**LIVER DISEASE**

Fatigue is associated with high circulating leptin levels in chronic hepatitis C

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**Background and aims:** Fatigue is a frequent and disabling symptom reported by patients with chronic hepatitis C (CHC). Its mechanism is poorly understood. Recent attention has focused on the role of leptin and energy expenditure in CHC. Our aim was to analyse fatigue in CHC and to determine its relationship with disease activity, resting energy expenditure (REE), circulating leptin, and tumour necrosis factor α (TNF-α).

**Methods:** Seventy-eight CHC patients, 22 healthy controls, and 13 primary biliary cirrhosis (PBC) patients underwent measurements of REE, body composition, leptin, and TNF-α. All subjects completed the fatigue impact scale (FIS) questionnaire. A liver biopsy and viral load measurements were performed in all patients.

**Results:** Thirty-eight of 78 CHC patients considered fatigue the worst or initial symptom of their disease. The fatigue score of patients was significantly higher than that of controls (53.2 (40.1) vs 17.7 (16.9); p=0.001) and was more pronounced in females (p=0.003). Leptin was increased significantly in CHC patients compared with controls (15.4 (20.7) vs 9.6 (4.1) μg/ml; p=0.05). In CHC patients, the fatigue score correlated significantly with leptin corrected for fat mass (r=0.30, p=0.01). This correlation increased when the physical domain of fatigue was included (r=0.39, p=0.009). Furthermore, a similar positive correlation was found in PBC patients (r=0.56, p=0.04). No correlation was found between fatigue and age, REE, liver function tests, viral load, or the METAVIR score in CHC patients.

**Conclusions:** Fatigue is present in CHC patients and is more pronounced in females. The FIS questionnaire is clinically relevant and may be useful for future therapeutic trials aimed at reducing fatigue. Fatigue may be partly mediated by leptin.

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Fatigue is one of the most common complaints of patients with chronic hepatitis C (CHC). Despite its significance as a crucial issue for health care in CHC, few studies have investigated this symptom. One of the major obstacles to research is the highly non-specific nature of fatigue. Accordingly, all experimental approaches have been hampered by the lack of any serious objective tools to assess the subjective experience of fatigue. In 1994, the fatigue impact scale (FIS) was developed to improve our understanding of the effects of fatigue on quality of life. In contrast, leptin has been implicated in the pathogenesis of chronic liver diseases. Increased cytokine production has also been related to fatigue in various conditions. However, Gershon et al failed to detect any relationship between fatigue and TNF-α levels in patients with CHC.

We therefore designed a prospective study to evaluate fatigue in patients with CHC using the FIS questionnaire, and to determine the relationship between fatigue and the severity of liver disease, REE, serum leptin, and TNF-α levels.

**METHODS**

**Subjects**

Seventy-eight consecutive patients admitted to our liver unit for compensated liver disease due to CHC were included in the study. The diagnosis of CHC was based on the association of: (a) elevation of serum alanine aminotransferase (ALT) above normal levels; (b) risk factors for chronic viral hepatitis such as alcohol abuse, injection drug use, and sexual transmission; and (c) histological evidence of chronic active hepatitis.

**Abbreviations:** CHC, chronic hepatitis C; FIS, fatigue impact scale; REE, resting energy expenditure; FM, fat mass; FFM, fat free mass; TNF-α, tumour necrosis factor α; ALT, alanine aminotransferase; BMI, body mass index; HCV, hepatitis C virus; PBC, primary biliary cirrhosis.
40 U/l (upper normal limit) for six months or longer, (b) presence of anti-hepatitis C virus (HCV) antibodies, (c) presence of HCV viraemia, and (d) exclusion of other causes of chronic liver disease (alcoholism, chronic hepatitis B, Wilson’s disease, hepatotoxic drugs, haemochromatosis, α-antitrypsin deficiency, autoimmune chronic active hepatitis). A liver biopsy was performed in all patients, none of whom was cirrhotic. These patients had no evidence of dehiscence or overhydration or any other acute or chronic disease suspected of causing hypermetabolism (including human immunodeficiency virus). Furthermore, none had renal failure, thyroid disease, or clinically apparent depression, and none was treated with beta blockers or had received antiviral therapy.

