

# Transcatheter aortic valve implantation and cognitive function in elderly patients with severe aortic stenosis



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## KEYWORDS

- aortic stenosis
- chronic heart failure
- TAVI

## Abstract

**Aims:** The aim of this study was to examine the mechanisms of cognitive impairment and reversibility in elderly patients with severe aortic stenosis (AS) after transcatheter aortic valve implantation (TAVI) with special reference to cerebral blood flow (CBF).

**Methods and results:** We examined 15 elderly patients with severe AS (mean age 83.2±4.5 years, 12 female) who underwent TAVI. Before and three months after TAVI, we evaluated cognitive function with the Logical Memory II test (LM II), cardiac output (CO) with echocardiography, and CBF with <sup>99m</sup>Tc single-photon emission computed tomography (SPECT). LM II score and CO were significantly increased after TAVI compared with baseline (p<0.01 for LM II, p<0.005 for CO). Notably, CBF in the local regions, including that in the right hippocampus, was significantly increased after TAVI (p<0.005 at each voxel). The patients with increased CO after TAVI also showed significantly increased CBF in the right hippocampus compared with those without it (p<0.01). Importantly, CBF in the right hippocampus was positively correlated with LM II scores (p<0.05).

**Conclusions:** These results provide the first evidence that TAVI may improve cognitive functions associated with increased cerebral perfusion especially in the hippocampus in elderly patients with severe AS.

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## Abbreviations

<b>AS</b>	aortic stenosis
<b>CBF</b>	cerebral blood flow
<b>CO</b>	cardiac output
<b>DW-MRI</b>	diffusion-weighted magnetic resonance imaging
<b>eNOS</b>	endothelial nitric oxide synthase
<b>GDS</b>	Geriatric Depression Scale
<b>LM II</b>	Logical Memory II test
<b>MCI</b>	mild cognitive impairment
<b>MMSE</b>	Mini-Mental State Examination
<b>MoCA</b>	Montreal Cognitive Assessment
<b>NYHA</b>	New York Heart Association
<b>SPECT</b>	single-photon emission computed tomography
<b>TAVI</b>	transcatheter aortic valve implantation

## Introduction

Severe aortic valve stenosis (AS) is the most common valvular heart disease in the elderly in Western countries and Asia that gradually leads to progression of valve calcification and eventually causes heart failure<sup>1,2</sup>. The interaction between the heart and the brain is important in the elderly with multiple comorbidities<sup>3</sup>, because the two important organ systems share many pathophysiological mechanisms<sup>3</sup>. Indeed, cognitive impairment is frequently noted in patients with AS<sup>4-6</sup>. Although severe AS is conventionally treated with surgical aortic valve replacement, the less invasive transcatheter aortic valve implantation (TAVI) has been developed for such elderly frail patients at high surgical risk<sup>7</sup>.

Previous studies have examined cognitive functions and diffusion-weighted magnetic resonance imaging (DW-MRI) in patients with severe AS who underwent TAVI<sup>4,8</sup>. Notably, recent studies have demonstrated that some patients with severe AS showed improved cognitive functions after TAVI<sup>5,9</sup>. However, detailed mechanisms for the improvement after TAVI remain to be examined. Notably, cerebral perfusion has been regarded as an important pathophysiological factor of the heart and brain interactions<sup>3</sup>. We and others have previously demonstrated that brain perfusion single-photon emission computed tomography (SPECT) is a useful imaging technique to evaluate regional cerebral perfusion and its relevance to cognitive impairment or stress cardiomyopathy<sup>10,11</sup>.

In the present study, we tested our hypothesis that TAVI increases CBF associated with increased cardiac output (CO) with a resultant improvement of cognitive functions in elderly patients with severe AS, using brain perfusion SPECT imaging before and three months after TAVI.

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## Methods

The present study protocol was approved by the ethics committee of the Tohoku University Graduate School of Medicine (No. 2018-1-329) and was performed in compliance with the Declaration of Helsinki (UMIN000034203).

## STUDY PATIENTS

From January 2017 to September 2018, we examined 57 consecutive patients with severe AS at the Tohoku University Hospital as candidates for TAVI. Inclusion criteria were 1) heart failure with New York Heart Association (NYHA) functional Class II to III symptoms, and 2) patient consent to undergo cognitive function tests for at least one hour. Exclusion criteria were 1) acute decompensated heart failure and heart failure with NYHA Class IV symptoms, 2) refusal of cognitive tests, and 3) insufficient quality of <sup>99m</sup>Tc SPECT. Based on these criteria, we excluded 35 patients in advance, including acute decompensated heart failure and heart failure with NYHA Class IV symptoms in 15, and refusal to undergo cognitive function tests for at least one hour in 20. Thus, we initially included 22 patients, seven of whom were then excluded because of dropout owing to refusal to undergo follow-up cognitive tests (n=2), and insufficient quality of <sup>99m</sup>Tc SPECT (n=5). Finally, we enrolled 15 patients in the present study with special reference to the association of cerebral blood flow (CBF) with cognitive functions (**Supplementary Figure 1**). Before and three months after TAVI, we measured cognitive functions with the Logical Memory II test (LM II)<sup>12</sup>, Mini-Mental State Examination (MMSE)<sup>13</sup>, and the Geriatric Depression Scale (GDS)<sup>14</sup>, CO with echocardiography, and CBF with <sup>99m</sup>Tc SPECT.

