#### **ORIGINAL ARTICLE**



# Communication-type smartphone application can contribute to reducing elapsed time to reperfusion therapy

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#### **Abstract**

**Background and objectives** Communication-type medical smartphone applications enable text, neuroimaging, photos, and videos to be shared securely among the stroke team. Our aim was to investigate whether use of a smartphone application would shorten the duration from admission to reperfusion therapy in patients with hyper-acute ischemic stroke.

**Methods** Enrolled were acute ischemic stroke patients who underwent reperfusion therapy (intravenous tissue plasminogen activator (IV t-PA) and mechanical thrombectomy (MT)) at our hospital between October 2012 and September 2018. We divided the patients into two groups based on the date of availability on smartphones of communication-type medical application: (1) Control group, conventional communication prior to September 2015, and (2) App group, communication via the smartphone app from October 2015 onwards. We compared door-to-image time (DIT), image-to-needle time (INT), door-to-needle time (DNT) for thrombolysis, and DIT, image-to-puncture time (IPT), and door-to-puncture time (DTP) for thrombectomy between the groups. **Results** We retrospectively enrolled 139 patients (68% male; median age, 69 years; median NIHSS score, 7) who were assigned into the App group (n = 86) and Control group (n = 53). Of the overall patients, 109 underwent IV t-PA (IV t-PA alone, 79 patients), and 63 underwent MT (MT alone, 30 patients), and 33 patients underwent combined IV t-PA and MT. There was no significant difference in DIT between the App and Control groups (23 min vs. 22 min, p = 0.493). DNT, DPT, INT, and IPT were significantly shorter in the App group than in the Control group (DNT, 62 min for the App group vs. 72 min for Control group, p = 0.038; INT, 42 vs. 48 min, p = 0.009; DPT, 106 vs. 129 min, p = 0.046; IPT, 89 vs. 117 min, p = 0.004).

**Conclusion** The present findings indicate that communication-type medical smartphone apps have potential for shortening the time elapsed between admission and reperfusion therapy, especially INT and IPT.

**Keywords** Smartphone application · Reperfusion therapy

## Introduction

The effectiveness of administration of intravenous tissue plasminogen activator (IV-tPA) and mechanical thrombectomy

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(MT) has been proven for treatment of patients with acute ischemic stroke [1–5]. A meta-analysis found that the benefits are highly time dependent: earlier IV-tPA and MT are associated with reduced disability, mortality, and risk of complications such as symptomatic intracranial hemorrhage [6–10]. Therefore, after the decision has been made to administer this therapy, the stroke team must initiate the treatment as soon as possible.

At hospitals where reperfusion therapy is performed, it is extremely important to establish a stroke care system to shorten the time taken at each of the steps between onset and reperfusion, i.e., from admission to imaging, imaging to puncture, and puncture to recanalization. The Target: Stroke initiative [7] developed a stroke algorithm system designed to reduce the time taken until treatment that codes single-call activation systems, stroke tools, rapid brain imaging, and laboratory testing.



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Recently, social networks using smart device such as smartphones and tablet terminal have become widespread and indispensable tool for patients and neurologists [11, 12]. Furthermore, it has been suggested that this digitization process may be accelerated in the stroke care after a global emergency represented by the COVID-19 pandemic [13]. Already, several stroke centers have trialed a medical information and communication technology (ICT) smartphone applications [14, 15] with the aim of shortening IV-tPA and MT times from onset. These applications perform task management and time tracking for reperfusion therapy, and present this information visually for sharing among the stroke team.

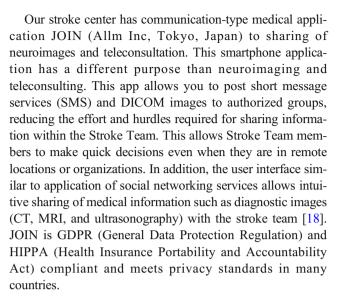
In the present study, we used a communication-type medical ICT smartphone application to build a seamless system for information sharing within the stroke team, similar to a social networking service. Within the application, the team can share texts, neuroimaging, photos, and videos with high security. This information be referred to independently at any time by each member of the specialized stroke team, whether they are physically at the hospital or not, for decision making regarding reperfusion therapy. Some previous studies have reported a reduction in reperfusion therapy time by using a smartphone application as a communication tool, but the applications did not incorporate a time tracker. The aim of the present study was to evaluate the ability of a communication-type medical application to reduce the time from hospitalization to reperfusion therapy, and thus improve patients' functional outcomes.

