

No Signs of Neuroinflammation in Women With Chronic Fatigue Syndrome or Q Fever Fatigue Syndrome Using the TSPO Ligand [¹¹C]-PK11195

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Neurol Neuroimmunol Neuroinflamm 2022;9:e1113. doi:10.1212/NXI.0000000000001113

Abstract

Background and Objectives

The pathophysiology of chronic fatigue syndrome (CFS) and Q fever fatigue syndrome (QFS) remains elusive. Recent data suggest a role for neuroinflammation as defined by increased expression of translocator protein (TSPO). In the present study, we investigated whether there are signs of neuroinflammation in female patients with CFS and QFS compared with healthy women, using PET with the TSPO ligand [¹¹C]-(R)-(2-chlorophenyl)-N-methyl-N-(1-methylpropyl)-3-isoquinoline-carboxamide ([¹¹C]-PK11195).

Methods

The study population consisted of patients with CFS (n = 9), patients with QFS (n = 10), and healthy subjects (HSs) (n = 9). All subjects were women, matched for age (±5 years) and neighborhood, aged between 18 and 59 years, who did not use any medication other than paracetamol or oral contraceptives, and were not vaccinated in the last 6 months. None of the subjects reported substance abuse in the past 3 months or reported signs of underlying psychiatric disease on the Mini-International Neuropsychiatric Interview. All subjects underwent a [¹¹C]-PK11195 PET scan, and the [¹¹C]-PK11195 binding potential (BP_{ND}) was calculated.

Results

No statistically significant differences in BP_{ND} were found for patients with CFS or patients with QFS compared with HSs. BP_{ND} of [¹¹C]-PK11195 correlated with symptom severity scores in patients with QFS, but a negative correlation was found in patients with CFS.

Discussion

In contrast to what was previously reported for CFS, we found no significant difference in BP_{ND} of [¹¹C]-PK11195 when comparing patients with CFS or QFS with healthy neighborhood controls. In this small series, we were unable to find signs of neuroinflammation in patients with CFS and QFS.

Trial Registration Information

EudraCT number 2014-004448-37.

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Go to [Neurology.org/NN](https://www.neurology.org/NN) for full disclosures. Funding information is provided at the end of the article.

The Article Processing Charge was funded by the Radboud University Medical Center.

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Glossary

[¹¹C]-PK11195 = ¹¹C-(R)-(2-chlorophenyl)-N-methyl-N-(1-methylpropyl)-3-isoquinoline-carbox-amide; BDI-PC = Beck Depression Inventory for Primary Care; BP = binding potential; CDC = Centers for Disease Control; CFS = chronic fatigue syndrome; CIS = Checklist Individual Strength; HS = healthy subject; Ig = immunoglobulin; QFS = Q fever fatigue syndrome; SIP-8 = Sickness Impact Profile 8; TSPO = translocator protein.

Chronic fatigue syndrome (CFS) is characterized by a debilitating fatigue without a known somatic cause that lasts for at least 6 months and is often accompanied by headache, sore throat, musculoskeletal pain, and neuropsychological symptoms, mainly impairments in memory and concentration.¹ There is a strong female preponderance in CFS (~ 75%). Previous research on CFS investigating metabolism, hormones, microbes, the immune system, and neuropsychology failed to discover a unifying pathogenesis.²⁻⁶ Many patients with CFS had a previous infectious disease and are considered as postinfectious fatigue syndromes.^{7,8} Q fever fatigue syndrome (QFS) is such a postinfectious fatigue syndrome that is characterized by a state of prolonged fatigue following approximately 20% of acute Q fever infections.^{9,10} The fatigue lasts for at least 6 months and usually coincides with musculoskeletal complaints, neurocognitive problems, sleeping problems, headache, respiratory tract symptoms, and mood disorders.⁹ In many ways, complaints of QFS are similar as to those reported by patients with CFS, and like in CFS, the pathophysiology of QFS is still unclear.¹¹

Given the complaints that patients with CFS and patients with QFS have, it is conceivable that both an inflammatory and neurologic component contribute to their pathophysiology.^{5,12,13} A hypothesis that connects these pathophysiologic components is that of chronic low-grade neuroinflammation.⁵ Within the context of this hypothesis, a peripheral inflammatory response, for example, initiated by Q fever, ultimately extends to resident tissue macrophages, that is, microglia, of the brain.¹⁴ Trained immunity of microglia occurs following an inflammatory or noxious stimulus. This initial stimulus elicits long-term changes that enable these cells to produce an enhanced inflammatory response on a second, nonspecific, stimulus. In this way, chronic low-grade neuroinflammation may persist following a transient single infectious insult.¹⁵ Microglia may be indirectly primed through active and passive transport of cytokines across the blood-brain barrier and stimulation of peripheral chemoreceptors of the vagal nerve.¹⁶

Through investigation of inflammatory markers in CSF, MR spectroscopy, and PET, several studies point toward neuroinflammation occurring in CFS.^{5,17-19} As microglial activation is its trademark characteristic, visualizing neuroinflammation is best done by PET neuroimaging using a radioligand that binds to the, increased expression of, 18-kD translocator protein (TSPO) in activated microglia and astrocytes.¹⁶ In recent years, Nakatomi et al.⁵ showed that compared with healthy subjects (HSs), patients with CFS exhibit an

increased PET signal, especially at the thalamus, showing positive correlation with pain scores, when using the ¹¹C-(R)-(2-chlorophenyl)-N-methyl-N-(1-methylpropyl)-3-isoquinoline-carbox-amide ([¹¹C]-PK11195) ligand for TSPO. These and other findings^{20,21} suggest that patients with CFS exhibit neuroinflammation and that this phenomenon warrants further investigation in the pathophysiology of CFSs. Given the overlap in symptoms with patients with CFS and apparent inflammatory etiology, we expect patients with QFS to exhibit similar or more signs of neuroinflammation than patients with CFS.

