2022 흉부외과 전공의 연수교육



다한증, 흉벽기형, 흉곽출구증후군

연세대학교 강남세브란스병원 흉부외과 문덕환

Contents



Hyperhidrosis

Pathologic condition of excessive sweating in amounts greater than physiologically needed for thermoregulation

Hyperhidrosis - Pathogenesis

Eccrine sweat glands are responsible for hyperhidrosis

mixture of the two [apo/eccrine] glands may play a role in axillary hyperhidrosis

A sympathetic signal is carried to sweat glands by cholinergic autonomic neurons

Idiopathic (focal) hyperhidrosis

Sweat glands are usually histologically and functionally normal. Abnormal central response to emotional stress

Genetic component

Hyperhidrosis - Pathogenesis

Types of hyperhidrosis	Commonly associated diseases and conditions	
Primary hyperhidrosis	Idiopathic (focal) hyperhidrosis	
	Gustatory hyperhidrosis (Frey's syndrome)	
Secondary hyperhidrosis	Endocrine: hyperthyroidism, hyperpituitarism, diabetes mellitus, menopause, pregnancy, pheochromocytoma, carcinoid syndrome, and acromegaly	
	Neurological: Parkinson's disease, spinal cord injury, and stroke	
	Neoplastic: Hodgkin's disease and myeloproliferative diseases	
	Infectious: tuberculosis and septicemia	
	Drugs: fluoxetine, venlafaxine, doxepin	
	Toxicity: alcoholism and illicit substance abuse	

Focal or Primary HH – Craniofacial, palmar, axillary, or plantar

Hyperhidrosis - Treatment

Treatment	Cost ^a	Side Effects
Topical, 20% to 35% aluminum chloride	\$288+/year	Skin irritation, localized burning, stinging, desquamation, poor efficacy, temporary (lasts about 48 hours per application)
Iontophoresis (usually 20 mA 3 to 4 treatments a week for 30 to 40 minutes each)	\$500/device	Irritation, dryness or peeling of skin, burning or stinging during therapy, temporary (one treatment lasts 1 to 4 weeks). Not recommended for women who are pregnant or for persons with pacemakers or substantial implants (eg, joint replacements)
Oral therapy (glycopyrrolate, atropine, acetylcholine inhibitors)	\$240+/year	Dry mouth, dry eyes, constipation, mydriasis, difficulty urinating, blurry vision
Botulinum toxin (Botox A or B)	\$2,250/session	Pain from injections, muscle weakness, headache, hematoma, swelling, need for repeat procedures
Liposuction/VASER	\$3,000/session	Hematoma, superficial skin erosion, alopecia, paresthesia
Endoscopic thoracic sympathotomy	\$15,000	Compensatory hyperhidrosis, bradycardia, pneumothorax, postoperative pain, Horner's syndrome

Table 2. Comparison of Therapies for Primary Hyperhidrosis

^a Approximate cost in US dollars.

+ Endoscopic Lumbar Sympathectomy (ELS)

Hyperhidrosis - Nomenclature for ETS

Rib- oriented nomenclature

- Too many patients having mediastinal fat that can obscure clear identification of the specific ganglia
- Many anatomical variations in the ganglion anatomy

Type of interruption

- Clipped, cut, or cauterized, or a segment removed

For example

- Clipped R5, top
- cauterized, top R4, bottom R4

Hyperhidrosis - "Type of interruption"

Transection? Resection? Ablatation with a cautery? Division with a harmonic scalpel? or Clipping?

> No clear differences(but clipping shows recurrence) If the correct level division was achieved

 Enough separation between the ends of the chain Regrowth is impossible

Hyperhidrosis - Patient Selection

Surgical consultation should include

- Secure diagnosis of primary focal hyperhidrosis
- Anatomic locations involved
- Amount of hyperhidrosis
- Full discussion of the options to surgery and potential complications

The patients should be made aware that the most satisfied patients are those with palmar or palmar-axillary hyperhidrosis, or both in ETS.

Endoscopic Lumbar Sympathetomy (ELS) for plantar hyperhidrosis seems to be an effective procedure, however this procedure is a challenging surgery.

