Original Article

Clinical characteristics of cerebral infarction in the perioperative period of colorectal cancer excision

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Abstract: Objective: This study aimed to investigate clinical characteristics as well as potential pathogenesis of cerebral infarction in the perioperative period of colorectal cancer excision. Methods: Retrospective analysis was performed using the clinical data of 11 cases with perioperative cerebral infarction, screened out from 4,879 patients that received colorectal cancer excision. Results: A total of 4,879 patients that received colorectal cancer excision were included in the screening, and 11 (0.22%) were found to have developed perioperative cerebral infarction. Among the 11 cases, 5 presented (while the other 6 did not) histories of conventional stroke risk factors such as hypertension, diabetes, hyperlipidemia, brain stroke etc. Pathologic findings confirmed well-differentiated adenocarcinoma in screened-out patients (5 with colon cancer, and 6 with rectal cancer). Cerebral infarction was observed in 2, 3, 2 and 4 cases on day 1, 2, 3 and 4-6 after surgery, respectively. Single intracranial high-intensity infarction signal was confirmed in 4 cases (36.36%), while multiple high-intensity signals from areas with various blood supplies in 7 cases (63.64%). Increased levels of carcino-embryonic antigen (CEA), CA19-9, CA125 and D-dimer were observed in 6, 4, 7 and 8 cases, respectively. Conclusion: Cerebral infarction in the perioperative period of colorectal cancer excision usually appears in patients within 3 days after surgery, mostly with abnormal levels of serum tumor markers, while some with multiple infarction signals from intracranial areas had overlapped blood supplies. The pathogenesis may be related to hypercoagulability.

Keywords: Colorectal cancer, cerebral infarction, perioperative period, clinical characteristics

Introduction

Somatic malignant tumor may be the direct cause of vascular oppression and infiltration, hypercoagulability, non-bacterial thrombotic endocarditis (NBTE) etc., which brings about cerebral infarction in patients with cancer [1-3]. The risk of cerebral infarction is dramatically elevated in such patients [4-6]. And malignant tumor-targeted treatments including surgery, chemotherapy, radiotherapy etc. will also affect the incidence of cerebral infarction. Colorectal cancer is a common type of malignant tumor, and its incidence is gradually increasing along with the changes in economic development, lifestyle, eating patterns and dietary structure [7]. Surgical excision is presently one of the major treatments for colorectal cancer. Therefore, cerebral infarction may probably appear with an increasing incidence due to the perioperative stress-related blood pressure fluctuation, changed coagulative state etc. [8]. Cerebral infarction is a rare but severe perioperative neurological complication, with a reported incidence varying between 0.08% and 2.90% [9]. However, so far there have been few researches on the pathogenesis and clinical characteristics of perioperative cerebral infarction in patients with colorectal cancer.

In this retrospective study, 11 patients with perioperative cerebral infarction were screened out from those that underwent colorectal cancer excision during hospitalization in the First Affiliated Hospital of Guangxi Medical University between January 2006 and January 2015. Clinical data of selected patients were analyzed to explore clinical characteristics as well as potential pathogenesis of perioperative cerebral infarction in those with colorectal cancer.
Subjects and methods

Subjects

Patients with perioperative cerebral infarction were screened out from those that received surgical excision for colorectal cancer during hospitalization in the Frist Affiliated Hospital of Guangxi Medical University from January 2006 to January 2015. The inclusion criteria included localized symptoms of neurological deficit such as limb asthenia, numbness, hypaesthesia and dysarthria etc., which could be explained by the fresh infarction foci revealed by intracranial CT and MRI scanning and DWI (diffusion-weighted imaging) sequence scan, as was verified by MRI scanning 21 days later. The exclusion criteria consisted of non-perioperative cerebral infarction in patients with colorectal cancer; patients with colorectal cancer complicated by other malignant tumors, or with brain metastasis; hematological diseases; cerebral hemorrhage and shortage of reliable encephalic imaging data. The diagnostic criteria for colorectal cancer and acute cerebral infarction were as per the National Comprehensive Cancer Network (NCCN) guideline [10] and the American College of Cardiology (ACC) newly-updated standard [11], respectively. The definition of perioperative period was in accordance with what Leemingsawat et al. [8] had described, that’s from the day of the surgery to the seventh day thereafter. Case inclusion was jointly decided by a neurologist blinded to this study and a specialist of gastrointestinal surgery.

