Secondary stroke prevention: challenges and solutions

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Abstract: Stroke is the leading cause of disability in the USA and a major cause of mortality worldwide. One out of four strokes is recurrent. Secondary stroke prevention starts with deciphering the most likely stroke mechanism. In general, one of the main goals in stroke reduction is to control vascular risk factors such as hypertension, diabetes, dyslipidemia, and smoking cessation. Changes in lifestyle like a healthy diet and aerobic exercise are also recommended strategies. In the case of cardioembolism due to atrial fibrillation, mechanical valves, or cardiac thrombus, anticoagulation is the mainstay of therapy. The role of anticoagulation is less evident in the case of bioprosthetic valves, patent foramen ovale, and dilated cardiomyopathy with low ejection fraction. Strokes due to larger artery atherosclerosis account for approximately a third of all strokes. In the case of symptomatic extracranial carotid stenosis, surgical intervention as close as possible in time to the index event seems highly beneficial. In the case of intracranial large artery atherosclerosis, the best medical therapy consists of antiplatelets, high-dose statins, aggressive controls of vascular risk factors, and lifestyle modifications, with no role for intracranial arterial stenting or angioplasty. For patients with small artery occlusion (ie, lacunar stroke), the therapy is similar to that used in patients with intracranial large artery atherosclerosis. Despite the constant new evidence on how to best treat patients who have suffered a stroke, the risk of stroke recurrence remains unacceptably high, thus evidencing the need for novel therapies.

Keywords: stroke mechanisms, stroke risk, hypertension, diabetes, dyslipidemia

Introduction

Stroke is defined as clinical, radiological, or pathological evidence of ischemia or hemorrhage, involving a defined cerebral vascular territory. In the USA, there are approximately 800,000 strokes per year, approximately 600,000 of which are recurrent events. Stroke is now the fifth leading cause of death, but it remains the number one cause of disability in the USA. While there has been a steady decline in stroke incidence in developed countries, incidence in low-to-middle-income countries continues to increase – accounting for 85% of the worldwide stroke burden.

Once a stroke has occurred, treatment options are limited and only available for a short time immediately after the symptom onset. As a result, stroke prevention has been considered the mainstay in stroke management for over half a century, and despite decades of research in stroke prevention, there remain basic challenges in secondary stroke prevention. The literature addressing secondary stroke prevention is vast and impossible to thoroughly discuss in a single paper. This review should be viewed as a summary of the more frequent stroke mechanisms and the available evidence from randomized clinical trials for the best therapies to prevent stroke recurrence.
Ischemic stroke characterization

Knowing the stroke etiology is a critical step in the entire process because both risk-factor modification and stroke prevention begin with appropriate characterization of the stroke mechanism. Depending on the stroke mechanism, the risk of recurrence and suggested algorithms to prevent stroke vary. The most commonly used classification scheme in ischemic stroke is the Trial of Org10172 in Acute Stroke Treatment (TOAST) classification, which includes the following five subtypes: large artery atherosclerosis (LAA), cardioembolic, small vessel occlusion (SVO), stroke of undetermined cause (ie, cryptogenic stroke), and stroke of “other” cause. Although other classification systems have been suggested, the TOAST classification remains the most frequently used in large epidemiological studies (Table 1).

To determine stroke etiology, a thorough workup is needed to exclude potential stroke mechanisms that may need a change in treatment. In epidemiological studies, the proportion of cryptogenic strokes varies from 13% to 50%.

**Table 1** Summary of studies from different ethnic and racial groups that disclosed stroke subtype rates

<table>
<thead>
<tr>
<th>Reference(s)</th>
<th>Country/study</th>
<th>Population composition</th>
<th>Stroke types</th>
<th>Ischemic stroke subtypes (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Prospective, population-based studies</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>White, Boden-Albala et al 2005</td>
<td>USA/NOMAS</td>
<td>64% Hispanic</td>
<td>77% IS</td>
<td>Hispanic</td>
</tr>
<tr>
<td>Collaborators, Sacco et al 1998</td>
<td>Follow-up</td>
<td>13% Black</td>
<td>17% ICH</td>
<td>CE</td>
</tr>
<tr>
<td>Sacco et al 1998</td>
<td>=4 years</td>
<td>22% White</td>
<td>6% SAH</td>
<td>LAA</td>
</tr>
<tr>
<td>Morgenstern, Smith et al 2004</td>
<td>USA/BASIC</td>
<td>48% White</td>
<td>83% IS</td>
<td>SVO</td>
</tr>
<tr>
<td>Smith et al 2004</td>
<td>project</td>
<td>53% Mexican</td>
<td>14% ICH</td>
<td>Other</td>
</tr>
<tr>
<td>Uchino, Risser et al 2004</td>
<td>Follow-up</td>
<td>American</td>
<td>3% SAH</td>
<td>Crypto</td>
</tr>
<tr>
<td>Broderick, Schneider, Kissela et al 2004</td>
<td>USA/GCNSK</td>
<td>18.5% Black</td>
<td>84% IS</td>
<td>Other</td>
</tr>
<tr>
<td>Broderick, Brott et al 1998</td>
<td>Follow-up</td>
<td>72.5% White</td>
<td>10% ICH</td>
<td>Mexican American</td>
</tr>
<tr>
<td>Schneider, Kissela et al 2004</td>
<td>=2–4 years</td>
<td>4% SAH</td>
<td>2% undetermined</td>
<td>White</td>
</tr>
<tr>
<td>Ohira, Shahar et al 2006, 2011</td>
<td>USA/ARIC</td>
<td>25% Black</td>
<td>88% IS</td>
<td>CE</td>
</tr>
<tr>
<td>Petty, Brown et al 1999</td>
<td>Follow-up</td>
<td>75% White</td>
<td>12% ICH</td>
<td>LAA</td>
</tr>
<tr>
<td></td>
<td>=4 years</td>
<td>Not available</td>
<td></td>
<td>SVO</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Crypto</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Other</td>
</tr>
<tr>
<td>Hajat, Heuschmann et al 2010</td>
<td>England/SLSS</td>
<td>71% White</td>
<td>80% IS</td>
<td>Whites (95% of the sample)</td>
</tr>
<tr>
<td>Hajat, Dundas et al 2001</td>
<td>Follow-up</td>
<td>20% Black</td>
<td>14% ICH</td>
<td>CE</td>
</tr>
<tr>
<td></td>
<td>=7 years</td>
<td>(13% Caribbean</td>
<td>6% SAH</td>
<td>LAA</td>
</tr>
<tr>
<td>(7% African)</td>
<td></td>
<td>Other</td>
<td></td>
<td>SVO</td>
</tr>
<tr>
<td>Feigin, Carter et al 2006</td>
<td>New Zealand/ARCSS</td>
<td>66% White</td>
<td>73% ischemic</td>
<td>Crypto</td>
</tr>
<tr>
<td></td>
<td>Follow-up</td>
<td>20% Maori/Pacific Islander</td>
<td>12% ICH</td>
<td>Other</td>
</tr>
<tr>
<td></td>
<td>=1 year</td>
<td>14% Asian/other</td>
<td>6% SAH</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>9% undetermined</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Maori/Pacific Islander</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Asian/other</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>White</td>
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<td></td>
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</table>

