

## เวชสารแพทย์ทหารบก Royal Thai Army Medical Journal

กรกฎาคม-กันยายน 2547 ปีที่ 57 ฉบับที่ 3

### นิพนธ์ต้นฉบับ

# Comparison of Phenol and Ethanol in Decreasing Spastic Calf Muscle in Cerebral Palsy\*

A Khunphasee, S Thinsheelong, S Hemtasilpa, T Sirimattayapant, P Aimpun\*, G Teeranet

Department of Physical Medicine and Rehabilitation Phramongkutklao Hospital, \*Department of Military and Community Medicine, Phramongkutklao College of Medicine

Objective: To determine the efficacy of phenol and ethanol for treatment of spastic gastrosoleus muscles in cerebral palsy. Design: Double-blinded, randomized clinical trial. Setting: Department of Physical Medicine and Rehabilitation, Phramongkutklao Hospital. Material and Methods: The 51 individuals (28 boys,23 girls) with mean age of 88.7 months were recruited and randomly assigned into 2 groups. Intervention: Intramuscular neurolysis of spastic gastrosoleus muscles managed by 5% phenol and 50% ethyl alcohol in group I and II, respectively and followed-up immediately and then at 7 days, 28 days, and 84 days. Outcome Measures: The severity of spasticity as assessed by Modified Ashworth Scale (MAS) score and the ankle passive range of motion (PROM). Results: Both groups had significant improvement in all outcomes at follow up statistically significant within 84 days. Conclusion: Phenol and ethanol preliminary provide good relief of spastic gastrosoleus in cerebral palsy in this limited study. The results emphasize the additional investigation in a long term follow-up.

**Key Words:** ● Intramuscular neurolysis ● Phenol ● Ethanol ● Spasticity ● Cerebral palsy RTA Med J 2004;57:127-34.

In cerebral palsy children, one of their important problem is muscular spasticity effects poorer life quality from various complications, especially, difficulty in standing and walking.

In management of spasticity, Biner and Eng<sup>1</sup> men-

tioned that spasticity treatment by taking medicine can temporary reduce spasticity with some results i.e. being weak, drowsy and disturbed daily activity and learning of patients. Local treatment by peripheral nerve block and motor point block by 5% phenol and 50% ethanol is more suitable for those patients.

So, for severe spasticity patients, treatment by chemical neurolysis such as phenol, ethanol and botu-

**เวชสารแพทย์ทหารบก** ปีที่ 57 ฉบับที่ 3 กรกฎาคม-กันยายน 2547

ได้รับต้นฉบับเมื่อ 18 สิงหาคม 2547 ได้ให้ตีพิมพ์เมื่อ 14 กันยายน 2547 ต้องการสำเนาต้นฉบับติดต่อ พ.อ.อารมณ์ ขุนภาษี กองเวชศาสตร์พื้นฟู โรงพยาบาลพระมงกุฎเกล้า ถนนราชวิถี เขตราชเทวี กทม. 10400

<sup>\*</sup>Approved by the Ethic Committee of Phramongkutklao Hospital

linum toxin is the option before surgery<sup>2</sup>. Phenol solution<sup>3</sup> contains a lot of chemical substances which are crystalline phenol, sodium bisulfate, sodium EDTA and water. It is different from ethanol solution<sup>4</sup> as it contains of ethanol in water only. The resemblance is phenol effected by reducing fat and denaturing tissue protein<sup>5</sup>; ethanol effected by inducing protein coagulation and protoplasm dehydration<sup>6</sup>. Five percent of more concentration of phenol effects some protein coagulation and tissue necrosis causing segmented demyelination and thrombosis of blood vessel supplies the nerve and development of Wallerian degeneration afterwards and finally muscular spasticity reduction. Ethanol effects by separating fat from nerve and precipitation of protein<sup>7</sup>, which effects swelling of Schwan cell and separation from myelin sheath causes Wallerian degeneration resulting reduction of muscular spasticity.

Both solutions are useful for spasticity management because of their effectiveness and their cost are not expensive. There are a lot of previous studies reported about their usefulness<sup>1,4,7-18</sup> but there is no any studied comparing their effectiveness. Thus, the purpose of this research is to compare the efficacy for muscular spasticity treatment betweenphenol and ethanol.

