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The iron brain: Post-mortem and in vivo imaging of iron in brain diseases

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Pathological characterization of T_2^* -weighted MRI contrast in the striatum of Huntington's disease patients

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ABSTRACT

Previous MRI studies consistently reported iron accumulation within the striatum of patients with Huntington's disease (HD). However, the pattern and origin of iron accumulation is poorly understood. This study aimed to characterize the histopathological correlates of iron-sensitive ex vivo MRI contrast change in HD brains. To this end, T_2^* -weighted 7T MRI was performed on post-mortem tissue of the striatum of three control subjects and 10 HD patients followed by histological examination. In addition, formalin-fixed paraffin-embedded material of three control subjects and 14 HD patients was selected for only histology to identify the cellular localization of iron using stainings for iron, myelin, microglia and astrocytes. As expected HD striata showed prominent atrophy. Compared to controls, the striatum of HD patients was in general more hypointense on T_2^* -weighted high-field MRI and showed a more intense histopathological staining for iron. In addition, T_2^* -weighted MRI identified large focal hypointensities within the striatum of HD patients. Upon histological examination, these large focal hypointensities frequently co-localized with enlarged perivascular spaces and iron was found within the vessel wall and reactive astrocytes. In conclusion, we show that the striatum of HD patients has a distinctive phenotype on T_2^* -weighted MRI compared to control subjects. On ex vivo MRI, these contrast changes are heavily biased by enlarged perivascular spaces from which it is currently unknown whether this is a fixation artefact or a disease specific observation. Clinically, the observation of iron within reactive astrocytes is of importance for the interpretation and understanding of the potential underlying mechanisms of T_2^* -weighted MRI results in HD patients.

5.1. INTRODUCTION

Huntington's disease (HD) is an autosomal dominant neurodegenerative disorder caused by an expanded cytosine-adenine-guanine (CAG) trinucleotide repeat in the Huntingtin (HTT) gene [1]. The disease-causing HTT alleles show >39 CAG repeats in exon 1, with disease onset being earlier and progression faster the larger the number of CAG repeats is [2]. Clinically, HD is characterized by a variety of motor disturbances (typically chorea and dystonia), cognitive decline and behavioural changes [3]. The histopathological correlate of these signs and symptoms is progressive neuronal loss, especially of the medium spiny neurons of the striatum, followed by atrophy. During disease progression, other regions including the white matter and cortex are also affected, eventually resulting in global brain atrophy [4].

Although HD has a clear monogenetic origin, the pathological cascade that leads to neuronal death is complicated, affecting many cell types and biological processes, and non-invasive methods to accurately monitor disease progressions in vivo remain to be developed. In addition, HD is one of the many neurodegenerative diseases in which increased levels of iron have been reported in the striatum, the brain region most severely affected [5–8]. Although iron is essential for many fundamental biological processes including oxygen transport, DNA synthesis, mitochondrial respiration in general, as well as myelin and neurotransmitter synthesis and metabolism in the brain specifically, due to its ability to catalyze the generation of reaction oxygen species via the Fenton reaction, iron levels have to be tightly controlled to prevent oxidative stress and ultimately neuronal loss [7, 9, 10].

In HD the largest increase in iron is reported in the basal ganglia, and more specifically in the striatum [6, 8, 11–16]. Due to the magnetic properties of iron, its presence affects the homogeneity of the local magnetic field in tissue, and hence contributes to MRI contrast, especially on T_2^* -weighted images [17]. Previous neuroimaging studies have used a range of iron-sensitive MRI techniques to analyze cerebral iron accumulation in HD. These studies consistently reported MRI contrast changes, hypointensities and increased susceptibility values, indicative of iron accumulation in the basal ganglia [6, 8, 11–16]. This effect has already been found in premanifest HD gene-carriers [6, 15], showing positive correlations with calculated disease severity as a function of age and the number of supernumerary CAG repeats, the so-called disease burden score [15, 18] and actual clinical disease severity [6]. These effects were found to be independent from basal ganglia atrophy, suggesting a net increase in total iron in these structures and the potential of iron-sensitive MRI as a biomarker of HD disease progression [12].

Although these studies did show the relevance of iron-sensitive MRI scans in HD, the underlying pathological substrates have never been investigated. Several studies reported the high correlation between MRI and iron levels using quantitative and qualitative techniques in both healthy and diseased brain [19–25]. However, the exact pat-

tern of iron accumulation, the colocalization with specific cells and correlation with MRI in HD remains to be elucidated. To date, only one post-mortem study linked microglial activation with iron accumulation, suggesting that iron accumulation in HD is predominantly associated with neuroinflammation [26].

Therefore, the objective of this study was to gain more insight into the histopathological correlates of the well-known contrast changes on T_2^* -weighted MRI in the striatum of HD. We performed ultra-high field ex vivo MRI and histopathology on post-mortem tissue of the striatum of HD patients to further delineate the pattern of iron accumulation and the colocalization of iron with specific cells known to be associated with HD pathology such as microglia and astrocytes [27].

5.2. METHODS

5.2.1. STUDY DESIGN

Formalin-fixed post-mortem brain material of the striatum (including the caudate nucleus and putamen, see Figure 5.1) of three control subjects and 10 HD patients was included for both MRI and histological examination. In addition, formalin-fixed paraffin-embedded material of three control subjects and 14 HD patients was selected for only histology to investigate the cellular localization of iron. Histological stainings for iron, myelin, microglia and astrocytes were performed, microscopically examined and correlated with the MRI images.

