Potential impact of new oral anticoagulants on the management of atrial fibrillation-related stroke in primary care

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SUMMARY

Aim: Antiplatelet prophylaxis with vitamin K antagonists (such as warfarin) is effective in reducing the risk of stroke in patients with atrial fibrillation (AF). New oral anticoagulants have emerged as potential alternatives to traditional oral agents. The purpose of this review was to summarise the effectiveness and safety of rivaroxaban, dabigatran and apixaban in stroke prevention in patients with AF in phase III trials, evaluate their cost-effectiveness and consider the implications for primary care. Methodology: A literature search was performed between 2007 and 2012, selecting all phase III trials (ROCKET AF, RE-LY and ARISTOTLE) of new oral anticoagulants and relevant cost–benefit studies. Results: Evidence shows that all three agents are at least as effective as warfarin in the prevention of stroke and systemic emboli, with similar safety profiles. Cost–benefit studies of rivaroxaban and dabigatran further confirm their potential use as alternatives to warfarin in clinical practice. These observations may allow stratification of the general practice AF population, to help prioritise which patients may benefit from receiving a new oral anticoagulant. Conclusion: The clinical and economic benefits of the new oral anticoagulants, along with appropriate risk stratification, may enable a higher number of patients with AF to receive effective and convenient prophylaxis for stroke prevention.

Introduction

Each year, approximately 110,000 people in England have a stroke, at a cost to the economy of around £7 billion (1). Atrial fibrillation (AF) is a known risk factor for stroke, imparting a fivefold increase in risk (2), and AF has a prevalence of around 1–2.5% (3,4). The risk of AF is strongly associated with old age, and prevalence rises to over 9% in people in their 80s (5).

Anticoagulation with warfarin substantially reduces the risk of stroke in patients with AF (6,7), but is associated with an increased risk of intracranial haemorrhage and other bleeding (8). Warfarin continues to be ‘underused’ compared with guideline-recommended care. In a study of health maintenance organisation patients in North Carolina, USA, less than 60% of patients with one or more risk factors for stroke and no contraindications were receiving warfarin (9). Another large study of patients with AF discharged from hospital found that only 64.6% of ideal candidates were prescribed warfarin (10).

A review of 310 practice populations in the UK showed that only 27% of high-risk (CHADS2 score > 1) patients with AF not taking warfarin had a contraindication for its use (11). Low usage of warfarin is in part caused by high discontinuation rates. For example, in the Birmingham Atrial Fibrillation Treatment of the Aged (BAFTA) trial, a third of patients randomised to warfarin had stopped taking it after an average follow up of 2.7 years (12). In addition, physicians may not initiate warfarin therapy because of a fear of intracranial haemorrhage, especially in elderly patients (11,13). Warfarin may be associated with further complications in patients with comorbidities associated with haemorrhagic risk (13), and some patients are deterred by the need for regular coagulation monitoring and the risk of interactions with food and other drugs (14).

Until recently, alternatives to warfarin have been unsatisfactory. Antiplatelet agents, although easier to use, are significantly less effective (15), even when used in combination (16). However, three promising
new oral anticoagulants (OACs) (17–19) have emerged: dabigatran etexilate, a direct thrombin inhibitor, and rivaroxaban and apixaban, which are both direct Factor Xa inhibitors. Both rivaroxaban and dabigatran are approved in the EU and USA, as well as in other countries, for stroke prevention in patients with AF. A decision by the US Food and Drug Administration on the use of apixaban in patients with AF has been delayed until 2013. There is also a potential fourth agent (edoxaban), for which a phase II dose-finding trial has been conducted (20). In this article, we review the evidence of the effectiveness and safety of the three agents for which phase III trials have been published, and consider their cost-effectiveness and the potential implications for practice in primary care.

**Methods**

We searched MEDLINE using the MeSH terms ‘anticoagulants’ and ‘atrial fibrillation’ and ‘clinical trial’, limiting our search to publications from 2007 to May 2012. We then reviewed titles and abstracts to find all phase III trials of new OACs. We then ran a separate MEDLINE search using the MeSH terms ‘anticoagulants’ and ‘atrial fibrillation’ and ‘cost–benefit analysis’, again limiting our search to publications from 2007 to May 2012 and reviewing titles and abstracts to find relevant papers.

**Safety, effectiveness and side effect profile**

The key findings from the trials that tested these drugs against warfarin are summarised in Tables 1 and 2. Direct cross-trial comparisons are problematic owing to differences in trial designs (i.e. open-label vs. double-blind), statistical analyses and baseline stroke risk in the study populations. Nonetheless, all three new OACs were at least as effective as warfarin in preventing strokes and systemic emboli, and within the context of the individual trials, apixaban and higher dose dabigatran were superior to warfarin (17–19). Furthermore, the risk of death was reduced by approximately 10% compared with warfarin (21). The main safety concern for anticoagulation therapy is an increased risk of bleeding. Rivaroxaban and higher dose dabigatran were associated with similar risks of major bleeding compared with warfarin, whereas apixaban and lower dose dabigatran were associated with lower risks (17–19). All three new OACs were associated with a lower risk of intracranial haemorrhage compared with warfarin (17–19).

