

Childhood 'bronchitis' and respiratory outcomes in middle-age: a prospective cohort study from age 7 to 53 years

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ABSTRACT

Background Chronic bronchitis in childhood is associated with a diagnosis of asthma and/or bronchiectasis a few years later, however, consequences into middle-age are unknown.

Objective To investigate the relationship between childhood bronchitis and respiratory-related health outcomes in middle-age.

Design Cohort study from age 7 to 53 years.

Setting General population of European descent from Tasmania, Australia.

Participants 3202 participants of the age 53-year follow-up (mean age 53, range 51–55) of the Tasmanian Longitudinal Health Study cohort who were born in 1961 and first investigated at age 7 were included in our analysis.

Statistical methods Multivariable linear and logistic regression. The association between parent reported childhood bronchitis up to age 7 and age 53-year lung conditions (n=3202) and lung function (n=2379) were investigated.

Results Among 3202 participants, 47.5% had one or more episodes of childhood bronchitis, classified according to severity based on the number of episodes and duration as: 'non-recurrent bronchitis' (28.1%); 'recurrent non-protracted bronchitis' (18.1%) and 'recurrent-protracted bronchitis' (1.3%). Age 53 prevalence of doctor-diagnosed asthma and pneumonia (p-trend <0.001) and chronic bronchitis (p-trend=0.07) increased in accordance with childhood bronchitis severities. At age 53, 'recurrent-protracted bronchitis' (the most severe subgroup in childhood) was associated with doctor-diagnosed current asthma (OR 4.54, 95% CI 2.31 to 8.91) doctor-diagnosed pneumonia (OR=2.18 (95% CI 1.00 to 4.74)) and, paradoxically, increased transfer factor for carbon monoxide (z-score +0.51 SD (0.15–0.88)), when compared with no childhood bronchitis.

Conclusion In this cohort born in 1961, one or more episodes of childhood bronchitis was a frequent occurrence. 'Recurrent-protracted bronchitis', while uncommon, was especially linked to multiple respiratory outcomes almost five decades later, including asthma, pneumonia and raised lung gas transfer. These findings provide insights into the natural history of childhood 'bronchitis' into middle-age.

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ Severe early childhood bronchitis manifested by recurrent episodes of protracted wet cough is associated with asthma and bronchiectasis in later childhood.

WHAT THIS STUDY ADDS

⇒ This is the first very long-term prospective study that has examined the relationship between childhood bronchitis severity with adult lung health outcomes. 'Recurrent-protracted bronchitis' in childhood, while uncommon, was associated with multiple respiratory outcomes in middle-age including asthma, pneumonia and raised lung gas transfer on lung function measurement.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE AND/OR POLICY

⇒ This study's findings support the need for increased vigilance in monitoring young children who have multiple episodes of protracted bronchitis for potential asthma, pneumonia and bronchiectasis up to mid-adult life, confirmed with objective testing.

INTRODUCTION

'Bronchitis' implies airway inflammation. However, historically, the term bronchitis in childhood has been applied broadly to bouts of coughing without implying a specific cause or diagnosis, for example, asthma or bacterial bronchitis.^{1 2} Clinicians have previously believed that 'wheezy bronchitis,' on the other hand, signifies asthma.³ However, the terminology and understanding of the causes of paediatric cough has since evolved significantly and the paradigm has changed. It is now recognised and accepted that 'chronic cough' in children, that is, cough lasting >4 weeks, can be caused by a range of pathologies including protracted bacterial bronchitis (PBB), bronchiectasis, asthma and tracheo/

bronchomalacia. Furthermore, wheeze and cough can be common to all of these conditions.^{3,4}

PBB, also known as chronic bacterial bronchitis, is recognised as a discrete diagnostic entity by Respiratory Societies across Europe,⁵ the USA⁶ and Australia/New Zealand. PBB is defined as presence of a chronic ‘wet’ cough that lasts 1 month or more and resolves with 2–4 weeks of appropriate antibiotic therapy, in the absence of indicators to suggest another cause for the cough.^{7,8} Prospective paediatric cohort studies have described a link between PBB and both bronchiectasis and asthma, with 27% of children with PBB having an asthma phenotype 5 years later⁹ and 8%–10% having bronchiectasis within 2–5 years.^{9,10} Whether chronic bronchitis (CB) in childhood is associated with poor lung health in adulthood is unknown and represents an important research priority, with potential to improve outcomes through earlier recognition and management.⁵

Longitudinal cohort studies from early life to adulthood have provided important insights into lung disease development¹¹ and are closing the gap in understanding between paediatric and adult respiratory medicine.¹² While several studies have supported the role of early life exposures in adult lung function deficits and chronic obstructive pulmonary disease (COPD),^{13–18} the early origins of adult chronic productive cough (or CB) remain poorly understood and sparsely studied.¹⁹

Using population-based data from our Tasmanian Longitudinal Health Study (TAHS) longitudinal cohort, we aimed to investigate trends and associations between bronchitis severity at age 7 in childhood and respiratory health outcomes of asthma, CB, pneumonia and lung function in middle-age at age 53.