Methods

The study had collected demographic data such as gender, age etc., as well as histories of conventional stroke risk factors such as hypertension, diabetes, hyperlipidemia, atrial fibrillation, myocardial infarction, smoking and alcohol abuse, pulmonary or limb embolism, brain stroke etc. Neurological scoring was performed using the National Institutes of Health Stroke Scale (NIHSS) as per collected data such as symptoms and signs at the onset of the stroke. Thirty-day prognosis was evaluated using the modified Rankin Scale (mRS) in order to avoid the influence of colorectal cancer development on limb function. And unfavorable prognosis was suggested if mRS ≥ 2 points. Patient information collection was comprised of the site of the colorectal cancer, type of the tumor cells, metastasis, blood pressure fluctuation during surgery, surgical blood loss, anesthetic method and the onset time of cerebral infarction within or after the surgery. Pre-operational examinations included hematological tests such as routine blood test, biochemical examination, examination of four coagulation indices (PT, APTT, TT and FIB), D-dimer, tumor markers etc., as well as other tests such as electrocardiogram, ultrasonic cardiogram (UCG), B-mode ultrasound (BUS) scan of the neck and the abdomen, transcranial Doppler (TCD) imaging, chest X-ray scanning, abdominal and pelvic CT and MRI scanning (both unenhanced and enhanced), colonoscopy etc. Examinations after cerebral infarction included cranial CT and MRI scanning as well as DWI sequence scanning etc.

Results

Results on colorectal cancer

The study included a total of 4,879 patients who underwent colorectal cancer excision during hospitalization. And perioperative cerebral infarction was spotted in 11 patients (0.22%). One patient had family history of rectal cancer. All patients first complained of bloody stool, abdominal pain and distention, which were then hospitalized for surgery when tumor-like neoplasm was discovered via digital rectal examination and colonoscopy. Post-operative pathologic findings all suggested tubular adenocarcinoma, with 5 patients determined as colon cancer while 6 as rectal cancer. Hepatic metastasis was found in 3 patients, while metastasis was absent in other patients revealed by cranial CT and MRI scanning, enhanced CT scanning on chest and abdomen, and whole-body PET/CT. Radical resection was applied in 8 patients, and extended radical operation in 3. General anesthesia was adopted for all the 11 patients. Parametric levels of blood pressure fluctuation, blood loss and serum tumor markers are listed in Table 1.

Results on cerebral infarction

Five patients presented histories of stroke risk factors such as hypertension, diabetes, smoking and alcohol abuse, brain stroke etc. CTA/MRA examination suggested narrowed A1 segment of the right anterior cerebral artery in 1 patient. Scleroses of bilateral intracranial ver-
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### Table 1. Results of 11 patients with colorectal cancer