5.2.2. POST-MORTEM BRAIN MATERIAL

Formalin-fixed striatal tissue blocks of six control subjects and 24 HD patients was included (Table 5.1 for characteristics). HD material was selected from the local neuropathology tissue collection of the Leiden University Medical Center (The Netherlands). Brain material from normal controls was selected from the Netherlands Brain Bank (NBB, Netherlands Institute for Neuroscience Amsterdam, The Netherlands) and the Normal Aging Brain Collection (Amsterdam Neuroscience, Amsterdam UMC, Vrije Universiteit Amsterdam, The Netherlands). Anonymity of all subjects was preserved by using a coded system for the tissue samples following the Dutch national ethical guidelines (Code for Proper Secondary Use of Human Tissue, Dutch Federation of Medical Scientific Societies).

5.2.3. POST-MORTEM MRI ACQUISITION

As previous studies showed the potential effect of formalin fixation on absolute iron concentrations and tissue integrity, only material with a maximum fixation duration of four years was selected for post-mortem 7T MRI [29, 30]. No significant differences in

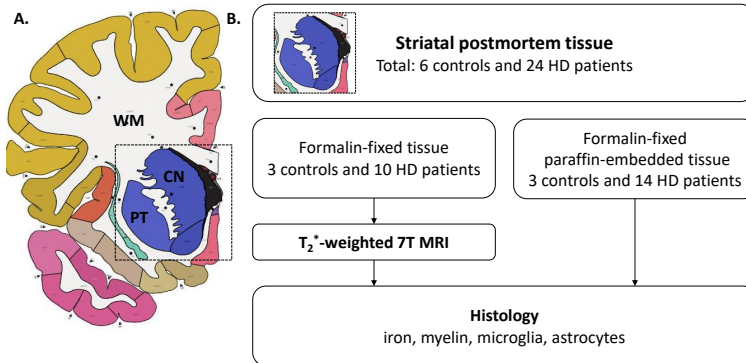


Figure 5.1: **Schematic overview of the study design.** From coronal formalin-fixed brain slaps (A), small tissue blocks including the striatum (level of the caudate nucleus and putamen) were dissected (dashed box in A, upper panel in B). Formalin-fixed tissue of three control subjects and 10 HD patients was included for both MRI and histological examination. Formalin-fixed paraffin-embedded tissue of three control subjects and 14 HD patients was selected for histology only. Histological stainings for iron, myelin, microglia and astrocytes were performed, microscopically examined and correlated with the MRI images. Anatomical atlas modified from the Allen Reference Atlas (human atlas, whole brain, 34 years) [28]. WM = white matter; CN = caudate nucleus; PT = putamen.

	Controls (n=6)	HD (n=24)
Age of death, mean \pm SD (range)	55.8 \pm 5.4 (49-63)	58.5 \pm 9.6 (36-74)
Male/Female	2 / 4	6 / 18
CAG repeat length, mean \pm SD (range)	n.a.	45.4 \pm 4.7 (36-57)
Vonsattel grade, (n)	n.a.	2.8 \pm 0.8 (1-4)
Post-mortem interval in hours, mean \pm SD	8.4 \pm 1.7	11.2 \pm 4.8

Table 5.1: **Case characteristics of controls and HD patients included.**

the duration of formalin fixation were found between tissue from HD patients and controls (3.2 years for HD tissue, 1.5 years for control tissue; $p=0.133$). The selected tissue blocks were put in a 50 ml tube (Greiner Bio-One). Before MRI the relaxation parameters were partially restored by rinsing the tissue with phosphate buffered saline (PBS) and soaking the tissue in PBS for 24 hours [31]. Before scanning, PBS was replaced with an MRI-invisible proton-free fluid (Fomblin LC08, Solvay). Care was taken to avoid the inclusion of air bubbles.

MRI scans were performed on a 7T horizontal bore Bruker MRI system equipped with a 38 mm transmit-receive volume coil and Paravision 6.0.1 imaging software (Bruker Biospin, Ettlingen, Germany). Multiple gradient echo scans with a total scan time of 10.5 hours were acquired. Each brain sample was scanned with the following parameters: repetition time 75 ms, echo times 12.5/23.2/33.9/44.6 ms, flip angle 25° at $100\mu\text{m}$ isotropic resolution with 14 signal averages.

5.2.4. HISTOLOGY

In addition to the tissue blocks scanned with MRI, formalin-fixed paraffin-embedded tissue blocks from three control subjects and 14 HD patients were included for histological examination. In total, tissue from six controls and 24 HD patients was included for histological examination. All tissue blocks were serially cut into $5\mu\text{m}$ -thick sections. Histochemical detection of iron was done following an adapted version of the Meguro protocol [2, 32]. In short, after deparaffinization sections were incubated for 30 minutes in 1% potassium ferrocyanide, washed followed by 60 minutes incubation in methanol with 0.01 M NaN_3 and 0.3% H_2O_2 . Subsequently, sections were washed with 0.1 M phosphate buffer followed by 30 minutes incubation in a solution containing 0.025% 3',3'-diaminobenzidine-tetrahydrochloride (DAB, DakoCytomation) and 0.005% H_2O_2 in 0.1 M phosphate buffer. The reaction was stopped by washing.

