Factors predisposing to a complicated initial febrile convulsion

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Wallace, S. J. (1975). *Archives of Disease in Childhood*, 50, 943. **Factors predisposing to a complicated initial febrile convulsion.** 131 consecutive admissions to hospital for a first febrile convulsion were studied to find which factors predisposed to a complicated fit—defined as one lasting more than 30 minutes, unilateral, or repeated within the same illness. A significant excess of complicated attacks occurred where the age of onset was less than 16 months, where both family history of convulsive disorder and perinatal abnormality were present, and, in females only, where it was suspected that neurological disorder preceded the first fit.

A febrile convulsion is defined here as any convulsion occurring in association with any febrile illness. It is further defined as complicated if it continues for more than half an hour, has unilateral features, or is repeated within the same illness. Such convulsions increase the likelihood of further febrile seizures, persisting neurological disorder, suboptimal mental development, and generalized or temporal lobe epilepsy (Aicardi and Chevrie, 1970; Bamberger and Matthes, 1959; Degen and Goller, 1967; Dooze *et al.*, 1966; Falconer, Serafetinides, and Corsellis, 1964; Gastaut *et al.*, 1960; Horstmann and Schinnerling, 1963; Hrbek, 1957; Lennox-Buchtal, 1973; Ounsted, Lindsay, and Norman, 1966; Schmidt, 1958; Wallace, 1974). It seems relevant to search for the causes of a complicated convulsion.

The present study compares 81 children whose initial convulsive episode was complicated, with 50 in whom there was a single generalized initial fit lasting less than half an hour. All 131 were seen at the Royal Hospital for Sick Children, Edinburgh.

Patients and methods

The 131 children were sequential admissions to hospital for febrile convulsions in whom family history, perinatal details, and developmental history were complete. All were aged more than 3 months and less than 7 years at the time of their first fit, and were febrile or had evidence of active infection. The temperature recorded as near the time of the fit as possible was noted.

A careful history of the convulsive episode was taken. Particular note was taken of the presence of unilateral clonic movements, even if the fit later became generalized, the duration of the tonic/clonic stage, and the number of separate tonic/clonic attacks. Only if a single completely generalized convulsion lasting less than half an hour had been present is it defined as simple. All other convulsions are classed as complicated.

The patients were divided into two groups. (a) Complicated fit; 50 males, 31 females, total of 81. (b) Simple fit, 32 males, 18 females, total of 50.

Histories of convulsive disorder in parents and sibs were sought. Note was made of which children had those perinatal problems found significantly more often in the same group of patients than in their sibs (Wallace, 1972). A detailed developmental history was taken and neurological examination performed. Neurological disorder before the first fit was suspected in those children who had favoured the limbs of one side early in either reaching or walking, showed unchanging abnormal neurological signs throughout their illness and follow-up, had dwarfing of the limbs on one side in association with a hemiplegia, or whose motor milestones were more than 2 SDs above the mean for age given by Neligan and Prudham (1969). No child was more than mildly affected, nor had any received previous medical attention for neurological symptoms or signs. Viral studies were available in 53 patients. Serum sodium levels were obtained within 6 hours of the fit in 120 children. Sibs who had not themselves had fits were examined neurologically (82 patients) and by electroencephalograph (EEG) (70 patients).

The children were categorized according to sex, family history of convulsive disorder, the presence of at least one significant perinatal problem and prior neurological disorder. In subsequent analysis, correction was made for these factors. The influences of the

TABLE I
Distribution of children with each type of fit

Sex	Type of fit					
	Simple	Complicated				
		Generalized		With focal features		
		Duration $\geq \frac{1}{2}$ h	Repeated within same illness	Duration $< \frac{1}{2}$ h	Duration $\geq \frac{1}{2}$ h	Repeated within same illness
Male	32	11	15	5	9	10
Female	18	5	12	6	5	3

following on the type of initial convulsive episode were examined: age at onset, viral infection, neurological abnormality in at least one sib, and EEG abnormality in at least one sib. The influence of previous feverish illness, temperature of at least 38.9°C , and serum sodium <130 mEq/l within 6 hours of the fit were examined after further correction for age at onset.

After summation of observed values for each factor in each category of patient, expected values were calculated and summed and the differences submitted to the χ^2 test of significance.