## **Methods**

The authors declare that all supporting data are available within the article.

## Study design

This retrospective study included consecutive ischemic stroke patients with acute ischemic stroke who underwent thrombolysis and/or mechanical thrombectomy between October 2012 and September 2018. Patients' clinical information, including cardiovascular risk factors (hypertension, diabetes mellitus, dyslipidemia, smoking status) and past history of atrial fibrillation, was obtained from the Jikei University School of Medicine Stroke Registry (Jikei Stroke Registry), a prospective database of patients with acute stroke admitted to Jikei University Hospital, a tertiary medical center located in the center of the Tokyo metropolitan area. Stroke severity on admission was assessed using the National Institutes of Health Stroke Scale (NIHSS) score [16]. Using the criteria of the Trial of ORG 10172 in Acute Stroke Treatment (TOAST) [17], stroke subtype was categorized into five groups: smallvessel occlusion, large-artery atherosclerosis, cardioembolism, other determined etiology, and undetermined.



This study conformed to the ethical principles established in the Declaration of Helsinki, and the Institutional Review Board of Jikei University School of Medicine, Japan, approved the study protocol (No. 8813). The board waived the need for patient consent.

# Statistical analysis

We divided the patients into two groups based on the date of availability on smartphones of communication-type medical application: (1) Control group, conventional communication prior to September 2015, and (2) App group, communication via the smartphone application from October 2015 onwards. We compared door-to-image time (DIT), image-to-needle time (INT), door-to-needle time (DNT) for thrombolysis, and DIT, image-to-puncture time (IPT), and door-topuncture time (DTP) for thrombectomy between the groups. Continuous variables were summarized using medians (interquartile range [IQR]) and categorical variables were expressed as counts (%). We compared the clinical backgrounds, elapsed time (DIT, INT, DNT, IPT, DPT), efficacy, and safety outcomes (recanalization rate at 24 h after reperfusion therapy, symptomatic intracranial hemorrhage at 24 h, neurologic deterioration at 24 h, modified Rankin scale [mRS] at discharge and 90 days, mortality at 90 days) between the two groups using the chi-square test or Mann-Whitney U test, as appropriate. Symptomatic intracranial hemorrhage was defined according to the European Cooperative Acute Stroke Study III criteria [2] as the presence of extravascular blood in the cranium that was associated with an increase in the NIHSS score of  $\geq$  4 points or with death, and judged to be the predominant cause of neurologic deterioration. Recanalization was defined as modified MORI grade > 2 on MR angiography [19]. Neurologic deterioration was defined as an increase in NIHSS score of  $\geq 4$  points within 24 h after stroke that was not attributed to intracranial hemorrhage or malignant cerebral



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edema. A two-tailed P < 0.05 was considered statistically significant. All statistical analyses were performed using SPSS for Windows version 25.0 (SPSS, Chicago, IL).

# Results

# **Clinical background**

We retrospectively enrolled 139 consecutive patients (68% male; median age, 69 years; median NIHSS score, 7). Table 1 lists patients' clinical characteristics. Of the 139 patients, 86 were assigned into the App group and 53 were assigned into the Control group. Of all patients, 109 underwent IV t-PA, of which 76 underwent IV t-PA alone; 63 underwent MT, of which 30 underwent MT alone; and 33 patients underwent combined IV t-PA and MT.

### IV-tPA after admission

Sixty-four patients in the App group and 45 patients in the Control group were treated with IV-tPA. There was no significant difference in DIT between the App and Control groups (23 min vs. 22 min, p = 0.493). In contrast, INT and DNT were significantly shorter in the App group than the Control group (INT, 42 min for the App group vs. 48 min for the

Control group, p = 0.006; DNT, 62 vs. 72 min, p = 0.038, Fig. 1).