METHODS

Study design and population

Our study sample included participants of the TAHS cohort (1968–2016) who took part in the original assessments at age 7 and were followed up at the mean age of 53 years (details published elsewhere^{18,20–22}). Tasmania is an island State of Australia located 240 km to the south of the Australian mainland, which is the least populated and marginally the poorest state by gross product per capita, with relatively high rates of smoking and historically, less accessible healthcare facilities for a developed country. Briefly, this population-based cohort of participants essentially from European descent were born in 1961 and first studied including retrospective questionnaire data in 1968 (aged 6–7 years, n=8583), was retraced (n=7312) and resurveyed by a postal survey questionnaire in 2004 at mean age 43 years (n=5729) with further clinical testing. Subsequently, 6127 (71.4%) of the full original cohort were retraced and invited to participate in a clinical follow-up between 2012 and 2016. At mean age=53, 3609 (58.9%) respondents completed the questionnaire and 2719 (43.9%) underwent lung function testing including prebronchodilator/postbronchodilator

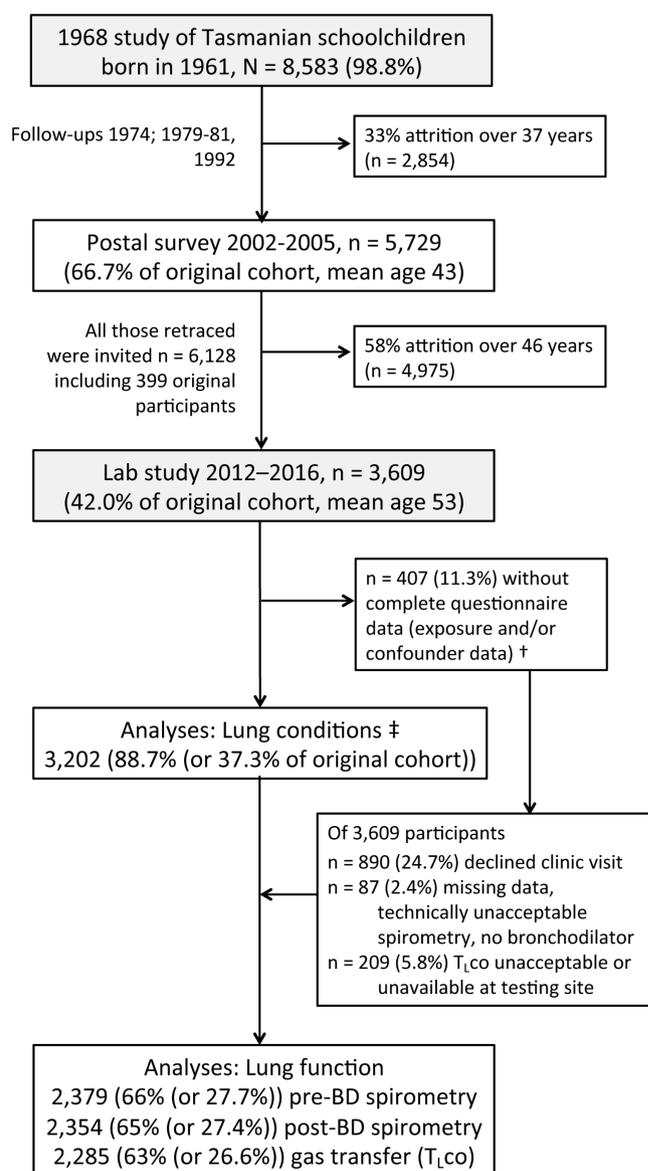


Figure 1 TAHS participants 1968–2016 study participants were those who attended both the 7 and 53 years follow-up visits and had complete questionnaire data and technically acceptable lung function measurements. Regression numbers varied with each outcome (n=3062–3196) ††. post-BD, post-bronchodilator; TAHS, Tasmanian Longitudinal Health Study; T_Lco, transfer factor for carbon monoxide of the lung.

spirometry and lung gas transfer measurements, on average 46 years later (figure 1). To minimise lost to follow-up, methods of tracing included computer linkage to the National Death Index, Commonwealth Electoral Roll, Medicare database, Tasmanian marriage records,²³ and via siblings and parents who were also part of the original 1968 survey, and multiple attempts at contacting participants may have been required. Over 70% of the original participants still resided in Tasmania, with migration mainly to the East Coast of Australia and fewer than 0.5% now living overseas.

Data collection methods

Details of lung function testing have been described²² and summarised in online supplemental methods E1a. Lung function testing was standardised across testing sites in accordance with the 2005 American Thoracic Society/ European Respiratory Society (ATS/ERS) standards.^{24 25} For this analysis, lung function data at age 7 and 53 years were converted to z-scores (ie, SD or SD units) using established standard reference values.^{26 27} Airflow obstruction was defined by forced expiratory volume in 1 s (FEV₁)/forced vital capacity (FVC) <lower limit of normal (LLN), and spirometric restriction by post-BD FVC <LLN and FEV₁/FVC ≥LLN.

Definition

Exposure

Childhood bronchitis (or ‘wet’ cough as current clinical definitions) was defined by an affirmative response by parents/guardian to the 1968 survey question by age 7: ‘Has he/she at any time in his/her life suffered from attacks of bronchitis or attacks of cough with sputum (phlegm) in the chest (‘loose’ or ‘rattly’ cough)?’. Based on the number of episodes and duration of attacks, those with childhood bronchitis was divided into three severity groups based on predetermined categories at the time of the 1968 survey: the non-recurrent subgroup had between one and five episodes only of any duration; the recurrent group had ≥6 episodes lasting <1 month on average; and the recurrent-protracted subgroup had either ‘continuous’ symptoms or ≥6 episodes each lasting an average of a month or more.