#### Material and Methods

#### Sample groups

Patients of Department of Physical Medicine and Rehabilitation Department of Phramongkut-klao Hospital during December 1, 2002 -March 12, 2003 were sampling by stratified by stratified randomization. They were grouped according to Modified Ashworth Scale score (MAS score)<sup>19</sup>.

And random treatedby these two solutions. Patients of the research were between 5 - 16 years with equines or equinovarus foot fromspasticity of gastrosoleus muscle with MAS score between 2 to 4. There was neither history allergy to phenol and ethanol. Nor kidney or liver diseases, nor chemical neurolysis. The second injection must be done after the first at least six months. The patients had to stop taking anti-spasticity drug at least 1 week before injection and throughout the research period. One leg was deemed as a case. Before being a part of the research, the patients and their cousins or parents acknowledged through information sheet, explained and replied to their questions, and consented in written to be a part of this research.

#### Materials

- Five percents of phenol in water was equally contained 3 mL. It was clear, mentioned by code instead of its exact medicine name, which would be disclosed after the research end.
- EMG ( Medelec V premier plus). Electrode and stimulator
- 3. Teflon coated 22 gauge needle, and 5 mL. syringe
- 4. Goniometer

#### Procedure

The solutions were prepared by medical personal and the doctor who made intramuscular neurolysis<sup>8</sup> did not know what solution was given to the patient. And all injections were done by the same doctor. The procedure were done by using surface stimulator to find the proper area for injection. Then, intramuscular neurolysis was done with Teflon coated 22-gauge needle and the intensity of electricity is 1 mA., 0.1 ms. or less.

#### Measurement

The examiner who measure MAS score and PROM

will not know what solution that the patient was given. The measurement were done in supine position. PROM of ankle joint were done<sup>20</sup>. For grading MAS score were done in 0,1,2,3,4,5 instead of 0,1,1+,2, 3,4 for the convenient in calculation. The measurements were done in every subjects before injection, immediate after injection, 7<sup>th</sup> day, 28<sup>th</sup> day, and 84<sup>th</sup> day post injection.

#### Result

There were 51 subjects. They are 28 boys and 23 girls. The first group: 24 patients treated by phenol were 11 boys (45.80%) and 13 girls (54.20%) at average age  $93.46\pm24.75$  months (67-143 months), with average weight 18.34±4.29 kg (10.4-26 kg). MAS level 2 was found in 8 patients (33.30%), level 3 was 11 patients (45.80%), level 4 was 5 patients (20.80%). Average MAS score was between 3.88±0.74(3-5) while average PROM was 42.50±10.53 (20-65 degree). The second group: 27 patients treated by ethanol were 17 boys(63%) and 10 girls (37%) at average age of 83.89 ? 24.38 months (54-137 months). Average weight 17.47±3.67 kg (11-26 kg). MAS level 2 was found in 8 patients (29.60%), level 3 in 11 patients (40.70%), level 4 in 8 patients (29.60%). Average MAS score was between 4.00±0.78<sup>35</sup> while average PROM was 40.93±9.91 degree (20-60 degree).

There is no significant difference between phenol and ethanol group in age, body weight, and dose as show in table 1, 2 and 3

Comparison of average MAS score between phenol and ethanol group was found no significant difference before injection, immediate after injection, 7<sup>th</sup> day, 28<sup>th</sup> day, and 84<sup>th</sup> day post injection. And The MAS score between before treatment and after treatment is significant different in both groups (p<0.001) as shown in figure 1 and table 4.

The comparison of average ankle PROM between phenol and ethanol group showed no significant difference before injection, immediate after injection, 7<sup>th</sup> day, 28<sup>th</sup> day, and 84<sup>th</sup> day post injection. And the ankle PROM between before treatment and after treatment is significant different in both groups (p<0.001) as shown in figure 2 and table 5.

