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Evaluation of Parkinson Disease and Alzheimer Disease with the Use of Neuromelanin MR Imaging and ^{123}I -Metaiodobenzylguanidine Scintigraphy

F. Miyoshi, T. Ogawa, S.-i. Kitao, M. Kitayama, Y. Shinohara, M. Takasugi, S. Fujii, and T. Kaminou

ABSTRACT

BACKGROUND AND PURPOSE: Progressive changes in the substantia nigra pars compacta and locus ceruleus of patients with Parkinson disease and Alzheimer disease visualized by neuromelanin MRI and cardiac postganglionic sympathetic nerve function on ^{123}I -metaiodobenzylguanidine scintigraphy have not been fully evaluated. We compared the diagnostic value of these modalities among patients with early Parkinson disease, late Parkinson disease, and Alzheimer disease.

MATERIALS AND METHODS: We compared contrast ratios of signal intensity in medial and lateral regions of the substantia nigra pars compacta and locus ceruleus with those of the tegmentum of the midbrain and the pons, respectively, by use of neuromelanin MRI in patients with early Parkinson disease ($n = 13$), late Parkinson disease ($n = 31$), Alzheimer disease ($n = 6$), and age-matched healthy control subjects ($n = 20$). We calculated heart-to-mediastinum ratios on ^{123}I -metaiodobenzylguanidine scintigrams after setting regions of interest on the left cardiac ventricle and upper mediastinum.

RESULTS: The signal intensity of the lateral substantia nigra pars compacta on neuromelanin MRI was significantly reduced in early and late Parkinson disease, and that of the medial substantia nigra pars compacta was gradually and stage-dependently reduced in Parkinson disease. The signal intensity of the locus ceruleus was obviously reduced in late Parkinson disease. Signal reduction was not significant in the substantia nigra pars compacta and locus ceruleus of patients with Alzheimer disease. The heart-to-mediastinum ratio on ^{123}I -metaiodobenzylguanidine scintigrams was stage-dependently reduced in Parkinson disease and normal in Alzheimer disease. The signal intensity ratios in substantia nigra pars compacta and locus ceruleus on neuromelanin MRI positively correlated with the heart-to-mediastinum ratio on ^{123}I -metaiodobenzylguanidine scintigrams.

CONCLUSIONS: Both neuromelanin MRI and ^{123}I -metaiodobenzylguanidine scintigraphy can help to evaluate disease progression in Parkinson disease and are useful for differentiating Parkinson disease from Alzheimer disease.

ABBREVIATIONS: LC = locus ceruleus; MIGB = ^{123}I -metaiodobenzylguanidine; NmMRI = neuromelanin MR imaging; PD = Parkinson disease; SNc = substantia nigra pars compacta

Early Parkinson disease (PD) can easily be mistaken for any number of disorders, including other forms of parkinsonism, such as multiple system atrophy, progressive supranuclear palsy, corticobasal degeneration, and dementia with Lewy bodies. Other degenerative diseases, such as Alzheimer disease and primary lateral sclerosis, can also be mistaken for PD.¹ Improving diagnostic accuracy is critical for the early differentiation of PD and other neurode-

generative types of parkinsonism because their prognoses are very different and the choice of treatment strategy is extremely important.

Neuromelanin is a dark pigment that locates within certain catecholamine neurons of the human brain, such as the dopaminergic neurons of the substantia nigra pars compacta (SNc) and the noradrenergic neurons of the locus ceruleus (LC).² It is thought to be formed as a by-product of the catecholamine metabolism cascade through enzymatic and/or oxidative polymerization.^{2,3} PD is characterized by the progressive loss of dopaminergic neurons that contain neuromelanin in the SNc and of noradrenergic neurons in the LC. Several pathologic studies have shown the selective loss of ventral intermediate and lateral cell groups of the SNc in PD.⁴⁻⁶ Noradrenergic neurons are also lost in the LC of patients with Alzheimer disease.⁷

It is reported that neuromelanin MRI (NmMRI) could visualize decreased signal intensity in regions that reflect the loss of

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Table 1: Clinical characteristics of patient and control groups

