Mixed methods process evaluation of My Breathing Matters, a digital intervention to support self-management of asthma

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Abstract

This study aimed to explore user engagement with 'My Breathing Matters', a digital selfmanagement intervention for asthma, and identify factors that may influence engagement. In a mixed methods design, adults with asthma allocated to the intervention arm of a feasibility trial (n=44) participated in semi-structured interviews (n=18) and a satisfaction questionnaire (n=36) to explore their views and experiences of the intervention. Usage data highlighted that key intervention content was delivered to most users. The majority of questionnaire respondents (78%; n=28) reported they would recommend the intervention to friends and family. Interviewees expressed positive views of the intervention and experienced several benefits, mainly improved asthma control, medication use, and breathing technique. Factors that may influence user engagement were identified, including perceptions of asthma control, current self-management practices, and appeal of the target behaviours and behaviour change techniques. Findings suggested My Breathing Matters was acceptable and engaging to participants, and it was used as intended.

Keywords: asthma; mixed methods; digital intervention; process evaluation

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Introduction

Asthma is estimated to affect 358 million people worldwide¹. In the UK, 8 million people have been diagnosed with asthma². The goal of asthma management is optimal control asthma symptoms, to reduce the risk of exacerbations, and for individuals to be able to lead a full and productive life^{3,4}; however, this is not always achieved^{5,6}. Despite the availability of effective treatments, asthma outcomes remain sub-optimal, resulting in many avoidable deaths, hospital admissions, quality of life impairment, and societal costs^{5,7–9}. Clinical guidelines recommend promoting self-management through the provision of a personalised asthma action plan, attendance at annual asthma reviews, and correct inhaler technique use¹⁰. However, patient adherence to regular preventer medication, such as inhaled corticosteroids, is often low¹¹, patients' inhaler technique can be poor¹², personal asthma action plans are underused, and annual asthma reviews are underattended^{6,8}.

One potentially cost-effective method for promoting self-management is through digital interventions, which offer convenient 24-hour access to relevant and personalised self-management support. There is preliminary evidence that digital interventions for asthma self-management can lead to improvements in asthma control and quality of life, with no evidence of harm^{13,14}. However, there is currently a lack of robust evaluations of digital interventions for adults with asthma.

My Breathing Matters is an internet-based self-management intervention for asthma, which aims to improve the quality of life of adults with asthma through improved pharmacological (e.g. supporting medication adherence) and non-pharmacological (e.g. breathing retraining, stress reduction) self-management¹⁵. Other digital interventions for asthma focus on controlling asthma through pharmacological management or self-monitoring of physiological and behavioural data^{14,16}. Unique to this intervention was the integration of non-

pharmacological self-management, including breathing retraining¹⁷ and several previously evaluated interventions promoting healthy lifestyle behaviours (smoking cessation¹⁸, physical activity¹⁹, weight management²⁰, handwashing to prevent infections²¹). It was developed using theory-, evidence-, and person-based approaches to intervention development to maximise its effectiveness, feasibility and acceptability^{15,22,23}. Trial feasibility outcomes (such as recruitment, retention and randomisation) in our randomised controlled feasibility trial (RCT) with 88 adults with asthma showed that a definitive RCT is feasible²⁴. In addition, we observed consistent trends with improvements in asthma-related patient-reported outcome measures, including quality of life and asthma control. Before a definitive trial can be carried out, it is important to ensure that the intervention is acceptable to its target group and used as intended, to maximise its effectiveness²⁵.

To achieve this, we carried out a mixed methods process evaluation of My Breathing Matters embedded within the feasibility trial. Process evaluations can help support and refine an intervention's 'programme theory', which describes how an intervention is expected to lead to its effects (mechanisms of impact), the key intervention components, and how these interact with contextual factors (e.g. population, setting)^{26,27}. Users' 'engagement' with digital interventions has been hypothesised to moderate the intervention's influence on its mechanisms of impact²⁸. Engagement has been defined in terms of the extent to which an intervention is used (e.g. amount, frequency), the user's subjective experience of the intervention, and engagement with wider behavioural goals, such as behaviour change and self-management^{28,29}. Engagement is influenced by the digital intervention itself (content and delivery) and the context in which the intervention is used²⁸. In asthma, there is a lack of research on potential factors influencing engagement with digital interventions¹³.

We aimed to explore user engagement with My Breathing Matters by examining how participants in the feasibility study used the intervention, and exploring participants'

experiences of the intervention. To refine our programme theory, we sought to identify aspects of the intervention's delivery and content, and contextual factors (any external factors that might interact with the intervention to produce variations in the outcome) that may strengthen or impede users' engagement with the intervention.

Results

Participants

Intervention usage data were available for all 44 participants. The My Breathing Matters Satisfaction Questionnaire was administered to all 36 participants in the intervention group who registered with the intervention (eight participants were given a hyperlink to the intervention, but did not register). Seventeen intervention users and one non-user (n=18; 41%) agreed to be interviewed. Participants who did not take part either withdrew before their interview was due (n=4; 9%), could not be contacted by phone or email after multiple attempts (n=18; 41%) or were too busy (n=4; 9%). Table 1 provides the participants' demographics.

How did participants in the feasibility trial use the intervention?

Of the intervention participants, 81.8% (n=36) logged into My Breathing Matters at least once and between 1 and 25 times (Median = 4; IQR = 8). Those using the intervention more than once (n=27) used it between 1.89 to 337.85 days (Median = 120.96; IQR = 148.23). Each session (total sessions=231) lasted between 0.01-58.81 minutes (Median = 4.69; IQR = 8.33). Of the 34 participants who reached the core intervention content, most (73.5%; n=25) looked at both the pharmacological and non-pharmacological content and most (71%; n=24) chose to look at the non-pharmacological content first. Table 2 provides information on number of participants using each intervention component. The breathing retraining module was the most viewed component and over half of participants signed-up to the breathing retraining challenge tool. The other intervention tools were used by less than a third of participants.

What were intervention participants' experiences of the intervention?

In the My Breathing Matters Satisfaction Questionnaire, 86.1% (n=31) of the intervention users (n=36) reported that My Breathing Matters provided at least some benefit (Figure 1) and 69.4% (n=25) reported that there were 'no disadvantages at all' (Figure 2). A large majority of survey respondents (77.8%; n=28) reported that they would recommend My Breathing Matters to friends and family if they needed similar care and treatment (Figure 3). Content analysis of the free-text comments identified 14 benefits of using My Breathing Matters (n=28; Table 3) and nine disadvantages (n=13; Table 4). Information provision (n=12) and provision of breathing retraining (n=5) were the most commonly cited benefits. A dislike of the intervention's design (n=3) and that the intervention was not accessible on smartphones and computer tablets (n=3) were the most commonly cited disadvantages. Thematic analysis of the qualitative interviews identified four themes, which are outlined alongside the codes in Supplementary Table 1.

The first theme was '*Benefits of My Breathing Matters*'. Many participants reported how they noticed changes in their asthma symptoms since using My Breathing Matters, including reduced coughing, chest tightness, and breathlessness; improved peak flow; feeling more in control of their asthma; and fewer or no asthma attacks.

I'm not coughing when I wake up in the morning any more, or rarely. I'm not waking up in the night feeling tight-chested and that I can't breathe properly. (P14, 31-40 years old, female, asthma 21-30 years) This change was mainly attributed to the breathing retraining and improved medication use. In contrast, some interview participants said that they did not notice any changes to their asthma since using the intervention.