Twenty two healthy volunteers formed the control group. These subjects were considered normal on the basis of history, physical examination, and biochemical blood screening (normal transaminase activity, no antibodies for HCV or human immunodeficiency virus, and negative hepatitis B surface antigen). There were no differences between patients and controls for age, sex, or body mass index (BMI). The characteristics of the CHC patients and controls are presented in table 1.

Table 1 Characteristics of the study population

<table>
<thead>
<tr>
<th>Variable</th>
<th>Female (n=34)</th>
<th>Male (n=44)</th>
<th>Controls (n=11)</th>
<th>Controls (n=11)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>42.5 (12.6)</td>
<td>40.0 (10.1)</td>
<td>37.4 (13.2)</td>
<td>41.0 (15.1)</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>24.6 (5.2)</td>
<td>23.8 (4.0)</td>
<td>21.0 (5.2)</td>
<td>23.2 (2.0)</td>
</tr>
<tr>
<td>Body weight (kg)</td>
<td>65.3 (17.3)</td>
<td>73.6 (14.4)</td>
<td>56.9 (6.1)</td>
<td>68.3 (8.0)</td>
</tr>
<tr>
<td>FFM (kg)</td>
<td>41.9 (9.2)</td>
<td>55.4 (9.8)</td>
<td>42.8 (6.2)</td>
<td>55.7 (6.6)</td>
</tr>
<tr>
<td>FM (kg)</td>
<td>23.3 (9.8)**</td>
<td>18.3 (8.7)*</td>
<td>14.1 (2.3)</td>
<td>12.6 (4.4)</td>
</tr>
<tr>
<td>ALT (U/l)</td>
<td>69.0 (59.5)**</td>
<td>106.9 (68.0)**</td>
<td>10.0 (2.6)</td>
<td>20.3 (1.38)</td>
</tr>
<tr>
<td>Vireal load (10⁶ copies/ml)</td>
<td>11.7 (14.4)</td>
<td>—</td>
<td>10.3 (13.1)</td>
<td>—</td>
</tr>
<tr>
<td>METAVIR (range 0–3)</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Activity index</td>
<td>1.0 (0.5)</td>
<td>1.0 (0.6)</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Fibrosis index</td>
<td>1.3 (1.0)</td>
<td>1.4 (1.0)</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

BMI, body mass index; FFM, fat free mass; FM, fat mass; ALT, alanine aminotransferase.
*p<0.05; **p<0.01 versus controls.
—, data not evaluated.

Assessment of fatigue
The impact of fatigue on quality of life was measured by the FIS, previously validated by Fisk and colleagues. The questionnaire was translated into French by Huet and colleagues, and has been used to compare fatigue in patients with CHC and healthy blood donors in Montreal. FIS is a self administered questionnaire consisting of 40 statements that describe possible manifestations of fatigue; these statements are classed into three health categories (cognitive (n=10), physical (n=10), and psychosocial (n=20)). Each item is rated on a five point scale of distress, ranging from 0 (“no problem”) to 4 (“extreme problem”) with a maximum of 160 points. FIS also includes a visual analogue scale allowing evaluation of the “disabling effect” of fatigue on a scale of 1 (“no disabling effect”) to 10 (“severe disabling effect”). Both patients and controls agreed to answer the FIS questionnaire after receiving counselling from one of the authors (EG). Questionnaires were completed at the clinic one week before clinical investigations.

Resting energy expenditure and body composition
REE was measured in the Functional Explorations Unit of Archet Hospital in the morning, after a 12 hour overnight fast. Participants were instructed to abstain from alcohol (including alcohol containing products and drugs) and smoking for 12 hours. After a 30 minute rest in the supine position, patients were placed in a semirecumbent position and measurements were taken using a ventilated hood open circuit indirect calorimeter (Deltatrac; Datex Instruments, Helsinki, Finland). After equilibrium was reached (10–20 minutes), respiratory exchanges were monitored continuously over a 30 minute period; data were obtained every minute and averaged over the 30 minutes. The system was checked weekly by burning ethanol under standard conditions, and calibrated immediately before each measurement with two standard gases. REE was calculated from the oxygen consumption rate and carbon dioxide production rate, using the modified Weir formula and was expressed in kilocalories per 24 hours. REE was also expressed as a ratio of FFM (REE/FFM) in kilocalories per kilogram of FFM per 24 hours.

