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Posted Date: 29 January 2024

doi: 10.20944/preprints202401.2035.v1

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Article

Quantification of Circulating Cell-Free DNA in Idiopathic Parkinson's Disease Patients

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Abstract: Background: Parkinson's disease (PD) is one of the most common neurodegenerative disorders globally, and leads to an excessive loss of dopaminergic neurons in the substantia nigra of the brain. Circulating cell-free DNA (ccf-DNA) are double-stranded DNA fragments of different sizes and origins that are released into the serum and cerebrospinal fluid (CSF) due to cell death (i.e., necrosis and apoptosis) or are actively released by viable cells via exocytosis and NETosis. **Methods:** Using droplet digital polymerase chain reaction (ddPCR) we comprehensively analyzed and distinguished ccfDNA and ccf mtDNA in the serum and CSF of PD and control patients. **Results:** The quantitative analysis of serum ccf-DNA in PD patients demonstrated significant increase of ccf mtDNA and ccfDNA compared to healthy control patients and significantly higher copy of ccf mtDNA when compared to ccfDNA. Next, the serum ccf mtDNA levels significantly increased in male PD patients compared to healthy male controls. Furthermore, CSF ccf mtDNA in PD patients increased significantly compared to ccfDNA, and ccf mtDNA decreased in PD patients compared to healthy controls. These decreases were not statistically significant, but were in agreement with previous data. Interestingly, ccf mtDNA increased in healthy control patients in both serum and CSF as compared to ccfDNA. **Limitations:** the small sample size of serum and CSF were the main limitations of this study. **Conclusion:** To the best of our knowledge, this is the first comprehensive study on serum and CSF of PD patients using ddPCR and indicating the distribution of the copy number of ccf mtDNA, as well as ccfDNA. If validated, we suggest that ccf mtDNA has greater than ccfDNA potential to lead the development of novel treatments for PD patients.

Keywords: Parkinson's disease; mitochondria; circulating cell free DNA (ccf -DNA); mitochondrial (mtDNA); nuclear (ccfDNA); droplet digital PCR (ddPCR)

1. Introduction

Parkinson's disease (PD) is the second most common neurodegenerative disease globally. It leads to the excessive loss of dopaminergic neurons in the substantia nigra of the brain. Accurate diagnosis of the disease remains challenging, and methods of characterizing the earliest stages of the disease is a focus of ongoing research [1,2]. Like in many neurodegenerative diseases, PD symptoms occur well after pathology begins due to the compensatory potential of the brain. Furthermore, due to the massive death of neuronal cells, it is difficult to treat advanced-stage PD

patients. Therefore, a simple and non-invasive method of early diagnosis could increase the efficiency of the limited treatment options available [1–4].

Circulating cell-free DNA (ccf-DNA) are short, double-stranded DNA fragments present in various body fluids, such as the blood, urine, serum and cerebrospinal fluid (CSF) [5–8]. Depending on the type of DNA released, ccf-DNA has two main sources: nuclear (ccfDNA) and mitochondrial (ccf mtDNA). It is believed that ccf-DNA is released due to cell death (i.e., necrosis or apoptosis) and through active release by viable cells via exocytosis and NETosis [9–12]. For the last decade, ccf-DNA has become a subject of interest for the non-invasive analysis of tumor-derived genetic material. Both ccfDNA and mtDNA have been the focus of qualitative and quantitative investigations. Alterations in these two types of ccf-DNA have also been implicated in various types of cancer [13–16].

Limited reports have primarily focused on the quantification of ccf-DNA levels in CSF of PD patients. Most studies involving ccf DNA have focused on ccf mtDNA [18,19], and post-mortem studies [17]. In these studies, ccf mtDNA has been found in decreased levels among PD patients compared to healthy controls. Lowes (2020a) found that a positive correlation existed between increased levels of CSF ccf mtDNA and various comorbidities such as depression and insomnia, however this was only significant if measured in the absence of treatment [19,20]. Regarding serum, the study by Borsche et al. (2020) investigated the sporadic form of PD and found that patients with biallelic PINK1 and Parkin mutations had elevated levels of ccf mtDNA and IL-6 suggesting increased ccf mtDNA release, and neuroinflammation in these PD patients [21]. There are currently no studies presenting quantification of the serum and CSF ccf mtDNA or ccfDNA of idiopathic PD patients. Such data could be useful for diagnostic purposes and could provide better understanding of the association between serum and CSF ccf-DNA and neurodegeneration.

In this study, we performed quantitative and qualitative studies on ccf-DNA isolated from the serum and CSF of idiopathic PD and healthy control patients using droplet digital PCR (ddPCR). This method allowed for the precise specification of the copy number of ccf mtDNA and ccfDNA. We also correlated the level of ccf-DNA with gender in the studied groups.

2. Results

Our aim was to characterize ccf-DNA in the serum and CSF of PD patients to compare them against healthy control patients. To distinguish differentially derived DNA, we chose target DNA fragments of the mitochondrial *COX3*, a gene encoding mitochondrial cytochrome c oxidase III, and nuclear *KRAS*, an oncogene encoding a protein belonging to the GTPase superfamily. These have previously been used in analogue studies involving different groups of patients [22,23]. Thus, we analyzed ccf-DNA isolated from the serum of 30 PD and 15 healthy control patients and the CSF from 13 PD and five healthy control patients. The PD patient cohort was selected by neurologists from the Department of Neurology University of Medical Sciences Poznan, Poland, from patients diagnosed with PD (**Table 1**).

ddPCR was used here to enable analysis of small amounts of material. Unlike quantitative PCR (qPCR), ddPCR does not require reference gene copy analysis [24,25]. In ddPCR, amplification occurs in small volume (1 nl) droplets. The total number of droplets is then calculated as the number of positive droplets (i.e., those that fluoresce) less than the number of negative droplets (i.e., those with no signal). These are counted in a flow cytometry-like fashion to produce a ratio that is then subjected to Poisson distribution, resulting in the absolute quantification of starting template molecules [26–28]. The copy numbers of ccf mtDNA and ccfDNA were calculated in this manner and underwent statistical analysis using the Mann-Whitney U test.