Next, for each subject three consecutive sections adjacent to the section stained for iron were used for immunohistochemistry to visualize either myelin, microglia or astrocytes with an anti-proteolipid protein (PLP) monoclonal mouse antibody (1:1000; Serotec), an anti-Iba1 polyclonal rabbit antibody (1:1000; Wako Chemicals USA), or an anti-human GFAP monoclonal mouse antibody (6F2; 1:1000; DakoCytomation, Glostrup, Denmark), respectively. To this end, after deparaffinization all sections were treated with 0.3% H_2O_2 in methanol to block endogenous peroxidase activity. Depending on the primary antibody, an antigen retrieval step followed. To detect Iba-1, sections were boiled in EDTA, pH 8.5, for 15 minutes and cooled down for 1 hour. Sections stained for GFAP were boiled in citrate buffer, pH 6, for 15 minutes and cooled down for 1 hour. Primary antibodies were diluted in 1% bovine serum albumin (BSA) in PBS and incubated overnight at room temperature. Secondary antibodies, for PLP and GFAP: biotinylated rabbit anti-mouse (1:200; DakoCytomation, Glostrup, Denmark) and for

Iba-1: biotinylated swine anti-rabbit (1:400; DakoCytomation, Glostrup, Denmark), were incubated for one hour at room temperature followed by a 30 minutes incubation with avidin-biotin complex (ABC, diluted as recommended by the manufacturer, Vector Labs, CA, USA). Signal enhancement was completed by immersion in DAB solution. The sections were counterstained with Harris Haematoxylin, dehydrated, cleared and mounted with Entellan (Merck, Darmstadt, Germany). The slides were digitized using an automatic bright field microscope (Philips Ultra Fast Scanner, Philips, Netherlands) for microscopic evaluation.

Per subject, the caudate nucleus and putamen were scored by two observers (MB and BK) for the presence of enlarged perivascular spaces (absent or present), microglia morphology (homeostatic, dystrophic, mixed dystrophic or depleted) and presence of reactive astrogliosis (absent or present). Microglia morphology was based on the description by Streit et al [33]. Homeostatic microglia morphology was defined as microglia with thin, highly branched processes. Dystrophic microglia were observed as a range of dystrophic entities from moderate beaded and twisted processes to complete fragmentation of cells. Mixed dystrophic was defined by a combination of dystrophic microglia and microglia with thicker soma and branches (probably activated microglia). Iba-1 depletion of microglia was scored when only a single cell or very few cells could be observed, and when 95% of the tissue was completely devoid of staining. Classic homeostatic astrocyte morphology was characterized by a dense network of finely branched processes. Marked reactive astrogliosis was observable as a dramatic change in morphology, including prominent hypertrophy and altered ramifications. Representative images of the different morphologies are included in Figure 5.5.

5.2.5. STATISTICS

Demographic characteristics and duration of formalin fixation were normally distributed and compared using independent t-test and chi-square tests. All statistical analyses were performed with SPSS version 25.0 for Windows (IBM SPSS statistics, Chicago, IL, USA).

5.3. RESULTS

5.3.1. STRIATAL T_2^* -WEIGHTED MRI CONTRAST IS CHANGED IN HD

Previous in vivo studies showed altered T_2^* -weighted MRI contrast indicative of iron accumulation within the striatum of HD patients compared to controls. We first confirmed these in vivo findings by examining the striatal anatomy and T_2^* -weighted MRI contrast of post-mortem tissue from control subject and HD patients. Gross examination of the striatum of all HD patients showed a varying degree of atrophy compared to controls, also represented by the neuropathological severity as indicated by the Vonsattel grade (Fig.5.2).

As shown in Figure 5.2, on T_2^* -weighted MRI the anatomy of the striatum was well visible, with clear contrast differences between the substructures of the striatum. The myelin-rich white matter and internal capsule were visible as most hypointense, followed by the caudate nucleus and putamen. In some cases, the globus pallidus was included in the tissue block and could be observed as most hypointense, with even lower signal intensities than the white matter, caudate nucleus and putamen.

Additionally, specifically in control subjects, small focal hypointensities were observed within the caudate nucleus and putamen (Fig. 5.2, box). HD patients also showed lowest signal amplitude in the myelin-rich areas of the white matter, however, both the caudate nucleus and putamen showed in addition to the small focal hypointensities also larger focal hypointensities, which were only rarely observed in control subjects. These large focal hypointensities sometimes completely lacked signal (Fig. 5.2, arrow), and sometimes filled with a hyperintense core probably as a result of residual PBS trapped within these spaces (Fig. 5.2, arrow head). In general, the striatum of HD patients was atrophic and due to the large hypointensities more hypointense compared to control subjects, like previously reported for in vivo MRI.

5.3.2. MACROSCOPIC COLOCALIZATION OF IRON, MYELIN, BLOOD VESSELS AND MRI

Histological sections stained for iron and myelin were investigated to establish the origin of the observed MRI contrast changes (Fig. 5.3 and 5.4). Macroscopically, the iron staining showed highest staining intensity in the white matter, internal capsule followed by the caudate and the putamen in both control subjects and HD patients. Comparison of MRI with the iron and myelin staining in control subjects showed that the small focal hypointensities on MRI originated from myelinated fiber bundles traversing the caudate nucleus and putamen. These myelinated fibers were also characterized by high iron concentrations (Fig. 5.3A and 5.4A).

