

Aspirin for Primary and Secondary Prevention of Cardiovascular Disease: Time to Stop?

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Recent trials suggest that aspirin does not reduce cardiovascular events or increase longevity in people who are not known to have cardiovascular disease,¹ leading to draft recommendations in the United States against the use of aspirin for primary prevention in people aged >60 years.² This will cause much dismay to those with a dogmatic belief in aspirin's benefits but, perhaps, much rejoicing amongst patients who have one less pill thrust upon them.^{3–5} The failure of aspirin for primary prevention should now lead to a re-examination of the evidence for aspirin for secondary prevention.^{1,6–11}

Calderone et al¹² provide an updated meta-analysis of 21 randomized trials of aspirin for the primary prevention of cardiovascular events, including 173,810 patients and almost one million patient-years of follow-up. Of the 21 trials, 16 were placebo-controlled, including a recent trial of an aspirin-containing polypill.¹³ An effective intervention does not require a large trial to show benefit, but confirming that an intervention has no effect does. The authors suggest that aspirin might reduce the risk of myocardial infarction and transient ischemic attacks (TIAs), but not stroke, and provide no evidence that this translates into a reduction in chronic disability. If the reduction in nonfatal events was real and these events were important, then they should translate into a reduction in mortality.⁶ However, aspirin did not reduce all-cause, cardiovascular, or noncardiovascular mortality. Most myocardial and cerebral vascular events are missed.⁶ Less than a third of myocardial infarctions and one-fifth of cerebral vascular events are clinically obvious.^{14–16} Many patients with a myocardial or cerebral infarction will die before a clinical diagnosis is possible.¹⁷ The failure of aspirin to reduce mortality suggests that it might simply change how events present rather than prevent them.

Calderone et al also report that aspirin increased the risk of major bleeding events by approximately 50%; use of lower doses did not diminish this risk. The authors hypothesize that the effects on vascular events and risk of bleeding may vary

with age, with the balance being less favorable in older people, even though they are more likely to have unrecognized coronary artery disease. Indeed, they found an interaction between age and mortality, with an excess mortality of approximately 10% in those aged >70 years, but neither an interaction between age and vascular events nor, surprisingly, the risk of major bleeding.

For people aged <65 years, death (47 per 1,000 people over 5 years without aspirin) was at least twice as common as myocardial infarction (22 events), stroke (16 events), or major bleeds (18 events). If 1,000 people aged <65 years took aspirin for 5 years, the authors predicted that this would lead to two fewer deaths, one cardiovascular and one noncardiovascular, about three fewer myocardial infarctions, and one less stroke, but one more intracerebral bleed and nine more major bleeding events. Even if the estimated effect of aspirin is true, is it worth it? The great majority of events would not be prevented. Do we not have more effective interventions to consider, such as antihypertensive and lipid-lowering agents, smoking cessation, and a healthy lifestyle?

For people aged ≥65 years, death (62 per 1,000 people over 5 years without aspirin) was more than twice as common as myocardial infarction (19 events), stroke (23 events), or major bleeds (29 events). If 1,000 people aged ≥65 years took aspirin for 5 years, the authors predicted that this would have no effect on cardiovascular death but result in five additional noncardiovascular deaths, four more intracerebral bleeds, and seven more major bleeding events, although it might lead to three fewer myocardial infarctions and two fewer strokes. Thus, despite being at higher risk of both cardiovascular disease and cancer, the balance of risk and benefit of primary prevention for aspirin looks rather unfavorable for people aged ≥65 years.