Outcomes

Adult lung conditions, current CB, which was a subset of those reporting ever-CB, was defined by the presence of a ‘usual’ cough with sputum (phlegm) for at least 3 months for the last two consecutive years at age 53. Asthma—current at age 53 was defined by a past doctor diagnosis of asthma associated with asthma-like symptoms, asthma medication use and/or related healthcare utilisation within 12 months prior to testing.

Online supplemental methods E1b provides further details for the exposure, other outcomes, confounders and effect modifier/s.

Statistical methods

Analyses were carried out using Stata (V.16, Stata). Our exposure, CB, included four categories as defined above, that is, no CB, non-recurrent bronchitis, recurrent non-protracted bronchitis, and recurrent-protracted bronchitis. Univariable trends of continuous/categorical respiratory outcomes were examined across these ordered categories (non-parametric trend tests (nptrend)²⁸/univariable logistic regression, respectively). This latter trend estimate approach assumes the relationship between the exposure expressed as an ordered variable

attached to numerical values is linearly related to the outcome.

Using complete cases, multivariable linear and logistic regression were used to examine associations between CB and with continuous and categorical respiratory outcomes respectively. For the lung function outcomes, the regression coefficients represented the associations between exposure categories and lung function values expressed as z-scores (SD units) which accounted for age, sex and height (online supplemental methods E2a).^{26 27}

We chose ‘no childhood bronchitis’ as the reference exposure category so differences between categories of childhood bronchitis severity were only relative to the reference group and not compared directly. To interpret differences between childhood bronchitis categories, the absolute numbers of participants estimated to have the lung condition at age 53 were calculated using the ‘margins’ command following the syntax for the separate multivariable regression analyses.

Informed by conceptualising the models using a directed acyclic graph,^{29 30} regression models were adjusted for paternal occupation, parental bronchitis, rurality of primary school, number of siblings and parental smoking when participants were aged 7, plus male/female for the non-lung function analyses (online supplemental methods E1b and E2b).

Childhood asthma, pneumonia-pleurisy and pre-BD FEV₁ at 7 years (ie, non-discriminatory for obstructive and restrictive lung function deficits) partly mediated relationships of corresponding adult outcomes as per causal mediation analyses (online supplemental methods E2c, table E1), thus were not included as model covariates. Mediation by FEV₁/FVC was possible but was not additionally performed. As childhood asthma/wheezing, ever-smoking and participant sex could plausibly strengthen the exposure/outcome relationships, corresponding interaction terms were used for the following reasons: to exclude the possibility that adult asthma was only related to participants who also wheezed in childhood; to assess whether the association with adult CB was related to adult cigarette smoking; and, because of known sex differences in lung function and asthma development. Sensitivity analyses that excluded participants who self-recalled asthma-ever and doctor-diagnosed pneumonia-ever before age 7 were performed (referred to as ‘recalled’ asthma and pneumonia, respectively).

A cut-off of $p < 0.10$ for interactions was used to assess for effect modification as this was a low powered statistical test. A z-score of > 0.5 SD²⁷ and absolute increase in FEV₁ of 200 mL and 12% post-BD were considered clinically significant.³¹

Patient and public involvement

Patients, TAHS participants or the public were not involved in the design, or conduct, or reporting, or dissemination plans of our research.

Table 1 Sociodemographic and clinical characteristics at age 7 of the middle-aged TAHS participants who formed the study sample for the current analysis

Childhood clinical characteristic in 1968	TAHS participants attending follow-up at age 53 (N=3202 (%))*				P trend
	Chronic bronchitis groups at age 7 years				
	None (n=1680 (52.5%))	Non-recurrent (n=902 (28.1%))	Recurrent, non-protracted (n=578 (18.1%))	Recurrent-protracted (n=42 (1.3%))	
Age (years, SD)	6.50 (0.3)	6.51 (0.3)	6.50 (0.3)	6.54 (0.3)	0.659
Sex (% male)	796 (47)	432 (48)	327 (56)	23 (55)	0.004
Familial bronchitic symptoms	–	–	–	–	–
One parent only	428 (25)	317 (35)	258 (45)	23 (55)	<0.001
Both parents	30 (1.8)	40 (4.4)	47 (8.1)	5 (12)	<0.001
≥3 Siblings, of any age	796 (47)	366 (41)	211 (36)	15 (36)	<0.001
Mother who regularly smoked	539 (32)	313 (35)	217 (37)	13 (31)	0.027
>5 cigarettes/day	436 (26)	257 (29)	184 (32)	7 (17)	0.040
Father who regularly smoked	1003 (60)	501 (56)	351 (61)	27 (64)	0.987
>5 cigarettes/day	882 (52)	445 (49)	313 (54)	24 (57)	0.694
Paternal occupation	–	–	–	–	–
Highest SES	401 (24)	237 (27)	150 (26)	7 (17)	0.118
Lowest SES	212 (13)	89 (9.8)	73 (13)	6 (15)	0.196
Remote rurality of school	60 (3.6)	20 (2.2)	16 (2.8)	0 (0)	0.086
Asthma/wheezing-ever	92 (5.5)	166 (18)	269 (46)	25 (60)	<0.001
>5 episodes	26 (1.6)	27 (3.0)	208 (36)	17 (44)	<0.001
Childhood infections	–	–	–	–	–
Pneumonia and/or pleurisy	108 (6.5)	153 (17)	151 (26)	16 (38)	<0.001
Measles (non-immunised)	1154 (69)	623 (69)	379 (65)	32 (76)	0.423
Pertussis (non-immunised)	30 (1.7)	14 (1.6)	7 (1.2)	0 (0)	0.220
Pertussis (immunised)	90 (5.3)	79 (8.8)	42 (7.3)	4 (9.5)	0.050

*Percentages in brackets compares the prevalence between childhood bronchitis groups. SES, socioeconomic status; TAHS, Tasmanian Longitudinal Health Study.

