Brain creatine kinase in blood after acute brain injury

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SYNOPSIS Severe cold injury of the brain increased significantly both total creatine kinase and the corresponding brain isoenzyme (CK_{BB}) activity in confluens sinuum samples. CK_{BB} could be detected also in peripheral blood a few hours after severe brain injury in eight of 12 patients. Finding of CK_{BB} in human plasma may prove a useful indicator of severe brain injury.

Acute tissue damage usually releases intracellular enzymes into the circulation. Acute brain damage is followed by increased creatine kinase (CK) activity in the serum (Acheson *et al.*, 1965; Langton *et al.*, 1967; Eisen and Sherwin, 1968; Wolintz *et al.*, 1969). Efforts to determine the source of increased CK activity have given surprising results: typical brain tissue isoenzyme, CK_{BB}, has not been found at all in the serum. Rather the increased total CK activity was composed of the isoenzymes of skeletal muscle and heart; CK_{MM} and CK_{MB} (Dubo *et al.*, 1967; Cao *et al.*, 1969). Serum enzyme diagnosis has therefore been considered useless for the diagnosis of brain damage.

New and more sensitive CK isoenzyme methods have been developed (Roe *et al.*, 1972; Somer and Konttinen, 1972) offering new possibilities for organ specific enzyme analysis (Klein *et al.*, 1973; Konttinen and Somer, 1973; Somer *et al.*, 1973). We therefore studied whether these more refined methods would reveal release of CK_{BB} into blood after acute brain injury in both experimental animals and in human cases.

METHODS

EXPERIMENTAL SERIES Healthy rabbits were used. The first blood sample was taken from an auricular vein (BA sample) and the animals were then anaes-

thetized with 30-35 mg sodium pentobarbitone (Nembutal) per kg body weight. Atropine was always used. Local anaesthesia for tracheostomy was induced with a 1-2% solution of prilocaine (Citanest). The animals breathed spontaneously throughout the experiment. Procaine penicillin (300 000 iu) was given intraperitoneally to prevent wound infection. Peripheral blood samples were taken through a catheter inserted into a jugular vein with the cephalad end of the vessel ligated. For cerebral blood samples a cannula was attached to the confluens sinuum (Trouppe *et al.*, 1966). One peripheral and one cerebral sinus sample were taken immediately after the surgical preparation was completed (AP sample). A funnel with a diameter of 15 mm was attached to the left of the sagittal suture of the rabbit skull, just behind the coronary suture. A severe cold injury was induced in 15 animals by pouring liquid nitrogen into the funnel (Kaste and Trouppe, 1972). The freezing time was one and a half minutes. Blood samples were drawn three, six, and 12 hours after the injury. Twelve control animals underwent all procedures except the brain injury.

CLINICAL SERIES Twelve patients with severe brain injury suffered within the previous eight hours, and seven patients undergoing neurosurgical operations were studied (see Table 2). The indications for the neurosurgical procedures were: middle cerebral aneurysm (two patients), deep meningioma (two patients), malignant intracerebral tumour (two patients), and one acoustic neuroma. In all instances there was some trauma to brain tissue during the operation. Serial peripheral blood samples were taken, but a rigid time schedule could not always be achieved with the brain injured patients.

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TABLE 1
PLASMA TOTAL CK AND CK_{BB} ISOENZYME ACTIVITY AFTER EXPERIMENTAL BRAIN INJURY*

	Cerebral sinus				Peripheral blood			
	Total CK (iu/l)	(No.)	CK _{BB} isoenzyme (iu/l)	(No.)	Total CK (iu/l)	(No.)	CK _{BB} isoenzyme (iu/l)	(No.)
Before anaesthesia								
Exp.					515 ± 197	15	383 ± 161	15
Control					406 ± 95	12	312 ± 99	12
After preparation								
Exp.	384 ± 84	15	236 ± 45	15	350 ± 55	15	238 ± 58	15
Control	401 ± 110	12	215 ± 29	12	399 ± 139	12	292 ± 38	12
Hours after injury								
3								
Exp.	881 ± 182	15	502 ± 122	15	495 ± 70	15	226 ± 39	15
Control	449 ± 57	12	229 ± 29	12	412 ± 48	12	206 ± 39	12
6								
Exp.	1 029 ± 223	14	662 ± 234	14	505 ± 92	12	212 ± 46	12
Control	623 ± 96	7	256 ± 68	7	530 ± 82	7	223 ± 49	7
12								
Exp.	1 248 ± 79	6	658 ± 236	6	1 173 ± 181	4	328 ± 16	4
Control	1 765 ± 1 020	5	358 ± 200	5	1 798 ± 1 514	4	281 ± 134	4

* Mean ± SE; no. = number of samples.

TABLE 2
PLASMA CREATINE KINASE ISOENZYMES IN CLINICAL BRAIN INJURY

Case	Diagnosis	Clinical data	Samples (hours after trauma)	CK _{BB} isoenzyme (iu/l)	Total CK* (iu/l)
1	Cerebral contusion	Intellectual impairment	5 6 12	— — —	15 18 26
2	Cerebral contusion	Unconscious for 10 d	3 6 12	14 3-10 —	274 494 1 030
3	Concussion	Conscious when brought in	3 17	— —	64 187
4	Cerebral contusion, intra-cerebral haematoma	Unconscious for 2 m	4	3-10	35
5	Cerebral contusion	Unconscious for 2 m, slight improvement later	5 6	3-10 3-10	230 363
6	Gunshot wound of head	Dead 22 h after injury	6	3-10	36
7	Concussion	Conscious; a few seizures after injury	1.5 8	— 3-10	53 147
8	Cerebral contusion	Unconscious for 5 w	2.5 6 12	88 15 —	445 740 1 575
9	Cerebral contusion	Dead 10 h after injury	8	—	215
10	Cerebral contusion	Intellectual impairment	8	—	140
11	Brain injury	Unconscious for 3 w. Intellectual impairment	6 9 12	12 3-10 3-10	26 59 157
12	Acute subdural haematoma	Dead 15 h after injury	1.5 12	3-10 3-10	76 66

* Normal values for total CK: 0-50 iu/l.