Methods of quantifying mtDNA copy numbers with ddPCR using purified genomic DNA have been developed previously [22,26,27,29,30]. Here, ccfDNA was purified immediately after blood donation by isolation from the frozen serum and CSF of 0.2 ml samples according to the Qiagen procedure.

Table 1. Data from Parkinson’s disease patients in the study.

Data of tested patients			
Serum			
Control patients (n=15)		Parkinson’s Disease patients (n=30)	
Age			
Mean	39	Mean	62
Median	37	Median	66
Min-Max	19-54	Min-Max	40-75
Gender			
Females	4	Females	12
% of females tested	27%	% of females tested	40%
Males	11	Males	18
% of males tested	73%	% of males tested	60%
Data of tested patients			
CSF			
Control patients (n=5)		Parkinson’s Disease patients (n=13)	
Age			
Mean	52	Mean	57
Median	56	Median	61
Min-Max	35-69	Min-Max	37-75
Gender			
Females	3	Females	6
% of females tested	60	% of females tested	46%
Males	2	Males	7
% of males tested	40	% of males tested	54%

2.1. Quantification of serum ccf-DNA in PD patients

2.1.1. Serum ccf mtDNA and ccfDNA in PD patients versus control patients

We analysed and compared the ccf DNA isolated from PD and healthy control patients, as shown in Figure 1. In the serum from the control patients, ccf mtDNA levels were significantly higher than the ccfDNA levels (*p-value* = 0.000052***; Figure 1a). In PD patients, ccf mtDNA levels were higher than ccfDNA levels (*p-value*: 1.691123e-17; Figure 1b). The ccf mtDNA levels of PD patients were significantly increased compared to the control patients (*p-value* = 0.001729**; Figure 1c). Interestingly, ccfDNA levels among PD patients were also significantly higher than those in the control patients (*p-value* = 0.0048926*; Figure 1d). All presented analyses were statistically significant.

Because the age of the PD and control healthy patients differs (median 37 and 66, respectively) we have performed ANOVA analysis with age as a covariant to assess the relationship of the Parkinson's disease presence and ccf mtDNA. In this case, multiple linear regression was performed. The overall association was statistically significant ($F = 5.542$, $p = 0.007309$), with the model explaining 17.11% of the variance. However, the observed increase of the ccf mtDNA copy number among PD patients was not statistically significant at $\alpha = 0.05$ ($t = 1.908$, $p = 0.0633$; Figure 1e). Obtained results revealed the age is not associated with ccf mtDNA levels in PD versus control healthy patients ($t = 0.362$, $p = 0.719$; Figure 1f). Analogous analysis was conducted for ccfDNA levels, although there was no significant overall association ($F = 2.173$, $p = 0.1264$) (Supplementary File 2. Table 1 and 1).

