Endovascular Therapy for Concurrent Cardio-Cerebral Infarction in a Patient With Trousseau Syndrome

Kenichi Sakuta 1*, Taiji Mukai 1, Asako Fujii 2, Kentaro Makita 2 and Hiroshi Yaguchi 1

1 Department of Neurology, The Jikei University School of Medicine, Kashiwa Hospital, Kashiwa, Japan, 2 Department of Cardiology, The Jikei University School of Medicine, Kashiwa Hospital, Kashiwa, Japan

Only a few patients have been reported to undergo endovascular therapy for Trousseau syndrome. This is the first report of a patient with Trousseau syndrome who developed synchronous cardiocerebral infarction and underwent endovascular therapy for both. A 55-year-old woman with Trousseau syndrome arising from stage IV ovarian cancer presented with consciousness disturbance, aphasia, and right hemiparesis. Magnetic resonance imaging showed acute cerebral infarction limited to the left basal ganglia and occlusion of the left middle cerebral artery (MCA). Electrocardiography showed ST elevation in leads II, III, and aVF with reciprocal change. Mild elevation of myocardial enzymes was observed in laboratory data. She was diagnosed with synchronous cardiocerebral infarction. Both infarctions were considered as appropriately indicated for endovascular therapy. Since her vital signs were stable, a decision was made to treat the cerebral infarction first. Thrombectomy with a stent retriever was performed, which achieved complete recanalization of the left MCA. Percutaneous coronary intervention successfully recanalized the occluded right coronary artery. She suffered no recurrence of stroke or acute coronary syndrome upon heparin administration. Cardiocerebral infarction caused by Trousseau syndrome is rare and demands optimal planning of endovascular therapy.

Keywords: cardiocerebral infarction, Trousseau syndrome, endovascular therapy, myocardial infarction, ischemic stroke

INTRODUCTION

Trousseau syndrome is a condition characterized as hypercoagulation disorder related to malignancy (1). It is considered a common cause of ischemic stroke in terminally ill patients, and several affected patients have been reported to undergo mechanical thrombectomy (2–4). Here, we report a rare case of synchronous cardiocerebral infarction (CCI) caused by Trousseau syndrome in a patient who underwent endovascular therapy to treat both infarctions.

CASE REPORT

A 55-year-old woman with a history of ovarian carcinoma and multiple metastasis to lymph node, liver, and bone (Figure 1) was admitted to our hospital with consciousness disturbance. Three months prior to admission, she had suffered deep vein thrombosis and pulmonary embolism attributable to Trousseau syndrome and was prescribed direct oral anticoagulants (DOACs).
She was expected to survive at least 6 months if chemotherapy could be continued, and Karnofsky Performance Status (KPS) was 90. On examination, her Glasgow Coma Scale score was 10 (E3, V1, M6), body temperature 36.0°C, blood pressure 133/78 mmHg, and heart rate 100 beats/min. Neurological examination revealed motor aphasia, right hemiparesis, and conjugate deviation of eyes to the left. The National Institutes of Health Stroke Scale score was 23. She reported no chest pain. Blood tests showed mild elevation of myocardial marker enzymes such as aspartate aminotransferase (45 U/L), lactate dehydrogenase (425 U/L), creatine kinase (777 U/L), creatine kinase MB (35 U/L), and troponin T (0.21 ng/mL; normal, <0.05 ng/mL). Tumor makers were elevated, especially CA-125 (190 U/mL; normal, <35 U/mL), and D-dimer was 11.4 µg/mL on admission, which gradually increased despite DOAC continuation. We performed brain magnetic resonance (MR) imaging and MR angiography immediately after admission. Acute cerebral infarction was detected in the left basal ganglia, and the left middle cerebral artery (MCA) was occluded (Figures 2A,B). Electrocardiography on admission revealed normal sinus rhythm with ST-segment elevation in leads II, III, and aVF, with reciprocal change in leads aVL and V2–V6. Bedside transthoracic echocardiography revealed almost normal wall motion without evidence of intracardiac thrombus, aortic dissection, or vegetation. These results led us to the diagnosis of synchronous CCI.

Both infarcts were considered suitably indicated for endovascular therapy, although tissue plasminogen activator was a contraindication because the time of onset was unknown. Since the vital signs were relatively stable, a decision was made to treat the MCA occlusion before the coronary artery occlusion. A cerebral angiogram showed distal MCA occlusion (Figure 2D, arrow), and the thrombus was successfully retrieved with a stent retriever (Figure 2D, arrowhead). The white-colored thrombus was collected. Since the awareness of the patient was delayed, time from last known well to recanalization was 19 h. Percutaneous coronary intervention (PCI) was subsequently performed using the same femoral sheath. A coronary angiogram showed complete occlusion of the distal right coronary artery at the atrioventricular branch (Figure 2C), and after plain old balloon angioplasty the occluded artery was recanalized (Figure 2F). After the intervention, ST elevation on electrocardiography was normalized. Since treatment with heparin injection for secondary stroke prevention was scheduled, stent placement was not performed as this requires dual-antiplatelet therapy, raising the fear of bleeding complications. After the procedures the patient was started on heparin injection instead of DOACs. Brain MRA on the second hospital day showed a recanalized left MCA (Figure 2E). The patient achieved a functional status of modified Rankin score 3 and KPS 70 at 3-month follow-up with no recurrence of infarctions under continuation of heparin subcutaneous injection. Such a good recovery made it possible to restart her chemotherapy.