RESULTS

Childhood sociodemographic and clinical characteristics

Among 3202 participants who had complete questionnaire data when aged 53, 47.5% had one or more episodes of childhood bronchitis (table 1). The prevalence of non-recurrent and recurrent but non-protracted bronchitis was 28.1% (95% CI 26.6% to 29.7%) and 18.1% (95% CI 16.7% to 19.4%), respectively. The prevalence of 1.31% (95% CI 0.94% to 1.77%) for recurrent-protracted bronchitis was similar to that of the 2379 participants with complete lung function data (1.18% (95% CI 0.78% to 1.70%), online supplemental table E2) and the original 1968 cohort (1.21% (95% CI 0.98% to 1.47%), where n=101 of n=8583 = 1.18%).

Minor differences in baseline characteristics at age 7 for those who did not participate in the subsequent

follow-up and/or who had missing data at age 53 were seen, namely males of lower socioeconomic status were less likely to have data available (online supplemental results E1, table E2a). The differences between exposure groups and subsequent participation were also only modest, especially for the recurrent-protracted subgroup for which the percentages participating and not participating varied between 1.2% and 1.3%. There was little difference in the prevalence of doctor-diagnosed asthma as a proxy outcome at mean age 43 years, and the reduced participation by those with self-reported CB was not significant (online supplemental table E2b).

At age 7, participants with recurrent childhood bronchitis, with or without protracted episodes, were more likely to be male (table 1). Children with a history of bronchitis trended to have fewer siblings and fewer

mothers of the recurrent-protracted group smoked regularly. Positive trends were seen across the childhood bronchitis severity subgroups with regard to an increasing prevalence of parental bronchitis and/or symptoms, coexistent asthma/wheezing (ever and recurrent) and doctor-diagnosed pneumonia and/or pleurisy (p-trend <0.001, [table 1](#)).

Middle-age sociodemographic and clinical characteristics

Similar positive trends across the childhood bronchitis severity subgroups were observed between the actual and estimated prevalence of asthma-ever, current asthma and pneumonia-ever by middle-age (p-trend <0.001, [tables 2 and 3](#), respectively). While the proportions of usual cough, usual phlegm at age 53 were higher in the recurrent-protracted group, trends did not attain statistical significance. Nevertheless, the prevalence for current CB was 2.4-fold higher in this severe compared with the no childhood bronchitis subgroup (12.0% compared with 5.0%, respectively, p-trend=0.07).

The prevalence of current smoking in individuals at age 53 was similar across childhood subgroups (~17%), although a disproportionately high percentage of participants in the recurrent-protracted group had ever-smoked (83% compared with ≤60%, [table 2](#)). Clear positive trends across childhood bronchitis severity-subgroups were also seen for transfer factor of the lung for carbon monoxide (T_{1co}) and its coefficient (K_{co}) with a modest inverse trend for alveolar volume. There were similar trends for lower pre- and post-bronchodilator FEV_1 and FEV_1/FVC at age 53 (p-trend between 0.005 and 0.042, [table 2](#)). These trends were somewhat stronger when participants underwent spirometry when aged 7 (pre-BD FEV_1 p-trend=0.004; FEV_1/FVC p-trend <0.001).

Main associations

Using multivariable regression, compared with no childhood bronchitis, all three childhood bronchitis (non-recurrent, recurrent non-protracted and recurrent-protracted) subgroups were associated with both doctor-diagnosed current and ever-asthma in middle-age. These associations increased in strength in a dose response manner as severity increased ([table 4](#)). Compared with no bronchitis, increasing childhood bronchitis severity was associated with recalled doctor-diagnosed pneumonia after 7 years (p=0.032, p<0.001, p=0.05, respectively).

For spirometry outcomes at age 53, compared with no childhood bronchitis, associations were observed for the recurrent (non-protracted) childhood bronchitis subgroup with regards to a significant association with pre-BD airflow obstruction and lower post-BD FEV_1/FVC taken as a continuous variable and paradoxically, a borderline association with spirometric restriction ([table 5](#)). Recurrent-protracted childhood bronchitis was associated with significantly higher levels of T_{1co} and K_{co} but not alveolar volume in middle-age.

Effect modification

Among those without childhood asthma/wheezing, the odds for current asthma at age 53 were higher for participants with recurrent-protracted childhood bronchitis (OR 4.29 (95% CI 1.45 to 12.6), p=0.008), p-interaction=0.063, online supplemental table E3) although case numbers in the recurrent-protracted subgroup were relatively few (5-10 cases).