	Early Parkinson Disease	Late Parkinson Disease	Alzheimer Disease	Control
Patients, <i>n</i>	13	31	6	20
Hoehn and Yahr stage	I:3, II:10	III:16, IV:12, V:3	/	/
Male/female	5/8	14/17	3/3	5/15
Age, y, mean $\pm$ SD	68.3 $\pm$ 5.88	71.8 $\pm$ 8.95	75.7 $\pm$ 9.52	74.8 $\pm$ 5.41
Duration, y, mean $\pm$ SD	4.30 $\pm$ 5.37	9.48 $\pm$ 6.86	2.5 $\pm$ 3	/
Hasegawa Dementia Scale-Revised, mean $\pm$ SD	/	/	12.8 $\pm$ 4.76	/

Note:—Age does not differ between patient and control groups ($P = .082$; 1-way analysis of variance).

neurons containing neuromelanin^{8,9} and the signal intensity in the SNc and LC was greatly reduced on NmMRI from patients with PD. Several investigators subsequently reported that NmMRI could show a reduction in the contrast ratio and volume of the SNc.^{10,11}

The physiologic analog of noradrenaline, ¹²³I-metaiodobenzylguanidine (MIBG), traces uptake and transport both in noradrenaline presynaptic sympathetic nerve terminals and in subsequent vesicular storage.¹² Postganglionic presynaptic cardiac sympathetic nerve endings can be noninvasively assessed by MIBG scintigraphy because a reduction of MIBG uptake indicates postganglionic sympathetic dysfunction. Cardiac MIBG uptake is reduced in patients with Lewy body diseases such as PD, as well as dementia with Lewy bodies, and MIBG scintigraphy can also help to differentiate PD from other types of parkinsonism.^{13,14}

However, progressive changes in PD and Alzheimer disease have not been fully evaluated by NmMRI and MIBG scintigraphy. We determined the usefulness of these modalities for differentially diagnosing PD and Alzheimer disease by analyzing changes in signal intensity in the SNc and LC and MIBG uptake in the left cardiac ventricle and evaluated correlations between NmMRI and MIBG scintigraphy findings.

MATERIALS AND METHODS

Patients

We investigated patients who had been tentatively diagnosed with suspected PD and who were finally confirmed as having PD or Alzheimer disease between December 2008 and April 2012. Both MRI and MIBG scintigraphy acquired within 1 month were retrospectively evaluated. Probable PD and Alzheimer disease were diagnosed according to the criteria of the United Kingdom Brain Bank and National Institute of Neurologic and Communicative Disorders and Stroke and the Alzheimer Disease and Related Disorders Association, respectively. Patients with PD were assigned to groups with either early or late PD on the basis of Hoehn and Yahr staging.¹⁵ Early PD comprised stages I and II, and late PD comprised stages III, IV, and V.

Patients with symptomatic cerebrovascular diseases and other central nervous system disorders were strictly excluded from the NmMRI evaluation, and those with cardiac diseases, diabetes mellitus, and/or medications that can interfere with MIBG uptake were excluded from the MIBG scintigraphy assessment.

We finally enrolled 13 (male, $n = 5$; female, $n = 8$), 31 (male, $n = 14$; female, $n = 17$), and 6 (male and female, $n = 3$ each) patients with early PD, late PD, and Alzheimer disease, respectively (Table 1) [ages (mean $\pm$ SD with range) 68.3 $\pm$ 5.88 (59–85), 71.8 $\pm$ 8.95 (59–83), and 75.7 $\pm$ 9.52 (58–84) years, respectively]. The mean ($\pm$ SD with range) duration of early PD, late PD, and Alzheimer

mer disease was 4.30 $\pm$ 5.37 (0–9), 9.48 $\pm$ 6.86 (2–27), and 2.5 $\pm$ 3.00 (1–3) years, respectively. We used the Hasegawa Dementia Scale-Revised¹⁶ to determine cognitive impairment or dementia, which is similar to the Mini-Mental State Examination and has a total score of 30. Hasegawa Dementia Scale-Revised scores in patients with Alzheimer disease ranged from 6–17 (mean $\pm$ SD, 12.8 $\pm$ 4.76).

Twenty age-matched patients (male, $n = 5$; female, $n = 15$) [ages (mean $\pm$ SD with range) 74.8 $\pm$ 5.41 (64–87) years] without a history of motor or cognitive impairment and significant abnormalities on brain MRI during the same period served as a control group of NmMRI. However, we did not obtain MIBG scintigraphic data from the control group during this period.

