Neurofilament proteins in axonal regeneration and neurodegenerative diseases

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Abstract

Neurofilament protein is a component of the mature neuronal cytoskeleton, and it interacts with the zygosome, which is mediated by neurofilament-related proteins. Neurofilament protein regulates enzyme function and the structure of linker proteins. In addition, neurofilament gene expression plays an important role in nervous system development. Previous studies have shown that neurofilament gene transcriptional regulation is crucial for neurofilament protein expression, especially in axonal regeneration and degenerative diseases. Post-transcriptional regulation increased neurofilament protein gene transcription during axonal regeneration, ultimately resulting in a pattern of neurofilament protein expression. An expression imbalance of post-transcriptional regulatory proteins and other disorders could lead to amyotrophic lateral sclerosis or other neurodegenerative diseases. These findings indicated that after transcription, neurofilament protein regulated expression of related proteins and promoted regeneration of damaged axons, suggesting that regulation disorders could lead to neurodegenerative diseases.

Key Words: axonal regeneration; nerve injury; neurodegenerative diseases; neurofilament protein; post-transcriptional regulation; reviews

Abbreviations: NF, neurofilament; NF-L, light neurofilament; NF-M, midsize neurofilament; NF-H, heavy neurofilament; hnRNA, heterogeneous nuclear RNA

INTRODUCTION

In the 1830s, the structure of neurofilament (NF) protein was first reported¹, but the shape and accurate positioning in neurons was further confirmed with the development of electron microscopy. In recent years, NF structure, expression, modification, protein interactions, and other biological characteristics, as well as post-transcriptional gene regulation, have become increasingly important in the fields of neurodegenerative diseases and neural regeneration. NF expression is closely associated with axonal growth and maintenance of neuronal homeostasis. Current evidence indicates that NF plays a key role in axonal regeneration and is dependent on post-transcriptional mRNA transport, translation, and stability². Recent studies have demonstrated that post-transcriptional regulation increases NF gene transcription during axonal regeneration, ultimately resulting in NF protein expression patterns³. However, disordered expression of post-transcriptional regulatory proteins and other variant proteins can lead to amyotrophic lateral sclerosis⁴. Recent studies have shown that NF protein expression is closely associated with continued growth of normal axons and the regeneration of impaired axons. These NF functions contribute to the transcription process, as well as the regulation of mRNA transport, translation, and stability.

Post-transcriptional regulation plays an important role in maintaining stability of adult neurons, and regulation disorders may be the cause of some neurodegenerative diseases⁵. These previous studies proposed explanations regarding how neurons coordinate with function-related proteins to promote axonal regeneration, revealing that regulation disorders can lead to neurodegenerative diseases. This paper summarizes current knowledge on NF biology, including NF structure, expression, modification, protein interactions, and post-transcriptional gene regulation related to neural regeneration and neurodegenerative diseases.

NF STRUCTURE, EXPRESSION, MODIFICATION, AND INTERACTIONS WITH PROTEIN

NF structure

The neuronal cytoskeleton is composed of actin filament protein (16 nm diameter), tubulin (6 nm diameter), and intermediate...
filament protein (10 nm diameter). Intermediate filaments are unique, because they are made of different proteins in different tissues. NFs are abundant in neurons, especially in large-diameter axons. In vitro studies have shown that axonal NFs are a parallel array arranged between NFs and microtubules, or membranous organelles, 10 nm long. The light neurofilament (NF-L) subunit itself is a 10-nm core filament. Although these three types of protein structures have been previously described, midsized neurofilament (NF-M) and heavy neurofilament (NF-H) have a long, highly variable carboxyl-terminal tail (NF-M 439 amino acids, NF-H 660 amino acids), which contains several specific protein phosphorylation targets.