Average MAS score in phenol group and ethanol group were found significant difference between before injection and after injection (immediate after injection, 7<sup>th</sup> day, 28<sup>th</sup> day, and 84<sup>th</sup> day post injection) as shown in table 6 and 7 (p<0.001)

Average ankle PROM in phenol group and ethanol group were found significant difference between before injection and after injection (immediate after injection, 7<sup>th</sup> day, 28<sup>th</sup> day, and 84<sup>th</sup> day post injection) as shown in table 8 and 9 (p<0.05)

From this study we found 3 allodynia cases which spontaneous recovery in 2 cases and cure after treatment in 1 case.

#### Discussion

Because of various differences of technique means

**Table 1** Age (months) of group1 (phenol) and 2 (ethanol)

Age	group	N	x	SD	t-value	df	p-value
I	Phenol	24	93.46	24.75	1 200	40	0 171
E	Ithanol	27	83.89	24.38	- 1.389	49	0.171

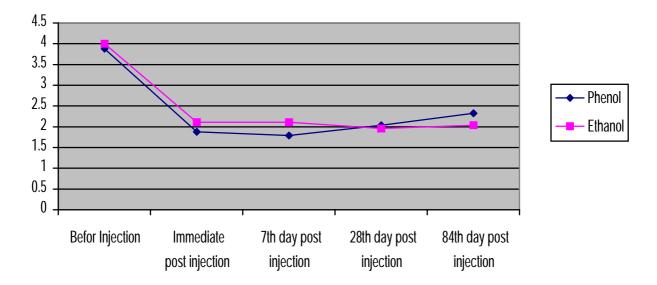


Table 2 Weight (Kg) of group 1(phenol) and 2 (ethanol)

Weight	group	N	X	SD	t-value	df	p-value
Phe	enol	24	18.34	4.29	0.700	49	0.427
Eth	anol	27	17.47	3.67	- 0.780	49	0.437

Table 3 Quantity of both substances (CC) for treatment of group 1 (phenol) and 2 (ethanol)

Quantity	group	N	x	SD	t-value	df	p-value
Pher	nol	24	3.48	0.906	0.270	40	0.707
Etha	nol	27	3.40	0.735	- 0.378	49	0.707

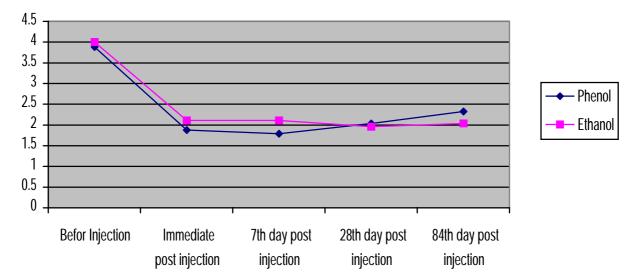


Figure 1 Show MAS score in average of group 1 and 2 by following up for a period after treatment test

Table 4 Show an analysis of variance of MAS score between group 1 (phenol) and 2 (ethanol)

ANOVA Source	SS	df	MS	F	p-value
Between subject	214.737	50			
Between groups	0.237	1	0.237	0.054	0.817
Error within group	214.500	49	4.378		
Within subject	224.54	204			
Between time	149.121	4	37.280	101.121	0.000
Between time with group	3.160	4	0.790	2.143	0.077
Between time with error within group	72.259	196	0.369		
Total	439.277	254			

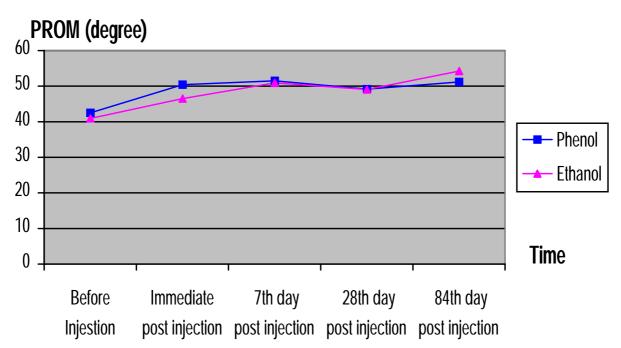


Figure 2 Show average scores of ankle PROM from of group 1 and group 2 by following up for a period after treatment test