This study aimed to confirm and further investigate a neuroinflammatory substrate in patients with CFS and QFS by using the TSPO ligand [¹¹C]-PK11195 for PET neuroimaging. Through analysis of questionnaires, we intended to correlate findings of neuroinflammation with psychiatric and physical well-being.

Methods

Study Population

The study population consisted of patients with CFS (n = 9), patients with QFS (n = 10), and HSs (n = 9). For reasons of homogeneity, all subjects were women and matched for age (±5 years) and neighborhood. All subjects were aged between 18 and 59 years, did not use any medication other than paracetamol or oral contraceptives, and were not vaccinated in the last 6 months. None of the subjects reported substance abuse in the past 3 months or showed signs of underlying psychiatric disease, that is, depression, bipolar disorders, anxiety, schizophrenia, psychosis, or eating disorders on Mini-International Neuropsychiatric Interview.

All patients with CFS were diagnosed with CFS at the Department of Internal Medicine and Expert Center for Chronic Fatigue of the Radboud University Medical Center, Nijmegen, the Netherlands, after a uniform workup according to the Fukuda 1994 criteria for CFS.^{1,22} They all had a score ≥40 on the subscale fatigue severity of the Checklist Individual Strength (CIS) questionnaire²³ and a score ≥700 on the Sickness Impact Profile 8 (SIP-8) questionnaire.²⁴ None of them experienced an acute Q fever infection or were vaccinated against Q fever in the past. *Coxiella* PCR and immunoglobulin G (IgG) were not tested, and no data were collected on whether an infection preceded CFS complaints.

All patients with QFS were diagnosed at the Radboud Expert Center for Q fever, Nijmegen, the Netherlands, after a uniform workup according to the Dutch guideline on QFS diagnosis.²⁵ All patients with QFS met the following diagnostic criteria: (1) fatigue lasted ≥ 6 months; (2) sudden onset of severe fatigue (defined as a score ≥ 40 on the subscale fatigue severity of the CIS) or significant increase in fatigue, both related to a symptomatic acute Q fever infection; (3) chronic Q fever and other somatic or psychiatric causes of fatigue were excluded; and (4) fatigue resulted in significant functional impairment (defined as a total score ≥ 700 on the SIP-8 questionnaire). All patients with QFS tested negative on *Coxiella* PCR and had IgG phase I or phase II titers $\geq 1:16$, but IgG phase I $\leq 1:512$, and none of them showed serologic signs of an acute or recent Q-fever infection, reflected by IgM antibodies in the absence of IgG antibodies.

HSs were recruited based on age, sex, and neighborhood that matched with both patients with QFS and CFS and had a score ≤ 35 on the subscale fatigue severity of the CIS questionnaire and a score ≤ 450 on the SIP-8 questionnaire. Similar to patients, HSs did not use any medication other than paracetamol or contraceptives and were screened for psychiatric disease. None of the HSs had experienced acute Q fever infection or were vaccinated for Q fever. *Coxiella* PCR and IgG were not tested.

The sample size needed was estimated using the effect size (Cohen *d*) as calculated from a previous [¹¹C]-PK11195 study by Nakatomi et al. in 9 patients with CFS vs 10 HSs⁵ and the [¹¹C]-PK11195 binding potential (BP_{ND}) in HSs from a previous study performed by our group.²⁶ Effect sizes of the BP_{ND} for the different brain regions included by Nakatomi et al.⁵ ranged from 1.4 (hippocampus) to 2.4 (midbrain). Using these effect sizes, the BP_{ND} in patients with CFS was estimated from the BP_{ND} in HSs. For example, the BP_{ND} of the hippocampus in HSs was 1.37 ± 0.30 , leading to an estimated BP_{ND} of 1.79 (effect size of 1.4) or 2.84 (effect size of 2.4) in patients with CFS. Based on these estimations, and using an alpha of 0.05 and a power of 0.80, it was estimated that a group size of 9 (effect size of 1.4) to 3 (effect size of 2.4) was sufficient for finding significant differences in the BP_{ND}. Based on these calculations and the study by Nakatomi et al., we have chosen a group size with a minimum of $n = 9$.

Questionnaires

All subjects were asked to fill out questionnaires, used in our expert centers,²³ on CFS aspects previously associated with neuroinflammation, that is, depression, concomitant CFS complaints, and fatigue:^{5,27}

CIS, subscale on fatigue severity, assesses the severity of fatigue, which is part of the inclusion criteria.²³

SIP-8 assesses the influence of disease and/or health complaints on functioning in daily life, which is part of the inclusion criteria.²⁴

BDI-II-NL-PC Beck Depression Inventory for Primary Care (BDI-PC, shortened) assesses depressive symptoms.²⁸

Centers for Disease Control (CDC) CFS Symptom Inventory Questionnaire, subscale on active complaints, assesses concomitant CFS symptoms.²⁹

