123I-metaiodobenzylguanidine myocardial scintigraphy in Parkinson’s disease

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Abstract

Objectives—123I-metaiodobenzylguanidine (MIBG) myocardial scintigraphy is clinically used to estimate local myocardial sympathetic nerve damage in some forms of heart disease, autonomic nerve disturbance in diabetic neuropathy, and disturbance of the autonomic nervous system in neurodegenerative disease. In the present study, examinations were performed to clarify (1) the proportion of cardiac sympathetic nerve disturbance in Parkinson’s disease, (2) the usefulness of 123I-MIBG myocardial scintigraphy to detect sympathetic nerve disturbances compared with autonomic function tests, (3) cardiac function in patients who have a decreased MIBG uptake in 123I-MIBG myocardial scintigraphy, (4) the usefulness of 123I-MIBG myocardial scintigraphy to differentiate Parkinson’s disease from the other neurological diseases mimicking it.

Methods—123I-MIBG myocardial scintigraphy was performed, together with autonomic function tests and cardiac examinations in 46 patients with Parkinson’s disease and 25 patients with vascular parkinsonism, essential tremor, or multiple system atrophy.

Results—In an anterior image study, the average count per pixel in heart to mediastinum (H/M) ratio decreased in 80% of the patients with Parkinson’s disease in the early phase and 84% in the late phase. The mean H/M ratio in Parkinson’s disease was significantly lower than that in controls and the other diseases. The H/M ratio tended to decrease with the disease progression. In almost half of the patients in Hoehn and Yahr stage I, the H/M ratio was already decreased. The sympathetic skin response in upper and lower limbs, head up tilt test, and coefficient of variation of R-R interval were abnormal in 17%, 31%, 30%, and 17% of the patients, respectively. All the patients with abnormal autonomic functions were in Hoehn and Yahr stage III, IV, or V. Echocardiography showed normal left ventricular function. Twenty four hour Holter electrocardiography detected no serious arrhythmias except for one patient with non-sustained ventricular tachycardia.

Conclusions—123I-MIBG myocardial scintigraphy might detect early disturbances of the sympathetic nervous system in Parkinson’s disease and might give useful diagnostic information to differentiate vascular parkinsonism, essential tremor, and multiple system atrophy from Parkinson’s disease.

Keywords: Parkinson’s disease; 123I-metaiodobenzylguanidine myocardial scintigraphy; autonomic function tests

Patients with idiopathic Parkinson’s disease may have several symptoms of autonomic dysfunction including constipation, anhidrosis, salorrhoea, seborrhoea, postural hypotension, and urinary disturbances. The cause of autonomic dysfunction in Parkinson’s disease may be due to the pathological changes in the centres of autonomic regulation. Lewy bodies, sometimes associated with neuronal loss, can be found in the sympathetic as well as the parasympathetic nervous system—namely, in the hypothalamus, the dorsal vagal motor nucleus, the Edinger-Westphal nucleus, the intermediolateral spinal column, and the sympathetic ganglia. Clinically various autonomic function tests have been carried out in patients with Parkinson’s disease. However the prevalence and severity of autonomic dysfunction vary, depending on the researchers. Magalhães et al reported that constipation, oesthatic hypotension, and bladder dysfunction were found in about one third of the patients with pathologically verified disease. On the contrary, van Dijk et al reported that no evidence of autonomic dysfunction was found in unmedicated patients and that mild autonomic dysfunctions were found in advanced or medicated patients. Now autonomic dysfunction in Parkinson’s disease is considered not to be rare, and may become apparent with progression of the disease or medication.