When major bleeding occurs in patients taking warfarin, anticoagulation can be reversed by administering vitamin K and fresh frozen plasma. Specific antidotes for the new OACs do not exist and management is largely supportive, given the relatively short half-lives of these drugs (22). Nevertheless, this may be a source of concern that will influence clinicians’ and patients’ choice of anticoagulant.

Regular international normalised ratio (INR) monitoring is needed for optimal anticoagulation using warfarin, and this can be a source of concern and inconvenience for patients (14,23). The three new OACs are either once-daily (rivaroxaban) or twice-daily (dabigatran and apixaban) regimens that do not require routine anticoagulation monitoring.

The main side effects of the new OACs relate to minor bleeding events, as may be expected; these may not be dangerous, but could impact on patient quality of life. Rates of gastrointestinal bleeding were similar or higher for all three new OACs compared with warfarin. Rivaroxaban showed an increased incidence of haematuria and epistaxis in ROCKET AF (Rivaroxaban Once daily, oral, direct factor Xa inhibition Compared with vitamin K antagonism for prevention of stroke and Embolism Trial in Atrial Fibrillation) (18), and in RE–LY (Randomized Evaluation of Long-term anticoagulation therapY) dabigatran showed higher rates of dyspepsia (17) compared with warfarin. A breakdown of adverse events is not available for apixaban, but the ARISTOTLE (Apixaban for Reduction In STroke and Other ThromboemboLic Events in atrial fibrillation) trial investigators reported that the total was similar to that seen with warfarin (19). In the three trials, the discontinuation rates of the new OACs were similar to or higher than (in the case of dabigatran) the discontinuation rates observed in patients receiving warfarin; however, it is difficult to extrapolate from trial conditions what discontinuation rates would equate to in clinical practice. More patients taking dabigatran in the RE–LY trial had a myocardial infarction than those receiving warfarin (17), although a subsequent re-evaluation after detection of several additional primary efficacy and safety outcome events found that the difference was not statistically significant, with a revised relative risk of myocardial infarction in the dabigatran 150 mg twice-daily group of 1.27 (95% confidence interval 0.94–1.71; p = 0.12) compared with warfarin (24). Nevertheless, a recent meta-analysis involving trials of use of dabigatran for other indications suggested an increase in observed rates of myocardial infarction compared with the controls used (25). As noted above, comparisons between the different new OACs are not straightforward because the studies had different populations and designs, and there have been no head-to-head comparison trials. Therefore, in this article, we have emphasised differences between
Table 1 Overview of efficacy data from phase III trials investigating rivaroxaban, dabigatran or apixaban compared with warfarin for the prevention of stroke and systemic embolism in patients with AF

| Study          | Method                                      | Population                                                                 | Mean CHADS2 score | Drug             | Hazard ratio (intention to treat; 95% CI) | Stroke (all types) or systemic embolism | Myocardial infarction | Deaths      |
|---------------|---------------------------------------------|-----------------------------------------------------------------------------|-------------------|------------------|-------------------------------------------|----------------------------------------|-----------------------|-------------|
| ROCKET AF (18)| Double-blind, double-dummy, randomised, non-inferiority trial | 14,264 patients with non-valvular AF with either a history of stroke, TIA or systemic embolus, or a CHADS2 score of ≥ 2 from 1178 participating sites in 45 countries. Mean age 73 years | 3.5               | Rivaroxaban 20 mg od* | 0.88 (0.75–1.03) p < 0.001 for non-inferiority, p = 0.12 for superiority | 0.81 (0.63–1.06) p = 0.12† | 0.92 (0.82–1.03) p = 0.15 |
| RE-LY (17)    | Open-label, randomised, non-inferiority trial | 18,113 patients with AF and risk factors for a stroke, from 951 centres in 44 countries. Mean age 71 years | 2.1               | Dabigatran 150 mg bid | 0.66 (0.53–0.82) p < 0.001 for non-inferiority, p < 0.001 for superiority | 1.38 (1.00–1.91) p = 0.048 | 0.88 (0.77–1.00) p = 0.051 |
| ARISTOTLE (19)| Double-blind, double-dummy, randomised, non-inferiority trial | 18,201 patients with AF and ≥ 1 additional risk factor for stroke from over 1000 centres in 39 countries. Median age 70 years | 2.1               | Apixaban 5 mg bid‡ | 0.79 (0.66–0.95) p < 0.001 for non-inferiority, p = 0.01 for superiority | 0.88 (0.66–1.17) p = 0.37 | 0.89 (0.80–0.99) p = 0.047 |

* Patients with creatinine clearance 30–49 ml/min received rivaroxaban 15 mg od. † This hazard ratio is for the as-treated safety population. The hazard ratio for myocardial infarction for the intention-to-treat population was not presented in ROCKET AF. All p-values are for superiority unless stated otherwise. All three trials express the end-point event rates as % per year. ‡ Patients with serum creatinine levels of > 1.5 mg/dl received apixaban 2.5 mg bid. AF, atrial fibrillation; bid, twice daily; CI, confidence interval; od, once daily; TIA, transient ischaemic attack.
warfarin and the new OACs, rather than differences between these agents.