Hyperhidrosis - ETS interruption level

Palmar HH : <u>R4 only</u> - limits the degree of CH

- may lead to moister hands
- R3,4 completely dry hands
 - Higher risk of CH

Axillary HH : R4,5 – less successful and has higher "regret rates" than ETS for palmar HH

CFH : <u>R2 vs. R3</u>

KEY LEVEL – R4 in ETS

Hyperhidrosis - ETS interruption level

Original Article

Early results of new endoscopic thoracic sympathectomy for craniofacial hyperhidrosis

Duk Hwan Moon¹, Du-Young Kang², Dong Won Kim³, Min Kyun Kang¹, Sungsoo Lee¹

¹Department of Thoracie and Cardiovascular Surgery, Canguan Severance Hospital, Yonsai University Cellege of Medicine, Saoal, Korea, ¹Department of Cardiovascular and "Thoracie Surgery, Kangluk Sunsang Hospital, Surghynnkwn University School of Medicine, Seoal, Korea, ¹Department of Thoracie Surgery, Congam Institute of Raliological and Medical Sciences, Busan Korea

Coursestant: (1) Conception and design: DH Moon, S Let; (11) Administrative support: None; (11) Provision of study materials or patients: Nene; (3V) Collection and assembly of data: DY Kang, DW Kin, MK Kang, (3) Data analysis and interpretation: DH Moon, S Lee; (31) Manuscript writing: All authors; (7) Final approval of memorying: All authors;

Correspondence tre Sungson Lee, MD, PhD. Department of Thoracic and Cardiovascular Surgery, Gangnam Severance Hospital, Yonsei University College of Medicine, Seoral, Korea, Email: CHESTLEE@yuhs.ac.

Background: Endocopic thoracic sympathectomy (ETS) has been considered as a definitive treatment for hyperhidrosis. However, despite its well-established success rate, surgical treatment for crariofacial hyperhidrosis (CFH) is rarely performed due to the possibility of fatal complications and compensatory sweating. The aim of this study was to evaluate the safety and efficacy of our newly developed method of ETS for CFH, based on early results.

Methods: Between June 2016 and October 2017, a total of 70 patients underwent ETS with our new technique for CFEL All patients were placed under double-lumen insubation anesthesia with CO2 gas installation. We utilized two ports, one for 2-mm endoscope and asother for 3-mm instrument. Our technique involved R2 and R4-R7 sympatheterony with R4-R7 runnel ablation.

Results: There were 55 males and 15 famales, with a mean age of 48 years (range, 22–75 years). The median operation time was 38 minutes (range, 24–75 minutes). There was no operative mortality and morbidity. During the short follow-up period (average 7 months; range, 1–17 months), symptoms were improved in all patients and compensatory hyperhidrosis was observed 68 patients (714%), moderate in 15 patients (18.6%), and severe in 5 patients (71.4%).

Conclusions: In select patients, our technique of ETS appears to be a safe and effective treatment method for treating CFH. However, a study with long-term follow-up is still necessary to confirm our findings.

Neywords: Craniofacial hyperhidrosis (CFH), compensatory hyperhidrosis; endoscopic thoracic sympathectomy (ETS)

Sabmitted Jar. 16, 2018. Accepted for publication May 23, 2018. doi: 10.21037/jtd.2018.05.150 View this article at: http://dx.doi.org/10.21037/jtd.2018.05.190



Hyperhidrosis - Compensatory HH after ETS

The most common side effect

- which occurs in the literature from 3% to 98%

The most common risk factor

- T2 ganglion interruption(R2, R3
- The number of levels interrupted has been inconclusive as a risk factor

Preoperative testing? controversial

- Injecting bupivacaine
 - reversibly achieve sympathetic nerve blockade observe for CH

Treatment

- Ditropan or other anticholinergic medications in escalating doses

Hyperhidrosis - Compensatory HH after ETS

Reversal Surgery

:Nerve Reconstruction

:R5, 6, 7 ± 8 , 9







#1. " Go to Brazil..."



15:30 Strategies in Compensatory Sweating Avoid Compensatory Hyperhidrosis After Sympathetic Surgery for Craniofacial Hyperhidrosis



#2 " Go to PISA, ITALY..."

Moon Duk Hwan (Republic of Korea)

08:40

Abstracts Sessions Hyperhidrosis, Endoscopic Thoracic Sympathectomy, and Cardiovascular Outcomes: A Korean Health Insurance Review and Assessment Service Database Cohort Study

Moon Duk Hwan (Republic of Korea)







Funnel chest is an oval depression which involves the sternum as well as the costal cartilages.

Usually it is alrea child reaches mat

The degree of the xiphoid angle, to a lower sternum tou.



e marked as the

n the sternonest wall with the

Pectus excavatum is a relatively common anomaly

- occurs in about one in 300-400 live births
- three times more frequent in males
- often associated with connective tissue disorders, such as Marfan's disease or Ehlers-Danlos syndrome

Symptoms

- palpitation, exertional dyspnea, fatigue and dull precordial pain, paradoxical breathing, exercise intolerance

The deformity is also often emotionally disturbing, especially in adolescents, who often avoid active sports and become shy and retiring.