Metaiodobenzylguanidine (MIBG) is a physiological analogue of noradrenaline (norepinephrine) and is actively transported into noradrenaline granules of sympathetic nerve terminals by the noradrenaline transporter. 123I-MIBG myocardial scintigraphy can be performed very safely and is clinically used to estimate local myocardial sympathetic nerve damage in some heart diseases, autonomic nerve disturbance in diabetic neuropathy, disturbance of the autonomic nervous system in neurodegenerative diseases, pure autonomic failure, and familial amyloidotic polyneuropathy. In the present study, we performed 123I-MIBG myocardial scintigraphy together with autonomic function tests and cardiac examinations in patients with Parkinson’s disease, vascular parkinsonism, essential tremor, and
formed simultaneously with 201thallium chloride (TlCl) myocardial scintigraphy. After being in the supine position for 20 minutes, 111mBq $^{123}$I-MIBG (Daichi Radioisotope Laboratories Co, Tokyo, Japan) and the same amount of $^{201}$Tl (as chloride; Daiichi Radioisotope Laboratories Co, Tokyo, Japan) were injected intravenously. A single photon emission computed tomography (SPECT) and a planar image of the chest were obtained using a double headed gamma camera (PRISM-2000, Shimadzu Co, Japan) after 20 minutes (early phase) and 3 hours (late phase). Background subtraction was not performed from any ROI count. The normal value of the H/M ratio in this hospital, obtained in 10 healthy volunteers (seven men and three women, mean age 58.8 (SD 13.6) (range 38–73) years) is 2.26 (0.16) (1.94–2.58) (early phase) and 2.30 (SD 0.22) (range 1.86–2.74) (late phase).

**Head up tilt test**
A head up tilt test was performed using a tilt table. After being in the supine position for 10 minutes, each patient was lifted in the head up position, at least 60°, using a tilt table. When a reduction of systolic blood pressure of at least 20 mm Hg or diastolic blood pressure of at least 10 mm Hg occurred within 3 minutes after the upright position, the patient was diagnosed as having orthostatic hypotension.

**Sympathetic skin response**
The sympathetic skin response was performed according to the method described by Yokota et al. Briefly, standard EMG disc electrodes were covered with conducting paste and attached to each site, as well as to the sole bilaterally. The skin temperature in all patients was kept above 31°C. Recordings were made on Neuropac Four (Nihon Kohden Co, Japan). Stimuli consisting of single square pulses of 200 ms duration, 20 mA intensity, were applied to the supraorbital nerves bilaterally on the forehead. More than 5 stimuli were administered at irregular intervals. Peak to peak amplitude of each response was measured. The following responses were considered to be abnormal as described previously: (1) absence of response, (2) absence of response at one site when responses at the other sites were continuously recorded. When the amplitude of the responses was <1000 µV for the palm or <300 µV for the sole, we considered it as a low response.

**Coefficient of variation in R-R interval (CVR-R)**
Electrocardiography was recorded by Cardiofax A (Nihon Kohden Co, Japan). After being in the supine position for 10 minutes, the ECG of each patient was recorded for 3 minutes. The CVR-R was calculated as 1 SD of R-R interval/mean value of R-R interval ×100 (%). The age matched normal value in this hospital, obtained from 20 healthy volunteers (10 men and 10 women, mean age 66.8 (SD11.6) (range 42–84) years of age) is 2.84 (SD 0.92) (range 1.00–4.67)%.

**Cardiac examinations**
Echocardiography was recorded at rest to evaluate left ventricular function and valvular diseases. Twenty four hour Holter ECG was also recorded.

**STATISTICAL ANALYSIS**
The results are expressed as means (SD). Differences of the variances and averages were tested by Student’s t test and one way analysis of variance.
of variance (ANOVA). Correlations between the two groups were assessed by Pearson’s correlation test. p Values <0.05 was considered to indicate statistical significance.

Results

201TL MYOCARDIAL SCINTIGRAPHY

One patient with Parkinson’s disease and two patients with vascular parkinsonism disclosed an abnormal defect of the inferior or apex wall in the early phase. The rest of the patients examined were within normal limits.

CLINICAL CHARACTERISTICS

We excluded one patient with Parkinson’s disease and two patients with vascular parkinsonism with abnormal Tl scintigraphy and evaluated 45 patients with Parkinson’s disease (17 men and 28 women, mean age 68.8 (SD 10.2) (range 41–84) years). The mean duration of illness was 4.3 (3.9) (0.25–19) years and Hoehn and Yahr stage was I in eight patients, II in three, III in 21, IV in nine, and V in four.

