Timing of Brain Expansion and Recurrence after Surgery of Chronic Subdural Hematoma

Sarun Nunta-aree MD, PhD*, Tewajetsada Paruang MD**, Bunpot Sitthinamsuwan MD, MSc*

* Division of Neurosurgery, Department of Surgery, Faculty of Medicine Siriraj Hospital, Mahidol University, Bangkok, Thailand

** Division of Surgery, Somdech Phra Pinklao Hospial, Bangkok, Thailand

Background: Chronic subdural hematoma (CSDH) is common in neurosurgical practice. The exact timing of brain expansion is still unknown.

Objective: To study prospectively the timing of full brain expansion and timing of recurrence after surgery of CSDH.

Material and Method: After burr hole and closed-system drainage for the treatment of CSDH, serial CT brains were done at 24 hours after removal of draining system, 1 week, 2 weeks, 4 weeks, 8 weeks postoperatively or whenever abnormal neurological symptoms being suspicious for recurrence occurred.

Results: 102 CSDHs from 75 patients were included in the study. A total 450 serial CT brains were done and reviewed. Full brain expansion after the surgery occurred 1% between the 1st and 2nd weeks, 5.9% between the 2nd and 4th weeks, 80.4% between the 4th and 8th weeks and 12.7% after the 8th weeks. The incidence of recurrent CSDH was 9.8%. In patients with recurrent CSDH, the recurrence was 10% before removal of draining system, 70% between the 1st and 2nd week, 10% between the 2nd and 4th week and 10% after the 8th week.

Conclusion: Brain expansion after surgery of CSDH is a very slow process. There is a discrepancy between the timing of full brain expansion and timing of recurrence. Recurrent CSDH mostly occurred between the 1st and 2nd week while full brain expansion mostly occurred after the 4th week. The period between the 1st and 2nd week should be the crucial period of clinical follow-up. Persistence of subdural space on a random postoperative CT brain, without any symptoms, should be considered as a normal process of brain expansion.

Keywords: Chronic subdural hematoma, Brain expansion, Recurrence

J Med Assoc Thai 2017; 100 (Suppl. 3): S59-S64 Full text. e-Journal: http://www.jmatonline.com

Chronic subdural hematoma (CSDH) is a common neurosurgical disease. Its conventional treatment is burr hole with or without closed-system drainage. The method is the most widely accepted surgical option⁽¹⁾ and has recurrence rates reported between 3 to $20\%^{(2-6)}$. Typically, the brains do not expand immediately after the surgery but gradually expand over certain periods, which, to our knowledge, the timing of brain expansion has never before been adequately investigated. If postoperative CT or MRI brain is done during this time, persistence of subdural collection will be seen. Similarly, the timing of recurrent CSDH after the operation has not been sufficiently investigated. These make for uncertainty whether the persistence of subdural space seen on the

Correspondence to:

Nunta-aree S, Division of Neurosurgery, Department of Surgery, Faculty of Medicine Siriraj Hospital, Mahidol University, Bangkok 10700, Thailand.

Phone: +66-2-4198003, Fax: +66-2-4113006

E-mail: sarunsisna@gmail.com

postoperative images is a normal evolution process of brain expansion or an abnormal evolution process of recurrent CSDH. If there is a discrepancy between the timing of brain expansion and recurrence after surgery of CSDH, a more accurate prediction and identification of patients who have a risk of recurrent CSDH can be done.

The primary objective of the present study is to determine the timing of brain expansion and recurrence after surgery of CSDH. The secondary objective is to determine the incidence of recurrent CSDH and factors associated with the recurrence. The findings will be used for the postoperative surveillance of the high risk group of recurrence.

Material and Method

Patient population and data collection

A prospective controlled study of CSDH was done at our institute during 2006 to 2009. Patients' data were collected which included age, sex, occupation, characteristic of injury, symptoms, alcohol abuse,

underlying diseases, previous medications, duration between head injury to initial CT scan of the brain. Relevant laboratory, such as platelet count and coagulogram, were also recorded.

Characteristics of CSDH on CT brain, including location, size, side and density, were collected. Internal architecture of hematoma was stratified into homogenous stage, separated stage (hematoma containing two components of different densities with a clear boundary lying between the lower densities component located above a higher density component), laminar stage (having a thin high density layer along the inner membrane), trabecular stage (having high density septums running between the inner and outer membrane on a low to isodensity background)⁽⁶⁾. Severity of brain atrophy on CT scan was also taken into account. Mild brain atrophy was defined as mild cerebral and/or cortical atrophy with slight enlargement of the lateral ventricle. Moderate brain atrophy was well established cerebral and/or cortical atrophy, ballooning of the lateral ventricle and/ or involvement of the other ventricles and severe brain atrophy was defined as extreme atrophy and/or extreme dilatation of the ventricular system⁽⁷⁾.

