INSTITUT CHIARI & SIRINGOMIELIA & ESCOLIOSIS DE BARCELONA

N Sinternational Congress Fascia a Ser Kurhaus Schever Ber JaRusry 1sth, 2020 N A

The Filum terminale

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TESTED ATA

Plan of presentation

- 1.1. Anatomy of Filum terminale
- 1.2. Histology of Filum terminale
- 1.3. Physiology of Filum terminale
- 1.4. Embryology of Filum terminale
- 2.1. Definition of Filum disease
- 2.2. Definition of Neuro-cranio-vertebral syndrome
- 2.3. Basic statistics
- 2.4. Clinical picture
- 2.5. Image features
- 3.1. The Filum System
- 3.2. Sectioning of Filum terminale
- 3.3. Other interventions
- 4. Results of Filum System
 - 5. Conclusions.





BARCELO NA

1.1. Anatomy of Filum terminale

- Fibroelastic ligament between the *Conus medullaris* and the dorsal surface of the tailbone.
- About 23 cm in length.
- Thickness of 1-2 mm.
- Accompanied by a visible arterial blood vessel over its entire length.
- Comprises a longer initial intradural segment (*Filum terminale internum*) and a shorter distal extradural segment (*Filum terminale externum*), the latter covered by a prolongation of the *dura mater*.

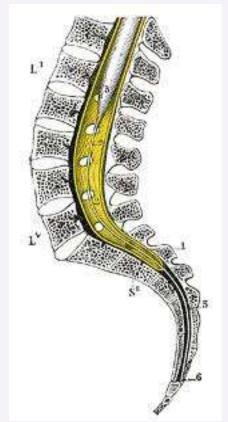








Conus medullaris and Filum terminale in anatomy textbooks:

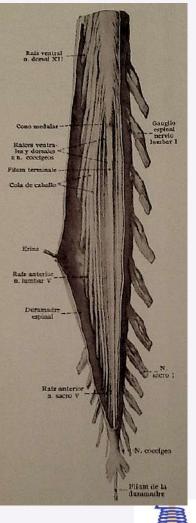


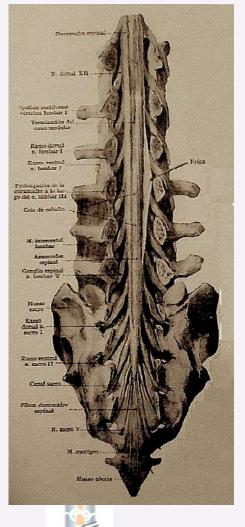
L. Testut y A.Latarjet 1902-1928

Werner Spalteholz 1895









FILUM ACADEMY BARCELONA"



Filum terminale in cadaver dissections:













Filum terminale in individual with scoliosis (prepared by Professor Alfonso Rodriguez Baeza, Chair of Human Anatomy and Embriology, Faculty of Medicine, Universidad Autónoma de Barcelona, 2017).











1.2. Histology of *Filum terminale*

- Collagen fibers
- Elastin fibers
- Atrophic neurons
- Atrophic neural fibers
- Fibrous and adipous tissue with multiple small blood vessels
- Scarce neural filaments positive for S100 Protein
- Various types of glial cells.

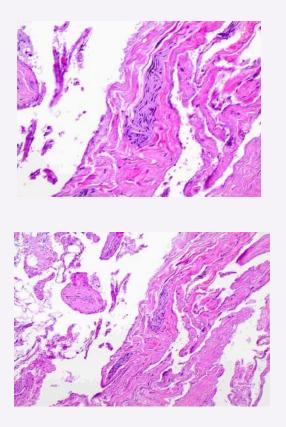


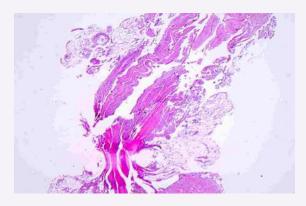


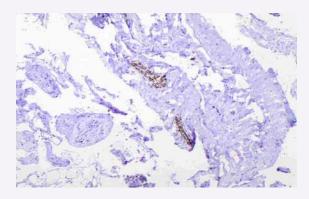














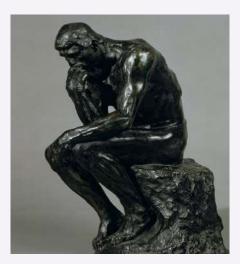






1.3. Physiology of *Filum terminale*

• Maintains the alignment of the spinal cord within the spinal canal.











