Blood spot glucocorticoid concentrations in ill preterm infants

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SUMMARY The adrenocortical response to stress was studied longitudinally in 10 ill preterm infants using measurements of cortisol and 17OH-progesterone concentrations in filter paper blood spots. Mean cortisol and 17OH-progesterone concentrations reached a peak of 2200 nmol/l and 65 nmol/l, respectively, between the third and fifth days of life. These concentrations far exceeded those observed in older children and adults subjected to stress as a result of surgery. Further pulses of endogenous cortisol production of 4000 nmol/l or more occurred in association with clinical complications such as intraventricular haemorrhage. These results indicate that infants undergoing intensive care are unduly stressed. Consideration should be given to providing enough sedation and appropriate analgesia for ill preterm infants during painful procedures such as insertion of venous cannulae and arterial puncture.

Increased plasma concentrations of 17OH-progesterone (17P) and cortisol in ill preterm infants have previously been reported.^{1 2} These observations, which indicated a presumed stress response, were based only on single random measurements of plasma steroid concentrations. Little is known about the magnitude and duration of the adrenocortical response to stress in the newborn. A study of ill infants during the first three days after birth by Gutai *et al* showed that there was no significant increase in plasma cortisol concentrations. Most of the infants studied were not unduly stressed when assessed by clinical criteria.³ In a study of ill preterm infants with hyaline membrane disease,⁴ however, the peak plasma glucocorticoid response achieved was related to the severity of respiratory distress.

Studies, in which serial measurements of plasma steroid concentrations in ill preterm infants have been made, have been limited by the difficulty of collecting venous samples from small infants. A steroid assay, which uses whole blood spotted on to filter paper, has recently been developed however,⁵⁻⁷ and this technique, which requires only a small volume of blood, has been used to obtain longitudinal data on 17P and cortisol concentrations as an index of the adrenocortical response to stress in a group of ill preterm infants.

Patients and methods

Ten preterm patients were studied longitudinally for the first 13 days of life. All were very ill and required assisted ventilation for severe hyaline membrane disease. Table 1 presents their clinical details. Gestational age was assessed by the method of Ballard *et al*.⁸ Arterial catheters (umbilical or radial) were inserted shortly after birth for measurements of blood gas and invasive blood pressure monitoring. When an arterial sample was collected additional drops were spotted on to a Guthrie card sufficient to fill completely four 10 mm circles. The date and time of each sample collection was recorded for each infant. The filter paper card was left to dry at room temperature before being sent to the laboratory for analysis. In five of the infants blood spots were obtained at about six hourly intervals for three to five days.

Blood spot 17P concentrations were determined by a specific radioimmunoassay.⁷ A similar method was used to determine cortisol concentrations in blood spots, but solvent extraction was not required. Briefly, 50 µl of ethanol followed by 200 µl of phosphate buffer solution were added to each 6 mm disc punched out of a blood spot. ¹²⁵I-labelled cortisol and an anticortisol serum that had been coupled to magnetisable cellulose were added to

Table 1 Clinical details of 10 infants studied

Case No	Sex	Birth weight (g)	Gestation (weeks)	Clinical problems
1	F	1860	32	HMD, IVH; died at 14 days
2	F	805	28	HMD, PDA; died at 6 weeks
3	M	1800	32	HMD, IVH, NEC
4	M	975	25	HMD, IVH, PDA
5	M	1850	30	HMD, IVH, ARF
6	F	1630	32	HMD, PDA, ARF, pneumothorax
7	M	780	25	HMD, PDA, NEC; died at 22 days
8	M	1140	31	HMD, PDA, DIC, ARF
9	M	850	25	HMD, IVH, pneumothorax; died at 12 weeks
10	M	1485	31	HMD, IVH, ARF; died at 15 days
Mean		1313	28.8	
Range		805-1860	24-32	

HMD=hyaline membrane disease; IVH=intraventricular haemorrhage; PDA=patent ductus arteriosus; NEC=necrotising enterocolitis; ARF=acute renal failure; DIC=disseminated intravascular coagulation.

100 µl of the ethanol-buffer solution and the mixture incubated for one hour. Antibody bound and free steroids were separated using a magnetic tray and the radioactivity was counted in a multiwell gamma counter. Unknown sample concentrations were interpolated from the dose-response curve. Blood spot cortisol and 17P values were converted to the equivalent concentration in plasma based on the assumption that each disc contained 4.5 µl plasma.⁷ For both assays the intra- and interassay coefficients of variation did not exceed 7.0% and 11.5%, respectively. Table 2 summarises the characteristics of the blood spot assays. Data were calculated as mean (SEM) and compared using Student's unpaired *t* test. In those infants from whom more than one sample was collected each day the mean steroid concentration was used to determine mean values for the total group of 10 infants studied.

Results

Fig 1 shows the mean (SEM) blood spot cortisol

Table 2 Relevant characteristics of blood spot radioimmunoassays

Characteristic	Steroid	
	17OH-progesterone	Cortisol
Sensitivity	1.0 nmol/l	28 nmol/l
Range	1.0-550 nmol/l	28-2450 nmol/l
Cross reactivity (%)		
	17OH-pregnenolone	11-deoxycortisol
	7.4	28
	Progesterone	Cortisone
	5.9	23
	21-deoxycortisol	Corticosterone
	1.5	20
	Pregnenolone	Prednisolone
	0.3	4.7

Cross reactivity=amount of steroid required to displace 50% binding of labelled 17P/cortisol at zero binding

concentrations determined serially in 10 infants from birth to 13 days of age. The mean cortisol concentration increased more than twofold during the first few days to reach a maximum of about 2000 nmol/l between days 3 to 5. Blood spot cortisol concentrations decreased during the ensuing four days, and thereafter stayed constant over the remainder of the study period. Fig 2 shows the

Fig 1 Mean (SEM) blood spot cortisol concentration in ill preterm infants.

