

Original research

Long-term survival benefit of ramipril in patients with acute myocardial infarction complicated by heart failure

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ABSTRACT

Aims ACE inhibition reduces mortality and morbidity in patients with heart failure after acute myocardial. infarction (AMI). However, there are limited randomised data about the long-term survival benefits of ACE. inhibition in this population.

Methods In 1993, the Acute Infarction Ramipril Efficacy (AIRE) study randomly allocated patients with AMI and clinical heart failure to ramipril or placebo. The duration of masked trial therapy in the UK cohort (603 patients, mean age=64.7 years, 455 male patients) was 12.4 and 13.4 months for ramipril (n=302) and placebo (n=301), respectively. We estimated life expectancy and extensions. of life (difference in median survival times) according to duration of follow-up (range 0-29.6 years).

Results: By 9 April 2019, death from all causes occurred in 266 (88.4%) patients in placebo arm and 275 (91.1%) patients in ramipril arm. The extension of life between ramipril and placebo groups was 14.5 months (95% Cl 13.2 to 15.8). Ramipril increased life expectancy more for patients with than without diabetes (life expectancy difference 32.1 vs 5.0 months), previous AMI (20.1 vs 4.9 months), previous heart failure (19.5 vs 4.9 months), hypertension (16.6 vs 8.3 months), angina (16.2 vs 5.0 months) and age >-65 years (11.3 vs 5.7 months). Given potential treatment switching, the true absolute treatment effect could be underestimated by

Conclusion For patients with clinically defined heart failure following AMI, ramipril results in a sustained survival benefit, and is associated with an extension of life of up to 14.5 months for, on average, 13 months treatment duration.

INTRODUCTION

Survival following acute myocardial infarction (AMI) remains a key objective for healthcare professionals and governments throughout the globe. Despite major advances in the emergency treatment and secondary prevention of AMI, a substantial proportion of patients with AMI develop heart failure, which in turn is associated with high morbidity, mortality and healthcare expenditure.2

Moreover, therapeutic inhibition of ACE has become an established strategy in the care of patients with AMI and heart failure.3 ACE inhibition reduces death and disease in patients with left ventricular systolic dysfunction, and coronary artery disease.6-8 In the Acute Infarction Ramipeil

Efficacy (AIRE) study, ACE inhibition at an average follow-up time of 15 months after randomisation significantly reduced all-cause mortality, and continued to confer a survival benefit at 3 years following study closure in the AIRE Extension (AIREX) study. 28 11 Further placebo-controlled survival studies testing the efficacy of ACE inhibition in patients with AMI and clinical heart failure are unlikely because withholding a treatment of accepted clinical value would be unethical. Therefore, in the AIRE survival (AIRE-S) study, we aimed to assess the long-term (29.6 years after randomisation) magnitude of impact on length of survival observed after treatment with ramipril (target dose 5 mg twice daily) when compared with placebo.

METHODS

Patients and design

The rationale, design, organisation, outcome definitions and results of the AIRE study have been published previously.18 12 This was an independent, multinational, multicentre, double-blind, randomised, placebo-controlled, parallel-group investigation into the potential benefits from treatment with the lipophilic ACE inhibitor ramipril (target dose 10 mg/day) following AMI complicated by clinical evidence of heart failure. When the trial was conducted, in 1993, all patients had an ECG and cardiac entryme confirmation of AMI, and also evidence of transient or persisting heart failure as judged by the attending physician (chest X-ray evidence of left ventricular failure, or auscultatory evidence of pulmonary oedema, or a third heart sound in the presence of pensistent tachycardia), usually sufficient to justify discretic or vasodilator therapy. Patients were excluded if they had evidence of clinical instability, contraindications to the use of ACE inhibitor therapy, heart failure of primary valvular or congenital origin or overt heart failure deemed to require treatment with an open-label ACE inhibitor. After applying these entry criteria to consecutive patients admirted to the UK centres (30) centres), 44% of patients approached declined to give their consent to take part in the study. All 603 of the remaining patients were then randomised by telephone to receive masked treatment with either ramipril (titrated over 4 days to 5 mg twice daily) or a matching placebo. Treatment was started between 2 and 9 days after the onset of AMI. Patients initially received either 1.25 or 2.5 mg ramipril or

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Peter L. Thompson

Long-Term Management of Patients After Myocardial Infarction C. Tissa Kappagoda, P. V. Greenwood, 1987-11-01 **Coronary Care Manual** Peter L. Thompson, 2010-12-03 The Coronary Care Manual has been designed as a practical manual for the management of the acute coronary patient and aims to achieve a balance between the large and rapidly changing evidence base and practical application in the CCU ICU ED and ambulance The book has been written by a group of Australian and international experts for medical nursing and paramedic staff working in these areas This edition covers a broad range of coronary care medicine in enough detail to eliminate the need to refer to a larger reference book providing a portable companion for a night on call Completely redesigned with new artwork the manual is organized to suit academics and practitioners alike covers a broad range of coronary care medicine provides specific advice on the management of common clinical problems eliminates the need to refer to a larger reference book features a consistent style and focus with standardised artwork for figures is now also available as an eBook A code inside the Coronary Care Manual enables a full text download allowing you to browse and search electronically make notes and bookmarks in the electronic files and highlight Cardiology E-Book Michael H. Crawford, John P. DiMarco, Walter J. Paulus, 2009-09-18 With your heavy case load material you can t afford to waste time searching for answers Cardiology 3rd Edition by Drs Crawford DiMarco and Paulus offers you just the practical problem based guidance you need to guickly overcome any clinical challenge 8 color coded sections cover the 8 major clinical syndromes of cardiovascular disease each section a virtual mini textbook on its topic 40 new chapters keep you up to date with the latest advances in the field while more than 2 000 lavish high quality illustrations color photographs tables and ECGs capture clinical manifestations as they present in practice It's current actionable information that you can put to work immediately for your patients Offers a problem based approach that integrates basic science diagnostic investigations and therapeutic management in one place for each cardiovascular disease so you can quickly find all of the actionable knowledge you need without flipping from one section to another Features introductory bulleted highlights in each chapter that present the most pertinent information at a glance Presents abundant algorithms to expedite clinical decision making Includes more than 2 000 lavish high quality illustrations color photographs tables and ECGs that capture clinical manifestations as they present in practice and promote readability and retention Includes 40 new chapters including Inherited Arrhythmia Syndromes Implantable Cardioverter Defibrillators and Cardiac Resynchronization Therapy in CHD Management of the Cyanotic Patient with CHD Special Problems for the Cardiology Consultant Dealing with Bariatric Gastric Bypass and many more that equip you with all of the latest knowledge Presents Special Problem sections many new to this edition that provide practical advice on problems that can be difficult to treat Cumulated Index Medicus, 1994

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