Within HD patients, the observed focal hypointensities were considerably larger than

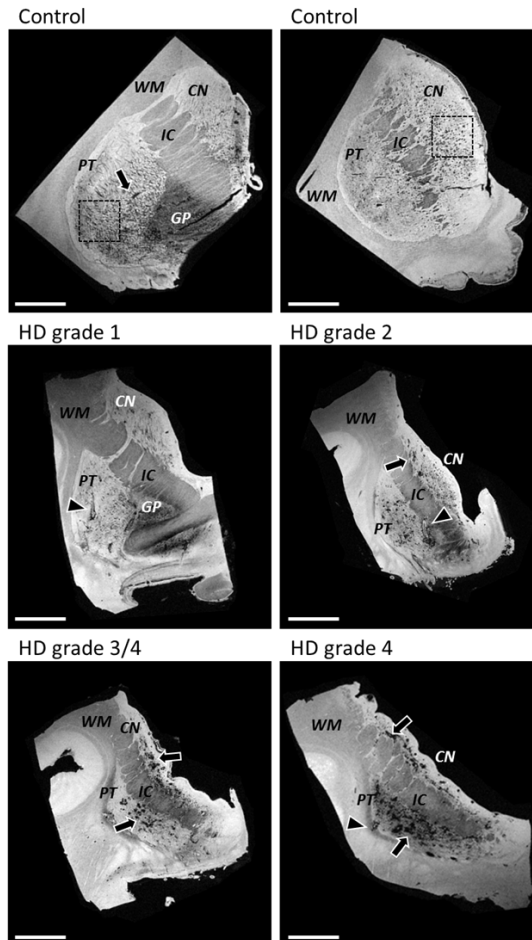


Figure 5.2: **Representative images of T_2^* -weighted MRI of striatal tissue blocks of two controls and four HD cases.** Gross examination of the HD patients showed a varying degree of atrophy which is also represented by the neuropathological severity as indicated by the Vonsattel grade. Small focal hypointensities were found within the caudate nucleus and putamen in control subjects (example within box, but also outside box present). In HD, both the caudate nucleus and putamen showed larger focal hypointensities, which were only rarely observed in control subjects. These larger focal hypointensities sometimes completely lacked any signal (arrow), and were sometimes filled with a hyperintense core probably as a result of residual PBS trapped within these spaces (arrow head). In general, the HD striata were atrophic and more hypointense compared to control striate in line with previously reported *in vivo* findings. CN = caudate nucleus; PT = putamen; IC = internal capsule; GP = globus pallidus; WM = white matter. Scale bar = 1 cm.

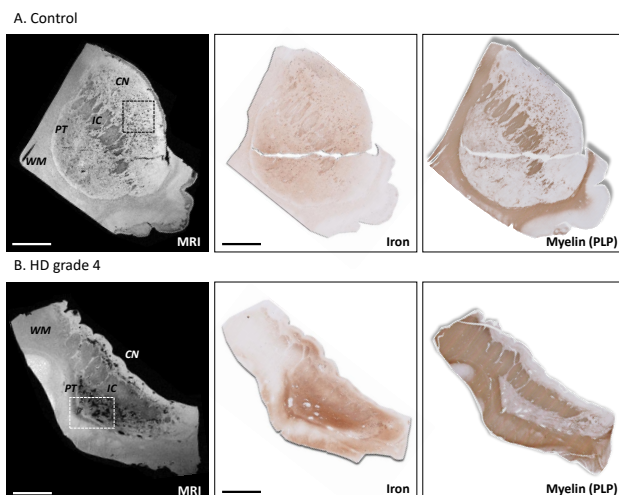


Figure 5.3: **Macroscopic colocalization of MRI and histological staining for iron and immunohistological visualization of myelin.** (A) Control subject showed macroscopic colocalization of the small focal hypointensities on MRI with small regions of increased staining intensity in both the iron and myelin staining. (B) HD patients exhibited larger focal hypointensities within the caudate nucleus and putamen. Histology showed high iron, but low myelin staining intensity within these regions. The hypointensities as observed on MRI frequently co-localized with vessels and enlarged perivascular spaces. See figure 5.4 for detailed information of the squares indicated. CN = caudate nucleus; PT = putamen; IC = internal capsule; WM = white matter. Scale bar = 1 cm.

the small hypointense dots representing fibre bundles. Histological examination revealed that these larger focal hypointensities frequently co-localized with grossly enlarged perivascular spaces of the lenticulostriate arteries. Enlarged perivascular spaces were observed in most HD patients (Table 5.2, 20/24), but only in two control subjects (2/6) as shown by the absence of large focal hyperintensities on MRI. This was confirmed by histology (Fig. 5.2 and 5.3). As can be seen in Figure 5.4B, these vessels were characterized by iron-rich vessel walls and iron-accumulating astrocytes surrounding the vessel.

Additionally, we observed that the border of the caudate nucleus, known as the subventricular zone, was hyperintense on MRI and showed low iron staining intensity on histology (Fig. 5.4C). This was most prominent in HD patients as the subventricular zone was large compared to the severely atrophic caudate nucleus. In control subjects only a very thin hyperintense line could be observed. Microscopic examination showed that this region was characterized by infiltration of dystrophic and activated microglia in both HD patients and control subjects (Fig. 5.4D).

5.3.3. MRI T_2^* -WEIGHTED CONTRAST IS MAINLY DRIVEN BY IRON WITHIN REACTIVE ASTROCYTES IN HD

Macroscopically, the MRI contrast was best explained by the presence of iron, myelin fiber bundles and enlarged perivascular spaces. Stainings for microglia and astrocytes were examined to determine cellular localization of iron.