Participants who used the 4-week medication challenge (see Table 5) explained how this component had helped them get into the habit of using their preventer inhaler and use their inhalers correctly.

I haven't been terribly good at using the brown [preventer] inhaler. But I have pretty much got into the habit now and I would put that very much down to the website reminders. (P4, 61-70 years old, male, asthma 21-30 years).

Others reported how, since using My Breathing Matters, they had not needed to use their reliever inhaler as often. This was because they had not had any exacerbations, were using their preventer inhaler as prescribed, or started to practice the breathing techniques provided in the intervention when they were having symptoms instead.

Sometimes, I forget, you know, and I think, 'Oh actually, perhaps I should have taken it [reliever inhaler]', but then I think let's do my breathing techniques. Sometimes I haven't needed to take it...the website's been good for that. (P1, 41-50 years old, female, asthma 21-30 years)

One participant reported how the intervention reassured them that it was acceptable to use their reliever when they need to (rather than just tolerating symptoms), while another had been told by a health professional that their asthma had improved to a point that meant they no longer needed to use their preventer inhaler.

Many participants spoke how My Breathing Matters improved their breathing awareness, technique and posture.

I just feel that, sort of, before I used to breathe a lot through my mouth . . . And I find that, obviously, that now I'm breathing through my nose, my asthma's not as bad . . . I find that I'm not coughing as much. (P12, 21-30 years old, female, asthma <5 years)

Interview participants reported how My Breathing Matters had helped them to better identify, and deal with, asthma triggers (e.g. air pollution); gave them breathing and relaxation techniques to manage chest tightness and breathlessness; and prompted them to engage in healthy lifestyle changes (e.g. physical activity, healthy eating). A few participants explained how the intervention could help them to decide whether to seek health professional advice, and help them avoid unnecessary GP visits or burdening their healthcare team.

A few participants mentioned that their understanding of asthma and its treatment had improved. One participant learned how she should have had an asthma action plan, which she had printed and intended to take to her to asthma clinic.

It might have been useful if I'd had one of these [an action plan] years ago. Then I might have known what to do at the time [I had an asthma attack]. So that was extremely useful. (P8, 61-70 years old, female, asthma 5-10 years).

The action plan also prompted another participant to have conversations with their family about what they should do if he had an exacerbation and could not explain this to them at the time. On the other hand, some participants commented how they already knew a lot of the information, felt there was nothing new in the intervention, found some of the content repetitive, or believed the advice was common sense.

Some participants explained how the breathing retraining and stress management techniques helped them relax or stay calm, in particular when they were feeling tight chested, panicking when having an asthma attack, and for trying to get to sleep. A few participants explained how My Breathing Matters could make people think more positively about asthma, especially if you have just been diagnosed.

I think maybe that's what I've really gained from it [My Breathing Matters], I've thought about it [asthma] more and if you think about problems or if you think about different things then that's a good thing to, you know, you're actively trying to improve something about it and, yeah, so I'm definitely thinking more positively. (P11, 41-50 years old, female, asthma 21-30 years)

Other benefits included addressing any asthma concerns you might have (e.g. side effects of medication, symptoms); providing reassurance that there are things that can help them cope; and highlighting that people with asthma are not alone and that there are other people with asthma or similar problems.

The second theme was '*Views on the intervention content*'. Participants particularly valued the breathing retraining, with many finding this the most helpful component. Most participants liked the videos and found the techniques relatively easy to learn. A few people found some of the techniques difficult to learn, including slow breathing and controlled breath holding, with one person preferring to have received the training in person. Another participant did not understand why the breathing exercises were beneficial and found the video irritating.

Some participants did not want to rely solely on their asthma medication to manage their asthma and liked that My Breathing Matters provided alternative management strategies, mainly the breathing retraining.

Anything that helps you only take the amount of medication you really need and helps you to self-control asthma in some way. And if My Breathing helps you to do that, that's got to be a good thing. (P2, 61-70 years old, female, asthma 11-20 years)

Now, after using the website, it's made me think about, well, what other things can I do to help myself, so that I don't have to rely on my inhaler so much? (P12, 21-30 years old, female, asthma <5 years)

In the 4-week medication challenge (see Table 5), participants valued the email reminders, the advice about keeping their inhalers somewhere accessible as a reminder, and the realisation that it was benefiting them. The other intervention components (action plan, annual asthma review, stress management, friends and family) were used to a lesser extent. Most participants either had not yet used the component or found that these components were not relevant to them. None of the interview participants reported contacting the Asthma UK helpline when asked about this.

The third theme was 'Views on the intervention design'. Participants expressed positive views on the intervention design and found the content easy to understand. Some participants liked that it was designed by an experienced team and that it was associated with a national charity (Asthma UK), and felt that the information was authentic and high quality. Generally, people found the intervention easy to navigate. However, a few people experienced navigation and technical difficulties, including logging on, following URLs in emails, and accessing the intervention by phone or tablet, or on their workplace computer. Participants expressed mixed views regarding the unlocking feature of My Breathing Matters, whereby new content was made available to users over time. Some liked this feature as it structured their learning and stopped them from feeling overwhelmed by too much information, but others found it frustrating or did not understand the reasoning behind the feature.

I'd have been bombarded with it all if it was too much at once, so it was quite nice it came in sections slowly . . . it's too much to take in otherwise. (P1, 41-50 years old, female, asthma 31-40 years)

I wanted to look through other bits that weren't enabled and then had to wait for them ... I think that probably would discourage me from using it. (P4, 61-70 years old, male, asthma 21-30 years)

Participants liked regularly receiving regular emails with additional behavioural content from My Breathing Matters because they reminded them to take their medication and use the website, provided encouragement and additional advice, and facilitated quick access back to the website. A few people expressed negative views of the emails, including finding the email content irritating or not useful, or that it made them feel guilty for not using the website.

The forth theme was '*Contextual factors influencing intervention engagement*'. Participants' engagement with My Breathing Matters was influenced by their perceptions of their asthma control. Participants explained how they did not engage with the intervention or specific components (e.g. the Asthma UK helpline, action plan, or the medication section) because they did not think their asthma was severe enough.

I possibly briefly looked at the sort of action plan thing, but decided that, actually, I didn't think it was gonna be beneficial for me . . . I just thought that probably my asthma wasn't severe enough that it was something that I needed to do at that moment in time. (P12, 21-30 years old, female, asthma <5 years)

Likewise, participants explained that they were more likely to use the intervention when their asthma symptoms were bad (e.g. in the winter or during allergy season) and less likely to use it when their asthma was well controlled. A few participants explained how, most of the time, they simply 'forgot' or tried not to 'dwell' on their asthma unless it was significantly restricting their lives.

My asthma is fairly well controlled, I haven't needed to refer to the [My Breathing Matters] site...*I'm very much a kind of person that, actually, I don't dwell on, you*

know, things that might inhibit you in life, and just get on with life . . . You know, I've had far more worse than asthma. (P15, 61-70 years old, female, asthma 31-40 years)

On the other hand, two people were unlikely to use it if it their asthma was bad, instead choosing to seek medical attention.

Some participants explained that they did not use certain components because they did not consider them relevant. For example, they already practiced the recommended behaviours (e.g. taking medication, attending reviews, being more active), were not stressed (relating to the stress management techniques), or their family or friends already knew about asthma. Participants explained how they thought My Breathing Matters would be most useful at the beginning of their asthma journey, once you have been diagnosed with asthma. Likewise, some people who have had asthma for a long time reported it was less useful. A few participants explained how they were not confident with using computers or expressed a dislike towards them. The non-user we interviewed was keen to use the intervention, but felt he lacked the computer skills to log onto it. Other reported reasons for low usage or not using certain components included lack of time or being busy with other priorities (e.g. work, family); and comorbidities that made some of the intervention behaviours (physical activity, breathing exercises) challenging.