BIOCHEMICAL METHODS Blood samples were taken into heparinized tubes. Some haemolysis occurred in most tubes. Glutathione was added to achieve a 10 mmol/l concentration in plasma. The samples were either analysed immediately or stored at -20°C and analysed within two months.

Total CK activity was measured by means of test kits (CPK activated, Boehringer, Mannheim), with a normal range of 0–50 iu/l in human plasma. Creatine kinase isoenzymes were separated electrophoretically and determined by a fluorescence technique (Somer and Konttinen, 1972).² The method detects an isoenzyme of 3–5 iu/l. A linear quantitation is achieved within 10–250 iu/l, if a short incubation time (0.5 h) is used. For higher activities corresponding dilutions were made. The results are expressed as CK_{BB} activity (iu/l).

RESULTS

The cold injury caused a rapid rise in CK activity in cerebral sinus blood. Within three hours total CK activity rose by 397 iu/l in the injury group but only by 48 iu/l in the control group. The difference is statistically significant

² The published method contains an error: incubation fluid should contain also 40 mg glucose in 5 ml glycylglycine buffer in addition to the other reagents mentioned.

FIG. 1 Plasma creatine kinase_{BB} isoenzyme levels (mean $\pm$ SEM) in the rabbit after a severe local freeze injury to the brain.

($P < 0.01$, Student's t test). The rise is mainly due to CK_{BB} activity, which rose by 266 iu/l in the injury group and by only 14 iu/l in the control group ($P < 0.02$). The highest CK_{BB} values were seen in the six hour and 12 hour samples from the confluens sinuum. No rise in CK_{BB} activity was observed in peripheral blood samples collected at the same time (Fig. 1, Table 1).

Peak total CK activity was measured at 12 hours. The animals surviving this long showed a wide scatter in enzyme values and no clear difference was noted between experimental and control animals, either in cerebral sinus or in peripheral blood samples. CK_{MM} , the isoenzyme of skeletal and heart muscle, was usually responsible for high total CK values in the 12 hour samples.

FIG. 2 Plasma creatine kinase isoenzyme in a patient with cerebral contusion. The CK_{BB} isoenzyme is typical of brain tissue, and CK_{MB} and CK_{MM} are typical of heart and skeletal muscle. The first sample (left) was taken 2.5 hours after the injury. Total CK activity was 445 iu/l (normally less than 50 iu/l), and CK_{BB} made up 20% of the total CK activity. The second sample (middle) was taken six hours after the injury. Total CK activity was 740 iu/l, and CK_{BB} was still visible. The third sample (right) was taken 12 hours after the injury. Total CK activity was 1 575 iu/l, and CK_{BB} could no longer be demonstrated.

CLINICAL In patients with acute brain injury CK_{BB} isoenzyme was found in peripheral blood in eight of 12 patients. All eight patients had had severe brain injury, usually a cerebral contusion, causing severe disturbance of consciousness (Table 2). CK_{BB} activity was usually at its peak in the first sample and then disappeared quickly (Fig. 2). At most it was 23% of total CK activity. The peak total CK activity was found later when there was little or no CK_{BB} activity. No CK_{BB} activity was demonstrated in samples collected after the neurosurgical operations.

DISCUSSION

In the experimental rabbits the rise in total CK activity in cerebral venous blood was largely due to CK_{BB} activity, the isoenzyme which occurs mainly in the brain. Peripheral blood did not show this rise in CK_{BB} activity; perhaps the amount of CK_{BB} released from the damaged brain was too small to cause an appreciable rise in CK_{BB} in pooled peripheral blood. The rise of CK_{BB} in cerebral venous blood alone, and not in pooled peripheral blood, excludes lungs, spleen, kidneys, red muscle or thyroid as sources of this CK_{BB}, although in the rabbit these organs contain some CK_{BB} (Brody and Hatcher, 1967; Sherwin *et al.*, 1967). Total CK activity increased in quite a few samples of both cerebral and pooled peripheral blood 12 hours after injury, but this was due to a rise in CK_{MM} or CK_{MB}, the isoenzymes presumably released by surgical trauma, anaesthesia, or impaired ventilation (Dixon *et al.*, 1971; Phornphutkul *et al.*, 1974).

The clinical results tally with the experimental ones. Some CK_{BB} is released from brain to blood soon after a severe brain injury. Since normal human plasma does not show any CK_{BB} activity, small amounts can easily be detected in analyses of peripheral blood. To find CK_{BB} in peripheral blood after clinical brain injury requires a sensitive method as well as well-timed sampling. The fluorescence technique (Somer and Kontinen, 1972) is clearly more sensitive than the methods used in previous studies (Dubo *et al.*, 1967; Cao *et al.*, 1969). When present, CK_{BB} isoenzyme could usually be detected even in the earliest samples collected. No association between the presence of CK_{BB} and total CK

activity could be observed. This again shows the uselessness of total CK measurements as an indicator of brain damage.

The clinical value of plasma CK_{BB} determinations has so far not been determined. Although CK_{BB} occurs in some other human organs too (Dawson and Fine, 1967), it has not been found previously in human plasma in any other clinical conditions except in a few cases of malignant hyperpyrexia (Zsigmond and Starkweather, 1973). It seems that a severe injury is needed to cause the release of CK_{BB} into peripheral blood and so, when it does, this may possibly offer new criteria for the early assessment of the severity of brain damage.

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