(Continued)
Table I (Continued)

<table>
<thead>
<tr>
<th>Reference(s)</th>
<th>Country/study</th>
<th>Population composition</th>
<th>Stroke types</th>
<th>Ischemic stroke subtypes (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Turin, Kita et al 2010</td>
<td>Japan/TSR</td>
<td>100% Japanese</td>
<td>67% IS</td>
<td>Japanese</td>
</tr>
<tr>
<td>Kita, Turin et al 2007</td>
<td>Follow-up</td>
<td>16 years</td>
<td>22% ICH 10% SAH 1% undetermined</td>
<td>CE 23 SVO 54 Non-SVO 21 Other 2</td>
</tr>
</tbody>
</table>

Cross-sectional, hospital-based studies

<table>
<thead>
<tr>
<th>Reference(s)</th>
<th>Country/study</th>
<th>Sample size</th>
<th>Population composition</th>
<th>Stroke types</th>
<th>Ischemic stroke subtypes (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gutierrez, Koch et al 2013</td>
<td>USA/JMH Sample size = 473</td>
<td>53% Hispanic</td>
<td>83% IS</td>
<td>Hispanic</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>23% African American</td>
<td>17% ICH</td>
<td>Black</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>13% Black</td>
<td>CE 38 LAA 22 SVO 17 Crypto 13 Other 5</td>
<td>Caribbean Black 23 28 13 16</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>11% White</td>
<td>4</td>
<td>White</td>
<td></td>
</tr>
<tr>
<td>Sharma, Tsivgoulis et al 2012</td>
<td>Singapore Sample size = 481</td>
<td>74.2% Chinese</td>
<td>Not available</td>
<td>Chinese</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>16.6% Malay</td>
<td>30</td>
<td>Malay</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>9.2% Indian</td>
<td>7</td>
<td>Indian</td>
<td></td>
</tr>
<tr>
<td>Saposnik and Del Brutto 2003</td>
<td>South America N=912 (Meta-analysis of hospital-based samples)</td>
<td>Mix of mestizos, whites and nates</td>
<td>65% IS</td>
<td>CE 14 LAA 7 SVO 43 Crypto 30 Other 6</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>34% ICH</td>
<td>33</td>
<td>Whites 20</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>1% Other</td>
<td>13</td>
<td>Natives 22 15 19</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>18</td>
<td></td>
</tr>
</tbody>
</table>

Notes: Stroke subtypes vary across different populations. Demographic information may be helpful in guiding workup for stroke subtypes. The different proportions of stroke subtypes in each of the reported groups underscore the burden of certain risk factors in these populations and how it may lead to targeted strategies for secondary stroke prevention from a public health perspective. With kind permission from Springer Science+Business Media: Curr Cardiovasc Risk Rep; An epidemiological perspective on race/ethnicity and stroke, 9, 2015, 19, Suvash-Garcy J, Gutierrez J, Table 1.9

Abbreviations: CE, cardioembolism; crypto, cryptogenic; ICH, intracranial hemorrhage; IS, ischemic stroke; LAA, large artery atherosclerosis; SAH, subarachnoid hemorrhage; SVO, small vessel occlusion.

according to the workup undertaken.10,11 This underscores the need to carry out an exhaustive workup of patients with strokes.

Stroke attributed to LAA is defined as infarction distal to a large vessel stenosis. Frequent sites for atherosclerotic plaques include the aortic arch and ascending aorta, the extracranial carotid artery, and the proximal arteries in the Circle of Willis (Figures 1 and 2).12-14 Ischemia most commonly results from distal embolization of thrombotic products, a so-called artery-to-artery embolism, but it can also be caused by hypoperfusion of brain tissue distal to a severely stenotic or occluded vessel or from branch occlusive disease (Figures 1–3).

