

1 **Research Article**

2 **How Are FIM Gains Improved after Intensive Rehabilitation for**
3 **Cerebrovascular Diseases?**

4 **FIM gains after rehabilitation in cerebrovascular diseases**

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15 Abstract

16 *Objective:* Not only functional independence measure (FIM) gain after intensive
17 rehabilitation for cerebrovascular diseases but also the effect of dementia and higher brain
18 dysfunction to FIM gain are not well to be understood. The purpose of this study was to
19 determine FIM gains based on clinical factors after intensive rehabilitation for
20 cerebrovascular diseases. *Patients:* A number of 181 patients were suffered from stroke 161
21 (cerebral infarction 107, cerebral hemorrhage 41, subarachnoid hemorrhage 12, subdural
22 hematoma 1), traumatic brain injury 12, and others 8. *Methods:* Dementia, higher brain
23 dysfunction, etiology, sex, age, history of cerebrovascular diseases, and location involvement
24 were analyzed using FIM. *Results:* There was a significant relationship of the correlation
25 coefficient 0.772 between FIM-total gain 4 weeks after admission and FIM-total gain at
26 discharge. FIM-total gain (23.6 ± 18.4) in patients without dementia was significantly higher
27 than that (14.1 ± 12.4) of patients with dementia. FIM-total gain (23.5 ± 18.4) in patients
28 with higher brain dysfunction tended to be higher than that (18.5 ± 16.2) of patients without
29 higher brain dysfunction. Patients with FIM-total (41-80) on admission showed significantly
30 higher FIM-total gain of (28.6 ± 17.2) than that (18.7 ± 22.2) of patients with FIM-total (40
31 or lower), or than that (17.2 ± 8.69) of patients with FIM-total (81 or higher). FIM-total gains
32 were decreased according to increasing ages. FIM-total gain (29.3 ± 19.8) of cerebral
33 hemorrhage was significantly higher than that (19.2 ± 15.5) of cerebral infarction. FIM-total
34 gain decreased according to incidents of cerebrovascular diseases in the past. *Conclusion:*
35 FIM gain scores after rehabilitation in cerebrovascular diseases were correlated with no
36 dementia, higher brain dysfunction, younger ages, intermediate FIM score (41-80) on
37 admission, cerebral hemorrhage, and no history of cerebrovascular diseases. FIM-total gain
38 at discharge can be extrapolated from FIM-total gain 4 weeks after admission.

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40 **1. Introduction**

41
42 The Global Burden of Diseases, Injuries, and Risk Factors Study (GBD) 2017 reported that
43 stroke was the third-leading cause of death and disability combined and the second-leading
44 cause of death in the world in 2017 [1,2]. Stroke rehabilitation has therefore become the
45 most important treatment in caring for stroke patients [3]. These patients show improvement
46 after rehabilitation programs, however, the quality and rate of this improvement vary in these
47 patients. Stroke represents the most common cause of adult disability and the second major
48 cause of dementia. Significant functional recovery may develop in the first 3 months
49 following the episode [4]. Afterward, recovery is associated with cerebral plasticity and
50 cortical reorganization, in great part stimulated by rehabilitation programs [5]. The
51 functional independence measure (FIM) is the most widely used standardized outcome
52 measure for rehabilitation in the world. Most patients with severe stroke do not achieve FIM
53 motor ≥ 70 after inpatient rehabilitation [6]. Older patients and patients with lower admission
54 FIM motor require more attention [7].

55 In this article we aim to take a comprehensive look at not only the rehabilitation of stroke
56 patients but also of patients with other cerebrovascular diseases, traumatic brain injuries,
57 brain tumors, and meningoencephalitis. We determined the characteristic of FIM gain after
58 intensive rehabilitation for stroke, traumatic brain injury, brain tumor or meningoencephalitis.
59 We also evaluated the effect of dementia and higher brain dysfunction to FIM gain after
60 rehabilitation.

61 **2. Methods**

62 *2-1. Design*

63 The ethical approval of the study was obtained from Shimada Hospital Ethics Committee
64 (No.2208). Informed consents were obtained from the patients who participated in the study.
65 The research was conducted in accordance with the 2008 Helsinki Declaration of Human
66 Rights. It was a retrospective research project and conducted at a single institution. Patients
67 who suffered from strokes, traumatic brain injuries, brain tumors or meningoencephalitis and
68 received intensive rehabilitation by qualified physical therapists, qualified occupational
69 therapists and qualified speech-language-hearing therapists from May 2021 to October 2022
70 were included in this study.