Twenty five patients were given antiparkinsonian drugs at the time of examination: 432 (138) (200–600) mg levodopa/carbidopa in 14 patients, 467 (103) (400–600) mg levodopa in six, 4.5 (0.9) (4–6) mg trihexyphenidyl in 15, 115 (24) (100–150) mg amantadine in 13, 435 (339) (50–1000) µg pergolide in 13, 5.0 (3.5) (2.5–7.5) mg bromocriptine in two, and 500 (245) (200–800) mg l-threo-DOPS in five. The patients with other neurological diseases were as follows. Eleven patients had vascular parkinsonism (nine men and two women, mean age 72.8 (6.2) (62–83) years). Mean duration of illness was 2.8 (3.0) (0.25–5) years. Two patients were given 100 mg levodopa/carbidopa.

123I-MIBG MYOCARDIAL SCINTIGRAPHY

In all the patients on whom 123I-MIBG myocardial scintigraphy was performed, no adverse reactions were found. Figure 1 shows short axis views of SPECT of 201Tl and 123I-MIBG myocardial scintigraphy in a healthy control and a 57 year old woman with Parkinson’s disease (Hoehn and Yahr stage I). The myocardial uptake of both Tl (a) and MIBG (b) is normal in a healthy control. Although the myocardial uptake of Tl is normal (c), no uptake of MIBG (d) is seen in a patient with Parkinson’s disease.

In the patients with Parkinson’s disease, the H/M ratio decreased in 36 patients (80%) in the early phase and in 38 patients (84%) in the late phase. The mean H/M ratio in the early and late phase was 1.71 (0.36) and 1.53 (0.36) in patients with Parkinson’s disease, 2.26 (0.16), 2.30 (0.22) in normal controls, and 2.20 (0.16) and 2.16 (0.22) in disease controls. The H/M ratio in the early/late phases was significantly less than in normal (p<0.001) in patients with Parkinson’s disease, 2.26 (0.16), 2.30 (0.22) in normal controls, and 2.20 (0.16) and 2.16 (0.22) in disease controls. The H/M ratio in the early/late phase tended to decrease with the progression of Hoehn and Yahr stage and the H/M ratio in the early/late stage in Hoehn and Yahr stage I was significantly high compared with stages III, IV, and V. In the early stage of Parkinson’s disease (Hoehn and Yahr stages I and II), the H/M ratio had already decreased in six patients (55%) in the early phase and eight patients (73%) in the late phase (fig 3). The H/M ratio in the early/late phase was correlated with the progression of Hoehn and Yahr stage and the H/M ratio in the early/late stage in Hoehn and Yahr stage I was significantly high compared with stages III, IV, and V. In the early stage of Parkinson’s disease (Hoehn and Yahr stages I and II), the H/M ratio had already decreased in six patients (55%) in the early phase and eight patients (73%) in the late phase (fig 3). The H/M ratio in the early/late phase was correlated with the progression of Hoehn and Yahr stage and the H/M ratio in the early/late stage in Hoehn and Yahr stage I was significantly high compared with stages III, IV, and V. In the early stage of Parkinson’s disease (Hoehn and Yahr stages I and II), the H/M ratio had already decreased in six patients (55%) in the early phase and eight patients (73%) in the late phase (fig 3).
However the mean H/M ratio of early/late phase was not different between the two groups.

A CVR-r was obtained in 39 patients. Three patients were excluded because of atrial fibrillation (two patients) and frequent premature atrial contraction (one patient). The CVR-r was abnormal in four out of 36 patients (17%).

All the patients with abnormal autonomic functions were in Hoehn and Yahr stage III, IV, or V.

**CARDIAC EXAMINATIONS**

Echocardiography (36 patients) showed normal left ventricular function in all the patients examined. Mild or very mild valvular abnormalities were seen in nine patients. Twenty four hour Holter ECG detected non-sustained (eight beats) ventricular tachycardia in one patient. This patient was an 82 year old woman (Hoehn and Yahr stage III). The early phase of the H/M ratio (1.37) was the third lowest and the late phase of the H/M ratio (1.17) was the lowest of all. No serious arrhythmias were detected in the remaining patients and ST changes were not seen in any patients examined.