There are three kinds of cytoskeleton, namely intermediate filaments, microtubules, and microfilaments, and each have unique structure and assembly properties. NFs are the main component of the cytoskeleton in mature neurons, as well as the intermediate filaments of neurons. NFs are abundant at myelinated axons within cytoskeleton polymers in the vertebrate. The estimated molecular weights of human NF-L, NF-M, and NF-H subunits in DNA sequences are 61.5, 102.5, and 112.5 kDa, respectively. However, the presence of a high content of negatively charged amino acids (glutamic acid), as well as wide post-translational modification (phosphorylation and glycosylation), allows for higher molecular weight in sodium dodecyl sulfate polyacrylamide gel electrophoresis. NF-M or NF-H alone cannot synthesize fibroin if NF-L is not involved. NF-L transfection alone in human cells is able to synthesize the homopolymer in vitro, but such an efficacy is not available in mice and other rodents. In rodents, NFs synthesize heteropolymers according to a 4:2:1 ratio of NF-L: NF-M: NF-H subunits, and this ratio is continuously altered during neuronal formation. Currently, it is widely held that these proteins are integral components of NFs, along with NFL, NFM, and NFH. In addition to NF-L, NF-M, and NF-H, Yuan et al. showed that α-tubulin in the central nervous system acts as a fourth type of NF subunit. Different NF subunits form intermediate filament proteins; although nucleotide binding or hydrolysis is not required, the formation depends on ionic strength, pH values, and temperature. The first step for NF formation consists of NF-L and NF-H forming a parallel coiled dimer with a relatively stable bar area; this coiled dimer produces an anti-parallel tetramer in a half-staggered manner. Accordingly, the tetramer is incorporated into the protofilament and subsequently assembles 10-nm filaments. In vitro studies have demonstrated that the interaction between NFs results in the deformation of a visco-elastic network, which highlights the significance of microfilaments on NF mechanical properties. Kreplak et al. verified the NF mechanical properties using a mechanical microscope, showing that NFs are extended over 3 folds, with an average of 2.6 folds, indicating that NF functions to absorb mechanical shock in vivo.

NF expression
NF-L and NF-M genes are closely linked with chromosome 8 in humans, while the NF-H gene is located on chromosome 22. In mice, NF-L and NF-M genes are located on chromosome 14, while the NF-H gene is located on chromosome 11. Expression of the three NF subunits is regulated by nervous system development and ongoing neuronal differentiation. Expression levels modulate location, stability, and conversion efficiency at both the transcriptional level and mRNA post-transcriptional level. To ensure axonal growth, NF-L, together with intein and peripherin, is the first expressed subunit, followed by NF-M expression. NF protein mRNA sequential expression occurs in the order of NF-L, NF-M, and NF-H. In the peripheral nervous system and central nervous system, NF expression is significantly down-regulated at the early period following axonal transaction, leading to reduced NF transport at injured neurons. In contrast, NG subunit expression is significantly increased during the regeneration of damaged neurons. However, this increase does not occur when regeneration is prevented or in mammalian central nervous system axons that normally do not regenerate.

NF translational modification
NF modification includes phosphorylation and glycosylation, which contributes to NF morphology and function.

Phosphorylation
NF phosphorylation regulates NF axonal transport in neurons, plays a significant role in NF translocation, morphology, and function, and is involved in the pathogenesis of some neurodegenerative diseases. NF protein phosphorylation most widely occurs in the area from 51 site to NF-H C-terminal region. Phosphorylation is dense in the axonal head-domain, but scarce in the cell body and dendrites. Phosphorylation mainly occurs in the N-terminal and carboxyl terminal of three subunits, which are second messenger-dependent kinase and non-dependent kinase targets.

Head-domain phosphorylation occurs soon after NF synthesis in cells, while tail-domain phosphorylation begins when it transfers to axons. NF phosphorylation helps to maintain a dynamic balance between kinases and phosphatases in different nerve segments. Because head-domain phosphorylation inhibits NF assembly, dephosphorylation is necessary prior to NF transport into the axons. A wide range of enzyme dephosphorylation events result in a gradual loss of NF functions, a loss of interconnecting in vitro into a reticulated network, measured by the formation of highly viscous gels in purified preparations of NFs. Finally, dephosphorylation of the NF termination contributes to degradation and regulates their interactions with other cytoskeletal proteins. NF subunit dephosphorylation is catalytically achieved via phosphatase 2A and is assisted by phosphatase 1. Lobsiger et al. studied NF-H and NF-M phosphorylation in gene knockout mice, demonstrating...
that amytrophic lateral sclerosis is delayed compared to normal animal models with identical total NF levels, suggesting that abnormal NF phosphorylation induces amytrophic lateral sclerosis.

**Glycosylation**

NFs are modified by O-linked N-acetyl glucosamine or by post-translational serine and acetyl glucosamine residues, which regulate protein stability, subcellular localization, and protein-protein interactions[34]. Similar to phosphorylation, O-glycosylation is a dynamic process that alternates with phosphorylation. Glycosylated NF is present around peripheral nerves in diabetic patients, which may be related to peripheral neuropathy[35]. Post-translational modified NFs may be significant in family and single amytrophic lateral sclerosis[36].