71 *2-2. Participants*

72 All the patients with cerebrovascular diseases who were treated in the full-time treatment
 73 program of rehabilitation 7 days/wk from May 2021 to October 2022 were eligible for this
 74 research. Patients who were 78 years or younger underwent 3 hours of professional stroke
 75 rehabilitation per day, (physical therapy 1 hour, occupational therapy 1 hour, speaking
 76 therapy 1 hour). Patients who were 79 years or older underwent 2 hours of professional
 77 stroke rehabilitation per day (physical therapy 40 minutes, occupational therapy 40 minutes,
 78 speaking therapy 40 minutes). Patients who died from complications during their
 79 hospitalization, or whose FIM on discharge were lower than on admission, were excluded.
 80 Finally 181 patients were registered in the study (Table 1). Ninety patients were male and 91
 81 were female. Their mean age was 76.7 years old (range 40 to 102). There were 161 strokes
 82 (cerebral infarction 107, cerebral hemorrhage 41, subarachnoid hemorrhage 12, subdural
 83 hematoma 1), 12 traumatic brain injuries, 6 brain tumors and 2 meningoencephalitis.

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Table 1. Patients characteristics

Age		40 ~ 102 (Mean ± S.D. : 76.7 ± 12.0)			
Sex	Male	90	Past history of cerebrovascular disease	No incident	127
	Female	91		One incident	47
Etiology	Cerebral infarction	107	Side involvement	Two incidents	7
	Cerebral hemorrhage	41		Right-sided	75
	Subarachnoid hemorrhage	12		Left-sided	83
	Subdural hematoma	1		Not determined	23
Location involvement	Traumatic brain injury	12	Dementia	Positive	33
	Brain tumor	6		Negative	148
	Meningoencephalitis	2		Higher brain dysfunction	Positive
Location involvement	Over the tent	141	Discharge	Negative	59
	Under the tent	26		Home	127
	Not determined	14		Hospital, nstitution	54

87

88 2-3. Data collection

89 Clinical and demographic features including these etiology, sex, age, history of
 90 cerebrovascular diseases, side involvement, location involvement, disability severity,
 91 dementia and higher brain dysfunction were analyzed. Functional status was evaluated by
 92 using the functional independence measure (FIM) at admission. The FIM items are broadly

93 classified into total, motor and cognitive categories (FIM-total, FIM-motor, FIM-cognition).
94 FIM scores were assigned according to a 7-point scale, and the score indicated the amount of
95 assistance required to perform each item (7 = totally independent and 1 = totally dependent or
96 not testable). FIM scores were recorded at admission, 4 weeks, 8 weeks, and 12 weeks after
97 admission.

