Brain Structure and Function in Patients after Metal-on-Metal Hip Resurfacing


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ABSTRACT

BACKGROUND AND PURPOSE: Hip prostheses that use a metal-on-metal articulation expose the brain to elevated metal concentrations that, in acute excess due to prosthesis malfunction, is associated with neurologic damage, including visual and hearing loss and motor deficits. Here, we examined whether chronic exposure to lower elevated metal levels, typical of well-functioning prostheses, also affects brain structure and function.

MATERIALS AND METHODS: We compared brain volumes, metal deposition, and gray matter attenuation by MR imaging and clinical neurologic function in patients 8 years after receiving a metal-on-metal hip resurfacing versus a matched group of patients with the same duration exposure to a conventional hip prosthesis.

RESULTS: Twenty-nine patients (25 men; mean, age 59 ± 7 years) after metal-on-metal hip resurfacing and 29 patients (25 men; 59 ± 8 years) after total hip arthroplasty were compared. Whole blood cobalt and chromium concentrations were 5–10 times higher in the metal-on-metal hip resurfacing group (P < .0001). Occipital cortex gray matter attenuation tended to be lower (P < .005 uncorrected, P > .05 corrected), and the optic chiasm area tended to be lower (mean difference, −2.7 mm²; P = .076) in the metal-on-metal hip resurfacing group. Subgroup analyses in 34 patients (17 per group), after exclusion of primary ocular pathology, showed the same trend in gray matter attenuation in the occipital cortex and basal ganglia and a smaller optic chiasm in the metal-on-metal hip resurfacing group (mean difference, −3.9 mm²; P = .048). No other structural or functional differences were found between the groups.

CONCLUSIONS: Chronic exposure to metal-on-metal hip resurfacing is associated with subtle structural change in the visual pathways and the basal ganglia in asymptomatic patients.

ABBREVIATIONS: Co = cobalt; Cr = chromium; MoMHR = metal-on-metal hip resurfacing; PRIME = partially refocused interleaved multiple-echo sequences; THA = total hip arthroplasty; UPDRS = Unified Parkinson’s Disease Rating Scale

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From the Departments of Human Metabolism (M.J.C., J.R.P., J.M.W.) and Cardiovascular Science (N.H., M.N.P.), University of Sheffield, The Medical School, Sheffield, United Kingdom; Department of Neurology (M.H.), Sheffield Teaching Hospitals National Health Service Foundation Trust, Royal Hallamshire Hospital, Sheffield, United Kingdom; and Department of Orthopaedics (J.M.W.), Sheffield Teaching Hospitals National Health Service Foundation Trust, Northern General Hospital, Sheffield, United Kingdom.
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Co and Cr are toxic to neuronal cells in rats, causing brain cell apoptosis mediated by the α subunit of hypoxia inducible factor-1 pathway. Metal toxicity presenting with neurologic deficit is reported in association with malfunctioning hip prostheses and excessive circulating Co and Cr. These neurologic deficits include visual and hearing loss, peripheral sensory and motor defects, tremor, and cognitive decline.

In the United States, >500,000 patients have received hip replacements by using a metal-on-metal bearing, and most have well-functioning devices. Well-functioning prostheses also release metal species into the systemic circulation, albeit at much lower concentrations than those associated with malfunctioning prostheses. Steady-state median blood Co and Cr concentrations during 10 years in patients with well-functioning devices are typ-
ically between 1.5 and 2.5 µg/L, and are 5 to 10-fold higher than normal physiologic concentrations. The effect on the brain of prolonged exposure to these moderately elevated concentrations of circulating metal for many years is unclear.

We examined whether prolonged exposure to circulating metal after metal-on-metal hip resurfacing (MoMHR) is associated with differences in brain structure and function assessed by MR imaging and clinical neurologic assessment in a group of clinically healthy patients after MoMHR versus an age-, sex-, diagnosis-, and time since surgery–matched group of patients who received a conventional total hip arthroplasty (THA) that did not use a metal-on-metal bearing to determine whether this choice of prosthesis has implications for brain health. We, therefore, chose demographically similar patients with conventional THA for the control group, rather than use background population controls, to minimize potential confounding due to brain structural change associated with underlying diagnosis or surgical treatment per se.