To contrast the above-mentioned finding, interactions were consistently seen between childhood bronchitis and childhood asthma/wheezing with regards to lung function measures used as continuous variables in middle-age, namely pre-BD FEV_1 , change in FEV_1 post-BD, pre-BD FEV_1/FVC , and post-BD FEV_1/FVC (p-interaction <0.1, online supplemental results E2, table E4). When the analysis was stratified by presence of childhood asthma/wheezing, compared with no childhood bronchitis, only participants with 1–5 episodes of any duration (ie, non-recurrent group) had non-obstructive lung function changes: reduced post-BD FVC (z-score -0.42 SD (95% CI -0.72 to -0.12), p=0.006), increased pre-BD FEV_1/FVC (+0.43 (95% CI +0.11 to +0.74), p=0.008) and post-BD FEV_1/FVC (+0.33 (95% CI +0.01 to +0.66), p=0.043) despite an increased pre-BD FEV_1 . There was also an associated threefold decrease in the odds for pre-BD airflow obstruction (OR 0.34 (95% CI +0.14 to +0.87), p=0.024), but there were too few cases to assess spirometric restriction as the outcome.

Despite few case numbers for the recurrent-protracted childhood bronchitis subgroup, associations with pneumonia-ever by age 53 were still seen for females (online supplemental results E3, table E5) and with current CB for middle-aged non-smokers (online supplemental results E4, table E6).

DISCUSSION

Using data from our whole-of-community cohort of European descent born in 1961, parent-reported childhood ‘bronchitis’ (and/or coughing phlegm, or ‘loose or rattly’ cough) was associated with consequential lung outcomes in middle-age, with significant trends for doctor-diagnosed asthma and pneumonia across the spectrum of cough-severity. Associations with doctor-diagnosed asthma and pneumonia were especially evident for the most severe subgroup termed ‘recurrent-protracted childhood bronchitis’. The findings of a low prevalence of CB and substantially higher lung gas transfer in mid-adult life for this subgroup have not previously been described. Similarly, the main association with a mixed obstructive/restrictive lung function pattern of modest magnitude for the subgroup with non-recurrent bronchitis episodes was also unexpected and probably not clinically important. Overall, we provide evidence to support childhood bronchitis as a contributor to clinical lung disease in mid-adult life, especially when chronic, recurrent and protracted.

**Table 2** Sociodemographic and clinical characteristics at age 53 of TAHS participants

Clinical characteristic at age 53 years	TAHS participants attending follow-up at age 53*†				P trend
	Childhood bronchitis by age 7 years			P trend	
	None	Non-recurrent	Recurrent, non-protracted		
Clinical feature (n (%)), N=3202	1680 (52.5)	902 (28.1)	578 (18.1)	42 (1.3)	–
Age (years, SD)	53.0 (1.0)	53.0 (0.9)	53.0 (0.9)	53.2 (1.0)	0.100
Sex (% male)	796 (47)	432 (48)	327 (56)	23 (55)	0.004
Ever-smokers	1008 (60)	520 (58)	325 (56)	35 (83)	0.045
Pack-years (median (IQR))	12.9 (3.9–29)	14.7 (3.4–29)	12.8 (2.5–30)	11.1 (7.5–37)	0.394
Current smokers	313 (19)	145 (16)	91 (16)	7 (17)	0.075
Usual cough ≥3 months	158 (9.4)	92 (10.3)	62 (10.7)	8 (19)	0.117
Usual phlegm ≥3 months	94 (5.6)	47 (5.3)	38 (6.6)	5 (12)	0.231
Chronic bronchitis-current	81 (5.0)	40 (4.6)	32 (5.9)	5 (12)	0.070
Chronic bronchitis-ever	137 (8.5)	72 (8.3)	59 (11)	7 (17)	0.245
Doctor-diagnosed current asthma	151 (9.0)	116 (13)	114 (20)	14 (33)	<0.001
Doctor-diagnosed asthma-ever	317 (19)	221 (25)	238 (41)	28 (67)	<0.001
Aged ≥7 years‡	313 (19)	221 (25)	218 (38)	28 (67)	<0.001
Doctor-diagnosed pneumonia-ever	238 (14)	174 (19)	146 (25)	15 (36)	<0.001
Aged ≥7 years‡	204 (12)	135 (15)	111 (19)	9 (21)	<0.001
Hospitalisation-ever	73 (4.4)	59 (6.7)	63 (11)	2 (4.8)	<0.001
Spirometry (n (%)) N=2379§	1238 (52.0)	686 (28.8)	427 (17.9)	28 (1.2)	–
Pre-BD zFEV ₁	–0.04 (2.0)	–0.08 (2.6)	–0.24 (1.2)	–0.19 (1.0)	0.016
Post-BD zFEV ₁	+0.16 (1.0)	+0.07 (1.0)	–0.02 (1.2)	+0.06 (0.9)	0.005
BDR Δ FEV ₁ (mL (SD))	109 (129)	104 (135)	111 (135)	110 (127)	0.766
Pre-BD zFVC	+0.10 (1.5)	+0.07 (2.1)	–0.03 (1.0)	+0.08 (1.0)	0.144
Post-BD zFVC	+0.10 (0.9)	+0.03 (0.9)	+0.01 (1.0)	+0.09 (0.9)	0.100
BDR Δ FVC (mL (SD))	19 (147)	13 (160)	33 (157)	13 (148)	0.310
Pre-BD zFEV ₁ /FVC	–0.28 (0.9)	–0.32 (0.9)	–0.39 (1.0)	–0.49 (0.7)	0.042
Pre-BD airflow obstruction¶	88 (7.1)	54 (7.9)	48 (11)	1 (4)	0.033
Post-BD zFEV ₁ /FVC	+0.07 (0.9)	+0.04 (0.9)	–0.07 (1.0)	–0.09 (0.7)	0.016
Post-BD airflow obstruction¶	48 (3.9)	30 (4.4)	24 (5.7)	1 (4)	0.077
Post-BD spirometric restriction¶	23 (1.9)	17 (2.5)	13 (3.1)	0 (0)	0.242
Transfer factor (n (%)) N=2281§	1194 (52.3)	647 (28.4)	412 (18.1)	28 (1.2)	–
CO transfer factor (zT _{lco})	+0.14 (0.9)	+0.17 (1.0)	+0.23 (1.0)	+0.67 (1.0)	0.008
CO transfer coefficient (zKco)	+0.23 (1.0)	+0.28 (1.0)	+0.41 (1.0)	+0.79 (1.1)	<0.001
Alveolar volume (zVA)	–0.09 (1.0)	–0.11 (1.0)	–0.19 (0.9)	–0.07 (1.0)	0.071

*Clinical and lung function variables are expressed by n(%) and z-score (SD), respectively, unless otherwise specified.