### Cost-effectiveness

We found five cost–benefit analyses for dabigatran, including a UK National Institute for Health and Care Excellence (NICE) technology appraisal, and for rivaroxaban, a NICE technology appraisal only (Table 3). Incremental cost-effectiveness ratio (ICER) estimates for dabigatran 150 mg compared with warfarin ranged from £5609 to £56,911 per quality-adjusted life-year (QALY) (26–29). The cost-effectiveness of dabigatran increased in groups more at risk of stroke, such as older patients: Freeman et al. (26) used the youngest patient group and estimated the second-highest ICER/QALY (for 150 mg twice-daily dose). Drug costs for dabigatran used in these analyses varied with a range of £1.99–€8.30 (based on 27 May 2012 exchange rates from http://markets.ft.com), with Shah et al. (28) and Freeman et al. (26) assuming the highest daily drug cost. The balance of cost-effectiveness is sensitive to the assumed cost of stroke treatment and follow up, which is much higher in the report of Sorensen et al. (29), which partly accounts for the much lower ICER estimate for dabigatran than in the other studies. The NICE technology appraisal for rivaroxaban found that it was likely to be cost-effective for adults with AF and one or more risk factors for stroke, with ICERs of less than £29,500/QALY (30).

The dabigatran technology appraisal by NICE considered the use of dabigatran in a two-tier regimen of 150 mg twice daily before the age of 80 followed by a switch to 110 mg twice daily after the age of 80 for stroke prevention in patients with AF and at least one additional risk factor for stroke. This strategy was found to be cost-effective for the UK National Health Service (NHS), with ICERs of less than £18,900/QALY in patients starting treatment younger than age 80, assuming a monitoring cost of £241.54.

### Table 2 Overview of safety data from phase III trials investigating rivaroxaban, dabigatran or apixaban compared with warfarin for the prevention of stroke and systemic embolism in patients with AF

| Study      | Drug          | Discontinuation: study drug vs. warfarin (%) | Hazard ratio (95% CI) | Side effects occurring significantly more in study drug vs. warfarin (%) |
|------------|---------------|---------------------------------------------|-----------------------|-----------------------------------------------------------------------|
|            |               |                                             | Major bleeding        | Intracranial bleeding | Gastrointestinal bleeding | Epistaxis 10.14 vs. 8.55, p < 0.05; haematuria 4.16 vs. 3.40, p < 0.05 |
| ARISTOTLE  | Apixaban      | 25.3 vs. 27.5                               | 0.69 (0.60–0.80)      | p < 0.001               | 0.89 (0.70–1.15)          | Dyspepsia 11.3 vs. 5.8, p < 0.001 |
|            | 5 mg bid      |                                             |                       |                         |                          | Dyspepsia 11.8 vs. 5.8, p < 0.001 |
| RE-LY (17) | Dabigatran    | 21.2 vs. 16.6                               | 0.93 (0.81–1.07)      | p = 0.31                 | 1.50 (1.19–1.89)          | No breakdown of adverse events provided, but total adverse events occurred in almost equal proportions |
|            | 150 mg bid    |                                             |                       |                         |                          |                                      |
|            | 110 mg bid    |                                             |                       |                         |                          |                                      |
| ROCKET AF  | Rivaroxaban   | 23.7 vs. 22.2                               | 1.04 (0.90–1.20)      | p = 0.58                 | 1.45†                    | Epistaxis 10.14 vs. 8.55, p < 0.05 |
|            | 20 mg od‡     |                                             |                       |                         |                          | haematuria 4.16 vs. 3.40, p < 0.05 |

* Patients with creatinine clearance 30–49 ml/min received rivaroxaban 15 mg od. †Relative risk calculated from data in supplementary table; 224 bleeding events (3.2%) in rivaroxaban group compared with 154 events in the warfarin group (2.2%, p < 0.001). ‡Patients with serum creatinine levels of > 1.5 mg/dl received apixaban 2.5 mg bid. AF, atrial fibrillation; bid, twice daily; CI, confidence interval; od, once daily.

Definitions of bleeding: RE-LY: Major bleeding was defined as a reduction in the haemoglobin level of ≥ 2 g/dl, transfusion of ≥ 2 units of blood or symptomatic bleeding in a critical area or organ.

ROCKET AF: Major bleeding was defined as clinically overt bleeding associated with any of the following: fatal outcome, involvement of a critical anatomical site, fall in haemoglobin concentration ≥ 2 g/dl, transfusion of ≥ 2 units of whole blood or packed red blood cells or permanent disability. Non-major clinically relevant bleeding was defined as overt bleeding not meeting the criteria for major bleeding, but requiring medical intervention, unscheduled contact (visit or telephone) with a physician, temporary interruption of study drug (i.e. delayed dosing), or causing pain or impairment of daily activities.