# Mechanical thrombectomy after admission

Forty-two patients in the App group and 21 patients in the Control group underwent mechanical thrombectomy. There was no significant difference in terms of DIT between the two groups (22 vs. 23 min, p = 0.493). However, IPT and DPT were significantly shorter in the App group than in the Control group (IPT, 75 vs. 94 min, p = 0.002; DPT, 100 vs. 129 min, p = 0.022; Fig. 2).

# Efficacy and safety outcomes

Table 2 shows the efficacy and safety outcomes for the two groups. There was no statistically significant difference between the App and Control groups from onset to 24 h after onset in terms of recanalization rate, symptomatic intracranial hemorrhage, or neurologic deterioration (recanalization, 66% vs. 72%, p = 0.53; symptomatic intracranial hemorrhage, 1% vs. 4%, p = 0.264; neurologic deterioration, 9% vs. 15%, p = 0.216). In terms of outcome, there was no difference in mRS at discharge and 90 days between the App and Control groups (mRS at discharge, 2 vs. 2, p = 0.215; mRS at 90 days, 1 vs. 2, p = 0.554). Of note, mortality at 90 days was lower in the App

Table 1 Patient characteristics

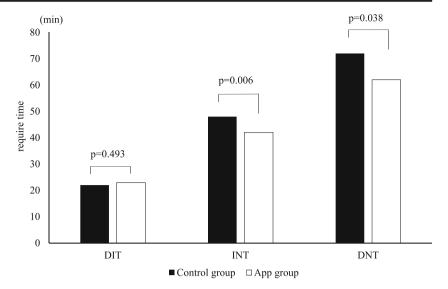
	App group $(n = 86)$	Control group $(n = 53)$	p
Age (years), median (IQR)	66 (54–79)	71 (64–78)	0.184
Male, <i>n</i> , %	57 (66)	37 (70)	0.405
Past history			
Hypertention, <i>n</i> , %	52 (61)	38 (72)	0.122
Hyperlipidemia, n, %	28 (33)	26 (49)	0.04
Diabetes melllitus, n, %	20 (23)	12 (23)	0.553
Atrial fibrillation, n, %	19 (22)	14 (26)	0.351
Prestroke mRS, median (IQR)	0 (0–1)	0 (0-0)	0.062
NIHSS score on admission, median (IQR)	6 (3–15)	9 (5–18)	0.027
TOAST classification, n, %			< 0.01
Large-artery atherosclerosis, n, %	8 (9)	15 (28)	
Small-vessel occlusion, n, %	4 (5)	2 (4)	
Cardioembolism, n, %	33 (38)	25 (47)	
Other determined ethiology, n, %	3 (4)	5 (10)	
Undetermined, n, %	28 (33)	6 (11)	
Major artery occlusion, n, %	62 (72)	43 (81)	0.692
Treatment			0.332
tPA, n, %	44 (51)	32 (60)	
Mechanical thrombectomy, n, %	22 (26)	8 (15)	
tPA+mechanical thrombectomy, $n$ , %	20 (23)	13 (25)	

mRS modified Rankin scale, NIHSS national institute health stroke scale, TOAST Trial of ORG 10172 in Acute Stroke Treatment



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Fig. 1 Time from admission to initiation of IV-tPA. INT and DNT were significantly shorter in the App group than the Control group. DIT door to image time, INT image to needle time, DNT door to needle time



group than the Control group (2% vs. 13%, p = 0.014). There was no difference between the groups in terms of mRS at 90 days in patients who underwent tPA alone. However, the percentage of good outcome at 90 days showed a tendency to be higher in patients who underwent MT alone, IV t-PA with MT, or MT with or without IV t-PA (supplementary 1).

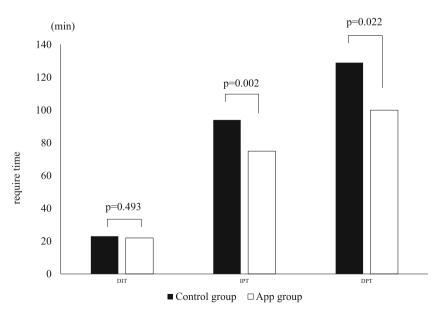
## Discussion

The results of the present study showed shorter DNT and DPT values for patients in the App group. In particular, treatment was most rapid for patients in this group who underwent MT. In addition, there were no difference DPT between the App group and Control group, but INT and IPT was shorter in the App group. The mRS at discharge and 90 days was similar

between the groups, although mortality at 90 days was lower in the App group.

