

BMJ Open Preventing adverse cardiac events (PACE) in chronic obstructive pulmonary disease (COPD): study protocol for a double-blind, placebo controlled, randomised controlled trial of bisoprolol in COPD

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ABSTRACT

Introduction Heart disease in chronic obstructive pulmonary disease (COPD) is a common but neglected comorbidity. Patients with COPD are frequently excluded from clinical trials of treatments aimed at reducing cardiac morbidity and mortality, which has led to undertreatment of cardiovascular disease in patients with COPD. A particular concern in COPD is the underuse of beta (β)-blockers. There is observational evidence that cardioselective β -blockers are safe and may even reduce mortality risk in COPD, although some evidence is conflicting. There is an urgent need to answer the research question: Are cardioselective β -blockers safe and of benefit in people with moderately severe COPD? The proposed study will investigate whether cardioselective β -blocker treatment in patients with COPD reduces mortality and cardiac and respiratory morbidity.

Methods and analyses This is a double-blind, randomised controlled trial to be conducted in approximately 26 sites in Australia, New Zealand, India, Sri Lanka and other countries as required. Participants with COPD will be randomised to either bisoprolol once daily (range 1.25–5 mg, dependent on tolerated dose) or matched placebo, in addition to receiving usual care for their COPD over the study duration of 24 months. The study will enrol 1164 participants with moderate to severe COPD, aged 40–85 years. Participants will be symptomatic from their COPD and have a postbronchodilator forced expiratory volume in 1 s (FEV₁) $\geq 30\%$ and $\leq 70\%$ predicted and a history of at least one exacerbation requiring systemic corticosteroids, antibiotics or both in the prior 24 months.

Ethics and dissemination The study protocol has been approved by the Sydney Local Health District Human Research Ethics Committee at The Concord Repatriation General Hospital.

Trial registration numbers NCT03917914; CTRI/2020/08/027322.

Strengths and limitations of this study

- This will be the first adequately powered randomised controlled trial of a cardioselective β -blocker, bisoprolol, in addition to usual care in participants with chronic obstructive pulmonary disease (COPD) with all-cause mortality, cardiopulmonary hospitalisations and major adverse cardiac events as a primary, composite endpoint.
- The trial is multicentre and multicountry and participants will have a spectrum of severity of COPD and be included as long as they have had a COPD exacerbation (ECOPD) in the prior 2 years.
- The study has an independent endpoint adjudication committee to determine the underlying cardiac or respiratory cause of death.
- Weaknesses of the study include the impact of the COVID-19 pandemic on both exacerbation rates and COPD hospitalisations, the exclusion of very severe patients with COPD and those who are oxygen dependent.
- The inclusion of hospitalised ECOPD with the other components of the primary outcome may mean that we cannot draw firm conclusions from the results as to whether β -blockers reduce ECOPD.

INTRODUCTION

Chronic obstructive pulmonary disease (COPD) is the third-leading global cause of death behind ischaemic heart disease and stroke and the fifth-leading cause of disability affected life years.¹ In randomised controlled trials (RCTs) of participants with COPD, 30%–50% of deaths are due to cardiac causes, and COPD participants have

high, often undiagnosed levels of cardiovascular disease (CVD).^{2,3} COPD and airflow limitation are risk factors for development of atherosclerosis and CVD, independently of cigarette smoking and air pollution. Even in a population with mild COPD, the Lung Health Study showed that for every 10% decrease in FEV₁% predicted, there was a 28% increase in fatal coronary events and 20% increase in non-fatal coronary events.⁴ In addition, a recent study demonstrated that COPD exacerbations (ECOPD) more than double the risk of myocardial infarction in the subsequent 5 days.⁵ Subclinical ischaemic events and cardiac arrest occur more frequently after an exacerbation⁶ and cardiac events are strong predictors of mortality in this setting.⁷ In patients hospitalised with an ECOPD in Waikato, New Zealand,⁸ elevated cardiac biomarkers were associated with a 15-times greater likelihood of dying within 30 days of admission than those without raised biomarkers and an 88% total mortality at 5 years. Despite this, CVD is often undiagnosed and undertreated in patients with advanced COPD.