To define a stroke as cardioembolic, a clear cardiac source of embolism must be established. The most common etiologies of cardioembolic stroke are atrial fibrillation, valvular heart disease, large myocardial infarction (MI), or dilated cardiomyopathy.15-18 On the other hand, strokes that have an associated proximal large vessel stenosis, or that appear most consistent with a lacunar syndrome, can also result from cardioembolism, highlighting the challenges in determining a precise stroke mechanism.19,20 Atrial fibrillation, if paroxysmal, may be elusive. Prolonged cardiac monitoring in patients with stroke deemed “cryptogenic” may reveal occult atrial fibrillation in up to a fifth of patients, compared to only 2% with 7-day cardiac monitoring.21,22 Consequently, a true cryptogenic stroke should typically exclude the presence of occult atrial fibrillation, particularly in subjects >60 years and those with evidence of prior cortical infarcts.23

Stroke from SVO is typically referred to as lacunar stroke, which in modern practice is not limited to a specific size cutoff, but rather defined by anatomical location and a typical lacunar syndrome.19,24 The typical underlying arterial pathology in lacunar infarcts includes microatheroma of the penetrating arteries (most common in the largest infarcts), lypohyalinosis (more frequent in small infarcts), microembolism, or branch occlusive disease (Figure 3).
More recent evidence suggests, however, that SVO and LAA may be the different phenotypic expression of the same underlying intracranial arterial disease, with SVO representing perhaps an earlier form of the disease.²⁵–²⁷ This fact may also explain why the medical therapy for those with SVO and intracranial LAA is similar, despite the higher risk of stroke recurrence noted for those with LAA.²³,²⁵ It is also important to recognize that a transient ischemic attack (TIA) calls for the same urgency in evaluation as a stroke. In one study, the risk of stroke after TIA was approximately 5% in the first 2 days and 10% within 90 days.²⁹
Secondary stroke prevention: challenges and solutions

Figure 2 Stroke and intracranial atherosclerosis.
Notes: A woman in her 60s came to the hospital for left-sided weakness and headache. On exam, she was found to have mild left pronator drift and visuospatial neglect. (A) The brain magnetic resonance image showed evidence of a right parietal lobe cortical infarct and (B) the brain magnetic resonance angiograph showed evidence of diffuse luminal narrowing of her brain arteries (arrows), with some of these stenoses located proximal to the area of her infarction (small arrow), suggesting artery-to-artery embolism from intracranial large artery stenosis as the most likely stroke mechanism.

Figure 3 Small artery disease versus branch occlusive disease.
Notes: (A) An example of a small infarct in the left putamen and subcortical white matter (arrow), most likely due to the occlusion of a lenticulostriate artery branching of from the middle cerebral artery. (B) The brain magnetic resonance angiograph in this same patient showed no evidence of large artery stenosis in the proximal middle cerebral artery. (C) A patient with evidence of an infarct (arrow) involving the lenticular nucleus and the head of the caudate nucleus. (D) Contrary to the case presented in A, this patient shows evidence of a high-degree of stenosis in the middle cerebral artery in the brain computed tomography angiograph (large arrow), suggesting branch occlusive disease and “pure” small artery disease as the underlying etiology of the infarct.
General prevention strategies

Antiplatelets

As a group, antiplatelets offer an absolute risk reduction of 2% in vascular events per year, at the cost of a 0.1%–0.3% increase in major extracranial hemorrhages (Table 2). Aspirin is the most extensively studied, cheapest, and most commonly used agent in secondary stroke prevention. The US Food and Drug Administration currently recommends doses between 50 and 325 mg daily for stroke prevention. There is no evidence that clopidogrel is superior to aspirin alone for secondary stroke prevention, but clopidogrel seems superior to aspirin in preventing combined vascular outcomes, which is mostly driven by a reduction in leg amputation among patients with peripheral arterial disease. Aspirin with extended-release dipyridamole, compared to aspirin alone, produced an approximate annual absolute risk reduction of 1% in two large clinical trials. Twice-a-day dosing and headache are drawbacks of aspirin with extended-release dipyridamole. On the other hand, a clinical trial investigating the use of clopidogrel vs the combination of aspirin with extended-release dipyridamole failed to show any significant reduction in events that would favor either agent.

Table 2 Summary of some of the major studies comparing antiplatelets against each other or against placebo in stroke prevention

