLIP170. Dynactin participates not only in centering the centrosome in interphase cells or positioning the mitotic spindle, but also pulls the ends of MTs to the cell cortex (Dujardin & Vallee, 2002; Gomes et al, 2005). Cytolinkers are also involved in crosslinking actin and MTs. ACF7 is a spectraplakin that accumulates at the tips of MTs through a direct interaction with EB1. Thereby it contributes to the control of cell polarity and migration, and stabilizes MTs (Kodama et al, 2003).
There are several non-exclusive hypotheses to describe actin-MT crosstalk, involving MT- and actin-binding proteins and signaling proteins. One possible mechanism to explain it might be a model where structural interaction of MTs and actin would be coupled with signaling cascades (Figure 5). MT breakage and depolymerization occurs as a result of actin retrograde flow (Waterman-Storer & Salmon, 1997). Depolymerizing MTs are thought to release the MT-bound Rho guanine nucleotide exchange factor (GEF)-H1 that is responsible for RhoA activation (Krendel et al., 2002). RhoA activation results in myosin phosphorylation leading to actin contraction and promoting the local stabilization and polymerization of MTs (Cook et al., 1998; Ren et al., 1999). MT polymerization activates Rac and Cdc42 leading to actin polymerization and protrusion formation (Waterman-Storer et al, 1999).

\[\text{Figure 5. Scheme representing actin-MT crosstalk.} \] Upon actin retrograde flow, MT breakage frees GEFs activating RhoA signaling pathway. This leads to actin contraction, MT polymerization and activation of Rac1, promoting new actin polymerization.

\textit{IF-MT crosstalk}

In contrast to actin, there is no direct evidence of IF-MT crosstalk. Although the cytolinker proteins plectin and BPAG1 have been identified as crosslinker between IFs and MTs (Svitkina
et al, 1996; Yang et al, 1999), no clear effects of IFs on MTs have been described yet. BPAG1 deficiency leads to MT destabilization independently of whether IFs are present or not.

1.1.2 Plectin

Plectin is one of the best characterized cytolinkers and the most versatile one (Wiche & Winter, 2011). Plectin was first purified in association with IFs (Pytela & Wiche, 1980). The discovery of plectin’s widespread expression in tissues and cells, especially at junctional sites, suggested a possible role of plectin in the formation of cell junctions and anchorage of cytoplasmic filaments (Wiche & Baker, 1982; Wiche et al, 1983). The crosslinking activity of plectin was first elucidated in studies on plectin’s molecular properties (Foisner & Wiche, 1987) and plectin-binding to IFs (Foisner et al, 1988). The plectin gene was first cloned and sequenced from rat (Wiche et al, 1991), and then from human (Liu et al, 1996). A detailed analysis of exon-intron organization of the rat gene locus was described in the following year by Elliott et al (1997), but it was not until 1999 that a detailed analysis of the murine plectin gene locus was published (Fuchs et al, 1999). A better understanding of plectin’s exact role came with the generation of plectin-deficient mice (Andrä et al, 1997), and the discovery that defects in the plectin gene cause epidermolysis bullosa simplex associated with muscular dystrophy (EBS-MD) (Gache et al, 1996; Smith et al, 1996), a severe skin blistering disease.

Molecular properties

Plectin is a large cytolinker protein of ~500 kDa that is composed of a central rod domain flanked by globular N-terminal head and C-terminal tail domains (Foisner & Wiche, 1987) (Figure 6). The central ~200 nm long α-helical coiled-coil rod structure is encoded by a single exon. It contains five subregions, each ~200 residues long, with a strict period of charged amino acids at 10.4 residues that might be involved in association between plectin molecules (Wiche et al, 1991). Indeed, it has recently been shown that dimeric rod domains can
associate laterally, forming stable polymers (Walko et al., 2011). The C-terminal globular domain (also encoded by a single exon) contains five PRD of the B type and one PRD of the C type. The IF-binding region of plectin consists of \( \sim 50 \) amino acid residues linking PRDs 5 and 6 (Nikolic et al., 1996). The globular C-terminal domain contains an additional binding site for integrin \( \beta 4 \) (Rezniczek et al., 1998) and a unique phosphorylation site for cdk1 kinase (Malecz et al., 1996).

**Isoform diversity**

Plectin is the most versatile cytolinker protein mainly due to its unique and complex exon-intron organization. Alternative splicing of the 5'-end of plectin's gene gives rise to at least 16 different transcripts (Figure 7): Eleven alternative first exons are spliced into a common exon 2. Eight of them are first coding exons (1-1g), and three of them are non-coding (1h, 1i, 1j). Three additional non-coding exons (E0, E0a and E-1) splice into exon 1c, and another two (2\( \alpha \) and 3\( \alpha \)) are optionally spliced within the exons encoding the ABD (Fuchs et al., 1999). This broad variety of sequences determines the cellular targeting of the different isoforms (Rezniczeck et al., 1997).
Plectin isoforms show preferential binding to an assortment of cellular structures, including hemidesmosomes (HDs), focal adhesions (FAs), costameres, mitochondria, MTs, nuclear/endoplasmic reticulum membranes, and Z-disks. Being expressed in the various cell types and tissue in different combinations and proportions, plectin isoforms connect the cellular structures targeted with the corresponding cell-type-specific IF network. By anchoring IFs at distinct cellular structures plectin isoforms control IF cytoarchitecture and thereby basic and other cell type-specific features of cells (Wiche & Winter, 2011).

**Figure 7. Transcripts generated by alternative splicing at the 5´-end of the plectin gene.** Eleven alternative exons splice into exon 2. Eight of them are first coding exons (1-1g), three of them are non-coding (1h-1j). In addition, there are three non-coding exons that splice into exon 1c (E0, E0a and E-1). Two optionally spliced exons (2α and 3α) are indicated. Black boxes indicate coding regions and white boxes designate non-coding sequences (Fuchs et al, 1999).

**Plectin in connective tissue**

First immunolocalization studies revealed plectin throughout the cytoplasm of immortalized fibroblast cell cultures (Wiche & Baker, 1982), and in cells of mesenchymal origin in various rat tissues examined (stomach, kidney, small intestine, liver, uterus and urinary bladder) (Wiche et al, 1983). However, most of the features known about connective tissue plectin came from studies of plectin-deficient fibroblasts. One of plectin’s major isoforms in fibroblasts is plectin 1
(P1). P1\textsuperscript{-/-} fibroblasts show impaired migration and alterations of the actin cytoskeleton (Abrahamsberg \textit{et al}, 2005), reminiscent of plectin-null (P0) fibroblasts (Andrä \textit{et al}, 1998). Another isoform of plectin prominently expressed in fibroblasts is plectin 1b (P1b). P1b targets mitochondria via its N-terminal part and anchors them to IF networks via its C terminus. P1b-deficient fibroblasts show abnormal mitochondrial shape and distribution (Winter \textit{et al}, 2008).

Plectin has also been shown to have an influence on vimentin IF cytoarchitecture of fibroblasts. Plectin is associated with vimentin from the early stages of filament assembly, is required for the formation of IFs and their directional growth towards the cell periphery, and plectin-deficient fibroblasts undergo faster mitosis than their wild-type (wt) counterparts (Spurny \textit{et al}, 2008). Moreover, plectin participates in the regulation of the de novo IF formation (Burgstaller \textit{et al}, 2010). Plectin 1f (P1f), one of the major isoforms expressed in fibroblasts is a component of FA-evolved fibrillar adhesions (FbAs) that are centrally located. P1f recruits vimentin intermediates (squiggles) to these sites and stabilizes FbAs, thus acting as a nucleation center for filament formation. As a result, vimentin filaments form a cage around the nucleus of the cell that in plectin-deficient fibroblast is distorted (Figure 8). Plectin deficiency in fibroblasts results also in increased stability of actin filaments, compromised signaling (decreased Src and FAK activities) and decreased migration potential (Andrä \textit{et al}, 1998; Osmanagic-Myers \textit{et al}, 2004; Osmanagic-Myers \textit{et al}, 2006; Gregor \textit{et al}, 2006).

\textit{Plectin in muscle}

Plectin is expressed in the three major types of muscle, localizing along the sarcolemma in smooth muscle cells (Wiche \textit{et al}, 1983; Tanaka \textit{et al}, 2001), at intercalated disks and Z-disks of cardiac muscle (Wiche \textit{et al}, 1983; Zernig & Wiche, 1985; Konieczny \textit{et al}, 2008), and at the sarcolemma and Z-disks in skeletal muscle (Wiche \textit{et al}, 1983; Konieczny \textit{et al}, 2008). However, the influence of plectin on IF network organization becomes most noticeable in
skeletal muscle. In this type of muscle four major isoforms of plectin are expressed, namely P1, P1b, P1d, and P1f. P1d is associated with Z-disks, P1f with the dystrophin-glycoprotein complex at the sarcolemma, P1b with mitochondria, and P1 with the outer nuclear/sarcoplas-

Figure 8. Simplified representation of plectin’s function as organizer of fibroblast cytoarchitecture. Vimentin filaments form a cage around the nucleus through their P1f-mediated attachment to FbAs and centrally located FAs in polarized fibroblasts. Anterogradely transported (red arrow) vimentin filament intermediates (“squiggles”) are “captured” by FA-associated P1f and by tandem-fusion will extend the filaments forming the centrally located cage-like network. Plectin-deficient cells are rounded, not polarized and their vimentin network extends to the outermost boundary of the cell (Burgstaller et al., 2010).

mic reticular membrane system (Rezniczek et al., 2007; Konieczny et al., 2008) (Figure 9). P1d and P1f are crucial for linking the contractile apparatus via desmin IFs to the sarcolemmal costameric protein skeleton. Disruption of either P1f or P1d leads to the loss of muscle fiber integrity (Konieczny et al., 2008). Albeit already newborn plectin-deficient (null) mice revealed abnormalities in skeletal and cardiac muscle, they were unusable for more detailed studies because they die within 2-3 days after birth (Andrä et al., 1997). Hence, the phenotypic analysis of plectin-deficient muscle had to be carried out with conditional (MCK-Cre) striated muscle restricted knock out and isoform-specific knock out mice (Konieczny et al., 2008).
Figure 9. Model depicting the role of plectin in skeletal muscle fibers. The four major plectin isoforms expressed in muscle, P1, P1b, P1d, and P1f anchor desmin IFs to the outer nuclear/SR membrane system, mitochondria, Z-disks, and the sarcolemmal dystrophin-glycoprotein complex, respectively. Plectin-deficiency causes aggregation of desmin IFs and misalignment of Z-disks (Wiche & Winter, 2011).

Epithelial plectin

Early studies on plectin distribution revealed plectin expression in practically all mammalian tissues. Antibodies to plectin were found to prominently decorate the basal surface membrane of stratified and simple epithelial cell layers (Wiche et al, 1983). In basal keratinocytes, keratin IFs are linked to hemidesmosomal integrin β4 via plectin isoform 1a (P1a) (Andrä et al, 2003; Kostan et al, 2009). Plectin-null (P0) and conditional (K5-Cre) plectin knockout mice, both dying early after birth, show blistering on their extremities and on the oral epithelium after initial nursing (Andrä et al, 1997; Ackerl et al, 2007). Although HDs are not structurally affected in P0
keratinocytes, they are reduced in number and their mechanical stability is compromised. More recently it was shown that the selective degradation of HD-associated P1a, by proteases activated specifically in keratinocytes, results in reduced numbers and dysfunction of HDs (Walko et al., 2011). Selective proteolytic degradation of P1a might be required also for epithelial differentiation (Kostan et al., 2009).

A second major isoform of keratinocytes, P1c has been shown to partially colocalize with MTs and cosediment with pre-assembled MTs (see below). Reminiscent of IF network alterations observed in plectin-deficient fibroblasts, keratinocytes lacking plectin display keratin networks where filaments appear to be bundled and extending to the periphery. In contrast, wt cells exhibit keratin networks that are more delicate and juxtapositioned to the nucleus, leaving a filament-free ring-shaped zone at the cell margins (Osmanagic-Myers et al., 2006). However, contrary to plectin-null fibroblasts, plectin-deficient keratinocytes show increased migration rates, corresponding to elevated basal activities of mitogen-activated protein (MAP) kinase Erk1/2 and of the membrane-associated upstream protein kinases c-Src and PKCδ (Osmanagic-Myers et al., 2006). Plectin deficiency leads to the skin blistering disease EBS (Gache et al., 1996; Smith et al., 1996) and EBS-Ogna (see chapter below).

**Plectin in the nervous system**

Little is known about the role of plectin in the nervous system. As shown by analyses of plectin transcripts, P1c is a major isoform not only of epidermal cells but also of neural cells (Figure 10). This is not unexpected considering that epidermal and neural cells share a common developmental origin, the ectoderm.

Thanks to P1c-specific antibodies and the availability of P1c-deficient (P1c<sup>-/-</sup>) mice to be used as optimal negative control, P1c was found to be expressed late in development and
associated with postsynaptic dendrites of central nervous system neurons, spinal cord motor neurons, sciatic nerve axons, and Schwann cells. In central nervous system, P1c was preponderantly expressed in the hippocampus. P1c deficiency affects the behavior of mice as shown by their reduced exploratory activity and voluntary wheel running (Michael Zörer, PhD thesis). In addition, lack of P1c leads to reduced nerve conduction velocity in motor neurons combined with a reduction in motor neuron calibers (Fuchs et al, 2009).

**Plectin and disease**

Most of the plectin mutations reported lead to epidermolysis bullosa simplex associated with muscular dystrophy (EBS-MD) (Gache et al, 1996; Smith et al, 1996). EBS-MD patients suffer from severe skin and mucous membranes blistering and late-onset muscular dystrophy. In addition, other plectin mutations were reported to cause EBS-MD with myasthenic syndrome (EBS-MD-MyS) (Banwell et al, 1999; Forrest et al, 2010; Selcen et al, 2011), limb-girdle muscular dystrophy type 2Q (LGMD2Q) (Gundesli et al, 2010), and EBS with pyloric atresia (EBS-PA) (Rezniczek et al, 2010; Natsuga et al 2010). Moreover, a dominant mutation in plectin has been reported to cause the rare skin blistering disease EBS-Ogna, which was originally identified in a Norwegian kindred and in an unrelated German family (Koss-Harnes et al, 2010).
al, 2002) (Figure 11). Except for EBS-Ogna and most of the EBS-PA cases, plectinopathies are characterized by pathological desmin-positive aggregates, myofibril degeneration, and mitochondrial abnormalities (Schröeder & Schoser, 2009; Winter et al, 2012). Insufficient amounts of plectin have been found to promote the aggregation of GFAP, an astrocyte-specific IF protein, in a rare but fatal neurological disorder known as Alexander disease characterized by cytoplasmic inclusion bodies, which contain GFAP, plectin and other components. Alexander disease patients suffer from macrocephaly and episodes of severe seizures, leading to progressive disability or early death (Tian et al, 2006).

**Figure 11. Scheme representing positional mapping of plectin mutations reported up to 2011.** Plectin mutations responsible for EBS-MD are depicted in black, EBS-PA in blue, EBS-OGNA in green, EBS-MD-MyS in red, and LGMD2Q in purple (Winter et al, 2012).

**Plectin and MTs**

The role of plectin as cytoskeletal networking element involving MTs became undisputed when G. Borisy’s group published images visualizing plectin molecules as linkers between MTs and...
vimentin IFs (Svitkina et al., 1996). Additionally, unpublished data from our group show P1c immunofluorescence microscopy to partially colocalize with MTs (Figure 12A), and to cosediment with pre-assembled MTs (Figure 12B) (G. Walko, unpublished; Andrä et al., 2003). Albeit the C-terminal tail contains GSR repeats that were reported to be a MT-binding site, it was shown by G. Walko (PhD thesis) that the C-terminal domain of plectin is not able to associate with MTs.

Figure 12. P1c interacts with MTs. A) Immortalized wt and P0 keratinocytes were immunolabeled using anti-α-tubulin and anti-P1c isoform-specific antibodies. Note even distribution of P1c (upper panels), and speckled decoration of MTs (lower panels). Scale bars, 50 µm (upper row); 10 µm (lower row). B) Wt keratinocyte cell homogenates were separated into high-speed supernatant (HS-SN) and pellet (HS-P) fractions, and the HS-SN fraction was incubated with (+MT) or without (-MT) taxol-stabilized MTs and centrifuged. Pellet (P) and supernatant (SN) fractions were analyzed by immunoblotting using isoform-specific antibodies. Note a fraction (~30%) of endogenous P1c cosedimented with pre-assembled MTs (G. Walko, unpublished).

The ABD of plectin is followed by a plakin domain containing spectrin repeats and a SH3 domain. Interestingly, preliminary data from our laboratory demonstrated that the SH3 domain present within the plakin domain is able to bind to high molecular weight (HMW)-MAPs and antagonize their MT-stabilizing effect (unpublished). At the molecular level, previous solid-phase binding studies had revealed binding of full-length plectin isolated from rat glioma C6
cells to HMW-MAPs purified from brain (Herrmann & Wiche, 1987). Afterwards, the laboratory strategy was to identify which molecular domain(s) of plectin were involved in MAP-binding. Using various fragments of P1c, expressed as fusion proteins with N-terminal GST tags, it was found that those containing the central part of plectin’s N-terminal plakin domain preceding the rod, showed binding to at least one of the HMW-MAPs. The minimal fragment showing MAP-binding, p20-21, corresponded to plectin’s putative SH3 domain (Ortega et al., 2011), raising the possibility that plectin bound to MAPs via this domain. A similar type of interaction has previously been shown for tau, another major MAP, which binds to the SH3 domain of the non-receptor tyrosine kinases Fyn and Src (Lee et al., 1998). P1c-MAP binding has further been demonstrated by co-sedimentation of HMW-MAPs with P1c immunoprecipitated from mouse brain lysates using anti-P1c isoform-specific antibodies (Figure 13C; E. Mihailovska, unpublished data).