<table>
<thead>
<tr>
<th>Serial number</th>
<th>Gender</th>
<th>Age</th>
<th>Type of tumor cells</th>
<th>Cancer diagnosis</th>
<th>Duration of hypotension in surgery &gt; 1 h</th>
<th>Blood loss in surgery &gt; 1000 ml</th>
<th>Stage of the tumor</th>
<th>CEA (ng/ml)</th>
<th>CA19-9 (U/ml)</th>
<th>CA125 (U/ml)</th>
<th>D-dimer (ng/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Male</td>
<td>55</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>+</td>
<td></td>
<td>T2N0M0</td>
<td>&gt; 1500</td>
<td>186.90</td>
<td>185.70</td>
<td>1091</td>
</tr>
<tr>
<td>2</td>
<td>Male</td>
<td>71</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer with hepatic metastasis</td>
<td>+</td>
<td></td>
<td>T3N1M1</td>
<td>2.95</td>
<td>2.65</td>
<td>32.13</td>
<td>420</td>
</tr>
<tr>
<td>3</td>
<td>Female</td>
<td>47</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Ascending colon cancer with hepatic metastasis</td>
<td>+</td>
<td></td>
<td>T2N0M0</td>
<td>4.68</td>
<td>5.60</td>
<td>131.60</td>
<td>890</td>
</tr>
<tr>
<td>4</td>
<td>Male</td>
<td>65</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Sigmoid colon cancer</td>
<td>+</td>
<td></td>
<td>T3N0M0</td>
<td>25.41</td>
<td>482.10</td>
<td>101.90</td>
<td>3734</td>
</tr>
<tr>
<td>5</td>
<td>Male</td>
<td>70</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>+</td>
<td></td>
<td>T2N1M0</td>
<td>13.53</td>
<td>18.39</td>
<td>7.60</td>
<td>230</td>
</tr>
<tr>
<td>6</td>
<td>Female</td>
<td>56</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>+</td>
<td></td>
<td>T2N0M0</td>
<td>7.37</td>
<td>56.10</td>
<td>349.60</td>
<td>2347</td>
</tr>
<tr>
<td>7</td>
<td>Female</td>
<td>49</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer with hepatic metastasis</td>
<td>+</td>
<td></td>
<td>T3N1M1</td>
<td>0.64</td>
<td>5.30</td>
<td>14.50</td>
<td>460</td>
</tr>
<tr>
<td>8</td>
<td>Male</td>
<td>60</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>+</td>
<td></td>
<td>T3N0M0</td>
<td>12.50</td>
<td>22.00</td>
<td>191.40</td>
<td>1297</td>
</tr>
<tr>
<td>9</td>
<td>Male</td>
<td>55</td>
<td>Stage 1-2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>T2N0M0</td>
<td>6.79</td>
<td>25.12</td>
<td>87.13</td>
<td>1100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>Male</td>
<td>55</td>
<td>Stage 1-2 tubular adenocarcinoma</td>
<td>Rectal cancer</td>
<td>+</td>
<td></td>
<td>T2N0M0</td>
<td>10.90</td>
<td>21.70</td>
<td>10.90</td>
<td>260</td>
</tr>
<tr>
<td>11</td>
<td>Female</td>
<td>60</td>
<td>Stage 2 tubular adenocarcinoma</td>
<td>Ascending colon cancer</td>
<td>+</td>
<td></td>
<td>T2N0M0</td>
<td>0.89</td>
<td>78.60</td>
<td>105.70</td>
<td>790</td>
</tr>
</tbody>
</table>

+: The described condition exists.

