Stroke Care Delivery in Institutions Participating in the Registry of the Canadian Stroke Network

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Background and Purpose—Guidelines and performance indicators have been established for acute stroke care. However, little is known about the process of stroke care delivery in Canada.

Methods—The Registry of the Canadian Stroke Network (RCSN) captured detailed clinical data on patients with stroke and transient ischemic attack seen at 21 acute care institutions across Canada. Data from phase 1 of the RCSN (June 2001 to February 2002) were used to determine the use of evidence-based acute stroke care interventions in participating institutions.

Results—Overall, 4439 patients were seen during the study time frame and 1701 (38%) consented to full data collection. Thirty-one percent received care on a stroke unit or from a mobile stroke team. Among patients with ischemic stroke, 7% received thrombolysis, 80% underwent carotid imaging, 89% received antithrombotic agents, and 54% of those with atrial fibrillation received warfarin. There were significant intersite variations in the delivery of all of these interventions except for the use of antithrombotic agents, and these persisted after adjustment for age, sex, stroke type, and other comorbid conditions.

Conclusions—Patients in institutions participating in the RCSN received high-quality stroke care based on a number of performance measures. However, gaps exist in the provision of other elements of stroke care, particularly organized inpatient stroke care and warfarin for atrial fibrillation. Future research should explore explanations for these findings and focus on solutions to deficiencies in care. (Stroke. 2004;35:1756-1762.)

Key Words: stroke ■ quality of health care

Clinical trials have established the efficacy of a variety of interventions for acute stroke management, with improved outcomes seen with the use of thrombolysis, early treatment with aspirin, and multidisciplinary care on a stroke unit. In addition, clinical trials have established the efficacy of a number of therapies for secondary stroke prevention, including antiplatelet therapy, anticoagulation for atrial fibrillation, and carotid endarterectomy for carotid stenosis. Based in part on these trials, expert panels have established guidelines and performance indicators for quality stroke care. Despite this, studies have demonstrated significant gaps and regional variations in the provision of quality stroke care in a number of countries. It is not known whether similar variations exist in Canada.

The Registry of the Canadian Stroke Network (RCSN) was established in 2001, with the goal of measuring, monitoring, and improving stroke care in Canada. Twenty-one institutions from across Canada participated in the RCSN; all were urban tertiary care institutions with specific stroke care resources and expertise. The RCSN recruited consecutive patients with acute stroke presenting to these institutions and recorded data on stroke presentation, type, and severity, prehospital and emergency care, and in-hospital management and outcomes.

We used data from the first 8 months (phase 1) of the RCSN to determine the use of evidence-based acute stroke care interventions in stroke patients seen at participating Canadian hospitals. The stroke care performance indicators selected were based on published standards for in-hospital...
acute stroke care and included interventions such as thrombolysis and antiplatelet agents in eligible patients, warfarin in those with atrial fibrillation, organized inpatient stroke care (either a stroke ward or a mobile stroke team), and testing for carotid stenosis.12 We determined the proportion of patients who received each of these interventions and compared the use of stroke care interventions among the 21 participating institutions.

Materials and Methods

Data Source: Registry of the Canadian Stroke Network

The RCSN was established by the Canadian Stroke Network (www.canadianstrokenetwork.ca), which is funded by the Canadian Networks of Centres of Excellence (www.nce.gc.ca). Phase 1 of the RCSN took place between June 2001 and February 2002, and included 21 participating sites from 8 Canadian provinces. Potential sites were identified by soliciting applications from stroke neurologists who were members of the Canadian Stroke Consortium. The Steering Committee then selected sites based on their anticipated volumes and commitment to the RCSN and also attempted to ensure representation from most Canadian provinces. These sites had more resources and stroke expertise than the typical Canadian acute care institution. All participating sites were urban tertiary care institutions with specific stroke care resources: all had a neurologist with expertise in stroke, 81% were teaching hospitals, 67% had a mobile interdisciplinary acute stroke team, and 57% had an acute stroke ward. Although these institutions represented only 3% of acute care institutions in Canada, analyses of Canadian administrative data seen at these Registry institutions (J.V. Tu, unpublished data, 2000).

The literature was reviewed for relevant performance indicators for stroke care. All Registry patients (consented and nonconsented) were included in the analysis of thrombolysis rates; the remainder of the analyses included only those patients who consented to full data collection.