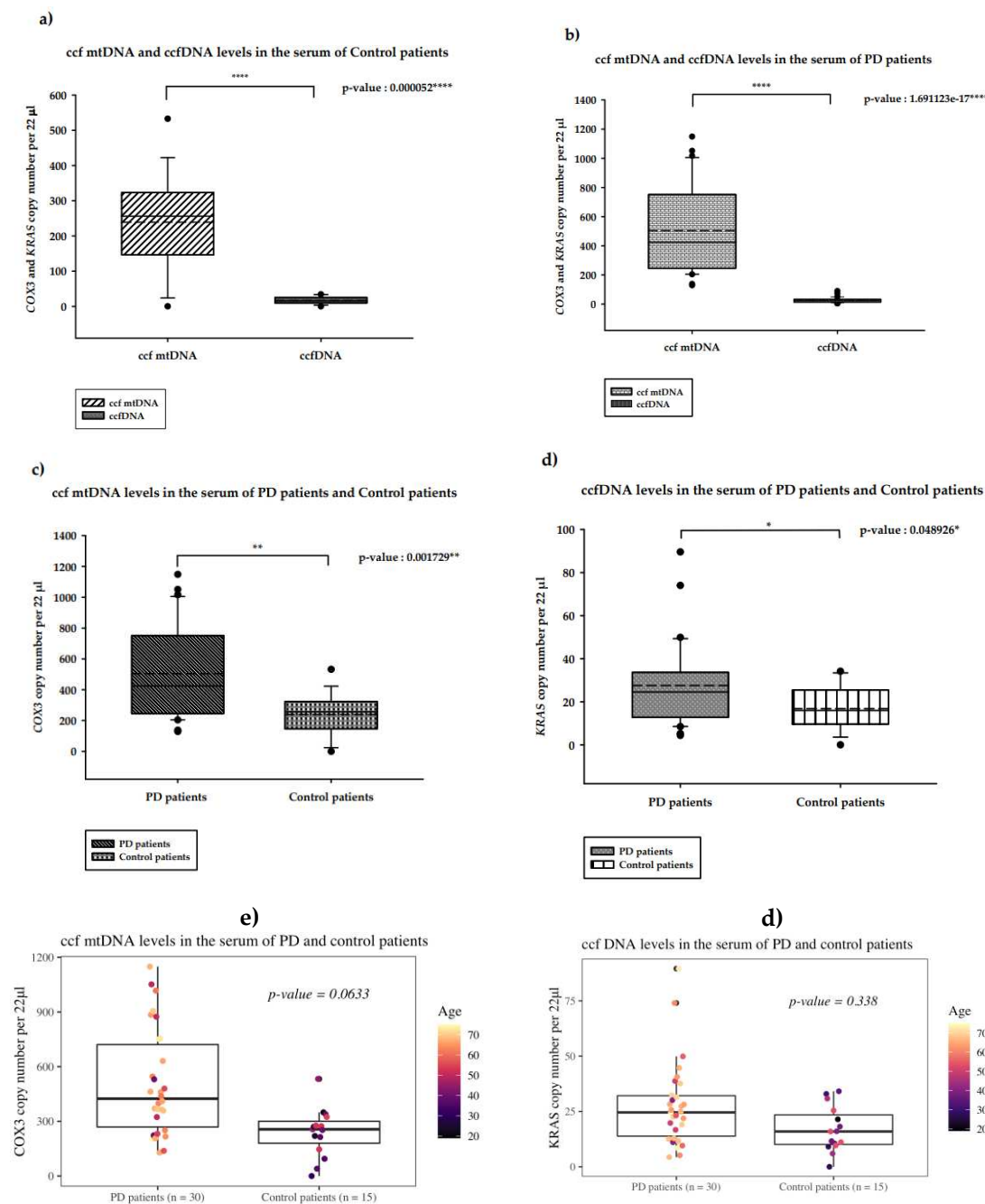


Figure 1. Results of the ddPCR analysis on serum ccf DNA from PD and healthy controls. a. ccf mtDNA and ccfDNA copy numbers in the serum of healthy controls; b. ccf mtDNA and ccfDNA copy numbers in the serum of PD patients; c. ccf mtDNA copy numbers in the serum of PD and healthy controls; d. ccfDNA copy numbers in the serum of PD and healthy controls. e. ccf mtDNA copy numbers in the serum of PD and healthy controls used by ANOVA with age as a covariant; f. ccfDNA

copy numbers in the serum of PD and healthy controls used by ANOVA with age as a covariant. Boxplots represent medians (solid lines) and means (dashed lines) with min and max values. Mann-Whitney U tests were performed for comparison. A probability of ≤ 0.05 was considered statistically significant; * $p \leq 0.05$, ** $p \leq 0.01$, *** $p \leq 0.001$, **** $p \leq 0.0001$.

2.2. Quantification of CSF ccf-DNA in PD patients

2.2.1. CSF ccf mtDNA and ccfDNA in PD patients versus healthy controls

We analyzed and compared the ccf DNA isolated from the CSF of PD and healthy controls (Figure 2). In the CSF of healthy controls, ccf mtDNA levels were significantly higher than ccfDNA levels (p -value = 0.007937**, Figure 2a). The same pattern was observed in PD patients, with the difference also being significant (p -value = 0.000006***, Figure 2b). In contrast to the results obtained for the serum levels, CSF ccf mtDNA levels among PD patients were lower than in the control patients (p -value = 0.094538; Figure 2c). ccfDNA levels were higher in PD patients compared to control patients; but, this difference was not statistically significant (p -value = 0.288749; Figure 2d).