**MATERIALS AND METHODS**

**Subjects**

We undertook this single-center cross-sectional study at a teaching hospital in the United Kingdom. Patients were recruited as part of a study examining the systemic effects of metal exposure after MoMHR (n = 35 patients in each group). The study was approved by a National Research Ethics Service Research Ethics Committee, and all patients provided written informed consent before participation. All patients underwent clinical neurologic examination, including assessment of extrapyramidal tract function by using the Unified Parkinson’s Disease Rating Scale (UPDRS), Part 2 (activities of daily living) and Part 3 (motor). The UPDRS is a standardized scale used to assess the severity of Parkinson disease symptoms and has previously been applied to the quantitation of movement disorders associated with metal neurotoxicity. Blood and plasma Co and Cr levels were measured by inductively coupled plasma–mass spectroscopy, as described previously. Blood and plasma Co and Cr levels were measured by inductively coupled plasma–mass spectroscopy, as described previously.

**MR Imaging Acquisition**

MR images were obtained by using a 3T system (Intera; Phillips Healthcare, Best, Netherlands) with an 8-channel sensitivity encoding head coil. The imaging protocol comprised structural, functional, and partially refocused interleaved multiple-echo (PRIME) sequences. 3D T1 magnetization-prepared rapid acquisition of gradient echo images were acquired for volumetric analysis (TR = 10.5 ms, TE = 4.8 ms, flip angle = 8°, 187 sagittal sections with 0.8-mm thickness, 256 × 205 mm FOV, acquired voxel size = 0.8 × 0.8 × 0.8 mm, scan time = 5 minutes 30 seconds). MR images were acquired using the Brain Extraction Tool. A standardized scan used to assess the severity of Parkinson disease symptoms and has previously been applied to the quantitation of movement disorders associated with metal neurotoxicity. Blood and plasma Co and Cr levels were measured by inductively coupled plasma–mass spectroscopy, as described previously.

The PRIME acquisition sequence was used to simultaneously measure quantitative relaxation rates, R2 and R2*, and hence R2*, which is related to magnetic inhomogeneity caused by metal deposition. The sequence comprised 5 spin-echoes with 9 gradient recalled-echoes per spin-echo, leading to a total of 45 images per section (relaxation time = 3000 ms; spin-echo times = 30, 60, 90, 120, and 150 ms; gradient-echo times at 3-ms increments for 9 echoes spaced symmetrically around the spin-echo; recon matrix = 512 × 512; flip angle = 90°; acquired voxel size = 1.44 × 1.8 × 2.5 mm; coronal sections with a section thickness of 2.5 mm; scanning time, 5 minutes 0 seconds). All analyses were made by using the fMRI of the Brain (FMRIB) Software Library (FSL, Version 4.1.1; http://www. fmrib.ox.ac.uk/fsl), apart from metal deposition relaxation time measurements made using the PRIME sequence and optic chiasm area made using in-house software.

**Data Analysis**

Brain volume and subcortical nuclear volumes were measured by using the FSL FMRIB Integrated Registration and Segmentation Tool (FIRST) and automated segmentation and bias field correction tool (FAST) after deletion of nonbrain tissue from images by using the Brain Extraction Tool. FAST was applied to estimate total brain gray and white matter, and FIRST was used to measure brain stem volumes and the volumes of 7 subcortical regions, including the amygdala, caudate nucleus, putamen, thalamus, pallidum, hippocampus, and nucleus accumbens. FSLstats (http://www2. warwick.ac.uk/fac/sci/statistics/staff/academic-research/nichols/ scripts/fsl/fslstats.sh) was used to output numeric data. All measured volumes were corrected for intracranial volume.

Metal deposition in the brain was assessed by using a relative-quantitative method with in-house Matlab software (Version 7.9; MathWorks, Natick, Massachusetts). We have previously used this method to measure brain iron concentration. Here we applied it to Co and Cr quantitation because both Co and Cr in their stable states (Co II and Cr III) are paramagnetic. Briefly, the method involves processing the images by using a nonlinear least-squares fitting routine to calculate values for R2*. The PRIME sequence provides a complete set of simultaneously acquired and registered T2 and T2* contrast-weighted images for each section, allowing calculation of the T2 and T2* relaxation times. T2 was calculated from the peak signal of the 5 acquired spin-echoes, and T2* was calculated from the peak signal from the 5 echoes, including the first spin-echo and subsequent 4 gradient echoes. R2* was calculated from the formula R2* = (1/T2* - 1/T2) by using the difference in the fits to the spin-echo and gradient-echo exponential decay curves from selected regions of interest. This measure indicates the additional inhomogeneity generated by the metal susceptibility difference from normal tissue and minimizes effects due to changes in the T2 relaxation time from other potential sources.

