INTRODUCTION

Treatment options for primary spontaneous pneumothorax range from simple observation to surgery (1-3). Despite all efforts, recurrence has always been a lingering problem, and many intraoperative methods to induce pleural symphysis have been introduced. We analyzed the effects of chemical pleurodesis during thoracoscopic procedures. Between August 2003 and July 2005, 141 patients among indicated surgical treatment for primary spontaneous pneumothorax in two hospitals of our institution allowed this prospective study. The patients were randomly assigned to 3 groups: thoracoscopic procedure only (group A, n=50), thoracoscopic procedure and pleurodesis with dextrose solution (group B, n=49), and thoracoscopic procedure and pleurodesis with talc-dextrose mixed solution (group C, n=42). There was no significant difference in demographic data among the three groups. The two groups that underwent intraoperative pleurodesis had significantly longer postoperative hospital stays (A/B/C: 2.50±1.85/4.49±2.10/6.00±2.58 days; p=0.001) and a higher incidence of postoperative fever (A/B/C: 10.0/22.45/52.38%; χ²=21.598, p=0.00). No significant differences were found for recurrence rates or the number of postoperative days until chest tube removal. Therefore, the results of our study indicate that intraoperative chemical pleurodesis gives no additional advantage to surgery alone in deterring recurrence for patients with primary spontaneous pneumothorax. Thus, the use of such scarifying agents in the operating room must be reconsidered.

Key Words: Pneumothorax; Thoracic Surgery, Video-Assisted; Pleurodesis
without any identifiable underlying lung disease, trauma, or iatrogenic injury. All patients were positively identified to have pleural/subpleural blebs by thoracoscopic examination (with a 2-mm thoracoscope) (6) or by high-resolution computed tomography. In this prospective study, the patients were randomly assigned to 3 different groups by two surgeons: thoracoscopic bleb resection or electrical coagulation (TBR) only (50 patients; group A), TBR and pleurodesis with a 20% dextrose solution (200 cc) (49 patients; group B), and TBR and pleurodesis with a talc (2 g of asbestos free sterilized talc powder) -20% dextrose mixed solution (200 cc) (42 patients; group C).

In the early stage of this study, 31 operations were performed with a 10-mm thoracoscope, but all remaining 110 consecutive operations were performed by needlescopy (2-mm thoracoscopy) (7). According to their assigned groups, the patients received adjuvant pleurodesis after resection of blebs with mechanical suturing devices. The chemicals were instilled into the pleural cavity, mainly at the thoracoscopic procedure sites and apex of the lungs. In group C, talc powder was mixed with the dextrose solution to form a slurry before instillation. The solutions were allowed to drain freely after the patient was transferred to the general ward (average time of pleurodesis, 2 hr). No form of mechanical pleural abrasion was conducted in any of the patients. Postoperative pain was managed by oral nonsteroidal anti-inflammatory drugs (NSAIDs) and intermittent parenteral NSAIDs and meperidine for breakthrough pain. Patients were discharged 2 days after chest tube removal according to our institutional protocol.

Postoperative data was recorded included recurrence of pneumothorax, postoperative hospital stay, days until chest tube removal, occurrence of postoperative fever, and frequency of parenteral analgesics use. Patients follow-up was conducted by outpatient clinic visits and telephone interviews. Statistical analysis was done using SPSS for Windows (Ver. 10.0.7., Chicago, IL, U.S.A.). Analysis of variance (ANOVA) was used to compare patient characteristics, postoperative hospital stay, days until chest tube removal, and chest tube drainage amount. Chi-square test was used to compare categorical variables. All p values <0.05 were considered statistically significant.

### RESULTS

Patient characteristics are summarized in Table 1. The mean ages of patients were 22.26 yr for group A, 24.18 for group B, and 23.02 for group C. There were 46 males and 4 females in group A, 46 males and 3 females in group B, and 42 males and no females in group C. The three groups did not differ significantly in age, sex, or surgical approach. There were no operative deaths, and all procedures were uneventful. No patient underwent conversion to open thoracotomy.

Postoperative data are summarized in Table 2. Days until chest tube removal were 2.96±1.56 days for group A, 2.95±2.31 for group B, and 3.60±1.65 for group C (p=0.188) and did not differ significantly. Postoperative hospital stay and frequency of parenteral analgesics use per patient differed significantly among the 3 groups. The durations of postoperative hospital stay (days) were 2.50±1.85 in group A, 4.49±2.10 in group B, and 6.00±2.58 in group C (p=0.001), and frequencies of parenteral analgesics use per patient were 2.37±3.51 for group A, 3.42±3.48 for group B, and 5.02±2.72 for group C (p=0.001). Post hoc studies (Duncan) showed that group C differed significantly from group A and B in postoperative stay and parenteral analgesics use.