Table 5 Show ANOVA result of ankle PROM. between group 1 (phenol) and 2 (ethanol)

ANOVA Source	SS	df	MS	F	p-value
Between subject	11,729.608	50			
Between groups	24.816	1	24.816	0.104	0.749
Error within group	11,704.792	49	238.873		
Within subject	17,108.274	204			
Between time	3,643.764	4	910.941	13.585	0.000
Between time with group	322.195	4	80.549	1.201	0.312
Between time with error within group	13,142.315	196	67.053		
Total	28,837.882	254			

Table 6 Compare the MAS score differences at any determined times for group 1 (Phenol)

Time	Mean	Standard	Sig.
	difference	error	
Immediate post injection	2.00	0.31	0.000
7 <sup>th</sup> day post injection	2.08	0.31	0.000
28 <sup>th</sup> day post injection	1.83	0.31	0.000
84 <sup>th</sup> day post injection	1.54	0.31	0.000
	Immediate post injection  7 <sup>th</sup> day post injection  28 <sup>th</sup> day post injection	Immediate post injection 2.00 7 <sup>th</sup> day post injection 2.08 28 <sup>th</sup> day post injection 1.83	Immediate post injection2.000.317th day post injection2.080.3128th day post injection1.830.31

Table 7 Compare the MAS score differences at any determined times for group 2 (Ethanol)

Time	Mean	Standard	Sig.
	difference	error	
Immediate post injection	1.89	0.30	0.000
7 <sup>th</sup> day post injection	1.89	0.30	0.000
28 <sup>th</sup> day post injection	2.04	0.30	0.000
84 <sup>th</sup> day post injection	1.96	0.30	0.000
	Immediate post injection  7 <sup>th</sup> day post injection  28 <sup>th</sup> day post injection	Immediate post injection 1.89 7 <sup>th</sup> day post injection 1.89 28 <sup>th</sup> day post injection 2.04	Immediate post injection1.890.307th day post injection1.890.3028th day post injection2.040.30

Table 8 Compare the PROM differences of ankles at any determined times for group 1 (Phenol)

Time	Mean	Standard	Sig.
	difference	error	
Immediate post injection	-7.92	2.93	0.008
7 <sup>th</sup> day post injection	-8.96	2.93	0.003
28 <sup>th</sup> day post injection	-6.67	2.93	0.025
84 <sup>th</sup> day post injection	-8.75	2.93	0.003
	Immediate post injection  7 <sup>th</sup> day post injection  28 <sup>th</sup> day post injection	Immediate post injection -7.92 7 <sup>th</sup> day post injection -8.96 28 <sup>th</sup> day post injection -6.67	Immediate post injection-7.922.937th day post injection-8.962.9328th day post injection-6.672.93

Table 9 Compare the PROM differences of ankles at any determined times for group 2 (Ethanol)

	Time	Mean	Standard	Sig.
		difference	error	
Before injection	Immediate post injection	-5.56	2.72	0.043
	7 <sup>th</sup> day post injection	-10.00	2.72	0.000
	28 <sup>th</sup> day post injection	-8.15	2.72	0.003
	84 <sup>th</sup> day post injection	-13.33	2.72	0.000

and concentration of solution for spasticity treatment so there are various differences of results from the previous of studies<sup>1,4,8,9,10,11,12,13,14,15,16</sup>. Aim of this research is to study only on efficiency of a treatment of 5% phenol comparing with 50% ethanol by intramuscular neurolysis technique.

From this study, we found that the results of 5% phenol and 50% ethanol throughout the period of follow up (84days) are good from MAS score that decreased (p<0.001) and ankle PROM that increased (p<0.05) comparing with the data before injection. However, from Kong and Chua study<sup>4</sup> found that in stroke patients whom treated by 50% ethanol in management for elbow flexor spasticity, the effect could last for 6 month. In this study we could not study so long as in Kong and Chua study because, for these spastic diplegic children, we should not wait too long to see the effect from only single modality, it is better to combine treatment with other modalities such as physical therapy, orthotic device and surgical management to make the best profit to the patients.