The baseline, TAVI procedure, and follow-up data were all collected in a dedicated database. Details of the TAVI procedure are shown in **Supplementary Appendix 1**.

## ECHOCARDIOGRAPHY

The details of echocardiography are shown in **Supplementary Appendix 2**.

## CBF IMAGE ACQUIREMENT AND PRE-PROCESSING

CBF can be measured not only by SPECT but also by MRI<sup>15,16</sup>. Since we and others have previously demonstrated that SPECT is a useful imaging technique to evaluate regional cerebral perfusion and its relevance to cognitive impairment or stress cardiomyopathy<sup>10,11</sup>, we selected SPECT for measuring CBF. H. Suzuki, who was blinded to the results of the imaging studies before and three months after TAVI, analysed and reported the SPECT scans. <sup>99m</sup>Tc-SPECT CBF images were acquired with a dual-head gamma camera (Symbia E; Siemens Healthineers, Erlangen, Germany). The following CBF image pre-processing and analyses were performed using SPM 12 software<sup>17</sup>. First, before CBF image analysis, we co-registered CBF images at three months to their corresponding baseline images. Second, the baseline and co-registered CBF images were normalised to the standard Montreal Neurological Institute space, using the SPECT template available in SPM 12. Finally, the normalised images were smoothed with an isotropic Gaussian kernel by convolving a 12 mm full width at half maximum to produce CBF maps. These pre-processing steps were described in detail in our previous reports<sup>16,18</sup>.

## ASSESSMENT OF COGNITIVE FUNCTIONS

A standardised cognitive assessment with the LM II, MMSE, and GDS was performed by a single experienced staff member blinded to the results of the imaging studies before and three months after TAVI. The LM subtest of the Wechsler Memory Scale-Revised is internationally used as an operational definition to identify individuals with mild cognitive impairment (MCI). In particular, the LM II test (a 30-minute delayed test of prose recall) is an indicator to discriminate between healthy older adults and individuals with very mild cognitive impairment<sup>12</sup>. MMSE is a widely used screening tool for cognitive impairment<sup>13</sup>. GDS is a screening instrument for late-life depression that demonstrates good accuracy<sup>14</sup>. In addition, GDS is based mainly on behavioural and cognitive aspects of depression and is not heavily weighted towards somatic complaints<sup>14</sup>. Thus, GDS is supposed to differentiate depressed from non-depressed elderly adults suffering from physical illness reliably.

## STATISTICAL ANALYSIS

Continuous variables are presented as mean±standard deviation (SD). Normality was assessed using the Shapiro-Wilk test. Continuous variables were compared by the Wilcoxon signed-rank test. Statistical analysis was performed using JMP® Pro 14 (SAS Institute Inc., Cary, NC, USA) at a significance threshold of  $p < 0.05$  except for voxel-wise CBF analyses.

We explored which brain areas showed CBF changes after TAVI by conducting a voxel-wise comparison between CBF maps before and three months after TAVI at an exploratory significance threshold of  $p < 0.005$ . CBF within the areas which changed after TAVI were calculated and were then used for a paired t-test between baseline and three months. A repeated measures linear mixed-model analysis was performed to evaluate changes in CBF and those in cognitive function tests. The details of the SPECT image pre-processing and analysis are shown in **Supplementary Appendix 3**.

## Results

### PATIENT CHARACTERISTICS

Clinical characteristics of the included and excluded patients are shown in **Supplementary Table 1** and **Supplementary Table 2**. There were no significant differences in the results of cognitive function tests at baseline between the included and excluded patients. In the present study, the mean age was  $83.2 \pm 4.5$  years, and 80% were female. On the basis of a cut-off of  $< 24$  points for MMSE, five patients (33.3%) were considered cognitively impaired, whereas no patients were diagnosed as having dementia that required treatment with acetylcholine esterase inhibitors. No patients had luminal narrowing  $> 25\%$  in the carotid arteries, although we did not evaluate the status of the posterior artery.

### PROCEDURAL CHARACTERISTICS AND CLINICAL OUTCOME

Procedural characteristics and clinical outcomes are shown in **Table 1**. No patients needed implantation of a second valve or

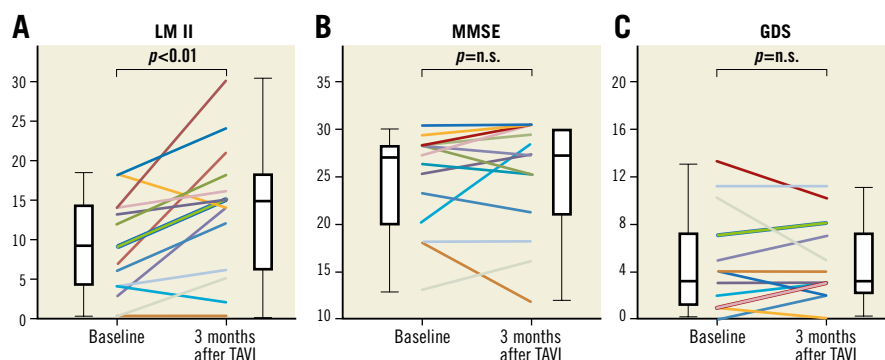
**Table 1. Procedural characteristics and clinical outcomes of the study population.**