To better understand the molecular mechanism involved in MT destabilization through plectin, the effects of plectin’s SH3 domain on MAP-promoted in vitro assembly of tubulin into MTs has previously been studied in this laboratory (L. Janda, unpublished data). For these assays MAP-free tubulin was used, along with fragment p20-21, and recombinant MAP2c. Upon incubation of MAP-free tubulin with MAP2c, under conditions where tubulin itself polymerized only poorly (if at all), ~65% of the tubulin was found to form polymers in the absence of fragment p20-21 (Figure 13A, lane 4). When fragment p20-21 at increasing concentrations was mixed with MAP2c prior to induction of MT formation, increasing amounts of MAP2c were found in the soluble fractions, indicating reduced MT-binding of MAP2c in the presence of fragment p20-21. Ultimately, reduced MAP2c-binding to MTs led to a shift of tubulin from the insoluble to the soluble fraction, indicating an inhibition of MAP2c-promoted MT assembly. Also previous results of the group showed that fragment p20-21 could compete with MTs for tau-binding under conditions where the polymeric state of MTs was maintained.
Figure 13. Plectin’s SH3 domain (fragment p20-21) compromises recombinant MAP-MT interaction. 

A) MTs were assembled in vitro from purified samples of tubulin and MAP2c in the presence of fragment p20-21 (at concentrations indicated) and sedimented by centrifugation. Resulting pellets (p, containing polymerized MTs and MT-bound MAP2c), and supernatants (s, containing soluble tubulin and unbound MAP2c), and p20-21 were analyzed by SDS-PAGE. Coomassie-stained gel bands corresponding to MAP2c in supernatant (white bars in bar diagram) and tubulin in pellet fractions (grey bars) were quantified. Error bars, ± SD (n=5). (L. Janda, unpublished data) 

B) The detachment of endogenous tau from MTs contained in brain lysates was measured by SDS-PAGE of the sedimented MT fractions after incubation of lysates with p20-21 at concentrations indicated. Protein bands corresponding to MT-bound tau, tubulin (representative of MT polymers), and NF protein M (NF-M, loading control) are shown. Quantitation (graph) as in (A). Error bars, ± SEM (n=5). (E. Mihailovska, unpublished data). 

C) Co-immunoprecipitation of P1c and HMW-MAPs from brain lysates. Note MAPs-P1c coprecipitation when anti-P1c antibodies, but not IgGs alone, were used (n=3) (E. Mihailovska, unpublished data).
(without disassembly of the polymer) (E. Mihailovska, unpublished data). For this, increasing concentrations of fragment p20-21 were incubated with MTs contained in brain lysates under conditions that were favorable for their polymerized state even after removal of MAPs. The analysis of polymeric MTs (i.e. sedimentable by high–speed centrifugation) and their co-assembling (bound) proteins by immunoblotting clearly revealed a decrease in MT-bound tau protein with increasing concentrations of fragment p20-21 (Figure 13B). These data suggested that plectin’s SH3 domain could detach MAPs from MTs.
2 AIMS OF THE THESIS

A number of observations made in previous studies, including the targeting of overexpressed P1c to MTs, the partial colocalization of P1c with MTs in cultured cells, and the in vitro binding of neural plectin to MAPs, pointed towards a potential interaction of plectin with MTs. Moreover, ACF7 and BPAG1, other members of the plakin cytolinker protein family, have been shown to be involved in the regulation of MT dynamics, both acting as MT stabilizers. Based on these previous observations and additional unpublished data from this laboratory, one of the major goals of my thesis was to analyze whether plectin plays a role in regulating MT dynamics. Of particular interest in this context was the question whether plectin isoform P1c antagonizes MAP-mediated MT stabilization, and if so, what was the underlying molecular mechanism.

A second major goal of my thesis was to search for a mechanistic link between plectin-related changes in MT dynamics and alterations in MT-dependent processes of physiological significance. Since P1c is a major isoform in keratinocytes and neurons, and the preliminary data obtained in the laboratory showed that P1c colocalizes with MTs, the most convenient system for this investigation was to use keratinocytes and neurons isolated from P1c-deficient mice. To analyze whether P1c-dependent MT organization could alter basic cellular functions in keratinocytes, processes including glucose uptake, cell division and growth, FA turnover, shape and polarized migration of cells were to be studied. P1c-deficient primary dorsal root ganglia (DRG) neurons should be assessed as a system in which to study the consequences of altered MT-dynamics for neurite cell functions, such as neurite outgrowth and branching, synaptic and other vesicle transport along neurites, and neuronal membrane depolarization.
3 RESULTS

This section is subdivided into two parts. Both address plectin isoform P1c’s potential role as a regulator of MT dynamics and the consequences such regulation might have for MT-dependent cellular functions. While part I addresses the keratinocyte cell system, part II is focused on neuronal cells.

Part I - Plectin 1c in keratinocytes

Preliminary results obtained in previous studies of this group showed partial colocalization of P1c with MTs in cultured keratinocytes (see Introduction). However, the biological significance of P1c’s presumptive interaction with MTs or its effects on the dynamic behavior of MTs had not been investigated.

3.1.1 Lack of P1c in keratinocytes leads to increased stability of MTs

Proteins that bind along MTs often alter the mechanical properties of the polymer, either by stabilizing (e.g. MAPs, ACF7/MACF) or destabilizing (e.g. Op18/stathmin, SCG10) (Conde & Cáceres, 2009). To investigate whether P1c shares such properties and to analyze its specific role in MT network organization, I chose to study primary P1c-deficient keratinocytes (derived from P1c isoform-specific knockout mice, Fuchs et al, 2009) in comparison to P0 keratinocytes and wt keratinocytes. This combination of cell types represents an optimal system to distinguish between isoform P1c-specific and other plectin isoform deficiency-related phenotypes. First, I investigated whether the absence of P1c affected the stability of MTs by tracing and quantifying MTs that remained polymerized after treatment of P1c<sup>−/−</sup> and wt primary keratinocytes with low doses of the MT-depolymerizing drug nocodazole (for details see chapter 5, Material and methods). Unexpectedly, I found MTs in P1c<sup>−/−</sup> keratinocytes to be
more, rather than less stable compared to those in wt cells (Figure 14), and a similar phenomenon was observed in P0 keratinocytes (Figure 14).

As stable populations of MTs are usually enriched in post-translationally acetylated α-tubulin (Piperno et al., 1987; Bulinski et al., 1988), I investigated the distribution and quantified the levels of acetylated tubulin present in MTs of primary P1c<sup>+/−</sup> keratinocytes and compared them to those of wt and P0 cells. Subjecting cells to double immunofluorescence microscopy, using anti-tubulin antibodies not discriminating between the modified (acetylated) and unmodified forms of the protein and antibodies that were specific for the acetylated version, I found that in wt keratinocytes only the central part of the cells was stained for acetylated tubulin, whereas in both mutant cell types (P1c<sup>+/−</sup> and P0) acetylated MTs were present at the cell center as well as at the periphery. A quantification of acetylated tubulin-positive areas (in

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**Figure 14. P1c deficiency affects drug resistance of keratinocyte MTs.** A) Nocodazole-treated primary keratinocytes (wt, P1c<sup>+/−</sup>, and P0) were immunolabeled using antibodies to α-tubulin. Intact MTs remaining after nocodazole treatment were traced (MT tracing). Note the abundant presence of intact MTs in nocodazole-treated P1c<sup>−/−</sup> and P0 cells (MT tracing, lower row) in contrast to wt keratinocytes. Scale bar, 20 µm. B) Statistical evaluation of data shown in A by one-way ANOVA-test and post-hoc Tukey correction compared to wt values (n=5; ~20 cells/experiment) *P<0.05; **P<0.01; ***P<0.001. Error bars, ± 95% CI.
pixels) versus unmodified (total) tubulin-positive areas (for details see Material and methods) showed the acetylated form to be ~3-fold increased in plectin-deficient (P1c⁻/⁻ and P0) keratinocytes compared to wt cells (Figure 15A). Increased acetylation of MTs in plectin-deficient keratinocytes was confirmed by immunoblotting analysis of tubulin present in extracts from proliferating cells (Figure 15B). Overall, these data were fully consistent with the observed higher nocodazole-resistance of MTs from mutant cells (see Figure 14).

![Figure 15](image)

**Figure 15. P1c deficiency affects acetylation of keratinocyte MTs.** A) Indicated cell types were analyzed by immunofluorescence microscopy using rat monoclonal anti-tubulin (red) and mouse monoclonal anti-acetylated tubulin (green) antibodies. Note more prominent signal to acetylated tubulin antibodies in P1c⁻/⁻ and P0 keratinocytes compared to wt cells. Scale bar, 20 µm. B) Statistical evaluations were done by a one-way ANOVA-test and post-hoc Tukey correction compared to wt values (n=5; ~20 cells/experiment). *P<0.05; **P<0.01; ***P<0.001. Error bars, ± 95% CI. C) Quantitation (immunoblotting) of acetylated tubulin present in cell lysates from immortalized wt and P0 keratinocytes prior to (0.05 mM Ca²⁺) and after exposure (3h) to 1.8 mM Ca²⁺. Numbers, quantified relative levels of acetylated tubulin.

### 3.1.2 Reversal of MT stabilization in P1c⁻/⁻ keratinocytes requires full-length P1c

To assess whether MT stabilization in mutant cells was directly linked to P1c deficiency, we performed rescue experiments where primary P1c⁻/⁻ keratinocytes were transiently transfected...
with cDNA expression plasmids encoding EGFP fusion proteins with either full-length P1c-EGFP (P1c-EGFP), or one of two N-terminal fragments of P1c; a shorter one (P1c-8-EGFP) comprising the N-terminal ABD preceded by the isoform-specific exon 1c-encoded sequence, and a similar longer one (P1c-30-EGFP) extending to plectin’s central $\alpha$ helical coiled-coil rod domain. Upon forced expression, P1c-EGFP was found accumulated at filamentous structures around the nucleus but was visualized also in a dotted pattern throughout the cytoplasm (Figure 16 A, upper panels). In contrast, P1c-8-EGFP colocalized with stress fibers (Figure 16A, middle panels) as has previously been described (Rezniczek et al., 2003). P1c-30-EGFP

![Figure 16.](image)

**Figure 16.** P1c variants lacking the IF-binding domain fail to rescue nocodazole sensitivity in P1c-deficient keratinocytes. A) Nocodazole-treated primary P1c$^{+/+}$ keratinocytes transfected with full-length P1c-EGFP (upper row), P1c8-EGFP (middle row), and P1c-30-EGFP (lower row) were immunolabeled using antibodies to $\alpha$-tubulin (tubulin). Drug-resistant MTs remaining in cells were traced (MT tracing). Scale bar, 15 µm. B) Bar graph shows quantification of MT length normalized to total cell area; red and black broken lines indicate corresponding values measured for wt and P1c$^{+/+}$ cells, respectively (see Fig. 14) (n=3; 6 cells/experiment). *P<0.05; **P<0.01; ***P<0.001. Error bars, ± 95% CI.
was found associated with filamentous structures throughout the cytoplasm (Figure 16A, lower panels). When MTs in transfected cells were exposed to nocodazole to assess their resistance towards drug-induced disassembly, we found that full-length P1c expression not only led to a reversal of the phenotype but even to an overshooting effect (Figure 16B, bar graph), i.e. destabilization of MTs to levels below that of wt cells, probably due to overexpression of P1c to levels higher than normal. Contrary to full-length P1c, expression of the truncated versions P1c-8 and P1c-30 did not effect a statistically significant reduction of MT stability, as revealed by the abundant presence of MTs remaining after nocodazole treatment (Figure 16A, middle and lower panels). Similarly, when MT acetylation was measured upon transfection of cells, only full-length P1c led to a reduction of acetylated MT signals, corresponding to a partial restoration of the phenotype (Figure 17A).

These data strongly suggested that P1c indeed was destabilizing rather than stabilizing MTs, contrary to what had been observed with other cytolinker protein family members, such as ACF7/MACF and BPAG1 (Kodama et al., 2003; Yang et al., 1999). Moreover, the results obtained with truncated forms of P1c suggested that the full-length protein, or at least a version of it containing the IF-binding site, was required in order to reset MT stability to the less stable state of wt cells.

3.1.3 P1c deficiency affects MT dynamics

To explore whether the increased stability of MTs observed in plectin-deficient cells was reflected in their dynamic behavior, we transfected keratinocytes with EGFP-tagged tubulin (Krylyshkina et al., 2002), enabling the visualization of rapidly shortening (catastrophe) and regrowing (rescue) MTs at the cell margins. Because of limited transfection rates obtained with primary keratinocytes, for these experiments immortalized wt and P0 keratinocytes were used. In addition to measuring catastrophe
Figure 17. P1c variants lacking the IF-binding domain fail to reverse increased MT acetylation in P1c-deficient keratinocytes. A) The proportion of acetylated tubulin present in primary P1c<sup>-/-</sup> keratinocytes expressing full-length or truncated versions of P1c was determined by immunofluorescence microscopy. B) Channels: red, tubulin; green, acetylated tubulin; blue, EGFP. Bar graph represents statistical evaluations as in Figure 16 B. Scale bars, 15 µm (upper row); 10 µm, middle and lower rows.

and rescue frequencies of MTs, we quantified the proportions of MTs that were growing perpendicular towards the membrane versus those bending or sliding along the membrane (compare to Kodama et al., 2003). The results revealed a nearly identical rescue frequency in wt and P0 cells, but a ~2.5 fold decrease of catastrophe frequency in P0 compared to wt cells.
(Figure 18 A and B); this was consistent with the higher stability of MTs towards nocodazole, as observed in mutant cells (see Figure 14). Interestingly, while in wt cells most of the MTs were growing perpendicularly towards the membrane and upon reaching it underwent rapid disassembly (catastrophe) (Figure 18A), P0 cells showed a higher percentage of MTs that upon reaching the membrane were bending or sliding along it instead of disassembling (Figure 18C).

![Figure 18. P1c affects dynamic properties of MTs. A) Images taken from a time-lapse recording of immortalized wt and P0 keratinocytes transfected with EGFP-tubulin and outlining (MT-tracing) of representative MTs. MTs outlined in red and black denote two examples of shrinking MTs in a wt cell, while lines in blue represent MTs in a P0 cell that failed to undergo catastrophe after reaching the membrane. Scale bar, 5 µm. B) Bar graphs showing analyses of rescue and catastrophe frequencies (n=3; >10 cells/experiment). *P<0.05. Error bars, ± 95% CI. C) Graph represents proportions (%) of MTs in wt versus P0 cells that were growing perpendicularly towards the plasma membrane (without bending), or bending and/or sliding along the membrane (n=3; 10 cells/experiment). ***P<0.001. Error bars, ± 95% CI.]

To investigate the growth parameters of MTs in more detail, we performed time-lapse microscopy of MTs labeled with GFP-tagged +TIP protein EB1 (Stepanova et al, 2003). In
accordance with the lower MT catastrophe frequency of P0 cells (Figure 18B), the number of EB1 comets quantified in GFP-EB1-transfected P0 keratinocytes was ~2.7 times higher than in wt cells (Figure 19).

Moreover, P0 MTs failed to undergo catastrophe after reaching the cell margins and continued to grow and bend at the periphery, contrary to wt MTs which paused upon reaching the cell membrane before disappearing (Fig. 20). Furthermore, by tracing single EB1 comets, MT tips were found to grow faster in P0 compared to wt cells (Fig. 21).

3.1.4 Glucose uptake is increased in P1c-deficient keratinocytes

To assess the physiological significance of plectin-modulated MT regulation I measured glucose uptake of mutant (P1c^{-} and P0) and wt keratinocytes. First, because glucose transport involving glucose transporter recycling is a MT-dependent process (Fletcher et al, 2000; Zaid et al, 2008), and second, because there is evidence for plectin's implication in wound healing (Abrahamsberg et al, 2005). Compromised wound healing and reduced skin proliferation are some of the severe complications associated with impaired glucose utilization in hyperglycemia and diabetes. When keratinocytes were incubated with 2-NBDG, a
Figure 20. Sequence of single frame images taken from a time-lapse recording of GFP-EB1-expressing wt and P0 keratinocytes. Coloured lines mark trajectories of different EB1 comets (EB1-tracing). Green broken line marks cell margins. Coloured EB1 traces in wt images (upper rows) represent individual MTs that grow towards the periphery and disappear after reaching the margin; outlines trajectories in P0 cells mark MTs that continue to grow upon reaching the membrane and either meander (red line) or bend and grow parallel to the membrane (blue and black lines). Scale bar, 5 µm.