ARISTOTLE: Major bleeding was defined as clinically overt bleeding associated with any of the following: fatal outcome, occurring at a critical site, decrease in the haemoglobin level of ≥ 2 g/dl, transfusion of ≥ 2 units of blood or symptomatic bleeding in a critical area or organ.
The original manufacturer’s submission to NICE estimated an INR cost of £414.90: this was based on the assumption that dabigatran would completely replace warfarin, so includes the fixed overheads of running anticoagulation clinics. The monitoring cost per patient used in the NICE final appraisal determination was estimated using the NICE AF costing report from 2006 (32) and the NHS reference costs for 2008/09 (33), both inflated to 2009/10 prices, but because dabigatran would be unlikely to completely replace warfarin, this figure was reduced to account for the fixed costs of running anticoagulation clinics (34).

As we performed our search, three further cost-effectiveness analyses have been published comparing warfarin with dabigatran in the general AF population, set in Denmark (35), Sweden (36) and the UK (37). All concluded that dabigatran was cost-effective in comparison with warfarin, with an ICER of €7000 per QALY in the Danish study (35), €7700 in the Swedish study (36) and approximately £4800–£7000 per QALY in the UK study (37). There has also been a US cost-effectiveness study that compared rivaroxaban with warfarin, and showed rivaroxaban to be cost-effective with an ICER of $27,500 per QALY (38).

Warfarin has previously been found to be cost-effective for stroke prevention in patients with AF (39,40). This was confirmed in an older age group by a recent economic evaluation conducted alongside the BAFTA trial (41). Cost-effectiveness for warfarin is dependent on the level of INR control achieved. Good INR control can be achieved in routine care with values within the therapeutic range around 68% of the time (42,43). The NICE technology appraisal for dabigatran estimated an ICER cost of £47,000/QALY for the subset of patients with the best INR control, although this figure was substantially reduced if the higher INR monitoring costs of £414.90 were assumed (31). There is a substantial group of patients for whom anticoagulation is indicated, but who prefer not to take warfarin. The ACTIVE A (Atrial fibrillation Clopidogrel Trial with Irbesartan for prevention of Vascular Events) trial used a population of patients with AF who were deemed unsuitable for vitamin K antagonist therapy; however, one of the main reasons for this was patient preference not to take warfarin, which was

| Drug         | Study                  | Population                                                                 | Sequential regimen of 150 mg bid dabigatran before age 80 followed by 110 mg bid afterwards | Dabigatran 110 mg bid | Dabigatran 150 mg bid | Rivaroxaban 20 mg or 15 mg od |
|--------------|------------------------|----------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------|-----------------------|-----------------------|-----------------------------|
| Dabigatran   | Sorensen et al. (29)   | Patients with AF and at ≥ 1 additional risk factor for stroke or impaired left ventricular ejection fraction. Mean CHADS2 score 2.1; mean age at starting 69 years | 18,608                                                            | 5609                  |                       |                             |
| Dabigatran   | Pink et al. (27)       | Patients at moderate to high risk of stroke with AF and a baseline CHADS2 score of 2.1; mean age at starting 71 years | 43,074                                                            | 23,082                |                       |                             |
| Dabigatran   | Freeman et al. (26)    | Patients aged 65 years at starting with non-valvular AF and CHADS2 score of ≥ 1 | 32,710                                                            | 28,970                |                       |                             |
| Dabigatran   | Shah et al. (28)       | Patients aged 70 at starting with AF at moderate risk of stroke (CHADS2 score of 1 or 2) | 95,775                                                            | 56,911                |                       |                             |
| Dabigatran   | NICE technology appraisal guidance (31) | Population reflects that of RE-LY trial, i.e. adult patients with AF and ≥ 1 additional risk factor for stroke and eligible for anticoagulation | 18,900                                                            |                       |                       |                             |
| Rivaroxaban  | NICE technology appraisal guidance (30) | Population reflects that of ROCKET AF, i.e. adult patients with AF who were at moderate to high risk of stroke (CHADS2 score ≥ 2) | < 29,500                                                          |                       |                       |                             |

*Exchange rates based on 27 May 2012 rates from http://markets.ft.com
AF, atrial fibrillation; bid, twice daily; NICE, National Institute for Health and Care Excellence; od, once daily; QALY, quality-adjusted life-year.
the case in approximately one in four patients enrolled (44). In these patients, new OACs are likely to be highly cost-effective compared with using an antiplatelet agent or no therapy.

**Implications for primary care**

The new OACs offer genuine alternatives to warfarin and exhibit similar or better efficacy, better safety in some parameters (e.g. intracranial haemorrhage) and are easier to use. The safety data from the trials show a profile of non-inferiority to warfarin, but drugs that have appeared to have a good safety profile in clinical trials in the past have later been withdrawn (45–47), whereas warfarin is a well-established drug that can be difficult to use. Key determinants of cost-effectiveness are quality of INR control and cost of INR monitoring on warfarin (14,29). Different models of INR monitoring have different costs, for example warfarin dosing in primary care costs less than in secondary care (32).