FFM was measured by bipolar bioelectrical impedance analysis with an alternating electric current (50 µA) at two frequencies: 1 MHz and 5 kHz, as previously described and validated by Boulier and colleagues. The portable impedance analyser (IMP BO 1; L’impulsion, Caen, France) was equipped with a microprocessor; a computer was used to calculate impedance and body composition. Measurements were taken in the morning under the conditions described below. Subjects had been supine for 30 minutes, arms relaxed at the sides but not touching the body. Two stainless steel needles were
inserted subcutaneously: one on the antero-internal side of the foot, the other in the first intermetacarpal space of the dorsal aspect of the contralateral hand. FFM and fat mass (FM) were expressed in kilograms.

**Leptin and TNF-α assays**

Serum samples were stored at −80°C until assayed; all assays were performed in duplicate. Serum leptin levels were measured using a specific radioimmunoassay for human leptin (Linco Research, St Charles, Missouri, USA) with a limit of detection of 0.5 ng/ml. Serum TNF-α concentrations were determined by a highly sensitive quantitative immunoassay (R&D Systems, Minneapolis, Minnesota, USA) which is linear to 32 pg/ml and has a sensitivity of 0.18 pg/ml. For both leptin and TNF-α, intra- and interassay coefficients of variation were less than 8%.

**Statistical analysis**

Quantitative data are expressed as mean (SD) and were compared using the Mann-Whitney U test and Kruskal-Wallis test. The χ² test was used to compare qualitative data. Relationships between fatigue score, and the visual analogue scale used to rate the disabling effect of fatigue (r=0.83, p<0.0001). The severity of fatigue in patients with CHC was similar to that in PBC (49.4 (32.3)).

**Energy expenditure**

REE/FFM was significantly increased in patients with CHC compared with controls (32.4 (5.1) v 28.0 (3.3) kcal/kg FFM/24 hours; p<0.0001). Females were more hypermetabolic than males (34.0 (5.4) v 31.2 (4.6) kcal/kg FFM/24 hours; p=0.02). REE/FFM was similar in patients with PBC (29.9 (4.5)) and CHC.

**Leptin levels**

Mean fasting serum leptin level was significantly higher in patients with CHC than in controls (15.4 (20.7) v 6.4 (4.1) ng/ml; p<0.05). Figure 2 shows serum leptin levels expressed as both absolute values and after adjustment per kilogram of FM. Absolute leptin levels were significantly higher in females than in males both in patients and controls (p<0.0001) and not in controls (p=0.09). Compared with controls, absolute leptin values were significantly increased in females (27.4 (25.0) v 8.7 (3.6) ng/ml; p=0.01) but not in males (6.9 (9.5) v 4.1 (3.2) ng/ml; NS). When expressed per kilogram of FM, leptin levels were increased in females (1.1 (0.8) v 0.6 (0.2) ng/ml/kg) but the difference was not statistically significant (p=0.06). No difference was observed in males (0.3 (0.2) v 0.3 (0.3) ng/ml/kg). Leptin adjusted for FM was significantly higher in PBC patients compared with CHC patients (1.3 (1.2) v 0.7 (0.7) ng/ml/kg; p=0.02).

**TNF levels**

Serum TNF-α was similar in patients with CHC and controls (1.7 (1.6) v 1.6 (1.7) pg/ml; NS). In females, but not in males, serum TNF-α levels were significantly higher in patients with CHC compared with controls (1.7 (0.7) v 1.0 (0.7) pg/ml in females (p=0.03); 1.7 (0.6) v 2.0 (1.2) pg/ml in males (NS)) (fig 3). Serum TNF-α was significantly higher in PBC patients (3.4 (1.7) pg/ml) compared with both patients with CHC and controls (p<0.01).