β-blockers are indicated for a range of CVDs, but, are frequently withheld from patients with COPD,⁹ due to uncertainty about the potential risk of bronchospasm. To date, most large RCTs of β-blockers for heart failure and other cardiac diseases have excluded patients with COPD, so their efficacy in COPD is also an area of uncertainty. However, there have been advances in knowledge regarding the distinct airway pathophysiology of COPD, and most observational studies and meta-analyses indicate that patients with COPD do not suffer adverse effects on lung function or symptoms when prescribed cardioselective β-blockers,^{10–12} even when there is a reversible component to their airflow limitation. Database analyses and retrospective case series suggest that cardioselective β-blockers are safe¹⁰ and may be beneficial^{13–18} in patients with COPD. However, the recent β-Blockers for the Prevention of Acute Exacerbations of COPD (BLOCK COPD) study suggested that the β-blocker metoprolol may increase the risk of severe and very severe ECOPDs.¹⁹ Importantly, however, the population in the BLOCK COPD study had a high proportion (40%) of oxygen-dependent participants with severe COPD, who may be more sensitive to adverse effects of β-blockade.²⁰ In addition, metoprolol is known to be less cardioselective than some other β-blockers such as bisoprolol.^{21–23} Faced with this uncertainty about the benefits of β-blockers in COPD, the fact remains that many people with COPD have cardiovascular indications for the use of β-blockers, and many patients are currently being treated with them.^{16,24} Therefore, it is important that the safety and efficacy of β-blocker use in people with COPD are further examined.

Rationale

Since a high proportion of people with COPD die from cardiac events, it is rational to propose that treatments should be trialled that target cardiac comorbidity, whether clinical or subclinical.²⁵

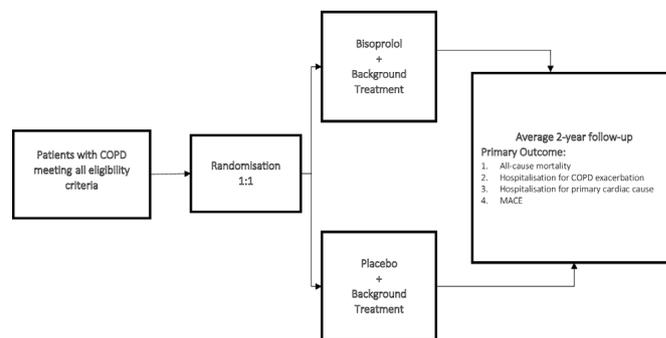


Figure 1 PACE study design. COPD, chronic obstructive pulmonary disease; PACE, preventing adverse cardiac events.

Our group has recently completed a feasibility study assessing the safety and efficacy of commencing the highly cardioselective β₁-blocker, bisoprolol, in people with COPD, and found that it was well tolerated and that a large RCT of cardioselective β-blockers would be feasible in this participant population.²⁶

The hypothesis of the preventing adverse cardiac events in COPD (PACE in COPD) clinical trial is that bisoprolol, taken in addition to usual care in participants with COPD over 24 months, will reduce combined all-cause mortality, cardiac and respiratory hospital admissions and major adverse cardiac events (MACE) compared with matched placebo.

METHODS AND ANALYSIS

Trial design and registration

The PACE in COPD study is a double-blind, multicentre RCT conducted in Australia, New Zealand, India and Sri Lanka with potential for expansion to other countries such as South Africa. Regulatory and ethical approval has not yet been obtained for these additional countries at this point in time. Participants with COPD will be randomised to one of two treatment arms: bisoprolol plus usual COPD care (intervention) or matched placebo plus usual COPD care (control). The design of the study and flow of participants are presented in [figure 1](#)

This study will be conducted, analysed and reported according to the Consolidation Standards of Reporting Trials statement for RCTs.²⁷ The George Institute for Global Health will act as study sponsor. The trial has been prospectively registered on the ClinicalTrials.gov and the Clinical Trial Registry—India, and other country registries as appropriate.

Participants

Inclusion and exclusion criteria

Participants will be eligible for this study if they qualify based on *all* of the criteria detailed in [box 1](#).