1.4. Embryology of *Filum terminale*

- Initiated by involution of the distalmost spinal cord through retrogressive differentiation;
- It becomes visible in the 9th gestational week, as a consequence of the differential growth of the spine and spinal cord;
- It contains both ectodermal and mesodermal-origin components.









9-week-old human embryo, at the beginning of the differential growth of the spine and spinal cord













Conus medullaris level in humans:



- Starts to ascend during the 9th week;
- At 3 months: 2nd coccygeal vertebra;
- At 5 months: 1st sacral vertebra;
- At 6 months: **L**₃-**L**₄.
- At birth: inferior part of L₃;
- During adulthood: **between D**₁₂L₁ and L₁L₂.









Normal *Conus medullaris* level in adults:

- Th₁₂-L₁ (Roth 1981, Royo-Salvador 1996)
- L₁ (Arai el al. 2001, Pou Serradell 1981)
- L₁-L₂ (Williams et al. 1995)
- Above upper L₂ (Nieuwenhuys et al. 2009)
- Above L₂ (Heimberger 1950)
- Above inferior third of L₂ (McCotter, 1916)
- Between Th₁₁ and L₃ (Neel, 2016)









Conus medullaris level

In MRI of 55 control subjects (Royo Salvador, 1993)



- Th₁₁-Th₁₂: 1 (2%)
- Th₁₂L₁ disc: 25 (50 %)
- L₁ vertebral body: 21 (42 %)
- Inferior margin of L₁: 1 (2%)
- L₁L₂ disc: 2 (4%)









2.1. Definition of Neuro-Cranio-Vertebral Syndrome

- Group of subjective and objective clinical manifestations, affecting the encephalon, the spinal cord, the skull and the vertebral column, referable to an abnormal axial traction transmitted by a *Filum terminale* apparently normal;
- It can be associated to known conditions of unknow etiology so far, as the Arnold-Chiari type I syndrome, Idiopathic scoliosis, Idiopathic syringomyelia, Platibasia, Odontoid retroflexion, Brainstem kinking, etc.









2.2. Definition of *Filum* disease

- Congenital form of Neuro-Cranio-Vertebral Syndrome with an apparently normal *Filum terminale*;
- It can be manifested through different pathologies: Arnold-Chiari type I syndrome, idiopathic scoliosis and syringomyelia, platybasia, odontoid retroversion, brainstem kinking, etc.











Estimated worldwide prevalence of Filum disease

- <u>CONVENTIONAL DIAGNOSES</u>: 616 millions idiopathic scoliosis + 7 millions Chiari I + 580,000 idiopathic syringomyelia
 =623,588,000 patients or 8.9 % of world population
- <u>ACTUAL ESTIMATE: + scolioses < 11^o + subclinical Chiari I and syringomyelia + related pathologies (occipito-cervical junction malformations)</u>
 - > 20 % of world population