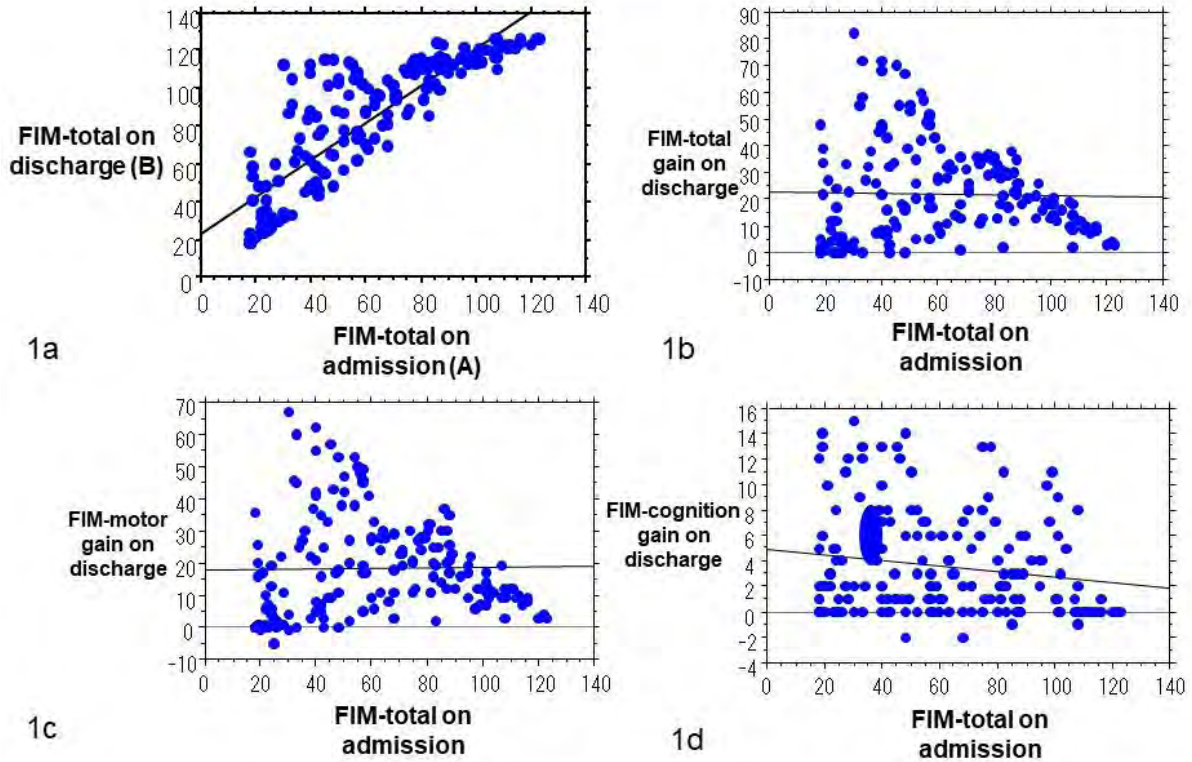
98 2-4. *Statistical Analysis*

99 The data is presented as the mean \pm standard deviation. A non-parametric test (Mann–Whitney
100 U test) was applied to compare the mean value of the two groups. The statistical analyses were
101 performed on StatView for Windows (Version 5.0; SAS Institute Inc. Cary, NC, USA). A *p*-
102 value of < 0.05 was defined as statistically significant.

103 3. Results

104 Length from onset to the start of rehabilitation were 0 to 41 days (2.2 ± 4.4). The
105 hospitalization at the first hospital that treated the patients in the acute phase was 6 to 114
106 days (27.3 ± 14.8). The hospitalization at our convalescent hospital for intensive
107 rehabilitation for patients was 10 to 178 days (75.0 ± 34.7).

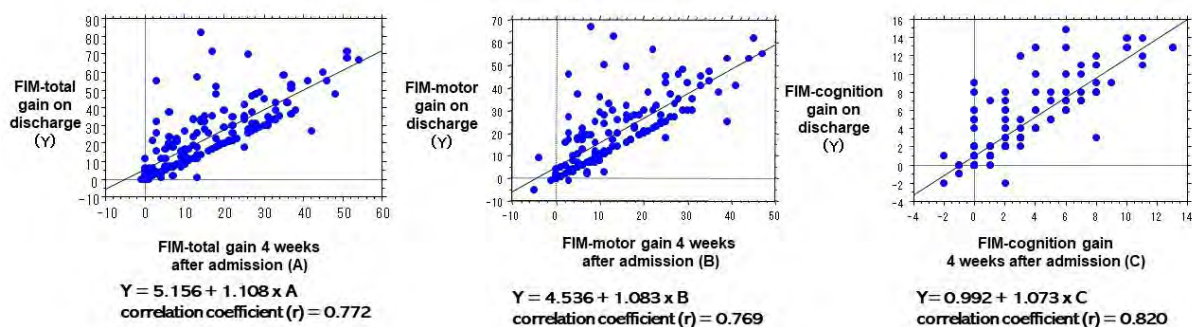
108 For 181 patients, FIM-motor gain was 18.3 ± 15.4 , FIM-cognitive gain was 3.5 ± 4.0 and
109 FIM-total gain was 21.8 ± 17.8 . There was a significant correlation between FIM-total on
110 admission (A) and FIM-total on discharge (B) ($B = 22.62 + 0.99 \times A$, the correlation
111 coefficient ($r = 0.857$) (Figure 1a). There were not any correlations between FIM-total on
112 admission and FIM-total gain, FIM-motor gain, or FIM-cognition gain on discharge (Figure 1
113 b-d).