Our institutional review board approved the study, and written informed consent was waived.

Image Acquisition

All MR images were acquired by means of a clinical 3T MR scanner (Signa Excite HD, GE Healthcare, Milwaukee, Wisconsin). Axial images were acquired parallel to the anterior/posterior commissure line. T1-weighted fast spin-echo sequences were applied to NmMRI with the following parameters: TR/TE, 600/13 ms; echo-train length, 2; section thickness, 2.5 mm with 1-mm intersection gaps; matrix size, 512 $\times$ 512; FOV, 220 mm; acquisition time, 12 minutes. The scan covered the area from the upper border of the midbrain to the inferior border of the pons. We excluded other coexisting central nervous system disorders by use of axial T1- and T2-weighted images, fluid-attenuated inversion recovery images, and diffusion-weighted images of the entire brain according to the following standard protocol for adult brain imaging at our hospital: T1-weighted spin-echo sequence, TR/TE, 600/15 ms; section thickness, 5 mm; FOV 220 mm; matrix 512 $\times$ 512; T2-weighted fast spin-echo sequence, TR/TE, 4000/90 ms; section thickness, 5 mm; FOV 220 mm; matrix 512 $\times$ 512; fluid attenuated inversion recovery sequence, TR/TE/IR, 4000/90/20; section thickness, 5 mm; FOV 220 mm; matrix 512 $\times$ 512; and diffusion-weighted imaging sequence, TR/TE, 4000/90 ms; section thickness, 5 mm; FOV 220 mm; matrix 512 $\times$ 512; maximum b factor, 1000 mm²/s.

The patients received an intravenous injection of 111 MBq of ¹²³I-MIBG, and static planar images of the chest were acquired 30 minutes later for 4 minutes in a 256 $\times$ 256 matrix by use of a dual-head gamma camera with a large field of view and a low-energy, high-resolution collimator (e.cam; Siemens, Erlangen, Germany).¹⁷

Image Analysis

Signal intensity was measured for quantitative NmMRI by setting regions of interest. We equally divided the SNc into medial and lateral regions at the level of the inferior colliculus and defined

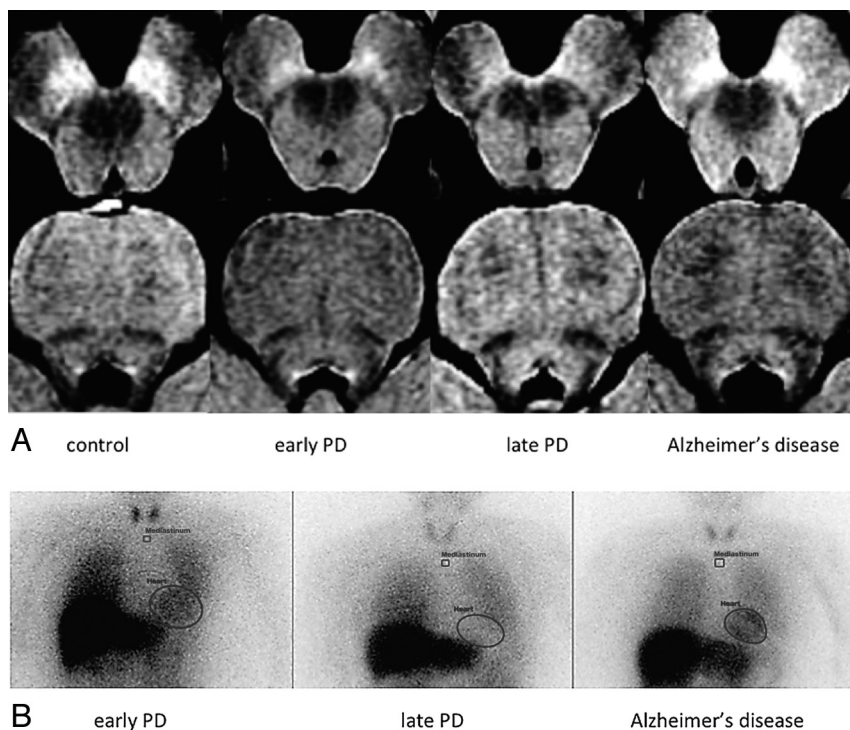


FIG 1. Findings of neuromelanin MRI and ^{123}I -metaiodobenzylguanidine scintigrams of patients and control subjects. **A**, NmMRI: Signals are reduced in NmMRI of substantia nigra pars compacta and locus ceruleus according to Parkinson disease stage. **B**, MIBG scintigraphy: MIBG scintigrams show PD stage-dependent cardiac MIBG reduction.

