Prognostic significance of serum iron levels in cases of Japanese encephalitis

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Summary: Forty-one children with acute laboratory confirmed Japanese encephalitis were studied. Serum iron concentrations were consistently low following Japanese encephalitis virus infection, with the levels being of prognostic significance.

Introduction

Japanese encephalitis virus (JEV), an arthropod borne flavivirus is one of the major causes of acute encephalitis and is responsible for thousands of cases each year, mainly in the Far East. In Uttar Pradesh, India. Japanese encephalitis is probably endemic with high mortality. Little is known about the pathogenesis of infection in man.

Alteration in iron metabolism is an accompaniment to a variety of infections and inflammatory states. Pekarek et al. have demonstrated a significant fall in serum iron concentration before the onset of illness in volunteers exposed to bacterial infection and in asymptomatic subjects after exposure to live attenuated Venezuelan equine encephalomyelitis virus vaccine. A reduction in plasma iron has been shown to occur during a number of bacterial infections or after injection of bacterial lipopolysaccharide. This response to infections has been regarded as a host-mediated attempt to control the infection by limiting the iron availability to organisms.

Recent studies in mice revealed a low serum iron concentration following JEV infection caused by the action of macrophage derived factor, which blocks the release of iron from splenic reticuloendothelial cells. To date no information is available on changes in iron levels during JEV infection in man.

Patients and methods

Ninety-five children with acute encephalitic illness (acute, nontransient alteration of consciousness with or without fever or other neurological symptoms) were admitted to G.M. and Associated Hospital, Lucknow, India between July to December 1989. Their age was between 3 to 12 years. The duration of illness on admission was less than 5 days. Forty-one children had one or more indicators of JEV infection. A diagnosis of JEV infection was confirmed by virus isolation from cerebrospinal fluid (CSF), by demonstration of JEV antigen in CSF cells (JEV monoclonal antibody was provided by Dr E.A. Gould, Oxford, UK), by demonstration of anti-JEV IgM in CSF or by rising antibody titre against JEV in paired sera as described previously. The control group comprised of 21 normal healthy children of matched age and sex. All the children were bled in the morning at approximately the same time. Blood was collected in polyethylene syringes and test tubes to prevent iron contamination. Blood specimens could only be collected once in fatal cases, but in others at 1–4 days, 5–8 days and 9–12 days of illness, on the basis of first day of appearance of fever.

Serum iron and total iron binding capacity were detected colorimetrically according to the modified method of Beale et al. with bathophenanthroline sulphate as the colour reagent as described earlier. The total protein and albumin in serum were measured colorimetrically.

Results

Forty-one children aged 3 to 12 years having confirmed Japanese encephalitis were included in the study. Thirteen children were severely ill and rapidly progressed to coma and death within 48 hours of admission. Twenty-eight children survived.

The serum iron levels in JEV patients and control groups are presented in Table I. The mean serum iron level of 34 ± 3 μg/dl in fatal Japanese encephalitis patients was significantly less (P < 0.001) than the mean (82 ± 9 μg/dl) observed in normal
healthy controls. There was no significant difference in serum iron values in the initial stage of illness in patients with non-fatal encephalitis when compared with control (Table I). In these patients, the later serum iron levels were 40 ± 3 μg/dl (P<0.001) between 5 to 8 days and 48 ± 4 μg/dl between 9 to 12 days of illness. The total iron binding capacities in JE patients was similar to that of control children.

There was a transient decrease in the mean haemoglobin values in the non-fatal encephalitis patients between 5 to 8 days (7.0 ± 1.5 g/dl) of illness when compared with control (11.5 ± 2.5 g/dl). No significant difference was noted in haemoglobin values in cases of fatal encephalitis and the controls (P>0.05). The total serum protein and albumin did not show any deviation from normal values (Table II).

**Discussion**

The initial response of serum iron concentration after fatal JEV infection was a decrease. Significant decrease in serum iron concentration was observed within 48 h in severely ill fatal patients. The investigations revealed that the serum iron levels did not show any marked alterations in the initial period in non-fatal encephalitis patients. Thus the reduction in serum iron level was directly related to the severity of illness. Out of the 13 fatal JE patients, in 5 the JE virus was isolated from CSF and in 8 patients JE virus-specific antigen in large numbers of CSF cells was detected by indirect immunofluorescence while no virus was isolated from CSF in non-fatal cases and the immunofluorescence positive cells were few in CSF (data not included).

Following JEV infection hypoferraemia with transient anaemia had been demonstrated in mice. The degree of decrease was associated with the severity of infection. The hypoferraemia is achieved by a block in the release of iron from reticulo-endothelial cells in spleen by a macrophage derived factor.

The serum iron level decreases during inflammation which causes reduction in synthesis of hepatic export proteins, albumin and transferrin, resulting in hypoferraemia. Simultaneously, the release occurs of certain acute phase proteins.
formed by activated monocytes of liver with their increased concentration in serum. A fall in serum iron due to alterations in plasma proteins is a general marker of inflammation. In the present study there was no apparent alteration in total protein or albumin, which suggests that there was no change in protein synthesis in liver during JEV infection.

Once the specific diagnosis of JE in patients has been made on the day of admission either by indirect immunofluorescence examination of cells in CSF or by examination of IgM antibodies in CSF, it may now be possible to predict the prognosis of JE in confirmed cases by measuring the serum iron on the same day.

References

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