114

115 **Figure 1a:** Relationships between FIM-total on admission (A) and FIM-total on discharge (B) There was a
 116 significant correlation between FIM-total on admission (A) and FIM-total on discharge (B): $B = 22.62 +$
 117 $0.99 \times A$, the correlation coefficient $r = 0.857$. **1b-1d:** Relationships between FIM-total on admission and FIM-
 118 total gain, FIM-motor gain, or FIM-cognition gain on discharge. There were not any correlations between them.
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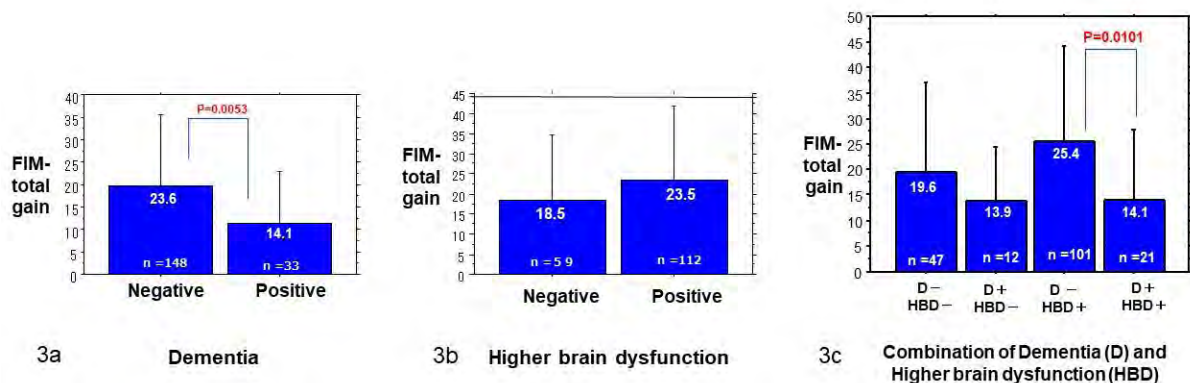
120 There were significant relationships of the correlation coefficient 0.772 between FIM-
 121 total gain 4 weeks after admission and FIM-total gain on discharge, the correlation
 122 coefficient 0.769 between FIM-motor gain 4 weeks after admission and FIM-motor gain on
 123 discharge, and the correlation coefficient 0.820 between FIM-cognition gain 4 weeks after
 124 admission and FIM-cognition gain on discharge (Figure 2). There were significant
 125 relationships between them.



126

127 **Figure 2:** Relationships between FIM gain 4 weeks after admission and FIM gain on discharge There were
 128 significant relationships between them.

129 For dementia, FIM-total gain (23.6 ± 18.4) in patients without dementia was significantly
 130 higher than that (14.1 ± 12.4) of patients with dementia ($p=0.0053$) (Figure 3a). FIM-motor
 131 gain (19.8 ± 15.8) in patients without dementia was significantly higher than that ($11.5 \pm$
 132 11.4) of patients with dementia ($p=0.0053$). FIM-cognition gain (3.8 ± 4.2) in patients
 133 without dementia tended to be higher than that (2.4 ± 2.7) of patients with dementia
 134 ($p=0.069$). Conversely, FIM-total gain (23.5 ± 18.4) in patients with higher brain dysfunction
 135 tended to be higher than that (18.5 ± 16.2) of patients without higher brain dysfunction
 136 ($p=0.076$) (Figure 3b). FIM-motor gain (19.4 ± 15.8) in patients with higher brain
 137 dysfunction tended to be higher than that (16.0 ± 14.5) of patients without higher brain
 138 dysfunction ($p=0.167$). FIM-cognition gain (4.0 ± 4.2) in patients with higher brain
 139 dysfunction were significantly higher than that (2.4 ± 3.4) of patients without higher brain
 140 dysfunction ($p=0.011$). FIM-total gain (25.4 ± 18.7) was highest in patients with dementia
 141 plus without higher brain dysfunction (Figure 3c). FIM-motor gain and FIM-cognition gain
 142 were highest in patients without dementia plus with higher brain dysfunction. FIM-total gain
 143 (13.9 ± 10.6) was lowest in patients with dementia plus without higher brain dysfunction.