What factors may influence user engagement with My Breathing Matters?

Across the qualitative findings, we identified several contextual factors and aspects of the intervention's content and delivery that may influence user engagement with the intervention (Figure 4). Contextual factors were derived from the interview data (theme 4) and included pre-existing beliefs (e.g. perceptions of asthma control/asthma-related quality of life, beliefs about medication), knowledge of asthma management and skills (e.g. confidence with computers), current self-management practices, environmental factors (current season, lack of

time), and health status (time since diagnosis, comorbidities). The interview data and qualitative data from the My Breathing Matters Satisfaction Questionnaire highlighted aspects of the intervention's content and delivery that may influence engagement including appeal and perceived ease of the target behaviours (e.g. breathing retraining); appeal of the behaviour change techniques (e.g. email reminders) and design (e.g. content released over time, instructional videos); novelty, relevance and clarity of the intervention content; and ease of use (navigation and accessing the website). Users reported both positive and negative aspects, and both are summarised along with the perceived benefits of the intervention (derived from theme 1 of the interview data and the qualitative questionnaire data) in Figure 4.

Discussion

This mixed methods process evaluation study explored users' engagement with My Breathing Matters, an internet-based self-management intervention for asthma. Overall, engagement with the intervention was high, it was used as intended, and people with asthma expressed positive views of the intervention, its intervention components, and its design features; thus, demonstrating that it was acceptable to participants. Users reported experiencing several benefits of the intervention, mainly improved asthma control, medication use, and breathing technique. These perceived benefits were in line with the hypothesised intervention mechanisms of impact and outcomes outlined in our original logic model. Our study findings also extended our current programme theory by identifying aspects of the intervention (content and delivery), and contextual factors that may influence user engagement with the intervention.

Despite our attempts to engage those who did not perceive themselves as having active asthma and only recruiting those with impaired asthma-related quality of life, users still

questioned the relevance of the intervention and its components, and did not believe that their asthma was severe enough for the intervention. This mirrors other studies that have demonstrated disparities between perceived and objective measures of asthma control, with patients overestimating how well their asthma was controlled^{30,31}. Notably, user engagement in this study was high despite such beliefs. This may be due to our use of 'positive illness contexts' as a key intervention design feature (promoting health rather than preventing illness). In this way, even when users considered the intervention not specifically necessary for asthma control, My Breathing Matters still provided self-management support. Users reported several benefits of the intervention, and our feasibility study observed trends with improvement across a range of asthma outcomes²⁴. This demonstrates that interventions developed using theory-, evidence-, and person-based approaches that target likely barriers to behaviour change can lead to effective user engagement and positive outcomes among individuals with different health beliefs, such as those in heterogeneous chronic disease populations.

Uniquely, My Breathing Matters integrated breathing retraining alongside established pharmacological self-management approaches. Consistent with other qualitative evaluations of breathing retraining^{32,33}, users valued how the non-pharmacological approaches in My Breathing Matters could help reduce their reliance on medication, which is an important goal for people with asthma³⁴. Most participants were satisfied with the online delivery of breathing retraining, with just a few users finding the exercises difficult to learn and only one participant reporting that they would have preferred to receive their training face-to-face with a health professional; thus further demonstrating the feasibility of delivering breathing retraining via an unguided digital intervention. A trial of breathing retraining demonstrated that face-to-face delivery was no more effective than DVD delivery¹⁷.

In an attempt to maximise user engagement and ensure all core content was accessed³¹, we implemented a design feature whereby new content was made available to users over time. Although this feature had been used successfully in other interventions³⁵ and some study participants found this feature helpful, others found this feature frustrating and did not understand the rationale behind it. It may be that by restricting users' access to specific content, the intervention may have impaired their sense of control and autonomy, which are important factors for maximising engagement²⁸. In future versions of the interventions, it would be helpful to provide users with a strong rationale for this feature (e.g. to encourage people to practice the techniques they have already accessed before trying new techniques), but allow users to unlock additional content themselves if they wished to maximise user autonomy^{22,28} and avoid disengaging active users.

One strength of this study was its mixed methods design. The triangulation of questionnaire measures with qualitative interviews and usage data enabled us to explore different aspects of intervention engagement and to increase the credibility of the research. Even though some questionnaires such as the My Breathing Matters Satisfaction Questionnaire were not formally validated, we could examine the extent to which the intervention is used, and users' subjective experiences of using the intervention and enacting its target behaviours (e.g. breathing retraining). Due to the limited sample size of the feasibility trial (*n*=88), we were not powered to do a more in-depth analysis of the usage data. A fully powered RCT is needed to explore how process measures, such as perceptions of asthma, pre-intervention levels of medication adherence, and time since diagnosis, is associated with user engagement and asthma outcomes. It would also be worthwhile exploring how usage might change across the seasons, given that some participants explained how they were more likely to use the intervention during certain seasons, when their asthma symptoms were worse. Although we endeavoured to recruit participants across a broad demographic range, participants were

generally older and white, and had high levels of educational attainment. They were also recruited from a feasibly trial sample, so are unlikely to be representative of the wider asthma population³⁶. A wider reach would avoid further worsening the digital divide and health inequalities. Moreover, the small sample size of the feasibility study meant that we could not purposively sample participants based on their usage and were, therefore, only able to recruit one non-user. A larger sample size would have allowed us to better target and capture the views of non-users and those who were less engaged with the intervention. Interviews with the control group would have allowed us to explore their experiences with usual care, in order to explore which perceived benefits are unique to the intervention and not from the feasibility trial itself. Interviews with those who declined to take part in the trial would have also given us useful insights into their reasons for this and how user engagement might be improved³⁷.

Our findings demonstrated that My Breathing Matters is acceptable and engaging to its target group, and the intervention was delivered and worked as intended. The person-based approach to intervention development was key to maximising intervention engagement and acceptability for adults with asthma. Along with the findings from the feasibility trial, the current study supports the move towards a fully powered RCT, including a mediation and moderation analysis, with only minor modifications to the intervention content required. More broadly, our findings highlight aspects of intervention content and delivery (such as targeting key issues using person-based approaches, providing non-pharmacological selfmanagement approaches), and contextual factors (such as perceptions of asthma control, current self-management practices) that may influence user engagement with digital asthma interventions. These should be considered when implementing the intervention or when developing asthma behaviour change interventions.

Methods

Design

A convergent mixed methods design was used for the process evaluation in which qualitative and quantitative methods are implemented in the same research phase and given equal weight, but the data is analysed separately³⁸. The process evaluation was embedded in a feasibility RCT of My Breathing Matters. Trial participants were randomised into an intervention group who were given access to My Breathing Matters or a usual care group. Outcome measures were assessed at baseline, 3 months and 12 months. Further details on the trial methods and feasibility outcomes are available elsewhere²⁴. Quantitative usage data were collected to describe patterns of intervention usage over the 12-month study period. The My Breathing Matters Satisfaction Questionnaire was devised for this study and administered to intervention participants at 12-month follow-up to assess their satisfaction with the intervention. Qualitative interviews were carried out to explore intervention participants' views and experiences of My Breathing Matters. Ethical approval was granted by the University of Southampton and South Central - Berkshire Research Ethics Committee (REC reference: 16/SC/0614). To increase the transferability of the research, the COREQ checklist³⁹ was used to guide reporting the qualitative research (Supplementary Table 2) and ensure a rich description of the participants and the research process.