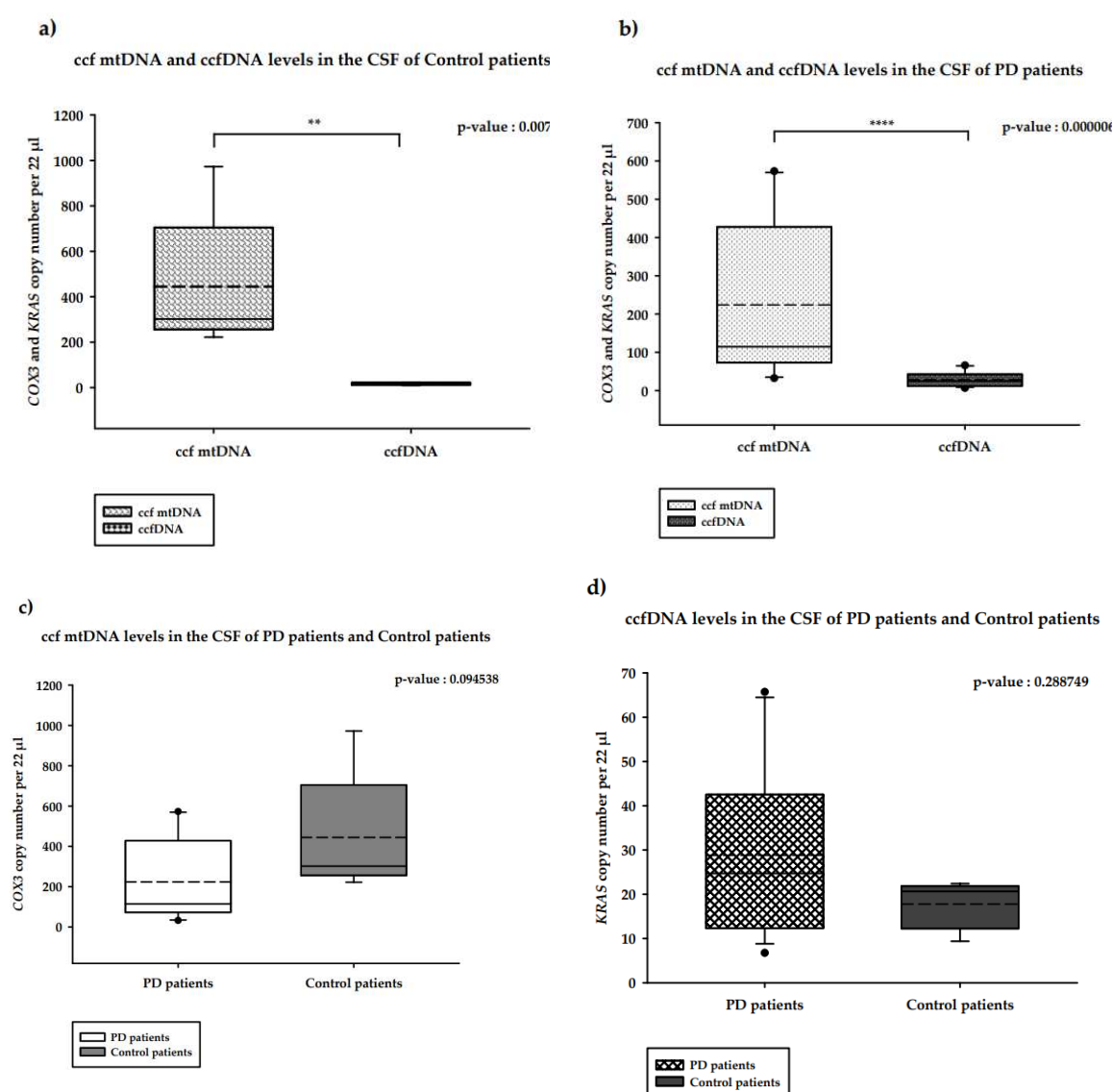


Figure 2. Results of the ddPCR analysis on the CSF ccfDNA of PD patients and healthy controls.

a. ccf mtDNA and ccfDNA copy numbers in the CSF of healthy controls; **b.** ccf mtDNA and ccfDNA copy numbers in the CSF of PD patients; **c.** ccf mtDNA copy numbers in the CSF of PD patients and healthy controls; **d.** ccfDNA copy number in the CSF of PD patients and healthy controls. Boxplots represent medians (solid lines) and means (dashed lines) with min and max values. Mann-Whitney

U tests were performed for comparison. A probability of ≤ 0.05 was considered statistically significant; * $p \leq 0.05$, ** $p \leq 0.01$, *** $p \leq 0.001$, **** $p \leq 0.0001$.

2.3. Distribution of the copy number of serum ccf-DNA and gender in PD patients

2.3.1. Distribution of the copy number of serum ccf mtDNA and gender in PD versus healthy controls

The correlation between serum ccf mtDNA and gender is presented in Figure 3. The median serum ccf mtDNA copy number among healthy males ($n = 11$) and females ($n = 4$) were 219.73 and 336.43 copies/22 μ l, respectively. This difference was statistically significant, with increased copy numbers observed among the healthy females (Mann-Whitney U test, p -value = 0.026374*; Figure 3a). The median serum ccf mtDNA copy number among female PD patients ($n = 12$) and male PD patients ($n = 18$) were 402 and 435.47 copies/22 μ l, respectively, although this difference was not statistically significant (Mann-Whitney U test, p -value = 0.573247; Figure 3b). The median serum ccf mtDNA copy numbers among female PD patients ($n = 12$) and healthy females ($n = 4$) were 402 and 336.43 copies/22 μ l, respectively and this difference was also not statistically significant (Mann-Whitney U test, p -value = 0.598901; Figure 3c). The median serum ccf mtDNA copy numbers among male PD patients ($n = 18$) and healthy males ($n = 11$) were 435.47 and 219.73 copies/22 μ l, respectively. This difference was statistically significant, with double the copy numbers seen in male PD patients (Mann-Whitney U test, p -value = 0.000857***; Figure 3d).