In most control subjects (4/6), predominantly homeostatic microglia were observed in both the caudate nucleus and putamen (Fig. 5.5A & Table 5.2). Two cases showed microglia with a range of dystrophic entities from moderate beaded and twisted processes to complete fragmentation of cells (Fig. 5.5B). Astrocytes showed in all control subjects a classic homeostatic morphology characterized by a dense network of finely branching processes and were especially found close to vessels. Within the iron staining, we could occasionally identify cells closely resembling microglia and astrocytes. No differences regarding the iron staining could be observed between cases with homeostatic or dystrophic microglia (Fig. 5.5A&B).

Within the HD cohort, microglia showed a larger morphological variety compared to control subjects (Table 5.2). Whereas in all control subjects microglia could be detected, 50% (13/24) of the HD patients showed very few microglia, with 95% of the tissue within both the caudate nucleus and putamen completely devoid of Iba-1 staining, an observation defined as microglia depletion (10/24) (Fig. 5.5C). Some HD patients showed a combination of microglia depletion and presence of dystrophic microglia (3/24). Other cases were characterized predominantly by dystrophic microglia (4/24) (Fig. 5.5D), or by a mixture of both dystrophic microglia and microglia with thicker branches (probably activated microglia), defined as mixed dystrophic mor-

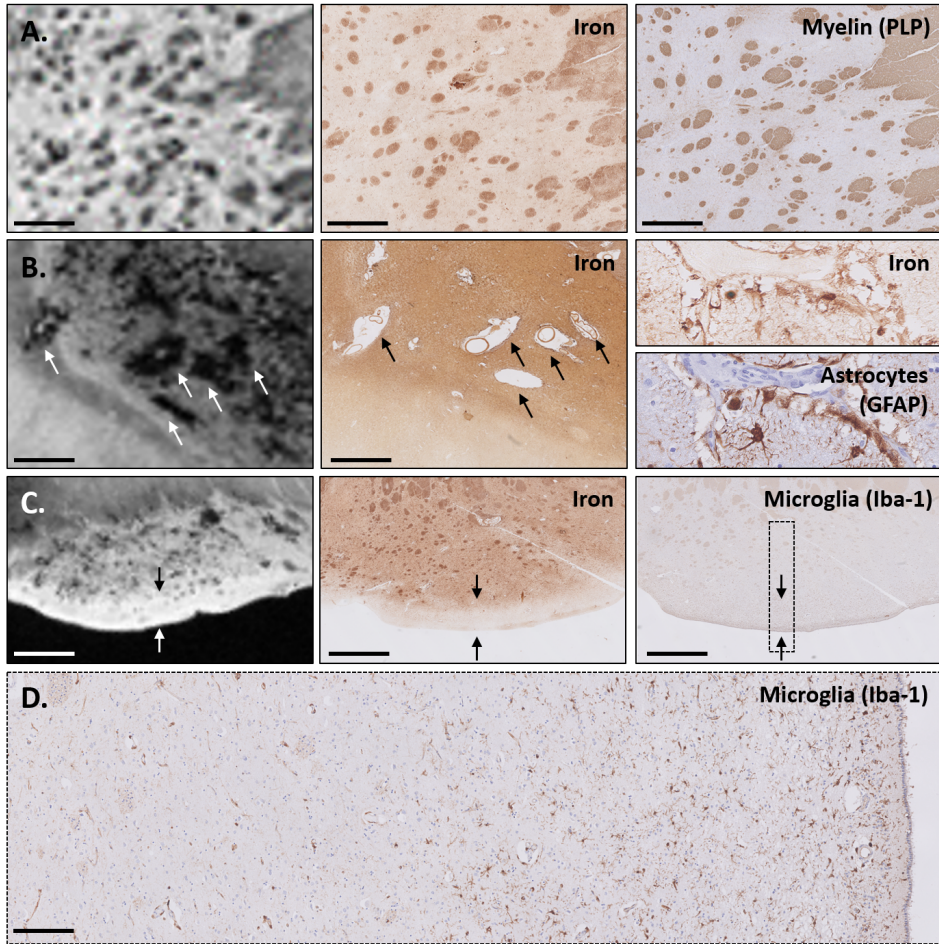


Figure 5.4: **Macroscopic structures affecting the MRI contrast in control subjects and HD patients.** (A) Small focal hypointensities on MRI originated from myelinated fiber bundles traversing the caudate nucleus and putamen, which were also characterized by high iron concentrations. (B) Within HD patients, large hypointense regions were found (arrows), frequently co-localized with vessels and enlarged perivascular spaces. The vessel walls were iron rich and surrounded by iron-accumulating astrocytes. (C) The hyperintense border (arrows), known as the subventricular zone, colocalizes with low iron staining intensity and (D) local infiltration of dystrophic and activated microglia. In control subjects only a very thin hyperintense line could be observed. Scale bar in A = 2 mm; Scale bar in B = 1 mm; Scale bar in C = 200 μm .

phology (6/24) (Fig. 5.5E). Marked reactive astrogliosis was present in all HD patients, observable as a dramatic change in morphology, including prominent hypertrophy and altered ramifications (Fig. 5.5C-E).

