Brain specific proteins in posthaemorrhagic ventricular dilatation

A Whitelaw, L Rosengren, M Blennow

Abstract

Median neurofilament and glial fibrillary acidic protein concentrations in the cerebrospinal fluid of 18 infants with posthaemorrhagic ventricular dilatation were 20–200 times higher than control values. S-100 protein in cerebrospinal fluid was four times higher than control values. Glial fibrillary acidic protein concentrations correlated with death or disability and with parenchymal lesions but not with shunt dependence.

(Arch Dis Child Fetal Neonatal Ed 2001;84:F90–F91)

Keywords: cerebrospinal fluid; glial fibrillary acidic protein; posthaemorrhagic ventricular dilatation; brain; neurodevelopment

Division of Child Health, University of Bristol, Bristol, UK
A Whitelaw

Department of Neurology, Sahlgrenska Hospital, Gothenburg, Sweden
L Rosengren

Department of Paediatrics, Huddinge Hospital, Huddinge, Sweden
M Blennow

Correspondence to:
Professor Whitelaw, Division of Child Health, University of Bristol Medical School, Southmead Hospital, Bristol BS10 5NB, UK
email: andrew.whitelaw@bristol.ac.uk

Accepted 21 January 2000

Table 1 Median (ranges) for cerebrospinal fluid concentrations of brain specific proteins

	PHVD – parenchymal lesion	PHVD + parenchymal lesion	p Value
n	7	11	
NFL (ng/l)	2563 (713–107000)	36446 (1846–168000)	0.02
GFAP (ng/l)	24849 (6808–67236)	196500 (15041–1100000)	0.002
S-100 (µg/l)	3.56 (2.19–9.73)	9.85 (2.19–36)	0.036
	PHVD normal at 12 months	PHVD disability at 12 months	p Value
n	7	11	
NFL (ng/l)	2563 (713–107000)	38092 (1846–168000)	0.0154
GFAP (ng/l)	24849 (6808–67236)	196500 (15041–1100000)	0.002
S-100 (µg/l)	3.56 (2.19–9.73)	9.85 (2.19–36)	0.036
	PHVD no shunt	PHVD shunt or dead	p Value
n	6	12	
NFL (ng/l)	22093 (1448–168000)	7622 (713–88168)	0.437
GFAP (ng/l)	58797 (10354–154000)	149250 (6808–1100000)	0.291
S-100 (µg/l)	4.78 (2.19–19)	8.76 (2.19–36)	0.30

Normal preterm control: NFL, median (range) 125 (<125–138) ng/l (n = 4); GFAP, mean (SD) 106 (15–362) ng/l (n = 17); S-100, median (range) 0.82 (0.39–1.09) µg/l (n = 4). p Values obtained using the Mann-Whitney U test.

PHVD, posthaemorrhagic ventricular dilatation; NFL, neurofilament triplet protein; GFAP, glial fibrillary acid protein.

its presence in CSF is related to astroglial damage.³ GFAP in CSF of preterm infants is increased fivefold in preterm infants with an abnormal neonatal course or abnormal neurological signs.⁴

S-100 is an acidic calcium binding protein produced in both protoplasmic and fibrillary astrocytes.⁵ These proteins are not normally synthesised outside the central nervous system and are not present in erythrocytes, leucocytes, or platelets.

The objects of this study were to determine whether the brain specific proteins GFAP, NFL, and S-100 in the CSF of preterm infants with posthaemorrhagic ventricular dilatation (PHVD) differed from control preterm infants and were related to neurodevelopmental outcome, the presence of parenchymal brain lesions on ultrasound, and the need for ventriculoperitoneal shunting.

Methods

CSF surplus to diagnostic requirements was available after clinically indicated ventricular puncture in 18 preterm infants of mean (SD) birth weight 1287 (490) g with PHVD defined as IVH followed by progressive enlargement of ventricular width reaching 4 mm over the 97th centile for each lateral ventricle.¹

Control values for GFAP were previously obtained in our laboratory from 17 preterm infants (mean (SD) birth weight 1242 (360) g).⁴ Control values for NFL and S-100 were obtained from CSF samples surplus to requirements from four preterm infants with birth weights 939–1520 g. All the control infants were lumbar punctured on clinical indications to exclude infection but were retrospectively found to have no evidence of infection or IVH and had normal outcome on follow up.

CSF samples were centrifuged, and the supernatants frozen and transported to the laboratory. NFL and GFAP concentrations were measured using sandwich enzyme linked immunoassays.^{2–6} S-100 protein was measured using a commercially available luminometric immunoassay (Sangtec Medical, Bromma, Sweden).

Results

Table 1 shows that CSF concentrations of GFAP, NFL, and S-100 from infants with PHVD were higher in those with parenchymal brain lesions and those who died or were disabled at 12 months (Ruth Griffiths developmental quotient < 70). The differences for GFAP were highly significant, but those for NFL and S-100 were of borderline significance. Infants with PHVD but no parenchymal

lesions and normal development had a median CSF concentration of GFAP that was over 200 times that in the normal control infants, with no overlap. In the case of NFL, the factor was almost 20 times that in the normal control infants. Median S-100 concentration in the PHVD group was over four times the median of the control values with no overlap. There was no significant difference in NFL, GFAP, and S-100 concentrations between the group who survived without shunting and the group who required shunt surgery or died with progressive hydrocephalus.

Discussion

The sampling technique in the infants with PHVD involved ventricular tap in all but one case. The one infant from whom samples were taken by ventricular reservoir had brain specific protein concentrations comparable with those of the other infants with PHVD. We cannot rule out the possibility that ventricular puncture introduced brain proteins into the CSF, although such a process would not account for the further increases in brain proteins in infants with parenchymal lesions and infants with later disability. We have not had the opportunity to compare simultaneous ventricular and lumbar samples so we cannot rule out the possibility that ventricular concentrations are consistently higher than lumbar concentrations.

The extremely high GFAP concentrations in the 18 infants with PHVD are comparable with those observed after acute parenchymal destruction in adult brain, such as herpes encephalitis and cerebral infarctions, but much higher than those observed in chronic neurodegenerative disorders with concurrent astrogliosis.³⁻⁶ The NFL levels in the present study

are much higher than those observed in CSF from patients with chronic neurodegenerative disorders but closer to those observed after large cerebral infarctions and herpes encephalitis.² The CSF concentrations of S-100 in the infants with PHVD were comparable with those in adults with acute brain tissue damage.³

The full extent of periventricular white matter injury in the preterm infant cannot be assessed by either repeated cranial ultrasound examinations or neurodevelopmental assessment at 12 months. Grossly elevated concentrations of brain specific proteins, especially GFAP, in the CSF provide further evidence of periventricular white matter damage in infants with PHVD.

This study was supported by the Cornell Foundation.

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