		Patients (n=15)
<b>Procedural characteristics</b>		
Approach	Transfemoral	14 (93)
	Subclavian	1 (7)
	Transapical	0
Valve type	Edwards SAPIEN 3	6 (40)
	Medtronic CoreValve or Evolut R	9 (60)
	Need for second valve	0 (0)
<b>Clinical outcomes: 30 days</b>		
Stroke		0 (0)
Myocardial infarction		0 (0)
Major or life-threatening bleeding		1 (7)
New-onset atrial fibrillation		1 (7)
New pacemaker implantation		1 (7)
Acute kidney injury stage 2 or 3		0 (0)
Major vascular complication		1 (7)
<b>Echocardiographic characteristics</b>		
Peak velocity at discharge, m/s		2.28±0.45
Mean transprosthetic gradient at discharge, mmHg		10.80±4.60
Aortic valve area at discharge, cm <sup>2</sup>		1.73±0.29
Cardiac output, l/min		5.10±1.14
Moderate or severe aortic regurgitation at discharge		0 (0)
<b>Clinical outcomes: from 30 days to 3 months</b>		
Stroke		0 (0)
Myocardial infarction		0 (0)
Major or life-threatening bleeding		0 (0)
New-onset atrial fibrillation		0 (0)
Categorical variables are expressed as n (%) and continuous variables as mean±SD.		

showed myocardial infarction or cardiovascular death at 30 days after TAVI. Notably, no patients showed clinical symptoms or signs of transient ischaemic attack or stroke after TAVI. In addition, CO was also significantly increased at three months after TAVI compared with baseline (baseline,  $4.03 \pm 0.88$  vs 3 months,  $5.10 \pm 1.14$  L/min,  $p = 0.0045$ ).

### CHANGES IN COGNITIVE FUNCTIONS AFTER TAVI

At baseline, the mean scores of LM II, MMSE and GDS were  $8.7 \pm 1.5$ ,  $24.6 \pm 1.3$ , and  $4.3 \pm 1.1$ , respectively. LM II was significantly improved at three months after TAVI compared with baseline (baseline,  $8.7 \pm 6.0$  vs 3 months,  $13.8 \pm 8.1$ ,  $p < 0.01$ ). In contrast, there were no significant differences in MMSE or GDS during the study period (MMSE, baseline,  $24.6 \pm 1.3$  vs 3 months,  $25.2 \pm 1.5$ ,  $p = 0.42$ ; GDS, baseline,  $4.3 \pm 1.1$  vs 3 months,  $4.2 \pm 0.9$ ,  $p = 1.0$ ) (**Figure 1**). Among five patients (one third of the patients in the present study) with cognitive impairment at baseline, three



**Figure 1.** Changes in cognitive functions after TAVI in patients with severe AS. A) Logical Memory II (LM II) score was significantly improved at three months after transcatheter aortic valve implantation (TAVI) compared with baseline. B) & C) There were no significant differences in Mini-Mental State Examination (MMSE) or Geriatric Depression Scale (GDS) at three months after TAVI compared with baseline. n.s.: not significant

showed that LM II was improved at three months after TAVI. In these three patients with MMSE scores 23, 18 and 13 at baseline, LM II scores increased at three months after TAVI from 6 to 12, 4 to 6, and 0 to 5, respectively.

#### CHANGES IN CEREBRAL BLOOD FLOW AFTER TAVI

There were no significant differences in the whole CBF during the study period (baseline,  $39.3 \pm 1.0$  vs 3 months,  $39.2 \pm 1.0$  ml/100 g/min,  $p=0.76$ ). However, CBF in specific regions was significantly increased after TAVI compared with baseline (baseline,  $51.2 \pm 1.0$  vs 3 months,  $53.3 \pm 1.0$  ml/100 g/min,  $p<0.001$ ) (**Figure 2A-Figure 2F**). All five patients with cognitive impairment at baseline showed that CBF increased at three months after TAVI. Indeed, in these five patients with MMSE scores 23, 20, 18, 18 and 13 at baseline, CBF (ml/100 g/min) increased at three months after TAVI from 54.9 to 56.1, 48.3 to 50.6, 50.6 to 52.4, 53.6 to 56.3, and 51.0 to 53.3, respectively. This correlation between right hippocampal CBF and LM II scores was supported by the results from repeated measures linear mixed-model analysis ( $p=0.017$ ) (**Figure 2G**). Moreover, the patients with increased CO after TAVI had significantly increased CBF in the right hippocampus compared with those without it (with increased CO,  $1.06 \pm 0.07$  vs without,  $0.94 \pm 0.04$ , for changes in CBF in the right hippocampus after TAVI,  $p<0.01$ ) (**Figure 2H**). Importantly, there was no significant difference in blood pressure during the study period (systolic blood pressure, baseline,  $120.6 \pm 15.4$  vs 3 months,  $121.6 \pm 14.4$ ,  $p=0.57$ ; diastolic blood pressure, baseline,  $62.9 \pm 12.4$  vs 3 months  $64.9 \pm 9.2$ ,  $p=0.63$ ).