**NF interactions with proteins and organelles**

As a complex, dynamic network, the interactions between NF and its yzosome are mediated by NF-associated proteins, which regulate enzyme function and NF linker protein structure. Linker proteins are responsible for the interaction between actin and cell organelles, while kinase and phosphatase are involved in regulating NF structure, assembly, and spacing. According to Tashiro et al[37], studies of the rat sciatic nerve showed that phosphorylated NF significantly stabilizes microtubules within axons, thereby promoting axonal regeneration. Other results have shown that dynein and kinesin in molecular motors together with C-terminal NF phosphorylation, promote NF transport along axons and dendrites[38]. Using a yeast double-hybrid and affinity chromatography model, dynein intermediate chain binding NF-M was shown to bilaterally participate in NF transport in axons[39]. This study summarized and analyzed well-known or likely proteins, enzymes, and receptors that interact with NF (Table 1).

**NF GENE EXPRESSION, HIERARCHICAL CONTROL, AND POST-TRANSCRIPTIONAL REGULATION**

NF gene expression plays an important role in nervous system development, as well as repair of damaged nervous system and neurodegenerative diseases[40]. Results have suggested that the regulation of NF gene transcription is crucial for the control of NF expression, especially in neural regeneration and neurodegenerative diseases[41].

**Effect of altered NF expression on axon growth, regeneration, and neurodegenerative diseases**

NF subunit expression is continuously changing in the developing vertebrate, indicating that expression correlates with nervous system regulation. As a protein expressed in the peripheral nervous system, NF-L maintains a high level in axons[37]. In the central nervous system, type III intermediate filament protein and vimentin expressions disappear in newly differentiated neurons, which is similar to the role of peripheral proteins[42]. Unexplained type III intermediate filament protein expression is conducive to axonal growth, and the majority of organisms survive due to wide axonal regeneration following severe nerve injury. The appropriate regulation of NF expression delays neural aging, and immunoblotting assays have shown that neuronal growth in aged rats is less than in young rats, because NF-L expression decreases in distal axons, while NF-M and NF-H mRNA expression is also selectively reduced[43]. Therefore, NF dysfunction leads to neurodegenerative diseases.

**Table 1** Protein interactions with neurofilament (NF)

| Proteinpanke | Known or possible functions | Source |
|--------------|-----------------------------|--------|
| Tubulin      | Integrity of cytoskeleton   | Miyasaka et al[44] Frappier et al[45] |
| Fodrin       | Interaction between cytoskeleton and organelles | |
| H1 histone   | Regulating DNA synthesis    | Young et al[46] |
| G-actin      | Maintaining neuronal structure | Hao et al[47] Haddad et al[48] |
| Hamartin     | Interaction between NF and actin | |
| Microtubule-associated protein 2 | Bridge between NF and MT | Perkins et al[49] |
| Synapsin 1   | Interaction between NF and synaptic non-secretory vesicle | Loureiro et al[50] |

**Motor proteins**

| Name          | Function                            | Source       |
|---------------|-------------------------------------|--------------|
| Dynnei        | NF transport in axon                | Shah et al[51] |
| Kinesin       | NF transport in axon                | Yabe et al[52] |

**Kinase and phosphokinase**

| Name          | Function                            | Source       |
|---------------|-------------------------------------|--------------|
| Stress activated protein kinase | NF-H C-terminal phosphorylation | Ackerley et al[53] |
| Casein kinase | NF-L and NF-M C-terminal phosphorylation | Nakamura et al[54] |
| Cyclin-dependent kinase 5 | NF-M and NF-H phenylalanine | Sharma et al[55] |

**Extracellular signal-regulated kinase**

| Name          | Function                            | Source       |
|---------------|-------------------------------------|--------------|
| Extracellular signal-regulated kinase | NF-M and NF-H phenylalanine | Guidato et al[56] |

**Protein kinase A, C**

| Name          | Function                            | Source       |
|---------------|-------------------------------------|--------------|
| Protein kinase A, C | NF triplex N-terminal phosphorylation | Sihag et al[57] |

**Receptors**

| Name          | Function                            | Source       |
|---------------|-------------------------------------|--------------|
| Dopamine receptor | Regulating cell surface expression and desensitization | Kim et al[58] |
| N-methyl-D-aspartate receptor | Reticular nucleus anchoring and positioning | Ehlers et al[59] |

Hierarchical control of NF protein expression

Regulation of gene expression was originally found to occur at the transcriptional level. Clearly, the transcriptional controlling effect is effective for NF expression and specificity of neuronal growth[60]. Gene expression includes transcriptional and post-transcriptional control, and the latter is of equal significance in NF expression. Post-transcriptional regulation of gene expression makes protein synthesis more effective and can coordinate related proteins. With a better understanding of eukaryotic gene expression transcription and post-transcriptional regulatory mechanisms, current studies focus on regulation points of gene expression (Figure 1).