Intervention

My Breathing Matters was systematically developed using person-, evidence-, and theorybased approaches, drawing upon primary mixed methods research^{17,31,40}, quantitative¹⁴ and qualitative⁴¹ systematic reviews, and consultation with Public and Patient Involvement (PPI) representatives and clinical and intervention development experts.

Following a person-based approach²², guiding principles were created, including intervention design objectives and design features to address key issues, needs and behavioural challenges

of the target population identified in the evidence synthesis stage. One key behavioural issue that emerged from the literature search conducted in the intervention development phase is that some people with non-optimal asthma control do not consider themselves as people with active asthma^{42–44}. Therefore, one intervention design objective was to specifically engage this group. To do this, the intervention maintained a positive illness context throughout (referring to 'keeping breathing healthy' rather than 'preventing asthma symptoms'), provided optional and flexible support only when needed, and promoted the belief that impaired quality of life can be improved (Supplementary Figure 1). To target influences on asthma control that are not often acknowledged, such as anxiety, stress and lifestyle (e.g. smoking, obesity, avoidance of physical activity), other design objectives aimed to encourage users to engage in non-pharmacological (e.g. breathing retraining, stress management, lifestyle changes), as well as pharmacological self-management, to improve asthma control (see Yardley et al. 2015¹⁵ for this process in more detail).

Theory-based behaviour analysis was used to identify the influences on target behaviours and the intervention components that could address these, and describe the intervention in terms of existing theory and programme level theory. A logic model was created to illustrate the hypothesised mechanisms of impact that explain how My Breathing Matters is expected to lead to improvements in asthma-related quality of life (Figure 5).My Breathing Matters is hypothesised to improve asthma-related quality of life through behavioural adherence (improved pharmacological and non-pharmacological management, engagement with the intervention), improved physiological outcomes (asthma control, lung function, exacerbations), and improved psychological outcomes (stress, mood, enablement). Table 5 outlines the components of the intervention in more detail.

An intervention prototype was developed and, consistent with a person-based approach, the views and experiences of adults with asthma who used the intervention were explored using

iterative qualitative methods (think aloud and retrospective semi-structured interviews) and the intervention was modified in response to this feedback.

On each unique login, users were asked to complete a brief quality of life assessment measuring activities, sleep, stress, illness, and reliever medication use (Supplementary Figure 2). Based on their answers, users were signposted to relevant content. Content was not available all at once, rather different content was 'unlocked' at various time points after the user's first visit to the website to encourage long-term engagement with the intervention (Supplementary Figure 3). The intervention is self-directed, but the contact details for the Asthma UK helpline were given to provide additional support if required. The intervention is available at mybreathingmatters.co.uk.

Participants and recruitment

Participants were eligible for the feasibility trial if they were aged 18 years or over, had physician-diagnosed asthma managed in primary care, had received at least one anti-asthma medication prescription in the previous year, and could use the Internet (self-judged). Anti-asthma medication included all commonly used inhaled and oral preparations for asthma treatment (both regular medication and as-required reliever preparations), such as inhaled corticosteroids, long and short acting beta agonists and leukotriene receptor antagonists. No patients were receiving injected biological treatments or maintenance oral corticosteroids. Participants also needed to have an impaired asthma-related health status at baseline, defined as a Mini Asthma Quality of Life Questionnaire⁴⁵ score of less than 5.5. Full trial inclusion and exclusion criteria are described elsewhere²⁴.

Eligible participants were identified and invited to take part in the trial by seven general practices from the Wessex, UK primary care research network. After the 3-month follow-up, all intervention group participants (n=44) were approached by phone or email by a member

of the study team and were invited to take part in a qualitative interview, irrespective of whether they used the intervention. Drawing on the guidelines on information power in qualitative interview studies⁴⁶, we aimed to recruit approximately 20 participants to the interview study. This number was deemed adequate given the study's narrow aim (views on one intervention), the small source population (n=44), the specificity of the experiences, knowledge and properties among the intervention trial participants, and the likely high quality of dialogue from using an experienced qualitative researcher. Informed consent was obtained for all trial participants. Participants received a £10 shopping voucher for submitting their follow-up questionnaires at 12 months. Interview participants did not receive any additional incentives for taking part.

Data collection

Intervention usage was automatically collected by the LifeGuide software (https://www.lifeguideonline.org), which was used to create and host the intervention. Data were collected on the number and duration of logins, date of last login, and pages visited. Participants were informed that they could use the intervention as much or as little as they liked.

The My Breathing Matters Satisfaction Questionnaire (Supplementary Note 1) was administered by paper at the 12-month follow-up appointment with a research nurse to those who registered with the intervention. Better understanding of the potential benefits and burdens of health interventions can help us to optimise these interventions and improve their effectiveness^{47,48}. To explore these two aspects, we devised two items to assess benefits gained from using the intervention and disadvantages of the intervention, and open questions allowed participants to report any benefits and disadvantages. These items were developed in discussion with our multidisciplinary intervention development team, consisting of experts in

intervention development and evaluation, behavioural science, and health economics. The one-item NHS Friends and Family Test⁴⁹ assessed how likely participants are to recommend the intervention to friends and family if they needed similar care and treatment using a 5-point Likert scale ranging from 'extremely likely' to 'extremely unlikely', with a 'don't know' option. This tool is used by NHS England to assess patient satisfaction across a wide range of services.

For the qualitative interviews, a semi-structured interview schedule was developed by experts in health psychology (KG, BA, LY) and asthma (MT, BA, AB), and a PPI representative with asthma (DR). Interview questions were designed to explore the key functions for process evaluation outlined in the Medical Research Council process evaluation guidelines⁵⁰: implementation (what was delivered), mechanisms of impact, and contextual factors. Specifically, the questions explored participants' experiences of the intervention and its components, how they used the intervention, their perceived advantages and disadvantages of the intervention, times they were more and less likely to use the intervention, and reasons for any non-usage (See Supplementary Note 2 for interview schedule). Open-ended questions were used to ascertain the most important issues or challenges for participants.

Interviews were carried out by telephone by KG (female health psychologist and research fellow who was experiences in qualitative research) who was not involved in intervention development and did not know the participants prior to the interviews. Participants were told that the interviews aimed to explore their view and experiences to help improve the research and intervention for future users. Interviews took place between July 2017 and January 2018, lasted between 21-65 minutes, were audio-recorded, and transcribed verbatim.

Data analysis

The intervention usage data and the closed questions of the My Breathing Matters Satisfaction Questionnaire were analysed using descriptive statistics to describe patterns of intervention usage. Content analysis⁵¹ was carried out on the open question data to identify benefits and disadvantages of using the intervention.

The qualitative interviews were analysed using inductive thematic analysis^{51,52}. Data analysis was assisted by QSR's NVivo 11 qualitative data analysis software (QSR International Pty Ltd., 2017). Analysis was informed by guidelines for establishing trustworthiness in qualitative research^{53–56}. KG familiarised herself with the data through repeated reading of the transcripts. Initial codes were generated that were grounded in the data and a coding manual was developed that listed all codes and themes, including descriptions and example quotes from the text. To increase the credibility of the research, the final coding manual was discussed and agreed with two other researchers (BA and LY) and the final interpretations in the results section were reviewed and agreed by all authors, as well as two PPI representatives. The constant comparison method⁵⁷, a grounded theory technique, was used to compare codes across different participants, contexts, and situations. Disconfirming case analysis⁵⁴ was used to actively identify data that did not fit with the identified themes. These two techniques were used to ensure the analysis was carried out with rigor and to increase its credibility. Participant quotes were used in the final write-up to illustrate the themes and pseudonyms used to refer to these participants. Data saturation was considered reached because participants in later interviews did not indicate any significant new benefits, concerns or barriers to engagement with My Breathing Matters.