Results

Eighty-one of the 131 patients (62%) had an initial attack which was multiple, had unilateral features, or lasted more than 30 minutes. The number of children having each type of fit is shown in Table I. Where differences between children with complicated and simple fits occurred, the trend was the same for each type of complicated fit. The findings for all complicated fits are therefore presented together (Table II).

Neither males nor females showed a greater propensity to a complicated initial convulsion ($\chi^2=0.07$, NS). A nonsignificant excess of males with a positive family history of convulsions had a simple initial fit ($\chi^2=2.09$, NS). When considered singly, a positive history for convulsions in parents and/or sibs or a perinatal abnormality were not enough in either sex significantly to increase the risk, but if both were present a significant excess of children had a complicated initial fit ($\chi^2=4.72$, $P<0.05$). Prior neurological disorder did not appear to influence the severity of the males' initial convulsions ($\chi^2=0.297$, NS), but was significantly associated with a complicated initial fit in females ($\chi^2=4.52$, $P<0.05$). Histograms showing the type of fit against the age of onset for males and females separately showed in both sexes, but particularly in the males, a significant tendency for children younger than 16 months to have complicated fits ($\chi^2=6.34$, $P<0.02$) (Fig. 1 & 2). A

FIG. 1.—Type of fit related to age at onset in 82 males. FH, family history.

surprisingly high proportion (65%) of the children had had previous feverish illnesses including measles, without having had a convulsion, and even after correction for age of onset there was no excess of children with a complicated fit among those who were thought to be suffering from their first fever ($\chi^2=0.2$, NS). Recorded temperatures of at least 38.9°C were not significantly related to complicated fits ($\chi^2=0.52$, NS). An excess of proven viral infections was found in the children with complicated fits, but was not significant ($\chi^2=3.16$, NS). Serum sodium levels of <130 mEq/l were not particularly associated with

TABLE II
Comparison between children with complicated and simple initial fits

Factor investigated	Initial fit		χ^2	P
	Complicated	Simple		
Sex (n = 131)				
Male	50	32	0.07	NS
Female	31	18		
Convulsions in sibs and/or parents (n = 131)				
Present	45	32	0.84	NS
Absent	36	18		
Convulsions in sibs and/or parents of male patients (n = 82)				
Present	28	22	2.09	NS
Absent	23	9		
Perinatal problem (n = 131)				
Present	54	26	3.28	NS
Absent	27	24		
If family history positive, perinatal problem (n = 77)				
Present	32	15	4.72	<0.05
Absent	13	17		
Prior neurological disorder: males (n = 82)				
Present	14	11	0.3	NS
Absent	36	21		
Prior neurological disorder: females (n = 49)				
Present	11	2	4.52	<0.05
Absent	20	16		
Age at onset (n = 131)				
<16 m	36	13	6.34	<0.02
16 m or more	45	37		
Previous feverish illness (n = 131)				
Present	52	33	0.2	NS
Absent	29	17		
Temperature $\geq 38.9^\circ\text{C}$ (n = 131)				
Present	34	24	0.52	NS
Absent	47	26		
Viral infection (n = 53)				
Present	19	9	3.16	NS
Absent	11	14		
Serum sodium within 6 h of fit (n = 120)				
<130 mEq/l	17	12	0.23	NS
130 mEq/l or more	56	35		
Nonconvulsing sib with abnormal CNS (n = 82)				
Present	30	16	0.03	NS
Absent	23	13		
Nonconvulsing sib with abnormal EEG (n = 70)				
Present	15	10	0.31	NS
Absent	28	17		

NS, not significant.

either type of fit ($\chi^2=0.23$, NS). Neurological abnormality in at least one nonconvulsing sib did not appear to influence whether the index case had a complicated or simple initial fit ($\chi^2=0.03$, NS). Nor did EEG abnormality in at least one nonconvulsing sib ($\chi^2=0.31$, NS).

Discussion

The study investigates the possibility that in those children where the initial convulsive episode is prolonged, multiple, or has unilateral features, there are factors predisposing to a complicated fit.

Compared with other reports, 62% is an unusually high proportion for complicated attacks. All patients in this series were admitted to hospital, but no other selection was made. Previous incidences for multiple, unilateral, or prolonged febrile convulsions have varied from 25 to 37% (Dooze *et al.*, 1966; Herlitz, 1941; Lennox, 1949; Lennox-Buchthal, 1973). The definition of severity is variable in these studies, making direct comparison difficult. It is also likely that full details of lateralizing features may not have been available.