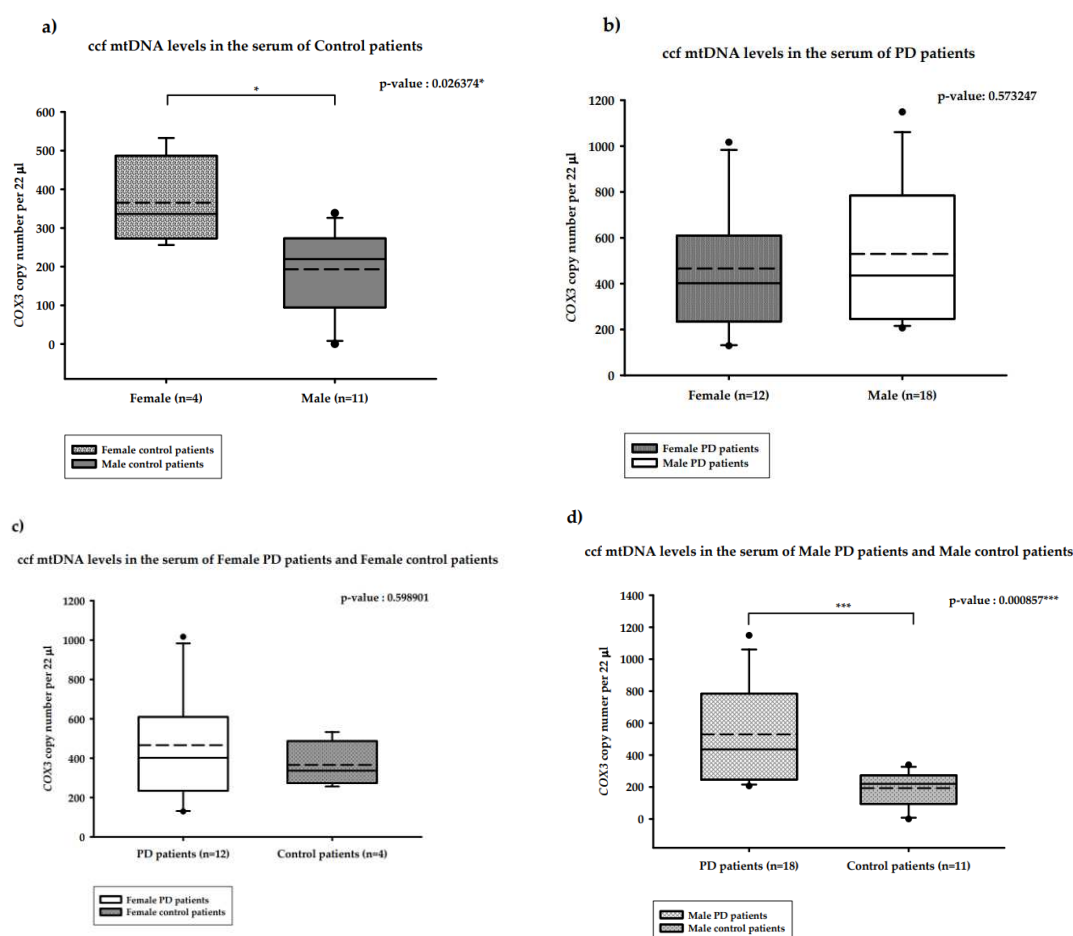


Figure 3. Distribution of copy numbers of serum ccf mtDNA and gender in PD patients versus healthy controls. a. ccf mtDNA copy numbers in the serum of female and male controls; b. ccf mtDNA copy numbers in female and male PD patients; c. ccf mtDNA copy numbers in female PD patients and female controls; d. ccf mtDNA copy numbers in male PD patients and male controls. Boxplots represent medians (solid lines) and means (dashed lines) with min and max values. Mann-Whitney

U tests were performed for comparison. A probability of ≤ 0.05 was considered statistically significant; * $p \leq 0.05$, ** $p \leq 0.01$, *** $p \leq 0.001$, **** $p \leq 0.0001$.

2.3.2. Distribution of the copy number of serum ccfDNA and gender in PD versus healthy controls

The correlations between serum ccfDNA and gender are presented in Figure 4. The median serum ccfDNA copy numbers in the healthy male ($n = 11$) and healthy female ($n = 4$) groups were 15.94 and 16.28 copies/22 μ l, respectively. This difference was not statistically significant (Mann-Whitney U test, p -value = 1.000000; Figure 4a). The median serum ccfDNA copy numbers among female PD patients ($n = 12$) and male PD patients ($n = 18$) were 24.95 and 24.02 copies/22 μ l, respectively and was also not statistically significant difference (Mann-Whitney U test, p -value = 0.631508; Figure 4b). The median serum ccfDNA copy numbers in female PD patients ($n = 12$) and healthy females ($n = 4$) were 24.95 and 16.28 copies/22 μ l, respectively; this difference was not statistically significant (Mann-Whitney U test, p -value = 0.446154; Figure 4c). The median serum ccfDNA copy numbers in male PD patients ($n = 18$) and healthy males ($n = 11$) were 24.02 and 15.94 copies/22 μ l, respectively; this difference was statistically significant (Mann-Whitney U test, p -value = 0.055121; Figure 4d).

