Hypopituitarism and hepatitis E genotype 3 infection - first case report and review on immunocompetence in hypopituitarism

Abstract

We studied the course of acute hepatitis E virus infection [HEV] in a German patient under stable hormonal replacement therapy for hypopituitarism. The 75-year-old male patient had acute hepatitis E with positive immunoglobulins [IgG and IgM] as well as HEV RNA Genotype 3c detection. Serum transaminases rose in 3/17 and spontaneously dropped to pre-infection levels after dose escalation of the hydrocortisone. Although there is data in the medical literature, that panhypopituitaric patients even under stable hormonal replacement regimens have immune deficiency, their immune competence seems to be sufficient to overcome acute hepatitis E infection.

Keywords: hepatitis e, hypopituitarism, panhypopituitarism, hydrocortisone, cortisone replacement therapy, hepatitis e genotype 3

Introduction

In the last decades, hepatitis E virus infection [HEV] had often been regarded as a disease limited to the tropics. Infections with the tropical HEV genotypes 1 and 2 are mainly transmitted fecal-orally by contaminated drinking water. In the Middle East and North African regions, the seroprevalence of HEV in the general population ranged from 2.0 to 37.5% and was higher in males than in females. HEV genotype 3 is also endemic in Europe and an under diagnosed and emerging public health issue. In industrialized countries, domestic pigs are the main carriers of hepatitis E and wild boars can also carry the virus. That said, the consumption of raw or undercooked pork meat and liver is the most common cause of hepatitis E infection in these regions. In the vast majority of patients, the infection is either clinically silent or has a benign self-limiting course. However, immunosuppressed individuals, such as transplant recipients or patients with human immunodeficiency virus [HIV]-infection, are at risk for developing a fulminating course of chronic hepatitis E, that may eventually lead to liver fibrosis and cirrhosis. The rise in hepatitis E infection in industrialized countries will lead to an increased number of infections in patients with other comorbidities, some of them with a negative impact on immune function and thus, at risk for a more complicated course of HEV infection. Among these could be hypopituitarism. The immune function and immunocompetence in hypopituitaric patients is still a matter of debate. There is a wide body of literature on immunological perturbations in animal models with deficiency or excess of pituitary hormones. However, in humans, the situation is not that clear. E.g. Mukherjee et al. reported significant humoral immune deficiency in twenty-one panhypopituitary patients although they were under stable pituitary hormone replacement. The reduced immune function accounted most strikingly in patients with low insulin-like growth factor 1 levels, which was not reproduced in more recent studies. The case of a patient with panhypopituitarism under pituitary hormone replacement with acute hepatitis E is reported. Such cases to our knowledge have not been reported before.

Case report

A 75-year-old male patient had maculopapulous exanthema in the gluteal region, as well as the extremities on 17.3.2017. The exanthema was non-itching and self-limiting the next day. General exhaustion, nausea and low appetite followed since 19.3.2017. Upon admission to our clinic he was fully conscious and not icteric. He had not stayed abroad since September 2016. Important in his medical history was well-controlled hypertension, treated with Valsartan 160mg/die and Hydrochlorothiazide 12.5mg/die. Furthermore, he underwent transsphenoidal pituitary surgery due to an endocrine inactive pituitary adenoma in 10/2003. Due to recurrence, stereotactic radiation therapy with 30x1.8 Gray was performed between 6-7/2004, leaving the patient panhypopituitaric. The present replacement therapy was levothyroxine 75µg daily, hydrocortisone 15-10-0mg daily, levothyroxine 75µg daily, hydrocortisone 15-10-0mg and testosteroneneanathate in a transdermal gel formulation with 25mg daily. Non-alcoholic fatty liver disease [NAFLD] with a slightly elevated gamma glutamyltransferase [Gamma GT] in a range between 2-4 U/l [normal value <1U/l] and steatosis hepatis in ultrasound scan was a suspected diagnosis since the nineties, liver biopsy had not been done. Due to dyslipidemia, he was also treated with Simvastatin 40mg/die since years. Although an unlikely culprit, this therapy was halted after admission. Physical examination showed an overweight (BMI 30kg/m²), non-icteric patient. He was normotensive and had a rhythmic heart action. He was normally hydrated and had no signs of exsiccosis. Furthermore, there were no pathological findings at the lung. The liver was enlarged, not painful. The neurological examination was normal.

Laboratory analysis of liver function 3/17: pathological were the levels of gamma GT, alanine aminotransferase [ALAT], aspartate aminotransferase [ASAT] and bilirubin. The course of the enzymes are shown in Figure 1. The serum was negative for antinuclear antibodies, and c -and p-anti-neutrophilic cytoplasmatic antibodies
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Results in the patient group

<table>
<thead>
<tr>
<th>Patients/Controls</th>
<th>Measurements</th>
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<tbody>
<tr>
<td>13 children with GH deficiency</td>
<td>Analysis of lymphocyte subset patterns before and during [6 and 12 months] therapy with biosynthetic GH</td>
<td>Absolute number of T-lymphocytes, T-cell subsets, natural killer cell activity and response of lymphocytes to polyclonal antigens all in the normal range before and during therapy.</td>
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</table>

Discussion

Several cells of the immune system have receptors for pituitary hormones suggesting interactions between the immune system and the pituitary. For example, receptors for IGF-1 and growth hormone are expressed in lymphocytes as well as receptors for prolactin. PRL has been associated with the modulation of a variety of actions in the immune response and inflammatory processes in several physiologic and pathologic conditions. Best characterized and well known to every physician is the effect of endogenous cortisone on immune function. However, studies in humans with isolated defects in pituitary hormone secretion or with panhypopituitarism are rare, have conflicting results and do not necessarily link an abnormal findings to clinical pathology. Furthermore, many studies have small patient numbers and no control group and focus on the impact of GH growth Hormone excess or deficiency. Table one gives an overview of the recent literature and the findings on the topic (Table 1).