#### Discussion

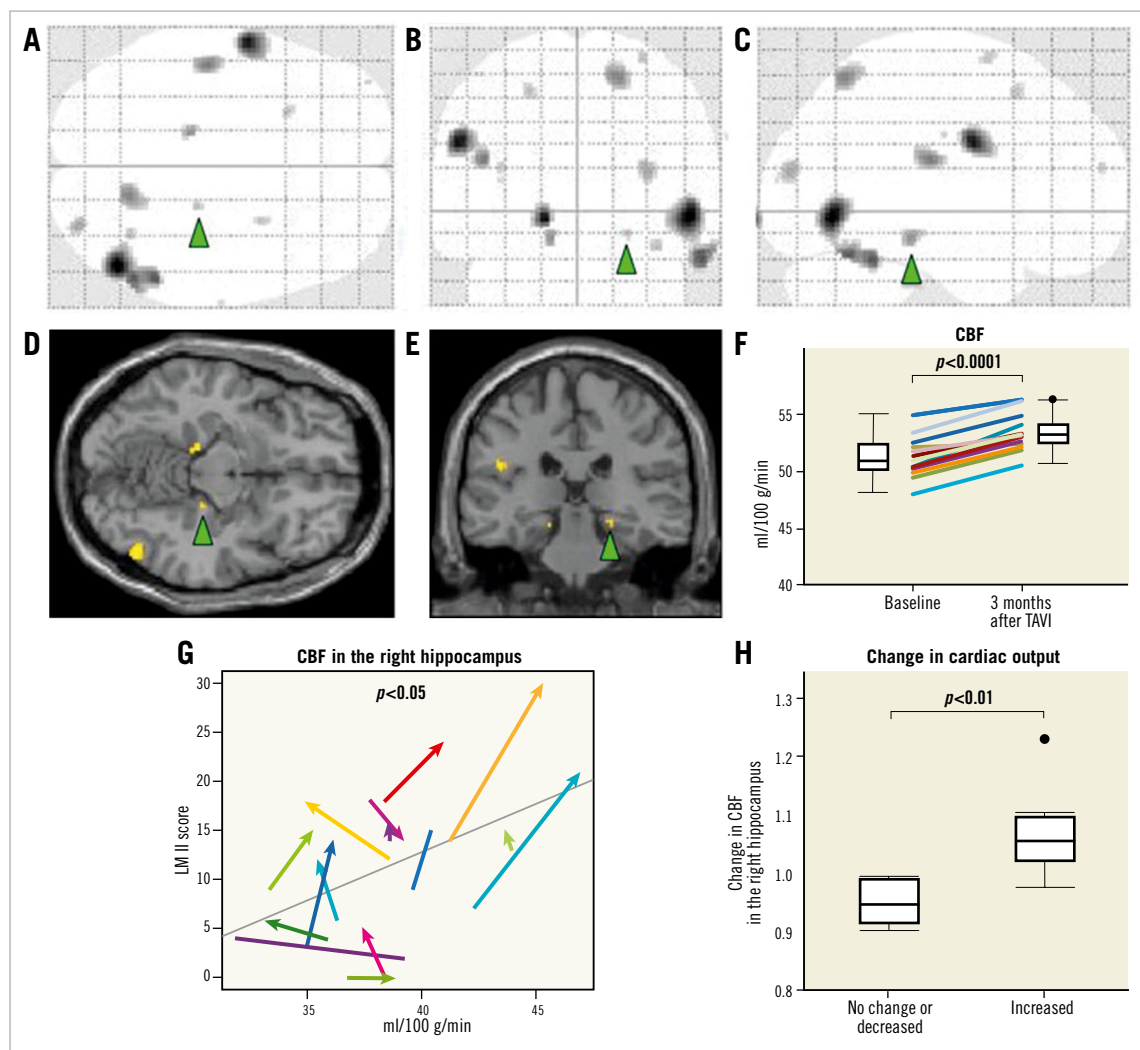
The major findings of the present study were that 1) LM II was significantly improved after TAVI, 2) CBF in the local regions, including the right hippocampus, was significantly increased at three months after TAVI, 3) increase in CO was associated with that in CBF in the right hippocampus, and 4) CBF in the right hippocampus was positively correlated with LM II. To the best

of our knowledge, this is the first study to demonstrate that TAVI may improve cognitive functions associated with increased cerebral perfusion especially in the hippocampus in elderly patients with severe AS.