**Correlation analysis**

Leptin levels were correlated with FM in both female and male patients with CHC (r=0.64, p<0.0001; r=0.73, p<0.0001,
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In patients with CHC, the total fatigue score was significantly correlated with leptin levels both when expressed as absolute values ($r=0.30$, $p=0.006$) and after correction for FM ($r=0.30$, $p=0.01$). The correlation was more pronounced for the physical domain of fatigue ($r=0.40$, $p=0.0003$ with leptin; $r=0.39$, $p=0.0009$ with leptin corrected for FM) (fig 4). Fatigue (and each subscale) was not correlated with age, weight, BMI, REE/FFM, or FM, or with standard liver function tests (bilirubin, ALT, aspartate aminotransferase, prothrombin index), viral load, or METAVIR score.

In PBC patients, a significant positive correlation ($r=0.56$, $p=0.0003$ with leptin corrected for FM) (fig 4).

**DISCUSSION**

Fatigue is a common and disabling symptom of CHC that impairs the quality of life of infected patients. However, its pathogenesis has been poorly investigated. Our findings confirm that fatigue is a major complaint in patients with CHC and has a negative impact on their quality of life. We also suggest that leptin may be implicated in the pathogenesis of fatigue in CHC and that this mechanism may be sex dependent.

Quantification of fatigue is difficult. In the present study, this subjective symptom was assessed using a validated 40 item self administered questionnaire (FIS). This questionnaire was translated into French, and has been validated in a cohort of patients with PBC. This method allows evaluation of the perceived impact of fatigue on patients’ lives, the factors that affect patients’ perceptions of fatigue, and the affect of fatigue on the mental and general health of patients. Determination of the effect of fatigue on activities is a more sensitive

**Figure 3** Comparison of serum tumour necrosis factor α (TNF-α) levels in male and female patients with chronic hepatitis C (CHC) and controls. *$p=0.03$.

**Figure 4** Relationship between leptin and fatigue in chronic hepatitis C patients. Correlation between absolute leptin levels and total score of fatigue (A) and its physical domain (B). Correlation between leptin levels corrected for fat mass and the total score of fatigue (C) and its physical domain (D).

**Figure 5** Correlation between leptin levels corrected for fat mass and the total score of fatigue in primary biliary cirrhosis patients.
which has been recognised as one of the major secretogogues previously reported data, heterogeneous (chronic hepatitis C and B) and the number of anxiety. Moreover, neither the level of viral load nor the relationship between fatigue and the degree of underlying leptin may at least partially contribute to fatigue in CHC. We therefore suggest that elevated in leptin in patients with CHC mediates this behavioural effect. Obviously, the underlying mechanisms of fatigue are multifactorial. It is generally recognised that fatigue occurs through changes in the central nervous system, including altered function of the hypothalamic-pituitary-adrenal axis and/or neurotransmission. It has also been suggested that altered serotoninergic pathways might contribute to fatigue. Leptin is known to exert its biological effect through the OB receptor that is present in several regions of the body, including the hypothalamus. It has also been demonstrated that human leptin levels are inversely related to pituitary-adrenal function and that leptin induced sympathetic tone exists in humans. Interestingly, leptin has been associated with behavioural changes in various conditions, such as sleep apnoea in humans, and has been shown to modify the functional capacity of the skeletal muscle mass in mice. Concerning the relationship between leptin and serotonin neurotransmission, both intracerebral and intraperitoneal leptin injections induce a significant increase in diencephalic 5-hydroxytryptamine content in mice. Taken together, these findings raise the possibility of a leptin dependent mechanism for fatigue. Whether circulating leptin acts in other liver diseases through a central and/or peripheral pathway requires further investigation.

In conclusion, we have demonstrated that fatigue, objectively assessed by the FIS questionnaire, is present in patients with CHC and that females are more susceptible to fatigue than males. The FIS questionnaire can thus be considered clinically relevant and may prove useful for future therapeutic trials. Study findings also suggest that leptin may play a role in the mediation of fatigue in CHC patients.