PET Imaging

Following testing for collateral blood flow and the injection of 1% lidocaine, a cannula was inserted in the radial artery to allow for arterial blood sampling. In the other arm, a cannula was placed in the antebrachial vein for the injection of [¹¹C]-PK11195. The PET scans were performed using the Biograph mCT (Siemens Healthineers, Germany). Head movement was minimized by using a head-restraining adhesive band. After positioning in the camera, a low-dose CT scan was made for attenuation and scatter correction. Hereafter, [¹¹C]-PK11195, produced under Good Manufacturing Practice conditions as described earlier,²⁶ was injected IV at a speed of 0.5 mL/s (total volume 8.3 mL). The injected dose of [¹¹C]-PK11195 was 367 ± 50 MBq (HSs, 370 ± 53 MBq; CFS, 375 ± 37 MBq, QFS, 356 ± 55 MBq) with a molar activity of $>12,000$ GBq/mmol. Simultaneously with the start of the injection, a 60-minute emission scan was started during which arterial blood radioactivity was continuously measured with an automated blood sampling system (COMECER Netherlands, the Netherlands). Five manual blood samples were collected at 10, 20, 30, 45, and 60 minutes after [¹¹C]-PK11195 injection to determine the amount of radioactivity in blood and plasma for calibration of the automated sampling system. The manual blood samples taken at 20, 45, and 60 minutes were additionally used for analysis of the percentage of intact [¹¹C]-PK11195 in plasma, according to the procedure described previously.²⁶ On the same day as the PET scan, a T1-weighted MRI scan, using a MAGNETOM Prisma (Siemens Healthineers, Germany), was made for anatomic reference.

PET Data Analysis

The list-mode data from the PET scans were reconstructed using the 3D OSEM algorithm (3 iterations and 24 subsets) into 24 successive frames (7*10, 2*30, 2*120, 2*180, 5*300, and 2*600 seconds). Image processing and pharmacokinetic analysis were performed with PMOD software v4.1 (PMOD Technologies Ltd., Switzerland). The summed PET image (frame 1–24) was used for rigid registration of the individual PET image to the individual MRI. The 6-tissue probability map normalization of the individual MRI into the Montreal Neurological Institute standard space was then performed and applied to the corresponding PET image. Predefined volumes of interest, based on the Hammers atlas,³⁰ were transformed back into individual PET space, and time activity curves were generated.

The 2-tissue compartment model was used to obtain the non-displaceable BP_{ND} of [¹¹C]-PK11195, using the metabolite-corrected plasma curve as the input function. The delay and the

blood volume were individually fitted. It was assumed that the distribution volume of the nondisplaceable compartment (K_1/k_2) and the dissociation rate (k_4) from the specific binding site were equal for all regions. A coupled fitting was performed for cortical regions that calculated a common K_1/k_2 and k_4 , which were then used to calculate an individual K_1 and k_3 for all regions. The BP_{ND} was defined as k_3/k_4 .

Statistical Analysis

Patient data were analyzed using GraphPad Prism (GraphPad Software Inc., version 5.03). An ANOVA was used to determine differences between groups. A Pearson correlation was used to determine correlations between BP_{ND} of [^{11}C]-PK11195 in various brain regions and symptom severity scores in patients with CFS and QFS. Statistical significance was attained if $p < 0.05$.

Standard Protocol Approvals, Registrations, and Patient Consents

All participants provided written informed consent, and the study was approved by the Medical Ethical Review Committee of the University Medical Center Groningen (UMCG NL51194.042.15).

Data Availability

The data sets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Results

Patients and Controls

At the time of PET imaging, patients with CFS had a significantly longer median duration of illness than patients with QFS (240 vs 84 months, $p = 0.01$). The median age of patients with CFS, patients with QFS, and HSs did not differ significantly (Table 1). Other than the fact that patients with CFS were significantly more functionally impaired than patients with QFS ($p = 0.02$), no significant differences in other scores were found between patients with CFS and QFS (Table 1).

BP_{ND} of [^{11}C]-PK11195 in Various Brain Regions

The BP_{ND} of [^{11}C]-PK11195 in various brain regions for patients with CFS and QFS compared with HSs was not significantly different (Figure 1 and Table 2). No differences were found in BP_{ND} values comparing means of patients with CFS and QFS with HSs for the cingulate (rostral anterior: mean difference -0.33 [95% CI, -0.82 to 0.17] and caudal anterior: mean difference -0.33 [95% CI, -0.83 to 0.16] and -0.20 [95% CI, -0.67 to 0.27] with $p = 0.29$ and 0.83 , respectively, posterior: mean difference -0.32 [95% CI, -0.85 to 0.22] and -0.18 [95% CI, -0.69 to 0.33] with $p = 0.43$ and 1.00 , respectively), hippocampus (mean difference -0.37 [95% CI, -0.90 to 0.16] and -0.23 [95% CI, -0.73 to 0.27] with $p = 0.25$ and 0.73 , respectively), thalamus (mean difference -0.32 [95% CI, -0.94 to 0.31] and -0.20 [95% CI,

-0.80 to 0.39] with $p = 0.61$ and 1.00 , respectively), midbrain (mean difference -0.40 [95% CI, -1.06 to 0.25] and -0.33 [95% CI, -0.95 to 0.29] with $p = 0.38$ and 0.54 , respectively), or pons (mean difference -0.44 [95% CI, -1.23 to 0.36] and -0.30 [95% CI, -1.05 to 0.45] with $p = 0.51$ and 0.95 , respectively). In fact, as seen by the mean difference, BP_{ND} values tended to be lower rather than higher for both patients with CFS and QFS compared with HSs.