Setting

Participants will be recruited from respiratory and cardiac outpatient clinics, pulmonary rehabilitation services, from

Box 1 Eligibility criteria

Inclusion criteria

- ▶ Have provided written informed consent.
- ▶ Have chronic obstructive pulmonary disease (COPD) defined by the 2019 Global Initiative for Chronic Obstructive Lung Disease diagnostic criteria.
- ▶ Aged ≥ 40 and ≤ 85 years.
- ▶ Forced expiratory volume in 1 s (FEV₁) $\geq 30\%$ and $\leq 70\%$ predicted postbronchodilator.
- ▶ FEV₁/forced vital capacity (FVC) < 0.7 postbronchodilator.
- ▶ Have had a COPD exacerbation (ECOPD) in the previous 24 months requiring oral corticosteroids, antibiotics or both.
- ▶ If taking maintenance oral corticosteroids, dosage is stable and ≤ 10 mg/day for 4 weeks prior to randomisation.
- ▶ Resting systolic blood pressure (SBP) ≥ 100 mm Hg.
- ▶ Participants' baseline FEV₁ must not fall $\geq 12\%$ and ≥ 200 mL at 2 hours after a test dose of 1.25 mg bisoprolol, and their SBP must be ≥ 100 mm Hg at 1 and 2 hours post-test dose.
- ▶ For New Zealand participants only, eligible participants must also have a history of CVD, including heart failure, ischaemic heart disease, tachyarrhythmia or hypertension.

Exclusion criteria

- ▶ Concurrent therapy with any other β -blocker.
- ▶ Resting heart rate < 60 bpm.
- ▶ Unstable left heart failure (ie, symptomatic and/or necessary change in management in the last 12 weeks or in investigator's opinion).
- ▶ Clinically significant pulmonary hypertension, which in the investigator's opinion would be a contraindication for β -blocker therapy.
- ▶ Severe end-stage peripheral vascular disease.
- ▶ Second or third-degree heart block.
- ▶ Currently using or have been prescribed long-term oxygen therapy or resting saturated oxygen level $< 90\%$ when stable.
- ▶ Expected survival is less than 12 months, or in the investigator's opinion, the person has such unstable disease (of any type) that maintaining 12 months' participation would be unlikely.
- ▶ Clinical instability since a MACE in the previous 12 weeks.
- ▶ Lower respiratory tract infection or ECOPD within the last 4 weeks.
- ▶ COPD not clinically stable as determined by the investigator.
- ▶ In the investigator's opinion, have asthma-COPD overlap or coexistent asthma; or an improvement in FEV₁ ≥ 400 mL postbronchodilator is observed on two occasions.
- ▶ Women of childbearing age and capability who are pregnant or breast feeding or those in this group not using adequate birth control.
- ▶ Coexistent illness that precludes participation in the study (eg, poorly controlled diabetes, active malignancy).
- ▶ Severe end-stage liver disease defined by International Normalised Ratio (INR) > 1.3 and albumin < 30 g/L or portal hypertension/ascites.
- ▶ High chance in the opinion of the investigator that the potential participant will not adhere to study requirements.

the inpatient setting and from specialist and primary care referrals.

Procedures

Potential participants who may be suitable based on the eligibility criteria outlined above will be provided with a current, approved participant information sheet and consent form, by the site personnel and given ample time

to review and consider participation before providing written informed consent. Participants will visit the study centres at 0, 1, 3, 6, 12, 18 and 24 months. The full schedule of study interventions is outlined in [table 1](#). Prebronchodilator (baseline only) and postbronchodilator (all visits) spirometry, blood pressure and heart rate (all visits), an ECG (baseline, 1, 3, 12 and 24 months) and 6-minute walk test (baseline and 12 months) will be conducted. Also, the COPD Assessment Test (CAT),²⁸ St. George Respiratory Questionnaire (SGRQ)²⁹ and EuroQoL Group 5-Dimension self-report questionnaire (EQ-5D-5L)³⁰ will be completed at baseline, 6, 12, 18 and 24 months. Participant-reported adverse events will be recorded at all visits. Telephone calls to participants to record adverse events and adherence will be made at 2 weeks, 8 weeks and approximately every 8 weeks in between clinic visits. Adherence to treatment is assessed via tablet counts and self-reported adherence.