Complications from this study were found only in 3 cases. All of them were allodynia whom 1 case from phenol injection and the other 2 cases from ethanol injection. And in these 3 cases, one case from phenol and the other one from ethanol injection developed spontaneous recovery in 1 week. For the third case, pain was cure after the treatment with oral prednisolone for 1 week. One of the most important factors for this low incidence of complication is the technique that we used which is intramuscular neurolysis technique. Comparing with peripheral nerve block technique that if major mixed nerves with many sensory axons are injected, there are 10% to 30% risk of painful paresthesias<sup>17,18</sup>.

#### Summary

Within 84 days of study, 5% phenol and 50% etha-

nol intramuscular neurolysis injection showed a good result in decreased spasticity of gastrocnemious muscle in spastic diplegic children.

#### Rererences

- Binder H,Eng GD. Rehabilitation Management of Children with Spastic Diplegic Cerebral Palsy. Arch Phys Med Rehabili 1989, 70:482-9.
- Little JW, Massagli TL. Spasticity and Associated Abnormalities of Muscle Tone. In: Delisa JA.ed. Rehabilitation Medicine 3<sup>rd</sup> ed. Philadelphia: Lippincott-Raven Publisher 1998:997-1013.
- Awad EA, Dykstra D. Treatment of Spasticity by Neurolysis.
   In: Kottke FJ, Lehmann JF. eds. Krusen's Handbook of Physical Medicine and Rehabilitation. 4<sup>th</sup> ed. Philadelphia: WB Saunder, co., 1990:1154-61.
- Kong KH, Chua KSG. Neurolysis of the Musculocutaneous Nerve with Alcohol to Treat Poststroke Elbow Flexor Spasticity. Arch Phys Med Rehabil 1999,80: 1234-6.
- Harvey SC. Antiseptics and Disinfectants. In: Gilman AG, Goodman LS,Rall TW,Murad F. eds. Goodman and Gilman's The Pharmacological Basis of Therapeutic 8<sup>th</sup> ed. New York: Macmillan Publishing, co., 1985:959-75.
- Richie JM. The Aliphatic Alcohols. In: Goodman LS, Gilman AG. Eds.Goodman and Gilman's The Pharmacological Basic of Therapeutics. 6<sup>th</sup> ed New York: Macmillan Publishing, co., 1980:376-90.
- Bell KR. The Use of Neurolytic Blocks for the Management of Spasticity. Physical Medicine and Rehabilitation Clinics of North America 1995,6:885-95.
- Easton JKM, Ozel T, Halpern D. Intramuscular Neurolysis for Spasticity in Children. Arch Phys Med Rehabil 1979,60:155-8.
- Cole TM, Tobis JS. Measurement of Skeletal Function. In:
   Kottke FJ, Lehmann JF. eds Krusen's Handbook Physical Medicine and Rehabilitation. 4<sup>th</sup> ed. Philadelphia: WB Sounders, co., 1990:20-71.
- Halpern D, Meelhuysen FE. Phenol Motor Point Block in the Management of Muscular Hypertonia. Arch Phys Med Rehabil 1966,47:659-64.
- Awad EA, Phenol Block for Control of Hip Flexor and Adductor Spasticity. Arch Phys Med Rehabil 1972:554-7.
- Braun RM, Hoffer MM, Mooney V, McKeever J, Roper B. Phenol Nerve Block in the Treatment of Acquired Hemiplegic in the Upper limb J Bone Joint Surg Am 1973,55 580-5.
- Garland DE, Lucie RS, Waters RL. Currant Used of Open Phenol Nerve Block for Adult Acquired Spasticity Clin Orthop

134

- 1982,165:217-22.
- Aimprasittichai, Tosayanonda O. Phenol Motor Point Block in Spastic Diplegia. J Thai Rehabil 1991,1:23-26.
- Khunphasee A, Aimprasittichai S, Intharakumhang P, Phatharawaratham S, Theranethara K, Tosayanonda O, Khunadorn F. Phenol Block in Spasticity. J Thai Rehabil 1991,1:15-18.
- Tunchavanit K, Khunphasee A, Khunadorn F. Phenol Intramuscular Neurolysis Longterm Study at Pramongkutklao Hospital. J Thai Rehabil 2001,10:117-26.
- 17. Glen MB. Nerve blocks. In: Glenn MB, Whyte J, eds. The