These results place further doubt on the value of long-term aspirin prophylaxis for secondary prevention. Many older people will have undiagnosed atherosclerotic disease,

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for whom this analysis suggests that the harms outweigh the benefits. Clinically overt hemorrhage may only be the tip of an iceberg of aspirin-related problems. Aspirin might also increase the rate of end-stage renal disease and microvascular cerebral hemorrhage, while increased rates of proton-pump inhibitor use may be fuelling an epidemic of iron deficiency.^{6,18}

Why do so many doctors believe that long-term aspirin for secondary prevention is effective?⁸ Historically, the most important reasons may be publication bias and meta-analyses of trials that appeared positive but only due to the inclusion of small, unrealistically positive trials.^{10,19} More recent versions of the secondary prevention meta-analysis have excluded trials which previously appeared to show that aspirin could achieve resurrection, but the damage that such bias creates to perceptions and guidelines is hard to reverse.^{7,10,19}

A short course of aspirin after a vascular event does appear beneficial, as might an antibiotic for pneumonia. Just because a course of treatment is effective it does not mean it should be continued lifelong. A definitive trial, ISIS-2, showed that aspirin reduced recurrent infarction and mortality when given immediately after a myocardial infarction.²⁰ The course of treatment was only 4 weeks but the legacy of that course of treatment lasted at least 10 years, despite the fact that most patients must have stopped taking aspirin after 4 weeks; after all, only 5% of patients in ISIS-2²¹ were

discharged on an antiplatelet agent and no intervening trial could have changed clinical practice. Also, if there was enough equipoise to have a placebo group in ISIS-2, then there was no valid argument for starting aspirin at the end of the 4-week double-blind period. Similarly, aspirin given immediately after a stroke/TIA for 6 to 12 weeks reduces the risk of recurrent stroke, disability, and death, but without evidence of benefit thereafter.²² There is little evidence that continuing aspirin beyond 12 weeks after a myocardial infarction or a stroke is beneficial (►Fig. 1).

The only substantial randomized, placebo-controlled, long-term trial of aspirin after a myocardial infarction at a dose of <300 mg/day exists only in the imagination of doctors. The long-term trials that do exist used much larger doses and showed no effect on mortality, or even a trend to excess²³ (►Fig. 1). Probably the strongest evidence for aspirin (dose 75 mg/day) for secondary prevention comes from the SAPAT trial that, between 1985 and 1989, enrolled 2,035 patients diagnosed with angina by a primary care physician.²⁴ This was before the widespread introduction of statins. The study found a significant reduction in the composite of myocardial infarction or sudden death (124 events on placebo compared with 81 on aspirin; $p=0.003$) with a similar trend for all-cause mortality (106 and 82 respectively; $p=0.103$). This is a slim amount of evidence upon which to make strong recommendations. These results also contrast with those of the largest, long-term secondary

Trials comparing long-term aspirin to placebo (where $n > 1,000$) in chronic 'stable' vascular disease published between 1980-2021 (there are only 3.5 trials!)

Trial	Year	Country	Prior Event	Average Time from Event to Randomisation	Aspirin Dose	Randomised		FU (months)	CV Deaths		Non-CV Deaths		All-Cause Mortality	
						Placebo	Aspirin		Placebo	Aspirin	Placebo	Aspirin	Placebo	Aspirin
PARIS-I*	1980	USA/UK	Post-MI	>6 months	972mg	406	*405	41	45	*37	7	*5	52	*43
AMIS	1980	USA	Post-MI	>12 months	1,000mg	2257	2267	38	181	197	38	48	219	245
SAPAT	1992	Sweden	Angina	Prior MI excluded	75mg	1026	1009	50	70	51	36	31	106	82
Total						3689	3681	>40	296 (8.0%)	285 (7.7%)	81 (2.2%)	84 (2.3%)	377 (10.2%)	370 (10.0%)
Effect of Aspirin														
CARDIFF-I#	1974	UK	Post-MI		300mg	560	566	12					61	47
				<4 weeks		157	138						15 (9.6%)	9 (6.5%)
				4-5 weeks		139	145						24 (17.3%)	13 (9.0%)
				6-13 weeks		136	150						14 (10.3%)	15 (10.0%)
				>14 weeks		128	133						8 (6.3%)	10 (7.5%)

*Randomisation was 2:1; accordingly, numbers in the aspirin group have been halved. # CARDIFF-I trial suggested benefit only when aspirin was given within 6 weeks of myocardial infarction (MI). For references to trials see reference 6.