Our study found that use of the smartphone app reduced the DNT and DPT values. Specifically, shortening INT and IPT contributed to shortening DNT and DPT. A previous study has reported that use of a medical acute care coordination app decreased mean DNT by 40 min [20, 21]; however, they did not investigate the cause of shortened DNT in detail. We consider that DNT and DPT were shorter with use of the smartphone app for the following reasons. Cooperation within the professional stroke treatment team is very important in reperfusion therapy. After rapid medical examination and neuroimaging have been performed, the stroke patient is transported to the ward, the operating room/angiography room is prepared, and the cerebrovascular specialist, anesthesiologist, and ICU/SCU physicians are contacted. Arranging each of these steps conventionally by telephone is time intensive; in

Fig. 2 Time from admission to initiation of mechanical thrombectomy. IPT and DRT were significantly shorter in the App group than the Control group. DIT door to image time, INT image to needle time, DNT door to needle time





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**Table 2** Efficacy and safety outcomes in the Smartphone app and Control groups

	App group $(n = 86)$	Control group $(n = 53)$	p
Recanalization at 24 h*, n, %	57 (66)	38 (72)	0.53
Symptomatic intracranial hemorrhage at 24 h**, n, %	1 (1)	2 (4)	0.264
Neurologic deterioration at 24 h***, n, %	8 (9)	8 (15)	0.216
discharge mRS	2 (1–4)	2 (1–4)	0.215
90 days mRS	1 (0-4)	2 (0-4)	0.554
90 days mRS < 2	55 (65)	32 (63)	0.480
90 days mortality, $n$ , %	2 (2)	7 (13)	0.014

<sup>\*</sup>Recanalization was defined as modified MORI grade > 2 with MR angiography

mRS modified Rankin Scale

comparison, communication via a smartphone app using a server-based network can instantly share patient medical information among the stroke team, including text and neuroimaging, in one simple action. Because it is available immediately after sharing, medical information can be evaluated via common chat room interfaces, thereby facilitating real-time discussions by the team. Task manager-type and time tracker-type apps cannot perform the function of communication among the stroke team. We consider that using the smartphone app saved process and communication time in coordinating reperfusion therapy, which requires collaboration of an interdisciplinary team.

Use of the smartphone app shortened DNT and DPT but was not related to favorable outcome at 3 months, for the reason that our cohort contained many patients with non-severe stroke. The favorable outcome rate was very high in both of the present groups compared with those reported previously [9, 22]. Previous studies have reported that in-hospital medical apps [15, 20], telestroke evaluation [23], and treatment by a mobile stroke unit [23, 24] reduced DNT, but to the best of our knowledge, none studied the relationship between favorable outcome and improvements in the stroke care system.

There are some limitations of the present study. We speculate that our results may have been affected by its small sample size, single-center design, retrospective cohort, and efforts to reduce time for IV t-PA and MT for code stroke system. Further large-scale trials are required to confirm the effect of the smartphone app on reducing time to initiation of reperfusion therapy.

# Conclusion

Use of a communication-type medical smartphone app reduced the time from admission to initiation of reperfusion

therapy compared with the conventional communication method. The time saving effect was higher for MT than for IV t-PA.

**Supplementary Information** The online version contains supplementary material available at https://doi.org/10.1007/s10072-021-05132-2.

#### **Declarations**

**Ethical approval** This article does not contain animals performed by any of the authors. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional research ethics committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

**Informed consent** Informed consent was obtained from all individual participants included in the study.

**Conflict of interest** The authors declare that they have no conflict of interest.

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<sup>\*\*</sup>Symptomatic intracranial hemorrhage was defined according to European Cooperative Acute Stroke Study III criteria as the presence of extravascular blood in the cranium that was associated with an increase in the NIHSS score of 4 points or more or death and was judged to be the predominant cause of neurologic deterioration

<sup>\*\*\*</sup>Neurologic deterioration was defined as an increase in the NIHSS score of 4 or more points within 24 h after stroke that was not attributed to intracranial hemorrhage or malignant cerebral edema

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