The NICE technology appraisals have major potential implications for primary care because they support the use of dabigatran and rivaroxaban in most circumstances. However, even if new OACs are cost-effective, this does not mean that they are affordable and total cost may be more important to local commissioning groups than cost-effectiveness, making the introduction of a new drug problematic in the current economic environment. For example, one costing study suggested that the annual cost of anticoagulation is about seven times higher with dabigatran than it is with warfarin (48). With this in mind, we have divided the general practice AF population into seven groups according to risk of stroke and previous experience of warfarin, and prioritised these groups in order of cost-effectiveness of prescribing a new OAC. We have quantified the number of people who would be in each group for an average-sized practice (Figure 1). This would suggest that for a typical practice of 6600 people, starting a new OAC should be considered in 14/65 (22%) of patients on the AF register – i.e. in those people who have previously tried warfarin, or in those for whom the INR control is poor. It is worth noting that we have assumed that 88% of patients have good INR control, which is based on information from the ACTIVE-W trial (49). It is likely that this figure

![Figure 1](https://example.com/figure1.png)

**Figure 1** Quantifying workload in general practice for introducing new anticoagulants by different categories of AF patient. Patients were categorised into seven groups (A–G), and assigned an order of priority (1–4) for receiving a new OAC. The following assumptions were made: the average General Practice population size in England is 6600 (54), AF has a population prevalence of approximately 1% (3), the incidence of newly diagnosed cases of AF is 0.6 per 1000 (55), approximately 90% of patients with AF are at high or moderate risk of a stroke/transient ischaemic attack using the CHADS2 score (50); approximately 47% of patients who should be receiving warfarin are not (58); patients with CHADS2 scores ≥ 2 are not receiving warfarin (59,60), 88% of UK participants have a mean time in therapeutic range of > 65% (49); discontinuation rates are > 25% in the first year for patients with AF started on warfarin (62). AF, atrial fibrillation; OAC, oral anticoagulant.
A patient’s risk of stroke can be quantified using CHADS₂ or CHA₂DS₂-VASc scores (Table 4) (50,51). Previously, patients were stratified into high-, medium- or low-risk groups, with only patients in the high-risk group (CHADS₂ score ≥ 2) recommended to receive anticoagulation (1). However, there is a reduction in stroke risk from anticoagulation even for patients with only one risk factor for stroke (52). We are now moving towards a binary system involving consideration of anticoagulation in patients with a CHADS₂ score of ≥ 1 if the CHA₂DS₂-VASc score is ≥ 1. This system results in better identification of a low-risk group for whom the risk of stroke is small (Table 5). We note, however, that while the NICE technology appraisal recommends dabigatran for patients with AF and one or more additional risk factors for stroke, neither female sex nor peripheral vascular disease constituted one of the risk factors (31) (both are included as additional risk factors in the CHA₂DS₂-VASc score, Table 4). For the purposes of this review, we have defined moderate to high risk as the presence of one additional risk factor for stroke and assumed consideration of anticoagulation in these individuals, with the benefits of anticoagulation increasing as risk of stroke increases.

Table 4 Comparison of CHADS₂ and CHA₂DS₂-VASc stroke risk scoring systems

| Risk factor                  | CHADS₂ score (50) | CHA₂DS₂-VASc score (51) |
|------------------------------|-------------------|--------------------------|
| Congestive heart failure     | 1                 | 1                        |
| Hypertension                 | 1                 | 1                        |
| Age ≥ 75 years               | 1                 | 2                        |
| Diabetes mellitus            | 1                 | 1                        |
| Previous stroke or TIA       | 2                 | 2                        |
| Vascular disease             | –                 | 1                        |
| Age 65–74                    | –                 | 1                        |
| Female sex category          | –                 | 1                        |

TIA, transient ischaemic attack.

Assumptions made to arrive at the numbers for the different population groups in Figure 1

- Average GP population size is 6600 in England (54)
- AF has a population prevalence of approximately 1% (3,4)
- The incidence of a new case of AF is 0.6 per 1000 (55); this has been estimated at 1.7 per 1000 for chronic AF in general practice (56) and 2.9 per 1000 for all cases of AF (57)

Approximately, 90% of patients with AF are at high or moderate risk of a stroke/transient ischaemic attack using the CHADS₂ score (50). This number may be slightly less if only patients with a CHADS₂ score of ≥ 2 were included

A 2006 costing report by NICE estimated that 47% of patients who should have been receiving warfarin were not, based on an assumption that all high-risk and half of moderate-risk patients with AF should be on anticoagulation (58).
More recent NHS improvement audits and a study of the QResearch database have found that similar percentages of patients with CHADS2 scores of ≥ 2 are not receiving warfarin (59–61). If we include all patients with ≥ 1 additional risk factor for stroke, it is likely that no more than 50% will be receiving warfarin.

In a post hoc analysis of patients on warfarin in the ACTIVE-W trial, 88% of UK participants had a mean time in therapeutic range of > 65%; this figure may be lower in the general population (49).

Fang et al. (62) found discontinuation rates of more than 25% in the first year for patients started on warfarin for AF in the ATRIA (Anticoagulation and Risk factors In Atrial fibrillation) cross-sectional study. Assuming this rate of discontinuation and that 30 patients per GP are currently taking warfarin, 10 of the patients currently not taking warfarin at a GP surgery may have previously tried it. In the AVERROES (Apixaban VERsus acetylsalicylic acid to prevent stRokEs) trial, which compared apixaban with acetylsalicylic acid in patients with AF with an additional risk factor for stroke but deemed unsuitable for warfarin, 40% of patients had previously tried a vitamin K antagonist, which helps corroborate our estimate (63).