### Table 2. Results of 11 patients with cerebral infarction

<table>
<thead>
<tr>
<th>Serial number</th>
<th>Gender</th>
<th>Age</th>
<th>Stroke risk factor</th>
<th>Site and number of cerebral infarction</th>
<th>Head-and-neck CTA &amp; MRA</th>
<th>Onset time of cerebral infarction</th>
<th>NIHSS score</th>
<th>mRS score 30 d after infarction</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Male</td>
<td>55</td>
<td>None</td>
<td>Multiple lesions in the left frontal, temporal, parietal and occipital lobes, as well as the head of caudate nucleus on the right side</td>
<td>Narrowed A1 segment of the right anterior cerebral artery</td>
<td>1</td>
<td>7</td>
<td>1</td>
</tr>
<tr>
<td>2</td>
<td>Male</td>
<td>71</td>
<td>Histories of brain stroke and hypertension</td>
<td>Multiple lesions in bilateral areas of corona radiate, basal ganglia and thalamus</td>
<td>Bilateral scleroses of intracranial vertebral artery</td>
<td>2</td>
<td>9</td>
<td>2</td>
</tr>
<tr>
<td>3</td>
<td>Female</td>
<td>47</td>
<td>None</td>
<td>One lesion in the right pons area</td>
<td>No abnormality</td>
<td>6</td>
<td>5</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
<td>Male</td>
<td>65</td>
<td>Histories of brain stroke and hypertension</td>
<td>Multiple lesions in the right frontal and parietal lobes, as well as the right basal ganglia area</td>
<td>No abnormality</td>
<td>3</td>
<td>5</td>
<td>6</td>
</tr>
<tr>
<td>5</td>
<td>Male</td>
<td>70</td>
<td>Histories of hypertension, smoking and alcohol abuse</td>
<td>Multiple lesions in the right cerebellar hemisphere, bilateral basal ganglia areas, left thalamus and pons</td>
<td>No abnormality</td>
<td>1</td>
<td>11</td>
<td>6</td>
</tr>
<tr>
<td>6</td>
<td>Female</td>
<td>56</td>
<td>None</td>
<td>Multiple lesions in the right temporal and occipital lobes, as well as the right thalamus</td>
<td>No abnormality</td>
<td>4</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>7</td>
<td>Male</td>
<td>49</td>
<td>None</td>
<td>Multiple lesions in the right cerebellum; left temporal, occipital and parietal lobes; left corona radiate and right frontal lobe</td>
<td>No abnormality</td>
<td>6</td>
<td>21</td>
<td>0</td>
</tr>
<tr>
<td>8</td>
<td>Female</td>
<td>53</td>
<td>None</td>
<td>Multiple lesions in bilateral frontal lobes, corona radiate, splenium of the left corpus callosum, and the basal ganglia area</td>
<td>No abnormality</td>
<td>2</td>
<td>3</td>
<td>6</td>
</tr>
<tr>
<td>9</td>
<td>Male</td>
<td>60</td>
<td>Diabetes</td>
<td>One lesion in the right basal ganglia area</td>
<td>No abnormality</td>
<td>3</td>
<td>11</td>
<td>0</td>
</tr>
<tr>
<td>10</td>
<td>Male</td>
<td>55</td>
<td>None</td>
<td>One lesion in the left temporal lobe</td>
<td>No abnormality</td>
<td>5</td>
<td>9</td>
<td>3</td>
</tr>
<tr>
<td>11</td>
<td>Female</td>
<td>60</td>
<td>Histories of brain stroke and hypertension</td>
<td>One lesion in the left basal ganglia area</td>
<td>No abnormality</td>
<td>2</td>
<td>7</td>
<td>3</td>
</tr>
</tbody>
</table>
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tebral arteries were also discovered in 1 patient. While no vascular or cardiac abnormalities were spotted in other patients via cranial CTA/MRA, UCG, TCD and neck BUS scanning. All patients presented acute neurological deficit symptoms to varying degrees, such as limb asthenia, numbness and dysarthria etc. And somnolence and obnubilation etc. were found in 3 patients. Cerebral infarction was observed in 2, 3, 2 and 4 cases on day 1, 2, 3 and 4-6 after surgery, respectively. Cerebral hemorrhage was excluded via cranial CT and MRI scanning as well as DWI sequence scanning. Cerebral infarction was confirmed by T1, T2 and DWI high-intensity signals. All patients received another round of MRI (both unenhanced and enhanced) and DWI sequence scanning 21 days later, to verify cerebral infarction and rule out brain tumor or metastasis. Single intracranial high-intensity infarction signal was confirmed in 4 patients (36.36%), while multiple high-intensity signals from areas with various blood supplies in 7 (63.64%) Figure 1. Neurological scoring was performed using the National Institutes of Health Stroke Scale (NIHSS). Thirty-day prognosis after stroke was evaluated using the modified Rankin Scale (mRS), and results had shown 4 cases with favorable prognosis (mRS = 0-1 point), 4 with unfavorable prognosis (mRS = 2-5 points), and 3 deaths (mRS = 6 points) (Table 2).

Figure 1. A-E. A 53-year-old female. Diffusion-weighted imaging: Multiple infarctions in supply areas of cerebral arteries. F-J. A 49-year-old male. Diffusion-weighted imaging: Multiple infarctions in supply areas of cerebral arteries.

Treatment and outcome

General nutritional support was provided and fasting required for all patients before surgery. Appropriate fluid infusion, blood transfusion and hemostatic treatment were adopted for patients suffering from blood pressure fluctuation and hemorrhage in surgery. Targeted treatments after stroke included dehydration therapy to reduce intracranial pressure, brain cell nourishment and neural rehabilitation, etc. No patient received intravenous thrombolysis or anti-platelet therapy. One out of the 3 deaths was caused by cerebral infarction, the other 2 by multiple organ failure (MOF).

Discussion

Malignant tumor will lead to increased risk of cerebral infarction, which suggests a correlation between cerebral infarction and malignant tumor [4-6]. A few researches have unraveled explicit characteristics of partial cerebral infarction patients with cancer. In the study carried out by Kim et al. [12] in 2012, 348 from 2,562 patients with acute cerebral infarction included along the continuum were screened out as complicated by active cancer. Seventy-one in those 348 patients (20.40%) presented increased D-dimer levels and multiple infarctions from intracranial areas with various arterial supplies, but no conventional stroke risk
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factors. Also, similar reports on cancer complicated by cerebral infarction are gradually increasing [13, 14]. Most patients in our study did not have conventional stroke risk factors, but had increased CEA, CA19-9, CA125 and D-dimer levels in peripheral circulation, as well as multiple infarctions from intracranial areas with various arterial supplies for a single onset. Such findings are consistent with prior researches, suggesting that there exist characteristics specifically related to colorectal cancer complicated by cerebral infarction.