Once the qualitative analysis was complete, we reviewed key findings from the interviews and My Breathing Matters Satisfaction Questionnaire to identify contextual factors and

aspects of the intervention's content and delivery that may have influenced user engagement with the intervention.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Competing interests

Neither MT nor any member of his close family has any shares in pharmaceutical companies. In the last 3 years, he has received speaker's honoraria for speaking at sponsored meetings or satellite symposia at conferences from the following companies marketing respiratory and allergy products: GSK, Novartis. He has received honoraria for attending advisory panels with; Boehringer Inglehiem, GSK, Novartis. He is a recent a member of the BTS SIGN Asthma guideline steering group and the NICE Asthma Diagnosis and Monitoring guideline development group. KG, BA, AB, EM, DR, and LY have no competing interests.

Author Contributions

All authors designed the study. KG was responsible for recruitment, data collection and analysis the data, with support from BA. KG drafted the manuscript with initial support from BA, LY, MT, and AB. All authors critically reviewed the manuscript, contributing important intellectual content and approved the final manuscript.

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Figure legends

Figure 1 Participant responses to single item relating to benefits of using My Breathing Matters.

Figure 2 Participant responses to item relating to disadvantages of using My Breathing Matters.

Figure 3 Participant responses to NHS friends and family test relating to how likely they would be to recommend My Breathing Matters to friends and family if they needed similar care and treatment.

Figure 4 Summary of the qualitative findings demonstrating the contextual factors and aspects of the intervention content and delivery that may influence engagement with My Breathing Matters and the perceived benefits.

Figure 5 Logic Model of My Breathing Matters intervention to improve quality of life in patients with asthma. Key: ^aUptake and engagement facilitation; ^bPharmacological support; ^cNon-pharmacological support.

Undergraduate qualification (e.g. Degree,	Postgraduate qualification (e.g. Masters, PhD)	Highert level of education (0/)	Single	Separated	Divorced	Widowed	Living with a partner	Married	Marital status (%)	Range	Mean (SD)	Time since asthma diagnosis (years)	Mean (SD)	Baseline asthma quality of life (AQLQ) score	Other	White	Ethnicity n (%)	Male	Female	Gender n (%)	Range	Mean (SD)	Age			
20 (45.5)	4 (9.1)		5 (11.4)	2 (4.5)	1 (2.3)	4 (9.1)	4 (9.1)	28 (63.6)		$1.3-64.0^{1}$	$25.2(17.2)^{1}$		4.9 (0.9)		2 (4.5)	42 (95.5)		17 (38.6)	27 (61.4)		20-78	57.0 (14.2)		participants ($n=44$)	All intervention	
19 (52.8)	3 (8.3)	•	4 (11.1)	1 (2.8)	1 (2.8)	3 (8.3)	3 (8.3)	24 (66.7)		$1.3-64.0^2$	$25.8(18.0)^2$		5.0 (0.9)		0	36 (100.0)		13 (36.1)	23 (63.9)		20-78	56.8 (15.1)			Users (n=36)	
1 (12.5)	1 (12.5)		1 (12.5)	1 (12.5)	0	1 (12.5)	1 (12.5)	4 (50.0)		3.0-43.0	22.8 (14.4)		4.2 (0.8)		2 (25.0)	6 (75.0)		4 (50.0)	4 (50.0)		43-77	57.9 (10.3)			Non-users (n=8)	
6 (33.3)	3 (16.7)		0	0	0	3 (16.7)	2 (11.1)	13 (72.2)		$1.3-64.0^3$	$24.6(19.7)^3$		5.0(1.1)		0	18 (100.0)		6 (33.3)	12 (66.7)		29-77	60.3 (13.2)			Interviewed (n=18)	
14 (53.8)	1 (3.8)		5 (19.2)	2 (7.7)	1 (3.8)	1 (3.8)	2 (7.7)	15 (57.7)		3.0-64.0	25.6 (15.8)		4.7 (0.8)		2 (7.7)	24 (92.3)		11 (42.3)	15 (57.7)		20-78	54.7 (14.7)		(n=26)	Not interviewed	

Table 1 Participant demographics at baseline

Key: ${}^{n}n=43$; ${}^{2}n=35$; ${}^{3}n=17$; ${}^{4}n=42$; ${}^{5}n=34$; ${}^{6}n=24$	Internet use per week (hours) Mean (SD)	Age left full-time education Mean (SD)	No formal educational qualifications	School leaver (e.g. GCSEs, O-levels)	Further education (e.g. A-Levels, ONC, OND)
; $GCSEs = General C$	13.1 (12.4)	19.4 (7.0) ⁴	5 (11.4)	10 (22.7)	5 (11.4)
ertificate of Secondar	13.2 (11.9)	19.5 (6.8) ⁵	2 (5.6)	7 (19.4)	5 (13.9)
y Education; O-Level	12.9 (15.0)	18.9 (8.3)	3 (37.5)	3 (37.5)	0
= The General Certific	12.4 (13.5)	20.7 (9.0)	2 (11.1)	5 (27.8)	2 (11.1)
ate of Education	13.7 (11.7)	18.4 (5.0) ⁶	3 (11.5)	5 (19.2)	3 (11.5)
	l		l		

Ordinary Level; A-Level = The General Certificate of Education Advanced Level; UNC = Ordinary National Certificate; UND = Ordinary National Diploma

Intervention component	Participants who viewed at	Participants who used the
	least one page of the session	main tool in the session ¹
	n (%)	n (%)
Pharmacological content		
Medication Advice	21 (58.3%)	n/a
4-week challenge	19 (52.8%)	$10 (27.8\%)^1$
Personal Asthma Action Plan	16 (44.4%)	$6(16.7\%)^2$
Annual Asthma Review	16 (44.4%)	$1 (2.8\%)^3$
Non-Pharmacological content		
Breathing Retraining	27 (79.4%)	20 (55.6%) ⁴
Stress Management	13 (36.1%)	6 (16.7%) ⁵
Friends & Family Support	10 (27.8%)	$1 (2.8\%)^6$
Lifestyle changes		
Weight management	3 (8.3%)	n/a
Physical activity	3 (8.3%)	n/a
Handwashing	2 (7.7%)	n/a
Smoking cessation	0	n/a

Table 2 Numbers (and percentages) of participants who used each intervention component (*n*=36)

Key: ¹Signing up to the 4-week challenge; ²Viewed blank plan or made online plan; ³Booked an

appointment with GP and recorded the appointment online; ⁴Signed-up to breathing retraining; ⁵Using the stress management tools; ⁶Emailed someone a link to the Friend & Family module.