Growth hormone is not replaced in our patient due to well-being, the age and the costs of this treatment. Prolactin is not replaced in hypopituitaric patients. Although data demonstrating significantly higher infection rates in hypopituitaric patients are lacking in the medical literature, an association with their premature mortality is sometimes discussed in this context and not only as a reason of cardiovascular disease. We were also not able to identify medical literature reporting more complicated courses of hepatitis types A, B or C in hypopituitaric patients. Data about hepatitis E in this context do not exist. As for other disorders, there is a trend in the studies demonstrating, that chronic or fulminant courses of HEV infection rather occur in severely immunosuppressed patients, such as transplant recipients or HIV infected patients. The courses in other disorders that have reduced immune response themselves or are treated with immunosuppressants seem to be more benign. E.g. Suzuki et al. describe the case of a patient with ulcerative colitis treated with prednisolone and infliximab. This patient had a benign self-limiting course of HEV infection. In a French study, 23 cases of HEV infection in patients with rheumatoid arthritis, axial spondyloarthritis, psoriatic arthritis, and other types of arthritides under immunosuppressive therapy regimens were studied. The treatment included discontinuation of immunosuppressants in 20 patients and ribavirin treatment in 5 patients. No chronification of the infection occurred. Although the literature gives no clear picture on the immune status of hypopituitaric patients, the replacement therapy with hydrocortisone is an important issue. It is well known, that patients with primary and secondary adrenal insufficiency require an adaptation of the hydrocortisone dose in case of infection. Since the patient was in good clinical condition, we decided to double the dose, when the transaminases rose again, which is a moderate dose adaptation. It remains a matter of speculation, to which degree the dose-escalation of hydrocortisone during the infection helped to overcome the hepatitis and what would have been the course without dose adaptation. Apart from a potential negative effect of panhypopituitarism on immune function, there is also data suggesting a worse course of Hepatitis E in underlying chronic liver disease (NASH), as can be assumed in diabetic patients. We could not identify literature on possible differences in the course of acute Hepatitis E in NASH only patients vs. patients with a healthy liver. Another factor with negative impact on the immune response is age, our patient was 75 years old.
Table continued...

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<tr>
<td>Lebl J et al.¹³</td>
<td>10 young GH deficient adults</td>
<td>Humoral [IgG, IgA, IgM, C3, C4, immune complexes] and lymphocyte subsets before GH therapy and 6 weeks, 3 and 6 months during therapy</td>
<td>„Trend“ to higher CD 4 and CD 8 lymphocytes with maximum change at 6 monthly of therapy. Decrease in IgA and C4. „Tendency“ to an increase in circulating immune complexes.</td>
</tr>
<tr>
<td>Colao A et al.¹⁶</td>
<td>100 patients with active acromegaly; 200 controls</td>
<td>Analysis of lymphocyte subset patterns</td>
<td>Increased levels of CD 3 and CD 4, decreased levels of CD 8 and CD 19 in patients [increase in T cell activity decreased B cell activity].</td>
</tr>
<tr>
<td>Sneppen SB et al.¹⁴</td>
<td>19 adults with severe GH deficiency; 110 healthy adults</td>
<td>Analysis of lymphocyte subsets and NK cell activity before and 18 months of randomized treatment with GH or placebo</td>
<td>NK cell activity [stimulated or unstimulated by Interferon Alpha or Interleukin 2] significantly impaired in GH deficient patients. No changes during therapy.</td>
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<tr>
<td>Mukherjee A et al.⁴</td>
<td>21 panhypopituitary patients under stable hormonal replacement incl. GH; 12 controls</td>
<td>Analysis of lymphocyte subsets, pneumococcal antibody levels, pre- and 1 month after polysaccharide vaccination; T-cell numbers, in vitro IFN gamma response</td>
<td>Lower B cell numbers in the patient group; lower antibody response to polysaccharide antigen in 9 of the patients.</td>
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<tr>
<td>Szynaka E et al.⁷</td>
<td>30 children and young people with GH deficiency; 25 healthy children</td>
<td>Lymphocyte subsets and concentration of IgA, IgG, IgM before and 6 months during GH therapy</td>
<td>Increase in percentage of active granulocytes after 6 months, significant decrease in IgM and IgA [but within normal range].</td>
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<tr>
<td>Campos VC et al.⁵</td>
<td>35 adults with isolated GH deficiency; 31 healthy controls</td>
<td>Skin tests and response to vaccination for hepatitis B, tetanus and bacillus Calmette-Guerin</td>
<td>Patients with GH deficiency had lower total IgG; no detectable clinical impact.</td>
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GH, Growth hormone

Conclusion

Although possibly compromised by hypopituitarism, by NAFLD and the age, the immune function of the patient described here, was obviously „good enough“ to overcome acute HEV infection. Other data on the course of acute hepatitis E in hypopituitarism have previously not been published. Whether the course of hepatitis E in patients with panhypopituitarism under insufficient hormonal replacement therapy is more complicated, remains to be elucidated.

Acknowledgements

None.

Conflict of interest

The authors declare, that there is no conflict of interest regarding the publication of this paper.

References


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