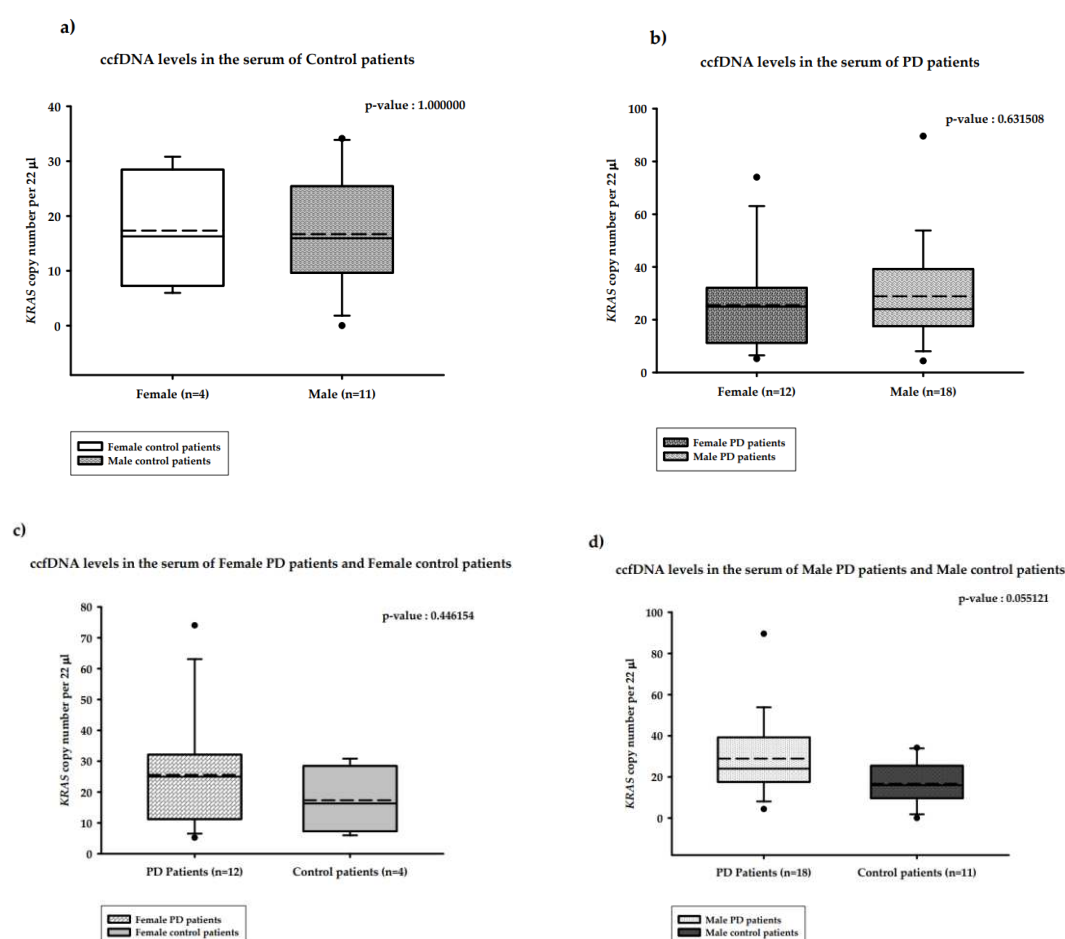


Figure 4. Distribution of copy numbers of serum ccfDNA and gender of PD patients versus healthy controls. a. ccfDNA copy numbers in the serum of female and male controls; b. ccfDNA copy numbers in female and male PD patients; c. ccfDNA copy numbers in the serum of female PD patients and female controls; d. ccfDNA copy number in male PD Patients and male controls. Boxplots represent medians (solid line) and means (dashed line) with min to max values. Mann-Whitney U tests were performed for comparison. A probability of ≤ 0.05 was considered statistically significant; * $p \leq 0.05$, ** $p \leq 0.01$, *** $p \leq 0.001$, **** $p \leq 0.0001$.

3. Discussion

To the best of our knowledge, this is the first comprehensive study of the serum and CSF of idiopathic PD patients performed using the precise technique of ddPCR to enable presentation of the distribution of the copy number of both ccf mtDNA and ccfDNA. Our data obtained for the serum revealed a significant increase in the copy number of ccf mtDNA versus healthy control patients. For both serum and CSF higher copy number of the ccf mtDNA compared to ccfDNA was observed (**Table 2**). In previous studies, it has been shown that mitochondria can control inflammation through the production of reactive oxygen species (ROS) and the release of mitochondrial components, including mitochondrial DNA (mtDNA), into the extracellular matrix, where they act as danger signals [31]. We found that ccf mtDNA dominated in the serum of PD patients; however, this does not reflect observation regarding CSF, where ccf mtDNA is reduced. Interestingly, unlike nuclear DNA, mtDNA contains unmethylated CpG sequences (a pattern common to bacterial DNA), which act as damage associated molecular patterns (DAMP) [32,33]. It is believed that this ccf mtDNA allows nonself recognition, further contributing to immune system activation [32] and stimulating an innate immune response through variety receptors expressed in neurons [34–36] and inflammatory response described for (PRKN-PINK) PD patients [37]. According to this study, increased ccf mtDNA in the serum of patients with a genetic form of PD (i.e., biallelic PD mutation PRKN/PINK1) is correlated with elevated IL-6 levels. Thus, we speculate that increased level of the ccf mtDNA detected in the serum of PD patients could be due to massive cell death, resulting in the release of ccf- DNA as a potential signal molecule for the cytokines responsible for the described above immunogenic response. Importantly, the level of nuclear derived ccfDNA was also increased however this difference among PD and healthy controls were not as significant as for of ccf mtDNA.

Regarding CSF, obtained results were in agreement with previously reported data [18–20] that revealed a reduced copy number of CSF ccf mtDNA in PD patients versus healthy controls. However, we found that, similarly to the serum, ccf mtDNA of PD patients significantly dominated when compared to ccfDNA, which we showed for the first time (**Table 2**) to reflect possible important function of ccf mtDNA under the ccfDNA in idiopathic PD pathogenesis. This cause of the reduction of ccf mtDNA in CSF is poorly understood. PD is linked to high levels of neuronal cell death within the substantia nigra; therefore, an increased level of CSF ccf mtDNA would be expected as a consequence of mitophagy process [8]. However, this pattern could also be caused by an overall decrease in the mitochondria pool of nerve cells, which is observed in the early stages of neurodegeneration in PD [18]. Studies that have reported decreased neuronal mtDNA copy numbers in neurodegenerative disorders have indicated that it is associated with a reduction in cell energy [31]. Reduced CSF ccf mtDNA levels in nerve cells have also been reported in other neurodegenerative diseases, such as Huntington's disease and Alzheimer's disease [8]. However, in Alzheimer's disease, the observation was not confirmed by the other studies [29]. In contrast, increased CSF ccf mtDNA levels have been reported in patients with multiple sclerosis. Studies have concluded that this occurs as a direct consequence of the increased activation of inflammatory cells, which release mtDNA into the CSF [39].

In the case of nuclear derived ccfDNA, we observed its decreased level in both the serum and CSF of PD patients. The mechanism driving this is unclear; however, it may be a consequence of the faster degradation of genomic DNA. Notably, ccfDNA has been reported to be more prone to nuclease degradation compared to ccf mtDNA [8]; this reflects the unknown mechanism in the CSF and serum of PD patients, which needs further study. We assumed that ccf mtDNA dominates over ccfDNA in the serum and CSF of PD patients, which may suggest some unknown mechanism in PD biogenesis in which mitochondria are engaged.

Importantly, our study has some limitations that need to be taken into account. Firstly, there was a rather small number of participants for serum (30 PD and 15 controls), and even more so for the CSF (13 PD and 5 controls). It is also known that obtaining a larger sample size for serum samples of the PD patients, the one of the most common neurodegenerative disease, should not be very difficult; however, despite the relatively small serum sample size, many observed results based on the precise

ddPCR method are statistically significant and novel (**Table 2**), and we believe that the presented data will inspire further follow-up studies.