The surgery was performed under general anesthesia. Two burr holes, drainage of hematoma, irrigation and insertion of closed-system drainage were done in all patients by nonspecific random surgeons.

All patients were investigated by serial postoperative CT scan before removal of draining system, at 1 week, 2 weeks, 4 weeks, 8 weeks postoperatively and whenever any abnormal neurological symptoms suspected for recurrence occurred.

The recurrence of subdural hematoma was defined as recurrence of clinical symptoms or development of new symptoms and presence of subdural space on the serial CT brain being larger than the previous CT brain.

Statistical analysis

Results were demonstrated as descriptive statistics, which continuous data were described as mean, standard deviation, median, range, and categorical data, was described as frequencies and in percentages. All data were analyzed using SPSS for Windows version 10.0 (SPSS Inc., Chicago IL, USA).

Results

A total number of 102 CSDHs from 75 patients and 450 serial CT brain were included in the study.

There were 79 lesions in men (77.5%) and 23 lesions in women (22.5%). The patients' age ranged from 1 to 88 years with a mean age being 60.67 years and median age being 66 years. There were 31 (30.4%) patients who had history of head trauma. The duration of injury ranges from 6 to 50 days with a mean duration of 21 days and median duration of 14 days.

The duration of symptoms ranged from 1 to 60 days with a mean time of 9.45 days and median time of 7 days. Headache was the most common presenting symptom, followed by weakness, alteration of consciousness, dizziness or unstable gait and seizure, respectively. There were 60 subjects (58.8%) who had coexisting diseases. Five patients (4.9%) had coagulopathy and 6 patients (5.9%) had thrombocytopenia. Thirty-one subjects (30.4%) took antiplatelet or anticoagulant. The most common drug was aspirin followed by warfarin and clopidigrel, respectively. Deterioration of consciousness and abnormal Glasgow Coma Scale score were found in 32.4% of patients. Demographic characteristics were summarized in Table 1.

Unilateral CSDH (48, 64%) was found more common than bilateral lesions (27, 36%) and was on the left side (60, 58.8%) more than the right side (42, 41.2%). The most common hematoma stage was separated (46, 45.1%) followed by homogenous (44, 43.1%), laminar (8, 7.8%) and trabecular stages (4, 3.9%), respectively. All hematomas were in the convexity area (102, 100%). Fronto-parieto-occipital was the most common location (83, 81.4%) followed by fronto-parietal (16, 15.7%), parieto-occipital (2, 2%) and fronto-temporal area (1, 1%). Brain atrophy was divided into 4 groups according to the severity as none (29, 28.4%); mild (35, 34.3%); moderate (24, 23.5%); and severe (14, 13.7%). Table 2 shows characteristics of CSDH and brain atrophy on CT brain.

The timing of full brain expansion and recurrence were observed by postoperative serial CT brain and were divided into 6 periods. Period 1 was before removal of the draining system. Period 2 was after removal of the draining system to the 1st week. Period 3 was the 1st week to the 2nd week. Period 4 was the 2nd week to the 4th week. Period 5 was the 4th week to the 8th week and Period 6 was after the 8th week. There were total 450 serial CT brain reviewed for the study. The brain expansion was found to be a delayed pattern in all patients and never been found before the 1st postoperative week. Indeed in most patients (80.4%), full brain expansion occurred after the 4th week with almost all patients (87.3%) experiencing full brain

Table 1. Demographic characteristics

Clinical symptoms Headache 69 (67.6%) Weakness 51 (50%) Alteration of consciousness 32 (31.4%) 25 (24.5%) Dizziness/unstable gait 3 (2.9%) Seizure Coexisting diseases Hypertension 30 (29.4%) Heart disease 15 (14.7%) Diabetes mellitus 11 (10.8%) Chronic renal failure 2 (2%) Alcoholism 2 (2%) Dyslipidemia 11 (10.8%) Other diseases 24 (23.8%) Bleeding disorders Coagulopathy 5 (4.9%) Thrombocytopenia 6 (5.9%) Use of drugs interfere platelet function or coagulation Aspirin 27 (26.5%) Warfarin 5 (4.9%) Clopidogrel 3 (2.9%) Glasgow Coma Scale score on admission 15 69 (67.6%) 13 to 14 18 (17.6%) 9 to 12 13 (12.7%) <9 2 (2%)

expansion by the 8th week (Table 3).