#### CHANGES IN COGNITIVE FUNCTIONS AFTER TAVI

In the current practice guidelines, management of cognitive impairment needs to be improved, as proven therapeutic options are still lacking<sup>3</sup>. Although the number of patients with cognitive impairment and heart failure has been rapidly increasing in Western countries and Asia<sup>1,19</sup>, heart failure-associated cognitive impairment may be underestimated. Indeed, in the present study, five patients (33.3%) actually had cognitive impairment (MMSE  $<24$ ). A recent study also demonstrated that 22~39% of patients with severe AS had impaired cognitive functions at baseline<sup>5,6</sup>. In the present study, although there were no significant differences in MMSE or GDS at three months after TAVI, LM II was significantly improved at three months after TAVI. Recent studies examined the global cognitive functions after TAVI, using MMSE and the Montreal Cognitive Assessment (MoCA)<sup>5,20,21</sup>. The MMSE, originally developed to screen for Alzheimer dementia, is currently widely used to assess post-stroke cognitive impairment<sup>22</sup>, although MMSE has been shown to lack sensitivity in the detection of very mild cognitive impairment<sup>22</sup>.

More recently, the MoCA has been developed to detect mild cognitive impairment with high sensitivity, which consists of seven cognitive domains, comprising orientation, attention, short-term memory, naming, visuospatial, language, and abstract reasoning<sup>5</sup>. LM II was developed specially to diagnose very mild cognitive impairment and episode memory<sup>12</sup>. In the present study, we used LM II instead of the MoCA for the following reasons. First, in a recent study, mean total MoCA score, especially short-term memory of the MoCA, was improved after TAVI<sup>5</sup>. Second, there was a significant improvement in the Immediate Recall Memory Test, with a trend towards an improved Delayed Recall



**Figure 2.** Changes in regional cerebral blood flow after TAVI and their associations with cognitive and cardiac functions. Glass brain representations showing TAVI-induced regional cerebral blood flow changes (black areas) from the coronal (A), axial (B), and sagittal (C) views ( $p < 0.005$  at each voxel). The coronal (D) and axial (E) slices including the right hippocampus are also presented. The green arrowheads indicate the right hippocampus. F) Local CBF was significantly increased after TAVI compared with baseline (baseline,  $51.2 \pm 1.0$  vs 3 months,  $53.3 \pm 1.0$  ml/100 g/min,  $W 60.0$ ,  $p < 0.0001$ ). G) Linear mixed-effects model showed that CBF in the right hippocampus was positively correlated with LM II scores. H) The patients with increased cardiac output (CO) after TAVI had significantly increased CBF in the right hippocampus compared with those without it. CBF: cerebral blood flow; CO: cardiac output; LM II: Logical Memory II; TAVI: transcatheter aortic valve implantation

Memory Test<sup>9</sup>. Third, LM II is a quantifiable neuropsychological test<sup>12</sup>. Taken together, it is possible that TAVI improves cognitive functions, especially LM II (episode memory), in the present study. In the present study, we had to exclude many patients, eventually analysing a relatively small number of patients, whose mean age was  $83.2 \pm 4.5$  years. A recent study has demonstrated that the risk and age of patients undergoing TAVI have become lower<sup>23</sup>. Thus, it remains to be elucidated whether TAVI improves cognitive function in younger patients with severe AS. Future studies with a large number of patients are needed to perform a multivariable analysis to adjust for possible factors contributing to the changes at follow-up.

#### ROLES OF INCREASED CEREBRAL BLOOD FLOW

Recent studies have shown that TAVI improves cognitive functions<sup>5,9,24</sup>. There were several hypotheses regarding the mechanisms of cognitive improvement after TAVI<sup>24-26</sup>. First, improvement of CBF due to improved CO after TAVI may contribute to the improvement of cognitive functions. Second, alleviation of physical symptoms and subsequent improvement in functional status may contribute to the improvement of cognitive functions. However, detailed mechanisms of the improvement of cognitive functions after TAVI remain to be examined.

Accelerated cognitive decline may result from chronic low cerebral perfusion in the long-term course of heart disease as



a pathophysiological consequence between the heart and brain interactions<sup>3</sup>. In the present study, TAVI significantly improved CO, local CBF especially in the right hippocampus, and LM II scores. Importantly, CO was associated with CBF in the right hippocampus, with a positive correlation with LM II scores. Thus, we were able to elucidate that TAVI increases CO and cerebral perfusion (especially that in the hippocampus) associated with improved cognitive functions, probably through the heart-brain interaction in elderly patients with severe AS.

Notably, we have recently demonstrated that whole-brain low-intensity pulsed ultrasound therapy markedly ameliorates cognitive impairment associated with improved CBF in mouse models of dementia, in which endothelial nitric oxide synthase (eNOS) activation plays a central role<sup>27</sup>. It is conceivable that increased CBF caused by upregulated eNOS may also be involved in the beneficial effects of TAVI.