ROIs in these areas that included high signal intensity on NmMRI. We also defined ROIs symmetrically in the ventral tegmentum located in the anterolateral areas of aqueduct as controls. Concerning the LC, we defined the ROIs in the anterolateral areas around the fourth ventricle at the level of upper pons and also defined ROIs symmetrically in the tegmentum located just behind the medial lemniscus as controls. The sizes of the region of interest were 8, 2, and 10 mm² on the SNc, LC, and tegmentum of the midbrain and pons, respectively. We calculated the contrast ratios of the 3 bilateral portions by dividing their signal intensity by that of control areas such as the tegmentum of midbrain and pons.

An ROI was drawn manually over the whole heart on MIBG scintigrams to assess the global myocardial kinetics of MIBG. A second rectangular region of interest over the upper mediastinum served as a background reference region. The attenuation of the MIBG count in the heart and the mediastinum and heart-to-mediastinum count ratios were calculated for the images.

Statistical Analysis

Differences in contrast ratios between early PD, late PD, Alzheimer disease groups, and control subjects in the medial SNc, lateral SNc, and LC on NmMRI and between early PD, late PD, Alzheimer disease groups, and control subjects¹⁷ on MIBG scintigraphy were then statistically analyzed. The medial SNc, lateral SNc, and MIBG scintigrams were analyzed by means of a 1-way analysis of variance and the Bonferroni post hoc test, and the LC was analyzed by means of the Kruskal-Wallis and Dunn post hoc tests. The level of statistical significance was defined as $P < .05$ for all

tests. The contrast ratios of NmMRI and MIBG scintigram in early PD, late PD, and Alzheimer disease were analyzed by means of Spearman rank-order correlation coefficient test.

RESULTS

The signal intensity of the lateral SNc on NmMRI was reduced in early and late PD, and that of the medial SNc was gradually and stage-dependently reduced in PD. The signal intensity of LC was obviously reduced in late PD. Signal reduction was not significant in the SNc and LC of patients with Alzheimer disease (Fig 1A). The heart-to-mediastinum ratio on MIBG scintigrams was stage-dependently reduced in PD and normal in Alzheimer disease (Fig 1B).

Quantitative NmMRI revealed smaller contrast ratios in the lateral SNc of patients with early PD than in that of control patients, and in that of patients with late PD than in that of patients with Alzheimer disease and control subjects ($P < .05$) (Fig 2A). The contrast ratios of the medial SNc were smaller in patients with early PD than in control subjects and in patients with late PD than in those with

early PD, Alzheimer disease, and control subjects ($P < .05$) (Fig 2B). Contrast ratios in the LC of patients with late PD were smaller than in control patients ($P < .05$) (Fig 2C). Signals were not significantly reduced in the SNc and LC on NmMRI of patients with Alzheimer disease compared with control subjects.

The heart-to-mediastinum count ratio on the MIBG scintigram was stage-dependently reduced in patients with PD ($P < .05$) and normal (mean $\pm$ SD, 2.15 ± 0.15^{17}) in those with Alzheimer disease (mean $\pm$ SD, 2.19 ± 0.40) (Fig 2D).

Signal intensity ratios in the SNc on NmMRI positively correlated with heart-to-mediastinum count ratios on MIBG scintigrams (Fig 3). Spearman rank-order correlation coefficients in the medial and lateral SNc were 0.358 and 0.398, respectively, which reached statistical significance (Table 2).