Ten patients (9.8%) developed recurrent CSDH. Seven patients (70%) developed recurrence between the 1st and 2nd weeks. There were only 3 patients who had recurrence of CSDH outside the period between the 1st and 2nd weeks. Two of them had coexisting thrombocytopenia. One patient had thrombocytopenia caused by acute hemorrhagic fever developed early recurrence before removal of drainage and another patient had thrombocytopenia due to aplastic anemia developed delayed recurrence after the 8th week. By excluding these 2 patients with coexisting thrombocytopenia, the possibility of recurrence between the 1st and the 2nd weeks increased to 87.5% and all recurrence (100%) happened before the 4th week.

Discussion

In studies of recurrent CSDH, several factors had been proposed as risk factors associated with recurrence such as advanced age⁽⁸⁾, coagulopathy^(9,10), atrophy of brain parenchyma^(8,11,12),

Table 2. Characteristic on CT brain

Thickness of CSDH	
Mean \pm SD (mm)	15.01 <u>+</u> 5.82
Median (mm)	13.5
Range (mm)	6-36
Laterality of CSDH $(n = 75)$	
Unilateral	48 (64%)
Bilateral	27 (36%)
Side of CSDH	
Right side	42 (41.2%)
Left side	60 (58.8%)
Internal architecture of CSDH	
Homogenous	44 (43.1%)
Separated	46 (45.1%)
Laminar	8 (7.8%)
Trabecular	4 (3.9%)
Location of CSDH	
Fronto-parietal	16 (15.7%)
Fronto-parieto-occipital	83 (81.4%)
Fronto-temporal	1 (1%)
Parieto-occipital	2 (2%)
Brain atrophy	
None	29 (28.4%)
Mild	35 (34.3%)
Moderate	24 (23.5%)
Severe	14 (13.7%)

Table 3. Timing of full brain expansion and recurrence of CSDH

D1. b.f	0 (00/)
P1: before removal of drainage	0 (0%)
P2: removal of drainage to 1 st week	0(0%)
P3: 1 st to 2 nd week	1 (1%)
P4: 2 nd to 4 th week	6 (5.9%)
P5: 4 th to 8 th week	82 (80.4%)
P6: after 8 th week	13 (12.7%)
Time of recurrent CSDH $(n = 10)$	
P1: before removal of drainage	1 (10%)
P2: removal of drainage to 1st week	0 (0%)
P3: 1 st to 2 nd week	7 (70%)
P4: 2 nd to 4 th week	1 (10%)
P5: 4 th to 8 th week	0 (0%)
P6: after 8 th week	1 (10%)

P = period

larger hematomas^(11,13), chronic alcoholism⁽¹⁴⁾, poor expansion rate of brain⁽¹⁵⁾, frequency of falling and head injury⁽¹³⁾, postoperative hematoma density⁽¹⁶⁾, separated type of CSDH^(6,17,18) and extension into cranial base⁽⁶⁾. Though there were total 102 CSDHs in our study, the

incidence of recurrence was low and we could not determine the significance of these risk factors.

Although we cannot clarify the definitive factors related to recurrent CSDH but the definitive timing of recurrent CSDH are observed. Most patients in the recurrent group, especially those who did not have coexisting bleeding disorder had recurrent CSDH happening between the 1st and 2nd weeks. Postoperative clinical observation in this period is crucial and CT brain should be done for any particular patients who develop symptom, even mild, in this period.

Full brain expansion after surgery of CSDH is usually considered as the goal of the operation and is the end point of postoperative follow-up. However, this issue has never been investigated before. The expansion rate and timing of full brain expansion are unknown. Our study clearly showed that brain expansion was always a delayed pattern. Most patients had their full brain expansion after the 4th week which was later than the timing of most recurrence. Several patients had even more delayed brain expansion after the 8th week. All these patients with much delayed brain expansion did well eventually, though the poor expansion rate was proposed as a risk factor for recurrence⁽¹⁵⁾. Thus, persistence of subdural space seen on postoperative CT scan without any clinical symptom should be considered as a normal process of brain expansion and should be managed conservatively, not to repeat the surgery or place a subduroperitoneal shunt. Concerning the significant difference of timing between recurrent CSDH and brain expansion and the insignificant prognostic value of delay brain expansion, benefit of a random postoperative CT brain, especially done after the 2nd week, without any clinical symptom cannot be identified. Only a subgroup of patients who develop recurrent symptom, especially between the 1st and 2nd weeks, that CT brain is highly recommended.