<table>
<thead>
<tr>
<th>Secondary stroke prevention setting</th>
<th>Trial</th>
<th>Placebo</th>
<th>ASA</th>
<th>ASA+D</th>
<th>C</th>
<th>ASA+C</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acute (0–2 weeks)</td>
<td>CAST</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td>12% RRR with ASA at 4 weeks with an ARR of 0.6% in early mortality and nonfatal stroke</td>
</tr>
<tr>
<td></td>
<td>IST+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td>20% RRR with ASA at 2 weeks with an ARR of 0.9% in stroke recurrence</td>
</tr>
<tr>
<td></td>
<td>CHANCE+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>32% RRR with dual antiplatelets in stroke with an ARR of 3.5% over 3 months in stroke</td>
</tr>
<tr>
<td>Subacute (3–6 months)</td>
<td>UKA TIA</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td>18% RRR with ASA in composite outcome</td>
</tr>
<tr>
<td></td>
<td>ESPS+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>38% RRR with ASA+D in stroke recurrence with a yearly ARR of 2.5%</td>
</tr>
<tr>
<td></td>
<td>ESPS2+</td>
<td>+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td>18% RRR in ASA group compared to placebo</td>
</tr>
<tr>
<td></td>
<td>ESPRIT+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>37% RRR in ASA+D with a yearly ARR of 2.5% in stroke/death compared to placebo</td>
</tr>
<tr>
<td></td>
<td>ProFESS+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>20% RRR with ASA+D over ASA alone with a yearly ARR of 1% in composite outcome</td>
</tr>
<tr>
<td></td>
<td>SPS3+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>No difference in stroke recurrence</td>
</tr>
<tr>
<td></td>
<td>CAPRIE+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>No benefit in stroke but increased risk of major hemorrhage with dual antiplatelets (2.1% vs 1.1% per year) and increased mortality (2.1% vs 1.5% per year)</td>
</tr>
<tr>
<td>Chronic (&gt;3 months)</td>
<td>AICLA+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>8.7% RRR of C over ASA in all groups with a yearly ARR of 0.5 in composite outcome (driven by PVD group)</td>
</tr>
<tr>
<td></td>
<td>MATCH+</td>
<td>+</td>
<td>+</td>
<td></td>
<td></td>
<td></td>
<td>6.4% RRR with dual antiplatelets in composite outcome, countered by an increase in life-threatening bleeds (2% vs 1%)</td>
</tr>
</tbody>
</table>

Note: Although antiplatelets are frequently prescribed for preventing non-cardioembolic strokes, the relatively low risk reduction in all trials underscores the need to target other risk factors that may increase stroke recurrence. + signifies that the agent was studied in the trial.

Abbreviations: ARR, absolute risk reduction; ASA, aspirin; ASA+D, aspirin plus dipyridamole; C, clopidogrel; PVD, peripheral vascular disease; RRR, relative risk reduction; ASA+C, aspirin plus clopidogrel.

Hypertension

Targeting hypertension carries the highest benefit in reducing stroke burden on a population level. The Perindopril Protection Against Recurrent Stroke Study (PROGRESS) trial was a randomized, double-blind, placebo-controlled trial of antihypertensive therapy among 6,105 patients with a history of hemorrhagic or ischemic stroke or TIA. Patients were treated with the angiotensin-converting enzyme inhibitor perindopril, with or without the addition of the thiazide diuretic indapamide. Therapy led to a mean blood pressure reduction of 9/4 mmHg and a 4% absolute risk reduction in recurrent stroke, with similar reductions in patients with and without a history of hypertension. The Secondary Prevention of Small Subcortical Strokes (SPS3) trial, aside from...
studying the effect of dual-antiplatelet therapy, also examined the role of strict vs standard blood pressure management (systolic <130 mmHg vs 130–149 mmHg) on the outcome of stroke recurrence and showed a trend toward benefit in stroke recurrence in the lower blood pressure group. Cur- rent American Heart Association guidelines recommend blood pressure goals of <140 mmHg and diastolic <90 mmHg; however, there is evidence that the benefit of lowering blood pressure extends to levels far below this cutoff, and as suggested in the PROGRESS trial, treatment should not be reserved for only those with history of hypertension.

Hyperlipidemia

Hypercholesterolemia is another general target in secondary stroke prevention. The Stroke Prevention by Aggressive Reduction in Cholesterol Levels (SPARCL) trial provided evidence of the benefit of statin therapy among patients with a TIA or stroke within the previous 6 months and no evidence of possible cardioembolic source (eg, atrial fibrillation). It enrolled 4,731 patients with a baseline low-density lipopro-tein cholesterol level between 100 and 190 mg/dL. Over a period of nearly 5 years, atorvastatin reduced the 5-year risk of recurrent stroke, from 13.1% to 11.2%. Although participants who took atorvastatin had a higher incidence of hemorrhagic stroke, there were no differences in mortality compared to placebo. There was also no difference seen in the efficacy of atorvastatin based on stroke mechanism. As a result, statins have become the mainstay in lipid reduction therapy after a stroke or TIA.

Diabetes

Diabetes is one of the most important vascular risk factors for stroke and a high-yield target for preventive measures. Among patients with diabetes, the risk of vascular events is increased thrice compared to nondiabetics, and in combinations with other risk factors, the risk increases exponentially compared to individuals with those risk factors without diabetes. Intensive glycemic control (defined as glycated hemoglobin <7%) was not associated with a significant reduction in the rates of stroke (fatal or nonfatal) among patients with diabetes, although there was a reduction of 16% in a combined vascular outcome. A recent meta-analysis of clinical trials comparing intensive glycemic control vs standard glycemic controls demonstrated a non-statistically significant 7% risk reduction among those in the intensive care group. An early, aggressive control of glycemia has lasting benefits in vascular events in patients with type I diabetes, and the long-term favorable results of intensive glucose lowering among those with type II diabetes argues for an early recognition of diabetes as the most effective way to reduce the risk of vascular events and stroke. Patients with diabetes frequently have other vascular risk factors that should be aggressively controlled. For example, selected patients with diabetes (ie, young patients) should have a blood pressure target of <130/80 mmHg if tolerated.