### IMPORTANCE OF HIPPOCAMPUS FOR COGNITION

In the present study, although the whole CBF was not significantly increased, local CBF, especially that in the right hippocampus, was significantly increased after TAVI. Notably, we recently demonstrated that patients with chronic heart failure frequently have cognitive impairment, where the hippocampus blood flow is significantly decreased<sup>16</sup>. A possible mechanism of cognitive impairment in chronic heart failure is abnormality of the hippocampus, which is the important brain area for memory<sup>28</sup>. Moreover, the hippocampus is one of the brain regions most vulnerable to cerebral hypoxia<sup>29,30</sup>. Importantly, patients with obstructive sleep apnoea who underwent continuous positive airway pressure had improved cognitive function associated with improved grey matter volume in the hippocampus but not in the whole brain<sup>31</sup>. It is possible that the hippocampus is one of the watersheds and may be the first area where CBF reduction or improvement occurs. In the present study, although the whole CBF was not significantly increased, local CBF, including that in the hippocampus, was significantly increased after TAVI. Notably, in the present study, CBF in the local regions, not only in the right but also in the left hippocampus, was significantly increased after TAVI. The lack of statistical association between the left hippocampal blood flow and LM II scores may be due to the small sample size. Thus, it is possible that haemodynamic improvement by TAVI increases the perfusion in these regions, although the effect of cerebral hypoxia on brain abnormality in patients with severe AS remains to be elucidated.

### Study limitations

Several limitations should be mentioned in relation to the present study. First, this study was a single-centre study with a relatively small number of patients. A fragility index value of 1 for our study indicated that an outcome change in a single patient would make the difference in the main outcome non-significant. Thus, future studies with a large number of patients are needed to perform a multivariable analysis to adjust for possible factors contributing to the changes at follow-up. Second, the present study focused

on the abnormality of the hippocampus blood flow based on our previous study in rats<sup>30</sup>. However, substantial anatomical differences including the prefrontal cortex may exist between rats and humans. Third, there was a lack of control AS patients (without TAVI), although it is ethically and practically difficult to recruit such patients. Fourth, although we performed the commonly used tests for cognitive functions as previously reported<sup>5,9,12-14</sup>, we were unable to exclude a possible involvement of the learning effect. Future studies are needed to elucidate this effect. Fifth, the present study did not verify the cerebral structure and the CBF measurement using other modalities such as MRI. However, as mentioned above, we and others have already demonstrated that brain perfusion SPECT imaging is useful for solid assessment of quantitative cerebral perfusion and its relevance to cognitive impairment<sup>10,11</sup>.

### Conclusions

In the present study, we were able to demonstrate for the first time that TAVI may improve cognitive functions associated with increased cerebral perfusion especially in the hippocampus in elderly patients with severe AS.

### Impact on daily practice

Recent studies suggest that cognitive decline may result from chronic low cerebral perfusion in the long-term course of heart disease as a pathophysiological consequence between the heart and brain interactions. Based on the present study, TAVI may improve cognitive functions associated with increased cerebral perfusion especially in the hippocampus in elderly patients with severe AS.

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### Conflict of interest statement

The authors have no conflicts of interest to declare.

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## Supplementary data

**Supplementary Appendix 1.** TAVI procedure.

**Supplementary Appendix 2.** Echocardiography.

**Supplementary Appendix 3.** SPECT image pre-processing and analysis.

**Supplementary Figure 1.** Study flow chart.

**Supplementary Table 1.** Clinical characteristics of the included and excluded patients.

**Supplementary Table 2.** Baseline clinical characteristics of patients with severe AS.

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## **Supplementary data**

### **Supplementary Appendix 1. TAVI procedure**

In accordance with the Japanese Circulation Society guidelines, our Heart Team, including cardiologists, at least two cardiac surgeons, and anaesthesiologists, determined the indication and approach for the TAVI procedure, and the type of transcatheter valve used [32].

Periprocedural events were defined according to the Valve Academic Research Consortium (VARC)-2 criteria [33].