Figure 21 Time-lapse images of representative single EB1 comets in wt and P0 cells. A) Note faster growth of EB1 comets (corresponding to growing tips of MTs) in P0 cell (red arrowheads) compared to its wt counterpart (white arrowheads) during the same time period. B) Graph shows analysis of MT growth rates (n=3; ~20 cells/experiment). ***P<0.001. Error bars, ± 95% CI. Scale bar, ~3 µm.
fluorescent derivative of D-glucose (Yamada et al., 2000), and fluorescence intensities measured by confocal microscopy, a significantly higher uptake of 2-NBDG by P1c−/− (~2.2-fold) and P0 (~1.5-fold), compared to wt cells was observed (Figure 22). Control experiments performed in the presence of the MT assembly blockers nocodazole (Figure 22) revealed a dramatic reduction in fluorescence intensity, confirming that 2-NBDG uptake was MT-dependent. Thus, a more efficient (MT-dependent) vesicular delivery of glucose transporters could readily explain the increase in glucose uptake shown by P1c−/− and P0 keratinocytes. Indeed, when cells were immunolabeled for GLUT1 and signal intensities quantitated (for details see Material and methods), a ~4.5 times higher level of glucose transporters was measured for P0 compared to wt keratinocytes (Figure 23). Glucose uptake

![Figure 22. Alterations of glucose uptake in plectin-deficient keratinocytes. A) Primary wt, P1c−/−, P0, and nocodazole-treated P0 keratinocytes were incubated with 2-NBDG and fluorescence intensity of cells was measured. Pictures obtained by immunofluorescence microscopy of 2-NBDG-treated keratinocytes show higher fluorescence intensity in P1c−/− and P0 keratinocytes than wt or nocodazole-treated P0 keratinocytes. Scale bar, 1 mm. B) Data were evaluated by a one-way ANOVA-test and post-hoc Tukey correction compared to wt values (n=3; 200 cells/experiment) ***P<0.001. Error bars, ± 95% CI.](image)
Figure 23. GLUT1 expression on keratinocytes. A) Keratinocytes were immunolabeled using antibodies to GLUT1. Note that GLUT1 staining remains in the perinuclear region in wt cells while in P0 cells the staining is spread over the cell. Scale bar, 10 µm. B) GLUT1 channel counts (normalized to total cell area) were subjected to a Student’s t-test. (n=3; ~10 cells/experiment) **P<0.01. Error bars, ± 95% CI.

might be additionally favored by the keratin network architecture of plectin-deficient keratinocytes, where keratin filaments typically extend all the way to the cell periphery instead of leaving a peripheral ring-shaped, filament-free zone characteristic of wt cells (Osmanagic-Myers, et al 2006). As recently suggested, keratins may participate in the regulation of cellular glucose uptake by properly localizing the glucose transporters GLUT1 and GLUT3, thereby increasing their density at the apical membrane (Vijayaraj et al, 2009).

3.1.5 Lack of P1c leads to aberrant mitotic spindles and diminished growth rate

As MTs are key players in cell division, I investigated whether their stabilization in P1c−/− keratinocytes affects the division process of these cells. Interestingly, the immunostaining of primary P1c−/− keratinocyte cultures using antibodies to α-tubulin, revealed a significant number of aberrant, multipolar as well as asymmetric, mitotic spindles (Figure 24A). A comparative quantitative analysis of aberrant mitotic spindles observed in primary P1c−/− and wt keratinocytes, showed an unexpectedly high fraction (40%) of P1c−/− mitotic cells to display
Figure 24. Alterations in mitotic spindle formation and cellular growth rate in plectin 1c-deficient keratinocytes. **A)** Spindle apparatuses of dividing cells were visualized by immunolabeling using antibodies to α-tubulin. Images display representative pictures of multipolar spindles (upper row) and asymmetric spindles (lower row) found in dividing P1c−/− keratinocytes. Scale bar, 10 µm. Abnormalities were statistically evaluated by a Student’s t-test **P<0.01. Error bars, ± 95% CI. (n=4; ~150 cells/experiment). **B)** Line graph represents growth rate of primary keratinocytes measured during 96 h (n=3). Error bars, ± SEM.

Multipolar or asymmetric spindles, compared to only ~3% of mitotic wt cells (Figure 24A, bar graph). The impairment of normal mitotic spindle formation correlated with a substantial delay in the growth rate of primary P1c-deficient compared to wt keratinocytes over the first 96 h after seeding of the cells (Figure 24B). Alterations in growth rate have already been observed in P0 fibroblasts (Spurny et al, 2008). Furthermore, plectin has been identified as a component of the mitotic spindle in several proteomic studies focused on the characterization of spindle-associated proteins (Mack & Compton, 2001; Sauer et al, 2005; Gache et al, 2010) and on the spindle phosphoproteome (Dephoure et al, 2008; Malik et al, 2009).
3.1.6 P1c-deficiency affects shape and directional migration of cells

It has previously been shown that primary as well as immortalized P0 keratinocytes migrate with higher velocities than their wt counterparts (Osmanagic-Myers, et al 2006), and it is known that polarized migration of mammalian cells requires stabilized MTs at the cell cortex (Watanabe et al, 2005; Kaverina & Straube, 2011). These observations prompted me to investigate a possible link between P1c-modulated MT dynamics and cellular features determining the shape and polarity of cells. Seeded at low densities to enable single cell monitoring, primary P1c-/- keratinocytes showed a reduction in size (cell area) but an increase in their perimeter compared to wt cells (Figure 25, a-b); P0 cells, too, showed a reduction in size, but hardly a perimeter change (Figure 25, a-b). This suggested that primary P1c-/- keratinocytes were displaying more protrusions than wt and P0 cells. Accordingly, a statistical analysis of calculated shape factors (SF = $4\pi \times \text{area} / \text{perimeter}^2$) and aspect ratios (AR = shortest diameter/largest diameter) for each cell type revealed wt and P0 keratinocytes to be similar in shape, but P1c-/- cells to have more protrusions and to be much more elongated (Figure 25 c-d, and Figure 26).

Time-lapse videos of single cells showed that primary P1c-/- keratinocytes migrated faster (~1 µm/min) compared to wt cells (~0.8 µm/min), albeit still slower than P0 cells (~1.2 µm/min) (Figure 25, e). One explanation why P1c-/- cells did not reach the velocity of P0 cells could have been that they were still expressing isoform P1a and thus were able to form intact HDs that were pinning them down, contrary to P0 cells that were lacking both isoforms. Interestingly, P1c-/- keratinocytes showed also a drastic loss of directional migration potential compared to wt and P0 cells (Figure 26, migration tracks, and Figure 25, f). Promotion of migration through stabilized MTs and simultaneous retention of cells via P1a anchorage, leading to the extreme elongation of cells and loss of directional migration potential, could again be the reason why P1c-deficient, but not P0 cells showed this phenotype.
3.1.7 Lack of P1c causes alterations in FA dynamics

To explore possible mechanisms underlying the higher migration rate of plectin-deficient keratinocytes, I examined whether the greater stability of MTs in the mutant cells could compromise the stability of FAs. MT targeting events to FAs have been shown to induce their dissociation by delivering FA-relaxing factors and promoting FA turnover (for review see Kaverina et al., 2002). Accordingly, one may expect that the increased number of stable MTs...
Figure 26. Loss of P1c affects polarized migration of keratinocytes. Phase contrast images of representative primary wt, P1c<sup>−/−</sup> and P0 keratinocytes (left columns) and migration tracks of single keratinocytes monitored by video microscopy (migration tracks) are shown. Elongated shape of P1c<sup>−/−</sup> keratinocytes (left column) and loss of polarization (migration tracks) can be observed. Scale bar, 100 µm.

encountered in peripheral cytoplasmic regions of mutant keratinocytes leads to a higher number of MTs converging at peripheral FAs, with consequences for their turnover and size. In addressing these issues I focused on P0 and wt cells as they showed the largest divergence in
migration velocities. In order to analyze FA turnover, p53<sup>−/−</sup> wt and P0 keratinocytes were transfected with EGFP-zyxin (Rottner et al., 1999) and subjected to time-lapse microscopy. In comparison to EGFP-zyxin-transfected wt keratinocytes, P0 cells display smaller and more numerous FAs (Figure 27A). After monitoring EGFP-zyxin labeled FAs, the ones of P0 cells showed a shorter life-time and thus increased turnover compared to wt cells (Figure 27A, bar graph). To test whether this difference was related to MT targeting, I analyzed in both cell types the proportions of MT-targeted versus not-targeted FAs by double staining for tubulin and vinculin (as FA marker) (Figure 27B). In contrast to wt cells, FA of P0 cells show more frequently colocalization of tubulin and vinculin (Figure 27B, arrowheads) than wt cells. I found that the cell type showing the higher proportion of MT-targeted FAs (Figure 27B, bar graph) was also the one displaying the more stable MT population (P0). To corroborate these data, I compared the size of vinculin-labeled FAs in wt and P0 keratinocytes and found that P0 FAs in general were smaller than wt (Figure 27C, bar graph), consistent with them being more frequently targeted by MTs and their higher turnover rates.

3.1.8 MTs in P1c-deficient keratinocytes show increased decoration by MAPs

To establish whether the data obtained in vitro showing MT destabilization using MAPs typical for neural cells (see Introduction) could be extrapolated to the situation prevailing in keratinocytes, I analyzed whether proteins of the MAP2/tau protein family were expressed in mouse epithelial cells; MAP2 expression in HaCat cells, an immortalized human keratinocyte cell line, had previously been reported by Liu <i>et al</i> (2007). As shown in Figure 25, when lysates of primary and immortalized mouse keratinocytes and of mouse epidermis were subjected to immunoblotting analysis using antibodies to MAP2, a ~70 kDa protein corresponding in size to MAP2c was detected in all samples tested (MAP2c is an isoform of MAP2 endowed with a potent MT assembly-promoting activity; Gamblin <i>et al</i>, 1996). In addition, proteins of ~95 kDa
Figure 27. Loss of P1c alters FA dynamics and stability. A) Images of EGFP-zyxin-expressing immortalized wt and P0 keratinocytes (scale bar, 10 µm) and bar graph showing FA turnover rates (n=3, 10 cells/ experiment). ***P<0.001. Error bars, ± SEM. B) Cells like in (A) double immunolabeled using antibodies to vinculin (green) and α-tubulin (red) to distinguish between MT-positive (arrowheads) and MT-negative FAs (scale bar, 15 µm). Bar graph, statistical analysis of MT-targeting of FAs in wt and P0 keratinocytes evaluated by a Chi-square test (n=3, 10 cells/experiment). *P<0.05. C) Detail images of vinculin-positive FAs at the periphery of cells (scale bar, 3 µm). Bar graph, statistical analysis of FA size in wt and P0 keratinocytes (n=3, 10 cells/experiment). ***P<0.001. Error bars, ± SEM.

and ~60 kDa were detected that probably corresponded to isoforms of tau that were recognized by anti-MAP2 antibodies due to the partial structural homology of tau and MAP2
(Al-Bassam et al., 2002). This notion was confirmed by incubating similar blots with monoclonal antibodies to tau, revealing the presence of the high-molecular weight form of tau (~95 kDa) as well as of 50–70 kDa tau isoforms (Figure 28). These results indicated that brain- and epithelia-derived cells express similar MAP species.

![Image](image1.png)

**Figure 28.** Tau and MAP2 are expressed in keratinocytes. Immunoblot showing the expression of MAP2 and tau in primary keratinocytes, immortalized (p53−/−) keratinocytes, and epidermis, with a brain lysate and purified MAP2c protein run as positive controls (n=3).

To confirm these results on the transcript level, analysis of tau and MAP2 mRNA expression was assessed by RT-PCR analysis of total RNA isolated from primary and immortalized keratinocytes as well as from epidermal tissue. As shown in Figure 29, using primers binding

![Image](image2.png)

**Figure 29.** Keratinocytes expressed tau and MAP2 at mRNA levels. RT-PCR results using primers specified in the text can be observed. Note expression of tau and MAP2 RNA in primary keratinocytes, immortalized keratinocytes, epidermis and brain used as a positive control.
to sequences in exons 9 and 11 of the tau gene (flanking the second MT binding domain repeat-encoding exon 10; Poorkaj et al., 2001; Dehmelt and Halpain, 2005; Andreadis, 2005), a DNA fragment of the expected size (390 bp) was amplified in all cases including adult mouse brain RNA (run as positive control). The 390 bp fragment corresponded to transcripts of the 4 MT-binding repeats contained in exons 9, 10 and 11 (Duff et al., 2000; Takuma et al, 2003). DNA sequencing of the PCR products confirmed that they had been derived from tau transcripts. To detect MAP2 transcripts, primers binding to sequences in exons 5 and 6, which are common to all MAP2 splice variants (Dehmelt & Halpain, 2005) were used. Single bands of the expected size (344 bp) were amplified from all tissues analyzed, including the positive control. Again, DNA sequencing of the PCR products confirmed their authenticity. Since among the samples analyzed at least the immortalized mouse keratinocyte cell line has been shown to be devoid of any contaminating fibroblasts or melanocytes (Osmanagic-Myers et al., 2006), these data clearly demonstrated that tau and MAP2 mRNAs were expressed in mouse keratinocytes. To examine whether tau and MAP2 are expressed in skin also on the protein level, sections of frozen foot pad skin from adult mice were subjected to immunofluorescence microscopy. A pronounced, predominantly cytoplasmic staining of all live keratinocyte cell layers of the epidermis was detected using specific antibodies to tau or to MAP2 (Figure 30). Similar staining patterns were observed on sections from other skin areas, e.g. ear and tail skin (data not shown). Controls without using primary antibodies were negative (Figure 30).

According to these results one could have expected higher levels of MAPs to be bound to MTs in plectin-deficient cells. To investigate this idea, primary wt, P1c−/−, and P0 keratinocytes were triple-immunostained using antibodies to MAP2, α-tubulin, and P1c. While in wt keratinocytes MAP2 showed a dotted staining pattern along MTs (similar to P1c) (Figure 31, upper panels), in both of the mutant cell types MTs were found decorated with MAP2 over much longer distances (Figure 31, lower panels, arrowheads), confirming our expectation.
Figure 30. Skin sections are immunoreactive with anti-tau and MAP2 antibodies. Frozen foot pad skin sections from adult wt mice were immunolabeled using antibodies to tau and MAP2 (left and center panels). Sections were immunolabeled using antibodies to tau or MAP2. Nuclei were stained with DAPI. In negative controls (right panel), primary antibodies were omitted. Note relatively strong immunofluorescence signals for both antigens in epidermis (e), and weaker signals in hair follicles (asterisk), and in a few scattered cells in the dermis (d). Scale bars, 25 µm.

Figure 31. MAP2-MT colocalization in wt and P1c<sup>−/−</sup> keratinocytes. Immunofluorescence microscopy of primary wt (upper row), P1c<sup>−/−</sup>, and P0 (lower row) keratinocytes was performed using antibodies to MAP2 (red), α-tubulin (green), and P1c (blue). Note that decoration of MTs with MAP2 can be found over longer distances on P1c<sup>−/−</sup> and P0 cells (arrowheads) than in wt keratinocytes. Scale bar, 2 µm.
However, in contrast to P1c\(^{-} \) keratinocytes, their P0 counterparts showed MAP2 staining not only along MTs but also in association with structures seemingly unlinked to MTs. Different levels of MT-unbound MAPs in P0 and P1c\(^{-} \) cells might be caused by elevated basal activities of Erk1/2 MAP and Src kinases (Osmanagic-Myers et al., 2006) in P0, but not in P1c\(^{-} \) keratinocytes. Since MT-binding of MAPs is regulated by several kinases including Src (Lim & Halpain, 2000; Lee, 2005), increased activity of Src in P0 keratinocyte might result in higher levels of MT-unbound MAP2.
Part II - Plectin 1c in neurons

Having identified a MT-destabilizing activity of P1c and a series of physiological consequences of P1c deficiency in keratinocytes, it was of particular interest to further analyze whether P1c is involved in biological functions of other cell types. Since P1c is the major plectin isoform expressed in neural cells, and MT organization and dynamics are essential for neuronal function, I selected P1c-deficient dorsal root ganglia (DRG) neurons as a second system in which to study P1c’s role in MT dynamics.

3.1.9 P1c-deficient neurons exhibit increased MT stability and altered MT dynamics

To investigate whether P1c causes changes in MT stability in neurons, DRG neurons were isolated from wt and P1c<sup>-/-</sup> mice, cultivated and treated with low doses of the MT-depolymerizing drug nocodazole. Since neuronal axons and neurites are densely populated with MT bundles, individual MTs remaining after nocodazole treatment could not be traced. Instead they were traced and quantified at the growth cone of neurons where it was possible to monitor single MTs. Figure 32 A shows images of the growth cone of wt and P1c<sup>-/-</sup> neurons immunostained with antibodies to tubulin after nocodazole treatment (upper row). To facilitate the visualization of nocodazole-resistant MTs, MTs were traced and their outlines together with growth cones boundaries are shown (Figure 32 A, lower row). P1c<sup>-/-</sup> DRG neurons displayed ~3.5 times more intact MTs in the growth cone after drug treatment than wt cells (Figure 32 B). These results were similar to those observed in keratinocytes (see Part I).