 Statistical Analyses

The main outcome measure was the proportion of stroke patients in the RCSN who received each of the performance indicators for stroke care. All Registry patients (consented and nonconsented) were included in the analysis of thrombolysis rates; the remainder of the analyses included only those patients who consented to full data collection.

Secondary analyses used χ² tests to compare the use of these interventions among participating institutions, with censoring of results from hospitals with <30 eligible cases or with >10% missing data for a given variable. For the use of thrombolysis, antithrombotic agents, carotid imaging, and organized stroke care, multiple logistic regression was used to compare care in different institutions with adjustment for age, sex, level of consciousness (as a surrogate for stroke severity), stroke type, and comorbidity. Analysis of use of thrombolysis, antithrombotic agents, and carotid imaging was limited to those with ischemic stroke. Regression modeling was performed using each performance indicator as the dependent variable, with predictor variables added using backward selection. The Charlson index was used to summarize comorbid illness.25 It was selected for this purpose because it is a widely used and well-validated index of comorbidity, which has been found to correlate with mortality in some studies. It provides a weighted summary score from 0 (no comorbid illness) to 31, based on the presence or absence of each of 17 medical conditions. Previous studies of individuals with stroke have found that the majority of patients (>75%) have Charlson scores of 0 or 1, indicating minimal comorbidity.30 SAS (version 8.02) was used for all analyses.

Results

During the study time period, 4439 patients were enrolled in the registry, and 1701 (38%) consented to full data collection. The mean age of patients in the study sample was 67 years and 46% were female (Table 1). There was a high prevalence of stroke risk factors documented on medical history, and there were interinstitutional variations in patient risk factors and stroke type and severity (Table 1). Compared with patients who did not consent to participate in the RCSN, participating patients were younger (median age of 69 versus 72 years, P = 0.0001), were more likely to speak English or French as a first language (90% versus 84%, P = 0.0017), and were more likely to be alert on admission (78% versus 66%, P = 0.0001).

Overall, 60% of patients were transported to hospital by ambulance, and 24% arrived within 2 hours of stroke onset (Table 2). The majority (86%) of Registry patients was admitted to hospital. Of these, 18% were admitted to an acute care stroke unit, 21% were seen by a mobile stroke team, and 31% received some form of organized stroke care (either ward or
TABLE 1. Baseline Characteristics of Patients in the Registry of the Canadian Stroke Network

<table>
<thead>
<tr>
<th>Variable</th>
<th>All Canada</th>
<th>Low</th>
<th>High</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>No. of consented patients</td>
<td>1701</td>
<td>9</td>
<td>240</td>
<td></td>
</tr>
<tr>
<td>Female (%)</td>
<td>47</td>
<td>35</td>
<td>58</td>
<td>0.2820</td>
</tr>
<tr>
<td>Mean age, y</td>
<td>67</td>
<td>63</td>
<td>71</td>
<td>0.0010</td>
</tr>
<tr>
<td>Past medical history</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stroke (%)</td>
<td>20</td>
<td>13</td>
<td>40</td>
<td>0.3928</td>
</tr>
<tr>
<td>TIA (%)</td>
<td>18</td>
<td>13</td>
<td>27</td>
<td>0.1979</td>
</tr>
<tr>
<td>Diabetes mellitus (%)</td>
<td>22</td>
<td>13</td>
<td>37</td>
<td>0.0207</td>
</tr>
<tr>
<td>Hypertension (%)</td>
<td>57</td>
<td>45</td>
<td>81</td>
<td>0.0020</td>
</tr>
<tr>
<td>Smoking (current) (%)</td>
<td>19</td>
<td>10</td>
<td>25</td>
<td>0.0436</td>
</tr>
<tr>
<td>Hyperlipidemia (%)</td>
<td>30</td>
<td>17</td>
<td>43</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Atrial fibrillation (%)</td>
<td>13</td>
<td>6</td>
<td>22</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Previous myocardial infarction</td>
<td>13</td>
<td>6</td>
<td>22</td>
<td>0.0140</td>
</tr>
<tr>
<td>Peripheral vascular disease (%)</td>
<td>5</td>
<td>1</td>
<td>15</td>
<td>0.1691</td>
</tr>
<tr>
<td>Charlson score &gt;1† (%)</td>
<td>27</td>
<td>15</td>
<td>44</td>
<td>0.1700</td>
</tr>
<tr>
<td>Stroke type</td>
<td></td>
<td></td>
<td></td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Ischemic stroke or TIA (%)</td>
<td>75</td>
<td>57</td>
<td>95</td>
<td></td>
</tr>
<tr>
<td>Intracerebral hemorrhage (%)</td>
<td>11</td>
<td>3</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>Subarachnoid hemorrhage (%)</td>
<td>9</td>
<td>0</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>Unable to determine (%)</td>
<td>4</td>
<td>0</td>
<td>26</td>
<td></td>
</tr>
<tr>
<td>Level of consciousness</td>
<td></td>
<td></td>
<td></td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Alert (%)</td>
<td>79</td>
<td>68</td>
<td>100‡</td>
<td></td>
</tr>
<tr>
<td>Confused (%)</td>
<td>7</td>
<td>0</td>
<td>20</td>
<td></td>
</tr>
<tr>
<td>Drowsy (%)</td>
<td>5</td>
<td>0</td>
<td>13</td>
<td></td>
</tr>
<tr>
<td>Unconscious (%)</td>
<td>9</td>
<td>0</td>
<td>24</td>
<td></td>
</tr>
</tbody>
</table>