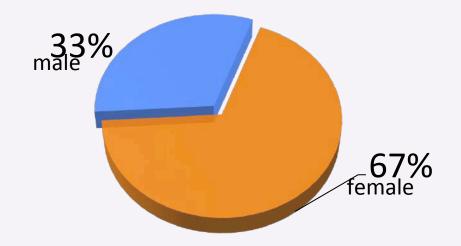






2.3. Basic statistics:

Sex distribution in 850 cases



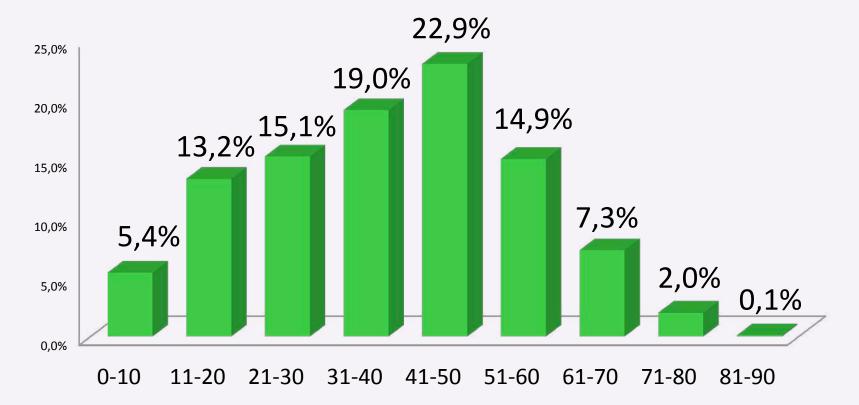








Age distribution in 850 cases



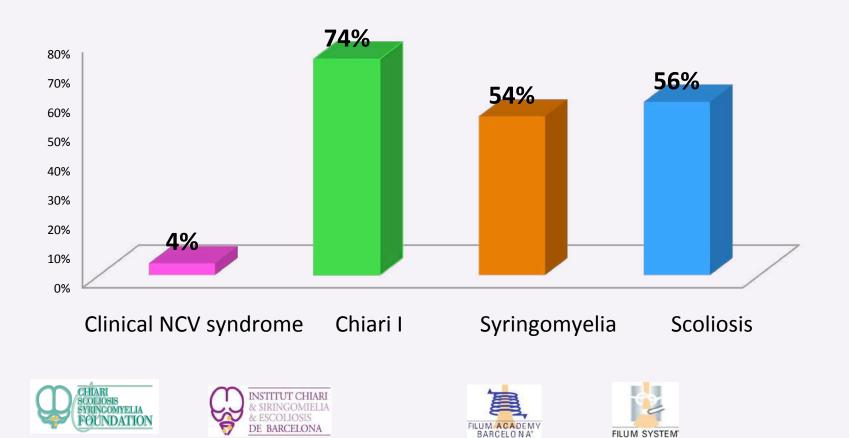




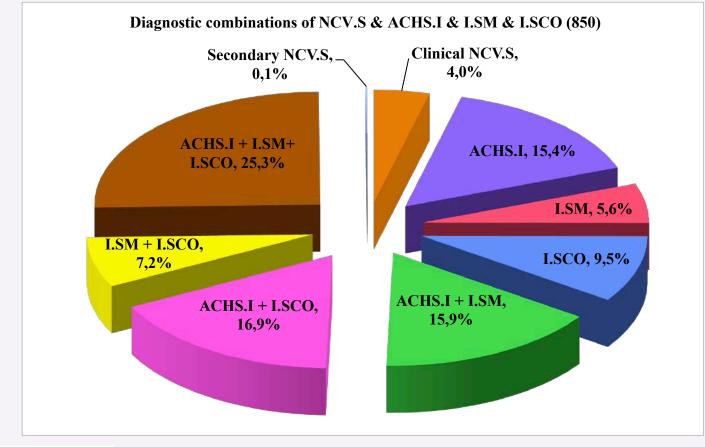




Ratio of classical diagnoses in 850 cases



Combinations of classical conditions in 850 cases



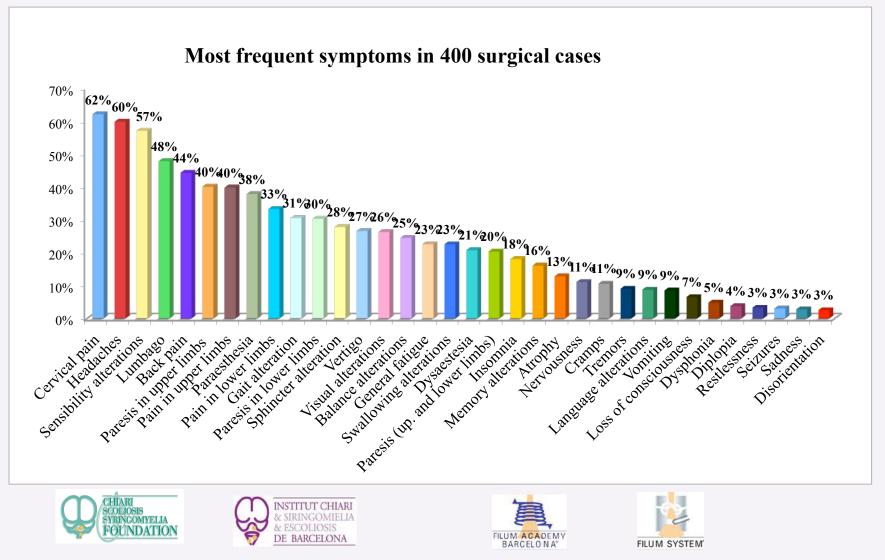








2.4. Clinical picture



Physical findings

- 1) Spontaneous nystagmus
- 2) Deviation of uvula and/or tongue
- 3) Diminished grip strength
- 4) Alterations of temperature sensibility
- 5) Alterations of touch sensibility
- 6) Altered deep tendon reflexes in upper limb(s)
- 7) Altered deep tendon reflexes in lower limb(s)
- 8) Altered abdominal cutaneous reflexes
- 9) Altered plantar cutaneous reflexes
- 10) Positive straight-leg raising test and reversed Lasègue's test
- 11) Positive Mingazzini's and Barré's tests
- 12) Positive Romberg's test
- 13) Difficulty standing up from a kneeling posture
- 14) Scoliotic attitude and associated signs.