Correlation Between Symptom Severity Scores and BP_{ND} of [^{11}C]-PK11195

Significant correlations between symptom severity scores and BP_{ND} of [^{11}C]-PK11195 in various brain regions of both patients with CFS and patients with QFS are shown in eTable 1 (links.lww.com/NXI/A662). For patients with CFS, the CDC questionnaire, subscale on complaints, and the CIS questionnaire, subscale fatigue severity, correlated with BP_{ND} in the caudate nucleus (-0.73 , $p < 0.05$ and -0.78 , $p < 0.05$, respectively) (eTable 1) (Figure 2 and 3). For patients with QFS, the CDC questionnaire, subscale on complaints, correlated with BP_{ND} in the brainstem (0.66 , $p < 0.05$), caudal anterior cingulate (0.64 , $p < 0.05$), insula (0.65 , $p < 0.05$), amygdala (0.71 , $p < 0.05$), and pons (0.69 , $p < 0.05$), and the CIS questionnaire, subscale on fatigue severity, correlated with BP_{ND} in the orbitofrontal cortex (0.70 , $p < 0.05$), middle frontal gyrus (0.83 , $p < 0.01$), inferior frontal gyrus (0.78 , $p < 0.01$), superior frontal gyrus (0.64 , $p < 0.05$), primary motor cortex (0.74 , $p < 0.05$), temporal lobe (0.78 , $p < 0.01$), primary somatosensory cortex (0.83 , $p < 0.01$), and parietal lobe (0.77 , $p < 0.01$) (eTable 1) (Figure 2 and 3). In HSs, no significant correlations were found between symptom severity scores and BP_{ND} of [^{11}C]-PK11195 (eTable 1).

Discussion

In this study, we aimed to investigate neuroinflammation in patients with CFS and QFS by using the TSPO ligand [^{11}C]-PK11195 for PET neuroimaging. No signs of neuroinflammation were seen in either patients with CFS or QFS. Our findings contradict previous findings in patients with CFS by Nakatomi et al.⁵ who found significantly increased BP_{ND} values in the cingulate, hippocampus, thalamus, midbrain, and pons. Although no signs of neuroinflammation were found, similar correlations between BP_{ND} of [^{11}C]-PK11195 and scores on questionnaires were found in the amygdala of patients with QFS, but not patients with CFS.

Although the setup of this study was similar to that of Nakatomi et al.⁵, using the same TSPO ligand ([^{11}C]-PK11195), a number of important differences can be discerned. First of all, for reasons of homogeneity, our study only included women. Around 75% of patients with CFS are female, and although the percentage of women in QFS is lower (52%),^{11,31} we felt that we should avoid a sex effect in a study with such a small sample size. Nakatomi et al. included 30%–40% males without presenting separate data for men

Table 1 Characteristics of Female Patients With QFS, Patients With CFS, and HSs

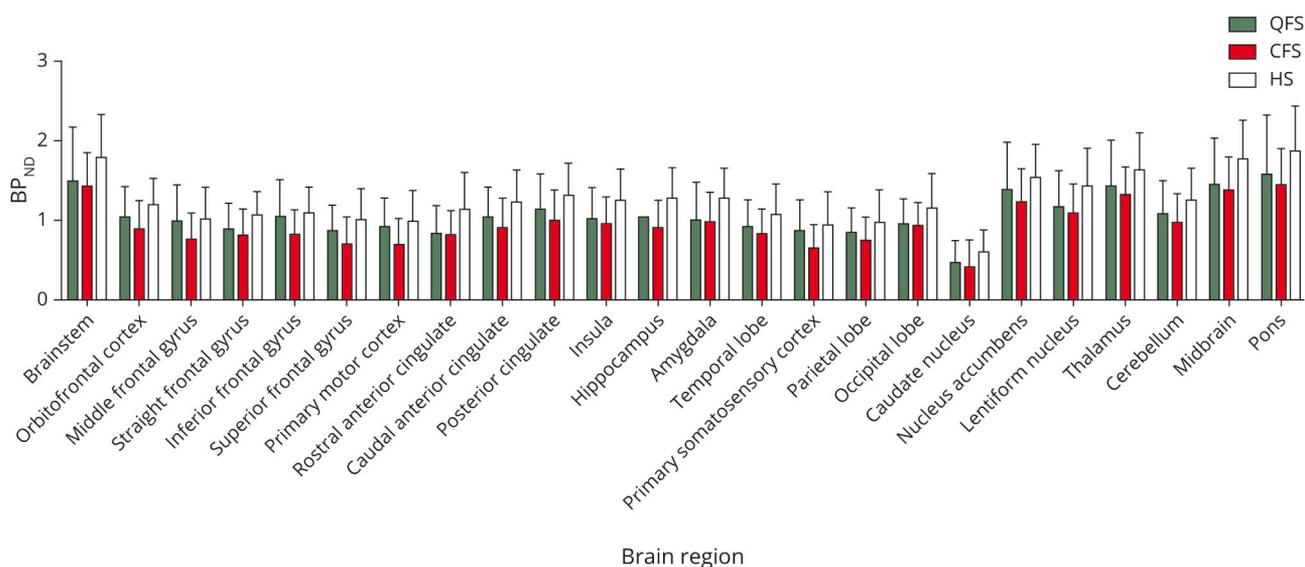
Characteristics	QFS (n = 10)	CFS (n = 9)	HSs (n = 9)
Age, y Median (IQR)	43 (32–48)	43 (30–52)	41 (27–47)
Duration of symptoms, mo^a Median (IQR)	84 (74–93)	240 (84–390)	—
CIS subscale fatigue severity score mean ± SD	50 ± 4.7	49 ± 5.0	12 ± 5.1
SIP-8 total score, mean ± SD	1,432 ± 362	1890 ± 395	0 ± 0
BDI-II-NL-PC score, mean ± SD	2.1 ± 1.5	3.8 ± 2.7	0.0 ± 0.0
CDC subscale on complaints score, mean ± SD	16 ± 4.1	17 ± 4.5	1 ± 1.7

Abbreviations: CDC = Centers For Disease Control; CFS = chronic fatigue syndrome; CIS = Checklist Individual Strength; HS = healthy subject; IQR = interquartile range; QFS = Q fever fatigue syndrome; SIP-8 = Sickness Impact Profile 8, BDI-II-NL-PC = Beck Depression Inventory for Primary Care.