The iron staining showed a very similar pattern to the staining for astrocytes. Apart from a general increase in the background intensity, iron was predominantly found in cells morphologically resembling reactive astrocytes. Although the morphological changes of microglia varied across patients, no clear differences in iron staining were found between patients with deplete, dystrophic or a mix of activated and dystrophic microglia, as the iron staining seemed to be dominated by iron accumulation within astrocytes.

The morphological changes in microglia and astrocytes were not correlated with the neuropathological severity as indicated by the Vonsattel grades. Deplete, dystrophic, and activated microglia were found in HD patients with low Vonsattel grades as well as with high Vonsattel grades.

5.3.4. HISTOPATHOLOGICAL CORRELATES OF AREAS WITHOUT MRI CONTRAST CHANGES

To confirm iron accumulation within reactive astrocytes as the main source of MRI contrast changes, other regions showing no abnormalities on MRI were further investigated (Fig. 5.6).

In the HD brain, the neocortex showed high signal intensity upon MRI and low iron staining intensity on histology. Microscopic examination of this region showed the presence of dystrophic microglia. However, in contrast to the caudate nucleus and putamen, astrocytes were characterized by a fine network of branches and thus showed significantly less dramatic changes. Within the iron staining, significantly less background was observed and occasionally we could identify iron-positive cells morphologically resembling microglia and astrocytes.

5.4. DISCUSSION

This study aimed to gain insight into the histopathological correlates of the previously reported T_2^* -weighted MRI contrast changes in the striatum in HD. Ultra-high field *ex vivo* MRI showed that the striatum of HD patients was characterized by large focal hypointense regions that frequently co-localized with enlarged perivascular spaces. Histopathology showed that reactive astrocytes are the predominant source of the general increase of iron within the striatum and hence the observed post-mortem MRI contrast changes.

As already hypothesized by previous *in vivo* MRI studies showing T_2^* -weighted MRI

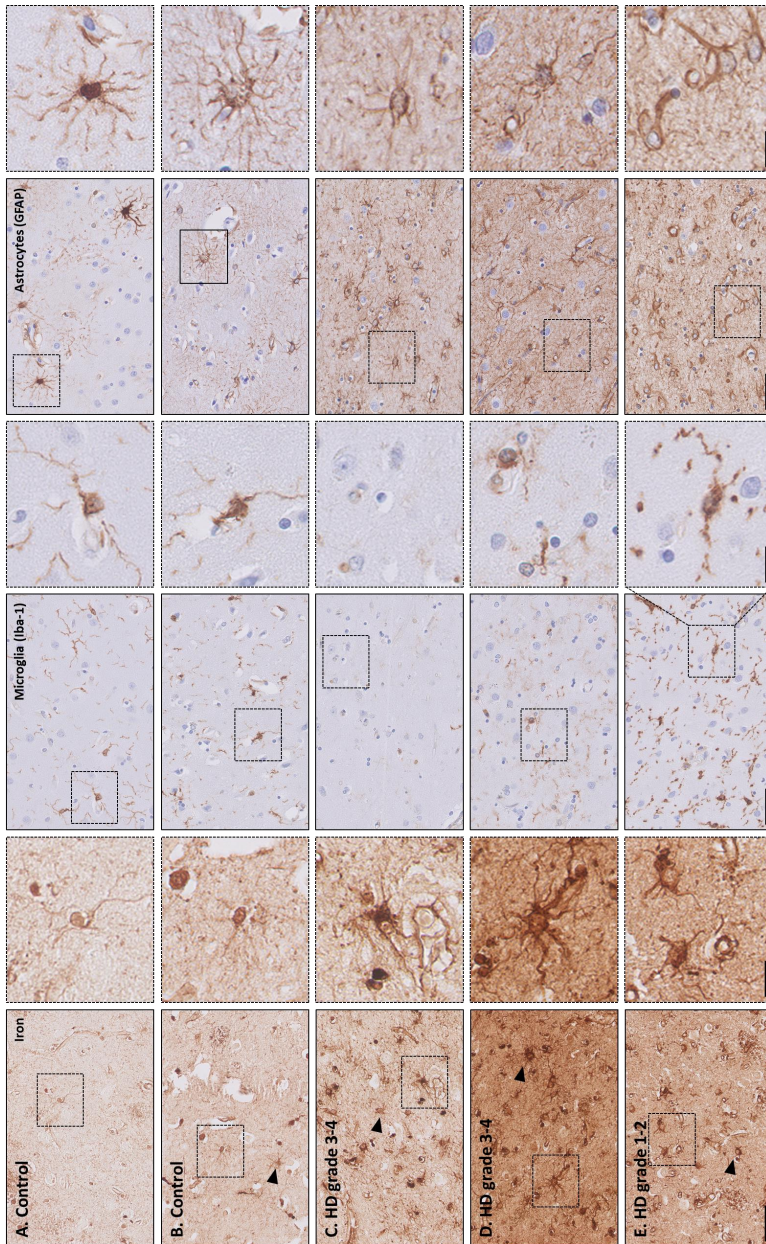
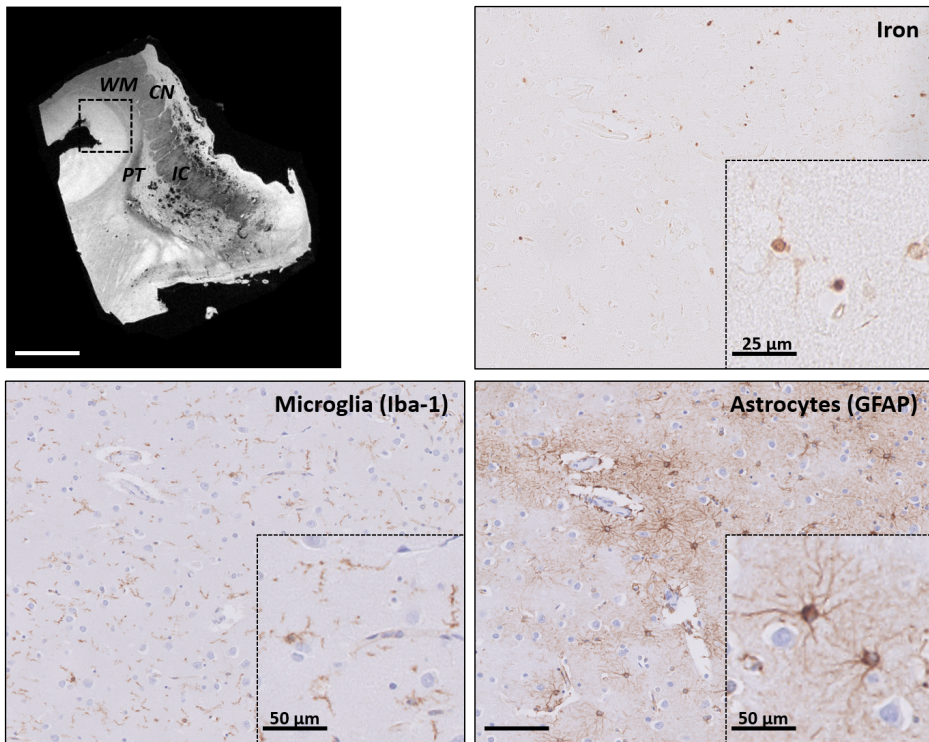