heredity :about 20 to 50% of patients have a family history of pectus deformities - Williams 1872

an overgrowth of the costal cartilages - Flesch 1873 - Lincoln Brown 1939(1596)

arrested growth of the sternum - Ebstein 1882

various intrauterine compressive forces such as pressure by the chin, knee or elbow

latent mediastinitis – Raubitsch

undue traction exerted upon the sternum by the diaphragmatico-sternal ligament - Lincoln Brown 1939(1596)

Chest Wall Deformity - Haller index

Pectus Severity Index

Used to assess severity of incursion of the sternum into the mediastinum Maximal transverse diameter/narrowest AP length of chest

Normal HI is 2.5

Significant PE has an index greater than $3.25 \rightarrow$ standard for determining candidacy for repair (cosmetic? function?)



less than satisfactory late outcomes



 Serial fixation

 with pectasion

Perichondrial

(c) transverse cuneiform osteotomy of the sternum at the upper level of the deformity(d) maintenance of the corrected position of the sternum













<u>Morphology – tailored bar shaping</u>

HJ Park





Double Compression and Complete Fixation bar system (DCCF)



Chest Wall Deformity - Vacuum Bell and 3MP



Chest Wall Deformity - Advice

Adults with severe complex chest wall deformity

Connective tissue disease

Stiff chest wall from prior thoracic or cardiac surgeries

Selection of proper indications and adequate surgical method is crucial for the successful and safe chest wall correction !!!

Things to remember...

Pectus carinatum is 16.7% of all chest wall deformities in the Boston children's hospital experience.

Chondrogladiolar type : most frequent form- anterior protrusion of the body of the sternum- protrusion of the lower costal cartilages

Chondromanubrial or "pouter pigeon" deformity

- : least frequent form
- protrusion of the upper costal cartilages
- relative depression of the body of the sternum.



Etiology : not clear

- an overgrowth of the costal cartilages with forward buckling of the cartilages and anterior displacement of the sternum

- genetic basis : 26% had a family history of chest wall deformity and 12% of scoliosis.
- more frequent in boys than in girls 3:1

PC is rarely present at birth

- deformity was not identified until after the eleventh birthday
- deformity often progresses during early childhood particularly in the period of rapid growth at puberty.

Chest Wall Defo

The current correction of Pect of costal cartilages and sterna invasive modifications using th

The majority of these operatio 1949 by Ravitch.



A B C

rinatum

often involving resection there are minimally

ocedure first described in

Localizador web Anticulo 79.720

NOTAS CLINICAS

Método miniinvasivo para la corrección



Dynamic Chest Compressor

Haje SA, Raymundo JLP.

Chest wall deformities: conservative treatment of

the anterior protrusion forms.

Brazilian J Orthop 1979;14:167-178.



Various Braces



Long-Term Results of Compressive Brace Therapy for Pectus Carinatum

Duk Hwan Moon¹ Min Kyun Kang¹ Hye Sun Lee² Sungsoo Lee¹

 Department of Thoracic and Cardiovascular Surgery, Gangnam Severance Hospital, Seoul, the Republic of Korea
 Biostatistics Collaboration Unit, Gangnam Severance Hospital, Seoul, the Republic of Korea Address for correspondence Sungsoo Lee, Department of Thoracic and Cardiovascular Surgery, Gangnam Severance Hospital, 211 Eonjuro, Gangnam-Gu, Seoul 06273, the Republic of Korea (e-mail: CHFSTLFR@vubs.ac).

Thorac Cardiovasc Surg 2019;67:67-72.

Abstract

Background Pectus carinatum (PC) is one of the most common types of congenital chest wall deformity. Recently, noninvasive compressive brace therapy has been more frequently used than invasive surgical correction to treat PC. Hence, the purpose of this study was to determine the long-term outcome of compressive brace therapy. Methods We retrospectively reviewed patients with PC who underwent compressive brace therapy between January 2014 and December 2016. All patients underwent a 2week compression period, in which braces were worn for 20 hours per day, followed by a 6-month maintenance period, in which braces were worn for 12 hours per day. Patient satisfaction was investigated via telephone survey.

Results A total of 320 patients were included in this study. The average age was 13 years, and 280 were males (87.5%). The median follow-up period was 42 months (13-68). Good compliance was observed in 286 patients (89.4%; compliance group). In this group, the initial Haller index significantly increased from 2.20 \pm 0.31 to 2.59 \pm 0.38 after the 6-month therapy period (p = 0.001). After the 6-month period, 255 patients (89.1%) and 31 patients (12.1%) in the compliance group were very satisfied and satisfied, respectively. Satisfaction at the last follow-up via telephone survey was very satisfied in 250 patients (87.4%) and satisfied in 36 (12.6%). In the compliance group, no patient needed compressive braces again after the therapy period.

Keywords

chest wall deformity
 conclusion
 Given the findings presented in this study, compressive brace therapy
 pectus carinatum
 appears to be a relatively simple and safe method with good long-term outcome in

compressive brace treating patients with PC.



