

EDITORIAL POINT OF VIEW

Current Clinical Practice of Nuclear Cardiology in Japan

Keiichiro Yoshinaga, MD, PhD, FACC, FASNC

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Abstract

For the numerous cardiology clinical practices in Japan, nuclear cardiology imaging tests are among the most important diagnostic tools. The Japanese nuclear cardiology community has developed a new application using ^{18}F -fluorodeoxyglucose (FDG) positron emission tomography (PET) to diagnose cardiac sarcoidosis, as well as new diagnostic imaging tests using ^{123}I -beta-methyl-p-iodophenyl-pentadecanoic acid (BMIPP) and ^{123}I -metaiodobenzylguanidine (MIBG). These new approaches have become popular worldwide. The Japanese Circulation Society (JCS) and the Japanese Society of Nuclear Cardiology (JSNC) have published clinical imaging guidelines and recommendations showing indications and standards for the new imaging tests. Current nuclear cardiology clinical practices in Japan may provide new insights for nuclear cardiology worldwide.

Keywords: Approval, Guidelines, Japanese Ministry of Health, Labor, and Welfare, Myocardial perfusion imaging, Positron emission tomography

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Based on the Japanese ISOTOPE Foundation survey, 1,170 facilities performed nuclear medicine tests in 2012. The population of Japan is currently 125 million, with 0.94 nuclear facilities per 100,000 population. Twenty-five percent of those facilities (296 of 1,170) performed PET studies. Among them, 104 performed PET only, and the remaining 192 performed both PET and SPECT (1,2).

Nuclear cardiology practice in Japan

The 2012 Japanese ISOTOPE Foundation survey showed the total number of nuclear cardiology studies performed as 31,475, an 11.0% decrease compared to the number in the previous survey in 2007. The Japanese Circulation Society (JCS) also conducts a clinical practice survey known as the Japanese Registry Of All cardiac and vascular Disease (JROAD) every year. Results of this survey also show a decreasing trend in the use of single-photon emission computed tomography (SPECT) myocardial perfusion imaging (MPI). In contrast, the overall numbers of PET studies are gradually increasing, possibly indicating that PET is gradually replacing conventional SPECT MPI, similar to the case in

North America (3).

Ministry's approval of nuclear cardiology in Japan

Almost all members of the Japanese population are covered by either governmental or company health insurance plans. Health insurance covers 70% of hospital visits, in-patient treatment, and drugs. In this regard, it is quite important to obtain Japanese Ministry of Health, Labor, and Welfare (JMHLW) approval and reimbursement for new diagnostic imaging tests. JMHLW has approved 3 MPI and two I-123 imaging tracers for SPECT (Table 1) (4). JMHLW currently approves ^{13}N -ammonia (NH_3) for MPI, ^{15}O -labeled gas for radionuclide angiography, and ^{18}F -fluorodeoxyglucose (FDG) for myocardial viability and cardiac sarcoidosis testing (Table 1) (2,5). Unlike the earlier approval of SPECT imaging, in the case of PET imaging, JMHLW approval was contingent upon some conditions. Currently standard hybrid PET/computed tomography (CT) scanners use CT for attenuation correction. However, for cardiac PET, only the PET part is reimbursed even when a PET/CT scanner is used, and the simultaneously performed CT study is not reimbursed. ^{13}N - NH_3 PET MPI is

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Keiichiro Yoshinaga

Director, Diagnostic and Therapeutic Nuclear Medicine National
Institute of Radiological Sciences 4-9-1 Anagawa, Inage-Ku, Chiba,
Japan 263-8555

E-mail: yoshinaga.keiichiro@qst.go.jp

Table 1 Japanese Ministry of Health, Labor, and Welfare's approval of the use of SPECT and PET in nuclear cardiology