In another approach to evaluate MT stability, the distribution of acetylated α-tubulin along neurites was investigated. P1c-deficient DRG neurons were compared to their wt counterparts by double immunofluorescence microscopy using anti-tubulin antibodies that did not discriminate between the modified (acetylated) and unmodified forms.
Figure 32. P1c affects drug resistance and stability of neuronal MTs A) Nocodazole-treated DRG neurons (wt and P1c<sup>-/-</sup>) were immunolabeled using antibodies to α-tubulin. Black lines outline the periphery of the growth cone and the tracing of nocodazole-resistant MTs. Note the presence of higher number of MTs remaining after drug treatment in P1c<sup>-/-</sup> growth cones (lower row, right panel). Scale bar, 4 µm. B) Bar graph showing statistical evaluation of results shown in (A) applying a Student’s t test (n=3). *P<0.05. Error bars, ± 95% CI.

of the protein and antibodies that were specific for the acetylated version. The growth cones of wt DRG neurons were found to be barely immunoreactive with antibodies to acetylated tubulin, whereas in the growth cones of P1c<sup>-/-</sup> DRG neurons acetylated MTs were clearly visible (Figure 33 A). A quantification of acetylated tubulin-positive areas (in pixels) versus unmodified (total) tubulin-positive areas showed the acetylated form to be ~2-fold increased in plectin-deficient DRG neurons compared to wt cells (Figure 33 B).

In order to confirm that the lack of P1c was directly responsible for the increased acetylation of MTs in P1c<sup>-/-</sup> DRG neurons, isolated neurons were transiently transfected with expression plasmids encoding a P1c-mCherry fusion protein using nucleofection. Based on
Figure 33. P1c affects MT acetylation in DRG neurons. A) Immunofluorescence microscopy was performed on wt, P1c<sup>−/−</sup>, and P1c-mCherry-transfected P1c<sup>−/−</sup> DRG neurons using rat monoclonal anti-tubulin (green) and mouse monoclonal anti-acetylated tubulin antibodies (blue). Note that only the growth cone of P1c<sup>−/−</sup> DRG neurons is prominently immunoreactive with antibodies to acetylated tubulin. Scale bars, 4 µm. B) Statistical evaluations (graph) of acetylated tubulin to total tubulin fluorescent signals were done by a one-way ANOVA-test and post-hoc Tukey correction compared to wt values (n=5). *P<0.05. Error bars, ± SEM.

Electroporation, this technique facilitates the transfer of cDNA constructs directly into the cell nucleus via short high-voltage pulses and optimized buffers to minimize damage to the cellular membranes (Zeitelhofer et al., 2007). As shown in Figure 33, the acetylated tubulin immunostaining of the P1c-mCherry-transfected P1c<sup>−/−</sup> DRG neuronal growth cone is similar to the one of the wt growth cone. Measurements of acetylated tubulin-positive areas in
comparison to total tubulin-positive areas led to a statistically significant reduction of MT acetylation in P1c-mCherry-transfected P1c⁻/⁻ DRG neurons, approximating the levels of wt neurons (Figure 33 B). Thus, these data were fully consistent with the observed higher nocodazole resistance of MTs in mutant cells (see Figure 32) and they confirmed that the lack of P1c⁻/⁻ leads to increased stability of MTs.

Previous results from this laboratory and my own data, including the observed accumulation of MAPs along MTs of P1c-deficient keratinocytes (Figures 13 and 31), suggested that P1c antagonizes MAPs and thereby causes destabilization of MTs. According to this model, one may expect to find more MAPs to be bound to MTs in the absence of P1c. To investigate whether this was the case, the levels of tau bound to endogenous MTs were quantified in wt and P1c⁻/⁻ mouse brain lysates prepared under MT-stabilizing conditions. This so called endogenous MT-binding assay (Planell et al, 2008) allowed polymeric MTs and their co-assembling (bound) proteins to be sedimented by high-speed centrifugation. Immunoblotting of total lysates, supernatant and pellet fractions revealed three bands corresponding in size to the most common tau isoforms (between 50 and 60 kDa) in all fractions (Figure 34). While P1c⁻/⁻ and wt samples showed similar levels of tubulin in all fractions, tau levels were visibly decreased in the soluble, and increased in the insoluble (MT-bound) fraction of P1c⁻/⁻ (compared to wt samples), at least in the case of the ~50 kDa tau isoforms. Quantitative immunoblotting analysis more clearly revealed increased levels of MT-bound tau in P1c⁻/⁻ brain lysates (Figure 34 B).

In order to investigate whether the lack of P1c leads to altered MT dynamics in DRG neurons, time-lapse video microscopy of GFP-tagged EB1 was performed. This technique allows the visualization of MT tips in the growth cone as well as along the axon of the neurons (Stepanova et al, 2003; Su et al, 1995; Akhmanova & Steinmetz, 2008). The image shown in Figure 35 A represents a growth cone of a GFP-EB1-transfected DRG neuron showing three
Figure 34. Tau is increased in the MT-bound fraction in P1c-deficient brain lysates. Endogenous MT-binding assay of tau was performed using wt and P1c\(^{-/-}\) brain lysates prepared under MT-stabilizing conditions. Tau and tubulin from total lysates, soluble, and the bound fractions were evaluated by immunoblotting analysis using anti-tau antibodies. Note that the \(~50\) kDa tau isoform is increased in the MT-bound fraction of P1c\(^{-/-}\) brain lysates, while tubulin levels are similar in both samples. Bar graph represents quantification of the immunoblotting data. Error bars, ± SEM.

GFP-EB1 comets (black dots). Analysis of GFP-EB1 tracks of transiently transfected DRG neurons revealed faster MT growth in plectin-deficient cells (Figure 35 B), similar to what I had previously found in keratinocytes.

3.1.10 P1c-deficiency affects growth cone extension and neuritogenesis

Neuronal polarization, which is essential for neuronal development, requires active participation of actin filaments and MTs for growth cone extension. The neurite elongation process can be divided into three stages: protrusion, engorgement, and consolidation (Goldberg & Burmeister, 1986). Neuronal growth cones are rich in MTs in their central domain and in actin filaments at their periphery. Peripheral actin arcs prevent MTs from invading the peripheral region but during protrusion, actin filaments reorient towards growth direction, creating an actin-free zone that can be invaded by MTs. Vesicles and organelles transported by MTs enter in this area during the engorgement phase. In the subsequent consolidation stage, actin polymerization and protrusion stop and the MTs are bundled at the growth cone wrist, causing the elongation of the neurite shaft (Stiess & Bradke, 2011). The elongation
process is positively regulated by enhanced MT stability and restrained actin arc formation (Neukirchen & Bradke, 2011). Moreover, MTs are also required for the accumulation of signaling molecules at the growth cone, including Src family kinases and Rho family GTP regulators involved in their cross-talk with actin filaments (Lowery & Vactor, 2009). Although the role of MTs during growth cone extension is still not well understood, it has been shown that dynamic MTs are indispensable for persistent growth cone advance (Tanaka et al, 1995).

In light of the fundamental role of MTs in neurite outgrowth, I tested whether the increased stability and growth rates of MTs in growth cones of P1c−/− DRG neurons were affecting neurite extension. For this I measured growth cone extension of DRG neurons (isolated from P1c−/− and wt mice) by phase-contrast and time-lapse video microscopy at 10 min intervals during at least 5 h. The comparison of neurite lengths reached by DRG neurons within the same time periods revealed a ∼25% faster neurite extension rate in P1c−/− compared to wt cells (Figure 36). Interestingly, also the morphology of P1c-deficient growth

Figure 35. P1c affects MT growth rates in DRG neurons. A) A single frame of a time-lapse recording of GFP-EB1 visualized in the growth cone of a DRG neurons is shown. GFP-EB1 complexes are seen as black dots inside the growth cone. Scale bar, 0.8 µm. B) Bar graph shows statistical analyses of MT growth rates subjected to a Wilcoxon-Mann-Whitney U-test (n=3; ~5 cells/experiment). *P<0.05. Error bars, ± SEM.
Figure 36. P1c-deficiency leads to faster growth cone extension. A) Growth cone extension of wt and P1c−/− neurons was visualized overnight using phase-contrast time-lapse microscopy. Images show time points indicated. Scale bar, 8 µm. B) Graph represents statistical analysis of growth cone extension velocity subjected to a Student's t-test (n=8) ***P<0.001. Error bars, ± 95% CI.

cones was found to be different from that of their wt counterparts. Growth cones of P1c−/− neurons appeared more robust compared to wt neurons, as indicated by its darker appearance. Thus unexpectedly, the higher stability of MTs in P1c−/− DRG neurons led to a more efficient elongation of the growth cone, and consequently to faster neurite extension. A possible explanation for this phenotype presumably is a more pronounced engorgement phase. Indeed, P1c-deficient neurons displayed a higher proportion of growth cones that were spread out (resembling the engorgement phase) (Figure 36, 2h, lower panel). Since MTs are more stable in P1c−/− neurons, they may invade the actin-free zone after protrusion, resulting in more vesicle and organelle transport into the growth cone required for the consolidation phase.

As neuronal morphology is dependent on cytoskeleton organization, and NFs, in
particular, have been reported to maintain the characteristic shape of axons during the structural changes that take place upon axonal elongation (Lin & Szaro, 1995), I investigated whether the changes in growth cone morphology of P1c-deficient DRG neurons were linked to changes in NF organization. Confocal immunofluorescence microscopy of wt and P1c<sup>−/−</sup> DRG neurons using antibodies to NF proteins revealed that in the case of P1c-deficient neurons ∼100% of the growth cones were filled with NFs, while this was the case for only ∼25% of the wt growth cones (Figure 37). This phenotype was reminiscent of plectin-deficient cell types, keratinocytes and fibroblast, where IF networks had been found to extend to the outermost parts of the cell instead of being restrained to its central part, as is typical for wt cells (Osmanagic-Myers <i>et al</i>, 2006; Burgstaller <i>et al</i>, 2010).

Being part of the mechanism underlying neuronal branching and neurite extension, MT dynamics are essential for neuronal development (Conde & Cáceres, 2009). To investigate whether MT stabilization effected by P1c deficiency had an influence on neuronal

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**Figure 37. Lack of P1c in DRG neurons affects IF localization.**

A) Growth cones of wt and P1c<sup>−/−</sup> DRG neurons were immunolabeled using antibodies to NFs (green) and stained for actin with phalloidin (red). Immunofluorescence microscopy images of P1c<sup>−/−</sup> DRG neurons show NF-positive growth cones (arrowhead, right panel) in contrast to wt cells that display more NF-negative growth cones (arrows, left panel). Scale bar, 10 µm. B) Graph shows quantification of NF-positive and NF-negative growth cones of wt and P1c<sup>−/−</sup> DRG neurons subjected to Chi-square test (n=4). ***P<0.001. Error bars, ± 95% CI.
development, I decided to analyze the neurite outgrowth of DRG explants. For this, DRG explants, isolated from wt and P1c−/− mice, were cultivated embedded in matrigel and neurite outgrowth was measured after 24 h. As shown in Figure 38, P1c-deficient DRGs showed less neurites growing out of explants than their wt counterparts. In addition, P1c-deficient neurites seemed to be less branched than those of wt.

Figure 38. Lack of P1c impairs neurite outgrowth. A) DRG explants of wt and P1c−/− mice were cultivated in matrigel. Phase contrast microscopy revealed less neurite outgrowth in P1c−/− DRG explants. Scale bar, 65 µm. B) Graph shows quantitative comparison of neurite outgrowth (normalized to the DRG surface length in µm) subjected to Student’s t-test (n ~20). ***P<0.001. Error bars, ± SEM.

To further analyze whether P1c−/− neurons undergo less branching, cultivated primary P1c−/− DRG neurons were compared with primary wt DRG neurons. According to their level of branching three different populations of neurons, showing low, medium or high branching, could be distinguished in both genotypes. While most wt neurons showed a medium level of branching, in P1c−/− neurons the category of low level branching was most frequent (Figure 39). These results suggested that in P1c-deficient DRG neurons branching was impaired. The reason for this could have been excessive amounts of tau bound to MTs, interfering with the severing activity of katanin, a protein triggering axonal branching (Yu et al, 2008).
Figure 39. P1c-deficiency leads to decreased neurite branching. A) Images obtained by immunofluorescence microscopy show representative examples of the three populations of DRG neurons sorted by their branching levels (branching points normalized to neurite length): low (<0.015), medium (0.015-0.030), and high branching (>0.030). Scale bar, 20µm. B) Line graph represents quantification of wt and P1c−/− DRG neurons classified according to the number of branches per neurite length (low, medium, and high). Results were subjected to Wilcoxon-Mann-Whitney U-test (n=20). *P<0.10.

The Rho family of GTPases plays an important role in various developmental processes of neurons, including differentiation and neurite outgrowth. Activation of Rho is generally associated with the inhibition of neurite formation (Jeon et al, 2011). To assess whether the impaired ability of P1c-deficient DRGs to form neurites correlated with an overactivation of Rho, I measured the RhoA activation status in brain lysates. GTP-RhoA was pulled down from brain lysates using a GST-Rhotekin-binding domain (GST-RBD) fusion protein, containing a binding domain for activated (GTP-bound) RhoA. GTP-RhoA such isolated was analyzed by quantitative immunoblotting using antibodies to RhoA, and values measured were subjected to a Student’s t test (Figure 40). While the total amount of RhoA was found to be similar in wt and
in P1c<sup>-/-</sup> lysates, a tendency towards a higher content of active RhoA was found for P1c<sup>-/-</sup> lysates, without being statistically significant, however.

![RhoA-GTP, RhoA-GDP, Total RhoA](image)

**Figure 40.** P1c-deficiency barely affects the activation of RhoA. A) Active RhoA pull down assays were performed using wt and P1c<sup>-/-</sup> brain lysates. GTP-bound RhoA was precipitated using GST-RBD fusion protein and quantified by immunoblotting analysis using antibodies to RhoA. B) Graph show statistical analysis of the signals obtained (n=3). Error bars, ± SEM.

### 3.1.11 Lack of P1c leads to abnormal vesicle transport in DRG neurons

One of the best-known functions of MTs is their role in mediating vesicle transport through the MT-based motor proteins kinesin and dynein. Alteration of MT organization and dynamics are shown to interfere with important neuronal functions such as synaptic vesicle transport or glucose uptake. Therefore, it was of interest to investigate whether the lack of P1c in DRG neurons and the stabilization of MTs had an effect on glucose uptake. Glucose is the major energy source for neurons, and the main glucose transporters identified in brain are GLUT1 and GLUT3 (Duelli & Kuschinsky, 2001), similar to keratinocytes. GLUT1 is the main isoform expressed in astrocytes and endothelial cells, while GLUT3 is mainly found in neurons. To test whether P1c-deficient DRG neurons, similar to P1c<sup>-/-</sup> keratinocytes, showed increased glucose uptake. DRG-neurons were incubated with 2-NBDG, and fluorescence signals within cells were monitored by confocal microscopy. Comparing wt and P1c<sup>-/-</sup> DRG neurons, increased glucose uptake was observed in P1c<sup>-/-</sup> neurons according to expectations (Figure 41). Glucose uptake of P1c-deficient DRG neurons was in fact ~3 times higher than that of wt DRG neurons, similar to the results obtained with P1c<sup>-/-</sup> keratinocytes.
Figure 41. Alterations of glucose uptake in P1c-deficient DRG neurons. A) Primary wt and P1c−/− DRG neurons were incubated with 2-NBDG and fluorescence intensity of cells was measured. Note higher fluorescence intensity of P1c−/− DRG neurons after 2-NBDG treatment. Scale bar, 30 µm B) Bar graph shows statistical analysis of data subjected to Student's t-test (n=3, ~20 cells/experiment). ***P<0.001. Error bars, ± 95% CI.

Another well-known MT-dependent vesicular transport event is the transfer of peroxisomes to the cell periphery as a defence mechanism against oxidative stress. Alteration of tau levels in neurons can enhance oxidative stress by inhibiting transport and redistribution of peroxisomes, eventually leading to the absence of catalase in neurites (Stamer et al, 2002). To study whether this was also the case in P1c-deficient DRG neurons, I exposed wt and P1c−/− DRG neuronal cultures to 250 µM H2O2 for 5 min. Then, I assessed the vulnerability of the neurons to oxidative stress by quantifying the proportion of collapsed (bulb-shaped) growth cones as an indicator of oxidative stress damage. Quantitative measurements revealed ~4 times more collapsed growth cones in P1c−/− DRG neurons compared to wt cells (Figure 42). These results suggested that enhanced levels of MT-bound MAPs, as is typical for P1c−/− neurons, interfere with the cellular machinery protecting the cell against oxidative stress, probably by impeding vesicle (peroxisome) transport.
MT-bound MAPs can spatially control MT-dependent axonal transport, by differentially regulating the activity of MT-dependent motor proteins like kinesin and dynein (Dixit et al., 2008). Since MT-bound tau levels were found to be increased in P1c^{-/-} brain lysates, it was intriguing to explore whether tau levels can interfere with synaptic vesicle transport in DRG neurons. To directly visualize synaptic vesicle transport in DRG neurons, cells were loaded with FM1-43, an amphiphatic dye used to monitor endosomal traffic, especially synaptic vesicle recycling at nerve terminals (Gaffield & Betz, 2006). FM dyes are non-fluorescent in aqueous media, but they become intensely fluorescent after their incorporation into a lipidic membrane.