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Translated mRNA in axons containing ribosomes is restricted to a specific segment of the axon. Willis et al. [60] discovered limitations of NF mRNA within growth cone axons. Similarly, post-transcriptional regulation of NF subtype expression was observed to be significant, and NF-L protein expression was significantly increased compared to NF-H in gene dosage tolerance in transgenic mice, which directly relates to post-transcriptional inhibition of NF-L mRNA expression [61].

**Translation and post-transcriptional interaction controls NF axonal regeneration in the central nervous system**

A number of studies on NF expression in Xenopus optic nerve regeneration have provided clear evidence for the role of transcriptional and post-transcriptional control on NF expression [62]. In these studies, heterogeneous nuclear RNA (hnRNA)-related mRNA changes were used as alternative measurements of gene transfer following optic nerve damage. hnRNA is an intermediate during transcription; it exists transiently and can be determined by introns that lack mature mRNA. Polysomal artherectomy serves as an analysis indicator of NF messenger RNA translation efficiency. Using introns and exons as probes, quantitative analysis of reverse transcription-polymerase chain reaction showed that NF-M transcription and linker protein subunit expression are similar to the normal side following optic nerve injury. In addition, NF mRNA expression is decreased in the surgical side of the optic nerve, although transcription was blocked during the initial regeneration phase. Even at the peak of axonal regeneration, hnRNA levels remain several times greater than mRNA levels [63]. During regeneration, NF information transcription efficacy fluctuates, with a greater amount of NF mRNA in polyribosomes at the operated side of the optic nerve [64]. Motor neurons from the normal side (no injury) mediate the injured side via the spinal cord, and similar signal transductions occur following retinal damage. Multi-path synaptic pathways are transferred from both eyes to the thalamus and visual nerve center. A small amount of NF-M mRNA is transformed to polyribosomes in the eyes and brain of the non-operated side, whereas about 85% of NF-M mRNA transform into polyribosomes in eyes from the surgical side when neural regeneration peaks, indicating that NF mRNA protein aggregation requires more NF-M structural proteins for neural regeneration [65]. Increased NF-M transcription on the normal and operated sides of the optic nerve reflects both optic nerve pathways in response to the initial fracture. For example, following peripheral nerve transection, instantaneous peripheral nervous system regeneration occurs in contralateral non-damaged neurons in rodent models. These neurons germinate towards the injury side, which is mediated by the spinal cord [66]. Similar signal transduction is also observed in the damaged retina, because signals are transferred from the thalamus and visual center to the eyes via a multi-synaptic pathway. Another possibility is that signals are mediated by body fluids, although the differences between these two mechanism require further studies [67].

Results have shown that NF expression regulates several genes that are involved in the repair of central nervous system injury [68], which suggests that post-transcriptional regulation meet requirements for NF subtypes in axonal regeneration, thereby promoting neural regeneration. Another study showed that NF post-transcriptional gene regulation mechanism is activated to promote repair of damaged nerves in the adult central nervous system, but also raises the question that inappropriate signal activation under the same mechanisms could be related to disease occurrence [59].

**NF post-transcriptional regulation in neurodegenerative diseases**

NF-containing polymers exhibit unique characteristics in various neurodegenerative diseases. Amyotrophic lateral sclerosis is closely related to out-of-control...
CONCLUSION

NF protein expression is closely linked with continuous growth of normal axons and impaired axonal regeneration. NF protein post-transcriptionally regulated changes in mRNA transport, translation and stability are now thought to work together with transcription to achieve the final expression pattern. Post-transcriptional regulation plays an important role in the maintenance of adult neuronal stability, and this regulation loss may result in neurodegenerative disease. NF post-transcriptional regulation is similar to axonal growth-related proteins, suggesting that post-transcriptional regulatory factors could act on critical axon proteins. In recent years, the use of transgenic mouse models allows researchers to determine the role of NF in axonal caliber during nerve growth. NF post-transcriptional gene regulation plays a role in axonal regeneration and neurodegenerative diseases.

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Conflicts of interest: None declared.

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