†Participant numbers differ by questionnaire, spirometry and transfer factor measurements as indicated in italics.

‡Doctor-diagnosed asthma | pneumonia recalled after childhood bronchitis was assessed that is, after age 7.

§By definition, z-scores that are measured in SD units have a mean of 0 and SD of 1.

¶Airflow obstruction was defined by FEV₁/FVC <LLN; spirometric restriction by post-BD FVC <LLN and FEV₁/FVC ≥LLN.

BD, bronchodilator; BDR, bronchodilator response; FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity; Kco, carbon monoxide transfer coefficient; LLN, lower limit of normal; TAHS, Tasmanian Longitudinal Health Study; T_{lco}, transfer factor for carbon monoxide of the lung; VA, alveolar volume; z, z-score.

Unexpectedly, a history of recurrent-protracted bronchitis in childhood was not strongly associated with CB in middle-age. Therefore, we postulate that only a relatively small subgroup at most may have had persistent productive cough episodes spanning childhood to mid-adult

life, with adult CB mainly influenced by two age-related factors. Analogous to asthma remission that is common in later childhood and adolescence, most children seem to ‘grow out of’ bronchitis during their late childhood or early adolescent years^{9 32} that could be in part explained

Table 3 Estimated prevalence of lung conditions by childhood bronchitis severity, adjusted for confounding variables

Clinical characteristic at age 53 years	TAHS participants attending follow-up at age 53 (% (95% CI))*				P trend
	Prevalence of childhood bronchitis by age 7 years				
	None (52.5%)	Non-recurrent (28.1%)	Recurrent, non-protracted (18.1%)	Recurrent-protracted (1.3%)†	
Chronic bronchitis-current	5.4 (4.2 to 6.5)	4.6 (3.2 to 5.9)	5.1 (3.3 to 6.8)	9.9 (1.5 to 18.2)	0.070
Chronic bronchitis-ever	8.9 (7.4 to 10.3)	8.2 (6.3 to 10.0)	9.5 (7.2 to 11.9)	14.8 (4.5 to 25)	0.245
Doctor-diagnosed current asthma	8.6 (7.2 to 10.0)	11.5 (9.4 to 13.6)	16.2 (13.1 to 19.2)	29.4 (15.7 to 43.1)	<0.001
Doctor-diagnosed asthma-ever	27.0 (24.9 to 29.0)	32.9 (30.0 to 35.8)	50.1 (46.4 to 53.9)	70.1 (57.1 to 83.0)	<0.001
Dr-diagnosed pneumonia-ever	14.4 (12.6 to 16.0)	19.2 (16.7 to 21.8)	25.1 (21.5 to 28.7)	35.0 (20.6 to 49.4)	<0.001

*Absolute numbers derived from multivariable regression analyses using the ‘margin’ command.
 †Prevalence estimates are based on n=42.
 TASH, Tasmanian Longitudinal Health Study.

by immune system maturation.^{33 34} Equally, CB most commonly develops during the productive working years of an adult resulting from environmental exposures e.g. cigarette smoking and/or air pollutants in the workplace,^{35 36} which is in keeping with CB-onset in adulthood.

Interestingly, among those most affected by childhood bronchitis, we found associations with current asthma in middle-age which were evident even for participants without a history of childhood asthma/wheezing at age 7. This suggests that a predisposition to multiple and protracted episodes of childhood bronchitis could lead to damaged airways, and that a degree of airway narrowing from remodelling could impair normal airway development to manifest as symptomatic asthma in later life.³⁷ As we were unable to tease out the precise overlaps between PBB, virus-associated wheeze and classic allergic asthma with regard to childhood wheezing, this prompted us to perform the analysis within strata of childhood asthma/wheezing. In this interaction analysis, we were still able to detect a significant association between the severe recurrent-protracted bronchitis subgroup and adult asthma for those who did not have parent-reported asthma and/or wheezing in childhood, suggesting the observed association could be independent of childhood asthma.