Code	Description	Frequency
Information provision	Intervention had improved awareness, improved or	12
	validated understanding about asthma and its	
	management. Participants liked the lifestyle advice and	
	tips on management. The information in the intervention	
	was reliable and clear.	
Provision of breathing	The breathing exercises were cited as a benefit of using	5
retraining	the intervention. The intervention helped them realise the	
	benefits of the breathing exercises/correct breathing,	
	learn correct breathing and be more aware of their	
	breathing.	
Medication adherence	The intervention helped people to build a medication	3
	habit, 'take notice' of medication, and made them realise	
	they should be using a preventer inhaler regularly.	
Lifestyle changes	The intervention provided lifestyle advice, including	3
	healthy eating, weight management, and physical	
	activity. Participants had lost weight and increased their	
	physical activity since using the intervention.	
Reassurance	The intervention reassured people that their asthma	3
	symptoms were normal, that they were doing the right	
	things to manage their asthma, and confirmed what they	
	already knew.	
Relaxation	The intervention helped people to relax. Participants	3
	started doing the relaxation techniques and they helped	
	one participant get to sleep.	
Access to information	The intervention provides access to information quickly	2
	and easily. The intervention can be accessed at home.	
Control of asthma	The intervention helped people to control their asthma	2
symptoms	symptoms or improved their lung function.	
Motivation for asthma	The intervention makes people think more about their	2
self-management	asthma and gives them to motivation to manage their	
	asthma.	
Provision of action	Being given access to an action plan/made aware of it.	2
plan		
Speaking to friends	Two participants had discussed asthma and its	2
and family	management with family and friends.	
Dealing with triggers	The intervention helped one person deal with asthma	1
	triggers.	

Table 3 Content analysis of free-text comments regarding benefits of using My Breathing Matters (*n*=28)

General health	The intervention made one person think about their	1
	general health, as well as asthma.	
Support	The intervention made one participant feel they were	1
	being taken care of.	

Code	Description	Frequency
Disliked design	Aspects of the design participants disliked, in particular the unlocking feature.	3
Not accessible on their device	The intervention could not be accessed on phone or computer tablet.	3
Difficulties logging on	Participants experienced difficulties logging on	2
Too many or too little emails	Participants received too many or not enough emails to keep engaged.	2
Annoying	The intervention was annoying or slow.	1
Boring	The intervention was short and became boring after a few months.	1
Lack of human contact	The intervention did not provide one-to-one human contact to allow the participant's questions to be answered.	1
Patronising	The intervention was patronising.	1
Time consuming	Participant did not have time to do the breathing exercises during the day.	1

Table 4 Content analysis of free-text comments regarding disadvantages of using My Breathing Matters (*n*=13)

Target behaviour	De	escription
Improved preventer	•	Information about the benefits of medication use for prevention
medication adherence		of asthma symptoms.
	•	Addressing 'common concerns' about asthma medication.
	•	A 4-week challenge (in which users were encouraged to engage
		in habitual optimal preventer inhaler use) to help people
		develop positive medication habits.
Appropriate healthcare	•	Tools to create and store a Personal Asthma Action Plan and
service use		provide encouragement for its use.
	•	Provide encouragement to attend an annual Asthma Review.
Engagement with breathing	•	A breathing retraining programme ¹⁷ to help control asthma
retraining		symptoms, including videos on how to improve your breathing
		technique.
Engagement with stress	•	Provision of stress management techniques, including
management		relaxation, and advice on stress management (e.g. time
		management) and adaptive ways of thinking (e.g. thought
		awareness, using positive thinking, talking through your
		worries), to reduce asthma-related stress.
Send information to friends	•	Ability to send friends and family a hyperlink to relevant
and family to encourage		information about asthma treatment and symptoms.
them to engage in asthma		
management		
Lifestyle changes	٠	Access to previously developed lifestyle change programmes
		adapted for asthma, including:
		• StopAdvisor ¹⁸ to support smoking cessation,

Table 5 Description of My Breathing Matters intervention components

- Getting Active¹⁹ to increase physical activity adapted for asthma,
- POWeR²⁰ to support weight management,
- \circ Germ Defence²¹ to promote handwashing to prevent

infections.



Figure 1

Figure 2





Figure 3



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understand

Ease of navigation and easy to

During the intervention



• •



Supplementary Material

Theme	Codes
Benefits of My Breathing Matters	Improved asthma & symptoms
	No noticeable change in asthma or breathing
	Reduction in medication use
	Improved medication adherence
	Provides an alternative to medication
	Improved breathing technique & posture
	Identifying, and dealing with, asthma triggers
	Prompts lifestyle changes
	Managing breathlessness and chest tightness
	Facilitates self-management
	Improved knowledge of asthma and its treatment
	Better use of healthcare resources
	Relaxation
	Thinking more positively about asthma
	Provides reassurance
	Feeling less alone
Views on the intervention content	Views on 4-week challenge
	Views on personalised asthma action plan (PAAP)
	Views on asthma review
	Views on breathing retraining
	Views on stress management

Supplementary Table 1 Themes and codes from data analysis for qualitative interviews

	Views on Friends & Family section
	Views on lifestyle modules
	Views on Asthma UK & helpline
Views on the intervention design	Views on website appearance
	Views on intervention credibility
	Views on delivery format
	Views on emails
	Views on information and advice
	Views on information novelty
	Views on interactive features
	Views on usability
	Views on information architecture
Contextual factors influencing	Relevance of intervention components
intervention engagement	Perceptions of asthma severity
	Time since diagnosis
	Confidence with, and dislike of, computers
	Season
	Other priorities
	Other health problems

No	Item	Guide	Comments	Location in
		questions/description		manuscript
Dom	ain 1: Research team	and reflexivity		
Perso	onal Characteristics			
1.	Interviewer/facilitat	Which author/s	KG	Methods (Data
	or	conducted the interview		collection)
		or focus group?		
2.	Credentials	What were the	KG – PhD, CPsychol.	Methods (Data
		researcher's		collection)
		credentials? E.g. PhD,		
		MD		
3.	Occupation	What was their	KG – Health Psychologist	Methods (Data
		occupation at the time of	& Research Fellow	collection)
		the study?		
4.	Gender	Was the researcher male	Female	Methods (Data
		or female?		collection)
5.	Experience and	What experience or	Experienced qualitative	Methods (Data
	training	training did the	postdoctoral researcher.	collection)
		researcher have?		
Relat	tionship with participa	nts		
6.	Relationship	Was a relationship	Participants were not	Methods (Data
	established	established prior to study	known to the researcher.	collection)
		commencement?		

Supplementary Table 2 COREQ checklist for qualitative interviews

No	Item	Guide	Comments	Location in
		questions/description		manuscript
7.	Participant	What did the participants	Participants were told that	Methods (Data
	knowledge of the	know about the	the interviews aimed to	collection)
	interviewer	researcher? e.g. personal	explore their view and	
		goals, reasons for doing	experiences to help	
		the research	improve the research and	
			intervention for future	
			users.	
8.	Interviewer	What characteristics	The researcher was not	Methods (Data
	characteristics	were reported about the	involved in intervention	collection)
		interviewer/facilitator?	development, although	
		e.g. Bias, assumptions,	she was part of the same	
		reasons and interests in	digital research team,	
		the research topic	which may have been a	
			source of bias.	
Dom	ain 2: study design	I	I	
Theo	retical framework			
9.	Methodological	What methodological	Thematic analysis.	Methods (Data
	orientation and	orientation was stated to		collection)
	Theory	underpin the study? <i>e.g.</i>		
		grounded theory,		
		discourse analysis,		
		ethnography,		
		phenomenology, content		
		analysis		
Parti	cipant selection	1	1	L