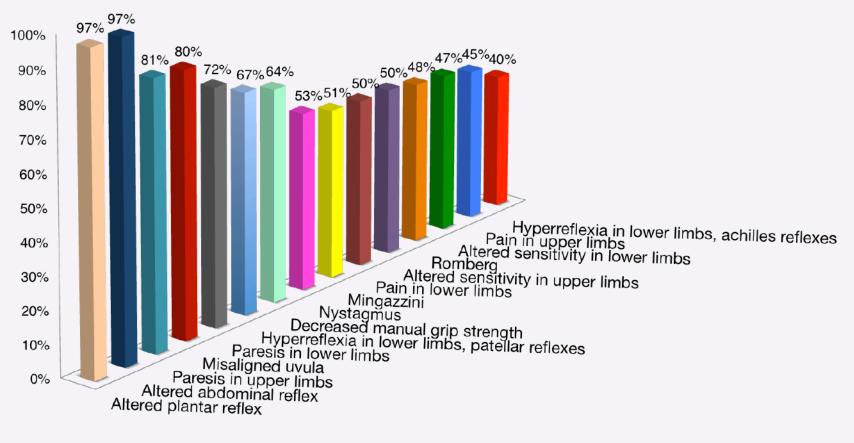












Most frequent clinical signs in 400 surgical cases

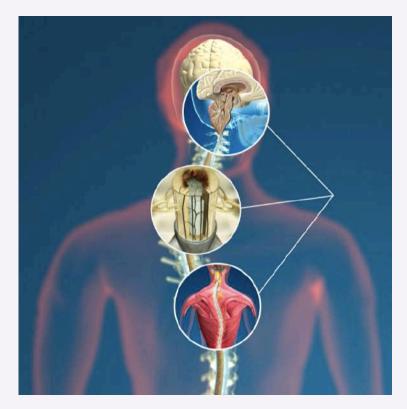








2.5. Image features of *Filum* disease: **Global view**











FILUM SYSTEM

Arnold-Chiari type I syndrome – caudal traction pathogenesis













Idiopathic syringomyelia





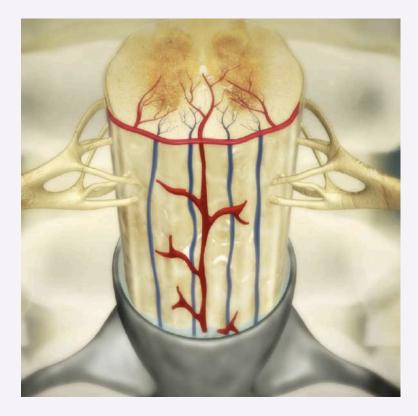


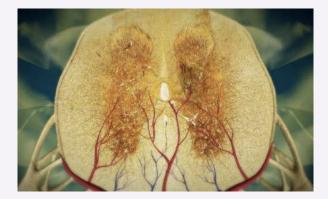






Idiopathic syringomyelia – ischemic pathogenesis







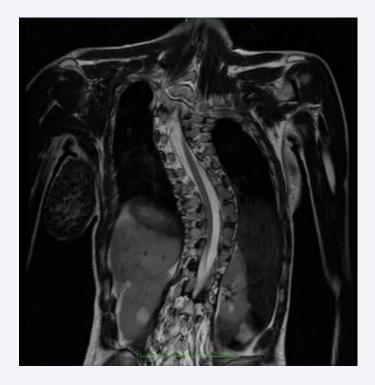


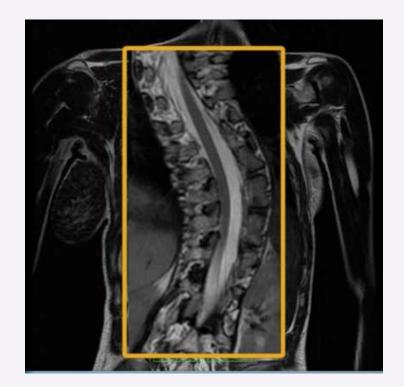






Idiopathic scoliosis





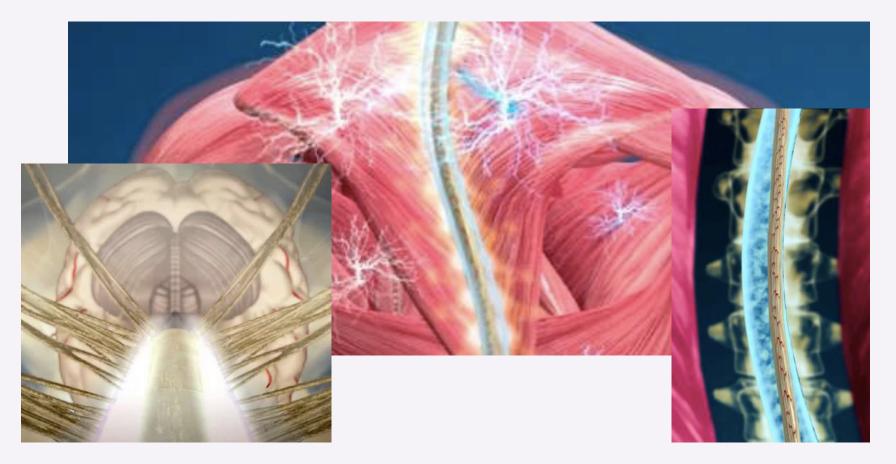








Idiopathic scoliosis – neurogenic pathogenesis











3.1. The FILUM SYSTEM[®] = 16 protocols

- I. Diagnosis
- II. Treatment
- III. Preoperative and Anaesthesia
- IV. Surgical
- V. Postoperative
- VI. Results