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method than simply asking patients to rate fatigue. Our findings are in accordance with those recently reported by Hassoun and colleagues who used the same French translation of the questionnaire in CHC patients. As expected, the degree of fatigue was markedly increased in CHC patients compared with healthy controls; this difference was noted in all three domains exploring quality of life (cognitive, physical, and psychosocial). In our study, the severity of fatigue, as evaluated by the FIS questionnaire (53.2 (40.1)), was comparable with that observed in patients with PBC but less than that reported in patients with multiple sclerosis (62.8 (36.3)) or the chronic fatigue syndrome (94.2 (28.1)). In contrast with the findings of Hassoun and colleagues, we found fatigue to be more severe in female than male patients with CHC.

In common with previous authors, we failed to find any relationship between fatigue and the degree of underlying hepatitis, including transaminase activity, prothrombin index, and bilirubin. Moreover, neither the level of viral load nor the degree of liver inflammation or fibrosis were correlated with the severity of symptoms.

It is well established that fatigued patients have psychometric scores suggestive of both depression and/or somatic anxiety. Although this point was not specifically evaluated in our study, none of our patients with CHC had clinically apparent severe depression. Consistent with our previous findings, REE/FFM was increased in patients with CHC, and this hypermetabolism was more pronounced in female patients. However, we did not find any relationship between REE/FFM and fatigue (or each subscale), which indicates that hypermetabolism per se probably has little or no influence on the severity of fatigue in such patients.

Attention has recently been focused on the role of leptin in chronic liver diseases. In cirrhosis, leptin levels have been found to be increased in some cases but decreased in others. In patients with chronic viral hepatitis (without cirrhosis), absolute leptin levels and leptin corrected for FM have been found to be lower than in normal controls. In our CHC patients, absolute leptin levels were significantly increased in females compared with controls, and this increase tended to persist when leptin was corrected for FM (p=0.06). Possible reasons for the discrepancy with the findings of Testa et al are that their study population was heterogeneous (chronic hepatitis C and B) and the number of subjects in each group was relatively small. In agreement with previously reported data, we found leptin levels to be sex dependent. The higher serum leptin levels observed in female patients cannot be attributed solely to the higher FM in the female groups because leptin levels tended to be higher (p=0.06) in women even after correction for FM. A large sample size is probably necessary to further investigate this dichotomy. However, our data suggest that this sex dependency may be explained by activation of the TNF-α system, which has been recognised as one of the major secretogogues for leptin. Indeed, a significant increase in serum TNF-α levels was observed in our young female patients with CHC. We can also speculate on the possible sensitising effect of oestrogen on release of TNF-α. However, the absence of any correlation between TNF-α and leptin levels (data not shown) indicates that other secretogogues such as insulin, corticosteroids, or nitric oxide may interact with release of TNF-α. Furthermore, we did not measure the bound leptin isoform which has been related to the TNF-α system in symptomatic human immunodeficiency virus infection.

Concerning the relationship between fatigue and circulating leptin, we found a slight positive correlation between fatigue (especially the physical domain) and circulating leptin levels, even after adjustment for FM. Both fatigue and leptin levels were higher in female patients with CHC. No correlation was found between fatigue and FM. We therefore suggest that leptin may at least partially contribute to fatigue in CHC patients. The higher FM in our CHC patients remains unclear. We can cautiously speculate on the possible role of a reduction in their physical activity due to chronic fatigue. However, this important observation of higher FM in CHC patients does not influence our findings and needs to be confirmed. An important question is whether the observed elevation in leptin in patients with CHC mediates this behavioural effect. The observation of the same correlation in PBC patients suggests that the elevation in leptin is not related solely to HCV infection itself. Obviously, the underlying mechanisms of fatigue are multifactorial. It is generally recognised that fatigue occurs through changes in the central nervous system, including altered function of the hypothalamic-pituitary-adrenal axis and/or neurotransmission. It has also been suggested that altered serotoninergic pathways might contribute to fatigue.
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