^aSymptom duration: time onset of symptoms until blood sampling.

and women.^{5,31} This is important as inflammatory responses are generally higher in males.³² Also, in experimental mouse studies of traumatic brain injury, male mice are more likely to exhibit neuroinflammation compared with female mice.³³ One could argue that neuroinflammation is more likely to occur, and perhaps even persist, in males compared with females. However, if neuroinflammation is indeed present, the high percentage of female patients with CFS contradicts with this hypothesis. A second difference between our study and that of Nakatomi et al. is that we distinguished patients with CFS, with often heterogenic etiologies,^{5,31} from patients with postinfectious fatigue syndrome, that is, patients with QFS. Third, we used a neighborhood control group with healthy women who were matched with patients with CFS and QFS in terms of age and geographical area to accomplish optimal

matching and avoid bias due to confounding. Also, patients, especially those with CFS, who were included in our study had a longer duration of illness than those included in the study by Nakatomi et al. (reported mean of 62.4 months).⁵ When using small numbers of included patients, as is the case in both studies, subtle differences like these might contribute to the different outcomes that are seen. This brings us to a fifth and final difference, that is, the method used for determining the binding of [¹¹C]-PK11195. We used pharmacokinetic binding with an arterial input function, whereas Nakatomi et al. used the cerebellum as a reference region in reference tissue modeling. We feel that the latter is methodologically less sound as no brain region is devoid of TSPO, meaning that the cerebellum is not an objective reference region, and the cerebellum may actually be involved in the disease process.

Figure 1 BP_{ND} of [¹¹C]-PK11195 for Various Brain Regions in Patients With QFS, Patients With CFS, and HSs

Graph showing BP_{ND} of [¹¹C]-PK11195 per brain region for patients with QFS, patients with CFS, and HSs. Data are depicted as mean ± SD. BP_{ND} = nondisplaceable binding potential; CFS = chronic fatigue syndrome; HS = healthy subject; QFS = Q fever fatigue syndrome.

Table 2 BP_{ND} of [¹¹C]-PK11195 for Various Brain Regions in Patients With QFS and CFS Compared With HSs

Brain region	QFS Mean (95% CI)	CFS Mean (95% CI)	HS Mean (95% CI)	QFS vs HS Mean difference (95% CI), <i>p</i> value ^a	CFS vs HS Mean difference (95% CI), <i>p</i> value ^a
Brainstem	1.49 (1.00–1.98)	1.42 (1.06–1.78)	1.79 (1.34–2.24)	–0.30 (–1.00 to 0.40), 0.84	–0.37 (–1.11 to 0.37), 0.63
Orbitofrontal cortex	1.04 (0.77–1.31)	0.89 (0.60–1.18)	1.20 (0.93–1.48)	–0.16 (–0.60 to 0.28), 1.00	–0.31 (–0.77 to 0.15), 0.28
Middle frontal gyrus	0.99 (0.66–1.32)	0.76 (0.48–1.04)	1.02 (0.69–1.35)	–0.04 (–0.53 to 0.46), 1.00	–0.27 (–0.79 to 0.26), 0.61
Straight frontal gyrus	0.89 (0.66–1.12)	0.81 (0.54–1.09)	1.07 (0.82–1.32)	–0.18 (–0.57 to 0.20), 0.70	–0.26 (–0.67 to 0.15), 0.35
Inferior frontal gyrus	1.05 (0.72–1.38)	0.81 (0.54–1.08)	1.09 (0.82–1.37)	–0.44 (–0.51 to 0.42), 1.00	–0.29 (–0.78 to 0.21), 0.44
Superior frontal gyrus	0.88 (0.66–1.10)	0.71 (0.42–0.99)	1.02 (0.70–1.33)	–0.14 (–0.56 to 0.28), 1.00	–0.31 (–0.75 to 0.14), 0.26
Primary motor cortex	0.92 (0.66–1.18)	0.70 (0.42–0.97)	0.99 (0.68–1.31)	–0.08 (–0.51 to 0.36), 1.00	–0.30 (–0.76 to 0.16), 0.33
Rostral anterior cingulate	0.84 (0.59–1.09)	0.81 (0.55–1.07)	1.14 (0.75–1.53)	–0.30 (–0.77 to 0.17), 0.33	–0.33 (–0.82 to 0.17), 0.30
Caudal anterior cingulate	1.04 (0.76–1.31)	0.91 (0.60–1.22)	1.24 (0.91–1.57)	–0.20 (–0.67 to 0.27), 0.83	–0.33 (–0.83 to 0.16), 0.29
Posterior cingulate	1.14 (0.81–1.46)	1.00 (0.68–1.32)	1.32 (1.00–1.65)	–0.18 (–0.69 to 0.33), 1.00	–0.32 (–0.85 to 0.22), 0.43
Insula	1.02 (0.74–1.30)	0.96 (0.69–1.23)	1.26 (0.93–1.58)	–0.23 (–0.69 to 0.22), 0.60	–0.29 (–0.77 to 0.19), 0.39
Hippocampus	1.04 (0.71–1.37)	0.90 (0.62–1.19)	1.27 (0.95–1.60)	–0.23 (–0.73 to 0.27), 0.73	–0.37 (–0.90 to 0.16), 0.25
Amygdala	1.01 (0.68–1.35)	0.98 (0.67–1.29)	1.28 (0.96–1.60)	–0.26 (–0.77 to 0.25), 0.60	–0.30 (–0.83 to 0.24), 0.50
Temporal lobe	0.92 (0.67–1.16)	0.83 (0.58–1.09)	1.08 (0.76–1.39)	–0.16 (–0.58 to 0.26), 1.00	–0.24 (–0.69 to 0.20), 0.52
Primary somatosensory cortex	0.87 (0.59–1.15)	0.65 (0.41–0.90)	0.95 (0.60–1.29)	–0.08 (–0.53 to 0.38), 1.00	–0.30 (–0.77 to 0.18), 0.37
Parietal lobe	0.84 (0.62–1.10)	0.75 (0.51–0.99)	0.97 (0.63–1.32)	–0.13 (–0.54 to 0.28), 1.00	–0.23 (–0.66 to 0.21), 0.58
Occipital lobe	0.95 (0.73–1.18)	0.94 (0.70–1.18)	1.15 (0.79–1.52)	–0.20 (–0.63 to 0.23), 0.73	–0.21 (–0.66 to 0.24), 0.71
Caudate nucleus	0.47 (0.27–0.66)	0.42 (0.14–0.70)	0.61 (0.38–0.84)	–0.14 (–0.50 to 0.22), 0.98	–0.19 (–0.57 to 0.19), 0.61
Nucleus accumbens	1.39 (0.96–1.81)	1.23 (0.87–1.58)	1.53 (1.19–1.88)	–0.15 (–0.76 to 0.46), 1.00	–0.31 (–0.95 to 0.34), 0.69
Lentiform nucleus	1.17 (0.84–1.50)	1.09 (0.77–1.40)	1.44 (1.04–1.83)	–0.26 (–0.80 to 0.27), 0.64	–0.35 (–0.91 to 0.22), 0.38
Thalamus	1.43 (1.01–1.84)	1.31 (1.02–1.61)	1.63 (1.24–2.02)	–0.20 (–0.80 to 0.39), 1.00	–0.32 (–0.94 to 0.31), 0.61
Cerebellum	1.08 (0.78–1.38)	0.98 (0.68–1.28)	1.26 (0.92–1.59)	–0.18 (–0.66 to 0.31), 1.00	–0.28 (–0.79 to 0.23), 0.52
Midbrain	1.45 (1.03–1.86)	1.38 (1.02–1.73)	1.78 (1.37–2.18)	–0.33 (–0.95 to 0.29), 0.54	–0.40 (–1.06 to 0.25), 0.38
Pons	1.58 (1.04–2.11)	1.44 (1.05–1.83)	1.88 (1.41–2.34)	–0.30 (–1.05 to 0.45), 0.95	–0.44 (–1.23 to 0.36), 0.51