In neurons that are actively recycling neurotransmitters, these dyes are internalized within recycled synaptic vesicles. After activating DRG neurons by exposure to high concentration of potassium to trigger synaptic vesicle recycling (Gaffield & Betz, 2006), a higher percentage of retrogradely moving vesicles were detected in P1c^{-/-} DRG neurons compared to their wt counterparts (Figure 43). These results suggested that the activity of kinesin (the motor protein responsible for anterograde transport) was decreased as a result of more tau bound to MT in P1c^{-/-} cells (Figure 34).
3.1.12  P1c deficiency causes abnormalities in membrane excitability of DRG neurons

It has been reported that tubulin in its acetylated form associated with Na⁺, K⁺-ATPase in vitro and in vivo and participates in the regulation of its enzymatic activity (Arce et al., 2008). Furthermore, it has been suggested that in cardiac myocytes modulation of Na⁺ current is a tubulin/GTP-coupled process (Casini et al., 2009). As acetylation of tubulin correlates with increased MT stability, it could be that the more stable MT population found in P1c⁻/⁻ DRG neurons has an influence on the Na⁺ influx into cells.

To investigate whether ion channel activity was altered in P1c-deficient DRG neurons, I measured the activity and expression levels of Na⁺ channels. Channel activity was measured using CoroNa green, a fluorescent Na⁺ indicator that allows the visualization of Na⁺ levels inside the cell. The fluorescence microscopy images suggested a slightly decreased channel activity in P1c⁻/⁻ DRG versus wt neurons, although after subjecting the data to a Student’s t test
the observed difference was found not to be statistically significant (Figure 44). However, it could be that the $\text{Na}^+$ indicator used was not sensitive enough to measure minor $\text{Na}^+$ fluctuations. When I analyzed the expression levels of $\text{Na}^+$ channels at the membrane of wt and $\text{P}1\text{c}^{-/-}$ DRG neurons by immunofluorescence microscopy, a homogeneous distribution of the channels at the surface of wt DRG neurons was revealed (Figure 45), whereas in $\text{P}1\text{c}^{-/-}$ DRG neurons, channels appeared in a clustered way and unevenly distributed over the cell surface.

![Figure 44](image)

**Figure 44. P1c deficiency does not affect intracellular Na$^+$ concentration.** A) Phase contrast (left column) and fluorescent images of wt and $\text{P}1\text{c}^{-/-}$ DRG neurons loaded with CoroNa green (right column) are shown. Scale bar, 15 µm. B) Graph represents fluorescent intensities measured using ImageJ software (NIH Image, Bethesda, MD). (n=3, ~20 cells/experiment). Error bars, ± SEM.

To directly investigate whether the ability of P1c-deficient DRG neurons to produce action potentials was affected, the membrane excitability of $\text{P}1\text{c}^{-/-}$ DRG neurons was measured
Figure 45. Clustering of P1c−/− Na+ channels in P1c+/+ A) Immunofluorescence microscopy of wt and P1c−/− DRG neurons immunolabeled using antibodies to Na+ channels. Note homogeneous dotted distribution of Na+ channels in wt DRG neurons contrasting their P1c−/− counterparts. Scale bar, 6 µm. B) Bar graph show statistical analysis of the Na+ channel clusters size. Data were subjected to Student’s t-test (n=3, ~25 cells/experiment). ***P<0.001. Error bars, ± 95%CI.

using current clamp electrophysiology. These experiments were performed in collaboration with A. Yousuf and K. Schicker in the laboratory of S. Böhm (Centre for Physiology and Pharmacology, Medical University of Vienna). The current clamp technique records whatever voltage the cell generates as a result of stimulation. When DRG neurons were subjected to these measurements, threshold levels reached by wt specimens were -35 mV and -42 mV (Figure 46). Although the data obtained with P1c−/− DRG neurons on average were not significantly different, the data distribution was remarkably wider in this case with a much higher difference between the minimum and maximum value compared to wt cells. These results were consistent with possible alteration of membrane excitability regulation.

3.1.13 Differential extractability of tau from hippocampal wt and P1c−/− lysates

Tau protein is very well known due to its implication in a group of neurodegenerative diseases known as tauopathies. The hallmark of tauopathies is the occurrence of tau aggregates and tau hyperphosphorylation (Kampers et al, 1999; Ballatore et al, 2007; Ittner et
Figure 46. **P1c deficiency causes a wide range of membrane potentials.** Membrane potentials of wt and P1c⁻/⁻ DRG neurons were measured in current clamp mode. Box and whisker diagrams represent the statistical distribution of threshold voltages measured (n= 4).

al, 2010; Fu et al, 2010). Since the levels of tau bound to MTs are increased in P1c⁻/⁻ brain lysates, I considered of high interest to investigate whether tau aggregates were present in P1c-deficient brain. Normal tau is a highly soluble protein that loses its solubility under pathological conditions. A potential tendency of tau protein to form aggregates in P1c-deficient hippocampus was first examined by sarcosyl extraction of the tissue, whereby one obtains a soluble and an insoluble fraction (Sydow et al, 2011). When tau is forming part of aggregates, it will be found in the insoluble fraction. The analysis of sarcosyl-insoluble fractions produced from 12 P1c-deficient hippocampuses revealed a significantly reduced amount of insoluble tau in comparison to wt samples (Figure 47). This result suggested that tau in P1c-deficient brain did not form aggregates, but was more soluble, probably because it remained bound to tubulin which ends up primarily in the soluble fraction of the sarcosyl extract.
Figure 47. Sarcosyl-insoluble P1c<sup>−/−</sup> fractions lysates contain reduced levels of tau. Sarcosyl soluble and insoluble fractions were subjected to immunoblotting analysis using antibodies to tau. Note the decreased tau-specific signal in the insoluble fraction of P1c<sup>−/−</sup> hippocampus, while comparable levels of tau were found in the sarcosyl soluble fractions. Tubulin levels were similar in all fractions.
4 DISCUSSION

Plectin is one of the most abundant and versatile cytolinkers expressed in mammalian cells. One of its outstanding features is its functional diversity that is mainly based on the alternative splicing of a series of different first coding exons (Fuchs et al, 1999). A variety of isoforms generated in this way just differ in short N-terminal sequences that specify distinct properties of the cytolinker, including its cellular targeting and N-terminal binding partners (Rezniczek et al, 2004; Abrahamsberg et al, 2005). In muscle, P1 is associated with the outer nuclear/ER membrane system, P1d with the Z-disks, P1f with the sarcolemmal dystrophin-glycoprotein complex and P1b with mitochondria (Konieczny et al, 2008; Rezniczeck et al, 2007). In epidermis, P1a links IFs to HDs and it undergoes selective degradation that seems to be required for epithelial differentiation (Walko et al, 2011). However, the role of P1c, a main isoform in the epidermis and the neural system, remained unclear up to date. In this thesis I provide evidence for a role of P1c in MT dynamics. Moreover, by studying MT-dependent basic cellular functions in keratinocytes and neuronal cells, the two cell types where P1c is most abundantly expressed, I could demonstrate that P1c-mediated MT regulation is of broad biological significance.

Plectin 1c acts as a MT destabilizer in a spatially-controlled manner

Based on three types of assays, resistance to MT depolymerizing drugs, assessment of acetylated tubulin levels, and direct observation of MT dynamics by video microscopy, this study provides evidence that P1c deficiency in keratinocytes and in DRG neurons results in increased MT stability. Increased stability was coupled with a decrease in the dynamics of MTs, manifesting as less frequent transitions between MT assembly and disassembly states. A destabilizing function of this kind was not expected for a cytolinker protein such as plectin. In fact, it was the opposite of what had been observed for other cytolinkers, such as ACF7 and
BPAG1, that have been shown to stabilize MTs (Kodama et al, 2003; Yang et al, 1999). Thus, plectin seems to be the first genuine cytolinker protein shown to act as a MT destabilizer. This finding, together with the observation that plectin can act also as a destabilizer of the actin cytoskeleton (Andrä et al, 1998), opens a new perspective on how cells maintain their cytoskeleton integrity. This study shows that this process is the result of a fine-tuned regulatory mechanism, in which cytolinker proteins are key players, not only by anchoring and providing stability to dynamic filaments, but also by destabilizing them and thus favoring their dynamic behavior, as demonstrated in this work for P1c.

My study demonstrates that abnormal MT stability in P1c−/− keratinocytes could be rescued, i.e. restored to normal levels, in P1c-deficient keratinocytes by forced expression of P1c-EGFP fusion proteins. However, from experiments where I transfected cells with truncated forms of P1c lacking the IF-binding domain, evidence emerged that the rescue potential of P1c was dependent on the localization of the protein, in particular its association with IFs. Unable to be recruited to IF networks, and instead associating with actin filaments, the truncated versions tested showed only a limited, if any, rescue potential. These findings point towards a fascinating new feature of cytoskeletal filament cross-talk, namely the potential of IFs to destabilize rather than stabilize MTs via an associated cytolinker protein, thereby stimulating MT dynamics. In previous studies, actin was found to reinforce and stabilize MTs at the cell periphery via ACF7, resulting in polarization and more effective migration of cells (Kodama et al, 2003; Wu et al, 2011).

**P1c destabilizes MTs by antagonizing MAP-mediated MT stabilization**

As a possible mechanism underlying MT destabilization through plectin, previous results obtained in our laboratory suggest that plectin’s SH3 domain interferes with MAP-MT binding and consequently antagonizes MAP-mediated MT stabilization (see model in Figure 48).
Figure 48. Model depicting plectin as MT destabilizer. Binding of P1c to MTs presumably occurs via its isoform-specific N-terminal sequence including the ABD. The SH3 domain located within the plakin domain binds to MAPs. Interference with the MT-stabilizing function of MAPs ensures a dynamic MT network in wt cells. When P1c is absent more MAPs can bind along MTs leading to their stabilization.

Several observations support such a mechanism. First, binding of plectin to HMW-MAPs was reported a few years ago (Herrmann & Wiche, 1987), and afterwards, using expression constructs encoding various domains of plectin, it could be shown that plectin’s SH3 domain is responsible for this interaction (G. Walko, unpublished data). Second, MT co-assembly assays performed previously in the lab by L. Janda demonstrated that increasing concentrations of plectin’s SH3 domain lead to decreased levels of MAPs binding to MTs and, as a consequence, less polymerized MTs (unpublished data). Third, based on an endogenous MT binding assay (see Results), it could be show that plectin’s SH3 domain was able to detach MAPs from stabilized MTs present in brain tissue lysates. Fourth, comparing wt, P1c−/−, and P0 keratinocytes in this thesis project, I found MAP2 to be associated with larger portions of cellular MTs in mutant compared to wt cells, where P1c’s presence interferes with MAP-MT binding. This observation was confirmed also by in vitro experiments performed with brain lysates where the amount of tau bound to MTs was analyzed. Comparison of wt and P1c−/− MT-bound tau fractions revealed a greater amount of tau bound to MTs in plectin-deficient samples.
Is MT destabilization P1c-specific?

Since plectin’s SH3 domain is expressed in all plectin isoforms known and is contained in many other proteins, including MT regulators, such as Fyn or Src (Lee et al., 1998), SH3 domain-mediated MT destabilization could be a function performed by plectin isoforms others than P1c as well as by many other proteins. However, although all plectin isoforms apparently have the potential to destabilize MTs via their SH3 domain, the destabilizing potential of P1c is likely to be more efficient than that of other isoforms. A possible reason for this could be a closer apposition of P1c to the MT surface accomplished by an additional tubulin-binding interface formed by its isoform-specific sequence and the succeeding ABD. The partial cosedimentation of endogenous keratinocyte P1c with MTs, but not of the equally abundant P1a, as previously observed in our laboratory (G. Walko, unpublished data), points in this direction. A weak, or even only transient binding of P1c to tubulin polymers might suffice to bring plectin’s SH3 domain in contact with MAPs. However, the validity of such a mechanism has to be rigorously tested.

The hypothetical model depicted in Figure 48 proposes that, by antagonizing MAP-mediated MT stabilization, IF-associated P1c leads to a MT network that is more susceptible to localized disassembly in the vicinity of IFs. Conversely, in a plectin-deficient system, the fraction of MAPs attaching along MTs is higher, leading to promotion of assembly and stabilization of the polymer. This model does not exclude additional or alternative ways of how plectin may regulate MT dynamics, e.g. through its scaffolding and platform function for other putative MT regulators.

P1c deficiency impairs neurite outgrowth and branching

Interestingly, isolated DRG neurons as well neurite outgrowth from DRG explants of P1c-deficient mice showed reduced branching and neurite outgrowth in comparison with their wt
counterparts. As the results obtained in this work indicates, there are increased levels of tau bound to the MT fraction of P1c-deficient brain lysates. According to Yu et al, 2008, the decrease in neurite branching observed in P1c-deficient cells could be the result of tau protection against the MT-severing protein katanin, impairing the formation of new branches. However, although P1c-deficient DRG neurons show impaired branching and DRG explants have decreased potential to regenerate newly formed neurites, MT stabilization caused by P1c-deficiency promotes growth cone extension. P1c-deficient growth cones extend faster than wt growth cones. A possible explanation for this could be that stable MTs of P1c−/− DRG neurons might act as a nucleation seed for the MT assembly and protrusion that is required during axon outgrowth (Conde & Cáceres, 2009). In addition they may provide tracks for MT-dependent motors that transport organelles and NFs to the growth cone during the outgrowth process (Lee et al, 2011). Together all these features would explain why growth cones observed in P1c-deficient DRG neurons are more spread out than their wt counterparts. They also correlate with the observed high percentage of P1c-deficient growth cones filled with NFs (~100%) in contrast to the much lower proportions (~35%) of NF-positive growth cones in the case of wt. The presence of NFs in P1c-deficient growth cones suggests an increased transport of NFs along MTs (Lee et al, 2011) that could result in more spread out growth cones as observed.

The neuronal outgrowth alterations observed in P1c-deficient DRG neurons suggest that P1c could be critical also in neuronal polarization. Since plectin has already been identified as a binding partner of ankyrin G (Maiweilidan et al, 2011), a marker for the axon initial segment (AIS) (Dzhashiashvili et al, 2007), it will be interesting to analyze whether ankyrin G localization at the AIS is impaired due to P1c deficiency. Moreover, tubulin acetylation (a characteristic feature of P1c−/− DRG neurons) has been found to impair the concentration of ankyrin G at the AIS as well (Tapia et al, 2010). However, to further
investigate neuronal polarization in P1c-deficient neurons, a more suitable system than DRG neurons, such as hippocampal neurons, should be used due to their pyramidal shape that allows the monitoring of axon formation.

**Is axonal transport altered in P1c-deficient DRG neurons?**

Based on three different types of experiments, glucose uptake, synaptic vesicle transport visualization, and oxidative stress resistance, this thesis shows alteration in MT-dependent transport. However, while the results of synaptic vesicle transport and resistance towards oxidative stress point to an impairment of MT-dependent transport, the outcome of the glucose uptake shows the opposite. This apparent contradiction might be due to differences in the cellular localization where these phenomena where analyzed. While glucose uptake was mainly monitored in the cell body, vesicle transport and resistance towards oxidative stress were measured at the distal end of the neurites, where tau levels are increased in comparison to the cell body. Kinesin motility is responsible for anterograde vesicle transport and might be spatially regulated by tau, detaching from MTs when tau concentration is high (Dixit *et al*, 2008). Since tau concentration increases following a proximal-distal gradient, kinesin anterograde transport at the end of the growth cones would be inhibited. However, data reported from Yuan *et al* (2008) contradict this hypothesis showing unaffected axonal transport rates in mice lacking or overexpressing tau protein. The results that I report here would support Dixit *et al*’s hypothesis, as synaptic vesicle anterograde transport and resistance towards oxidative stress mediated by peroxisomes transport along the neurites in P1c-deficient cells were found reduced, correlating with to the higher levels of tau associated with MTs along the neurite in comparison to wt DRG neurons.
Is membrane excitability affected in P1c-deficient neurons?

Na$^+$ influx in P1c-deficient cells showed a tendency to be reduced in comparison with wt cells, although the difference found was not statistically significant. The distribution of Na$^+$ channels was abnormal as well. Unlike wt cells Na$^+$ channels were not homogeneously distributed across P1c-deficient DRG neurons distributed (Figure 46). As described in the Results, wt neurons were stimulated between -35 mV and -42 mV, while the corresponding threshold levels reached by P1c-deficient DRGs were more varied. Since P1c-mediated MT destabilization appears to be a spatially controlled event, the diversity of results obtained from current clamp technique could be due to the different points where the current was applied, making it difficult to compare and evaluate the different measurements. Anyway, these results do not provide any explanation for the reduced motor nerve conduction velocity observed in P1c-deficient mice. Thus, it could be that the higher number of small caliber axons observed in these mice is the sole reason for the reported phenotype (Fuchs et al., 2009).

Are there any similarities between P1c deficiency and tauopathy or neurodegenerative disorder phenotypes?