*Two institutions with <30 eligible patients are censored in reporting the range among institutions.*
†Charlson score indicates the Charlson comorbidity index score, based on the presence or absence of 17 medical conditions; a score of zero indicates no comorbid illness and higher scores indicate more comorbid illness.
‡One institution did not permit surrogate consent; therefore, only alert patients capable of giving consent were included at this site.
TIA indicates transient ischemic attack.

Discussion

We found that patients seen at the institutions participating in the RCSN received high-quality stroke care based on a number of performance indicators. Specifically, 90% of “ideal” patients received antithrombotic agents at discharge from hospital, 80% underwent carotid imaging in hospital, and 14% of potentially eligible patients received thrombolysis. However, we also found gaps in the provision of other components of stroke care: only 18% of Registry patients were admitted to an acute stroke unit, only one third of Registry patients received any form of organized inpatient stroke care, and only 54% of those with atrial fibrillation received warfarin at discharge. In addition, there were significant variations among participating sites in the provision of organized stroke care, thrombolysis, and carotid imaging, and these did not appear to be fully explained by variations in case mix among the sites.

Variations and deficiencies in stroke care delivery have been documented in many other jurisdictions. An Australian stroke audit found that 78% of those with ischemic stroke or TIA received antithrombotic agents, with low rates of warfarin use for those with atrial fibrillation (33%), stroke unit care (23%), and thrombolysis (1%). A study of US Medicare patients hospitalized with stroke or transient ischemic attack found rates of discharge antithrombotic use ranging from 74% to 91% across the country, whereas studies from the Scottish Stroke Outcomes Study Group and the United Kingdom General Practitioner Research Database have found...
underuse and regional variations in the use of antithrombotic agents, multidisciplinary teams, and neuroimaging. More recently, the UK National Sentinel Stroke Audit found that 36% of patients across the United Kingdom were admitted to stroke units and 91% were prescribed antithrombotic agents at discharge, with wide interinstitutional variations in the organization and delivery of care.

Variations in care may be explained in part by appropriate patient selection resulting in exclusion of patients with contraindications to the therapy under consideration. The detailed clinical data in the Registry of the Canadian Stroke Network allowed us to examine the use of interventions among “ideal” patients without contraindications to therapy and to explore predictors of use for each intervention. The use of tPA was relatively high (14%) in those presenting to hospital within 3 hours of stroke onset, and the majority of patients who did not receive tPA were excluded on the basis of eligibility criteria derived from the NINDS tPA study. Similarly, the use of carotid imaging in 80% of patients may be appropriate, given that such imaging is not required for