Future study of the ccf DNA of the blood serum should also be considered. It is known that ccf-DNA that circulates in the blood originates from different tissues. Thus, this DNA has the same genome and cannot be associated with a specific source tissue through DNA sequencing [40–45]. In our study, we have shown that using target mitochondrial and nuclear genes to identify the origin of ccf-DNA in blood serum in PD patients by ddPCR enabled to quantify of ccf-DNA, which could be used in the future for optimisation PD therapy.

It is known that Parkinson's disease has a genetic origin (i.e., mutations in the PARK genes encoding alpha-Synuclein, DJ-1, PINK, LRRK2, etc.) in 5–10% of patients, causing so-called early onset PD and that most of these cases are idiopathic and associated with aging. In our study, an idiopathic group of PD patients was chosen as a group reflecting various features associated with aging, and consequently, with neuroinflammation [46]. For this reason, we decided to take healthy blood donors to compare our PD results with undoubtedly homogeneous controls. It should also be noted that a previous study revealed that ccf mtDNA of healthy individuals declined with the age of healthy individuals [47]. In line with this finding, we performed ANOVA analysis with age as a covariates. The obtained results revealed the age is not associated with ccf mtDNA and ccfDNA levels in PD patients versus control healthy patients although there was no significant overall association (Figure 1e-f), (Supplementary File 2. Table 1-2).

Ccf-DNA levels vary over time depending on increased physical activity, and the existence of various medical conditions, such as infectious diseases [48]. Thus, we also determined the copy number of ccfDNA in the serum of PD patients according to gender, although this was not performed for CSF due to a smaller sample size. The obtained results revealed that the level of ccf mtDNA was significantly increased in male PD patients as compared to healthy male control patients. However, serum ccf mtDNA levels were significantly higher among the female healthy patients compared to the male healthy patients. This could suggest that the presence of hormones influences the course of this neurodegenerative disease. This observation is particularly interesting, especially when considering data obtained by Patel and Kompoliti, which showed lower prevalence of PD among females [49]. Ultimately, we can only speculate on the influence of hormonal or menopausal status of our female participants. Undoubtedly, the impact of the level of female hormones on PD biogenesis requires further analysis.

It is worth highlighting certain results obtained for the serum and CSF of healthy controls. For both cases, the level of ccf mtDNA was statistically higher than ccfDNA, and differences were more significant in the serum. This may suggest that ccf mtDNA could be engaged in unknown regulatory processes, such as cell signaling, which again illustrates the need for further study of the role of mitochondria in ccf mtDNA biogenesis.

The results of our quantitative ccf-DNA analysis of the serum and CSF of PD patients may also be correlated with some other neurodegenerative aspects, such as, for example, protein aggregation. Indeed, for ccf mtDNA, it has previously been found that there is no significant correlation between CSF ccf mtDNA and α synuclein [18].

In summary, the results obtained here recommend serum for the studies on PD patients, which is a safer candidate for use in non-invasive diagnostic studies than CSF. Despite this and the other discussed limitations, if validated, we suggest ccf mtDNA to be used as a target for the optimization of PD therapies.

In the future, determining the precise mechanism of ccf DNA release in PD patients would add further value to ccf mtDNA's potential to serve as a reliable, non-invasive approach to monitoring responses to medical treatment during therapy.

Table 2. Comparative table of previously reported results and this study regarding ccf mtDNA and ccfDNA in human blood serum and CSF. EOPD (Early-onset Parkinson's disease); *results are not statistically significant; novel results obtained in this study are indicated in bold.

source of sample	gender male/female	type of PD	Ccf mtDNA/ healthy control	Ccf mtDNA/ ccfDNA	Ccf DNA/ healthy control	number of PD/number of control	method of analysis	reference
serum	-	mut+/+ PD PRKN/PINK1	increase	-	-	17/57	ddPCR	[21]
		mut+/- PD PRKN/PINK1	increase			17/55		
serum	-	idiopathic	increase	increase	increase	30/15	ddPCR	this study
serum	male	idiopathic	increase	increase*	no difference*	18/11	ddPCR	this study
serum	female	idiopathic	no difference*	no difference*	no difference*	12/4	ddPCR	this study
CSF	-	idiopathic	reduced	-	-	56/10	qPCR	[18]
CSF	-	EOPD	reduced	-	-	176/87	qPCR	[19]
CSF	-	idiopathic	reduced*	increase	increase*	13/5	ddPCR	this study

4. Materials and Methods

4.1. Collection of CSF and serum samples

The study was performed with the cooperation of the Department of Neurology, Division of Neurochemistry and Neuropathology and Department of Neurosurgery and Neurotraumatology, Poznan University of Medical Sciences. It was approved by the local institutional review board (206/17). Written consent forms were collected from all patients who were involved in this study.

4.2. Study cohort and sampling procedure

The blood samples used in this study were collected from 2016 to 2020 at the University of Medical Sciences. In total, 20 healthy and 34 PD patients were included in this study. All blood and CSF samples were taken before doctoral interventions. Patients were analysed for the assessment if they were suitable for deep brain stimulation treatment and on average five years after initial diagnosis of Parkinson's disease.

Patient data, including age and length of treatment, were obtained from the patients' pathological reports. The blood samples were processed, and the serum was isolated immediately from 1 ml of blood according to the procedure described by Al Amir Dache et al. [23]. Serum samples were immediately frozen in liquid nitrogen and stored at -80°C. The study cohort (n = 30) was divided into two groups: the first group consisted of PD patients (n = 34), while the second consisted of healthy control patients (n = 20). The diagnosis of PD was established based on the criteria of the United Kingdom PD Brain Bank. The healthy controls were recruited from the Regional Blood Centre and Blood Therapy in Poznan.

Regarding the CSF samples, we collected 13 from patients diagnosed with PD and 5 from healthy controls. All samples were obtained by lumbosacral puncture and frozen and stored immediately at -20°C.