Tauopathies and neurodegenerative disorders share some common features like protein aggregate formation, alterations of MT stability, and hyperphosphorylation of proteins (Lee et al., 2001; Avila et al., 2004). For example Alzheimer disease is characterized by the presence of neurofibrillary tangles or aggregates formed by hyperphosphorylated tau and NFs (Rudrabhatla et al., 2011). Similarly, the neurodegenerative disorder Charcot-Marie-Tooth disease manifests with NF aggregates, impaired MT-dependent transport, aberrant mitochondria distribution, reduced motor nerve conduction velocity, and increased MT stability (Schröder, 2005). Interestingly, P1c-deficient mice share some of these phenomena including, impaired MT-dependent transport, increased MT stability and reduced motor nerve conduction
velocity. Moreover increased NF-positive staining but no aggregates were found.

Results presented in this thesis show increased levels of tau bound to MTs in P1c-deficient brain lysates. Whether MT-bound tau in P1c−/− neurons is hyperphosphorylated is still an open question. Although hyperphosphorylation of tau (usually connected with impaired memory) has been found to be the cause for tau detachment of MTs (Biernat et al., 1993; Drewes et al., 1995), these studies were focused on human tau and not murine tau. Human differs from murine tau in the ratio of the isoforms, containing more 3R (3 MT-binding domain repeats) tau than 4R (4 MT-binding domain repeats) tau. In contrast, murine tau is almost all 4R tau and is more difficult to detach from MTs. Thus, it would be still possible that the increased levels of tau bound to MTs in P1c−/− brain lysates would be hyperphosphorylated. In fact, hyperphosphorylated tau bound to MTs present in brain lysates has been recently reported (Planel et al., 2008).

It has been suggested that as well as an increase in the amount of intracellular tau, structural changes to this protein, modifications by phosphorylation, or its aggregation could produce toxic effects in cells (Avila, 2010). Future investigations using P1c-deficient mice could help to clarify the mechanism underlying the toxic effects of tau. Questions to be addressed include whether toxicity is a result of tau detachment from MTs, or whether tau could still promote toxicity even when it remains bound to the polymers. Extracellular tau was also found to promote cellular toxicity by increasing the levels of intracellular calcium (Gómez-Ramos et al., 2006). In addition, tau overexpression results in its secretion via membrane vesicles as a mechanism to avoid intracellular toxicity (Simón et al., 2012). However, it remains to be elucidated how excess tau protein gets associated with these membrane vesicles. Interestingly, preliminary data from this laboratory suggest that the rodless isoform of P1 (an isoform targeted to membranes) might be actively transported to the exterior of differentiated myoblast. In fact, plectin was found to co-fractionate with Golgi stacks suggesting an
association of plectin with the Golgi apparatus (P. Möseneder, Master thesis). Thus, it might be of interest to study whether plectin is involved in tau association with membrane vesicles.

**Closing remarks**

In conclusion, my study adds a new facet to the already broad spectrum of plectin functions. The fact that plectin can act as a MT destabilizer opens a new perspective on the role of cytolinkers in regulating cytoskeletal integrity. In fact, P1c is the first cytolinker protein reported to possess a MT-destabilizing function. The analyses of P1c-deficient keratinocytes and DRG neurons combined with in vitro data using recombinant proteins, suggest that plectin antagonizes MAP-mediated MT-stabilization in a spatially controlled manner. As a consequence, P1c-deficient keratinocytes show aberrations in MT-dependent functions, such as mitotic spindle formation, glucose uptake, FA turnover, and polarized migration. Similarly, P1c-deficient DRG neurons show alterations in glucose uptake, axonal vesicle transport, Na⁺ channel distribution, membrane excitability, and reduced tau solubility. The broad spectrum of MT-dependent cellular processes affected by P1c deficiency makes P1c not only a uniquely versatile plectin isoform variant, but also an essential element in the orchestration and proper functioning of the MT network. It will be a challenging task for future investigations to dissect in detail the mechanism underlying MT destabilization mediated by plectin. Of particular interest will be to investigate whether plectin is involved in tau phosphorylation, and how it regulates NF network assembly and its extension into the growth cone of neurons. Eventually it will be a challenging task to analyze whether plectin-related phenotypes have consequences at the cognitive levels of P1c-deficient mice, and whether P1c-deficiency could prevent tau’s toxic effects by keeping tau attached to MTs, avoiding aggregate formation.
5 MATERIAL AND METHODS

Plasmids

Mammalian expression plasmids encoding GFP-EB1 (Stepanova et al., 2003), EGFP-tubulin and EGFP-zyxin (Rottner et al., 1999), and the bacterial expression plasmid pET3d/MAP2c encoding rat MAP2c (Ludin et al., 1996) were kindly provided by A. Akhmanova (Erasmus Medical Center, Rotterdam, The Netherlands), J. Wehland (Gesellschaft für Biotechnolo-gische Forschung, Braunschweig, Germany), M. Gimona (University of Salzburg, Austria), and A. Matus (Friedrich Miescher Institute, Basel, Switzerland), respectively. Mammalian expression plasmids encoding full-length mouse P1c and P1c-8 with C-terminal EGFP or mCherry tags have been described previously (Rezniczek et al., 2003; Burgstaller et al., 2010). Plectin fragments corresponding to exons 1c-8 (p1c-8, amino acids 1-299, NCBI Reference Sequence: NP_035247); exons 1c-30 (p1c-30, amino acids 1-1374); exons 16-24 (p16-24, amino acids 632-1018); exons 20-21 (p20-21, amino acids 815-889) were excised from existing plasmids (constructed as described in Rezniczek et al., 2003), and inserted into the EcoRI site of the bacterial expression vector pGEX-4T-1 (GST gene fusion system).

Antibodies, antisera and dyes

Table 1. Primary antibodies and dyes used for immunofluorescence microscopy (IFM) and immunoblotting (IB).

<table>
<thead>
<tr>
<th>Antibody/Dye</th>
<th>Antibody type</th>
<th>Antigen/Epitope</th>
<th>Dilution</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>#32</td>
<td>serum, rabbit</td>
<td>purified MAP2 of hog brain</td>
<td>1:500 (IFM) 1:2000 (IB)</td>
<td>G. Wiche</td>
</tr>
<tr>
<td>Acetylated tubulin, T6793, clone 6-11B-1</td>
<td>monoclonal, mouse</td>
<td>Chlamydomonas axonemal acetylated tubulin</td>
<td>1:300 (IFM) 1:1000 (IB)</td>
<td>Sigma-Aldrich®</td>
</tr>
<tr>
<td>Actin, A-2066</td>
<td>polyclonal, rabbit</td>
<td>C-terminal end of actin</td>
<td>1:100 (IFM)</td>
<td>Sigma-Aldrich®</td>
</tr>
<tr>
<td>Antibody/Reagent</td>
<td>Type</td>
<td>Target and Description</td>
<td>Concentration</td>
<td>Source</td>
</tr>
<tr>
<td>------------------</td>
<td>---------------</td>
<td>----------------------------------------------------------------------------------------</td>
<td>---------------</td>
<td>-------------------------------------------</td>
</tr>
<tr>
<td>Actin, A4700, clone AC-40</td>
<td>monoclonal, mouse</td>
<td>C-terminal end of actin</td>
<td>1:200 (IFM)</td>
<td>Sigma-Aldrich®</td>
</tr>
<tr>
<td>Fyn-59, sc-73388</td>
<td>monoclonal, mouse</td>
<td>amino acids 7-176 of human Fyn</td>
<td>1:100 (IFM) 1:2000 (IB)</td>
<td>Santa Cruz Biotechnology, Inc.</td>
</tr>
<tr>
<td>GAPDH, G9545</td>
<td>polyclonal, rabbit</td>
<td>amino acids 314-333 of mouse GAPDH</td>
<td>1:15000 (IB)</td>
<td>Sigma-Aldrich®</td>
</tr>
<tr>
<td>GLUT1, 07-1401</td>
<td>polyclonal, rabbit</td>
<td>C-terminal end of GLUT1</td>
<td>1:100 (IFM)</td>
<td>Millipore</td>
</tr>
<tr>
<td>GFAP (GA5), #3670</td>
<td>monoclonal, mouse</td>
<td>purified GFAP of pig spinal cord</td>
<td>1:300 (IFM) 1:1000 (IB)</td>
<td>Cell signaling technology®</td>
</tr>
<tr>
<td>GST, G1160, clone GST2</td>
<td>monoclonal, mouse</td>
<td>Schistosoma japonicum GST</td>
<td>1:1000 (IB)</td>
<td>Sigma-Aldrich®</td>
</tr>
<tr>
<td>Na⁺ channel α, cardiac (III-IV loop)</td>
<td>polyclonal, rabbit</td>
<td>Na⁺ channel intracellular III-IV loop</td>
<td>1:80 (IFM)</td>
<td>Upstate (Millipore)</td>
</tr>
<tr>
<td>Neurofilament 160 kD, MAB5254, clone NN18</td>
<td>monoclonal, mouse</td>
<td>neurofilament 160 kD</td>
<td>1:50 (IB)</td>
<td>Chemicon® International</td>
</tr>
<tr>
<td>Neurofilament 200kD, clone RT97</td>
<td>monoclonal, rat</td>
<td>neurofilament heavy chain of rat brain</td>
<td>1:10 (IFM)</td>
<td>J. Wood</td>
</tr>
<tr>
<td>Phalloidin conjugated with Texas Red®-X, T-7471</td>
<td>dye</td>
<td>actin</td>
<td>1:100 (IFM)</td>
<td>Molecular Probes</td>
</tr>
<tr>
<td>Plectin 1c</td>
<td>polyclonal, rabbit</td>
<td>N-terminal sequence M₁-S₁₇ of plectin 1c</td>
<td>1:100 (IFM) 1:800 (IB)</td>
<td>G. Wiche</td>
</tr>
<tr>
<td>Plectin #9</td>
<td>serum, rabbit</td>
<td>plectin (exons 9-12)</td>
<td>1:3330 (IB)</td>
<td>B. Nicolic, G. Wiche</td>
</tr>
<tr>
<td>Plectin#10F6</td>
<td>serum, mouse</td>
<td>plectin (rod-domain)</td>
<td>1:2 (IFM)</td>
<td>G. Wiche</td>
</tr>
<tr>
<td>Plectin #46</td>
<td>serum, rabbit</td>
<td>plectin (rod-domain)</td>
<td>1:100 (IFM)</td>
<td>G. Wiche</td>
</tr>
<tr>
<td>Antibody Description</td>
<td>Host Type</td>
<td>Dilution</td>
<td>Source</td>
<td></td>
</tr>
<tr>
<td>--------------------------------------------</td>
<td>------------</td>
<td>----------</td>
<td>---------------------------------------------</td>
<td></td>
</tr>
<tr>
<td>Synaptophysin-Prediluted</td>
<td>polyclonal, rabbit</td>
<td>C-terminus of human synaptophysin 1:5 (IFM) 1:50 (IB)</td>
<td>Invitrogen</td>
<td></td>
</tr>
<tr>
<td>Tau, A-0024</td>
<td>polyclonal, rabbit</td>
<td>amino acids 243-241 of human tau (containing 4R of MT-binding site) 1:100 (IFM) 1:2000 (IB)</td>
<td>Dako</td>
<td></td>
</tr>
<tr>
<td>Tau-1, MAB3420, clone PC1C6</td>
<td>monoclonal, mouse</td>
<td>purified denatured bovine MAPs 1:100 (IFM) 1:2000 (IB)</td>
<td>Chemicon® International</td>
<td></td>
</tr>
<tr>
<td>Tubulin, SM2202P</td>
<td>monoclonal, rat</td>
<td>C-terminal end of yeast α-tubulin 1:300 (IFM)</td>
<td>Acris antibodies</td>
<td></td>
</tr>
<tr>
<td>Tubulin, T5168, clone B-5-1-2</td>
<td>monoclonal, mouse</td>
<td>purified α-tubulin of sea urchin sperm axonemes 1:1000 (IFM) 1:100000 (IB)</td>
<td>Sigma-Aldrich®</td>
<td></td>
</tr>
<tr>
<td>Vinculin, clone VIN-11-5</td>
<td>Monoclonal, mouse</td>
<td>purified vinculin of chicken gizzard smooth muscle 1:50 (IFM)</td>
<td>Sigma-Aldrich®</td>
<td></td>
</tr>
</tbody>
</table>

Table 2. Secondary antibodies used in immunofluorescence microscopy (IFM) and immunoblotting (IB). Horseradish peroxidase (HRPO).
<table>
<thead>
<tr>
<th>Antibody Type</th>
<th>Species</th>
<th>Dilution</th>
<th>Supplier</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cy5 anti-rabbit IgGs</td>
<td>donkey</td>
<td>1:300 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Cy5 anti-rat IgGs</td>
<td>donkey</td>
<td>1:300 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>DyLight 649 anti-mouse IgGs</td>
<td>donkey</td>
<td>1:500 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>DyLight 649 anti-rabbit IgGs</td>
<td>donkey</td>
<td>1:500 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>DyLight 649 anti-rat IgGs</td>
<td>donkey</td>
<td>1:500 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>HRPO anti-mouse IgM (µ chain)</td>
<td>donkey</td>
<td>1:10000 (IB)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>HRPO anti-mouse IgGs</td>
<td>goat</td>
<td>1:10000 (IB)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>HRPO anti-rabbit IgGs</td>
<td>goat</td>
<td>1:20000 (IB)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Rhodamine red anti-mouse IgGs</td>
<td>donkey</td>
<td>1:200 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Rhodamine red anti-rabbit IgGs</td>
<td>donkey</td>
<td>1:100 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Texas red anti-mouse IgGs</td>
<td>goat</td>
<td>1:200 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Texas red anti-mouse IgGs</td>
<td>donkey</td>
<td>1:200 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Texas red anti-rabbit IgGs</td>
<td>donkey</td>
<td>1:200 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
<tr>
<td>Texas red anti-rat IgGs</td>
<td>goat</td>
<td>1:200 (IFM)</td>
<td>Jackson ImmunoResearch Laboratories, Inc.</td>
</tr>
</tbody>
</table>

**DNA methods**

5.1.1 Preparation of plasmid DNA

Plasmid DNA purification from bacteria was carried out using anion exchange JetStar™ resin
columns according to manufacturer’s instructions (Genomed). Endotoxin-free plasmid DNA suitable for transfection of primary cells was isolated from bacteria using JetStar™ Endotoxin-free Plasmid kit.

### 5.1.2 Determination of DNA concentration

DNA concentration was calculated according to its optical density. Spectrophotometric measurements were performed at 260 nm. An optical density of 1 corresponds to a concentration of 50 µg/ml of double-stranded DNA.

### Protein methods

#### 5.1.3 Determination of concentration

Protein concentration was determined by colorimetric detection of the cuprous cation (Cu^{1+}) by bicinechninic acid using the BCA™ Protein Assay kit (Pierce) according to manufacturer’s instructions.

#### 5.1.4 SDS-Polyacrilamide gel electrophoresis (PAGE)

Proteins were separated using standard SDS-6%, -10% or -12% PAGE (Laemmli, 1970). Gels were prepared using the Mini-PROTEAN® Electrophoresis System (Bio-Rad). First, the cassette was filled with resolving gel solution, overlaid with isopropanol and allowed to polymerize. Then, the stacking gel was poured on the top of the resolving gel, and a comb was inserted to allow sample loading. Samples were boiled at 95°C during 5 min prior to loading into the gel slots. Gels were run at 25 mAmp/gel and 500 V.

Buffers and solutions:

- 5x SDS Laemmli sample buffer:
  - Tris-HCl, pH 6.8 --------------------------- 125 mM
  - DDT ------------------------------------------ 250 mM
  - SDS ------------------------------------------ 5%
5.1.5 Coomassie staining

Gels were stained after electrophoresis by incubation with Coomassie staining solution. After adding the solution, gels were warm up in the microwave for 10 s and incubated at room temperature for 15 min. Then, gels were destained with destaining solution during 1 h at room temperature or at 4°C overnight.

Solutions:

- Coomassie staining solution:
  - Coomassie R-250 (Sigma-Aldrich) ----- 0.05% (w/v)
  - Methanol------------------------------- 40%
  - Acetic acid -------------------------- 7%
  - ddH₂O ------------------------------- 53%

- Destaining solution:
  - Acetic acid -------------------------- 10%
  - Methanol ---------------------------- 30%
  - ddH₂O ------------------------------- 60%

5.1.6 Immunoblotting

After electrophoresis, gels were electrophoretically transferred onto nitrocellulose membranes at 25 V, 4°C, overnight, or 100 V, 4°C for 2 h. Then, the transfer efficiency was tested by staining with Ponceau S (Sigma-Aldrich) and washed with ddH₂O. Membranes were blocked
for 30 min, incubated with primary antibodies diluted in PBST at room temperature for 1h or at
4°C overnight, wash 2 x 15 min with PBST, incubated with secondary antibodies diluted in
PBST at room temperature for 1h and washed 2 x 15 min with PBST. Detection was carried
out using the SuperSignal West Pico Chemiluminescent Substrate (Pierce) and exposing the
membranes to an X-ray sensitive film (Fuji).

