



# Brain Perfusion Scintigraphy in Evaluation of Pathogenesis of Fatigue in Patients with Primary Biliary Cholangitis (PBC) – A Pilot Study

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**Abstract:** Primary biliary cholangitis (PBC) is an autoimmune cholestatic liver disease characterized by a breakdown of immune tolerance to mitochondrial and nuclear antigens, causing injury to the biliary epithelial cells. Fatigue is the commonest reported symptom in PBC and has a negative impact on patients' perceived quality of life, often through social isolation. It is unrelated to the severity of liver disease and appears unresponsive to current therapies, including ursodeoxycholic acid and transplantation. Fatigue in PBC is complex, with numerous associated peripheral and central nervous system (CNS) features. Initially, cholestasis causes degenerative CNS change affecting areas of the brain regulating autonomic dysfunction and sleep, and these changes lead directly to some manifestations of fatigue and the associated cognitive impairment. The aim of the study was to examine global cerebral blood flow with brain perfusion scintigraphy SPECT in well-defined group of PBC Caucasian patients with verbally reported fatigue. Twenty consecutive PBC female patients (median age 58.9, ranges 38-80; 4 cirrhotic) with mean duration of the disease 3.3 years, were prospectively enrolled into the study. Fatigue Impact Scale (FIS) questionnaire was administered to every patients at the moment of brain examination. Brain perfusion scintigraphy SPECT was performed after intravenous injection of 760-800 MBq of technetium<sup>99m</sup> labeled exametazime (<sup>99m</sup>Tc –HMPAO). Then patients were examined using double head gamma camera system, and data were analyzed with dedicated nuclear medicine software. In analyzed cohort the median FIS score was 70.5 points (ranges 21-160), which was higher than previously reported. There were no correlation between age of patients at the SPECT/FIS examination, duration of the disease, the presence of liver cirrhosis, Mayo Risk Score, and FIS domains: *Cognitive*, *Physical* and *Social*, as well as with brain blood flow. However, positive correlation between *Cognitive* dimension of FIS measure and right frontal lobe perfusion impairment assessed with SPECT technic ( $p < 0.05$ ) was found. The results of this study showed that right lobe perfusion impairment might impact brain function in PBC. Cognitive dimension was stage-independent symptom in analyzed cohort, and cognitive impairment might be, in turn, associated with functional brain lesion.

**Keywords:** Brain Perfusion Scintigraphy, Primary Biliary Cholangitis (PBC), Fatigue, Cognitive Impairment

## 1. Introduction

Primary biliary cholangitis (formerly primary biliary cirrhosis, PBC) is a chronic, cholestatic, autoimmune liver

disease, affecting mainly middle age women. Management of PBC has traditionally focused on preventing advanced disease, liver failure, and cirrhosis. However, patients often experience profound fatigue, memory impairment, and excessive daytime somnolence which are seemingly

unrelated to the disease severity. The processes underpinning these symptoms are unknown, and there are currently no effective treatments. Fatigue affects between 60% and 80% of PBC patients, with impairment of their quality of life and their perception of their own mental health; it is not related to exercise, nor is it improved by rest [1-3]. The combination of fatigue, cognitive symptoms, and sleep disturbances points to central nervous system dysfunction as a contributing factors [4]. There are some evidences that fatigue in PBC is associated with abnormalities of autonomic function (e.g. baroreflex sensitivity, heart rate variability, blood pressure regulation, orthostatic intolerance) [5-8]. Brain change can occur in PBC, potentially as a result of cholestatic and/or inflammatory processes. This change is linked to systemic symptoms of fatigue and cognitive impairment. The fatigue phenotype appears to be highly stable and its presence is independently associated with a significantly risk of death in general, and cardiac death in particular [9, 10]. It might not disappeared after liver transplantation [11]. The aim of the study was to examine global cerebral blood flow with single-photon emission computed tomography (SPECT) brain perfusion scintigraphy in well-defined group of PBC Caucasian patients with verbally reported fatigue.

## 2. Patients and Methods

Twenty consecutive PBC female patients (mean age  $58.9 \pm 10.2$  years) with mean duration of the disease 3.3 years, were prospectively enrolled into the study. PBC was diagnosed according to the latest guidelines issued by the European Association for the Study of the Liver. All the participant verbally reported chronic fatigue. In four (20%) liver cirrhosis was diagnosed. Their clinical data are summarized in table 1.