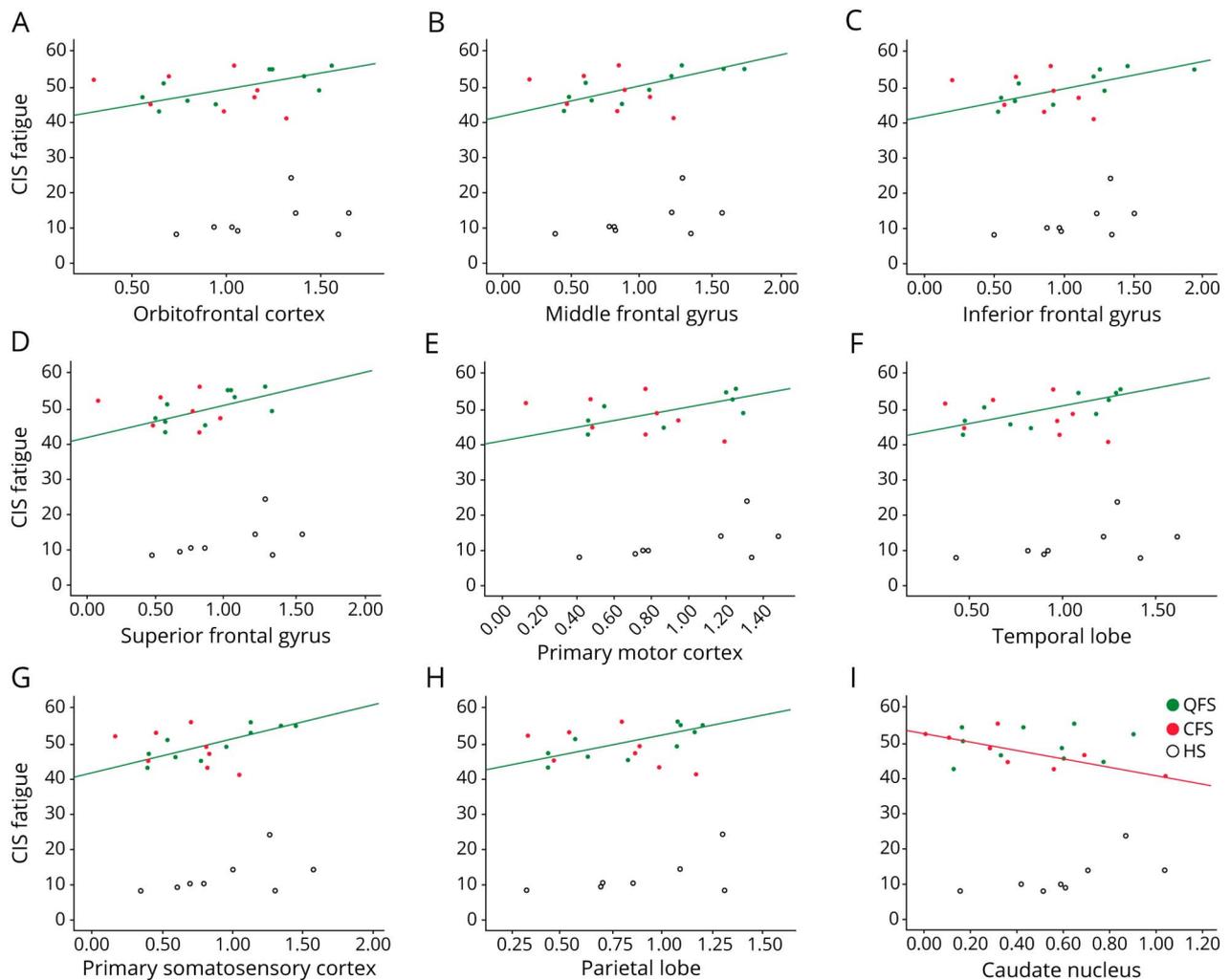
Abbreviations: BP_{ND} = nondisplaceable binding potential; CFS = chronic fatigue syndrome; HS = healthy subject; QFS = Q fever fatigue syndrome. Data are depicted as mean (difference) with 95% CI. A 1-way ANOVA with post hoc Bonferroni (^a) was used for comparing QFS and CFS with HS.