- practical management of spasticity in children and adults. Philadelphia: Lea & Febiger, 1990.
- Khalili AA, Betts HB. Peripheral nerve block with phenol in the management of spasticity: indications and complications. JAMA 1967:200:1155-7.
- Bohannon RW, Smith MB, Interrater Reliability of Modified Ashworth Scale of Muscle Spasticity. Phys Ther 1987,67:206-7.
- Carpenter EB, Seitz DG, Intramuscular alcohol as an Aid in Management of Spastic Cerebral Palsy. Dev Med Child Neurol 1980,22:479-501.

# การศึกษาเปรียบเทียบการลดการหดเกร็งของกล้ามเนื้อน่อง ในผู้ป่วยสมองพิการระหว่างฟินอลและเอธานอล\*

อารมย์ ขุนภาษี, เศรษฐพงศ์ ถิ่นชีลอง, สมเกียรติ เหมตะศิลป, ธิติมา ศิริมาตยาพันธุ์, พจน์ เอมพันธุ์\*, ไกรวัชร ธีรเนตร

กองเวชศาสตร์ฟื้นฟู, โรงพยาบาลพระมงกุฎเกล้า, \*ภาควิชาเวชศาสตร์ทหารและชุมชน วิทยาลัยแพทยศาสตร์พระมงกุฎเกล้า

วัตถุประสงค์: เพื่อเปรียบเทียบประสิทธิภาพของฟืนอลและเอธานอล ที่ใช้ในการรักษาการหดเกร็งของกล้ามเนื้อที่ทำหน้าที่กระดกข้อ เท้าลงในผู้ป่วยสมองพิการ รูปแบบการวิจัย: Double - blinded, randomized clinical trial สถานที่ทำการวิจัย: กองเวชศาสตร์ พื้นฟู โรงพยาบาลพระมงกุฎเกล้า กลุ่มที่ถูกทำการวิจัย: ผู้ป่วยสมองพิการ 51 ราย เด็กชาย 28 ราย เด็กหญิง 23 ราย อายุเฉลี่ย 88.7 เดือน แบ่งผู้ป่วยเป็น 2 กลุ่ม วิธีการ: กลุ่มที่ 1 จำนวน 24 ราย รักษาโดย 5%ฟืนอล กลุ่มที่ 2 จำนวน27 ราย รักษาด้วย เอธานอล รักษาการหดเกร็งด้วยวิธี Intramuscular Neurolysis วัดผลโดยการให้คะแนน Modified Ashworth Scale (MAS) และ พิสัยการเคลื่อนไหวข้อเท้าโดยใช้แรงของผู้ตรวจ (PROM) ติดตามผลการรักษาทันทีหลังฉีด, หลังฉีด 7 วัน, หลังฉีด 28 วัน, และหลัง ฉีด 84 วัน ผลการวิจัย: ทั้ง 2 กลุ่มมีผลการรักษาดีขึ้นอย่างชัดเจนตลอดเวลาที่ติดตามผล ไม่มีความแตกต่างกันอย่างมีนัยสำคัญทาง สถิติในช่วงเวลาติดตามผล 84 วัน ผลสรุป: ในเบื้องต้นฟืนอลและเอธานอลให้ผลการรักษาที่ดีในการลดการหดเกร็งกล้ามเนื้อน่อง ในผู้ป่วยสมองพิการ เนื่องจากมีข้อจำกัดในการคึกษาควรติดตามผลการรักษาที่นานขึ้น

**Key Words:** ● Intramuscular neurolysis ● Phenol ● Ethanol ● Spasticity ● Cerebral palsy เวชสารแพทย์ทหารบก 2547;57:127-34.

\*การศึกษานี้ได้ผ่านการพิจารณาเห็นชอบให้ดำเนินการได้โดยคณะอนุกรรมการจริยธรรมโรงพยาบาลพระมงกุฎเกล้า และสนับสนุนทุนการศึกษาวิจัยโดยโรงพยาบาลพระมงกุฎเกล้า