Quantitative measures of gray matter attenuation were made by using voxel-based morphometry (FSL-VBM, Version 1.1; http://fsl.fmrib.ox.ac.uk/fsl/fslvbm/index.html). Structural images had the brain extracted by using the Brain Extraction Tool. The gray matter was then segmented and registered to the Montreal Neurological Institute 152 standard space by using nonlinear registration. The resulting images were averaged to create a study-specific template to which the native gray matter images were nonlinearly registered and corrected for local expansion or contraction. The modulated gray matter images were then smoothed with an isotropic Gaussian kernel with a of 3 mm. Differences in gray matter attenuation were calculated by threshold free-cluster enhancement by using permutation-based nonparametric testing (5000 permutations) by using the Ran-
null agreement between 2 observers (linear regression
center by using a random sample of 20 subjects showed excel-
tions and were quantitated by using methods described previ-
gions outlining the optic chiasm were added to coronal sec-
s (Philips Healthcare) region-of-interest software. Manual re-
frselview (http://fsl.fmrib.ox.ac.uk/fsl/fslview/).
Image outputs were viewed by using
paired
All between-group analyses were completed by using the un-
Statistical Analysis
randomize tool within FSL (http://fsl.fmrib.ox.ac.uk/fsl/fslwiki/
FSLVBM/UserGuide). Image outputs were viewed by using

FIG 1. Voxel-based morphometric analysis of differences in brain
gray matter attenuation between the MoMHR and THA groups be-
fore exclusion of patients with primary ocular pathology. Compara-
tive areas of gray matter loss in the MoMHR group are highlighted
by using the color bar across the significance value range P < .05 to
P < .001 (uncorrected).

domize tool within FSL (http://fsl.fmrib.ox.ac.uk/fsl/fslwiki/
FSLVBM/UserGuide). Image outputs were viewed by using fsview (http://fsl.fmrib.ox.ac.uk/fsl/fsview/).
The optic chiasm area was measured by using the ViewForum
(Philips Healthcare) region-of-interest software. Manual re-
regions outlining the optic chiasm were added to coronal sec-
tions and were quantitated by using methods described previ-
ously.27 Interobserver repeatability of this method in our
center by using a random sample of 20 subjects showed excel-
ent agreement between 2 observers (linear regression r² =
0.97, P < .0001).

Statistical Analysis
All between-group analyses were completed by using the un-
paired t test or Mann-Whitney U test when appropriate (Statisti-
cal Package for the Social Sciences, Version 20; IBM UK, Woking,
United Kingdom). The results were considered significant at P <
.05. Voxel-based morphometry analyses were completed by using
the unpaired t test by using the Randomize software of FSL within

the framework of the General Linear
Model. Because this was an explor-
atory study, results were considered
significant at P < .05 corrected for the
number of comparisons, as described
previously.28-30

RESULTS
Twenty-nine of the 35 recruited patients with MoMHR and 29 of the 35 recruited patients with conventional THA under-
went brain MR imaging and neurologic assessment and are included in these
analyses. A key industry in our local population is metal grinding, and 5 subjects with a history of metal intraocular foreign bodies
that precluded brain MR imaging were excluded from the study, 3
subjects were unable to complete the scanning protocol due to
claustrophobia, and MR imaging data were lost in 4 subjects due
to an archiving failure.
The patient groups were well-matched for age at surgery, sex,
and time since surgery (Table 1, P > .05, all comparisons). Clinical
neurologic examination demonstrated no abnormal clinical
findings, with the exception of a benign essential tremor in 1 male
subject with MoMHR and reduced facial expression, bradykine-
sia, and mild spasticity consistent with a mild extrapyramidal dis-
order in 1 man in the conventional THA group. UPDRS scores for
dysfunction in Part 2 (activities of daily living) and Part 3 (motor
symptoms) were similar between the 2 groups (P > .05, both
comparisons). Median visual acuity at 6 meters in the MoMHR
group was 12 (interquartile range, 6.8–51) in the left eye and 12
(interquartile range, 6.8–51) in the right eye; and in the THA
group, it was 9 (interquartile range, 9–33) in the left eye and 12
(interquartile range, 9–33) in the right eye (P > .05, both com-
parisons). Median blood and plasma Co and Cr levels were 5- to
50-fold higher in patients with MoMHR versus those with THA
(P < .0001, all comparisons, Table 1).