In contrast, we also found lung function deficits, for example, decrease in the odds for airflow obstruction for the subgroup with non-recurrent bronchitis episodes and coexistent childhood asthma/wheezing, but there was limited power to detect coexistent spirometric restriction. While we cannot draw firm conclusions, these findings are consistent with the Tucson Children’s Respiratory Study (TCRS). The TCRS accurately defined ‘other lower respiratory illnesses (LRIs) by age 3’ after excluding radiologically confirmed pneumonia and found a main association with active wheeze but not physician-diagnosed asthma

by age 30 (n=738), however, did not stratify analyses by childhood asthma/wheeze status.³⁸ These authors also found consistent reductions in FEV₁ and FVC of around 120 mL prebronchodilator and postbronchodilator in those with LRIs in early life, which could predispose individuals to COPD with a normal age-related decline if peak lung function was not attained from impaired lung function growth due to adverse exposures in utero and early childhood.^{13–18 39} The latter lung function finding is also consistent with our previous study that investigated the long-term outcomes of doctor-diagnosed childhood pneumonia and/or pleurisy and showed a restrictive lung function pattern on measures of static lung volumes in middle-age.²²

The observed ‘physiologically relevant’ elevations in lung transfer factor levels²⁷ in the absence of increased alveolar volume for the severe recurrent-protracted subgroup could plausibly relate to the high observed prevalence of coexistent asthma/wheezing in childhood⁴⁰ and adulthood.^{41 42} However, this finding also raises the possibility of a vascular mechanism like that seen in bronchiectasis in which broncho-pulmonary anastomoses derived from an expansion of a collateral and high-pressure bronchial artery network diverts pulmonary artery blood flow away from diseased parts to better supply the normal lung,⁴³ although the increase in bronchitic symptoms was only modest. Interestingly, lung transfer factor (T_Lco) can be high for adult patients at their first clinical presentation with bronchiectasis, which could be a feature prior to an accelerated decline which typically accompanies disease progression.⁴⁴

Our detailed questioning of parents of the 7-year-old children from a whole-of-community population allowed us to identify a severe subgroup that probably resembles PBB as closely as was possible without having data on antibiotic therapy and its response, given PBB was not



Table 4 Multivariable main associations between increasing childhood bronchitis severity by age 7 and various lung conditions at age 53 years

Lung condition at age 53	Cases (N)	Model N	Childhood bronchitis severity at age 7 (N=3202)†‡							
			Non-recurrent (n=902)			Recurrent, non-protracted (n=578)			Recurrent-protracted (n=42)	
			Case (n)	OR (95% CI)	Case (n)	OR (95% CI)	Case (n)	OR (95% CI)	Case (n)	OR (95% CI)
CB-current	158	3062	40	0.90 (0.67 to 1.22)	32	1.07 (0.76 to 1.50)	5	1.75 (0.75 to 4.08)		
CB-ever	275	3085	72	0.84 (0.56 to 1.25)	59	0.93 (0.60 to 1.45)	7	1.94 (0.72 to 5.21)		
Asthma-current	395	3178	116	1.48 (1.14 to 1.92)**	114	2.44 (1.85, 3.21)***	14	4.54 (2.31 to 8.91)***		
Asthma-ever	804	3196	221	1.33 (1.11 to 1.57)**	238	2.75 (2.28 to 3.31)***	28	6.49 (3.44 to 12.2)***		
≥7 years§	783	3196	221	1.41 (1.16 to 1.72)**	218	2.74 (2.21 to 3.41)***	28	8.19 (4.22 to 15.9)***		
Pneumonia-ever	574	3188	174	1.43 (1.15 to 1.78)**	146	2.02 (1.58 to 2.57)***	15	3.19 (1.66 to 6.14)**		
≥7 years§	460	3188	135	1.30 (1.02 to 1.65)*	111	1.79 (1.37 to 2.33)***	9	2.18 (1.00 to 4.74)¶		
<7 years	114	3188	39	2.21 (1.38 to 3.56)**	35	3.39 (2.05 to 5.60)***	6	9.80 (3.70 to 25.9)***		

*P<0.05, **p<0.01, ***p<0.001.
†Parent-reported bronchitis by age 7 categorised into none (reference=1, not shown), non-recurrent, recurrent (>5 episodes of coughing phlegm or 'loose, or rattly' cough lasting <1 month on average), and recurrent-protracted (>5 episodes, ≥1 month average duration).
‡Multivariable models were adjusted for rurality of school, paternal occupation, family history of bronchitis, sibling number, maternal and paternal smoking, and participant sex.
§Doctor-diagnosed asthma | pneumonia recalled after childhood bronchitis was assessed that is, after age 7.
¶P=0.050.
CB, chronic bronchitis.

Table 5 Multivariable main associations between increasing childhood bronchitis severity by age 7 and lung function at age 53 years

Adult outcome at age 53	Childhood bronchitis severity at age 7(β (95% CI) N=2379)†‡§				
	Case (n)	Model (N)	Non-recurrent (n=686)	Recurrent, non-protracted (n=427)	Recurrent-protracted (n=28)
Pre-BD zFEV ₁	–	2379	–0.02 (–0.22, +0.17)	–0.15 (–0.39, +0.08)	–0.10 (–0.88, +0.68)
Post-BD zFEV ₁	–	2354	–0.09 (–0.19, +0.01)	–0.16 (–0.28, –0.05)**	–0.09 (–0.47, +0.29)
BDR Δ FEV ₁ (mL)	–	2344	–4.8 (–17, +8)	–0.05 (–15, +15)	–0.10 (–50, +50)
Pre-BD zFVC	–	2378	–0.02 (–0.18, +0.13)	–0.10 (–0.28, +0.08)	+0.01 (–0.59, +0.62)
Post-BD zFVC	–	2354	–0.08 (–0.16, +0.01)	–0.09 (–0.19, +0.01)	–0.005 (–0.34, +0.33)
BDR Δ FVC (mL)	–	2343	–3.3 (–17, +11)	+8.7 (–8.1, +26)	–10 (–66, +46)
Pre-BD zFEV ₁ /FVC	–	2378	–0.03 (–0.12, +0.06)	–0.09 (–0.19, +0.02)	–0.19 (–0.54, +0.16)
Pre-BD airflow obstruction††	191	2378	1.12 (0.79–1.61)	1.64 (1.11–2.40)*	§
Post-BD zFEV ₁ /FVC	–	2354	–0.03 (–0.11, +0.06)	–0.12 (–0.22, –0.02)*	–0.15 (–0.49, +0.19)
Post-BD airflow obstruction**	103	2342	1.16 (0.72–1.87)	1.46 (0.87, 2.47)	§
Spirometric restriction**	53	2239	1.67 (0.89–3.11)	1.99 (0.97–4.01)††	§
zT _L co	–	2285	+0.03 (–0.06, +0.12)	+0.09 (–0.02, +0.20)	+0.51 (+0.15, +0.88)**
zKco	–	2285	+0.05 (–0.04, +0.15)	+0.18 (+0.06, +0.29)**	+0.51 (+0.13, +0.89)**
zV _A	–	2285	–0.02 (–0.11, +0.07)	–0.09 (–0.20, +0.02)	+0.06 (–0.30, +0.42)