Whether binding of the [¹¹C]-PK11195 ligand is considered enhanced, normal, or even lowered may be explained by this difference in methodology.

Regarding the effect of disease duration, Hornig et al. previously reported that the inflammatory response, determined by cytokine measurements, is lower in patients with CFS with a long duration, that is, >39 months, of illness than in those with a short duration, that is, <39 months, of illness.³⁴ In a previous study, however, we were unable to confirm these findings.¹³ We found that patients with CFS, who had fatigue for a median of 240 months, show less signs of neuroinflammation than patients with QFS, who were fatigued for a median of 84 months. Given previous findings by Hornig et al.

and our observation that HSs generally showed a stronger signal of TSPO binding than patients, one could speculate that neuroinflammation wanes off over time and is followed by a refractory period with decreased expression of TSPO. Furthermore, studies on peripheral inflammatory cell metabolism in patients with CFS and QFS have repeatedly shown that mitochondria of these cells are likely to be affected.^{35–37} As TSPO is expressed in the outer mitochondrial membrane, it could be conceived that its expression is similarly affected in chronically fatigued patients. In lieu of further speculation, we implore a large longitudinal investigation of TSPO expression in the mitochondrial membrane of chronically fatigued patients and relate findings to symptom severity scores.

Figure 2 Correlation Between BP_{ND} of [¹¹C]-PK11195 in Various Brain Regions With CIS Questionnaire, Subscale on Fatigue Severity, Scores in Patients With QFS, Patients With CFS, and HS