Lifestyle modification

Apart from pharmacological therapy, lifestyle modification, including a healthy diet, regular physical activity, and weight loss in overweight or obese patients, may have substantial benefits on blood pressure and lipid levels and, ultimately, stroke recurrence. Diet is probably the best studied when it comes to stroke prevention and the Mediterranean diet has been shown to protect against cardiovascular disease and specifically stroke, decreasing 5-year stroke risk by approximately 30%. In general, a diet that encourages a high intake of plant-based nutrients, low salt intake, and a limited intake of saturated fats and simple sugars is likely to have significant cardiovascular benefits if adhered to for a long period of time. Tobacco use should be strongly discouraged and among smokers, smoking cessation leads to a significant reduction in stroke risk. Furthermore, obesity is an independent risk factor for stroke, even after adjusting for physical activity and diet. Behavioral risk factors may be the most difficult to control, making patient education and a multidisciplinary approach extremely important.

Secondary stroke prevention by specific mechanisms

Cardioembolism

Atrial fibrillation

The biggest modifier of stroke risk in persons with atrial fibrillation is anticoagulation (Table 3). When atrial fibrillation from a non-valvular cause is discovered, long-term stroke risk can be quantified using the CHADS2 or CHADS2-VASc prediction scoring systems, with CHA2DS2-VASc being preferred in recent American and European guidelines. Included in both scores are age >75 years, history of congestive heart failure, hypertension, diabetes mellitus, and previous stroke. The sole presence of one risk factor (other than history of stroke) places the patient at a low risk of systemic cardioembolism, while an annual stroke risk of >1% based on the CHA2DS2-VASc score is typically considered the threshold to discuss the initiation of anticoagulation. This must be weighed against the risk of bleeding associated with anticoagulation.
When used for stroke prevention from atrial fibrillation, warfarin is associated with a 60%–70% relative risk reduction in stroke and has been the gold standard in primary and secondary stroke prevention in patients with known persistent or paroxysmal atrial fibrillation. More recently, a set of novel oral anticoagulants (NOACs) have been approved specifically for thromboembolism prophylaxis in patients with non-valvular atrial fibrillation. More recently, a set of novel oral anticoagulants (NOACs) have been approved specifically for thromboembolism prophylaxis in patients with non-valvular atrial fibrillation. Currently, there are four NOACs approved for prevention of stroke. They fall into the categories of either direct thrombin inhibitors or factor Xa inhibitors. They are dabigatran, rivaroxaban, apixaban, and edoxaban. Although their relative ease of use and lack of need for routine blood checks have made them popular first-line choices among some, candidates on the horizon for an assay that can quickly and reliably track level of anticoagulation. This may be especially important in those with renal impairment, which has been shown to increase the risk of hemorrhagic complications with some of these agents. The lack of an antidote to reverse the therapeutic effects of the NOACs may be offset by lesser risk of ICH compared to warfarin. Further, individualized risk of in-hospital mortality from ICH was studied in a retrospective analysis, and no difference in outcome was seen in patients experiencing ICH secondary to warfarin compared to in patients who had been on dabigatran. This suggests that the lack of an antidote should not dissuade physicians from using dabigatran (and presumably any of the other NOACs) over warfarin. Currently, PER977 – small synthetic cationic molecule – is being investigated as a possible antidote to all of the NOACs alike.

The greatest challenge in the prophylactic use of anticoagulation in the setting of atrial fibrillation revolves particularly for developing countries. There are, however, candidates on the horizon for an assay that can quickly and reliably track level of anticoagulation. This may be especially important in those with renal impairment, which has been shown to increase the risk of hemorrhagic complications with some of these agents. The lack of an antidote to reverse the therapeutic effects of the NOACs may be offset by lesser risk of ICH compared to warfarin. Further, individualized risk of in-hospital mortality from ICH was studied in a retrospective analysis, and no difference in outcome was seen in patients experiencing ICH secondary to warfarin compared to in patients who had been on dabigatran. This suggests that the lack of an antidote should not dissuade physicians from using dabigatran (and presumably any of the other NOACs) over warfarin. Currently, PER977 – small synthetic cationic molecule – is being investigated as a possible antidote to all of the NOACs alike. The greatest challenge in the prophylactic use of anticoagulation in the setting of atrial fibrillation revolves