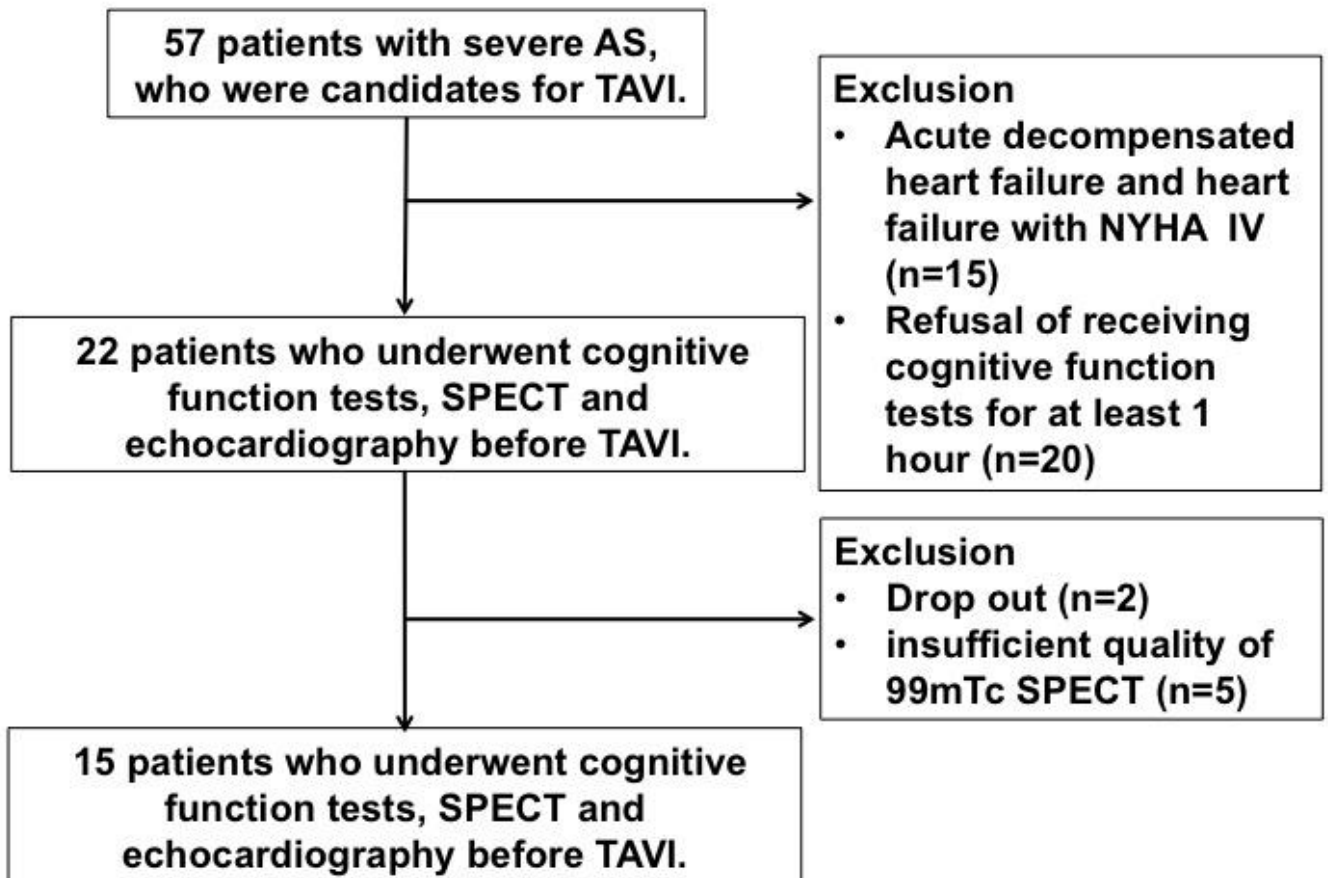
### **Supplementary Appendix 2. Echocardiography**

Before and three months after TAVI, echocardiography studies were performed by experienced sonographers blinded to the results of the cognitive tests and the imaging studies. Measurements were made from an average of three cardiac cycles in sinus rhythm and three to five cycles in atrial fibrillation. Assessments of the native AV and the bioprosthesis were made according to the original Valve Academic Research Consortium (VARC-1), American Society of Echocardiography, and European Association of Echocardiography [33,34]. Measurements of CO and left ventricular ejection fraction were performed according to the guidelines [35]. Carotid ultrasonography studies were performed by experienced sonographers who were blinded to the results of the cognitive tests and the imaging studies before TAVI.

### **Supplementary Appendix 3. SPECT image pre-processing and analysis**

The SPECT image pre-processing and analysis consisted of the following six steps: 1) co-registering CBF images at three months after TAVI to their corresponding baseline images before TAVI, 2) normalising these images to the standard Montreal Neurological Institute space, 3) smoothing the normalised images, 4) comparing these pre-processed baseline images with those at three months after TAVI, 5) calculating CBF within the brain regions derived from the step 4, and 6) testing an association of contrasts for cognitive test scores with those for CBF of step 5. Since all six steps do not require any arbitrary interventions from the analyst, there were no inter- or intra-observer variabilities.

## Supplementary Figure 1



**Supplementary Figure 1.** Study flow chart.

From January 2017 to September 2018, we examined 57 consecutive patients with severe AS at Tohoku University Hospital as candidates for TAVI. Inclusion criteria were 1) heart failure with New York Heart Association (NYHA) functional Class II to III symptoms, and 2) patient consent to receive cognitive function tests for at least one hour. Exclusion criteria were 1) acute decompensated heart failure and heart failure with NYHA Class IV symptoms, 2) refusal of cognitive tests, and 3) insufficient quality of 99mTc SPECT. Based on these criteria, we excluded 35 patients in advance, including acute decompensated heart failure and heart failure with NYHA Class IV symptoms in 15, and refusal to undergo cognitive function tests for at least one hour in 20. Thus, we initially included 22 patients, seven of whom were

then excluded because of dropout owing to refusal to undergo follow-up cognitive tests (n=2), and insufficient quality of  $^{99m}\text{Tc}$  SPECT (n=5). Finally, we examined 15 patients in the present study with special reference to the association of cerebral blood flow (CBF) and cognitive functions.