Uptitration protocol

PACE in COPD will use bisoprolol 2.5 mg tablets and matching placebo that will be taken as a half tablet, whole tablet or two tablets according to the uptitration protocol. From baseline to visit 2 (4 weeks), participants will be taking a daily dose of half a tablet of study medication (1.25 mg bisoprolol or placebo) per day. At the 4-week safety visit, participants will be uptitrated to one whole tablet (2.5 mg bisoprolol or placebo) per day. At visit 3 (12 weeks), the dose of study medication will be increased to two whole tablets (5 mg bisoprolol or placebo) per day, provided that participants have not experienced any of the adverse events listed in [box 2](#). If any adverse events occur, the lower dose can be maintained, or the dosage may be reduced. All dose changes are subjected to the investigator's discretion.

Patient and public involvement

The research question, protocol document and design of the study were informed by feedback obtained from the study's consumer representative, who is a member of the PACE Steering Committee (PSC). Our consumer representative has lived experience from being a carer for a person suffering from COPD and is a COPD patient advocate. Our consumer representative provides ongoing advice on behalf of patients with COPD regarding recruitment and ongoing management of the study. Once the study is complete and analysis has been finalised, a plain language statement will be drafted to explain the outcomes of the study for all randomised participants. This statement will be reviewed by the consumer representative and the central ethics committee prior to dissemination. The burden of the intervention will be assessed by participants via an anonymous qualitative questionnaire when each participant completes the study.

Randomisation, stratification and interventions

Randomisation will be performed according to a computer-generated algorithm carried out in the IBM



Table 1 Schedule of study assessments

Visit number	V1	P1	V2	P2	V3	P3	V4	P4	P5	V5	P6	P7	V6	P8	P9	V7	D1																				
Assessment description	Screening and baseline			Follow-up																																	
Time since randomisation	0 W			2 W		4 W		8 W		12 W		16 W		6 M		8 M		10 M		12 M		14 M		16 M		18 M		20 M		22 M		24 M		24 M+1-3 W			
Visit type	Clinic	Phone	Clinic	Clinic	Phone	Clinic	Clinic	Phone	Phone	Phone	Clinic	Phone	Clinic	Phone	Phone	Phone	Clinic	Phone	Phone	Phone	Phone	Phone	Phone	Clinic	Clinic												
Visit window	±0 W			±1 W		±2 W		±1 W		±2 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W		±1 W					
Informed consent/eligibility	X																																				
Reasons for non-participation	X																																				
Demographics, medical, clinical and smoking history	X																																				
Physical examination and anthropometrics	X																																				
Randomisation	X																																				
Vital signs and vital status	†																																				
BP, HR																																					
Spirometry																																					
ECG																																					
CAT, SGRQ and EQ-5D-5L [‡]																																					
Adverse events																																					
Concomitant medications																																					
6MWT ^{**}																																					
Medication dosage review																																					
Participant diary review																																					
Treatment adherence ^{††}																																					

* Weight only for anthropometrics at V7.
 †BP at screening will be measured at 0 minutes, 30 minutes, 1 hour and 2 hours after first test dose of bisoprolol.
 ‡Spirometry will be measured at 0 minutes, 1 hour and 2 hours after first test dose of bisoprolol.
 §Screening ECG will be 12-lead, all other measurements will be 3-lead.
 ¶CAT; SGRQ; EQ-5D-5L questionnaires.
 ** 6MWT (6 Minute Walk Test) is for a subset of the study population.
 †† Clinic visits will involve pill counts. At phone visits, participants will be asked how many doses they missed.
 CAT, COPD Assessment Test; ECG, Electrocardiogram; EQ-5D-5L, EuroQoL Group 5-Dimension self-report questionnaire; 6MWT, 6-minute walk test; SGRQ, St. George Respiratory Questionnaire.