**Table 3 Stroke mechanism and the estimated risk of stroke recurrence in the setting of prevention therapy**

<table>
<thead>
<tr>
<th>Stroke mechanism</th>
<th>Selected trials</th>
<th>Placebo/gold standard</th>
<th>Treatment</th>
<th>Outcome(s)</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Carotid stenosis</td>
<td>NASCET</td>
<td>Best medical</td>
<td>CEA</td>
<td>Stroke or death</td>
<td>ARR of 10% (6% vs 16%) with CEA vs best medical management at 1 year in patients with severe stenosis Composite outcome for CAS (7.2%) is non-inferior to CEA (6.8%) at 4 years</td>
</tr>
<tr>
<td>[23]</td>
<td>CREST</td>
<td>CEA</td>
<td>CAS</td>
<td>Stroke, MI, death</td>
<td>ARI of 2% (17% vs 15%) with warfarin compared to aspirin in patients with 50%–99% intracranial vessel stenosis ARI of 8% (20% vs 12%) with angioplasty/stenting compared to best medical practice alone in stenosis &gt;70%</td>
</tr>
<tr>
<td>Intracranial artery stenosis</td>
<td>WASID</td>
<td>Aspirin</td>
<td>Warfarin</td>
<td>Stroke or death</td>
<td>ARR of 9% (8% vs 17%) with warfarin compared to placebo/aspire</td>
</tr>
<tr>
<td>[12]</td>
<td>SAMMPRIS</td>
<td>Medical therapy</td>
<td>Angioplasty and stenting</td>
<td>Ischemic stroke</td>
<td>ARI of 8% (20% vs 12%) with angioplasty/stenting compared to best medical practice alone in stenosis &gt;70%</td>
</tr>
<tr>
<td>Cardioembolic</td>
<td>EAFT</td>
<td>Placebo/Aspirin</td>
<td>Warfarin</td>
<td>Stroke, death, MI, systemic embolism</td>
<td>ARR of 4% (15% vs 19%) with aspirin compared to placebo</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>EAFT</td>
<td>Placebo</td>
<td>Aspirin</td>
<td>Stroke or systemic embolism</td>
<td>ARR of 1.11% vs 1.71% per year favoring NOAC</td>
</tr>
<tr>
<td>[22]</td>
<td>RE-LY</td>
<td>Warfarin</td>
<td>Rivaroxaban</td>
<td>Stroke or systemic embolism</td>
<td>ARR of 1.27% vs 1.6% per year favoring NOAC</td>
</tr>
<tr>
<td>Heart failure</td>
<td>ARISTOTLE</td>
<td>Apixaban</td>
<td>Edoxaban</td>
<td>Death or stroke</td>
<td>2.1% vs 2.4% per year favoring NOAC</td>
</tr>
<tr>
<td>[15]</td>
<td>ENGAGE</td>
<td>Warfarin</td>
<td>Warfarin</td>
<td>Death or stroke</td>
<td>1.18% vs 1.5% per year favoring NOAC</td>
</tr>
<tr>
<td>Lacunar</td>
<td>SPS3</td>
<td>Aspirin + clopidogrel</td>
<td>Stroke</td>
<td>No difference in yearly risk (2.7% vs 2.5%) with CEA vs best medical management at 1 year in patients with severe stenosis Composite outcome for CAS (7.2%) is non-inferior to CEA (6.8%) at 4 years</td>
<td></td>
</tr>
<tr>
<td>[13]</td>
<td>CHANCE</td>
<td>Aspirin + clopidogrel</td>
<td>Stroke</td>
<td>ARR of 3.5% (8.2% vs 11.7%) with dual-antiplatelets compared to a single antiplatelet agent, over 3 months</td>
<td></td>
</tr>
</tbody>
</table>

**Notes:** The stroke recurrence rates vary by stroke mechanism, thus the need to investigate for a plausible cause of stroke. The natural history of the disease is modified with various therapies as shown in this table. Large artery atherosclerosis, intra- and extracranial, confers one of the highest risks of stroke recurrence, thus the need to intensively look for it. Atrial fibrillation, sometimes elusive, may be identified with prolonged cardiac monitoring after the acute phase of stroke. Given the dramatic change in treatment if atrial fibrillation is found, ruling out this arrhythmia is one of the most important steps in deciding the adequate treatment for stroke prevention. “Stroke” refers to both ischemic and hemorrhagic unless otherwise specified. Unlike the SPS3 trial, dual-antiplatelet therapy in CHANCE was initiated within 24 hours of symptoms onset.

**Abbreviations:** AR, atrial fibrillation; ARR, absolute risk reduction; CAS, carotid artery stenosis; CEA, carotid endarterectomy; CHF, chronic heart failure; EF, ejection fraction; MI, myocardial infarction; NOAC, novel oral anticoagulant; RRR, relative risk reduction.
around the risk of ICH vs the benefit of preventing ischemic stroke.59 The HAS-BLED (Hypertension, Abnormal renal/liver function, Stroke, Bleeding history or predisposition, Labile international normalized ratio, Elderly [≥65 years], Drugs/alcohol concomitantly) and ATRIA (Anticoagulation and Risk Factors in Atrial Fibrillation) prediction scoring systems are clinical decision-making tools that can aid the clinician in quantifying risk of hemorrhage, which can then be compared to yearly risk of stroke using CHADS2-VASc, to guide the clinician in how to best proceed.60,61

Until recently, patients with a high risk of bleeding had to be placed on less protective agents like aspirin, clopidogrel, or a combination of the two, as opposed to standard therapy with antiocoagulation. In March 2015, the WATCHMAN™ Left Atrial Appendage Closure Device was approved for use in the USA for the treatment of atrial fibrillation based on two trials showing non-inferiority to warfarin.62,63 Currently, the procedure is being offered to patients who carry a significant hemorrhagic risk with antiocoagulation or have other contraindications to antiocoagulation therapy. The caveat with using WATCHMAN is that patients need to be able to tolerate warfarin plus aspirin for at least 45 days (discontinued after there is proof of closure of the left atrial appendage) and dual antiplatelets for at least 6 months, which presents a challenge to the applicability of the WATCHMAN device to the FDA-approved high-risk population.63

Prosthetic cardiac valves
The risk of major embolism among patients with mechanical valves is estimated to be 4% per year without antithrombotic or antiocoagulation therapy, 2% with the use of aspirin, and 1% with the use of antiocoagulation.64 Anticoagulation with warfarin and occasionally with the additional low-dose aspirin (depending on comorbidities) is therefore indicated in patients with mechanical valves.65 Among patients with mechanical valves, NOACs are contraindicated.66 In patients with bioprosthetic valves, the risk of embolism is less compared to in those with mechanical valves, and a single antiplatelet is usually recommended for secondary stroke prophylaxis, although a short course of antiocoagulation after implantation may be reasonable.65,67