No	Item	Guide	Comments	Location in
		questions/description		manuscript
10.	Sampling	How were participants	All intervention	Methods
		selected? e.g. purposive,	participants from the	(Participants
		convenience,	feasibility trial were	and
		consecutive, snowball	approached.	recruitment)
11.	Method of	How were participants	All intervention group	Methods
	approach	approached? e.g. face-to-	participants were	(Participants
		face, telephone, mail,	approached by phone or	and
		email	email by a member of the	recruitment)
			study team and were	
			invited to take part.	
12.	Sample size	How many participants	18	Results
		were in the study?		(Participants)
13.	Non-participation	How many people	Participants who did not	Results
		refused to participate or	take part either withdrew	(Participants)
		dropped out? Reasons?	before their interview was	
			due (<i>n</i> =4; 9%), could not	
			be contacted by phone or	
			email after multiple	
			attempts (<i>n</i> =18; 41%) or	
			were too busy (<i>n</i> =4; 9%).	
Settin	ıg	1	1	1
14.	Setting of data	Where was the data	Telephone	Methods (Data
	collection	collected? e.g. home,		collection)
		clinic, workplace		

No	Item	Guide	Comments	Location in
		questions/description		manuscript
15.	Presence of non-	Was anyone else present	No	-
	participants	besides the participants		
		and researchers?		
16.	Description of	What are the important	Demographic information	Table 1;
	sample	characteristics of the	can be found in Table 1.	Methods (data
		sample? <i>e.g.</i>	Interviews took place	collection)
		demographic data, date	between July 2017 and	
			January 2018	
Data	collection			
17.	Interview guide	Were questions,	The interview schedules	Supplementary
		prompts, guides	can be found in	Note 2
		provided by the authors?	Supplementary Note 2. It	
		Was it pilot tested?	was reviewed a PPI	
			representative.	
18.	Repeat interviews	Were repeat interviews	Only single interviews	-
		carried out? If yes, how	were carried out.	
		many?		
19.	Audio/visual	Did the research use	Audio	Methods (Data
	recording	audio or visual recording		collection)
		to collect the data?		
20.	Field notes	Were field notes made	Field notes were not	-
		during and/or after the	made.	
		interview or focus		
		group?		

No	Item	Guide	Comments	Location in
		questions/description		manuscript
21.	Duration	What was the duration of	Between 21-65 minutes	Methods (Data
		the interviews or focus		collection)
		group?		
22.	Data saturation	Was data saturation	Data saturation was	Methods (Data
		discussed?	considered reached	analysis)
			because participants in	
			later interviews did not	
			indicate any significant	
			new benefits, concerns or	
			barriers to engagement	
			with My Breathing	
			Matters.	
23.	Transcripts	Were transcripts	No	-
	returned	returned to participants		
		for comment and/or		
		correction?		
Dom	ain 3: analysis and fi	ndings		
Data	analysis			
24.	Number of data	How many data coders	One (KG) but the coding	Methods (Data
	coders	coded the data?	manual was discussed	analysis)
			and agreed with two other	
			researchers (BA & Y)	

No	Item	Guide	Comments	Location in
		questions/description		manuscript
25.	Description of the	Did authors provide a	Yes in Supplementary	Supplementary
	coding tree	description of the coding	Table 1	Table 1
		tree?		
26.	Derivation of	Were themes identified	Derived from the data	Methods (Data
	themes	in advance or derived	(inductive analysis).	analysis)
		from the data?		
27.	Software	What software, if	QSR's NVivo 11 was	Methods (Data
		applicable, was used to	used.	analysis)
		manage the data?		
28.	Participant	Did participants provide	No, but the final	Methods (Data
	checking	feedback on the	interpretations were	analysis)
		findings?	reviewed and agreed with	
			two PPI representatives.	
Repo	rting			
29.	Quotations	Were participant	Participant quotations are	Results
	presented	quotations presented to	presented and each	(Qualitative
		illustrate the themes /	quotation is identified by	interviews)
		findings? Was each	a pseudonym and their	
		quotation identified? e.g.	gender, age and asthma	
		participant number	duration is noted.	
30.	Data and findings	Was there consistency	Yes	Results
	consistent	between the data		(Qualitative
		presented and the		interviews)
		findings?		

No	Item	Guide	Comments	Location in
		questions/description		manuscript
31.	Clarity of major	Were major themes	Yes	Results
	themes	clearly presented in the		(Qualitative
		findings?		interviews)
32.	Clarity of minor	Is there a description of	Diverse cases are	Results
	themes	diverse cases or	discussed.	(Qualitative
		discussion of minor		interviews);
		themes?	Coding tree presented in	Supplementary
			in Supplementary Table	Table 1
			1.	
1				1

No	Item	Description	Location in manuscript
Title	and abstract	I	
1.	Title	Concise description of the	Title identifies research as a mixed
		nature and topic of the study.	methods study, which includes
		Identifying the study as	qualitative research.
		qualitative or indicating the	
		approach (e.g., ethnography,	
		grounded theory) or data	
		collection	
		methods (e.g., interview, focus	
		group) is recommended.	
2.	Abstract	Summary of key elements of	Abstract formatted as per npj
		the study using the abstract	Primary Care Respiratory Medicine
		format of the intended	guidelines.
		publication; typically includes	
		background, purpose, methods,	
		results, and conclusions	
Intro	duction		
3.	Problem	Description and significance of	Given in introduction.
	formulation	the problem/phenomenon	
		studied;	
		review of relevant theory and	
		empirical work; problem	
		statement.	

Supplementary Table 3 SRQR checklist for qualitative interviews

No	Item	Description	Location in manuscript
4.	Purpose or research	Purpose of the study and	Aims given in last paragraph of
	question	specific objectives or questions	introduction.
Meth	ods		
5.	Qualitative	Qualitative approach (e.g.,	Mixed methods research; inductive
	approach and	ethnography,	thematic analysis. Approach
	research paradigm	grounded theory, case study,	detailed in the data analysis section.
		phenomenology, narrative	
		research) and guiding theory if	
		appropriate; identifying the	
		research paradigm (e.g., post-	
		positivist,	
		constructivist/interpretivist)	
		is also recommended;	
		rationale.	
6.	Researcher	Researchers' characteristics	Characteristics of the interviewer,
	characteristics and	that may influence	including credentials, relationship
	reflexivity	the research, including	with participants and involvement
		personal attributes,	in intervention development, given
		qualifications/experience,	in data collection section.
		relationship with participants,	
		assumptions, and/or	
		presuppositions; potential or	
		actual interaction between	
		researchers' characteristics and	
		the research questions,	

No	Item	Description	Location in manuscript
		approach, methods, results	
		and/or	
		transferability.	
7.	Context	Setting/site and salient	Feasibility trial participants
		contextual factors; rationale.	recruited from primary care.
			Interviews carried out by telephone.
			Detail given in participants and
			recruitment section.
8.	Sampling strategy	How and why research	Study was nested within a
		participants, documents, or	feasibility randomised controlled
		events were	trial study. All participants from
		selected; criteria for deciding	intervention arm were approached.
		when no further sampling was	Data saturation was considered
		necessary (e.g., sampling	reached because participants in
		saturation); rationale.	later interviews did not indicate any
			significant new benefits, concerns
			or barriers to engagement with My
			Breathing Matters. Detail given in
			participants and recruitment
			section.
9.	Ethical issues	Documentation of approval by	Ethical approval was granted and
	pertaining to	an	details are reported in the Design
	human subjects	appropriate ethics review	section.
		board and participant consent,	
		or explanation for lack thereof;	
		other	