Conclusions

Anticoagulant prophylaxis with warfarin is highly effective in reducing the risk of thromboembolic stroke risk in patients with AF; however, it continues to be underused for a variety of reasons, including patient and physician reluctance to initiate treatment, and problems incurred while on treatment. The availability of new OACs as cost-effective alternatives to warfarin will mean that in the future a higher proportion of people with AF and at high risk of stroke can receive effective stroke prevention medication.

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Author contributions

Both authors jointly conceived and designed this review. KH wrote the first draft, and subsequent drafts were co-written. Both authors approved the final version.

References

1 National Collaborating Centre for Chronic Conditions. Atrial fibrillation: national clinical guideline for management in primary and secondary care. 2006. http://www.nice.org.uk/nicemedia/pdf/cg036 fullguideline.pdf (accessed March 5, 2013).
2 Wolf PA, Abbott RD, Kannel WB. Atrial fibrillation: an independent risk factor for stroke: the Framingham Study. Stroke 1991; 22: 983–8.
3 Mant J, Wade DT, Winner S. Health care needs assessment. In: Health Care Needs Assessment: The Epidemiologically Based Needs Assessment Reviews, 2nd edn. Oxford: Radcliffe Medical Press, 2004: 141–244.
4 Andersson P, Londahl M, Abdon NJ, Terent A. The prevalence of atrial fibrillation in a geographically well-defined population in Northern Sweden: implications for anticoagulation prophylaxis. J Intern Med 2012; 272: 170–6.
5 Kannel WB, Wolf PA, Benjamin EJ, Levy D. Prevalence, incidence, prognosis, and predisposing conditions for atrial fibrillation: population-based estimates. Am J Cardiol 1998: 82: 2N–9N.
6 Hart RG, Pearce LA, Aguilar MI. Meta-analysis: antithrombotic therapy to prevent stroke in patients who have nonvalvular atrial fibrillation. Ann Intern Med 2007; 146: 857–67.
7 Atrial Fibrillation Investigators. Risk factors for stroke and efficacy of antithrombotic therapy in atrial fibrillation. Analysis of pooled data from five randomized controlled trials. Arch Intern Med 1994; 154: 1449–57.
8 van Walraven C, Hart RG, Singer DE et al. Oral anticoagulants vs aspirin in nonvalvular atrial fibrillation: an individual patient meta-analysis. JAMA 2002; 288: 2441–8.
9 Go AS, Hylek EM, Borowosky LH, Phillips KA, Selby JV, Singer DE. Warfarin use among ambulatory patients with nonvalvular atrial fibrillation: the Anticoagulation and Risk Factors In Atrial Fibrillation (ATRIA) Study. Ann Intern Med 1999; 131: 927–34.
10 Birman-Deych E, Radford MJ, Näläsenä DS, Gage BF. Use and effectiveness of warfarin in Medicare beneficiaries with atrial fibrillation. Stroke 2006; 37: 1070–4.
11 Cowan C, Fay M, Griffith K, Jenkinson D. Anticoagulation in AF. Anticoagulation uptake remains poor in high risk patients. BMJ 2011; 342: d1153.
12 Mant J, Hobbs FDR, Fletcher K et al. Warfarin versus aspirin for stroke prevention in an elderly community population with atrial fibrillation (the Birmingham Atrial Fibrillation Treatment of the Aged Study, BAFTA): a randomised controlled trial. Lancet 2007; 370: 493–503.
13 Gross CP, Vogel EW, Dhond AJ et al. Factors influencing physicians’ reported use of anticoagulation therapy in nonvalvular atrial fibrillation: a cross-sectional survey. Clin Ther 2003; 25: 1750–64.
14 National Institute for Health and Clinical Excellence. Final appraisal determination: dabigatran etexilate for the prevention of stroke and systemic embolism in atrial fibrillation, 2011. http://www.nice.org.uk/nicemedia/live/12225/56899/56899.pdf (accessed March 5, 2013).
15 Aguilar M, Hart R, Pearce L. Oral anticoagulants versus antiplatelet therapy for preventing stroke in patients with non-valvular atrial fibrillation and no history of stroke or transient ischemic attacks. Cochrane Database Syst Rev 2007; 3: CD006186.
16 ACTIVE Writing Group of the ACTIVE Investigators. Clopidogrel plus aspirin versus oral anticoagulation for atrial fibrillation in the Atrial fibrillation Clopidogrel Trial with Irbesartan for Prevention of Vascular Events (ACTIVE W): a randomised controlled trial. Lancet 2006; 367: 1903–12.
17 Connolly SJ, Ezekowitz MD, Yusuf S et al. Dabigatran versus warfarin in patients with atrial fibrillation. N Engl J Med 2009; 361: 1139–51.
18 Patel MR, Mahaffey KW, Garg J et al. Rivaroxaban versus warfarin in nonvalvular atrial fibrillation. N Engl J Med 2011; 365: 883–91.
19 Granger CB, Alexander JH, McMurray JJ et al. Apixaban versus warfarin in patients with atrial fibrillation. N Engl J Med 2011; 365: 981–92.
20 Weitz JI, Connolly SJ, Patel I et al. Randomised, parallel-group, multicentre, multinational phase 2 study comparing edoxaban, an oral factor Xa inhibitor, with warfarin for stroke prevention in patients with atrial fibrillation. Thromb Haemost 2010; 104: 633–41.
21 Mega JL. A new era for anticoagulation in atrial fibrillation. N Engl J Med 2011; 365: 1052–4.
22 Camm AJ, Lip GY, De Caterina R et al. 2012 focused update of the ESC Guidelines for the management of atrial fibrillation: an update of the 2010 ESC Guidelines for the management of atrial fibrillation. Developed with the special contribution of the European Heart Rhythm Association. Eur Heart J 2012; 33: 2719–47.
28 Shah SV, Gage BF. Cost-effectiveness of dabigatran for stroke prophylaxis in atrial fibrillation. *Circulation* 2011; 123: 2562–70.