- VII. Postoperative medical treatment
- VIII. Postoperative physical therapy and rehabilitation
- IX. Postoperative orthoses and prostheses
- X. Postoperative check-up and genetics
- XI. Information
- XII. Future
- XIII. Spinal surgery
- XIV. Institute of filum
- XV. Case series
- XVI. Actualizations and related documents





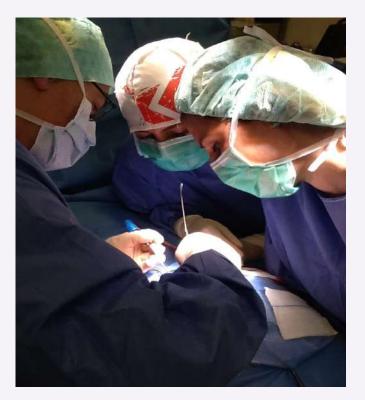






3.2. Sectioning of *Filum terminale*

original minimally invasive technique local anesthesia with sedation possible day surgery











3.3. Other surgical interventions

- Anterior cervical discectomy and fusion for cervical disc herniations.
- **Transarticular approach and discectomy** for thoracic disc herniations.
- Microsurgical or endoscopic discectomy for lumbar disc herniations.
- Laminectomy for spinal stenosis.
- Radiofrequency thermocoagulation for facet syndrome.