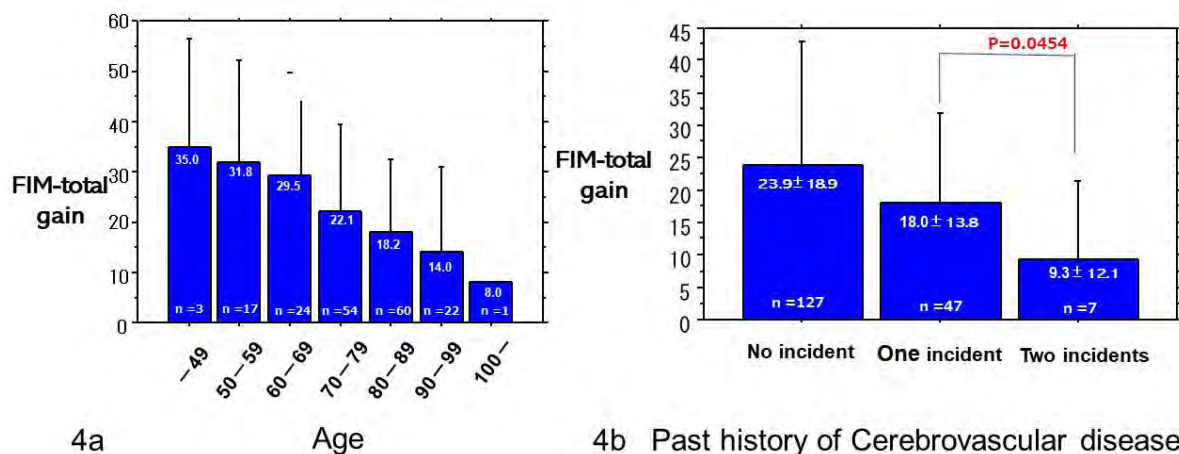
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145 **Figure 3a:** FIM-total gain in patients without dementia were significantly higher than that of patients with
 146 dementia. **3b:** FIM-total gain in patients with higher brain dysfunction was tended to be higher than that of
 147 patients without higher brain dysfunction. **3c:** FIM-total gain was lowest in patients with dementia plus
 148 without higher brain dysfunction. FIM -total gain was highest in patients without dementia plus with higher
 149 brain dysfunction.

150

151 For patients, length of stay in our convalescent rehabilitation ward varied: 10 to 178 days
 152 (75.0 ± 34.7). FIM-total scores were 60.3 ± 30.1 for admission ($n=181$). 75.0 ± 33.5 for 4
 153 weeks ($n=124$), 75.0 ± 33.5 for 8 weeks ($n=124$), and 70.0 ± 35.1 for 12 weeks ($n=69$). FIM-
 154 total gains were 15.2 ± 30.1 for 4 weeks ($n=179$), 22.9 ± 17.0 for 8 weeks ($n=124$), and
 155 26.1 ± 21.3 for 12 weeks ($n=69$). FIM-total gains significantly increased with the length of
 156 admission.

157 FIM-total gain (21.6 ± 18.6) of males were the same as that (22.1 ± 17.1) of females.
 158 FIM-total gains decreased in relation to an increase in age. FIM-total gains were 35.0 ± 21.7
 159 for patients of 49 years old or younger, 31.8 ± 20.4 for 50-59 years old, 29.5 ± 20.3 for 60-69
 160 years old, 22.1 ± 17.3 for 70-79 years old, 18.2 ± 14.3 for 80-89 years old, 14.0 ± 17.0 for 90-
 161 99 years old, and 8.0 for 100 years old or more (Figure 4a). FIM-motor gains decreased
 162 according to the increase in age as same as FIM-total gains. FIM-cognition gain (5.2 ± 5.1)
 163 was highest in patients of 60-69 years old. For past history, FIM-total gains were 23.9 ± 18.9
 164 in patients with no history of cerebrovascular disease, 18.0 ± 13.8 in patients with a history of
 165 one incident, and 9.3 ± 12.1 in patients with a history of two incidents (Figure 4b). FIM-total
 166 gain decreased according to the number of incidents of cerebrovascular diseases. FIM-total
 167 gain (22.6 ± 18.4) of left-side involvement tended to be higher than that (19.8 ± 16.1) of right-
 168 side involvement, but there was no significant difference. FIM-total gain (25.7 ± 20.8) of the
 169 involvement without the laterality tended to be higher than those of the involvement with the
 170 laterality. FIM-total gains (21.5 ± 18.2) in organs involved over the tent was as same as that
 171 (21.4 ± 14.6) in organs involved under the tent.