**Table 1.** Clinical characteristics of the study group.

	Median (ranges)	Mean $\pm$ SD
Age at diagnosis (years)	54 (37 – 78)	55.6 $\pm$ 9.6
Age at examination (years)	60 (38 – 80)	58.9 $\pm$ 10.2
Mayo Risk Score (points)	4.27 (2.42 – 7.72)	4.475 $\pm$ 1.448
Cognitive dimension FIS (points)	13.5 (3 – 40)	16.5 $\pm$ 9.6
Physical dimension FIS (points)	19.5 (4 – 40)	20.8 $\pm$ 8.9
Social dimension FIS (points)	28.5 (9 – 80)	33.2 $\pm$ 17.9
FIS total score (points)	61.5 (3 – 80)	70.5 $\pm$ 36.4

### 2.1. Fatigue Impact Scale

Fatigue Impact Scale (FIS) questionnaire was administered to every patients at the moment of brain SPECT examination. FIS measure was developed and validated in patients with chronic fatigue syndrome [12] and PBC [13], and consisted of 40 questions to evaluate the impact of perceived fatigue during the last month on 3 subscales: physical, cognitive and psychosocial. Each item uses a 5-grade scale with higher scores denoting increased fatigue. Fatigue Impact Scale was applied in many studies regarding fatigue in PBC patients [1, 10, 14-23].

### 2.2. Single-Photon Emission Computed Tomography (SPECT)

Brain perfusion scintigraphy was performed after intravenous injection of 760-800 MBq of  $^{99m}\text{Tc}$  - HmPAO. Prior to injection radiochemical purity was controlled using chromatography and yield of labeling was more than 95%. After 20-30 minutes post injection patients were examined using double head gamma camera system (Infinia Hawkeye 4, GE). Acquisitions parameters were: matrix 128 X 128 pixels, 20 seconds per view, 64 views. Raw data was reconstructed with Filtered Back Projection (FBP) and analyzed with nuclear medicine dedicated neurological software (Neuro Gam, Segami).

### 2.3. Ethics

Appropriate informed consent was obtained from each patient included in the study. The study protocol was approved by the ethics committee of Pomeranian Medical University and confronts with the ethical guidelines of the 1975 Declaration of Helsinki (6th revision, 2008).

### 2.4. Statistics

All values were shown as median (and ranges), and mean  $\pm$  SD. All statistical tests were performed in SPSS 15.0 program. To determine whether there is correlation between the different domains of FIS tool, and between age and laboratory measures, the Spearman's correlation coefficient was calculated and  $p$  values less than 0.05 were considered statistically significant.

## 3. Results and Discussion

### 3.1. Results

In analyzed cohort the median FIS score was 70.5 points (ranges 21-160), which was higher than previously reported [16-19, 22, 23]. There were no correlation between age of patients at the SPECT/FIS examination, duration of the disease, the presence of liver cirrhosis, Mayo Risk Score, and FIS domains: *Cognitive* (mean  $16.5 \pm 9.6$  points), *Physical* ( $20.8 \pm 8.9$  points) and *Social* ( $33.2 \pm 17.9$  points), as well as with brain blood flow. However, positive correlation between *Cognitive* dimension of FIS measure and decreased right frontal lobe perfusion assessed with SPECT technic ( $p = 0.026$ ) was found.

### 3.2. Discussion

SPECT brain perfusion scintigraphy showed the reduction of right frontal lobe brain perfusion. This finding was associated with cognitive dimension in PBC chronically fatigued female patients.