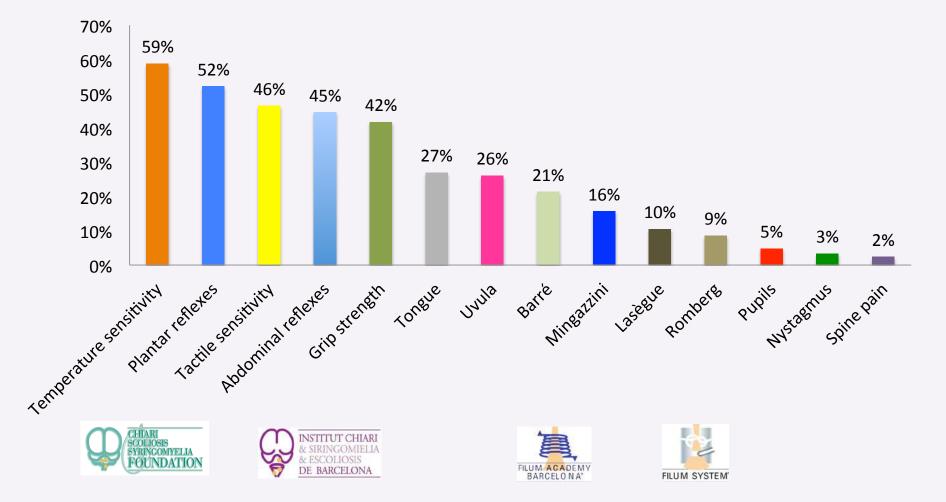




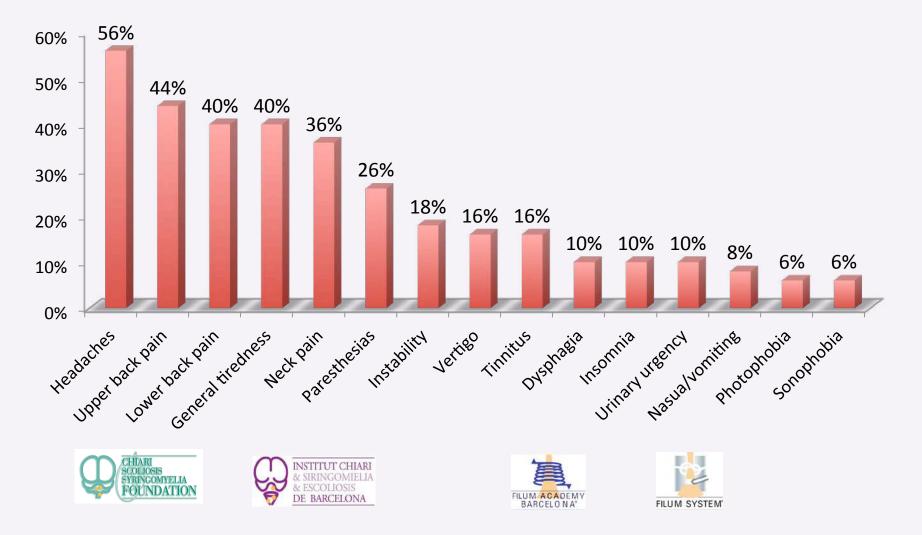


4. Results of FS[®]

Early improvements of clinical signs 8 hours postSFT in 211 cases:



Improvements of clinical symptoms one month postSFT in 50 cases:



Improvements of clinical signs one month postSFT in 50 cases:

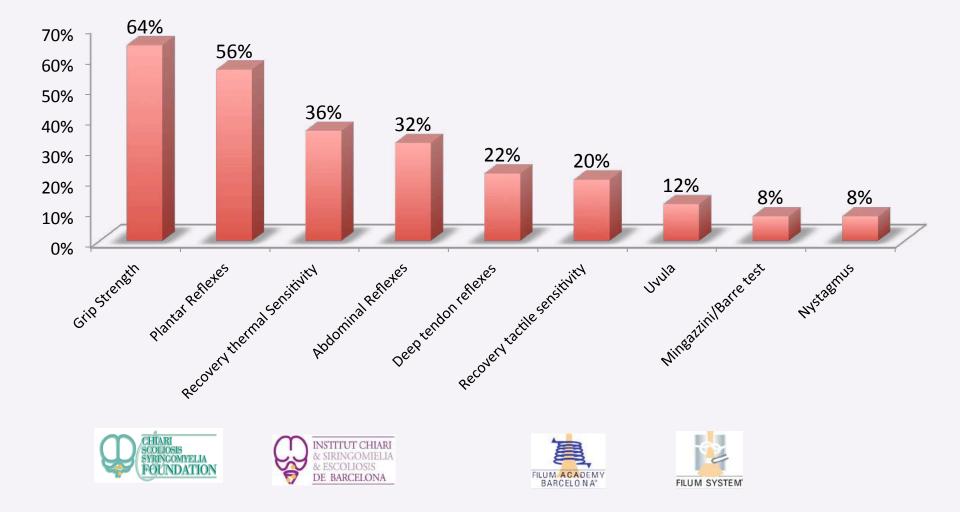


Image improvements after SFT: less tonsillar descent (1)



2011



2012





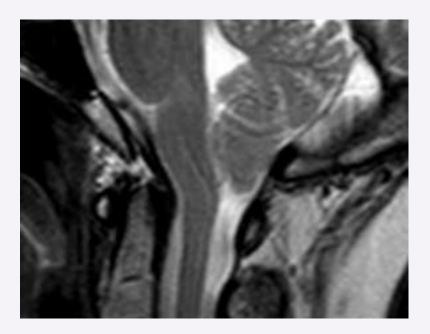




FILUM SYSTEM

Image improvements after SFT: less tonsillar descent (2)





2012









FILUM SYSTEM

Image improvements after SFT: less tonsillar descent (3)













FILUM SYSTEM

Image improvements after SFT: smaller syringomyelia (1)













Image improvements after SFT: smaller syringomyelia (2)



2009



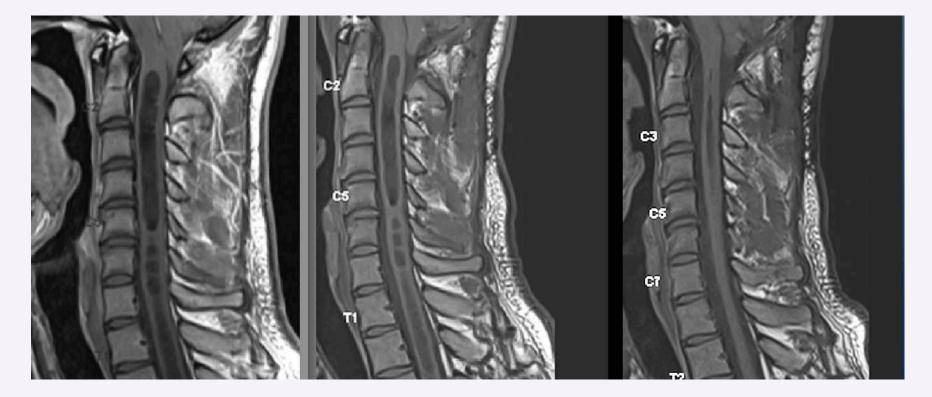








Image improvements after SFT: smaller syringomyelia (3)



2016

2017

2019









Image improvements after SFT: less scoliosis (1)



2001/12.14°D,16.29°L







2009/1.84°D, 10.54°L





Image improvements after SFT: less scoliosis (2)





2010



2009









Image improvements after SFT: less scoliosis (3)





2017

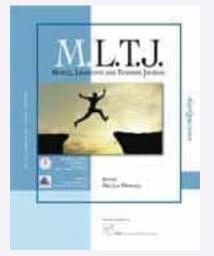








5. Conclusions: Manual Therapy in Filum Disease



Shortness of *filum terminale* represents an anatomical specific feature in fibromyalgia: a nuclear magnetic resonance and clinical study

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Summary

Background: we aimed to assess whether shortness of filum terminale (FT) can represent a specifKEY WORDS: anatomical modification, fibromyalgia impact questionnaire, fibromyalgia assessment status, mini-invasive surgery, musculo-skeletal disorder, physical therapy.