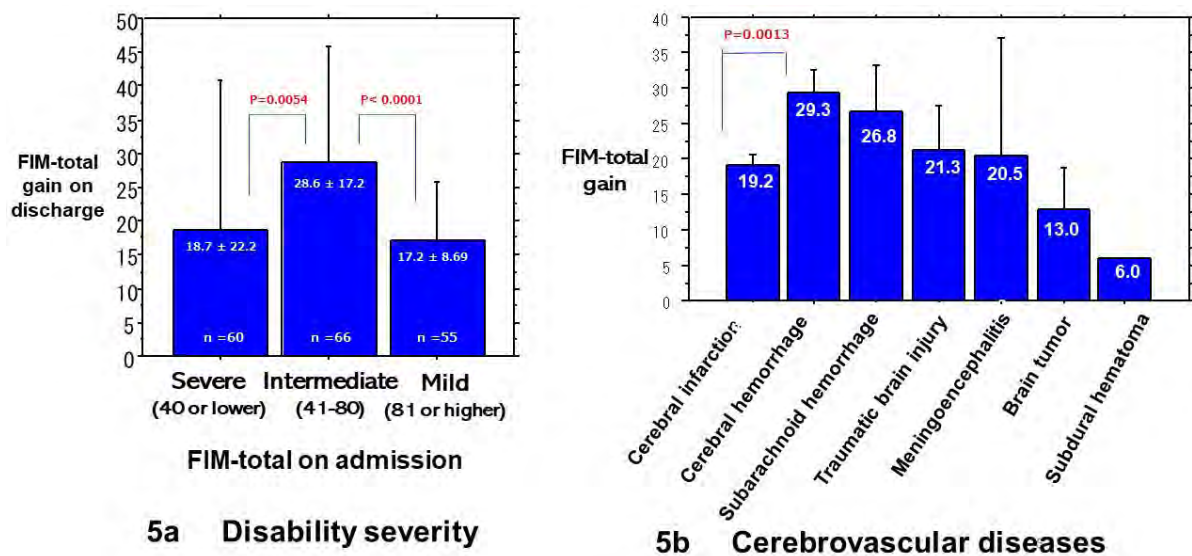


172 4a Age 4b Past history of Cerebrovascular diseases
 173 **Figure 4a:** FIM-total gains were decreased according to an increase in age. FIM-total gains were 35.0 ± 21.7 for
 174 patients of 49 years old or younger, 31.8 ± 20.4 for 50-59 years old, 29.5 ± 20.3 for 60-69 years old, 22.1 ± 17.3 for
 175 70-79 years old, 18.2 ± 14.3 for 80-89 years old, 14.0 ± 17.0 for 90-99 years old, and 8.0 of 100 years old or
 176 more. **4b:** The FIM-total gains decreased according to incidents of past history of cerebrovascular diseases.

177

178 For disability severity, FIM score on admission were classified in three groups: severe
 179 disability (FIM 40 or lower), intermediate disability (FIM 41-80) and mild disability (FIM 81
 180 or higher). Patients with intermediate disability (FIM 41-80) on admission showed
 181 significantly higher FIM-total gain of (28.6 ± 17.2) on discharge than that (18.7 ± 22.2) of
 182 patients with severe disability (FIM 40 or lower) on admission, or than that (17.2 ± 8.69) of
 183 patients with mild disability (FIM 81 or higher) on admission (Figure 5a). FIM-total gains

184 were 19.2 ± 15.5 for cerebral infarction, 29.3 ± 19.8 for cerebral hemorrhage, 26.8 ± 21.9 for
 185 subarachnoid hemorrhage, 21.3 ± 21.4 for traumatic brain injury, 20.5 ± 23.3 for
 186 meningoencephalitis, 13.0 ± 14.3 for brain tumor and 6.0 for subdural hematoma (Figure 5b).
 187 FIM-total gain of cerebral hemorrhage was significantly higher than that of cerebral
 188 infarction ($p=0.0013$). FIM-total gains tended to be higher in subarachnoid hemorrhage
 189 compared to cerebral infarction. FIM-motor gains were higher in cerebral hemorrhage and
 190 subarachnoid hemorrhage compared to cerebral infarction. FIM-cognition gains were
 191 significantly higher in meningoencephalitis compared with the other etiologies.
 192



193

194 **Figure 5a:** Relationships between disability severity (FIM-total on admission) and FIM-total gain on
 195 discharge. FIM-total (41-80) on admission was significantly higher than that of FIM-total (40 or
 196 lower) or FIM-total (81 or higher). **5b:** FIM-total gains were 19.2 ± 15.5 for cerebral infarction, 29.3
 197 ± 19.8 for cerebral hemorrhage, 26.8 ± 21.9 for subarachnoid hemorrhage, 21.3 ± 21.4 for intracranial
 198 trauma, 20.5 ± 23.3 for meningoencephalitis, 13.0 ± 14.3 for resected brain neoplasm and 6.0 for
 199 subdural hematoma.