4.3. ccfDNA isolation

Ccf-DNA was isolated from both the serum and CSF samples according to the Qiagen procedure using an isolation kit specific for isolating this DNA from plasma or serum (QIAamp MinElute ccfDNA Mini Kit). For each isolation procedure, 0.2 ml of serum or CSF were taken. The isolated ccfDNA was eluted in 20 µl and stored in 5 µl aliquots in standard Eppendorf tubes at -80°C. DNA was quantified using a nanodrop spectrophotometer (Thermo Scientific).

4.4. Quantification of serum ccf mtDNA and ccfDNA levels

DdPCR was used to quantify ccf mtDNA and ccf nuDNA levels. The ddPCR assay was performed using the QX200 ddPCR EvaGreen Supermix (Bio-Rad) [42,43]. Primers coupled with the EvaGreen dye were used to quantify fragments, with amplicons targeting the wild-type sequences of specific genes: the KRAS nuclear gene and the mitochondrial COX gene, MT-CO3 (Table 2). Quantification of the short and long amplicons provided an estimation of the concentrations of the total ccfDNA and ccf mtDNA, respectively (Supplementary Data File 1) [23]. The mitochondrial gene encodes the mitochondrial COX3, while the nucleus gene is an oncogene which encodes a protein belonging to the GTPase superfamily (KRAS). Primer sequences for the COX3 gene were based on previous analogue studies [23]. Primer sequences for the KRAS gene were designed using the PRIMER 3 program (<https://bioinfo.ut.ee/primer3-0.4.0/>). The specificity of primers was assessed using the Primer-BLAST program (<https://www.ncbi.nlm.nih.gov/tools/primer-blast/>), as well as in-silico PCR (<https://genome.ucsc.edu/cgi-bin/hgPcr>). DdPCR samples were prepared following the manufacturer’s instructions. The PCR was performed on the QuantStudio 6 albo 7 Flex (Applied Biosystems) [44]. For the ddPCR analysis of the serum samples, all samples were diluted 15 times. For the ddPCR analysis of the CSF samples, isolated ccf DNA was diluted five times. After preparing the correct dilutions, 1µl of ccf DNA was added to 21 µl of the mixture, which consisted of forward and reverse primers and EvaGreen dye. The droplets were generated using the Droplet Generator (Bio-Rad). To obtain information regarding the number of copies of the COX3 and KRAS genes, the QX200 Droplet Reader (Bio-Rad) was used.

Prior to the ddPCR analysis, a qPCR analysis was performed to confirm the presence of the COX3 and KRAS genes and, therefore, the presence of ccf mtDNA and ccfDNA in the CSF and serum samples. We used the Applied Biosystems QuantStudio 7 Flex System to perform the qPCR. The qPCR analysis allowed us to select the correct 5-fold dilution of ccf DNA for the ddPCR analysis. For qPCR analysis, all samples were diluted four times, and SYBR green dye was used.

Table 3. Primer sequences used for ddPCR and qPCR.

Gene	Sequence of the Forward primer (5'→3')	Sequence of the Reverse primer (5'→3')
COX3	GACCCACCAATCACATGC	TGAGAGGGCCCTGTTAG
KRAS	CCTTGGGTTTCAAGTTATATG	CCCTGACATACTCCCAAGGA

4.5. Statistical analysis

The results obtained from the ddPCR analysis were calculated using the Poisson equation. The step-by-step calculations are described below. The statistical analysis involved the Mann-Whitney U test (<https://astatsa.com/WilcoxonTest/>). The threshold for statistical significance was set to $p < 0.05$.

Multiple linear regression analyses were conducted in R using the lm package to assess the association of ccf mtDNA or ccfDNA levels with Parkinson's disease (PD), with age serving as a covariate. The regression models were formulated as $\text{ccf mtDNA} \sim \text{PD} + \text{age}$. The difference in the levels of ccf and mt ccfDNA in PD and control patients was assessed with paired t test, using t. test function in R.

4.6. Data presentation

The results collected from the ddPCR analysis were presented in the form of boxplot graphs generated using SigmaPlot 13.0 (Systat Software). For each generated boxplot graph, the median (solid line) and mean (dashed line) number of copies of ccf-DNA were calculated and presented. Due to the small number of PD patients and control patients tested, no outlier ccf-DNA results were removed during graph generation. Due to the small number of PD patients and control patients tested, no outlier copy number values (round, black dots above or below box plots) were removed during graph generation.

Supplementary Materials: The following supporting information can be downloaded at the website of this paper posted on Preprints.org.

Author Contributions: Conceptualization, M.W. and B.S.; methodology, M.W.; A.P.; software, A.P.; N.K.; validation, N.K.; K.P., A.P.; formal analysis, N.K, M.W, K.P.; investigation, K.P, N.K., M.W., resources, J.F-W., K.P-D., B.S, T.M.; data curation, J.F-W., K.P-D., B.S.N.K.,P.K.; writing—original draft preparation, M.W.; writing—review and editing, M.W.; N.K., K.P; visualization, N.K.,P.K.; supervision, B.S.; project administration, M.W.; funding acquisition, M.W., N.K., K.P., B.S. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded This studies was supported by IDUB - Initiative of Excellence - Research University" program conducted by Adam Mickiewicz University in Poznan 001/34/UAM/0012 and, 014/34/UAM/0003 (2020-2023).

Institutional Review Board Statement: The study was conducted in accordance with the Declaration of Helsinki, and approved approved by the local institutional review board (206/17). Written consent forms were collected from all patients who were involved in this study.

Data Availability Statement: No.

Acknowledgments: No.

Conflicts of Interest: The authors declare no conflict of interest.

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