The drawback of this study is limited period of postoperative follow-up. Long-term clinical outcome and radiographic finding after 8th week are required to investigate the correlation between delayed brain expansion and recurrence of CSDH.

Conclusion

Recurrent CSDH mostly occurred between the 1st and 2nd week while full brain expansion mostly occurred after the 4th week. Follow-up of clinical symptoms, especially between the 1st and 2nd week, is more valuable than postoperative CT brain for detection of recurrence. Persistence of subdural space on a random postoperative CT scan, even long-lasting,

without any neurological symptoms, should be considered as a normal process of brain expansion and managed conservatively.

What is already known from this topic?

Risk of recurrent CSDH was associated with old age, abnormal coagulation, cerebral atrophy, large hematoma, poor brain expansion, frequent head injury and separated stage of CSDH.

What this study adds?

Full brain expansion after surgical drainage of CSDH is usually found after 4 weeks post surgery. One to two weeks after surgery is a critical period of recurrent hematoma. In cases with residual subdural fluid collection visible on CT brain without clinical symptoms of recurrent CSDH before 4 weeks after drainage, follow-up of CT brain is recommended.

Potential conflicts of interest

None.

References

- Wakai S, Hashimoto K, Watanabe N, Inoh S, Ochiai C, Nagai M. Efficacy of closed-system drainage in treating chronic subdural hematoma: a prospective comparative study. Neurosurgery 1990; 26: 771-3.
- 2. Arbit E, Patterson RH Jr, Fraser RA. An implantable subdural drain for treatment of chronic subdural hematoma. Surg Neurol 1981; 15: 175-7.
- 3. Kotwica Z, Brzezinski J. Chronic subdural haematoma treated by burr holes and closed system drainage: personal experience in 131 patients. Br J Neurosurg 1991; 5: 461-5.
- 4. Asano Y, Hasuo M, Takahashi I, Shimosawa S. Recurrent cases of chronic subdural hematoma—its clinical review and serial CT findings. No To Shinkei 1992; 44: 827-31.
- 5. Ernestus RI, Beldzinski P, Lanfermann H, Klug N. Chronic subdural hematoma: surgical treatment and outcome in 104 patients. Surg Neurol 1997; 48: 220-5.
- 6. Nakaguchi H, Tanishima T, Yoshimasu N. Factors in the natural history of chronic subdural hematomas that influence their postoperative recurrence. J Neurosurg 2001; 95: 256-62.
- Kraus J, Selecki BR. Brain atrophy and assessment of intellectual deterioration on the Wechsler Adult Intelligence Scale. J Nerv Ment Dis 1965; 141: 119-22
- 8. Stroobandt G, Fransen P, Thauvoy C, Menard E.

- Pathogenetic factors in chronic subdural haematoma and causes of recurrence after drainage. Acta Neurochir (Wien) 1995; 137: 6-14.
- 9. Diamond T, Gray WJ, Chee CP, Fannin TF. Subdural haematoma associated with long term oral anticoagulation. Br J Neurosurg 1988; 2: 351-5.
- Mattle H, Kohler S, Huber P, Rohner M, Steinsiepe KF. Anticoagulation-related intracranial extracerebral haemorrhage. J Neurol Neurosurg Psychiatry 1989; 52: 829-37.
- Yamamoto H, Hirashima Y, Hamada H, Hayashi N, Origasa H, Endo S. Independent predictors of recurrence of chronic subdural hematoma: results of multivariate analysis performed using a logistic regression model. J Neurosurg 2003; 98: 1217-21.
- Markwalder TM, Steinsiepe KF, Rohner M, Reichenbach W, Markwalder H. The course of chronic subdural hematomas after burr-hole craniostomy and closed-system drainage. J Neurosurg 1981; 55: 390-6.
- 13. Oishi M, Toyama M, Tamatani S, Kitazawa T, Saito M. Clinical factors of recurrent chronic subdural hematoma. Neurol Med Chir (Tokyo) 2001;41:382-6.

- 14. Merlicco G, Pierangeli E, di Padova PL. Chronic subdural hematomas in adults: prognostic factors. Analysis of 70 cases. Neurosurg Rev 1995; 18: 247-51.
- 15. Mori K, Maeda M. Surgical treatment of chronic subdural hematoma in 500 consecutive cases: clinical characteristics, surgical outcome, complications, and recurrence rate. Neurol Med Chir (Tokyo) 2001; 41: 371-81.
- Ko BS, Lee JK, Seo BR, Moon SJ, Kim JH, Kim SH. Clinical analysis of risk factors related to recurrent chronic subdural hematoma. J Korean Neurosurg Soc 2008; 43: 11-5.
- 17. Fujioka S, Matsukado Y, Kaku M, Sakurama N, Nonaka N, Miura G. CT analysis of 100 cases with chronic subdural hematoma with respect to clinical manifestation and the enlarging process of the hematoma (author's transl). Neurol Med Chir (Tokyo) 1981; 21: 1153-60.
- Nomura S, Kashiwagi S, Fujisawa H, Ito H, Nakamura K. Characterization of local hyperfibrinolysis in chronic subdural hematomas by SDS-PAGE and immunoblot. J Neurosurg 1994; 81: 910-3.