200 4. Discussion

201 4-1. FIM gain 4 weeks after admission

202 We realized that improving the physical condition of patients is based on FIM gain 4 weeks
 203 after admission, not on the FIM score on admission. There were significant relationships of
 204 the correlation coefficient 0.772 between FIM-total gain 4 weeks after admission and FIM-
 205 total gain on discharge. FIM-total gain 4 weeks after admission was revealed to be a
 206 prognostic predictor of cerebrovascular diseases.

207 4-2. *Dementia and/or higher brain dysfunction*

208 FIM-total gain, FIM-motor gain and FIM-cognition gain in patients without dementia were
209 significantly higher than those of patients with dementia. Both total and motor FIM scores at
210 admission and discharge, and their respective FIM gain scores at discharge were higher in
211 non-pre-stroke dementia compared with pre-stroke dementia patients ($P < 0.001$) [7]. Despite
212 severe neurologic impairment(s) and disability, cognitively impaired stroke patients made
213 significant functional gains while undergoing rehabilitation and many can be regulated to
214 getting better at home [8]). The management of dementia can lead to better functional
215 recovery during rehabilitation [9]. Most of the cognitive improvement took place within 6
216 months [10]. Conversely, FIM-total gain, FIM-motor gain and FIM-cognition gain in
217 patients with higher brain dysfunction tended to be higher than those of patients without
218 higher brain dysfunction. Higher brain dysfunction refers to deficits in intellectual functions,
219 such as language, thinking, memory, behavior, learning, and concentration, resulting from
220 organic brain lesions. Damage to neuronal networks in the bilateral frontal and temporal lobe
221 appeared to play the most important role in higher brain dysfunction [11]. The results of
222 this study showed that patients with dementia had significantly reduced FIM gains, and
223 patients with higher brain dysfunction have a potential of cognition improvement and can
224 improve FIM gains by intensive rehabilitation. Post-stroke cognitive impairment is
225 associated with poorer outcomes and greater disability.

226 4-3. *Clinical factors*

227 The Intercollegiate Stroke Working Party (ICSWP) published the 5th edition of the *National*
228 *clinical guideline for stroke* in October 2016 [12]. It provides the most up to date and
229 comprehensive overview of the management of strokes available, covering the whole of the
230 pathway from acute care to longer term management. In our research, in the acute phase in
231 hospitals, early rehabilitation starts on 0 - 41 days (2.2 ± 4.4) after admission for patients with
232 cerebrovascular diseases. The patients are treated by an intensive rehabilitation program for
233 2-3 hours every day, 7 days/wk. ICSWP recommends that stroke rehabilitation should begin
234 24-48 hours after a stroke and treatment and care should be reviewed at six months [12].

235 In this article we determined the important factors which influence the functional
236 improvement of rehabilitation in cerebrovascular diseases: no dementia, higher brain
237 dysfunction, younger age, intermediate disability (FIM 41-80), cerebral hemorrhage, no
238 history of cerebrovascular diseases, and left-side involvement. Previous studies and
239 systematic reviews reported that age, marital status, time from stroke onset to rehabilitation,
240 aphasia, neglect, stroke severity presented in National Institutes of Health Stroke Scale,

241 cognitive function, and motor function such as walking distance were associated with the
242 gain score in FIM after stroke rehabilitation [13-15]. And, factors influencing the
243 rehabilitation results were age, cognitive impairment, unilateral spatial neglect, ADL before
244 the illness, and time to hospitalization from the onset [16]. The significant predictors for
245 “clinically significant functional gain” were also younger age <75 years old, higher Glasgow
246 Coma Scale score at admission, hemorrhagic stroke, intermediate FIM- motor measure (MM)
247 function group [17]. The important factors which influence the functional improvement of
248 rehabilitation in cerebrovascular diseases would be younger age, intermediate FIM score,
249 cerebral hemorrhage, no dementia and length from onset to the start of rehabilitation.