DISCUSSION

Symmetrical spots of high signal intensity were visualized from around the anterolateral region to the floor of the fourth ventricle corresponding to the LC on axial NmMRI of the control group, and symmetrical bandlike high signal intensity was located around the posteromedial region of the cerebral peduncle corresponding to the SNc. Sasaki et al⁸ reported that their distribution obviously correlated with those of neuromelanin in gross specimens of the SNc and the LC, suggesting that the high signal intensity areas corresponded to neurons containing neuromelanin. We proved that NmMRI can accurately reflect the amount of neurons containing neuromelanin in the SNc, on the basis of a direct comparison of histopathologic and NmMRI findings of autopsied brains (unpublished data).

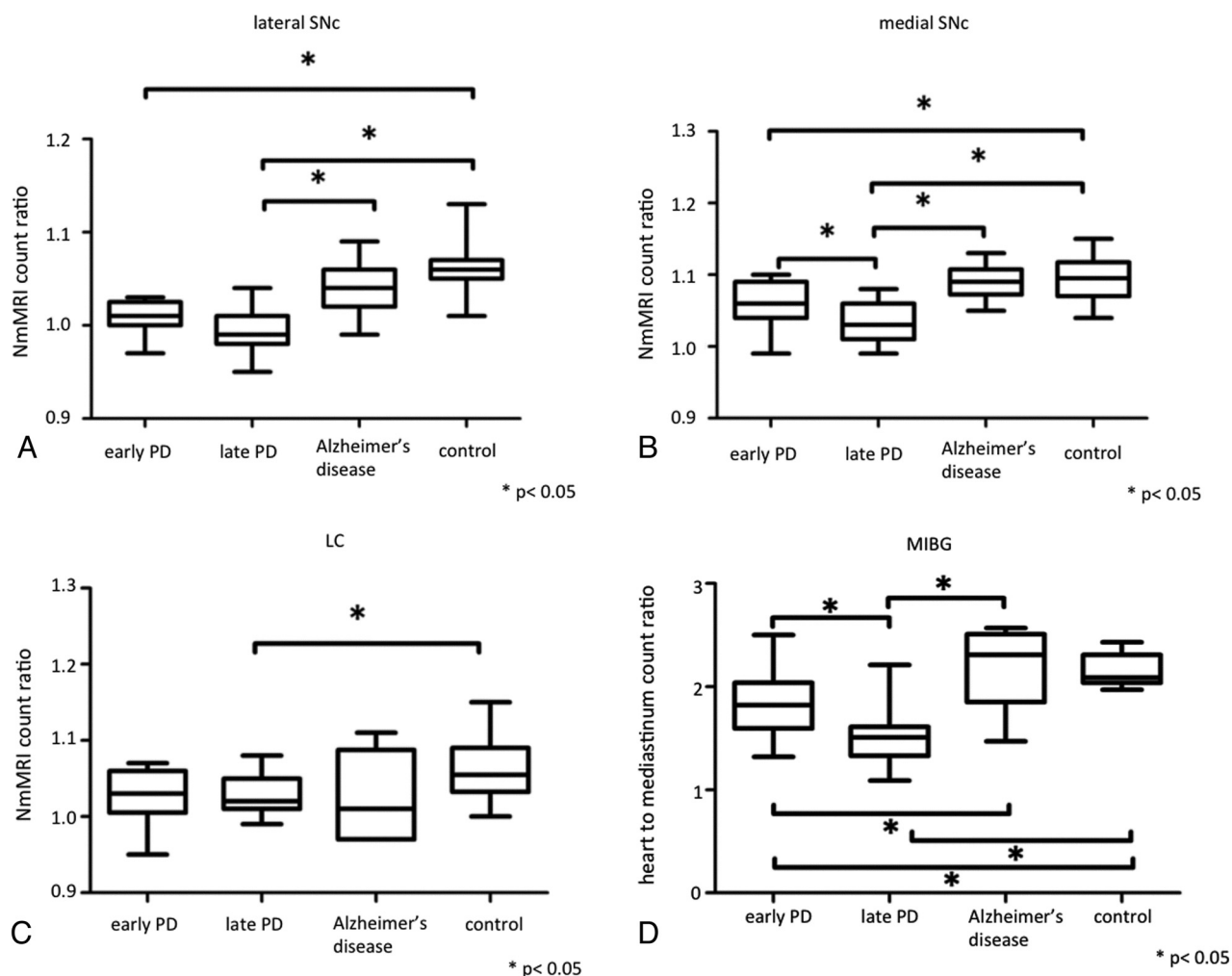


FIG 2. Box-and-whisker plots of signal intensity ratios on neuromelanin MRI and heart-to-mediastinum count ratios on ^{123}I -metaiodobenzylguanidine scintigrams in patients with Parkinson disease, patients with Alzheimer disease, and control subjects. Horizontal bars in boxes show median values. A, Lateral substantia nigra pars compacta; B, medial SNc; C, locus ceruleus; D, MIBG. * $P < .05$.