29 Sorensen SV, Kansal AR, Connolly S et al. Cost-effectiveness of dabigatran etexilate for the prevention of stroke and systemic embolism in atrial fibrillation: a Canadian payer perspective. *Thromb Haemost* 2011; 105: 908–19.

30 National Institute for Health and Clinical Excellence. Rivaroxaban for the prevention of stroke and systemic embolism in atrial fibrillation: Technology appraisal TA256. 2012. http://www.nice.org.uk/ta256 (accessed March 5, 2013).

31 National Institute for Health and Clinical Excellence. Dabigatran etexilate for the prevention of stroke and systemic embolism in atrial fibrillation: Technology appraisal TA249. 2012. http://www.nice.org.uk/ta249 (accessed March 5, 2013).

32 National Institute for Health and Clinical Excellence. Atrial fibrillation: the management of atrial fibrillation. NICE Clinical Guideline 36. London: National Institute for Health and Clinical Excellence, 2009. London: Department of Health, 2010. http://www.healthcareimprovementscotland.org/hcis/doc ashx?docid=05a9291c-f901-4a8e-85d0-de8699a31757&version=-1 (accessed March 5, 2013).

33 Department of Health. NHS Reference Costs 2008. London: Department of Health, 2010. http://webarchive.nationalarchives.gov.uk/20130107105354/http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DHF_1111591 (accessed March 5, 2013).

34 Department of Health. NHS Reference Costs 2009. London: Department of Health, 2010. http://webarchive.nationalarchives.gov.uk/20130107105354/http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DHF_1111591 (accessed March 5, 2013).

35 Spackman E, Burch J, Faria R, Corbacho B, Fox D, Woolacott N. Dabigatran etexilate for the prevention of stroke and systemic embolism in atrial fibrillation: Evidence Review Group Report. 2011. http://www.nice.org.uk/nicemedia/pdf/CG036nicegrading.pdf (accessed March 5, 2013).

36 Langkilde IK, Bergholdt Asmussen M, Overgaard M. Cost-effectiveness of dabigatran etexilate for stroke prevention in non-valvular atrial fibrillation. Applying RE-LY to clinical practice in Denmark. *J Med Econ* 2012; 15: 695–703.

37 Davidson T, Husberg M, Ianzon M, Oldgren J, Levin LA. Cost-effectiveness of dabigatran compared with warfarin for patients with atrial fibrillation in Sweden. *Eur Heart J* 2013; 34: 177–83.

38 Lee S, Angladre MW, Pham D, Pisacane R, Kluger J, Coleman CI. Cost-effectiveness of rivaroxaban compared to warfarin for stroke prevention in atrial fibrillation. *Am J Cardiol* 2012; 110: 845–51.

39 Thomson R, Parkin D, Eccles M, Sudlow M, Robinson A. Decision analysis and guidelines for anticoagulant therapy to prevent stroke in patients with atrial fibrillation. *Lancet* 2000; 355: 956–62.

40 Gage BF, Cardinali AB, Albers GW, Owens DK. Cost-effectiveness of warfarin and aspirin for prophylaxis of stroke in patients with nonvalvular atrial fibrillation. *JAMA* 1995; 274: 1839–45.

41 Jowett S, Bryan S, Mant J et al. Cost effectiveness of warfarin versus aspirin in patients older than 75 years with atrial fibrillation. *Stroke* 2011; 42: 1717–21.

42 Burton C, Isles C, Norrie J, Hanson R, Grubb E. The safety and adequacy of antithrombotic therapy for atrial fibrillation: a regional cohort study. *Br J Gen Pract* 2006; 56: 697–702.

43 Jones M, McEwan P, Morgan CL, Peters JR, Goodfellow J, Currie CJ. Evaluation of the pattern of treatment, level of anticoagulation control, and outcome of treatment with warfarin in patients with nonvalvar atrial fibrillation: a record linkage study in a large British population. *Heart* 2005; 91: 472–7.