*P<0.05, **p<0.01.
 †Parent-reported bronchitis by age 7 categorised into none (reference=1, not shown), non-recurrent, recurrent (>5 episodes of coughing phlegm or ‘loose, or rattly’ cough lasting <1 month on average), and recurrent-protracted (>5 episodes, ≥1 month average duration).
 ‡Multivariable models were adjusted for rurality of school, paternal occupation, family history of bronchitis, sibling number, maternal and paternal smoking, and participant sex.
 §Cells that had <5 cases; see table 2 for case numbers.
 ¶Lung function measured in z-scores which, by definition, has a mean of 0 and SD of 1.
 **ORs: pre-BD airflow obstruction (zFEV₁/FVC <LLN); spirometric restriction (post-BD zFVC <LLN and zFEV₁/FVC ≥LLN).
 ††P=0.050.
 BD, bronchodilator; BDR, bronchodilator response; CB, chronic bronchitis; FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity; Kco, carbon monoxide transfer coefficient; LLN, lower limit of normal; T_Lco, carbon monoxide transfer factor of the lung; V_A, alveolar volume.

defined until 2016.⁷ Notably, 8%–10% of children with PBB are complicated by bronchiectasis within the next 5 years,^{9 10} which raises the possibility that some positive auscultatory signs heard by a medical practitioner who diagnosed pneumonia during an acute respiratory illness could rather have actually been bronchiectasis in our most severe subgroup and high resolution CT scans would have ideally been performed to investigate this hypothesis.⁴⁵ Nonetheless, the awareness of PBB as a treatable and preventable cause of bronchiectasis is crucial especially for some socially disadvantaged communities, for example, Indigenous Australian populations.

The main strength of our TAHS cohort is the prospective, longitudinal data collection over five decades used to evaluate relationships between childhood respiratory health and long-term respiratory outcomes. The comprehensive questioning allowed us to create severity categories for childhood cough phenotypes that identified some severity–outcome relationships which strengthens the argument for a causal association. Although the severe subgroup with recurrent-protracted childhood bronchitis was small in this general population sample,

there was sufficient statistical power to detect significant associations with long-term outcomes enhanced by a large reference group without bronchitis in childhood.

Regarding limitations, our study relied on parental report of their children’s health at age 7 and self-report of doctor-diagnosed conditions at mean age 53. We acknowledge the potential for misclassification of childhood bronchitis (and asthma/wheezing), which could have been overestimated or underreported by parents according to its severity, but this misreporting of information across multiple exposure groups would bias estimates towards the null as such misclassification is more likely to be non-differential. There was also potential for differential misclassification of the self-reported adult lung conditions by exposure status. However, with the 46-year lag between exposure and outcome measurements and the likelihood of multiple intervening life events, such differential reporting by participants seemed unlikely. While a typical pattern of attrition was observed for males of lower socioeconomic class, the possibility of collider bias (from inappropriately controlling for a collider variable that is influenced independently by both the exposure

and outcome) is not likely because the prevalence of proxy outcomes 10 years earlier were largely unaffected and the loss to follow-up was similar between the exposure groups. Although several hypotheses were tested, these were predetermined by their biological plausibility based on previous literature^{9 10 19 22} and so we interpreted the general pattern of results rather than focusing on individual p values or using a Bonferroni correction. Finally, the Australian participants included in this study were almost exclusively of European ethnicity and in this era (born 1961) were not routinely immunised against common childhood pathogens, for example, measles, pneumococcus and *Haemophilus influenzae* type B. This may limit the generalisability of our findings to later birth cohorts and other ethnic groups in the vaccination era.

CONCLUSION

In summary, our findings suggest associations between childhood ‘bronchitis’ and doctor-diagnosed pneumonia and asthma in adulthood, especially evident for the recurrent-protracted subgroup, with trends across the spectrum of cough severity. The elevation of lung gas transfer in middle-age for this recurrent-protracted group is interesting and may be explained by asthma, chronic vascular changes and/or mild bronchiectasis; where possible, replication in other similar cohorts and further research to establish its clinical significance is warranted. Collectively, these findings provide novel insights into the potential natural history of severe bronchitis in children. Importantly, they support initiatives to enhance awareness among primary care physicians to promote the timely diagnosis and management of protracted episodes of cough in children with close monitoring for subsequent development of asthma and other complications such as bronchiectasis.

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