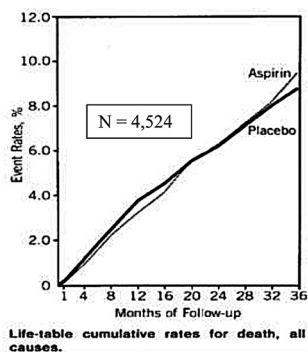


Fig. 1 All-cause mortality in the Aspirin Myocardial Infarction Study (AMIS), the largest trial of long-term aspirin administration both in terms of numbers of patients and events. The authors concluded "aspirin is not recommended for routine use in patients who have a survived an MI." Although the doses used are much higher than contemporary guidelines suggest, there is no substantial long-term, placebo-controlled trial of aspirin after myocardial infarction at a dose of <300mg/day. There is no placebo-controlled trial of late-initiation, long-term aspirin after a cerebro-vascular event.²³

prevention trial after myocardial infarction (AMIS; $n = 4,524$; aspirin dose 1,000 mg/day), which showed numerically more deaths on aspirin (245 deaths; 10.8%) than on placebo (219 deaths; 9.7%).²³ The largest long-term trial of aspirin after a stroke (UK TIA; $n = 2,449$; aspirin dose 300 mg or 1,200 mg/day) reported 119 strokes on placebo compared with 100 on aspirin 300 mg/day and, respectively, 90 compared with 81 coronary events and 122 compared with 109 deaths.²⁵ None of the differences in UK TIA were significant. Importantly, these trials were conducted before the era of effective treatments for hyperlipidemia, hypertension, and diabetes for which there is robust evidence of an effect on morbidity and mortality.

Aspirin, even at low doses, blocks the production of vasodilator prostaglandins that reduce endothelial platelet adhesion and have important effects on renal water and salt handling.^{6,26} P2Y₁₂ inhibitors have more selective effects on platelet function but clopidogrel has only marginally greater effects on cardiovascular events compared with aspirin and no greater effect on mortality.²⁷ The COMPASS trial did suggest that addition of rivaroxaban 2.5 mg twice daily to aspirin 100 mg/day might reduce both morbidity and mortality for patients with chronic atherosclerotic disease but created uncertainty over whether rivaroxaban alone might be as effective as the combination.²⁸ Preconceptions about the efficacy of aspirin interfered with the design, duration, and interpretation of the effects of rivaroxaban monotherapy in the COMPASS trial.

Theoretical considerations also influence practice. Vascular occlusion is considered by many to be primarily a thrombotic event. However, thrombosis is usually secondary to plaque rupture, which may often be caused by hemorrhage from neovascular proliferation from the vasa vasorum⁶; a pathology akin to diabetic retinopathy. For a patient with an ulcerated plaque presenting with a vascular event, the risk of thrombosis is high and the net effect of a short course of aspirin is beneficial. For patients with plaque that is not ulcerated, the risk of hemorrhage may balance or outweigh the risk of thrombosis.⁶ Percutaneous vascular interventions will cause plaque disruption for which a course of antiplatelet therapy is warranted but we have no evidence that life-long therapy is required. Indeed, recent trials investigating de-escalation of antithrombotic therapy suggest that withdrawal may be possible or even advisable, although this should be substantiated by further research.

In conclusion, aspirin taken long-term may “give with one hand but take away with the other,” leaving the individual only with indigestion, an increased risk of major bleeding, and other adverse consequences. However, there is good evidence that aspirin given for 4 to 12 weeks after a vascular event is beneficial; beyond that, surely we have better things to do for our patients and with our valuable time than prescribe aspirin.

Conflict of Interest

J.G.F.C. has received research funding and personal honoraia from Bayer Pharmaceuticals.

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