Box 2 Adverse events that preclude uptitration of bisoprolol

- ▶ Bronchoconstriction defined as $\geq 12\%$ and ≥ 200 mL forced expiratory volume in 1 s fall from baseline.
- ▶ < 100 mm Hg or HR < 60 bpm.
- ▶ Symptomatic postural hypotension.
- ▶ Increased wheezing, chest tightness.
- ▶ Marked exercise-related fatigue.
- ▶ Symptoms of peripheral arterial insufficiency.

Clinical Development database. The randomisation list will be concealed and generated centrally using a random combination of four by six permuted blocks within each stratum. Randomisation will be stratified by site, smoking status and previous diagnosis of CVD requiring treatment.

Active and matched placebo will be labelled using blinded kit numbers. Un-blinding will be unlikely, however the Site Principal Investigator has the ability to unblind, after confirmation with the Chief Investigator, within the database. Test dose tablets of bisoprolol, to be given to the participant during screening to test tolerability, will be packaged in a separate container of a half dose (1.25 mg) for each site.

Primary and secondary outcomes

The primary end point is the time to the first event of a composite outcome, which includes any of: all-cause mortality, hospitalisation for ECOPD or a primary cardiac cause (ischaemia, arrhythmia or heart failure), or a MACE. A MACE event includes cardiovascular death, non-fatal myocardial infarction, or non-fatal stroke. ECOPDs will be graded as moderate (requiring oral corticosteroids, antibiotics or both without hospital admission) or severe (requiring above treatment and hospital admission).

Exploratory outcomes include the annualised ECOPD rate (mild/moderate/severe), time to first moderate/severe ECOPD, quality of life assessed by the SGRQ, EQ-5D-5L and CAT, clinical spirometry, hospitalisation rate for respiratory and cardiac causes, total number of cardiac events and the number of composite (annualised) cardio-respiratory hospital admissions and MACE.

A within trial cost-effectiveness analysis of the intervention will also be conducted.

Data collection

All study data will be entered into the electronic case report form (eCRF) hosted by IBM Clinical Development, including clinic and phone visit attendances, study measurements, safety monitoring and all adverse events, as well as the independent adjudication of the primary end points. All data, including source data at sites will be stored for 15 years. Confidentiality will be ensured by only de-identified information being collected and stored centrally. Only authorised sponsor staff will have access to the trial dataset.

Consumer/community involvement

A consumer representative was involved in the design of this research protocol and has ongoing input into the operational conduct of the study as a member of the Steering Committee. Once the trial is finalised and published, participants will be informed of the results through a plain language statement along with information on the unblinded allocation they received during the trial.

Prohibited concomitant medications

Participants are not permitted to take other β -blockers, non-dihydropyridine calcium channel blockers (with the exception of diltiazem, which is at the investigator's discretion), class Ia or Ic anti-arrhythmic agents or topical acetylcholine during the study. If it is deemed clinically necessary to resume a prohibited medication by the treating physician, participants must cease their randomised intervention prior to recommencing the prohibited medication. They will continue with follow-up visits in the study.

Sample size estimation and recruitment target

The study will recruit 1164 participants to achieve a power of 90% to detect a risk reduction of 25% in the intervention vs the control arm (HR=0.75). Assuming a primary end point event rate of 30 per 100 person-years³¹⁻³³ in the control group and that every participant will be followed up for 2 years with up to 10% lost to follow-up, the study will continue until a total of 514 participants experience a primary endpoint event. The study will be conducted in multiple countries to ensure adequate recruitment is achieved.

Adverse events and safety monitoring

All adverse events will be recorded at study visits using the eCRF. Adverse events of special interest will include all potential adverse effects of the investigational treatment and COPD-related events, such as light-headedness or falls, increased wheezing, fatigability, abnormal heart rate, worsening respiratory symptoms, respiratory tract infections and including presentation to general practitioners, specialists, or emergency departments for respiratory symptoms. Testing positive for the Severe Acute Respiratory Syndrome Coronavirus-2 will be recorded as an adverse event. If the participant develops coronavirus disease 2019 (COVID-19) and is hospitalised, or if any other serious adverse event (SAE) criteria are met, then this will be classified as an SAE. The standard definition for all other SAEs will be used and data collected and submitted to regulatory agencies as required for each country.