### TABLE 2. Emergency Department Stroke Care Performance Indicators

<table>
<thead>
<tr>
<th>Variable</th>
<th>All Canada</th>
<th>Range Among Institutions*</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arrival by ambulance, n/n eligible patients (%)</td>
<td>60</td>
<td>24/64 (38)</td>
<td>43/45 (96)</td>
</tr>
<tr>
<td>Arrival within 2 hours of stroke onset, n/n eligible patients (%)</td>
<td>24</td>
<td>5/42 (13)</td>
<td>24/56 (42)</td>
</tr>
<tr>
<td>Thrombolysis given, n/n eligible patients† (%)</td>
<td>4</td>
<td>0/119 (0)</td>
<td>8/68 (12)</td>
</tr>
<tr>
<td>In subgroup with ischemic stroke</td>
<td>7</td>
<td>0/80 (0)</td>
<td>8/64 (12)</td>
</tr>
<tr>
<td>In subgroup with ischemic stroke and symptoms &lt;3 hours</td>
<td>14</td>
<td>0/44 (0)</td>
<td>11/38 (29)</td>
</tr>
</tbody>
</table>

*Two institutions with <30 eligible patients are censored in reporting the “range among institutions.”

†Ten institutions with <30 eligible patients or >10% missing data are censored in reporting the “range among institutions” for this variable.

‡Other reasons include recent surgery or trauma, bleeding disorders, and other unspecified reasons.

ER indicates emergency room.

### TABLE 3. In-Hospital Stroke Care Performance Indicators

<table>
<thead>
<tr>
<th>Performance Indicator</th>
<th>All Canada</th>
<th>Range Among Institutions*</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Care on acute stroke ward, n/n eligible patients (%)</td>
<td>18</td>
<td>0/199 (0)</td>
<td>43/51 (85)</td>
</tr>
<tr>
<td>Care on stroke ward or by stroke team, n/n eligible patients (%)</td>
<td>31</td>
<td>0/68 (0)</td>
<td>45/51 (88)</td>
</tr>
<tr>
<td>Care on stroke ward or by stroke team in subgroup of institutions where stroke ward or team care available,† n/n eligible patients (%)</td>
<td>44</td>
<td>4/41 (10)</td>
<td>45/51 (88)</td>
</tr>
<tr>
<td>Antithrombotic therapy at discharge,‡§ n/n eligible patients (%)</td>
<td>89</td>
<td>55/73 (76)</td>
<td>119/127 (94)</td>
</tr>
<tr>
<td>Antithrombotic therapy in “ideal” patients,§¶ n/n eligible patients (%)</td>
<td>90</td>
<td>54/68 (80)</td>
<td>80/83 (96)</td>
</tr>
<tr>
<td>Warfarin for atrial fibrillation§ (%)</td>
<td>54</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Warfarin for atrial fibrillation in “ideal” patients§¶ (%)</td>
<td>57</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Carotid imaging,‡§ n/n eligible patients (%)</td>
<td>80</td>
<td>39/63 (62)</td>
<td>193/199 (97)</td>
</tr>
</tbody>
</table>

*Three institutions with <30 eligible patients are not reported in the “range among institutions.”

†Analysis limited to 14 institutions with an acute stroke ward or stroke team.

‡Antithrombotic therapy includes aspirin, clopidogrel, ticlopidine, dipyridamole, and warfarin; carotid imaging indicates carotid Doppler, cerebral angiography or magnetic resonance angiography performed during the index hospitalization.

§Analysis limited to patients with ischemic stroke.

¶“Ideal” patients are those without a history of bleeding disorder, peptic ulcer disease, or cirrhosis.

||Seven institutions with <30 eligible patients of >10% missing data for the variable of interest are censored in reporting the “range among institutions.”
patients in whom the results are unlikely to change management, such as those who are not candidates for carotid endarterectomy because of disabling stroke or other comorbid illness. Additional patients may have undergone carotid imaging soon after discharge, and this would not have been captured by the database. Warfarin was used in only 54% of those with atrial fibrillation; discussion with neurologists participating in the Registry suggests that this may be explained in part by uncertainty about the relative benefits of warfarin versus antiplatelet agents for secondary stroke prevention in elderly patients with severe strokes and a higher perceived risk of complications. However, even after adjustment for stroke type and other prognostic factors, we observed persistent intersite variations in the use of stroke care interventions, suggesting that patient factors are not the sole explanation for the observed variations in stroke care delivery.