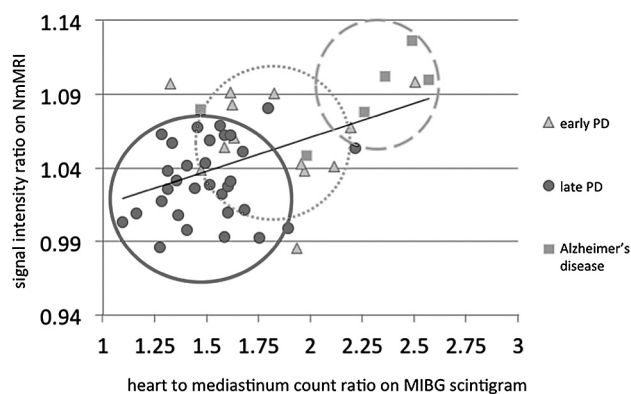


FIG 3. Correlation between signal intensity ratio in medial substantia nigra pars compacta on neuromelanin MRI image and heart-to-mediastinum count ratio on ^{123}I -metaiodobenzylguanidine scintigram. $\rho = 0.3984$.

Table 2: Correlation between signal intensity ratio in medial SNc on NmMRI and heart-to-mediastinum count ratio on MIBG scintigraphy

Spearman Coefficient Rate (ρ)	Lateral SNc	Medial SNc	LC
Heart-to-mediastinum count ratio	0.358	0.398	NS

SNC lesions extend from the lateral to the medial regions as PD progresses. Tanaka et al¹⁸ reported that a reduction in neuromelanin contrast started ventrolaterally and advanced medially in the substantia nigra on the basis of a visual assessment of NmMRI. However, the present study is the first effort to quantify medial and lateral SNc by use of NmMRI, to the best of our knowledge.

Braak et al¹⁹ proposed that α -synuclein-immunopositive Lewy neuritis and the accumulation of Lewy bodies that are characteristic pathologic features of PD emerge from the inferior brain stem and ascend to the nuclear gray and cortical areas. However, some studies have also indicated that Lewy bodies and neural cell loss are irrelevant in some regions of the brain in patients with PD. Noradrenergic neurons are relatively well preserved in the LC of patients with early PD, and they undergo different intracellular changes from the SNc.²⁰

Reports indicate that the signal intensity of the LC on NmMRI

We found that the contrast ratio of the lateral SNc was significantly decreased in early and late PD and that of the medial SNc gradually and stage-dependently decreased in PD, which corresponded to the pathologic lesions. These findings indicate that

is significantly reduced in PD.⁸ We identified significant signal reductions in the LC of patients with late PD, which suggests that lesions emerge in the LC during late PD. These findings are compatible with the findings of others indicating that the LC plays a compensatory role for the substantia nigra during early PD.^{21,22}

Noradrenergic neurons containing neuromelanin in the LC are disrupted in Alzheimer disease, but the clinical or functional importance of such disruption remains unclear.⁷ Some reports suggest that a reduction in the number of noradrenergic neurons affects the accumulation of β -amyloid, inflammation, or microcirculation in the LC.^{23,24} Our NmMRI study could not identify a significant signal reduction in the LC of patients with Alzheimer disease. Busch et al²⁵ reported that the LC begins to lose cells only during the later stages of Alzheimer disease. Therefore, we consider that this result can be attributed to the relatively short duration (2.5 years) of the disease and the small number of patients with Alzheimer disease in this study. Signals were significantly more reduced in the medial and lateral SNc of patients with late PD compared with Alzheimer disease, which is useful for differentiating these diseases.