An independent Data Safety and Monitoring Board (DSMB) will monitor the study for safety and efficacy. Five independent, senior clinicians and academics covering respiratory, cardiac and statistical expertise constitute our DSMB membership. The DSMB will perform an ongoing review of predefined safety parameters and overall study

conduct, with 6-monthly meetings and quarterly SAE line listings. The board will review blinded allocated data (ie, aggregated by treatment group) on participant characteristics and adverse events at regular intervals during the study. The board will be charged with monitoring total mortality, SAEs and making recommendations based on other outcomes such as cause-specific death or non-fatal SAEs. They will principally monitor safety and adverse events without including any formal testing of a treatment effect. Safety analyses presented at DSMB meetings will include summary reports of frequency and differences between treatment groups for deaths, cardiac and respiratory hospitalisations, ECOPDs, MACE and all SAEs.

An independent respiratory or cardiac adjudicator will assess all hospitalisations and deaths reported in the eCRF. If there is disagreement in the causality and/or relatedness of the event between the independent adjudicator and the site investigator, the event will require adjudication by the PACE Adjudication Committee (PAC). The PAC consists of five experienced clinicians who will adjudicate/confirm primary causality of all hospitalisations and deaths. The membership of this committee will be independent of the Chief Investigator team.

Data monitoring will be conducted remotely on a monthly basis and onsite annually at all clinical trial sites by the study sponsor.

Statistical analysis

The statistical analysis will be by intention-to-treat. The primary analysis will be an analysis of time to any event included in the composite endpoint that occurs first: all-cause mortality, cardiac and respiratory hospital admissions and MACE. The main analysis will consist of a Kaplan-Meier plot and a log-rank test comparing survival times between the β -blocker and the placebo groups. Sensitivity analyses adjusting for important prognostic factors such as age, sex, smoking status, acute ischaemic event, HEART (history, ECG, age, risk factors and troponin) score (where available), hospitalised ECOPDs or heart failure in the previous 12 months will be conducted using Cox models. In the secondary analyses, all-cause mortality, cardiac or respiratory hospitalisation and MACE will be separately analysed. The rate of ECOPDs will be annualised, analysed by Poisson regression with an overdispersion correction applied if necessary. Continuous variables, such as the CAT scores, SGRQ and FEV₁, will be compared by t-tests and mixed models to account for repeated measurements.

Economic evaluation

We will undertake an Australia-specific trial-based cost-effectiveness analysis. Costs will include treatment costs ascertained from participant and study records. Healthcare utilisation will be followed up in both arms of the study based on self-reported general practitioner or hospital attendances and changes in concomitant medications. Health state utilities will be estimated via the EQ-5D-5L and will be used to weight survival up to 24

months to determine quality-adjusted life years. Mean differences in costs and quality-adjusted life years between both treatment groups will be estimated to determine the potential incremental quality-adjusted life year gain from the intervention. Usual deterministic one-way and probabilistic sensitivity analyses will be conducted.

DISCUSSION

After more than 50 years of widespread use and proven benefit in many contexts, there is limited clinical trial evidence to clarify the effects of cardioselective β -blockers in people with COPD, with or without clinically evident CVD. To date, there are very few studies targeting cardiovascular risks in participants with COPD, and very few RCTs examining the role of cardioselective β -blockers to improve the management and outcomes of heart disease in participants with COPD. The 2019 BLOCK COPD¹⁹ trial investigated the effect of metoprolol compared with placebo on ECOPD, however, the study was stopped early because of futility and concerns of an adverse safety signal. All participants in the BLOCK COPD trial had moderate to severe airflow limitation with 40% requiring long-term oxygen therapy. Although no statistical association was found for this marker of severity with worse outcomes on metoprolol, these patients were at high risk of severe exacerbation and death. The study reported that metoprolol was associated with an increased risk of severe and very severe exacerbations. However, the design of this trial does not provide any guidance for the use of β -blockers in COPD populations with less severe disease to prevent cardiac events or ECOPDs. Hence, the target population for the PACE in COPD study are those with an FEV₁ $\geq 30\%$ and $\leq 70\%$ predicted postbronchodilator and excludes all participants dependent on long-term oxygen therapy, thus enrolling a less severe COPD population, but at similar or greater risk of cardiac events.^{34,35} Refer to [table 2](#) for a comparison of the eligibility criteria for the BLOCK COPD and PACE in COPD trials. The addition of lung function and blood pressure criteria after a test dose of bisoprolol to assess eligibility for the PACE trial ensures that participants sensitive to bisoprolol are excluded.