Figure 5.5: **Representative examples of the iron, microglia and astrocyte staining in two control and three HD striata.** Microglia staining showed a variety of morphologies within both control subjects and HD patients, ranging from (A, putamen) homeostatic microglial morphology to (B, putamen & D, putamen) dystrophic morphology or (E, caudate nucleus) mixed dystrophic, defined as a mixture of both dystrophic and activated microglia. (C, caudate nucleus) Microglia depletion was also frequently observed. (C-E) In all HD patients, marked reactive astrogliosis was observable as dramatic change in morphology including hypertrophic and altered ramifications. Interestingly, the iron staining showed a very similar pattern as the staining for astrocytes. No clear differences in iron staining were found between HD patients with deplete, dystrophic or a mix of activated and dystrophic microglia as the iron staining seemed to be dominated by iron accumulation within astrocytes. No correlation between iron accumulation as well as morphological changes in microglia and astrocytes with the Vonsattel grade was found. Scale bar = $50\mu\text{m}$; Scale bar in zoom (indicated with dotted lines) = $25\mu\text{m}$.

	Vonsattel	EPS	Microglia morphology				Astrogliosis
			Homeostatic	Deplete	Dystrophic	Mixed dystrophic	
HD#1	1	C-P			P	C	C-P
HD#2	1-2			C-P			C-P
HD#3	1-2	C-P		C-P			C-P
HD#4	2	C-P		C	P		C-P
HD#5	2	C-P		C-P			C-P
HD#6	2-3				C-P		C-P
HD#7	2-3	C-P				C-P	C-P
HD#8	2-3					C-P	C-P
HD#9	2-3	C-P		C		P	C-P
HD#10	3	C-P		P	C		C-P
HD#11	3	P	P		C		C-P
HD#12	3	C-P		C-P			C-P
HD#13	3	C-P		C-P			C-P
HD#14	3	C-P				C-P	C-P
HD#15	3	C-P		C-P			C-P
HD#16	3	P			C-P		C-P
HD#17	3	C-P			C-P		C-P
HD#18	3-4				C-P		C-P
HD#19	3-4	C-P				C-P	C-P
HD#20	3-4	C-P				C-P	C-P
HD#21	3-4	C-P		C-P			C-P
HD#22	4	C-P		C-P			C-P
HD#23	4	C-P		C-P			C-P
HD#24	4	C-P		C-P			C-P
Control#1	-	P	C-P				Absent
Control#2	-	Absent	C-P				Absent
Control#3	-	Absent	C-P				Absent
Control#4	-	Absent			P		Absent
Control#5	-	P		C	P		Absent
Control#6	-	Absent	C		P		Absent

Table 5.2: **Overview of microscopic results of enlarged perivascular spaces, microglia and astrocyte morphological examination per subject.** EPS = Enlarged perivascular spaces; C = Caudate nucleus; P = Putamen; C-P = caudate nucleus and putamen.