AS: aortic stenosis; NYHA: New York Heart Association; SPECT: single-photon emission computed tomography; TAVI: transcatheter aortic valve implantation

**Supplementary Table 1. Clinical characteristics of the included and excluded patients.**

	Included patients (n=15)	Excluded patients (n=7)	<i>p</i> -value
Age, years	83.2±4.5	84.0±6.1	0.41
Female, n (%)	12 (80)	6 (86)	0.74
Body mass index, kg/m <sup>2</sup>	22.8±2.2	21.4±2.1	0.19
Handedness - right-handed, n (%)	15 (100)	7 (100)	-
Hypertension, n (%)	14 (93)	6 (86)	0.57
Diabetes mellitus, n (%)	4 (27)	1 (14)	0.51
Dyslipidaemia, n (%)	12 (80)	5 (71)	0.66
Atrial fibrillation, n (%)	5 (33)	1 (14)	0.33
New York Heart Association functional			
Class II, n (%)	12 (80)	4 (57)	0.27
Class III, n (%)	3 (20)	3 (43)	0.27
Coronary artery disease, n (%)	3 (20)	4 (57)	0.086
Previous myocardial infarction, n (%)	0 (0)	1 (14)	0.12
Previous stroke/transient ischaemic attack, n (%)	4 (26.7)	0 (0)	0.063
Peripheral vascular disease, n (%)	2 (13)	1 (14)	0.95
Chronic obstructive pulmonary disease, n (%)	0 (0)	3 (43)	0.0048
Chronic kidney disease, n (%)	10 (67)	5 (71)	0.82
STS-PROM, %	6.7±3.1	8.7±3.7	0.19
MMSE <24	5 (33)	5 (71)	0.21
Echocardiographic characteristics			
Left ventricular ejection fraction, %	66.8±13.7	62.6±10.8	0.31
Peak velocity, m/s	4.62±0.74	4.98±0.98	0.50
Aortic mean gradient, mmHg	50.5±17.4	58.7±23.5	0.50



Aortic valve area, cm <sup>2</sup>	0.71±0.15	0.66±0.26	0.42
Cardiac output, l/min	4.03±0.88	3.59±1.28	0.31
Medical treatments, n (%)			
ACE-I or ARB, n (%)	9 (60)	7 (100)	0.018
Beta-blocker, n (%)	3 (20)	4 (57)	0.086
CCB, n (%)	8 (53)	3 (43)	0.64
Diuretic, n (%)	7 (47)	6 (86)	0.069
Statin, n (%)	10 (67)	2 (29)	0.092
Acetylcholine esterase inhibitors, n (%)	0 (0)	0 (0)	-
Cognitive function tests			
Mini-Mental State Examination	24.6±5.1	22.1±4.5	0.24
Geriatric Depression Scale	4.3±4.1	3.4±2.1	0.94
Logical Memory II	8.7±6.0	4.7±5.3	0.15

Categorical variables are expressed as n (%) and continuous variables as mean±standard deviation (SD).

ACE-I: angiotensin-converting enzyme inhibitor; ARB: angiotensin II receptor blockers; CCB: calcium channel blocker; MMSE: Mini-Mental State Examination; STS-PROM: Society of Thoracic Surgeons Predicted Risk of Mortality

**Supplementary Table 2. Baseline clinical characteristics of patients with severe AS.**

	Patients (n=15)
Age, years	83.2±4.5
Female, n (%)	12 (80)
Body mass index, kg/m <sup>2</sup>	22.8±2.2
Handedness - right-handed, n (%)	15 (100)
Hypertension, n (%)	14 (93)
Diabetes mellitus, n (%)	4 (27)
Dyslipidaemia, n (%)	12 (80)
Atrial fibrillation, n (%)	5 (33)
New York Heart Association functional	
Class II, n (%)	12 (80)
Class III, n (%)	3 (20)
Coronary artery disease, n (%)	3 (20)
Previous myocardial infarction, n (%)	0 (0)
Previous stroke/transient ischaemic attack, n (%)	4 (26.7)
Peripheral vascular disease, n (%)	2 (13)
Chronic obstructive pulmonary disease, n (%)	0 (0)
Chronic kidney disease, n (%)	10 (67)
STS-PROM, %	6.7±3.1
MMSE <24	5 (33)
Echocardiographic characteristics	
Left ventricular ejection fraction, %	66.8±13.7
Peak velocity, m/s	4.62±0.74
Aortic mean gradient, mmHg	50.5±17.4
Aortic valve area, cm <sup>2</sup>	0.71±0.15

Cardiac output, l/min	4.03±0.88
Medical treatments, n (%)	
ACE-I or ARB, n (%)	9 (60)
Beta-blocker, n (%)	3 (20)
CCB, n (%)	8 (53)
Diuretic, n (%)	7 (47)
Statin, n (%)	10 (67)
Acetylcholine esterase inhibitors, n (%)	0 (0)

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Categorical variables are expressed as n (%) and continuous variables as mean±standard deviation (SD).

ACE-I: angiotensin-converting enzyme inhibitor; ARB: angiotensin II receptor blockers; CCB: calcium channel blocker; MMSE: Mini-Mental State Examination; STS-PROM: Society of Thoracic Surgeons Predicted Risk of Mortality