Low ejection fraction
After an anterior wall MI, a third of patients develop a thrombus and a third of these have embolization.68 Predictors of left ventricular thrombus are low ejection fraction (EF), worsening EF after discharge, and wall dyskinesis.69 The most consistent predictor of embolization is thrombus mobility.58,69 It is considered reasonable to use short-term anticoagulation in patients with MI and evidence of left ventricular thrombus and in those with wall dyskinesis, but randomized data to universally recommend this intervention are lacking.70

Anticoagulation with warfarin in patients with dilated cardiomyopathy with an EF <35% and a sinus rhythm was studied in one large trial warfarin and aspirin in patients with heart failure and sinus rhythm (WARCEF) and while the warfarin group had a lower incidence of ischemic stroke, any benefit was offset by an overall increase in major hemorrhage.16 Among WARCEF participants <60 years old, however, there was a benefit with antiocoagulation vs aspirin, even after taking into account the rates of bleeding.71 In further subgroup analysis, those with prior stroke and an EF <15% had a stroke risk of 6% per year, suggesting that in this high-risk group, the preventive effects of warfarin may offset the risk of hemorrhage.72 As a result, warfarin is currently not indicated for the sole purpose of stroke prevention in dilated cardiomyopathy with low EF and its use in some of the mentioned subgroups should be carefully discussed with the patients and their families.

Aortic atheroma
Complex aortic arch plaques, defined by a protruding component >4 mm, presence of a mobile component, or intraplaque ulceration, are deemed high risk for embolization.73 The 2-year risk of stroke or death among individuals with aortic plaques <4 mm was 16.5% compared to 26.7% in those with plaques ≥4 mm.74 In this observational study, the rate of events was similar among those using antiplatelets or warfarin. Whether antiocoagulation reduces incident vascular events compared to antiplatelets in patients with aortic plaques >4 mm has been formally explored in a clinical trial.75 The primary endpoint (cerebral infarction, MI, peripheral embolism, vascular death, or ICH) in the group on antiocoagulation occurred in 11.0% vs 7.6% in the dual-antiplatelet group, failing to reach statistical significance (probably due to the low rate of events). Further, the group on antiocoagulation had a higher incidence of vascular death and ICH. There is therefore no clear evidence to support the systematic use of antiocoagulation in patients with aortic arch atheroma.

Patent foramen ovale
The role of antiocoagulation among patients with possible paradoxical embolism through a patent foramen ovale (PFO) is unclear, and current guidelines support the use of antiplatelet agents for secondary prevention in this setting.76
Three large clinical trials have investigated whether closure of PFOs using trans-cardiac devices can reduce secondary stroke risk compared to medical therapy. None of them was able to show benefit of PFO closure when compared to medical therapy alone, in the intention-to-treat analysis.\textsuperscript{77–79} Importantly, the yearly rates of stroke differed among these trials. In the closure or medical therapy for cryptogenic stroke with patent foramen ovale (CLOSURE) I trial, the 2-year rate of stroke recurrence was 7% in the medical group, while in the closure of patent foramen ovale versus medical therapy after cryptogenic stroke (RESPECT) and percutaneous closure of patent foramen ovale in cryptogenic embolism (PC) trials (with more stringent criteria for ruling out alternative stroke mechanisms), the rates were closer to 1% per year. This discrepancy is in itself a challenge to the interpretation of these trials, but suggests that in studies reporting a high rate of stroke recurrence attributed to a PFO, participants have alternative stroke mechanisms that are not modified by closing a PFO. Ideally, clinical trials assessing whether closing the PFO reduces the risk of stroke recurrence would benefit from using a probabilistic model like the RoPE (Risk of Paradoxical Embolism) score to identify those with the highest likelihood for a causal role of the PFO, after exhaustive workup to rule out alternative etiologies.\textsuperscript{80} Until stronger data exist to justify the use of a device to close the PFO and its potential risks, a single daily antiplatelet seems a reasonable recommendation in patients with truly cryptogenic strokes.

**Extracranial carotid atherosclerosis**

Extracranial carotid artery stenosis is most effectively treated with endarterectomy or stenting (Figure 1). There have been a number of studies comparing best medical management vs carotid endarterectomy (CEA) in patients with symptomatic carotid stenosis. The North American Symptomatic Carotid Endarterectomy Trial demonstrated a risk reduction of 17% in all ipsilateral stroke recurrence, and 11% in ipsilateral major or fatal strokes.\textsuperscript{81} Post hoc analysis of this study demonstrated that the greatest value of CEA comes if the procedure is done within 2 weeks of the stroke.\textsuperscript{82} The European Carotid Surgery Trial showed a comparable risk reduction in the surgical arm for patients with carotid stenosis >80%. An analysis of pooled data for symptomatic carotid stenosis trials demonstrated robust beneficial effects of surgical interventions across trials in stenosis of ≥70%, with more marginal effects with lesser degrees of stenosis or in those with carotid occlusion.\textsuperscript{83} Among patients with carotid occlusion and evidence of increased oxygen extraction in the ipsilateral hemisphere (ie, at the greatest risk of recurrent stroke), an extracranial–intracranial arterial bypass failed to reduce the risk of stroke recurrence in one study.\textsuperscript{84}

Three large trials have assessed whether carotid artery stenting (CAS) is as equally effective as CEA among patients with symptomatic extracranial carotid atherosclerosis.\textsuperscript{14,85,86} The results consistently show a slight increase in periprocedural stroke (30 days) with CAS compared to CEA. Combined adverse outcomes over a 2-year period are similar, however, establishing CAS as a viable option for treating severe symptomatic carotid stenosis in high-risk surgical patients or those with unsuitable anatomy.