Brain gray matter attenuation measured by voxel-based mor-
phometry was lower in the occipital cortex of the patients with
MoMHR versus those with THA, corresponding to the primary
visual cortex (Fig 1; P < .005, uncorrected; P > .05, corrected). In
view of the location of this difference in gray matter attenuation,
patients were invited to return for clinical ophthalmologic ex-
amination to exclude primary eye pathology as a confounding vari-
able for this analysis. Forty-eight male patients returned for this
examination. Fourteen of these subjects were identified as having
a primary ocular pathology that might contribute to visual cortical
dysfunction or optic atrophy, leaving 34 subjects (17 in each
group) for the subgroup analysis. Gray matter attenuation re-
mained reduced in the occipital region (Fig 2, top panel; P < .005,
uncorrected; P > .05, corrected) but was also reduced in the right
and left putamen and the left head of the caudate nucleus (Fig 2,
bottom panel; P < .005, uncorrected; all regions, P > .05, cor-
rected). The optic chiasm area, measured by using ViewForum, was
36.7 ± 4.7 mm² and 39.5 ± 6.7 mm² in the patients with MoMHR
and THA, respectively (mean difference = –2.7 mm²; 95% CI, –5.7–
0.2 mm²; P = .076). After we removed patients with primary ocular
pathologies and those who did not return for ophthalmic assessment,
the optic chiasm area was 36.3 ± 4.7 mm² in the MoMHR group and 40.3 ± 6.3 mm² in the THA group (n = 17 per group; unpaired mean difference, −3.9 mm²; 95% CI, −7.84 to −0.04; P = .048).

Total brain volume and white and gray matter volumes were similar in both patient groups (as measured by FAST, FSL), as were subcortical nuclei, basal ganglia, and brain stem volumes computed by using FIRST, FSL (Table 2, values corrected for intracranial volume; P > .05, all comparisons). R2 relaxation times in the caudate nucleus, subthalamic nucleus, pallidum, putamen, and white matter were also similar between groups; this finding indicates that any possible difference in the level of metal deposition between the subjects with MoMHR and those with THA was below the level of detection available by using the PRIME sequence (Fig 3; P > .05, all comparisons).

**DISCUSSION**

In this study, we found that healthy asymptomatic patients with confirmed chronic exposure to elevated circulating concentrations of Co and Cr after MoMHR had differences in brain structure compared with a group of age-, sex-, and time since surgery–matched patients with the same underlying disease, but who had received a conventional hip prosthesis and who had circulating metal concentrations that are similar to normal physiologic levels. Patients who had received MoMHR tended toward lower gray matter attenuation in the occipital cortex and basal ganglia and had a smaller optic chiasm area than those patients who had received a conventional prosthesis, after exclusion of patients with primary ocular pathologies from the analysis. We found no evidence of association between MoMHR exposure and changes in brain structural volumes or with metal accumulation within the subcortical nuclei or basal ganglia. We also found no differences in neurologic function, measured by clinical examination and UPDRS, and no differences in visual function, measured by ophthalmologic assessment, between the groups.

These structural trends, though small, are consistent with known toxicologic effects of Co and Cr on the visual system observed in patients with acute exposure to high metal concentrations associated with prosthesis malfunction.8–10 Our data suggest that moderately elevated circulating metal concentrations for 8 years after MoMHR are associated with imaging features in keeping with possible cell loss in the visual system. A chronic insult to the anterior visual pathways has been previously reported in humans to cause voxel-based changes in the primary visual cortex in the context of glaucoma.31 The rational underpinning of these findings is

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**FIG 2.** Voxel-based morphometric analysis of differences in brain gray matter attenuation between the MoMHR and THA groups, after exclusion of patients with primary ocular pathology. The top panel highlights occipital cortex differences between groups; the bottom panel highlights differences in basal ganglia between groups. Comparative areas of gray matter loss in the MoMHR group are highlighted by using the color bar across the significance value range P < .05 to P < .001 (uncorrected).