**OTHER NEUROLOGICAL DISEASES**

The mean H/M ratios (early/late) of the patients with vascular parkinsonism, essential tremor, and multiple system atrophy were significantly less (p<0.005/p<0.005) than that in the patients with Parkinson's disease. The mean H/M ratios (early/late) of the patients with Parkinson's disease was significantly lower than for the patients with vascular parkinsonism (p<0.005/p<0.005), essential tremor (p<0.001/p<0.0001), and multiple system atrophy (p<0.005/p<0.0001) (fig 2). The H/M ratios (early/late) of two patients of multiple system atrophy with orthostatic hypotension were 1.82/1.49 and 1.94/2.06, respectively.

**Discussion**

Hakusui et al first reported a decreased myocardial MIBG uptake in patients with Parkinson's disease by $^{123}$I-MIBG myocardial scintigraphy.\(^{(19)}\) After that several investigators reported that myocardial MIBG uptake often decreased in patients with Parkinson's disease.\(^{(20-23)}\) However, the clinical relevance of this, as shown by $^{123}$I-MIBG myocardial scintigraphy, remains to be elucidated. In the present study, we performed examinations to clarify (1) the proportion of cardiac sympathetic nerve disturbance in Parkinson's disease, (2) usefulness of $^{123}$I-MIBG myocardial scintigraphy to detect sympathetic nerve disturbance compared with autonomic function tests in Parkinson's disease, (3) cardiac functions in the patients with Parkinson's disease who have a decreased MIBG uptake in $^{123}$I-MIBG myocardial scintigraphy, and (4) the usefulness of $^{123}$I-MIBG myocardial scintigraphy to differentiate Parkinson's disease from the other neurological diseases mimicking Parkinson's disease.
Firstly, we discuss the safety of $^{123}$I-MIBG myocardial scintigraphy. It was reported to be safe as follows: in 981 patients studied with $^{123}$I-MIBG myocardial scintigraphy, there were no severe adverse reactions, except complaints of burning on the injection site of the agent, nausea, palpitations, and feeling ill from four patients (0.4%). In all the patients in whom $^{123}$I-MIBG myocardial scintigraphy was performed, no adverse reactions were found. Therefore we infer that it can be performed very safely compared with the autonomic function tests such as the head up tilt test, sympathetic skin response, and CVR-R.