In patients with asymptomatic carotid stenosis, older evidence suggested a benefit in CEA vs medical therapy, but the effect size was more modest and the benefit delayed, compared to the benefit observed for patients with symptomatic carotid stenosis.\textsuperscript{87,88} Furthermore, the advent of statins and more aggressive medical therapy may have reduced the differential effects of surgical intervention compared to modern medical therapy. Based on this presumed medical equipoise, a new trial will investigate the effect modification of CEA or carotid stenting against medical therapy in patients with asymptomatic carotid stenosis.\textsuperscript{89}

**Intracranial LAA**

Several trials have investigated the best possible treatment in stroke recurrence in the setting of intracranial vessel stenosis (Figure 2). The Warfarin and Aspirin Symptomatic Intracranial Disease (WASID) trial studied aspirin vs warfarin in patients with verified 50%–99% stenosis of a major intracranial artery (carotid, middle cerebral, vertebral, or basilar arteries).\textsuperscript{13} Even though the study was stopped early because of the high rate of adverse events in the warfarin group, it showed a 20% 2-year risk of overall stroke recurrence and a 15% 2-year risk of stroke recurrence in the vascular territory distal to the stenosis.\textsuperscript{13}

To directly address this high-risk group of patients, the Stenting and Aggressive Medical Management for Preventing Recurrent Stroke in Intracranial Stenosis (SAMPRI) trial attempted to follow the therapeutic model used in coronary artery disease by comparing percutaneous transluminal angioplasty and stenting to best medical therapy in patients with recent stroke and proximal intracranial arterial stenosis of >70%.\textsuperscript{90} Both groups received aggressive lifestyle modification counseling and intervention and were placed on dual-antiplatelet therapy with aspirin and clopidogrel for 3 months. The final outcome of stroke or death occurred in 20% in the stenting group and 12% in the medical group at
1 year. The majority of strokes in the intervention arm were perioperative, occurring in the first 30 days. Although a recurrence of 12% per year among those with medical therapy is intolerably high, aggressive medical therapy appears to be the only proven therapy recommended for patients with intracranial LAA. Novel therapies are evidently and urgently needed to further reduce the risk of stroke recurrence among this population.

Small artery occlusion/lacunar stroke
Antiplatelet therapy with clopidogrel and aspirin, compared to aspirin alone, was investigated as secondary prevention after lacunar stroke in the SPS3 trial. There was no evidence of a reduction in the rates of recurrent strokes using dual antiplatelets vs aspirin alone (2.5% vs 2.7% per year), but an increase in the risk of bleeding and mortality was noted in the dual-antiplatelet therapy group. Based on this, there is no evidence to support the use of dual antiplatelets to reduce the risk of recurrent stroke in the long-term.

In the short-term, however, there may be a role for the use of dual antiplatelets in “minor” strokes or TIA. The Clopidogrel in High-risk Patients with Acute Non-disabling Cerebrovascular Events (CHANCE) trial found that a 21-day course of dual-antiplatelet therapy with aspirin and clopidogrel among Chinese patients with TIA and minor stroke, given within 24 hours of symptom onset, lowered the 3-month stroke recurrence rate. The majority of the benefit from dual-antiplatelet therapy in CHANCE occurred within the first week of treatment. A short course of aspirin and clopidogrel in the early period may therefore be considered, but whether these results are applicable to non-Chinese populations will be further investigated in the Platelet-Oriented Inhibition in New TIA (POINT) trial.

Future directions/Conclusion
Preventing stroke recurrence is one of the major concerns of practitioners who treat patients with stroke, either in the hospital or in the clinic. Determining the potential causal mechanisms of a patient’s stroke offers the opportunity to establish tailored therapies, while in the hospital or shortly after, for extracranial carotid stenosis, intracranial large artery stenosis with flow failure, cardiac thrombus in the setting of low EF and/or atrial fibrillation. Defining stroke mechanism, however, is challenging and, as discussed, it is crucial that physicians exhaustively rule out alternative etiologies for stroke. In the case of mixed etiologies (eg, intracranial atherosclerosis in patient with atrial fibrillation), the proven treatment for the conditions with the highest risk of recurrence might be indicated. A comprehensive stroke workup is expensive, and it may not be within reach in developing countries where stroke represents an even greater challenge. Developing cheaper methods to more accurately ascertain stroke mechanism may be a way to overcome this challenge.

Furthermore, although clinical trials have framed and keep fine-tuning the major therapeutic axis for stroke prevention, there are still many uncertainties in specific conditions with higher risk of stroke. These less frequent, and consequently less well-studied, conditions or clinical scenarios lead to heterogeneous patterns of practice that may result in more expensive treatment or even worse – harm to patients. Collaborations across major academic centers and integrated electronic systems may offer an opportunity to study outcomes in some of these cases, which may lead to a more unified and better informed approach to treat patients with stroke.

Finally, in our view, the greatest challenge is that, even with the best medical therapies, the recurrence rates of stroke are not zero, and one out of four patients who have had strokes in the USA has a recurrent event. It is imperative to foster the development of newer therapies to reduce even further the risk of stroke recurrence. Given the unacceptably high risk of stroke recurrence with traditional vascular disease, a stronger effort is needed in controlling vascular risk factor on a population-based level.

Disclosure
The authors report no conflicts of interest in this work.

References


Secondary stroke prevention: challenges and solutions