**DISCUSSION**

The unique point of the present case is that simultaneous CCI occurred in a patient with Trousseau syndrome. Furthermore,
both occluded arteries were successfully recanalized with endovascular therapy.

This is the first report of dual endovascular intervention in a CCI patient with Trousseau syndrome. Endovascular therapy for cerebral infarction in Trousseau syndrome is rarely reported (2–4). All of these previous cases exhibited terminal stage of cancer, elevated D-dimer, and thrombus with white coloration consisting of fibrin and platelets. The present case is consistent with these characteristics. In cases where the prognosis is unfavorable because of terminal-stage cancer (3, 4), it is sometimes difficult to make a decision to undertake invasive surgery in patients in such poor condition. Recent article regarding endovascular therapy for cancer-related stroke revealed that the rate of good outcome at discharge was not significantly differ compared to those without cancer (5). In the present case we readily decided to perform surgery because the patient was expected to survive another 6 months in the absence of infarction.

Acute CCI is defined as synchronous infarction of the brain and heart, and is a life-threatening condition that requires immediate medical intervention (6). Frequency is reported as 1.6% in a cohort of acute ischemic stroke patients and 0.9% in a cohort of acute coronary syndrome patients (7, 8). In the case of CCI with indications for both endovascular therapies, no evidence-based treatment plan has been established. Some authors recommend administration of intravenous tissue plasminogen activator of stroke dose, followed by PCI (6, 9). Thrombectomy for cerebral infarction takes precedence over PCI only when there is basilar occlusion or non-ST elevation myocardial infarction (9). In the present case cerebral infarction of the anterior circulation was treated first, followed by the ST-elevation myocardial infarction. The reason why the outcome was favorable in this case may be because hemodynamics were maintained and arrhythmia did not develop despite the inferior wall infarction.

**CONCLUSION**

Trousseau syndrome rarely causes major vessel occlusion and in the case of CCI, hemodynamics stability may play an important role in determining the appropriate ordering of treatment.

**DATA AVAILABILITY**

All data containing relevant information to support the study findings are included in the manuscript.

**ETHICS STATEMENT**

The authors declare that ethics approval was not required for this case report. Written informed consent was obtained from the patient for publication of this case report and any accompanying data.

**AUTHOR CONTRIBUTIONS**

KS drafted the manuscript and prepared the figures. TM designed the study and analyzed the clinical data. AF collected the clinical data and interpreted the data. KM collected the clinical data and interpreted the data. HY helped to write and revise the manuscript. All authors have read and approved the final manuscript.
FUNDING

The authors received no financial support for the research, authorship, and/or publication of this article.

ACKNOWLEDGMENTS

We thank Hugh McGonigle, from Edanz Group (www.edanzediting.com/ac), for editing a draft of the manuscript.

REFERENCES

1. Ikushima S, Ono R, Fukuda K, Sakayori M, Awano N, Kondo K. Trousseau's syndrome: cancer-associated thrombosis. Jpn J Clin Oncol. (2016) 46:204–8. doi: 10.1093/jjco/hvy165

2. Matsumoto N, Fukuda H, Handa A, Kawasaki T, Kurosaki Y, Chin M, et al. Histological examination of trousseau syndrome-related thrombus retrieved through acute endovascular thrombectomy: report of 2 cases. J Stroke Cerebrovasc Dis. (2016) 25:227–30. doi: 10.1016/j.jstrokecerebrovasdis.2016.08.041

3. Shinohara T, Tsumura M, Hasegawa A, Yamanaka Y. A case of mechanical thrombectomy for middle cerebral artery occlusion with trousseau syndrome. J Neuroendovasc Ther. (2017) 11:485–91. doi: 10.5797/jnet.cr.2016-0133

4. Kuroda N, Hiramatsu H, Mori M, Tanaka T. Mechanical thrombectomy for trousseau syndrome in a terminally ill cancer patient. J Pain Symptom Manage. (2019) 57:688–94. doi: 10.1016/j.jpainsymman.2018.12.327

5. Jung S, Jung C, Hyoung Kim J, Se Choi B, Jung Bae Y, Sunwoo L, et al. Procedural and clinical outcomes of endovascular recanalization therapy in patients with cancer-related stroke. Interv Neurol Radiol. (2018) 24:520–8. doi: 10.1177/1591019918776207

6. Akinseye OA, Shahreyar M, Heckle MR, Khouzam RN. Simultaneous acute cardio-cerebral infarction: is there a consensus for management? Ann Transl Med. (2018) 6:7. doi: 10.21037/atm.2017.11.06

7. Alqahtani F, Aljohani S, Tarabishy A, Busu T, Adcock A, Alkhoulil M. Incidence and outcomes of myocardial infarction in patients admitted with acute ischemic stroke. Stroke. (2017) 48:2931–8. doi: 10.1161/STROKEAHA.117.018408

8. Budaj A, Flasinska K, Gore JM, Anderson FA Jr, Dabbous OH, Spencer FA et al. Magnitude of and risk factors for in-hospital and postdischarge stroke in patients with acute coronary syndromes: findings from a Global Registry of Acute Coronary Events. Circulation. (2005) 111:3242–7. doi: 10.1161/CIRCULATIONAHA.104.512806

9. Yeo LLL, Andersson T, Yee KW, Tan BYQ, Paliwal P, Gopinathan A, et al. Synchronous cardiocerebral infarction in the era of endovascular therapy; which to treat first? J Thromb Thrombolysis. (2017) 44:104–11. doi: 10.1007/s11239-017-1484-2

Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Copyright © 2019 Sakuta, Mukai, Fujii, Makita and Yaguchi. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.