### Table 1. Patient characteristics

| Group | A (total N=50) | B (total N=49) | C (total N=42) | Signif |
|-------|---------------|---------------|---------------|-------|
| Operation methods | TBR Only | TBR with 20% dextrose | TBR with talc-20% dextrose | |
| Male:female | 46 (92%):4 (8%) | 46 (93.9%):3 (6.1%) | 42 (100%):0 (0%) | NS |
| Mean age (Range: yr) | 22.26 (13-36) | 24.18 (16-45) | 23.02 (15-65) | NS |
| Site (right:left) | 24 (48%):26 (52%) | 23 (46.9%):26 (53.1%) | 18 (42.9%):24 (57.1%) | NS |

N, number of patients; Signif, statistical significance; NS, not significant.

TBR Only, thoracoscopic procedures without chemical pleurodesis; TBR with 20% dextrose, thoracoscopic procedures and chemical pleurodesis with a 20% dextrose solution; TBR with talc-20% dextrose, thoracoscopic procedures and chemical pleurodesis with a talc-20% dextrose solution.

### Table 2. Postoperative data

| Group | A (total N=50) | B (total N=49) | C (total N=42) | p value |
|-------|---------------|---------------|---------------|---------|
| Postoperative chest drainage (days) | 2.96±1.56 | 2.95±2.31 | 3.60±1.65 | 0.188 |
| Postoperative hospital stay (days) | 4.50±1.85 | 4.49±2.10 | 6.00±2.58* | 0.001* |
| Parenteral analgesics (frequencies/patient) | 2.37±3.51 | 3.42±3.48 | 5.02±2.72* | 0.001* |
| Rate of fever (>37.5°C) | 10% (N=5) | 22.45% (N=11) | 52.38% (N=22) | 0.01 |
| Recurrence rate | 6% (N=3) | 2.04% (N=1) | 2.38% (N=1) | 0.504 |

N, number of patients. *: Significantly different group according to post hoc studies; †, statistically significant.
Five patients in group A, 11 in group B, and 22 in group C experienced postoperative fever ($\chi^2=21.598, p=0.00$), showing a significant difference.

Mean follow-up durations were 20.22±9.71 months in group A, 23.65±2.51 months in group B, and 17.93±3.64 months in group C. During follow-up, recurrent ipsilateral pneumothorax was noted in 5 of the 141 patients (3.54%); 3 in group A (6.0%), 1 in group B (2.04%), and 1 in group C (2.38%). There was no statistically significant difference in recurrence rates among the three groups ($\chi^2=1.372, p=0.54$). Of the 5 patients with ipsilateral recurrence, 2 underwent reoperation by video-assisted thoracoscopic surgery (VATS), 2 were treated by chest tube drainage, and 1 patient received conservative management with inhaled oxygen.

**DISCUSSION**

VATS is currently well accepted as a safe and effective procedure for the management of primary spontaneous pneumothorax. After resection or ligation of blebs and electrocautery of subpleural blebs many adjuvant procedures have been used to achieve pleural symphysis in order to further reduce recurrence rates. Partial pleurectomy and pleural abrasion have shown recurrence rates of 2-4.4% (8, 9) and 3-3.6% (10, 11), respectively. Chemical pleurodesis has also been tried with talc, and minocycline and have shown recurrence rates of 1.73-5% (12, 13) and 2.9% (14), respectively. Other reported methods include electrocautery of the pleura and ablation with Nd:YAG laser.

At our institution, VATS is conducted for pneumothorax with a 2-mm thoracoscope and instruments from 2003. Surgical staplers are introduced via the 11.5 mm trocar site at the thoracotomy wound, which most of the patients already have before the operation. Although a recent study by Chang (15) showed a higher rate of recurrence by needlescopic VATS (procedure using instruments with an external diameter ≤3 mm [7]) compared to the same procedures performed by conventional VATS, our experience shows that recurrence rates are comparable to conventional VATS (16-19). We hypothesized that additional chemical pleurodesis may further decrease recurrence rates after bleb resection.

The selection of the agents for chemical pleurodesis was based on their low cost, ease of administration, and reported high effectiveness (20-22). Our results show that 20% dextrose solution and talc are effective in reducing recurrence rates, showing comparable recurrence rates with those of previous studies with other agents in the literature. However, there is still much concerns in regards to acute and chronic side effects of talc. During this study, we did not experience any cases of postoperative empyema, arrhythmia, or respiratory failure, all of which have been reported in the literature (23). Although chemical pleurodesis did not increase the duration of postoperative chest drainage, a significantly greater number of patients with fever was noted and due to more frequent chest pain more parenteral analgesics was used in the pleurodesis groups, especially the talc group. There is also the issue of possible oncogenic risk and difficulty in performing future thoracoscopy or thoracotomy caused by the creation of tight adhesions (11).

Our results showed that VATS procedures for primary spontaneous pneumothorax are safe and effective. However, we feel that practice of additional chemical pleurodesis for deter- ring recurrence should be reconsidered. Chest pain and fever associated with this procedure may cause unnecessary suffering for the patients and extension of postoperative hospital stay.

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