No	Item	Description	Location in manuscript
		confidentiality and data	
		security issues.	
10.	Data collection	Types of data collected; details	All data collection methods,
	methods	of data collection procedures	including details of the interview
		including (as appropriate) start	and start and stop dates, given in
		and stop dates of data	the data collection section.
		collection and analysis,	
		iterative process,	
		triangulation of	
		sources/methods, and	
		modification of procedures in	
		response to evolving study	
		findings; rationale.	
11.	Data collection	Description of instruments	The interview schedules can be
	instruments and	(e.g.,	found in Supplementary Note 2.
	technologies	interview guides,	Interviews were audio-recorded
		questionnaires) and devices	(details in data collection section).
		(e.g., audio recorders) used for	
		data collection;	
		if/how the instrument(s)	
		changed over the course of the	
		study	
12.	Units of study	Number and relevant	Demographic information can be
		characteristics of participants,	found in Table 1.
		documents, or	

No	Item	Description	Location in manuscript
		events included in the study;	
		level of participation.	
13.	Data processing	Methods for processing data	Transcription, use of pseudonyms,
		prior to and during analysis,	and data handling approach is
		including	outlined in the data collection and
		transcription, data entry, data	analysis sections.
		management and security,	
		verification of data integrity,	
		data	
		coding and anonymization /	
		de-identification of excerpts	
14.	Data analysis	Process by which inferences,	Content and thematic analysis
		themes, etc. were identified	approaches are outlined in the data
		and developed,	analysis sections. Structure of codes
		including the researchers	and themes provided in
		involved in data analysis;	Supplementary Table 1.
		usually references a specific	
		paradigm or	
		approach; rationale.	
15.	Techniques to	Techniques to enhance	Techniques to enhance
	enhance	trustworthiness and	trustworthiness is outlined in the
	trustworthiness	credibility of data	data analysis and strengths and
		analysis,(e.g., member	limitations section.
		checking, triangulation, audit	
		trail); rationale	
Resu	lts/Findings	1	1

No	Item	Description	Location in manuscript
16.	Synthesis and	Main findings (e.g.,	A diagram of the main qualitative
	interpretation	interpretations, inferences, and	findings is presented in Figure 4
		themes); might include	and the findings are discussed in
		development of a theory or	relation to prior research in the
		model, or integration with	discussion.
		prior research or	
		theory	
17.	Links to empirical	Evidence (e.g., quotes, field	Anonymised quotes are provided
	data	notes, text excerpts,	throughout the results section to
		photographs)	support the qualitative themes.
		to substantiate analytic	
		findings.	
Disci	ussion		
18.	Integration with	Short summary of main	The discussion explains how the
	prior work,	findings, explanation of how	findings support and build on
	implications,	findings and conclusions	previous research and highlights the
	transferability, and	connect to,	unique contribution of this research.
	contribution(s) to	support, elaborate on, or	
	the	challenge conclusions of	
	field	earlier scholarship; discussion	
		of scope of	
		application/generalizability;	
		identification of unique	
		contribution(s) to scholarship	
		in a discipline	

No	Item	Description	Location in manuscript
-		or field.	
19.	Limitations:	Trustworthiness and	The strengths and limitations are
		limitations of findings	outlined in the discussion. Further
			details of the steps taken to increase
			the trustworthiness of the research
			is outlined in the data analysis
			section.
Othe	φ*		
20.	Conflicts of interest	Potential sources of influence	Competing interests are declared at
		or perceived influence on study	the end of the manuscript.
		conduct and conclusions; how	
		these were managed.	
21.	Funding	Sources of funding and other	Sources of funding are detailed in
		support; role of funders in data	the acknowledgments section.
		collection,	
		interpretation, and reporting	



Supplementary Figure 1: Screenshot of My Breathing Matters pages to engage people who

do not view themselves as having active asthma



Supplementary Figure 2: Screenshot of the My Breathing Matters breath check



Breathing Retraining Main Menu

More sessions will become available shortly after you complete each one. In the meantime, why not practice the sessions that are available to you below? The more you practice, the better you'll be!



If you'd like to reread the information about about the challenge (what it is, and how it works) then click here. . If you want, you can track your progress using a Breathing Retraining Progress Chart - find out more here.

Supplementary Figure 3: Screenshot of My Breathing Matters breathing retraining with

'unlocking' feature

Supplementary Note 1: My Breathing Matters Satisfaction Questionnaire

INSTRUCTIONS: Please only answer the below questions **if you registered with the My Breathing Matters website in the last 12 months**. Please tick **one** answer for each question.

1. Did you think there were any benefits of using My Breathing Matters?

No benefit at all	
Very little benefit	
Some benefit	
Quite a bit of benefit	
A large amount of benefit	

If any benefits, please note them down below:

2. Did you think there were any disadvantages of using My Breathing Matters?

No disadvantages at all	
Very little disadvantages	
Some disadvantages	
Quite a bit of disadvantages	
A large amount of disadvantages	

If any disadvantages, please note them down below:

3. How likely are you to recommend My Breathing Matters to friends and family if they needed similar care and treatment?

Extremely likely	
Likely	
Neither likely or unlikely	
Extremely unlikely	
Don't know	

Supplementary Note 2: Interview schedule

Interview questions for intervention participants who have logged on

- Q1. Can you tell me what it's like to have asthma?
- Q2. I'm really interested in hearing about your experiences of using My Breathing Matters, can you tell me all about it?
- Q3. Can you tell me about anything you liked about My Breathing Matters?
- Q4. Can you tell me about anything you disliked about My Breathing Matters?
- Q5. Can you tell me about any advantages of using My Breathing Matters for you?
- Q6. Can you tell me about any disadvantages of using My Breathing Matters for you?
- Q7. The research will continue for another 9 months. Do you think you will keep on using My Breathing Matters over this time? [Prompts: Why/why not?]
- Q8. Would you recommend My Breathing Matters to other people with asthma? [Prompts: Why/Why not?]
- Q9. Since using My Breathing Matters, how do you feel about your asthma now?
- Q10. Can you tell me about anything that you feel has changed from using My Breathing Matters?
 - a. Can you tell me about what changed? (e.g. anything different in your day-to-day life, the way you are managing your asthma?)
 - b. Can you tell me how you came to notice things changing?
 - c. Why do you think these things changed?
- Q11. When do you think My Breathing Matters would be most helpful to you?
- Q12. When do you think My Breathing Matters would not be helpful to you?

For each component:

- Q13. **[If didn't use]** Can you tell me why you decided not to use this part of My Breathing Matters?
- Q14. **[If used]** Can you tell me about how you found this section? *[Prompts: What did you like/dislike? Can you tell me about any problems you came across when doing the challenge?]*

Emails:

- Q15. Can you tell me about how you felt when My Breathing Matters sent you emails?
- Q16. Can you tell me what you thought about what the emails said?
- Q17. Can you tell me your thoughts about how often you received the emails?
- Q18. Can you tell me about any other ways you would like My Breathing Matters to contact you?
- Q19. Can you tell me about any advantages of getting these emails?
- Q20. Can you tell me about any disadvantages of getting these emails?

Interview questions for intervention participants who have not logged on

- Q1. Can you tell me what it's like to have asthma?
- Q2. We are interested to hear from people who did not use My Breathing Matters, can you tell me why you have not used My Breathing Matters?
- Q3. What are your thoughts on using a website to help you to manage your asthma?
- Q4. The research will continue for another 9 months. Do you think you will use My Breathing Matters over this time? [Prompts: Why/why not?]