Long-Term Results of Compressive Brace Therapy for Pectus Carinatum

Duk Hwan Moon¹ Min Kyun Kang¹ Hye Sun Lee² Sungsoo Lee¹

 Department of Thoracic and Cardiovascular Surgery, Gangnam Severance Hospital, Seoul, the Republic of Korea
 Biostatistics Collaboration Unit, Gangnam Severance Hospital, Seoul, the Republic of Korea Address for correspondence Sungsoo Lee, Department of Thoracic and Cardiovascular Surgery, Gangnam Severance Hospital, 211 Eonjuro, Gangnam-Gu, Seoul 06273, the Republic of Korea (e-mail: CHFSTLF8/whk a.c)

Thorac Cardiovasc Surg 2019;67:67-72.

Abstract

Background Pectus carinatum (PC) is one of the most common types of congenital chest wall deformity. Recently, noninvasive compressive brace therapy has been more frequently used than invasive surgical correction to treat PC. Hence, the purpose of this study was to determine the long-term outcome of compressive brace therapy. **Methods** We retrospectively reviewed patients with PC who underwent compressive brace therapy between January 2014 and December 2016. All patients underwent a 2-week compression period, in which braces were worn for 20 hours per day, Followed by a 6-month maintenance period, in which braces were worn for 12 hours per day. Patient satisfaction was investigated via telephone survey.

Results A total of 320 patients were included in this study. The average age was 13 years, and 280 were males (87.5%). The median follow-up period was 42 months (13-68). Good compliance was observed in 286 patients (89.4%; compliance group). In this group, the initial Haller index significantly increased from 2.20 ± 0.31 to 2.59 ± 0.38 after the 6-month therapy period (p = 0.001). After the 6-month period, 255 patients (89.1%) and 31 patients (12.1%) in the compliance group were very satisfied and satisfied, respectively. Satisfaction at the last follow-up via telephone survey was very satisfied in 250 patients (87.4%) and satisfied in 36 (12.6%). In the compliance group, no patient needed compressive braces again after the therapy period.

Keywords

chest wall deformity
 Conclusion Given the findings presented in this study, compressive brace therapy
 pectus carinatum
 appears to be a relatively simple and safe method with good long-term outcome in
 compressive brace
 treating patients with PC.



Thoracic Outlet Syndrome (TOS)

TOS is a group of anatomically related, conditions caused by <u>compression of neurovascular</u> <u>structures that serve the upper extremity</u>.

Thoracic Outlet Syndrome (TOS)

TOS is a group of anatomically related, conditions caused by <u>compression of</u> <u>neurovascular structures that serve the upper extremity</u>.

- ✓ Scalene triangle
- ✓ Costoclavicular space
- ✓ Pectoralis minor space



Thoracic Outlet Syndrome (TOS) - Type

Туре	Characteristics
Neurogenic TOS	Caused from brachial plexus compression Symptoms include pain , dysesthesia , numbness , weakness – not localized in specific peripheral nerve distribution
Venous TOS	Caused from subclavian vein compression Symptoms include swelling , paresthesias in the fingers
Arterial TOS	Caused from subclavian artery compression Almost always associated with a cervical rib or anomalous rib Symptoms include hand ischemia with pain, pallor, paresthesia, coldness

Thoracic Outlet Syndrome (TOS) - Cause

Congenital abnormality

- **Cervical rib**
- Prolonged transverse process
- Muscular abnormality(ant. scalene m., sickle-shaped scalene m.)
- Fibrous connective tissue anomalies.

Trauma

Whiplash injury

Repetitive strain

Etc.

Tumor Hyperostosis Osteomyelitis



Thoracic Outlet Syndrome (TOS) - Surgery

Common Surgical Approaches for Thoracic Outlet Syndrome			
Surgical Approach	Characteristics/Proposed Advantages	Disadvantages	
Transaxillary	Most commonly used approach	Risk of iatrogenic brachial plexus injury ³²	
	Allows more complete exposure of first rib		
	More cosmetic scar		
	No retraction of neurovascular structures necessary for first rib removal ³¹		
Supraclavicular	Allows better exposure of the middle and upper trunks, neck of the first rib, and anterior and middle scalene muscles ³²	Retraction of brachial plexus and vascular structure necessary for complete first rib removal ³¹	
	May also allow effective first rib resection ³²		
	Allows vascular reconstruction ^{33,35}		
Posterior	Favored for recurrent TOS and in cases of prior anterior neck surgery	Requires extensive muscle dissection that can lead to postoperative shoulder dysfunction	
	May allow better exposure of proximal elements of the brachial plexus ³⁴	Risk of injury to the long thoracic, dorsal scapular, and accessory nerves ³⁴	