The PACE in COPD trial uses the cardioselective β -blocker, bisoprolol, with the goal to reduce morbidity and mortality from cardiovascular events and ECOPDs. Differences in cardioselectivity appear to advantage some drugs over others^{9,36,37} in relation to airway outcomes and potentially also to CVD outcomes. Sudden cardiac death is also more common in COPD and may be reduced by β -blocker therapy.¹⁸ People with COPD demonstrate evidence of sympathetic overdrive, a consequence of both their disease and acute and chronic β_2 -agonist therapy. In heart failure, cardiac β_1 -receptors are reduced in number and β_2 -receptors are increased,³⁸ thus β_2 -agonists may further increase adrenergic drive, tachycardia, inotropic contraction and myocardial oxygen consumption, at least in the short term. During an ECOPD, when patients are more hypoxaemic, very large doses of β_2 -agonists

Table 2 Comparison of the BLOCK COPD and PACE in COPD trial eligibility criteria

BLOCK COPD trial (¹⁸)	PACE in COPD trial
Inclusion	
40–85 years old	40–85 years old
FEV ₁ <80% predicted postbronchodilator	FEV ₁ ≥30% and ≤70% predicted postbronchodilator
FEV ₁ /FVC ratio <0.7 postbronchodilator	FEV ₁ /FVC ratio <0.7 postbronchodilator
Antibiotic/systemic glucocorticoids OR Emergency Department (ED)/hospitalisation OR prescription/or use of supplemental oxygen in previous 12 months	Have had a COPD exacerbation in the previous 24 months requiring oral corticosteroids, antibiotics or both
HR 65–120 bpm	HR ≥60 bpm
SBP >100 mm Hg	SBP ≥100 mm Hg
	Baseline FEV ₁ must not fall ≥12% and ≥200 mL at 2 hours after a test dose of 1.25 mg bisoprolol, and participant must have SBP ≥100 mm Hg at 1-hour and 2-hour post-test dose.
Exclusion	
Proven indication for use of a β-blocker, including history of myocardial infarction or revascularisation within 36 months or heart failure with a left ventricular ejection fraction <40%	Unstable left heart failure, clinically significant pulmonary hypertension, severe end-stage peripheral vascular disease, second or third-degree heart block
	Currently using or have been prescribed long-term oxygen therapy, or resting saturated oxygen level <90% when stable

BLOCK COPD, Beta-Blockers for the Prevention of Acute Exacerbations of COPD; COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity; PACE, preventing adverse cardiac events; SBP, systolic blood pressure.

are often used, and this is associated with persistent biochemical evidence of cardiac dysfunction.³⁹ We chose bisoprolol over other cardioselective β-blockers given its favourable β₁ selectivity and positive findings in database reviews, observational studies and small randomised trials in people with COPD.^{20 40 41}

Structural vascular changes have been demonstrated in people with COPD and plaque burden is significantly greater than in matched smokers without COPD or non-smokers.⁴² Functional vascular changes such as high arterial stiffness have also been found in people with stable COPD and have been shown to increase for up to 8 weeks after exacerbations. Both the presence of plaques and arterial stiffness are established risk factors for cardiovascular events. People with COPD who experience frequent exacerbations have a higher risk of post-ECOPD myocardial infarction and increased arterial stiffness, compared with people with COPD with infrequent exacerbations. This suggests that there may be a cumulative effect of vascular and cardiac damage with frequent exacerbations. Other mechanisms for the association include systemic inflammation, oxidative stress, physiological stresses, adrenergic sensitivity/overdrive during ECOPD and accelerated ageing.^{42–45}