250 In our patients, mean FIM-motor gain was 18.3 ± 15.4 , mean FIM-cognitive gain was 3.5
251 ± 4.0 and mean FIM-total gain was 21.8 ± 17.8 . Our results correlated with the results from a
252 Shannon Janzen et al. experiment that stroke rehabilitation FIM-motor gain was 18.9 ± 14.0 ,
253 FIM-cognitive gain was 2.7 ± 3.6 and FIM-total gain was 21.7 ± 15.5 [18]. Turner-Stokes L
254 et al. also reported that after stroke rehabilitation FIM-motor gain was 18.4, FIM-cognitive
255 gain was 4.2 and FIM-total gain was 22.6 [19]. Functional improvement was observed in all
256 the groups in our study, but the improvement in the patients with intermediate disability (FIM
257 score 41–80) was significantly higher than that in patients with severe disability (FIM 40 or
258 lower) or mild disability (FIM 81 or higher). Inouye et al. divided their patients into 3 groups
259 according to FIM admission score and found that the patients who were moderately affected
260 at admission showed higher FIM gain levels than those who were severely affected [20].
261 Stroke patients in the intermediate FIM-motor function group generally had the best FIM-
262 motor gain [21].

263 FIM-total gain (21.6 ± 18.6) of males were the same as that (22.1 ± 17.1) of females.
264 FIM gain was decreased according to repeating cerebrovascular diseases. No history of
265 previous cerebrovascular diseases was correlated with improved FIM-total gain. FIM-total
266 gains decreased according to increases in age. FIM-total gains were 35.0 ± 21.7 for patients
267 of 49 years old or younger, and 14.0 ± 17.0 for 90-99 years old. FIM-cognition gain was
268 highest (5.2 ± 5.1) in patients of 60-69 years old. There was a report that the age group 65–
269 74 years old were obviously not yet too old to be labelled with limited rehabilitation
270 potential; this age group was associated with better functional gain when being compared
271 with the age group 75-84 years old and the age group >85 years old [17]. Younger ages were
272 associated with better functional gain in both univariate and multivariate analysis [22,23]. It
273 is common that FIM improvement is constant in younger patients under 69 years old, and
274 decreasing linearly in older patients over 70 years old [24].

275 FIM-total gain of left-side involvement tended to be higher than that of right-side
276 involvement. FIM-total gain of the location without the laterality tended to be higher than
277 that of locations with the laterality. FIM-total gains were same in organs involved under the
278 tent and organs involved over the tent. Patients with bilateral hemiparesis had significantly
279 greater FR (functional recovery) after stroke than patients with unilateral hemiparesis [25].
280 The magnitude of change was significantly higher for patients with bilateral hemiparesis [25].

281 FIM-total gain of cerebral hemorrhage was significantly higher than that of cerebral
282 infarction. Krishnan RR et al. reported that functional improvements after rehabilitation of
283 cerebral hemorrhage was better than that of cerebral infarction (FIM-hemorrhage Δ 27 vs.
284 FIM-infarction Δ 21, $p = 0.05$), despite significantly worse initial stroke severity [26].
285 However, there is evidence that recovery was not limited to this time period alone. In the
286 past, the observation of spontaneous recovery after stroke has misled some authors to believe
287 that recovery of upper extremity function is intrinsic and that little can be done by therapists
288 to influence it.

289

290 **5. Limitations of the study**

291 We must keep in mind that this research has two limitations. First, it was a retrospective
292 research project and was conducted at a single institution. Second, the number of patients of
293 cerebrovascular diseases were limited to only 181. For a more accurate assessment,
294 additional cases of cerebrovascular diseases and a longer follow-up study are necessary.

295

296 **6. Conclusions**

297 Improved FIM gain scores after stroke rehabilitation in cerebrovascular diseases were
298 correlated with no dementia, higher brain dysfunction, younger ages. intermediate FIM score
299 (41-80), cerebral hemorrhage, no history of cerebrovascular diseases, left-side involvement.
300 We can extrapolate FIM-total gain at discharge from FIM-total gain 4 weeks after admission.

301

302 **Data Availability**

303 Data include sensitive and personal information and will not be shared as supplemental or
304 underlying data.

305 **Conflicts of interest**

306 The authors have no conflicts of interest to declare.

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309

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315

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