Cerebral infarction is a rare but severe perioperative neurological complication, which usually takes place on the day of the surgery or 7 days thereafter [15]. A study carried out in the US targeting perioperative cerebral infarction of non-cardiac and non-vascular surgeries includes 130,000 patients and lasts for 18 years [16]. According to its results, conventional stroke risk factors such as hypertension, hyperlipidemia, smoking, alcohol abuse and obesity etc. are also risk factors of perioperative cerebral infarction. The incidence to develop perioperative cerebral infarctions increases to more than twice if the patients have brain stroke history. And the incidence of brain stroke has been raised by 25% in diabetic patients. Among patients included in our study, 3 had history of brain stroke, and 5 had other risk factors such as hypertension, diabetes, smoking and alcohol abuse etc. And the pathogenesis of brain stroke in these patients might be probably the same as that of non-perioperative cerebral infarction. Multiple infarctions from intracranial areas with various blood supplies were observed in 6 patients without stroke risk factors, and the pathogenesis of such infarctions was worth studying.

Malignant tumor may be the direct cause for vascular oppression and infiltration, hypercoagulability, NBTE etc., which brings about cerebral infarction in patients with cancer [1-3]. And hypercoagulative state is already confirmed as one important pathogenesis for complicated cerebral infarction in patients with malignant tumor [17]. The study by Jovin et al. [18] has shown increased level of serum CA125 and hypercoagulability in patients with malignant tumor complicated by cerebral infarction, indicating that the mucoprotein characteristics of CA125 may be related to hypercoagulability, and thus participate in the genesis of cerebral infarction in patients with malignant tumor. Hypotension is also considered as one of the major causes of perioperative cerebral infarction. The stress state that perioperative patients are going through is relevant to factors such as persistent hypotension and hypercoagulative state etc., which may contribute to the incidence of cerebral infarction [8, 19]. It was found in our study that partial patients had hypotension and increased serum CEA, CA19-9, CA125 and D-dimer levels, all of which were potential factors participating in the development of perioperative cerebral infarction in patients with colorectal cancer.

No patient received intravenous thrombolysis or anti-platelet therapy in our study. One out of the 3 deaths was caused by cerebral infarction and the other 2 by multiple organ failure (MOF). Although ultra-early thrombolytic therapy is viewed as a very effective approach to treat cerebral infarction, recent surgical history within three months is still a contraindication. Therefore the fatality and disability rates of perioperative cerebral infarction are increased due to the limited treatment. As revealed by the study of Chalela et al. [20] who treat 36 cases of ultra-early cerebral infarction via arterial injection of recombinant tissue-type plasminogen activator (rt-PA), partial or complete recanalization of occluded vessels is achieved by arterial thrombolysis in 80% of the patients with perioperative cerebral infarction. No or merely mild neurological deficit symptoms are present in 38% of the patients. The mortality is similar to that of arterial thrombolysis in patients with non-perioperative cerebral infarction. The incision bleeding rate is 17%, with only slight blood loss (< 30 mL) in most patients. The incidence of intracranial hemorrhage increases to about 25% of the patients, but only 8% with aggravated symptoms. Mortazavi et al. [21] have performed arterial thrombolysis on 3 arthroplastic patients with perioperative cerebral infarction, and succeeded in restoring blood flow through the occluded vessels. Two of the 3 patients are discharged from the hospital without any disability, and 1 dies from hemorrhagic complication. The detailed clinical efficacy remains to be observed as a result of the limited number of cases, and the shortage of large-scale clinical trials. The treatment and prognosis of perioperative cerebral infarction in
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patients that have undergone colorectal cancer excision are still yet to be further investigated.

The retrospective nature of the study, relatively smaller number of cases included, and clinical data collection limited by in-patient treatment requirements have altogether restricted more in-depth discussion on the pathogenesis of perioperative cerebral infarction in patients with colorectal cancer, which can be better elucidated through active large-scale prospective multicenter studies in the future.

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Disclosure of conflict of interest

None.

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