In the present study, cardiac sympathetic nerve disturbances were detected in 84% of the patients with Parkinson's disease. The severity of the disturbances was correlated with the progression of the disease and the duration of illness. The decrease in MIBG uptake in the early phase was considered to be attributable to denervation of the postganglionic cardiac sympathetic nerve due to diabetic neuropathy or various heart diseases including myocardial infarction, cardiomyopathy, hypertensive heart disease, and congestive heart disease, disturbance of the sympathetic nerve in the CNS, disturbance of the noradrenaline transporter, and several drugs already mentioned. The decrease in MIBG uptake in the late phase was due to increased MIBG wash out from the myocardium in addition to the causes in the early phase. In the present study, no patients were receiving drugs that may have interfered with MIBG uptake by sympathetic nerve terminals or various heart diseases including myocardial infarction, cardiomyopathy, hypertensive heart disease, and congestive heart disease, disturbance of the sympathetic nerve in the CNS, disturbance of the noradrenaline transporter, and several drugs already mentioned. Although the disturbance of the noradrenaline transporter has been reported in heart diseases including ischaemic heart disease, left ventricular hypertrophy, and congestive heart disease, all the patients had no evidence of them. Therefore, we infer that the decreased MIBG uptake is due to disturbances of sympathetic nerves including the postganglionic cardiac sympathetic nerves or the sympathetic nerves in the CNS. Recently Goldstein et al presented a new clinical pathophysiological classification of dysautonomias using myocardial 6-$^{18}$F fluorodopamine PET (6-$^{18}$F-FDA on PET) and cardiac noradrenaline spill over. In the patients with Parkinson's disease with autonomic failure, both decreased cardiac concentration of 6-$^{18}$F-FDA on PET and cardiac noradrenaline spill over were shown to be similar to those in patients with pure autonomic failure, indicating the loss of myocardial sympathetic nerve terminals. This peripheral lesion which is responsible for autonomic failure in Parkinson's disease is similar to that in a previous report. We detected cardiac sympathetic nerve disturbances in 84% of the patients with Parkinson's disease and even in the early stage (Hoehn and Yahr stages I and II), the H/M ratio was already decreased in 73% of the patients. On the contrary, only 29%, 42% (abnormal and low response), and 11% of the patients showed abnormality in the head up tilt test, sympathetic skin response, and CVR-R, respectively. None of the patients in Hoehn and Yahr stages I and II showed abnormal autonomic functions. In previous reports, orthostatic hypotension was found in various proportions of the patients with Parkinson's disease, depending on the accuracy of the diagnosis for Parkinson's disease, disease severity, antiparkinsonian drugs, and criteria for orthostatic hypotension. Magalhães et al reported that orthostatic hypotension was found in about one third of the patients with pathologically verified Parkinson's disease; Hirashima et al reported that 36.1% of the patients with Parkinson's disease showed abnormal sympathetic skin response, and 12% showed a low response, and Braune et al reported that 48% of the patients showed an abnormal sympathetic skin response compared with age matched controls. These data are consistent with the present study. Kuroiwa et al reported that 19% of patients studied with Parkinson's disease had abnormal CVR-R at rest, which is similar to the present study. With the previous reports and the present study taken together, $^{123}$I-MIBG myocardial scintigraphy is one of the most useful methods of detecting autonomic nerve disturbances in patients with Parkinson's disease.

We also performed cardiac examinations. One patient exhibited non-sustained ventricular tachycardia on 24 hour Holter ECG. Because the H/M ratio of this patient was very low, serious arrhythmias might occur in a patient with extremely decreased myocardial MIBG uptake. But even such patients might not have serious arrhythmias because these were not found in the rest of the patients. Moreover, left ventricular function was normal in all the patients examined by echocardiography. Hypertrophic cardiomyopathy under exercise or pharmacological stress remains to be elucidated.

It is occasionally difficult to differentiate cardiovascular parkinsonism from Parkinson's disease especially when the disease is associated with multiple brain infarctions. Also, it is not always easy to differentiate essential tremor, multiple system atrophy, and progressive supranuclear palsy from Parkinson's disease especially in the early stage. Recently Yoshita reported that the mean value of H/M ratio in patients with Parkinson's disease was significantly lower than that in those with striatoni gracile degeneration and progressive supranuclear palsy, and that $^{123}$I-MIBG myocardial scintigraphy might be helpful in differentiating between these three diseases, especially in the early stage. In the present study, the mean ages of patients with vascular parkinsonism and essential tremor were matched with that of Parkinson's disease and the mean value of the H/M ratio in the patients with Parkinson's disease was significantly lower than that in those with vascular parkinsonism or essential tremor. Therefore $^{123}$I-MIBG myocardial scintigraphy might give us useful diagnostic information to aid in differentiation between vascular parkinsonism or essential tremor and Parkinson's disease.
For multiple system atrophy, the H/M ratio of the patients with orthostatic hypotension showed a slight decrease or lower limit. The ratio of the patients had no orthostatic hypotension and had normal MIBG uptake. Although the mean age of patients with multiple system atrophy was not matched with that of Parkinson’s disease and there were only a few cases, scintigraphy may give useful diagnostic information to differentiate Parkinson’s disease from multiple system atrophy without orthostatic hypotension, as reported by Yoshita et al.22

In conclusion, we infer that123I-MIBG myocardial scintigraphy can be performed very safely and that it might give us useful diagnostic information to differentiate the sympathetic nervous system in Parkinson’s disease and also that it might give us useful diagnostic information to differentiate vascular parkinsonism, essential tremor, and multiple system atrophy from Parkinson’s disease.