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Figure 5.6: **Region without MRI contrast changes showing normal astrocyte morphology and less intense iron staining.** Within HD, the neocortex showed high signal intensity on MRI and low iron staining intensity on histology. Microscopic examination of this region showed in this specific case the presence of dystrophic microglia. However, astrocytes showed a normal morphology. Within the iron staining significantly less background was observed and occasionally we could identify iron-positive cells morphologically resembling microglia and astrocytes. CN = caudate nucleus; PT = putamen; IC = internal capsule; WM = white matter. Scale bar in MRI = 1 cm; Scale bar in histology = 100 μm ; See zoom for scale bar size.

changes within the striatum of HD patients [6, 13, 15, 16, 21], the spatial co-localizations reported in this study indeed demonstrate a relationship between T_2^* -weighted MRI and increased iron accumulation. Apart from an increase of diffuse iron within the parenchyma, we showed that on ex vivo MRI a significant part of the MRI contrast changes is caused by enlarged perivascular spaces causing large focal hypointensities. Vessels within the striatum were positive for iron, which corresponds to previous reports on both healthy ageing as well as on neurodegenerative diseases [34, 35]. It is suggested that the high metabolic rate of the basal ganglia makes this structure more prone to the accumulation of minerals as calcium and iron [34]. Although being clearly a factor for MRI contrast changes ex vivo, evidence for the presence of enlarged perivascular spaces from in vivo HD studies is currently, apart from one study [36], lacking. In contrast, several studies reported increased brain vessel density in both mice and patients with HD with specifically an increased proportion of small compared to medium-sized blood vessels within the putamen [37–39]. At this moment, we cannot exclude the possibility that the observed enlarged perivascular spaces are a post-mortem artifact, e.g., pronounced parenchymal shrinkage of these structures upon fixation, due to a loss of tissue integrity as a result of neurodegeneration.

Next to these macroscopic structures affecting the MRI contrast, we investigated the cellular localization of iron within microglia and astrocytes, as both are known to be involved in HD pathogenesis [27]. Positron emission tomography (PET) suggests that microglial activation is present in HD patients and correlates with disease severity and striatal neuronal loss [40]. Several studies also showed that microglial activation is an early event in HD pathogenesis as microglial activation is already observed in gene-carriers prior onset of symptoms [41, 42]. In contrast, reactive astrocytes have been observed only after neurodegeneration has become evident [4, 43]. Importantly, the increased number of reactive astrocytes is spatially correlated with the gradient of neurodegeneration within the striatum [43].

We report here that 50% of the HD patients showed microglia depletion rather than dystrophy or activation, and we found no correlation of microglial abnormalities with Vonsattel grades. As microglia within the subventricular zone, internal capsule and cortex were observed within the same brain section of these cases, we do not attribute this observation to a staining artefact. In addition, these findings were confirmed using the antibody TMEM119, a microglial marker that discriminates resident microglia from blood-derived macrophages (not shown) [44]. However, as reported recently by Lier et al [45], a subgroup of microglia exist that exhibit a localized loss of Iba-1 in obese subjects, which was in part linked with obesity and hepatic dysfunction. These Iba-1 negative microglia were also negative for ferritin, but remained immuno-positive for markers expressed primarily by microglial cells (GPX1) [46], and intraparenchymal microglia (P2ry12) [47]. Whether the Iba-1-negative microglia in our study belong to the

same subgroup as described by the Lier needs further investigation. Nevertheless, Lier et al [45] also reported that the Iba-1 negative microglia also stained negative for ferritin, corresponding to the observations in the current study that the microglia did not significantly contribute to the iron staining as this was dominated by activated astrocytes. Alternatively, based on the descriptions by Streit et al [33], we hypothesize that microglia depletion might be a consequence of prolonged microglia activation, with activated microglia becoming dystrophic and undergoing cytorrhesis. Eventually, only scattered fragments and cell nuclei are left which can be observed as microglia depletion in our study.

We hypothesize that initially in HD, microglia likely play an important role in sequestering iron, but due to prolonged activation, microglia become dystrophic and degenerate. Following microglia activation and neurodegeneration, astrocytes are activated. Upon activation their ability to sequester iron increases. Finally, after microglial degeneration in end stage HD, activated astrocytes take over the role of microglia as the predominant iron-sequestering glial cells. This hypothesis is based on previous studies reporting that microglia are the first responders to iron accumulation within the brain parenchyma and are known to rapidly and effectively sequester excess of iron [33] resulting in an iron-positive dystrophic microglial phenotype. This phenomenon has been observed in several neurodegenerative diseases, including HD [26, 48]. Astrocytes are known to play a crucial role in iron handling as well by controlling iron uptake through the BBB and redistribution of iron to neuronal cells [49]. The competence of astrocytes to sequester iron is further increased upon activation in a neuroinflammatory and neurodegenerative environment [50, 51].

A limitation of our study is that the HD material included patients with relatively high Vonsattel grades. Whereas previous studies did report a correlation between microglia activation, reactive astrocytes and disease severity [40, 43], we were unable to reproduce this link. The availability of tissue from patients with very low disease severity is limited. Consequently, it also remains unclear whether the iron distribution in microglia and astrocytes is different in earlier stages of the disease. Translating our findings to pre-manifest HD patients remains challenging as these patients are expected to have microglia activation, iron accumulation but fewer or no reactive astrocytes. Correlating in vivo T_2^* -weighted MRI with biofluid markers for microglia and astrocytes might help interpret the MRI findings.

In conclusion, we show that compared to control tissues the HD striatum has a distinctive phenotype on T_2^* -weighted MRI. On ex vivo MRI, the contrast changes are heavily biased by enlarged perivascular spaces, from which it is currently unknown whether these are fixation artefacts or a disease-specific observation. Microscopically, iron was predominantly found within reactive astrocytes. Clinically, these results are important for the interpretation and understanding of potential underlying mechanisms of T_2^* -weighted MRI results in HD patients. However, the exact sequence of iron accumula-

tion within astrocytes and microglia during disease progression, and the implication of glial iron accumulation for disease progression needs to be further investigated.

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