Buffers and solutions:

- **Blot Buffer:**
  - Tris 48 mM
  - Glycine 40 mM
  - SDS 0.1%

- **PBS (phosphate buffered saline):**
  - NaCl 8.18 g
  - KCl 0.201 g
  - KH$_2$PO$_4$ 0.204 g
  - Na$_2$H$_2$PO$_4$ 1.132 g
  - pH 7.4
  - ddH$_2$O up to 1000 ml

- **PBST:**
  - PBS containing 0.1% Tween-20

- **Ponceau S staining solution:**
  - Ponceau S (Sigma-Aldrich) 0.5% (w/v)
  - Acetic acid 1%
  - ddH$_2$O

- **Blocking solution:**
  - 5% non-fat dry milk in PBST

### 5.1.7 Quantification of protein bands on immunoblots

Densitometrical quantification of protein bands was performed after scanning the X-ray films
on a flat-bed scanner using QuantiScan v1.5 software (Biosoft, Cambridge, UK).
5.1.8 Preparation of total cell and tissue lysates

Tissues were snap frozen in liquid nitrogen, pulverized and suspended in 1% Triton X-100, 0.5% NP-40, 150 mM NaCl, 20 mM Tris-HCl, pH7.5 supplemented with Complete Mini Protease Inhibitor Cocktail tablets (CMPI) (Roche). Cell lysates were prepared from confluent cultures and lyse in 1x PBS, 0.5% Triton X-100, 1 mM PMSF, 2 mM DTT.

5.1.9 Endogenous MT-binding assay

The protocol of this assay was based on a similar one described by Planel et al, 2008; except where indicated otherwise, all working steps were carried out above 20°C to prevent MT depolymerization. Brains isolated from mice were homogenized in 80 mM MES, pH 6.8, 1 mM MgCl₂, 2 mM EGTA, 30% glycerol, 0.1% Triton X-100, 1 mM PMSF, 1 mM Na₃VO₄, 1 mM NaF, and CMPI (Roche) using a mechanical tissue homogenizer. Lysates were centrifuged for 6 min at 16,000 x g to obtain supernatant fraction containing intact MTs. The MT fraction was centrifuged (20 min at 25°C) in a vacuum Optima™ TLX Ultracentrifuge (Beckman) at 40,000 rpm (100,000 x g) to obtain a MT-free supernatant and a MT-containing pellet fraction. MTs in the pellet fraction were resuspend in 100 mM MES pH 6.5, 0.5 mM MgSO₄, 30% glycerol, 1 mM EGTA, 0.1% Triton X-100, 1 mM Na₃VO₄, 1 mM NaF, 2 mM DTT, 1 mM PMSF, and CMPI (Roche). Samples of both fractions were mixed with O+ buffer (O’Farrel, 1975) supplemented with CMPI (Roche) and boiled 5 min prior to immunoblotting. Tau levels were analyzed by immunoblotting MT-free and MT-containing fractions.

5.1.10 RhoA pull down assay

pGEX-2T-RBD culture

100 ml LB medium were inoculated with pGEX-2T-RBD (GST-RBD) bacterial stock (DH5α), and let it grow at 37°C overnight. GST-RBD fusion protein encodes the Rho-binding domain of
Rhotekin, used as an activation specific probe for RhoA-GTP.

- **LB medium (Luria-Bertani medium):**
  - Bacto-tryptone: 1% (w/v)
  - Bacto-yeast extract: 0.5% (w/v)
  - NaCl: 1% (w/v)
  - pH 7.5
  - ddH₂O up to 1000 ml

**pGEX-2T-RBD: Growth and Induction**

300 ml LB medium were inoculated with 400 µl of overnight bacterial stock supplemented with corresponding antibiotics and 15 ml glucose and incubated at 37°C until the culture reaches OD₆₀₀ of 0.7 to 0.8. Protein expression was induced by adding 0.5 mM IPTG and culture was incubated at 30°C for 2-4 h. Cells were harvested by centrifugation at 10000 x g (Heraeus centrifuge) during 15 min at 4°C. Pellets were shock freeze in liquid nitrogen and store at -80°C.

**pGEX-2T-RBD: Cell lysis**

Bacteria pellets were resuspended in 4.5 ml ice-cold STE-Buffer supplemented with 1 mM PMSF and homogenized using a 19G needle. Suspensions were supplemented with lysozime, 5 mM DTT, 1% Tween-20, 0.03% SDS and centrifuged at 16000 xg for 20 min at 4°C.

- **STE Buffer:**
  - Tris, pH 8.0: 10 mM
  - NaCl: 150 mM
  - EDTA: 10 mM EDTA
Purification of GST-Rhotekin-BD (GST-RBD)

Glutathione Sepharose™ 4B (GE, Healthcare) is supplied as 75%, in order to prepare 50% slurry the bottle of Glutathione Sepharose 4B slurry was gently shaken, 399 µl of 75% slurry for 300 µl desired volume were taken and centrifuged at 500 x g for 5 min at room temperature. Supernatant was discarded and 1 ml STE Buffer was added to remove ethanol after 2x centrifugation at 500 x g for 5 min at room temperature. 5ml from the previously obtained lysates was added to the 300 µl 50% slurry and mixed gently for 30 min at 4°C. Then the mixed was washed 3 x with 5ml ice-cold STE Buffer by centrifuging at 500 x g for 5 min at 4°C. Pellets were resuspended in 120 µl of STE buffer, aliquoted into 30-35 µl and stored at -80°C until use on pull down assay.

RhoA pull down assay

Brain was lysed in 500 µl ice-cold pull down lysis buffer, transferred to pre-chilled eppendorf tubes and centrifuge at 14000 x g for 5 min at 4°C. Pre-cleared lysates were transferred into new pre-chilled tubes with 30 -35 µl of GST-RBD (30-45 µg) and incubated for 1 h at 4°C. Then, the samples were spun down at 700xg for 3 min at 4°C and washed 2x with 600 µl ice-cold 1 x Mg²⁺Buffer. RhoA was eluted from beads with 45 µl of 2x SDS sample buffer and boiled for 5 min at 95°C. Samples were loaded on SDS 15%-PAGE gels.

- Pull down lysis buffer:
  - Heps/HCl 25 mM
  - NaCl 150 mM
  - MgCl₂ 10 mM
  - NP-40 1%
  - PMSF 100 µM
  - CMPI (Roche)
  - ddH₂O
5.1.11 Extraction of sarcosyl-insoluble tau

Extraction of sarcosyl-insoluble tau was carried similarly as described (Greenberg & Davies, 1990). In brief, the brain tissue was homogenized in 3 volumes of cold Buffer H and centrifuged at 27000 g for 20 min at 4°C (21000 rpm with TLA-45 rotor). The supernatant was collected and the resulting pellet was homogenized in buffer H. The homogenization was centrifuged at 27000 g for 20 min at 4°C (21000 rpm with TLA-45 rotor). The supernatant obtained was collected and combined with the one resulting from the first centrifugation. N-lauroylsarcosine was added to the supernatant at 1% (w and/v) final concentration and then incubated at 37°C shaking for 60-90 min. The samples were centrifuged at 150000g for 35 min at 20°C (58800 rpm TLA 120.2 rotor) and the resulting pellets were resuspended in 50 mM Tris-HCl, pH 7.4, using 0.5 µl of buffer per mg of initial weight. The samples were diluted in SDS sample buffer, boiled at 95°C for 5 min and loaded in SDS-15% gels at a ration 1:2 (soluble: insoluble). Sarcosyl-insoluble tau was evaluated after immunoblotting using tau antibodies.

- **Buffer H:**
  - Tris-HCl ------------------------------ 10 mM
  - EGTA ---------------------------------- 1 mM
  - NaCl ---------------------------------- 0.8 M
  - Sucrose ------------------------------- 10%
  - pH = 7.4

- 1x Mg$^{2+}$ Buffer:
  - NaF -------------------------------------- 25 mM
  - Aprotinine ---------------------------------- 10 µg/ml
  - Benzamidine ---------------------------------- 10 µg/ml
  - Glycerol ---------------------------------- 10%
  - PMSF ---------------------------------- 100 µM
  - ddH$_2$O

- **Buffer H:**
  - Tris-HCl ---------------------------------- 10 mM
  - EGTA ---------------------------------- 1 mM
  - NaCl ---------------------------------- 0.8 M
  - Sucrose ---------------------------------- 10%
  - pH = 7.4

- 1x Mg$^{2+}$ Buffer:
Mammalian cell culture

5.1.12 Cultivation of p53⁻/⁻ keratinocytes

Immortalized plectin wt and P0 mouse keratinocytes cell cultures were obtained from plectin wt/p53⁻/⁻ and P0/p53⁻/⁻ mice (Andrä et al., 2003). Wt and P0 p53⁻/⁻ keratinocytes were cultured on 100 µg/ml bovine collagen I (Sigma-Aldrich®)-coated plastic dishes (Nunc) using keratinocyte growth medium (KGM). Cultivation was done at 37°C in a humidified atmosphere of 5% CO₂. Cell were let grow until 80-90% confluency before splitting. Splitting of cells was carried out using strong trypsin solution for 8-10 min at 37°C. Then, cells were spun down at 200 xg for 3 min at room temperature. Supernatant was discarded, cell pellet resuspended in KGM and plated.

- Keratinocyte growth medium:
  - KBM™ w/o Ca²⁺ (Lonza) --------------- 500 ml
  - KGM™ BulletKit™ w/o Ca²⁺ (Lonza):
    - BPE (bovine pituitary extract) --------- 2ml
    - hEGF (human epidermal growth factor) --- 0.5 ml
    - Insulin (recombinant human) -------------- 0.5 ml
    - Hydrocortisone ------------------------ 0.5 ml
    - GA-1000 (gentamicin, amphotericin B) ---- 0.5 ml
  - Chelex-treated FCS (Gibco®) --------------- 2%
  - ITS (Insulin-Transferrin-Selenium) (Gibco®) ------- 1%
  - CaCl₂ --------------------------------------- 0.05 mM

- Chelex- (Gibco®) treated fetal calf serum (FCS) (Sigma-Aldrich):
  - 20 g of Chelex to 200 ml ddH₂O
  - adjust pH to 7.4 overnight
  - place Chelex for 1h at 4°C to form a compact pellet
  - add Chelex to 50 ml FCS
  - stir for 1h at 4°C
  - place the mixture for 1h at 4°C to form a compact Chelex pellet
  - decant serum through a bottle top filter into an autoclaved glass bottle
5.1.13 Isolation of primary keratinocytes

Primary keratinocytes were isolated from newborn mice skin or adult tail skin. Skins were flatten and put dermis-side down on Dispase II solution (Roche) for 1h at 37°C or on strong trypsin solution at 4°C overnight. After incubation, epidermis was separated from dermis and minced into small pieces with two scalpels. Cells were dissociated from tissue by incubation in 3 ml strong trypsin solution under shaking conditions at 37°C for 8 min (newborn mices) or 6 min (adult mice). Epidermal pieces were resuspended by pipetting up and down 60 times and then filtered through a 70 µm mesh of a cell strainer attached to a 50 ml tube filled with 40 ml of KGM at 4°C to remove pieces of stratum corneum. Suspension was centrifuged 150 g for 7 min at room temperature. Supernatant was discarded, cell pellet was resuspend in KGM-Gold medium and cell suspension was plated on collagen-I-coated dishes. Primary keratinocytes cultivation was performed as described previously.

- Dispase II solution (Roche):
  - 10 mg/ml of dispase II (Roche) in KBM™ (CC-3112, Lonza)

- Keratinocyte growth medium (KGM)-Gold:
  - KBM-Gold™ w/o Ca²⁺(Lonza) --------------------------- 500 ml
  - KGM-Gold™ BulletKit™ (CC-3111, Lonza):
    - BPE (bovine pituitary extract) -------------- 2ml
    - hEGF (human epidermal growth factor) --- 0.5 ml
    - Insulin (recombinant human) --------------- 0.5 ml
    - Hydrocortisone --------------------------- 0.5 ml
    - GA-1000 (gentamicin, amphotericin B) ----- 0.5 ml
    - Epinephrin -------------------------- 0.25 ml
    - Transferrin ----------------------------- 0.5 ml
5.1.14 Isolation and cultivation of DRG neurons

DRG neuron isolation was performed similarly as described by Stroissnigg et al, 2007. DRGs of adult mice were dissected and harvested in ice-cold RPMI 1640 (Gibco) medium, cut into small pieces before enzymatic dissociation by 4000 U/ml collagenase solution (SERVA Electrophoresis, GmbH) for 90 min at 37°C followed by treatment with normal trypsin solution supplemented with DNase I (Sigma-Aldrich) for 7-15 min at 37 °C. Trypsinization was inhibited adding stop solution, and finally the DRGs were triturated several times through a narrowed Pasteur pipette. Suspension was spun down at 120xg 5 min, resuspended in growth medium and plated onto precoated coverslips with poly-L-lysine (10 µg/ml in ddH2O; Sigma-Aldrich) overnight at 37 °C, followed by laminin (10 µg/ml in PBS; at least 3 h at 37 °C; Sigma-Aldrich). DRG neurons were cultivated at 37°C in a humidified atmosphere of 5% CO2.

Media and solutions:

- Collagenase solution (1 ml/mice):
  - 200 µl collagenase (4000 U/ml, Sigma-Aldrich)
  - 20 µl horse serum (Gibco)
  - 780 µl RPMI 1640 (Gibco)

- Trypsin solution (1 ml/mice):
  - 500 µl normal trypsin solution
  - 20 µl DNAse I (Roche)

- Stop solution (2 ml/mice):
  - 400 µl horse serum (Gibco)
  - 1.6 ml RPMI 1640 (Gibco)
• Growth medium (2 ml/mice):
  o 10 µl penicillin/streptomycin (5000 U/ml-5000 µg/ml, Gibco®)
  o 100 µl horse serum (Gibco®)
  o 35.5 µl glucose (50%)
  o 35.5 µl N3 mix
  o 1840 µl RPMI 1640 Gibco®

• Normal trypsin solution:
  o 0.05% Trypsin (Gibco®)
  o 0.2 mg/mol EDTA
  o HBSS (Gibco®)

• 57x N3 mix (Bottenstein and Sato, 1979):
  o 1 mg/ml BSA fraction V in HBSS (PAA)
  o 10 mg/ml apotransferrin in HBSS (T1147, Sigma-Aldrich®)
  o 1 µg/ml Na-Selenit in HBSS (S9133, T1147, Sigma-Aldrich®)
  o 3.2 mg/ml putrescine in HBSS (P5780, Sigma-Aldrich®)
  o 1.25 µg/ml progesterone in EtOH (P6149, Sigma-Aldrich®)
  o 4 µg/ml corticosterone in EtOH (C2505, Sigma-Aldrich®)
  o 2 µg/ml tri-iodothyronine in 0.01% NaOH (T6397, Sigma-Aldrich®)
  o 1 mg/ml insulin (I6634, Sigma-Aldrich®)
  o HBSS - Ca²⁺ - Mg²⁺ (Gibco®)

5.1.15 Freezing and thawing of cells

Cells were frozen in freezing solution containing 90% FCS (Sigma-Aldrich) and 10% DMSO, kept at -80°C for 24h and then transfer to liquid nitrogen for long-term storage. Thawing of cells were done diluting the frozen aliquots in 15 ml of growth medium, spinning down at 200 xg for 3 min, resuspending the cell pellets in growth medium and plating the cells in appropriate medium.

5.1.16 Transient transfection

_FuGene6 (Roche)_

FuGEnE6 Transfection Reagent (Roche) is a lipid-base transfection reagent used to transfect cell lines. For transfecting 35 mm dishes, 200 µl of medium without serum or growth factors
were added into polysterene tubes (do not use polypropylene tubes). 8 µl of room temperature pre-warmed FuGENE6 reagent was added directly into the medium and tap gently to mix. Then, 5 µg of DNA were added and tap gently again to mix the contents. The mixture was incubated for 30 min at room temperature. After incubation, 300 µl of serum-free medium were added and the solution was dropwise added onto the cells. Cells were plated at allowed to grow to 60-70% confluency in the presence of growth medium.

**Nanofectin (PAA)**

Nanofectin (PAA) consists of a positively charged polymer with a high DNA-binding capacity, which is embedded into a porous nanoparticle that is optimized for the endocytosis machinery of the cell. Transfection of cell lines was done following manufacture’s instructions. DNA and nanofecting diluted in Diluent (1:50) were mixed in a 1:32 proportion. The mixture was incubated for 15 to 30 min at room temperature and added drop wise onto the cells. Prior transfection cells were plated and allowed to grow to 70-80% confluency in the presence of growth medium.

**Nucleofector™ Technology (Lonza)**

Nucleofector™ Technology (Lonza) is especially designed for primary cell transfection. It is a non-viral method, which is based on a combination of electrical parameters and cell-type specific solutions.

**Amaxa® human keratinocyte Nucleofector® kit**

Primary keratinocytes were transiently transfected using the Amaxa® human keratinocyte Nucleofector® kit similarly to manufacturer’s instructions. 7x10^5 cells per transfection were resuspended in 100 µl of human keratinocyte Nucleofector® solution and combined with 4-5 µg of endotoxin-free DNA. The cell/DNA suspension was transferred to a cuvette avoiding cell bubbles and the cuvette was inserted into the Nucleofector® cuvette holder. Program T-024 for high transfection efficiency was applied and immediately after transfection 500 µl of growth medium was added. The transfected cell suspension was gently plated onto plastic dishes.
Amaxa® basic neuron SCN Nucleofector® kit

Primary DRG neurons were transiently transfected using the Amaxa® basic neuron SCN Nucleofector® kit similarly to manufacturer’s instructions. 2x10^4 cells per transfection were resuspended in 20 µl of basic neuron SCN Nucleofector® solution and combined with 0.5-1 µg of endotoxin-free DNA. The cell/DNA suspension was transferred to a cuvette avoiding cell bubbles and the cuvette was inserted into the Nucleofector® cuvette holder. Program SCN basic neuro program 6 was applied and immediately after transfection 500 µl of growth medium was added. The cuvette containing the cell suspension was placed inside an incubator to avoid high mortality. After 5-10 min incubation the transfected cell suspension was gently plated onto coverslips.