The neurologic diseases with significant reduction in MIBG scintigraphy uptake include PD, dementia with Lewy bodies, and pure autonomic failure, and they are recognized as characteristic findings of Lewy-body diseases. On the other hand, heart-to-mediastinum count ratios are higher in Parkinson syndromes such as multiple system atrophy, progressive supranuclear palsy, and corticobasal degeneration than in PD. Therefore, MIBG scintigraphy is an important supportive tool for discerning Parkinson syndromes.²⁶ Sympathetic nerves in patients with PD are generally disturbed, and Lewy bodies are found in sinoatrial nodal ganglia, the myocardium, and paravertebral ganglia.^{27,28} A pathologic investigation has revealed that sympathetic nerve fibers are significantly reduced in patients with PD and well preserved in those with Alzheimer disease.²⁹ Although the heart-to-mediastinum count ratio is reduced on MIBG scintigrams of early PD, its relationship with progression remains controversial.^{29,30} In our present study, we found that the heart-to-mediastinum count ratio was significantly lower in late PD than early PD, indicating that signal reduction is stage-dependent. On the other hand, we could not exclude the effect of a possible relationship between heart-to-mediastinum signal reduction and duration. The heart-to-mediastinum count ratio was well-preserved in patients with Alzheimer disease, which helped to differentiate it from late PD.

To the best of our knowledge, this is the first comparison of brain NmMRI and MIBG scintigraphy of PD and Alzheimer disease. We identified a weak correlation in the SNc, implying that the findings of both modalities can serve as indicators of progressive PD, which involves both the central and peripheral autonomic nervous systems. Our results revealed that the autonomic nervous system gradually becomes disturbed over time.

The limitations of the present study are as follows. Because the study comprised very few patients with Alzheimer disease, further studies of a large population are required to validate our findings. Second, iron concentrations in the SNc that increase with age can mask signal alterations on NmMRI.³ Therefore, we could not exclude this effect when evaluating neuromelanin in elderly individuals. Third, the low spatial resolution of NmMRI did not allow 3D

image acquisition. Therefore, measurement errors might have arisen due to partial volume effects particularly when assessing changes in the LC.

CONCLUSIONS

The SNc becomes disturbed from the lateral to the medial region on NmMRI of the SNc as PD progresses and the LC is also disturbed in late PD. The sympathetic nerves in the left cardiac ventricle are also disturbed in MIBG scintigrams of PD. Thus, NmMRI and MIBG scintigraphy can be helpful tools to evaluate PD progression and to differentiate PD from Alzheimer disease.

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REFERENCES

1. Pahwa R, Lyons KE. **Early diagnosis of Parkinson's disease: recommendations from diagnostic clinical guidelines.** *Am J Manag Care* 2010;16(Suppl):S94–99
2. Fedorow H, Tribl F, Halliday G, et al. **Neuromelanin in human dopamine neurons; comparison with peripheral melanins and relevance to Parkinson's disease.** *Prog Neurobiol* 2005;75:109–24
3. Zecca L, Tampellini D, Gerlach M, et al. **Substantia nigra neuromelanin: structure, synthesis, and molecular behavior.** *Mol Pathol* 2001;54:414–18
4. Damier P, Hirsch EC, Agid Y, et al. **The substantia nigra of the human brain, II: patterns of loss of dopamine-containing neurons in Parkinson's disease.** *Brain* 1999;122:1437–48
5. Fearnley JM, Lees AJ. **Aging and Parkinson's disease: substantia nigra regional selectivity.** *Brain* 1991;114:2283–301
6. Halliday GM, McRitchie DA, Cartwright HR, et al. **Midbrain neuropathology in idiopathic Parkinson's disease and diffuse Lewy body disease.** *J Clin Neurosci* 1996;3:52–60
7. Förstl H, Burns A, Luthert P, et al. **Clinical and neuropathological correlates of depression in Alzheimer's disease.** *Psychol Med* 1992;22:877–84
8. Sasaki M, Shibata E, Tohyama K, et al. **Neuromelanin magnetic resonance imaging of locus ceruleus and substantia nigra in Parkinson's disease.** *NeuroReport* 2006;17:1215–18
9. Mann DM, Yates PO. **Possible role of neuromelanin in the pathogenesis of Parkinson's disease.** *Mech Ageing Dev* 1983;21:193–203
10. Schwarz ST, Rittman T, Gontu V, et al. **T1-weighted MRI shows stage-dependent substantia nigra signal loss in Parkinson's disease.** *Mov Disord* 2011;26:1633–38
11. Kashihara K, Shinya T, Higaki F. **Neuromelanin magnetic resonance imaging of nigral volume loss in patients with Parkinson's disease.** *J Clin Neurosci* 2011;18:1093–96
12. Wieland DM, Brown LE, Rogers WL, et al. **Myocardial imaging with a radioiodinated norepinephrine storage analog.** *J Nucl Med* 1981;22:22–31
13. Druschky A, Hilz MJ, Platsch G, et al. **Differentiation of Parkinson's disease and multiple system atrophy in early disease stages by means of I-123-MIBG-SPECT.** *J Neurol Sci* 2000;175:3–12
14. Spiegel J, Hellwig D, Farmakis G, et al. **Myocardial sympathetic degeneration correlates with clinical phenotype of Parkinson's disease.** *Mov Disord* 2007;22:1004–08
15. Hoehn MM, Yahr MD. **Parkinsonism: onset, progression and mortality.** *Neurology* 1967;17:427–42
16. Imai Y, Hasegawa K. **The revised Hasegawa's dementia scale (HDS-R) - evaluation of its usefulness as a screening test for dementia.** *J Hong Kong Coll Psychiatr* 1994;4:20–24
17. Tanabe Y, Harada H, Sugihara S, et al. **123I-metaiodobenzylguanidine**