เวลาของการขยายตัวของสมองและการกลับเป็นซ้ำของเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรังหลังการผาตัด

ศรัณย ์ นันทอารี, เทวเจษฎา ภาเรื่อง, บรรพต สิทธินามสุวรรณ

ภูมิหลัง: เลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรังเป็นโรคทางประสาทศัลยศาสตร์ที่พบบอย ยังไม่เป็นที่ทราบแน่ชัดเกี่ยวกับการขยายตัวของสมอง วัตถุประสงค์: การวิจัยนี้เป็นการศึกษาไปข้างหน้าเพื่อศึกษาเวลาของการขยายตัวของสมองและการกลับเป็นซ้ำของเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรัง วัสดุและวิธีการ: ภายหลังการผ่าตัดระบายเลือดและใส่สายระบายเลือดใต้เยื่อหุ้มสมอง ผู้ป่วยแต่ละรายจะได้รับการตรวจเอ็กซเรย์คอมพิวเตอร์สมอง ที่เวลาต่าง ๆ ดังนี้ ก่อนนำสายระบายเลือดใต้เยื่อหุ้มสมองออก หลังผ่าตัด 1 สัปดาห์ 2 สัปดาห์ 4 สัปดาห์ และ 8 สัปดาห์ หรือเมื่อมี อาการผิดปกติทางระบบประสาทที่สงสัยว่าจะมีการกลับเป็นซ้ำของเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรัง

ผลการศึกษา: การศึกษานี้รวบรวมเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรัง 102 ครั้ง ในผู้ป่วย 75 ราย เอกซเรย์คอมพิวเตอร์สมองทั้งหมด 450 ครั้ง ในผู้ป่วยทั้งหมดได้รับการศึกษาพบผู้ป่วยที่มีการขยายตัวของสมองเต็มที่ระหวางสัปดาห์ที่ 1 และ 2 เพียงร้อยละ 1 ระหวางสัปดาห์ที่ 2 และ 4 ร้อยละ 5.9 ระหวางสัปดาห์ที่ 4 และ 8 ร้อยละ 80.4 และหลังสัปดาห์ที่ 8 ร้อยละ 12.7 พบอุบัติการณ์ของการกลับเป็นซ้ำของเลือดออกใต้เยื่อหุ้มสมอง แบบเรื้อรังร้อยละ 9.8 โดยในผู้ป่วยที่มีการกลับเป็นซ้ำ ร้อยละ 10 เกิดก่อนนำสายระบายเลือดใต้เยื่อหุ้มสมองออก ร้อยละ 70 เกิดระหวางสัปดาห์ที่ 1 และ 2 ร้อยละ 10 เกิดระหวางสัปดาห์ที่ 2 และ 4 และร้อยละ 10 เกิดหลังสัปดาห์ที่ 8

สรุป: การขยายตัวของสมองหลังการผ่าตัดเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรังเป็นกระบวนการที่เป็นไปอยางช้า ๆ จากการศึกษาพบการไม่เข้ากัน ระหวางเวลาของการขยายตัวของสมองและการกลับเป็นซ้ำของเลือดออกใต้เยื่อหุ้มสมองแบบเรื้อรัง การกลับเป็นซ้ำสวนใหญ่จะพบระหวางสัปดาห์ที่ 1 และ 2 ในขณะที่การขยายตัวของสมองเต็มที่จะพบภายหลังสัปดาห์สัปดาห์ที่ 4 ช่วงเวลาระหวางสัปดาห์ที่ 1 และ 2 เป็นช่วงเวลาที่สำคัญสำหรับ ดิดตามอาการหลังผ่าตัด การที่พบช่องวางใต้เยื่อหุ้มสมองหลังผ่าตัดจากการตรวจเอกซเรยคอมพิวเตอร์สมองที่เวลาแบบสุ่มหลังผ่าตัดโดยที่ผู้ป่วย ในมีอาการผิดปกติถือว่าเป็นกระบวนการที่เกิดขึ้นปกติของการขยายตัวของสมอง