**FIG 3.** Quantitation of brain metal deposition by PRIME R2 values (second⁻¹). Analyses are MoMHR versus THA by Wilcoxon test. P > .05 for all comparisons.
that such visual cortex changes are produced by transneuronal degeneration in patients with optic nerve damage. Visual cortex abnormalities on voxel-based morphometry consistent with cortical atrophy have also been found in patients with macular degeneration. A recent animal study by Apostoli et al., showed rabbits with prolonged systemic exposure to Co or Cr in isolation or combined have optic and auditory neuropathy. Cobalt produces toxicity in rats, leading to glioma, neuroblastoma, and neuronal cell damage. In addition, cobalt has a dose-dependent effect on apoptosis and secondary necrosis in astrocytes, through the generation of reactive oxygen species and subsequent α subunit of hypoxia inducible factor-1 pathway activation.

Our FIRST data suggest that this concentration of metal exposure is not associated with a detectable loss in volume of the major brain structures. Both of our patient groups had brain and subcortical volumes similar to those found in the general population for individuals of similar age and sex. Similarly, although both Co and Cr may be deposited in the basal ganglia, especially in the pallidum, putamen, and caudate nucleus, our PRIME analysis of R2* data suggest that this exposure does not result in an increase in detectable deposition of paramagnetic ions within these structures. Our methods have been used previously to assess iron concentration within the basal ganglia of patients with HIV and Parkinson disease. Concentrations of iron up to 200µg/g tissue wet weight are found in the globus pallidus, red nucleus, substantia nigra, and putamen regions. A previous study showed a sensitivity of the PRIME sequence R2* relaxation time to iron deposition of approximately 0.02/s/µgFe/g wet weight of tissue. In a study to measure R2* in intact postmortem brains followed by quantitative postmortem brain measurement with inductively coupled plasma mass spectrometry, Langkammer et al found a formula for R2* (second-1) = 14.3 + 0.27 [iron concentration] (mg Kg-1 wet mass). The use of this approach to detect other metals such as Co and Cr is not so well-developed. Co, like iron, displays ferromagnetism, while Cr has a lower paramagnetic susceptibility of +167 µm/10-6 cm3 mol-1. As a sulfate, this increases to +11,800µm/10-6 cm3 mol-1. No differences were observed for R2* between the patient groups we studied, but this may reflect the expected lower sensitivity of the PRIME sequence to effects from Co and Cr compared with iron and the relatively low concentrations of Co and Cr in the brain. Typical normal levels in the cortex are 2 µg/Kg for Co and 10 µg/Kg for Cr, which are several orders of magnitude lower than those of iron (50–250 mg/Kg). If the serum concentration of these elements was raised by a factor of 50 and they crossed the blood-brain barrier, the maximum levels of Co and Cr in the brain would be ~500 µg/Kg, and thus our failure to detect these metals may be due to insensitivity of the PRIME sequence method to these relatively low metal concentrations compared with iron in brain tissue.

This is the first study to explore whether chronic exposure to moderate circulating metal is associated with brain structural and functional change. It was, by its nature, exploratory and not powered for analysis of specific brain areas or with a priori knowledge of potential effect sizes. While we observed some differences in brain morphometry between the groups, these did not remain after multiple-comparison correction, possibly related to sample size. Further study focusing analysis on the visual cortex and basal ganglia is warranted to confirm these preliminary findings in patients prescreened for potential confounders such as coexistence of primary ocular pathology.

CONCLUSIONS

This exploratory study was conducted in a clinically asymptomatic population and was cross-sectional in design. The patients with MoMHR who we studied had circulating metal concentrations that, though elevated, were not indicative of a failing prosthesis. We thus aimed to detect subclinical signals to indicate whether metal exposure due to prosthesis choice is associated with structural or functional changes in the brain and, if so, which systems show the greatest signal change. Our detection of subtle structural changes associated with metal exposure suggests that further examination of the relationship between metal exposure, brain structure, and clinical symptoms in a larger group of patients across a higher concentration range of metal exposure and for longer periods is warranted to definitively address the relationship between exposure and structural change and to determine their clinical relevance.

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REFERENCES

32. Gooddy W, Hamilton EI, Williams TR. Mitochondrial DNA damage and a hypoxic response are induced by CoCl(2) in rat neuronal PC12 cells. Nucleic Acids Res 2000;28:2135–40