The finding that neither the males nor the fe-

FIG. 2.—Type of fit related to age at onset in 49 females. FH, Family history.

males in this study appeared to be more prone to a complicated initial convulsion is at variance with some studies where girls seem more susceptible to prolonged or unilateral fits (Herlitz, 1941; Aicardi and Chevrie, 1970; Taylor and Ounsted, 1971). However, these findings are not strictly comparable with those of the present group since the authors did not restrict themselves to the initial convulsive episode and the larger the number of fits suffered, the more likely is a severe one to have occurred (Lennox-Buchthal, 1973). The Gentofte series also failed to show a female preponderance for prolonged or unilateral convulsions (Lennox-Buchthal, 1973).

The influence of family history on the severity of individual convulsions does not appear to have been examined previously, but it has been reported that males with a negative family history are a group least likely to suffer severe sequelae (Taylor and Ounsted, 1971). The present study showed that males with a positive family history were less likely to have a complicated fit. This suggests that inheritance of the febrile convulsion trait might protect males against a fit carrying a serious prognosis. Further evidence for the absence of simple positive correlation of hereditary factors with complicated fits is found in the analysis of the sibs' neurological and EEG data.

Although the children with a significant perinatal abnormality had a higher risk of a complicated initial fit, this risk was significantly increased only

where the family history was positive. It is probable that in these cases summation of two independent factors is responsible for perpetuation and/or lateralization of the convulsive process.

Prior neurological damage was associated with a complicated initial fit in the females only. Few studies of febrile convulsions report findings on neurological examination, and information which might suggest that neurological disorder existed before the convulsion is almost entirely lacking. However, Lennox-Buchthal (1973) reports personal communications which show a higher incidence of convulsions with fever in cerebral-palsied children than in the general population, but no relation to the type of fit or sex of the affected children is given. Bamberger and Matthes (1959) found that unilateral convulsions frequently arise as a result of brain damage regardless of its aetiology. It is not surprising that prior neurological disorder predisposes to prolongation or lateralization of a convulsion, but the explanation for this appearing in girls and not boys is obscure. The theory of differential cerebral maturation of Taylor and Ounsted (1971) might suggest that the effect of age and the presence of neurological disorder could combine to make those girls with abnormal symptoms and signs liable to convulse at an age at which they are prone to a complicated seizure. However, the present study did not confirm that girls more often had complicated seizures at a younger age than boys.

Although most authors agree with the present conclusion that younger children are more severely affected, the critical age below which convulsions are more likely to be complicated has varied. Lennox (1949) and Dose *et al.* (1966) suggest up to 12 months, the present study up to 15 months, and Aicardi and Chevrie (1970) up to 18 months. The finding that younger boys were more likely to have a complicated initial seizure contrasts with previous series where younger girls have seemed more vulnerable (Herlitz, 1941; Taylor and Ounsted, 1971).

Attention has been directed to the height and rapid rise in fever in the causation of febrile convulsions but correlation between these features and the type of fit has been infrequent. The present study, which fails to show a relation between a high fever and a complicated fit, agrees with Herlitz (1941). The finding that complicated fits occurred more often in children in whom positive evidence of viral infection was found agrees with the work of Lennox-Buchthal (1973). Evidence from studies in children who have positive viral findings suggests that their complicated fits are often symptoms of

encephalitis, rather than mere responses to fever (Wallace and Zealley, 1970).

Although Nyhan and Cooke (1956) have reported very low serum sodium levels in association with persisting convulsive activity, and Millichap, Madsen and Aledort (1960), and Melekian, Laplane, and Debray (1962) have found levels of less than 130 mEq/l in some children with febrile convulsions, Lennox-Buchthal (1973) finds insufficient evidence that hyponatraemia causes febrile convulsions. In addition, 21 of a control series of 33 children without convulsions whose temperatures were at least 37.8 °C were found to have serum sodium levels of less than 130 mEq/l (unpublished data). It is therefore not surprising that the type of fit appeared to be uninfluenced by the serum sodium level.

In conclusion, evidence is presented that those children whose first febrile convulsion occurs before the age of 16 months, those who have both a positive family history of convulsive disorder and a perinatal insult, and those females thought to have had neurological disorder before the fit, carry a significant risk that their first febrile convulsion will last longer than 30 minutes, have unilateral features, or be repeated within the same illness. It may therefore be postulated that some of the serious sequelae of febrile convulsions are determined before or at the time the first fit starts.

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