Other explanations for variations in care include the availability and organization of local resources. The use of thrombolysis and antithrombotic agents in RCSN patients was higher than that observed in some previous stroke audits, and likely reflects the specialized nature of the participating institutions. However, even among these highly selected institutions, some administered no thrombolysis at all. Discussions with the participating neurologists revealed that in some cases this was attributable to insufficient personnel with expertise in administering thrombolysis, or an institutional or departmental concern about the efficacy of tPA. Furthermore, only 57% of these selected institutions had an acute stroke ward and only 67% had either a stroke ward or a mobile stroke team, and the availability of acute stroke units among other institutions in Canada is almost certainly much lower: a 1999 survey of all Ontario hospitals found that an acute stroke unit existed in only 4% of acute care institutions.39

Other local initiatives and resources that may influence care include public awareness campaigns and training of emergency medical service personnel to decrease prehospital delays, as well as emergency department protocols to facilitate timely neuroimaging and use of thrombolysis. In this study, only 24% of all stroke patients arrived in hospital within 2 hours of stroke onset. In addition, 15% of patients arriving within 2 hours of stroke onset did not receive thrombolysis because of symptom duration >3 hours at the time of stroke team assessment, suggesting that in-hospital delays may have prevented some eligible patients from receiving this therapy. Further research is needed to determine whether stroke protocols and other local initiatives can result in improvements in these areas.

Variations and deficiencies in stroke care are important if they result in variations in stroke outcomes. Studies evaluating the link between stroke care process and outcome have yielded conflicting results, and we do not yet have long-term stroke outcome data on our study sample. In theory, however, assuming that 50% of stroke patients will be dead or dependent (defined as a Rankin grade of ≥3) 1 year after stroke, and that organized stroke care reduces the odds of death or dependency by 22%, the failure to provide organized inpatient stroke care in 70% of patients could contribute to an increase of 8 patients dead or dependent for every 100 stroke patients seen. Similarly, assuming a 15% annual recurrent stroke risk with atrial fibrillation and a 66% relative risk reduction with warfarin therapy, the failure to use warfarin in 46% of patients would be expected to contribute to an excess of 5 preventable strokes per year per 100 eligible patients. Given an estimated 50 000 stroke admissions per year in Canada, addressing such deficiencies in care could have a significant impact on reducing death and disability from stroke.

A number of study limitations merit comment. First, the current analysis captures only in-hospital data. Although this could lead to underestimates of rates of carotid imaging and medication usage, usual Canadian practice patterns suggest that it is unlikely that a significant proportion of patients would have such testing or treatment initiated after hospital discharge. Second, we did not evaluate all potentially relevant care process indicators, but rather chose to focus on selected, easily measurable indicators that would apply to a majority of patients. Third, we do not yet have data on long-term stroke outcomes such as mortality and functional status, so we cannot evaluate the association between the processes of stroke care and stroke outcomes. Fourth, our study sample includes only patients seen at highly specialized institutions with stroke care expertise and resources, and it is probable that deficiencies in stroke care delivery are more marked at other institutions. Finally, even at these institutions, our study sample did not capture all stroke patients. Because of the requirement of informed consent, patients with minor stroke were less likely to participate (they were sometimes discharged before the coordinator had an opportunity to approach them), as were those with severe or fatal stroke (the coordinators were sometimes unable to obtain surrogate consent). Although such selection bias could limit the generalizability of results of analyses of patient outcomes, it is unlikely to lead to systematic biases in the current analyses of in-hospital processes of care.

In conclusion, Canadian stroke patients at these specialized institutions received high-quality stroke care based on a number of performance measures. However, gaps and significant interinstitutional variations exist in the provision of other elements of evidence-based stroke care. These data provide a benchmark against which quality improvement in Canadian hospitals can be measured. Future research should explore explanations for these observed differences, evaluate their effect on stroke patient outcomes, and focus on solutions to deficiencies in care.
Appendix

The following persons and institutions participated in phase 1 of the Registry of the Canadian Stroke Network:

**Participating Centers**

- **Queen Elizabeth II Health Sciences Centre**, Halifax, NS: S. Phillips, MD (Principal Investigator), M. Desrochers, RN (Coordinator), L. Mercille, RN (Coordinator)
- **Saint John Regional Hospital**, St. John, NB: P. Bailey, MD (Principal Investigator), P. Cook, RN (Coordinator), S. Alward, RN (Coordinator)
- **Hopital Notre-Dame du CHUM**, Montreal, QC: L. Lebrun, MD (Principal Investigator), M. Desrochers, RN (Coordinator), L. Mercille, RN (Coordinator)
- **Hopital de l’Enfant-Jesus**, Quebec City, QC: D. Simard, MD (Principal Investigator), A. Mackey, MD (Principal Investigator), S. Dube, RN (Coordinator), B. Leger, RN (Coordinator)
- **Hopital Charles le Moyne**, Greenfield Park, QC: L. Berger, MD (Principal Investigator), L. Moisan, RN (Coordinator), Y. Ser-raspino, RN (Coordinator), D. Truong, RN (Coordinator)
- **Montreal General Hospital and SMBD-Jewish General Hospital**, Montreal, QC: R. Cote, MD (Principal Investigator), J. Minuk, MD (Principal Investigator), C. Wong (Coordinator)
- **Sunnybrook & Women’s College Health Sciences Centre**, Toronto, ON: S. Black, MD (Principal Investigator), N. Jiang, (Coordinator), J. Bray (Coordinator), M. Kerr-Taylor, RN (Coordinator)
- **University Health Network/Toronto Western Hospital**, Toronto, ON: F. Silver, MD (Principal Investigator), P. Urzua, RN (Coordinator), G. Gutierrez, RN (Coordinator), R. Wiegner, RN (Coordinator)
- **London Health Sciences Centre**, London, ON: V. Hachinski, MD (Principal Investigator), N. Absolon, RN (Coordinator), L. Cotton, RN (Coordinator)
- **The Ottawa Hospital**, Ottawa, ON: A. Douen, MD (Principal Investigator), M. Sharma, MD (Principal Investigator), N. Pageau, RN (Coordinator), M. Savage, RN (Coordinator)
- **Kingston General Hospital**, Kingston, ON: D. Howse, MD (Principal Investigator), D. Brunet, MD (Principal Investigator), S. Weatherby, RN (Coordinator)
- **Hamilton Health Sciences Centre**, Hamilton, ON: W. Ocz-kowski, MD (Principal Investigator), N. Pyette, RN (Coordinator), L. Gould, RPN (Coordinator)
- **Trillium Health Sciences Centre**, Mississauga, ON: D. Selchen, MD (Principal Investigator), H. Hinks, RN (Coordinator), T. Stokes, RN (Coordinator)
- **Winnipeg Regional Health Authority**, Winnipeg, MB: B. Anderson, MD (Principal Investigator), D. Gladish, RN (Coordina-tor), J. Gousseau, RN (Coordinator), P. Pisk, RN (Coordinator)
- **Royal University Hospital**, Saskatoon, SK: C. Vell, MD (Prin-cipal Investigator), S. Bishop, RN (Coordinator), L. Schmidt, RN (Coordinator), B. Kwiatkowski, RN (Coordinator)
- **Foothills Medical Centre**, Calgary, AB: M. Hill, MD (Principal Investigator), L. Sinclair, RN (Coordinator), M. Schebel, RN (Co-ordinator), A. Cole-Haskayne, RN (Coordinator)
- **University of Alberta Hospital**, Edmonton, AB: A. Shuaib, MD, (Principal Investigator), A. Nasser, RN (Coordinator)
- **Lion’s Gate Hospital**, Vancouver, BC: D. Cameron, MD (Principal Investigator), C. Tadey, RN (Coordinator)
- **Vancouver General Hospital**, Vancouver, BC: P. Teal, MD (Principal Investigator), T. Steele, BSN, RN (Coordinator)
- **St. Paul’s Hospital**, Vancouver, BC: D. Johnston, MD (Principal Investigator), M. Wong, MD (Principal Investigator), H. Connolly, RN (Coordinator)
- **Capital Health Region**, Victoria, BC: A. Penn, MD (Principal Investigator), M. Laporte, RN (Coordinator)

**Coordinating Centre at the Institute for Clinical Evaluative Sciences, Ontario**: M. Kapral, F. Silver, J. Fong, A. Laupacis, J. Richards, J. Tu

**Steering Committee**: F. Silver (Cochair, phase 1), A. Hakim, M. Hill, M. Kapral (Chair, phase 2), A. Laupacis, M. Lewis, N. Mayo, S. Phillips, (Cochair, phase 1), G. Taylor, J. Tu, K. Willis

**Data Privacy and Security Committee**: D. Willison, ScD (Chair), A. Buchan, MD, A. Laupacis, MD, P. Peladeau, A. Penn, J. Richards, F. Silver, J. Williams


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**References**

1762 Stroke July 2004


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