Fatigue is a significant problem in approximately 50% of primary biliary cholangitis patients, impacting their quality of life together with the presence of social dysfunction that accompanies fatigue. The pathogenesis of fatigue remains

unclear, this symptom is unrelated to the PBC severity and is unresponsive to ursodeoxycholic acid treatment as well as liver transplantation. The complex and multifactorial origin of fatigue is supported with association with depression, autonomic dysfunction and sleep disturbances. It is postulated that chronic cholestasis and inflammation impact autonomic centers in central nervous system leading to cognitive impairments and sleep disturbances. The neuroimaging abnormalities suggest that the brain changes seen in PBC occur early in the pathological process, even before significant liver damage has occurred [24, 25]. Cognitive impairment in PBC patients was previously reported by groups of Raszeja-Wyszomirska *et al.* [26] and Newton *et al.* [27] as the symptom independent of liver disease severity and associated with poorer performance on objective cognitive testing. Cognitive impairment was, in turn, associated with structural brain lesions and autonomic dysfunction, which might predict risk of cognitive decline in the study of Newton *et al.* [27]

Brain perfusion scintigraphy provides dynamic, functional measures of brain function based on regional cerebral blood flow and tracer accumulation in brain grey matter compared with the static, structural data provided by conventional computerized axial tomography or magnetic resonance imaging (MRI). Mental fatigue was related to brain activity during the fatiguing cognitive task and significant positive relationships were found for cerebellar, temporal, cingulate and frontal regions on MRI [28]. Neuroimaging research in chronic fatigue has found white matter lesions in the frontal area on MRI and cerebral hypoperfusion in the brain stem on SPECT [29], although hypoperfusion has been found in several brain regions, and the findings varied across research centers. Patients with fatigue and without comorbid psychopathology have been found to have more white matter lesions on MRI [30], and more brain stem hypoperfusion [31]. Fischler *et al.* found positive associations between frontal blood flow, objective and subjective measure of cognitive function, and depressive symptoms [32]. However, these findings were not supported by the result of the other study with SPECT brain perfusion scintigraphy [25]. SPECT abnormalities occur more frequently and in greater numbers than MR abnormalities in patients with chronic fatigue syndrome [33], but there are no study regarding SPECT scanning of cerebral blood flow in PBC patients with fatigue to date.

The study group consisted of females with chronic fatigue in the course of PBC, and, of importance, previous cerebral perfusion studies of normal subjects indicated that flow is higher in women than in men [34]. Patients with chronic fatigue had reduced global cerebral blood flow in Yoshiuchi *et al.* study [35], supported the previous results regarding global hypoperfusion using SPECT brain imaging [33, 36]. Chronically fatigued patients had decreased blood flow in bilateral middle cerebral artery territories with increased perfusion in cerebellum [25, 31]. Thus, the result of the recent study, together with the results of Carbone *et al.* study [37] pointed to possible new aspects of fatigue in PBC, i.e. higher brain sensitivity to reduced blood flow in female

patients. This hypothesis might be supported with the result of Ahboucha *et al.* study. They showed that plasma levels of progesterone metabolites were significantly higher in PBC patients and in those with fatigue in particular. Some of progesterone metabolites are positive allosteric modulators of the gamma-aminobutyric acid type A (GABA-A) receptor and readily cross the blood-brain barrier. It might be that increased inhibition through GABA-A receptors due to the accumulation of neuroinhibitory steroids may represent an important pathophysiological mechanism of fatigue in chronic liver diseases [14].

Paired-pulse trans-cranial magnetic stimulation showed, that PBC patients had a significantly lower central activation before fatiguing exercise and transplanted and non-transplanted patients show similar abnormalities [4]. This finding pointed to potentially irreversible and progressive brain impairment in PBC patients, and it was supported with the results of the other studies. Newton *et al.* and Firth *et al.* noticed that cognitive symptoms (memory and concentration) were prevalent in PBC independently of liver disease severity [27, 38], similarly to findings from this recent study. On the other hand cognitive impairment indicated the role of autonomic dysfunction. Hollingsworth *et al.* showed the presence of increased cerebral vascular resistance and abnormal cerebral autoregulation in PBC patients, with a potentially important association between the degree of abnormality in structural changes in the globus pallidus. These findings suggested that organic brain injury in PBC might be directly related to autonomic dysfunction [39].

## 4. Conclusion

Chronic fatigue is one of the most debilitating symptom in primary biliary cholangitis affecting a large number of patients. Cholestatic or inflammatory process is linked to systemic symptoms of fatigue and cognitive impairment in PBC. SPECT brain perfusion scintigraphy seems to be useful diagnostic tool in chronically fatigued patients. The results of this pilot study confirmed central nervous system origin of cognitive dimension in PBC patients, independently from severity of liver disease. Right lobe perfusion impairment might impact brain function, and cognitive impairment might be, in turn, associated with functional brain lesion.

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