Fig 2 Mean (SEM) blood spot 17OH-progesterone concentration in ill preterm infants.

Fig 3 Serial blood spot cortisol and 170H-progesterone concentrations in an ill preterm infant: stressful clinical events (arrows).

pattern of blood spot 17P concentrations measured over the same time period. Peak values were seen on day 5 with a mean 17P concentration of 65 nmol/l. The increased blood spot 17P concentrations in preterm ill infants compared with those measured in healthy full term infants (range 3–18 nmol/l) were still evident on day 13.

Fig 3 shows the individual blood spot 17P and cortisol concentrations in one infant who died. The peak cortisol concentration exceeded 4000 nmol/l; this coincided with the development of a large intraventricular haemorrhage shown on cranial ultrasonography. Extension of the blood into the brain parenchyma was accompanied by further pulses in cortisol secretion. The infant subsequently developed hepatic failure and died. Just before death the concentrations of cortisol fell. A similar pattern of secretion was seen in plasma 17P concentrations. Fig 4 shows the individual profiles of cortisol concentrations determined in serial blood spots collected from five infants. There was a large range of values, but most infants had cortisol concentrations consistently above 1000 nmol/l.

Discussion

Mixed cord blood cortisol concentrations in healthy term infants are relatively high (800 nmol/l) but fall rapidly within 24 hours of birth to about 270 nmol/l.⁹ The cortisol gradient may be the result of placental transfer of maternal plasma free cortisol, or it may be due to increased fetal adrenal activity because of the stress of delivery.¹⁰ There is no obvious difference in cortisol concentrations between healthy term and preterm infants during the first 10 days after birth.¹¹

The pattern of cortisol secretion in ill preterm infants seems to be quite different. In this study cord blood cortisol concentrations were not measured. Although individual cortisol concentrations varied widely during the first six days after birth, blood spot cortisol concentrations progressively increased reaching a peak on day 3. They remained constant for two days before decreasing gradually over the remaining eight days of the study. Even the smallest preterm infant seems to be able to secrete large amounts of adrenocortical hormones for a

Fig 4 Blood spot cortisol profiles in five ill preterm infants.

protracted period in response to stress. A similar response was observed in concentrations of 17P in response to stress. Interestingly, peak cortisol and 17P concentrations were delayed until days 3 and 5, respectively, even though the infants were generally most ill during the first 72 hours after birth.

Studies in newborn rats have shown that there is a clearly defined period when rats do not respond to stress.¹² This is an interval immediately after birth during which the adrenocortical neonatal response to a variety of stresses is absent.¹³ Corbier and Roffi¹⁴ showed that in rats the period of absolute unresponsiveness to stress was followed by a period of relative unresponsiveness. The authors proposed that increased concentrations of corticosteroid, the principal glucocorticoid in rats, accumulate in the brain and inhibit the production of corticotrophin releasing factor. Alternatively, it was suggested that the absence of response was the result of a transient defect in adrenocorticosteroid synthesis. This is unlikely in view of the findings of a previous study which showed that ill preterm infants attain adequate concentrations of plasma cortisol if given adrenocorticotrophic hormone (ACTH).² If a similar period of absolute or relative unresponsiveness to stress also occurs in human neonates, this

may be important in the early management of the very ill preterm infant.

The results of this study illustrate that small ill preterm infants experience considerable stress while undergoing intensive care. Furthermore, an added clinical complication such as the extension of an intraventricular haemorrhage (fig 3) results in a further pulse of endogenous cortisol secretion. The pronounced changes in cortisol concentrations seen over 24 hours (fig 4) indicate that frequent sampling is required to define the period when a surge in adrenocortical activity occurs. Blood spot sampling is ideal for this purpose.

Comparable studies in adults undergoing major and minor operations indicate that mean postoperative plasma cortisol concentrations reach 1080 nmol/l and 540 nmol/l, respectively,¹⁵ values considerably lower than those achieved in ill preterm infants. The neonatal adrenal gland has a remarkable capacity for cortisol production; and Forest reported concentrations as high as 11 556 nmol/l in infants given ACTH 500 µg/m² twice daily for three days.¹⁶ Newborn puppies subjected to haemorrhagic shock achieved cortisol concentrations up to 2000 nmol/l.¹⁷ It was notable that the highest cortisol concentrations occurred in those puppies that died first. In a study of term infants undergoing cardiac operations with cardiopulmonary bypass, the plasma cortisol concentration increased from 356 nmol/l before operation to 781 nmol/l postoperatively.¹⁸ When the stress response was reduced by the administration of analgesia, the infants were more stable postoperatively.

While the stress response may be beneficial in the short term there are some drawbacks. In the absence of hypovolaemia stress induces unwanted hypertension associated with the breakdown of protein and a negative nitrogen balance.¹⁹ Transcutaneous oxygen measurements have shown that many core orientated stimuli given to preterm infants produce a fall in pO₂.²⁰ The results of the present study indicate that infants being given intensive medical treatment are even more stressed than those who have had surgery. Painful procedures liable to cause stress, such as insertion of venous cannulae and arterial stabs, perhaps should be accompanied by better sedation and the provision of appropriate analgesia for ill preterm infants.

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