44 Connolly SJ, Pogue J, Hart RG et al. Effect of clopidogrel added to aspirin in patients with atrial fibrillation. *N Engl J Med* 2009; 360: 2066–78.

45 European Medicines Agency. European public assessment reports for rivoglitazone, 2012. http://www.ema.europa.eu/ema/index.jsp?curl=/pages/medicines/landing/epar_search.jsp&mid=WC0b01ac058001d1248&source=home&medSearch&keyword=rivoglitazone&category=human&isNewQuery=true (accessed May 3, 2013).

46 Albers GW, Diener HC, Frison I et al. Ximelagatran vs warfarin for stroke prevention in patients with nonvalvular atrial fibrillation: a randomized trial. *JAMA* 2005; 293: 690–8.

47 European Medicines Agency. Press release: AstraZeneca withdraws its application for ximelagatran 36-mg film-coated tablets. 2006. http://www.ema.europa.eu/docs/en_GB/document_library/Press_release/2010/02/WC500074073.pdf (accessed March 5, 2013).

48 Ali A, Bailey C, Abdelhafiz AH. Stroke prophylaxis with warfarin or dabigatran for patients with nonvalvar atrial fibrillation-cost analysis. *Age Ageing* 2012; 41: 681–4.

49 Connolly SJ, Pogue J, Eikelboom J et al. Benefit of oral anticoagulant over antiplatelet therapy in atrial fibrillation depends on the quality of international normalized ratio control achieved by centers and countries as measured by time in therapeutic range. *Circulation* 2008; 118: 2029–37.

50 Gage BF, Waterman AD, Shannon W, Boechler M, Rich MW, Radford MJ. Validation of clinical classification schemes for predicting stroke: results from the National Registry of Atrial Fibrillation. *JAMA* 2001; 285: 2684–70.

51 Lip GYH, Nieuwlaat R, Pisters R, Lane D, Crijns H. Refining clinical risk stratification for predicting stroke and thromboembolism in atrial fibrillation using a novel risk factor based approach: the Euro Heart Survey on Atrial Fibrillation. *Chest* 2010; 137: 263–72.

52 Healey JS, Hart RG, Pogue J et al. Risks and benefits of oral anticoagulation compared with clopidogrel plus aspirin in patients with atrial fibrillation according to stroke risk: the atrial fibrillation clopidogrel trial with ibradin for prevention of vascular events (ACTIVE-W). *Stroke* 2008; 39: 1482–6.

53 Healthcare Improvement Scotland. Prevention of stroke and systemic embolism in adult patients with atrial fibrillation. 2012. http://www.healthcareimprovementscotland.org/hsis/doc/ashx?docid=05a9291c-f901-4a8e-85d0-de8699a31757&version=-1 (accessed March 5, 2013).

54 The Health and Social Care Information Centre. General practice trends in the UK: The NHS Information Centre for health and social care - Part of the Government Statistical Service. 2011.

55 Stewart S, Hart CL, Hole DJ, McMurray JJ. Population prevalence, incidence, and predictors of atrial fibrillation in the Renfrew/Paisley study. *Heart* 2001; 86: 516–21.

56 Ruigomez A, Johansson S, Wallander MA, Rodriguez LA. Incidence of chronic atrial fibrillation in general practice and its treatment pattern. *J Clin Epidemiol* 2002; 55: 356–63.

57 Carroll K, Majed A. Consoribidity associated with atrial fibrillation: a general practice-based study. *Br J Gen Pract* 2001; 51: 884–91.

58 National Institute for Health and Clinical Excellence. Atrial fibrillation: the management of atrial fibrillation: Costing Report. Implementing NICE guidance in England. 2006. http://www.nice.org.uk/nicemedia/pdf/CG036costingreport.pdf (accessed March 5, 2013).

59 National Institute for Health Care and Clinical Excellence. Stroke prevention in primary care - the role of atrial fibrillation. 2009. http://www.improvement.nhs.uk/heart/Portals/0/documents/2009_AF_Commissioning_Guide_v2.pdf (accessed March 5, 2013).

60 National Health Service. Guidance on risk assessment and stroke prevention for atrial fibrillation (GRASP-AF). 2009. http://www.improvement.nhs.uk/graspdf/GRASPResources.html (accessed March 5, 2013).

61 Holt TA, Hunter TD, Gunnarsson C, Khan N, Cloud P, Lip GYH. Risk of stroke and oral anticoagulant use in atrial fibrillation: a cross-sectional survey. *Br J Gen Pract* 2012; 62: c710–7.

62 Fang MC, Go AS, Chang Y et al. Warfarin discontinuation after starting warfarin for atrial fibrillation. *Circ Cardiovasc Qual Outcomes* 2010; 3: 624–31.

63 Connolly SJ, Eikelboom J, Joyner C et al. Apixaban in patients with atrial fibrillation. *N Engl J Med* 2011; 364: 806–17.

64 Olesen JB, Lip GYH, Hansen M et al. Validation of risk stratification schemes for predicting stroke and thromboembolism in patients with atrial fibrillation: nationwide cohort study. *BMJ* 2011; 342: d124.