Original article

Introduction

Filum terminale (FT) is a strand of fibrous tissue about 20 cm long and made of two parts: *filum terminale internum* and *filum terminale externum*. The former is nearly entirely made of *pia mater*, the innermost meningeal layer, and extends from the *conus medullaris* of the spinal cord to the *cul-de-sac* of the most external meningeal layers, i.e., the arachnoid membrane and the *dura mater*. At this level, it fuses with those meningeal membranes thus making the coccygeal ligament, which fastens the most distal portion of the *medulla* to the *coccys.* Indeed, the coccygeal ligament is often referred as *filum terminale externum*.

Some diseases affecting FT have been described, including some forms of tumors, mineral deposition and *filum terminale* disease (FD), described in 1996 by Royo-Salvador^{1,2}. *Filum terminale* disease is an outcome of the abnormal traction exerted on the spinal cord by a FT shorter that usual. This condition is

MUSCLES, LIGAMENTS AND TENDONS JOURNAL 2015;5 (1):33-37









- Royo-Salvador MB (1996) [Syringomyelia, scoliosis and idiopathic Arnold-Chiari malformations: a common etiology]. Revista de Neurología 24:937-959
- Royo-Salvador MB (1996) [Platybasia, basilar groove, odontoid process and kinking of the brainstem: a common etiology with idiopathic syringomyelia, scoliosis and Chiari malformations] Revista de Neurología 24:1241-1250
- Royo-Salvador MB (1997) A new surgical treatment for syringomyelia, scoliosis, Arnold-Chiari malformation, kinking of the brainstem, odontoid recess, idiopathic basilar impression and platybasia. Revista de Neurología. 25:523-530
- Royo-Salvador MB Sole-Llenas J Domenech JM Gonzalez-Adrio R (2005) Results of the section of the filum terminale in 20 patients with syringomyelia, scoliosis and Chiari malformation. Acta Neurochirurgica. 14:515-523
- Royo-Salvador MB (2014) Filum System[®] A Brief Guide. Chiari & Scoliosis & Syringomyelia Foundation, Barcelona
- Royo-Salvador MB (2016) The MRI in Arnold-Chiari Syndrome I and Idiopathic Syringomyelia. Rare Disease Report. <u>https://institutchiaribcn.com/commons/pdf/publicaciones/dr-royo/The-MRI-in-Arnold-Chiari-Syndrome-I-and-Idiopathic-Syringomyelia.pdf</u>.
- Royo-Salvador MB (2017) Differentiating Tethered Cord Syndrome, Neuro-Cranio-Vertebral Syndrome, and Filum Disease. Rare Disease Report.
 <u>https://institutchiaribcn.com/commons/pdf/publicaciones/dr-royo/Differentiating-Tethered-Cord-Syndrome-Neuro-Cranio-Vertebral-Syndrome-and-Filum-Disease.pdf.</u>
- Royo-Salvador MB, Fiallos-Rivera M, Salca H (2019) Caudal Traction as a Pathogenetic Mechanism of Chiari Malformation Type I [Online First], IntechOpen, DOI: 10.5772/intechopen.90044.

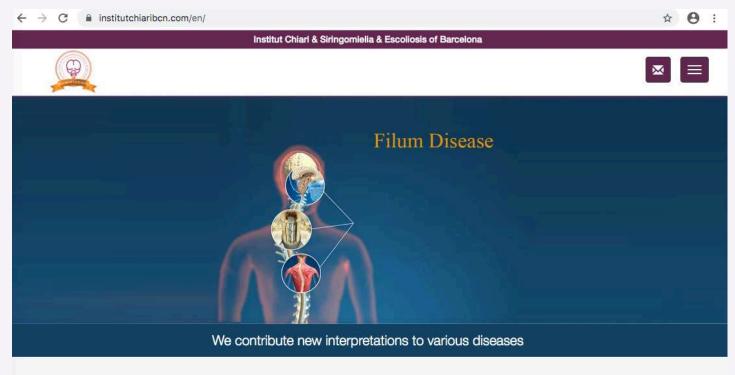








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