- dine myocardial scintigraphy in panic disorder. *J Nucl Med* 2004;45:1305–08
18. Tanaka M, Aihara Y, Ikeda S, et al. Neuromelanin-related contrast in the substantia nigra semiquantitatively evaluated by magnetic resonance imaging at 3T: comparison between normal aging and Parkinson disease. *Clin Neurol* 2011;51:14–20
 19. Braak H, Del Tredici K, Rub U, et al. Staging of brain pathology related to sporadic Parkinson's disease. *Neurobiol Aging* 2003;24:197–211
 20. Halliday GM, Ophof A, Broe M, et al. α -Synuclein redistributes to neuromelanin lipid in the substantia nigra early in Parkinson's disease. *Brain* 2005;128:2654–64
 21. Isaías IU, Marotta G, Pezzoli G, et al. Enhanced catecholamine transporter binding in the locus coeruleus of patients with early Parkinson disease. *BMC Neurol* 2011;11:88
 22. Rommelfanger KS, Weinshenker D. Norepinephrine: the redheaded stepchild of Parkinson's disease. *Biochem Pharmacol* 2007;74:177–90
 23. Weinshenker D. Functional consequences of locus coeruleus degeneration in Alzheimer's disease. *Curr Alzheimer Res* 2008;5:342–45
 24. Haneka MT, Nadrigny F, Regen T, et al. Locus coeruleus controls Alzheimer's disease pathology by modulating microglial functions through norepinephrine. *Proc Natl Acad Sci U S A* 2010;107:6058–63
 25. Busch C, Bohl J, Ohm TG. Spatial, temporal and numeric analysis of Alzheimer changes in the nucleus ceruleus. *Neurobiol Aging* 1997;18:401–06
 26. Nakajima K, Yoshita M, Matsuo S, et al. Iodine-123-MIBG sympathetic imaging in Lewy-body diseases and related movement disorders. *J Nucl Med Mol Imaging* 2008;52:378–87
 27. Okada Y, Ito Y, Aida J, et al. Lewy bodies in the sinoatrial nodal ganglion: clinicopathological studies. *Pathol Int* 2004;54:682–87
 28. Orimo S, Takahashi A, Uchihara T, et al. Degeneration of cardiac sympathetic nerve begins in the early disease process of Parkinson's disease. *Brain Pathol* 2007;17:24–30
 29. Rascol O, Schelosky L. ^{123}I -metaiodobenzylguanidine scintigraphy in Parkinson's disease and related disorders. *Mov Disord* 2009;24(Suppl 2):S732–41
 30. Watanabe M, Takeda T, Nakamagoe K, et al. Sequential imaging analysis using MIBG scintigraphy revealed progressive degeneration of cardiac sympathetic nerve in Parkinson's disease. *Eur J Neurol* 2011;18:1010–13