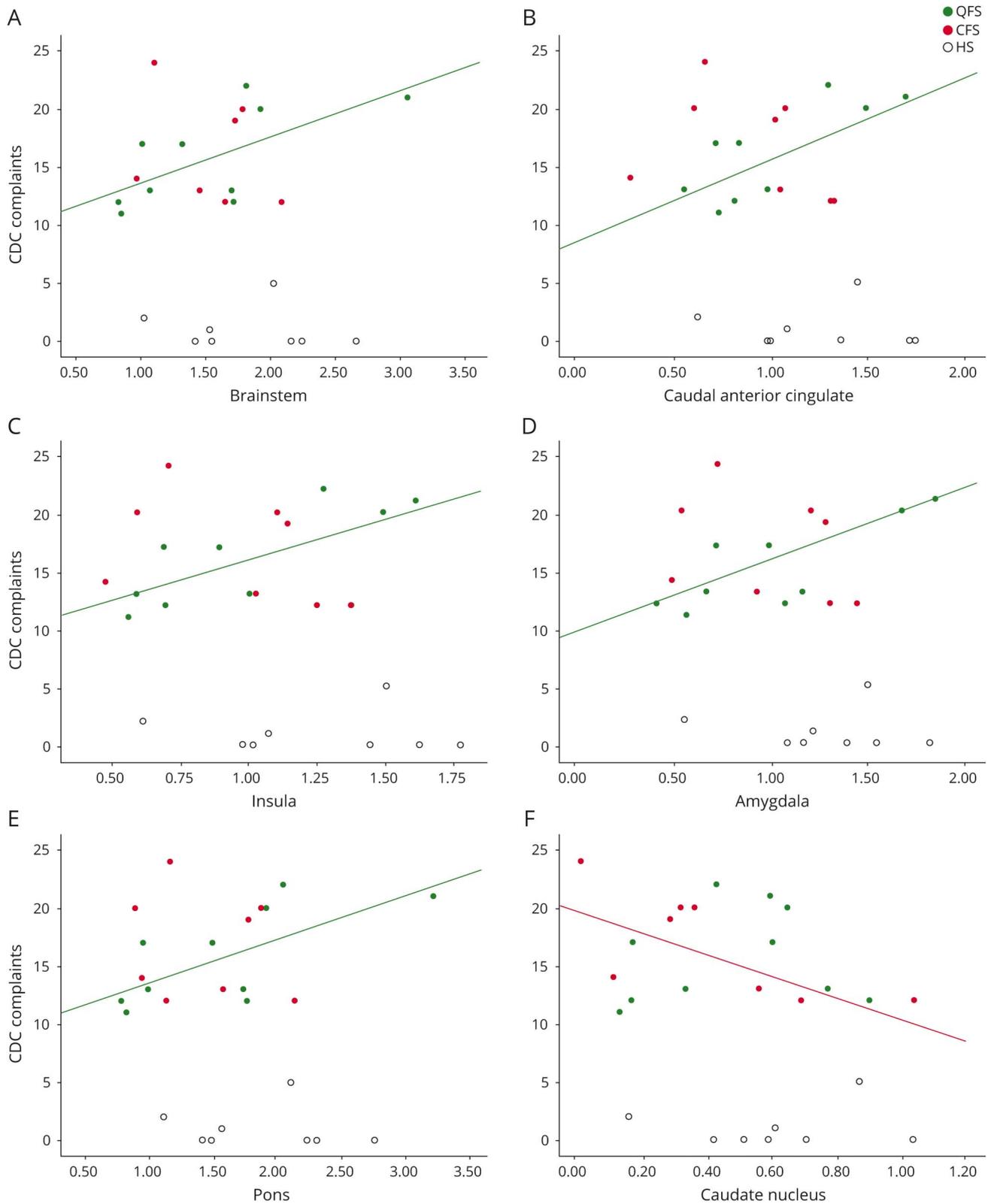


Scatter plots showing significant correlations between BP_{ND} of [¹¹C]-PK11195 and fatigue severity, measured by the CIS questionnaire, subscale on fatigue severity, for patients with QFS; in the orbitofrontal cortex (A, $R = 0.70^*$), middle frontal gyrus (B, $R = 0.83^{**}$), inferior frontal gyrus (C, $R = 0.78^{**}$), superior frontal gyrus (D, $R = 0.64^*$), primary motor cortex (E, $R = 0.74^*$), temporal lobe (F, $R = 0.78^{**}$), primary somatosensory cortex (G, $R = 0.83^{**}$), and parietal lobe (H, $R = 0.77^{**}$), and patients with CFS; in the caudate nucleus (I, $R = -0.78^*$). BP_{ND} = nondisplaceable binding potential; CFS = chronic fatigue syndrome; HS = healthy subject; QFS = Q fever fatigue syndrome. Statistical significance was attained if $p < 0.05$.

An inherent problem in CFS research is the presumed heterogeneity of the disorder. We addressed this problem in several ways. As mentioned above, we only enrolled adult women. Second, we used a validated test panel of instruments to assess fatigue and disability. Third, we included a group of patients with postinfectious fatigue related to antecedent Q fever (QFS). As in the latter group, an infectious, and therefore inflammatory, etiology was the precipitating factor, we would have expected this group in particular to exhibit signs of neuroinflammation. However, although BP_{ND} of [¹¹C]-PK11195 was generally higher than in patients with CFS, even in this well-defined group, we could not detect neuroinflammation. For future perspective, we should avoid previous mistakes in CFS research and continue investigating neuroinflammation in chronic fatigue by using strict and uniform in- and exclusion criteria, together with well-defined control groups.^{4,38}

Our study has some limitations. For reasons of homogeneity, we chose to use the [¹¹C]-PK11195 ligand as this was the ligand used by the only neuroinflammation PET imaging study in CFS by Nakatomi et al.⁵ Nowadays, a new generation of more sensitive ligands such as [¹¹C]-PBR28 and [¹⁸F]-DPA-714 are available, and perhaps even preferable, when taking allelic dependence of affinity into account.¹⁶ Using [¹¹C]-PBR28 for example, signs of neuroinflammation have been found in functional somatic syndromes such as fibromyalgia and Gulf War Illness.^{27,39} The former study included mostly women, whereas the latter included mostly men, but with complaints for up to 30 years. Another limitation is the large amount of correlations that were conducted, which increases the risk of a type 1 error (whereas post hoc analyses increase the risk of a type 2 error). A final limitation is the small number of subjects included in both our study and the study by Nakatomi et al. One could argue that a more sensitive new

Figure 3 Correlation Between BP_{ND} of [¹¹C]-PK11195 in Various Brain Regions With CDC Questionnaire, Subscale on Complaints, Scores in Patients With QFS, Patients With CFS, and HSs



Scatter plots showing significant correlations between BP_{ND} of [¹¹C]-PK11195 and fatigue severity, measured by the CDC questionnaire, subscale on complaints, for patients with QFS; the brainstem (A, R = 0.66*), caudal anterior cingulate (B, R = 0.64*), insula (C, R = 0.65*), amygdala (D, R = 0.71*), and pons (E, R = 0.69*), and patients with CFS; in the caudate nucleus (F, R = -0.73*). BP_{ND} = nondisplaceable binding potential; CDC = Centers for Disease Control; CFS = chronic fatigue syndrome; HS = healthy subject; QFS = Q fever fatigue syndrome. Statistical significance was attained if $p < 0.05$.

generation ligand would be better suited when using such small numbers.⁴⁰ Other than imploring a larger study or using more sensitive TSPO ligands, we should keep an eye on current investigations on other targets than TSPO for PET neuroimaging.⁴¹

In contrast to what was previously reported, our well-controlled study shows no significant difference in BP_{ND} of [¹¹C]-PK11195 when comparing both patients with CFS and QFS with HSs. A larger and preferably longitudinal study, including both men and women together with well-matched controls, is needed to confirm whether chronically fatigued patients exhibit neuroinflammation. For this study, we propose using a new generation ligand with kinetic modeling via an arterial input function.

Acknowledgment

The authors thank Professor I.E.C. Sommer for her contributions to the inclusion of subjects in our study.

Study Funding

This study was funded by the Q-support Foundation (UMCN-140928-00). There was no role for the funding body in the design of the study, collection of data, analysis, interpretation of data, or writing of the manuscript.

Disclosure

The authors declare that they have no competing interests or financial relationships relevant to the manuscript. Go to Neurology.org/NN for full disclosures.

Publication History

Received by *Neurology: Neuroimmunology & Neuroinflammation* May 20, 2021. Accepted in final form September 15, 2021.

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Megan Roerink, MD	Department of Internal Medicine, Radboud University Medical Center, Nijmegen, the Netherlands	Major role in the acquisition of data and analysis or interpretation of data
Stephan Keijmel, MD	Radboud Expertise Center for Q fever, Department of Internal Medicine and Radboud Center for Infectious Diseases, Radboud University Medical Center; Department of Internal Medicine, Radboud University Medical Center, Nijmegen, the Netherlands	Major role in the acquisition of data and analysis or interpretation of data

Appendix (continued)

Name	Location	Contribution
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Neurol Neuroimmunol Neuroinflamm 2022;9;

DOI 10.1212/NXI.0000000000001113

This information is current as of November 23, 2021

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