Test	Date of approval	Indication	Condition
SPECT			
²⁰¹ Tl-thallium	April 19, 1979	Diagnosis of heart disease	
^{99m} Tc sestamibi	April 2, 1993	Diagnosis of heart disease	
^{99m} Tc tetrofosmin	April 15, 1994	Cardiac function measurement using first pass method	
¹²³ I-BMIPP	January 19, 1993	Diagnosis of heart disease	
¹²³ I-MIBG	October 2, 1992	Diagnosis of heart disease	
PET			
¹⁸ F-fluorodeoxyglucose	March 8, 2002	Diagnosis of viability	In case of difficult to diagnose myocardial viability using myocardial SPECT
	March 5, 2012	Detection of inflammatory myocardial regions in cardiac involvement sarcoidosis	Approval of PET only. CT attenuation correction is not reimbursed.
¹³ N-ammonia	March 5, 2012	Diagnosis of ischemic heart disease	When other tests are unable to make diagnosis Approval of PET CT attenuation correction is not reimbursed

¹²³I-BMIPP: ¹²³I-beta-methyl-p-iodophenyl-pentadecanoic acid; ¹²³I-MIBG: ¹²³I-metaiodobenzylguanidine

Table 2 Indication of ¹²³I-BMIPP and ¹²³I-MIBG based on JCS guidelines

Test	Indication	Class	Evidence level
¹²³ I-BMIPP	Diagnosis of unstable angina	1	B
	Diagnosis of coronary vasospastic angina	2a	C
	Diagnosis of stable coronary artery disease	2a	C
¹²³ I-MIBG	Assessment of severity and prognosis of heart failure	1	B
	Assessment of treatment effects on heart failure	2a	C

¹²³I-BMIPP: ¹²³I-beta-methyl-p-iodophenyl-pentadecanoic acid; ¹²³I-MIBG: ¹²³I- metaiodobenzyl-guanidine

reimbursed only when other noninvasive diagnostic imaging tests such as SPECT MPI are unable to provide a diagnosis of coronary artery disease (CAD).

The JMHLW's approval of PET including the use of ¹⁸F-FDG to detect cardiac involvement of sarcoidosis is significant considering that it is not approved in other countries. This approval is likely based on the significant evidence in Japan showing the usefulness of ¹⁸F-FDG PET in detecting cardiac involvement of sarcoidosis (6). This approval may help other countries to obtain approval for the use of ¹⁸F-FDG PET in detecting cardiac sarcoidosis.

Japanese Circulation Society guidelines for nuclear cardiology and Japanese Society of Nuclear Cardiology recommendation

JCS guidelines, initially issued in 1989, provided clinical indications for nuclear cardiology. The current JCS guidelines were updated in 2010 (7). In accordance with JMHLW approval, the JCS guidelines include indications for the clinical use of ¹²³I-BMIPP and ¹²³I-MIBG (Table 2).

In 2014, the Heart Rhythm Society (HRS) issued an expert statement on the diagnosis of cardiac involvement of

sarcoidosis, in which it raised the importance of ¹⁸F-FDG PET imaging for that purpose (8). However, previous to that, no clinical standard for ¹⁸F-FDG PET in sarcoidosis patients existed. In this regard, the Japanese Society of Nuclear Cardiology (JSNC) established a subcommittee and issued the recommendation that ¹⁸F-FDG PET imaging be used for detecting cardiac sarcoidosis (6).

Conclusions

Current nuclear cardiology practice in Japan resembles that in North America but specifically differs in that ¹²³I-tracers imaging and ¹⁸F-FDG PET have been approved for the diagnosis of cardiac sarcoidosis, differences that may be worthy of consideration for nuclear cardiology practice worldwide.

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Conflicts of interest

None

Reprint requests and correspondence:

Keiichiro Yoshinaga, MD, PhD
Director, Diagnostic and Therapeutic Nuclear Medicine
National Institute of Radiological Sciences 4-9-1 Anagawa,
Inage-Ku, Chiba, Japan 263-8555
E-mail: yoshinaga.keiichiro@qst.go.jp

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