5.1.17 Nocodazole and colchicine treatment

To assess MT stability, cells were kept in culture medium supplemented with 1 µM nocodazole in DMSO (Sigma-Aldrich) or 10 µM colchicine in DMSO (Sigma-Aldrich) for 30 min in the case of keratinocytes or 3 min in the case of DRG neurons, before being fixed and processed for microscopy. Immunolabeled single MTs remaining in nocodazole-treated cells were traced, their length measured using LSM software (Zeiss), and normalized to total cell area.

5.1.18 Oxidative stress

Oxidative stress of DRG neurons was tested treating primary DRG neurons with 250 µM H2O2 for 5 min before being fixed and processed for microscopy. After visualization by confocal microscopy, oxidative stress was quantified as the proportion of collapse growth cones (bulb-shaped growth cones) to the total number of growth cones and comparing the results obtained from wt and P1c⁻ DRG neurons.

5.1.19 DRG explants

DRGs of adult mice were dissected and harvested in ice-cold RPMI 1640 (Gibco®) medium and cut in half. Half-cut DRG explants were placed on 5-10 µl drops of ice-cold matrigel (BD-Science) onto uncoated coverslips and completely covered with 10 µl matrigel drop.
5.1.20 2-NBDG uptake

Primary mouse keratinocytes and DRG neurons were incubated with 600 µM 2-NBDG (Invitrogen) for 15 min following the protocol described by Yamada et al, 2000. Fluorescence intensities of 2-NBDG were collected using a confocal microscope and after background subtraction, fluorescence intensity was measured using ImageJ software (NIH Image, Bethesda, MD).

5.1.21 Na⁺ uptake

Na⁺ uptake of primary DRG neurons was measured using CoroNa Green Na⁺ Indicator (Invitrogen). CoroNa Green dye is a Na⁺ indicator that exhibits an increase in fluorescence intensity upon binding Na⁺ (excitation/emission = 492/516 nm). Cells were loaded by adding 1 µM CoroNa Green Na⁺ in DMSO (Invitrogen), incubated for 10 min at 37°C and washed with dye-free medium. Fluorescence intensities of CoroNa Green Na⁺ Indicator (Invitrogen) were collected using a confocal microscope and after background subtraction, and measured using ImageJ software (NIH Image, Bethesda, MD).

5.1.22 Synaptic vesicle exocytosis and endocytosis with FM dyes

Visualization of synaptic vesicles exocytosis and endocytosis was performed similarly as described by Gaffield & Betz (2006). Cells were treated with 15 µM FM1-43 (Invitrogen) diluted in high K⁺ Ringer solution to trigger synaptic vesicle recycling for 10 min at 37°C and washed 3 x 5 min normal Ringer solution before performing time-lapse microscopy.

- High K⁺ Ringer solution:
  - NaCl ----------------------------------------- 31.5 mM
  - KCl ----------------------------------------- 90 mM
  - CaCl₂ --------------------------------------- 2 mM
  - MgCl₂ --------------------------------------- 2 mM
  - Glucose ------------------------------------- 30 mM
  - Hepes -------------------------------------- 25 mM
  - pH 7.3

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• Normal Ringer solution:
  o NaCl --------------------------------------------- 119 mM
  o KCl --------------------------------------------- 2.5 mM
  o CaCl₂ ------------------------------------------ 2 mM
  o MgCl₂ ------------------------------------------ 2 mM
  o Glucose ---------------------------------------- 30 mM
  o HEPES ----------------------------------------- 25 mM
  o pH 7.3

5.1.23 Current clamp electrophysiology

Electrophysiology experiments were performed by A. Yousuf and K. Schicker at S. Böhm laboratory (Centre for Physiology and Pharmacology, Medical University of Vienna). DRG neuronal current potentials were recorded by current clamp technique at room temperature after 1 to 2 days in culture. Measurements were carried out using Axopatch 200B amplifier and the pCLAP 8.1 hard- and software (Molecular Devices, Sunnyvale, CA) as described by A. Yousuf et al, 2011.

Microscopy

5.1.24 Immunolabeling for immunofluorescence microscopy

Keratinocyte cell cultures were rinsed (1 min) with pre-warmed (37°C) 60 mM PIPES, pH 6.9, 25 mM HEPES, 10 mM EGTA, and 2 mM MgCl₂ (MT-stabilizing solution). Cells were then fixed with 2.5% PFA, quenched with 0.1 M glycine, permeabilized with 0.1% Triton X-100, and block with 5% BSA in PBS for 1h at room temperature. After blocking cells were immunolabeled incubating with primary antibodies previously described diluted in 3% BSA in PBS for 1h at room temperature. Then, cells were washed 3 x 10 min with PBS and incubated with corresponging secondary antibodies diluted in PBS for 1h at room temperature. After 3 x 10 min washing with PBS, cells were mounted in mowiol 4-88 (Hoechst).
Primary DRG neurons were fixed with pre-warmed (37°C) 4% PFA in 22% sucrose solution added directly to growth medium for 45 min at room temperature. Cells were then washed 3 x 5 min with PBS and blocked with 10% BSA, 0.3% Triton X-100 for 1h at room temperature. After blocking cells were incubated with primary antibody diluted in 5% BSA, 0.15% Triton X-100 for 3h at room temperature or overnight at 4°C and then washed 3 x 5 min with PBS. Cells were then incubated with secondary antibody diluted in PBS for 2h at room temperature, washed 3 x 5 min with PBS and mount in mowiol 4-88 (Hoechst).

5.1.25 Confocal microscopy
Microscopy was performed at room temperature using a confocal microscope (Zeiss, LSM 510) equipped with a Plan-Apochromat 63x 1.4 NA or 100x 1.4 NA objective lens. Images were recorded using the LSM 510 module and the LSM software.

5.1.26 Wide-field microscopy
Wide-field microscopy was performed at room temperature using a DeltaVision Image Restoration microscope system (Applied Precision Instruments, LLC, Issaquah, WA, USA), equipped with a Plan-Apochromat 60x1.4 NA objective lens or using an AxioObserver Z1 microscope coupled to AxioCam MRm (Carl Zeiss MicrolImaging) equipped with phase contrast optics. Images were acquired with softWoRx software (Applied Precision Instruments) and Zeiss AxioVision 4.8.1. software respectively.

5.1.27 Deconvolution
Acquired pictures were deconvolved using 3D Huygens Deconvolution & Analysis software (Scientific Volume Imaging) or using softWoRx software (Applied Precision Instruments).
5.1.28 Time-lapse video microscopy

Visualization of MT dynamics

Cells expressing GFP-EB1 or EGFP-tubulin were kept in a closed POCmini cultivation system (Zeiss), and live-cell imaging was performed using an inverted microscope (Zeiss, Axiovert S100TV) at 37°C and in a humidified atmosphere of 5% CO₂. Frames of GFP-EB1 and EGFP-tubulin were collected with a Plan-Apochromat 100x 1.4 NA objective lens every 2 s during 5 min. Individual comets or MTs were traced using Metamorph 6.3 software (MDS Analytical Technologies).

Visualization of FA dynamics

Cells expressing EGFP-zyxin were kept in a closed POCmini cultivation system (Zeiss), and live-cell imaging was performed using an inverted microscope (Zeiss, Axiovert S100TV) at 37°C and in a humidified atmosphere of 5% CO₂. Frames of EGFP-zyxin images were acquired every 2 min during 30 min. Individual FAs were tracked using Metamorph 6.3 software (MDS Analytical Technologies).

Single cell migration

Migrating keratinocytes were recorded using an AxioObserver Z1 microscope coupled to AxioCam MRm (Carl Zeiss MicroImaging) equipped with phase contrast optics at 37°C and humidified atmosphere of 5% CO₂. Frames were taken with an EC Plan-Neofluar 10x/0.3NA objective lens in 7 min intervals over a period of 12 h. Images were processed with Zeiss AxioVision 4.8.1 image analysis software and further analyzed with ImageJ software (NIH Image, Bethesda, MD) for manual tracking of migrating cells. To track the whole cell trajectory, cell nuclei were marked for each frame throughout the entire time-lapse sequence.
Visualization of synaptic vesicles

Cells stained with FM1-43 (Invitrogen) were kept in a closed POCmini cultivation system (Zeiss), and live-cell imaging was performed using an inverted microscope (Zeiss, Axiovert S100TV) at 37°C and in a humidified atmosphere of 5% CO₂. Frames of FM1-43 images were acquired every 2 s during 15 min. Individual synaptic vesicles were tracked using Metamorph 6.3 software (MDS Analytical Technologies).

Growth cone extension

Extending growth cones were recorded using an AxioObserver Z1 microscope coupled to AxioCam MRm (Carl Zeiss MicroImaging) equipped with phase contrast optics at 37°C and humidified atmosphere of 5% CO₂. Frames were taken with a LD A Plan 32x/0.4 NA objective lens in 10 min intervals over periods from 5 h to 12h. Images were processed with Zeiss AxioVision 4.8.1 image analysis software and further analyzed with ImageJ software (NIH Image, Bethesda, MD) for manual tracking of migrating cells.

5.1.29 Image processing and semiquantification

Images were subjected equally across the entire image to minimum degree of Gaussian filter or histogram stretch necessary to facilitate visualization using ImageJ software (NIH Image, Bethesda, MD), LSM software (Zeiss), softWoRx software (Applied Precision Instruments) or AxioVision 4.8.1. software (Zeiss).

The proportion of acetylated to total tubulin in MTs was measured in double labeled specimens by selecting acetylated tubulin-positive and tubulin-positive areas (in pixels) using the magic wand toll in Adobe® Photoshop® CS2 and keeping the tolerance constant. Immunolabeled GLUT1 and Na⁺ clusters were counted and measured using ImageJ software (NIH Image, Bethesda, MD) and values normalized to cell area after GLUT1 immunolabeling.
Statistical analyses

Comparisons between values of two groups were made using Student’s t-test, Wilcoxon-Mann-Whitney U-test or Chi-square test ($\alpha=0.001-0.10$). Comparisons among values of multiple groups were performed using one-way analysis of variance (ANOVA) (alpha=$0.001-0.10$). The significance between the individual groups was subsequently determined using the Tukey post-hoc test ($\alpha=0.05$). Analyses were performed using SPSS Statistics v.19 (IBM®).
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<th>Full Form</th>
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<tbody>
<tr>
<td>2-NBDG</td>
<td>2-[N-(7-nitrobenz-2-oxa-1,3-diazol-4-yl) amino]-2-deoxy-D-glucose</td>
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<tr>
<td>ABD</td>
<td>actin-binding domain</td>
</tr>
<tr>
<td>ACF7</td>
<td>microtubule-actin crosslinking factor 7</td>
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<tr>
<td>AIS</td>
<td>axon initial segment</td>
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<tr>
<td>BCA</td>
<td>bicinchoninic acid</td>
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<td>BPAG1</td>
<td>bullous pemphigoid antigen 1</td>
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<td>cAMP</td>
<td>cyclic adenosine monophosphate</td>
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<td>cdc2</td>
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</tr>
<tr>
<td>CH</td>
<td>calponin-homology domain</td>
</tr>
<tr>
<td>CMPI</td>
<td>complete mini protease inhibitor</td>
</tr>
<tr>
<td>DDT</td>
<td>dithiothreitole</td>
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<td>dimethyl sulfoxide</td>
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<tr>
<td>DNA</td>
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<tr>
<td>DRG</td>
<td>dorsal root ganglia</td>
</tr>
<tr>
<td>EB1</td>
<td>end-binding protein 1</td>
</tr>
<tr>
<td>EBS</td>
<td>epidermolysis bullosa simplex</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Description</td>
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<tr>
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<td>------------------------------------------------------------------</td>
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<tr>
<td>EBS-MD</td>
<td>epidermolysis bullosa simplex with muscular dystrophy</td>
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<tr>
<td>EGFP</td>
<td>enhanced green fluorescent protein</td>
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<tr>
<td>GC</td>
<td>globular carboxi-terminal domain</td>
</tr>
<tr>
<td>GFAP</td>
<td>glial fibrillary acidic protein</td>
</tr>
<tr>
<td>GFP</td>
<td>green fluorescence protein</td>
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<tr>
<td>GLUT1</td>
<td>glucose transporter 1</td>
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<tr>
<td>GLUT3</td>
<td>glucose transporter 3</td>
</tr>
<tr>
<td>GN</td>
<td>globular amino-terminal domain</td>
</tr>
<tr>
<td>GST</td>
<td>glutathione S-transferase</td>
</tr>
<tr>
<td>GTP</td>
<td>guanosine-5’-triphosphate</td>
</tr>
<tr>
<td>hEGF</td>
<td>human epidermal growth factor</td>
</tr>
<tr>
<td>EDTA</td>
<td>ethylenediaminetetraacetic acid</td>
</tr>
<tr>
<td>EGTA</td>
<td>ethylene glycol tetraacetic acid</td>
</tr>
<tr>
<td>ER</td>
<td>endoplasmic reticulum</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Definition</td>
</tr>
<tr>
<td>--------------</td>
<td>------------------------------------------------</td>
</tr>
<tr>
<td>FCS</td>
<td>fetal calf serum</td>
</tr>
<tr>
<td>HBSS</td>
<td>Hank’s buffered salt solution</td>
</tr>
<tr>
<td>HD</td>
<td>hemidesmosome</td>
</tr>
<tr>
<td>Hepes</td>
<td>4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid</td>
</tr>
<tr>
<td>HMW</td>
<td>high molecular weight</td>
</tr>
<tr>
<td>HS-SN</td>
<td>high-speed supernatant</td>
</tr>
<tr>
<td>HS-P</td>
<td>high-speed pellet</td>
</tr>
<tr>
<td>IB</td>
<td>immunoblotting</td>
</tr>
<tr>
<td>IF</td>
<td>intermediate filament</td>
</tr>
<tr>
<td>IFM</td>
<td>immunofluorescence microscopy</td>
</tr>
<tr>
<td>IPTG</td>
<td>isopropyl β-D-1-thiogalactopyranoside</td>
</tr>
<tr>
<td>ITS</td>
<td>insulin-transferrin-selenium</td>
</tr>
<tr>
<td>KBM</td>
<td>keratinocyte basal medium</td>
</tr>
<tr>
<td>KGM</td>
<td>keratinocyte growth medium</td>
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<tr>
<td>LB</td>
<td>Luria-Bertani</td>
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<tr>
<td>MACF</td>
<td>microtubule-actin crosslinking factor 7</td>
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<tr>
<td>MAP</td>
<td>microtubule associated protein</td>
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<tr>
<td>MCK</td>
<td>muscle creatine kinase</td>
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<tr>
<td>MES</td>
<td>2-(N-morpholino)-ethanesulfonic acid</td>
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<tr>
<td>MT</td>
<td>microtubule</td>
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<td>Abbreviation</td>
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<td>--------------------------------------------------</td>
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<tr>
<td>NF</td>
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<td>NP-40</td>
<td>nonyl phenoxypolyethoxylethanol 40</td>
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<tr>
<td>RPMI</td>
<td>Roswell Park Memorial Institute</td>
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<tr>
<td>SDS</td>
<td>sodium dodecyl sulfate</td>
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<td>SDS-PAGE</td>
<td>SDS-polyacrylamide-gel electrophoresis</td>
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<tr>
<td>SH</td>
<td>Src homology domain</td>
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<tr>
<td>SR</td>
<td>spectrin repeat</td>
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<tr>
<td>STE</td>
<td>sodium chloride-tris-EDTA</td>
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<td>P</td>
<td>pellet</td>
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<tr>
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<td>plectin 1c</td>
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<td>plectin 1c-deficiency</td>
</tr>
<tr>
<td>P1d</td>
<td>plectin 1d</td>
</tr>
<tr>
<td>P1f</td>
<td>plectin 1f</td>
</tr>
<tr>
<td>PBS</td>
<td>phosphate buffered saline</td>
</tr>
<tr>
<td>PBST</td>
<td>phosphate buffered saline-tween</td>
</tr>
<tr>
<td>PFA</td>
<td>paraformaldehyde</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Full Form</td>
</tr>
<tr>
<td>--------------</td>
<td>-----------</td>
</tr>
<tr>
<td>PIPES</td>
<td>piperazine-N,N'-bis(2-ethanesulfonic acid)</td>
</tr>
<tr>
<td>PMSF</td>
<td>phenyl-methyl-sulfonyl-fluoride</td>
</tr>
<tr>
<td>PMT</td>
<td>post-translation modifications</td>
</tr>
<tr>
<td>PRD</td>
<td>plakin repeat domain</td>
</tr>
<tr>
<td>RBD</td>
<td>rhotekin-binding domain</td>
</tr>
<tr>
<td>S</td>
<td>supernatant</td>
</tr>
<tr>
<td>Tris</td>
<td>tris-(hydroxymethyl) aminomethane</td>
</tr>
<tr>
<td>Wt</td>
<td>wild-type</td>
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