There is an urgent need for additional therapies to reduce morbidity and mortality and improve clinical outcomes in people with COPD. Therapies directed at COPD do not improve CVD mortality: despite a marked reduction in ECOPDs, neither long-acting muscarinic antagonists nor combination therapy with inhaled corticosteroids and long-acting β₂ agonists reduce all-cause or

CVD-related mortality.⁴⁶ Moreover, there is an increased risk of fatal and non-fatal cardiovascular events immediately following ECOPDs and so a different strategy is needed, in conjunction with the conventional approach of maximising bronchodilator pharmacotherapy for airway disease. Interventions to reduce cardiac complications during ECOPD along with reducing cardiac risk during stable periods are urgently needed. β-blocker medications, which have such a major role in improving outcomes in heart failure, need further investigation in the COPD population, a population where their potential benefit is still unrealised, and the extent of cardiac disease remains unrecognised.

There is an urgent need to answer the research question: Are cardioselective β-blockers safe and of benefit in people with moderately severe COPD? The aim of the PACE in COPD clinical trial is to assess the efficacy and safety of 24 months' therapy with a cardioselective β-blocker, taken in addition to usual care, in participants with COPD to reduce all-cause mortality, cardiac and respiratory events.

ETHICS AND DISSEMINATION

The study protocol has been approved by the Sydney Local Health District Human Research Ethics Committee at The Concord Repatriation General Hospital (Protocol Version 7.0, 25 February 2021). In New Zealand, the study protocol has been approved by the Health and Disability Ethics Committee at the Ministry of Health, Wellington. In India, each clinical sites' Institutional Ethics committee

has individually approved the study protocol. All future amendments will be submitted to the ethics committees and circulated to all clinical sites for governance approval. Clinical trial insurance is provided by the study sponsor as per Medicines' Australia guidelines for compensation.

The trial is registered at ClinicalTrials.gov.

Study findings will be disseminated through presentations at national and international conferences and publications in peer-reviewed journals. Authorship will be determined by the PSC and as per the PACE publication policy.

ROLES AND RESPONSIBILITIES

Trial sponsor: The George Institute for Global Health, Australia. The Sponsor is responsible for drug procurement, contract development and operational oversight, database development and data collection, ethical conduct and safety reporting.

PSC: Christine Jenkins, Robert Hancox, Catherina Chang, Richard Beasley, Jeremy Wrobel, Vanessa McDonald, Claudia Dobler, Ian Yang, Claude Farah, Belinda Cochrane, Graham Hillis, Caroline Polak Scowcroft, Ashutosh Aggarwal, Allison Martin. The PSC is responsible for the study design, analysis, interpretation of results and writing reports and publications.

DSMB: Professor Eric Bateman (Chair), Professor Anthony Keech (Deputy Chair), A/Professor Katrina Sharples, Professor Gary Anderson, Professor Peter Calverley. The DSMB is an independent, multidisciplinary group including experts in respiratory medicine, cardiology, pharmacology clinical trial methodology and statistics. All members have experience in the conduct and monitoring of RCTs. THE DSMB will meet 6 monthly to ensure ongoing safety surveillance. Two formal interim analyses are planned to assess efficacy and safety, when approximately one-third and two-thirds of the primary endpoint outcomes have been observed. The first of these is expected to occur at approximately 1.5 years after start of the study. The Haybittle-Peto stopping rule based on 3 SDs (interim $\alpha=0.27\%$) will be used.

PAC: Professor Lorcan McGarvey (Chair), Professor Claus Vogelmeier, Dr Sundeep Salvi, Professor Michael Williams, Dr Geoffrey Clare. All deaths and cardiac/respiratory hospitalisations will be reviewed by an independent adjudicator. If the independent reviewer agrees with the causality indicated by the principal investigator, no further adjudication is required. If causality is discrepant between the principal investigator and the independent adjudicator, the adverse event is referred to the PAC.

Coordinating centre (Australia/India): Professor Christine Jenkins, Dr Allison Martin, Ms Grace Balicki, Dr Shane Galgey, Mallikarjuna Kunigari. The coordinating centre is responsible for the overall operational conduct of the study in all countries other than New Zealand.

Coordinating Centre (New Zealand): Robert Hancox, Catherina Chang, Christine Tuffery.

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