Interventions to enhance the adoption of asthma self-management behaviour in the South Asian and African American population

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Interventions to enhance the adoption of asthma self-management behaviour in the South Asian and African American population: a systematic review

Salina Ahmed, Liz Steed, Katherine Harris, Stephanie J. C. Taylor, and Hilary Pinnock

South Asian and other minority communities suffer poorer asthma outcomes, have a higher rate of unscheduled care and benefit less from most existing self-management interventions when compared to the majority population. Possible reasons for these differences include failure to implement asthma self-management strategies, or that strategies implemented were inappropriate for their needs; alternatively, they may relate to the minority and/or lower socioeconomic status of these populations. We aimed to synthesise evidence from randomised controlled trials for asthma self-management in South Asian and Black populations from different sociocultural contexts, and identify barriers and facilitators to implementing self-management. We systematically searched eight electronic databases, and research registers, and manually searched relevant journals and reference lists of reviews. Seventeen trials met the inclusion criteria and were analysed narratively. We found two culturally targeted interventions compared to fifteen culturally modified interventions. Interventions used diverse self-management strategies; education formed a central component. Interventions in South Asian and African-American minority communities were less effective than interventions delivered in indigenous populations in South Asia, though the latter trials were at higher risk of bias. Education, with continuous professional support, was common to most interventions. Facilitators to asthma self-management included: ensuring culturally/linguistically appropriate education, adapting to learning styles, addressing daily stressors/social support and generic self-management strategies. In conclusion, when developing and evaluating self-management interventions aimed at different cultures, the influence of sociocultural contexts (including whether patients are from a minority or indigenous population) can be important for the conceptualisation of culture and customisation of self-management strategies.

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INTRODUCTION

South Asian communities, along with other minority populations, have poorer asthma outcomes, higher rates of hospital admission, greater risk of rehospitalisation and a higher death rate compared to majority white populations. Asthma self-management, consisting of education, written Personalised Asthma Action Plans (PAAPs) and regular reviews (supported self-management) is known to improve health outcomes, and is recommended in national and international guidelines. Despite hopes that self-management offers a potential solution to address preventable health inequalities, there are concerns that asthma self-management interventions have produced little or no positive improvements on health outcomes for South Asians or other minority populations, further widening the gap of asthma inequalities. Possible explanations for these variations include differences in health-seeking behaviours related to health beliefs and attitudes to mainstream medicine, environmental or lifestyle factors, poor healthcare access and the quality of asthma care provided to these communities. These factors may be driven by cultural diversity, by the experience of being a minority and/or by socioeconomic status (SES). Thus, the way in which self-management is accessed and delivered to these various populations, need to be explored, and self-management strategies may need to be developed for the target population’s culture, ethnicity, SES or other needs.

There are distinctions between the way interventions can be made relevant to a population (see Table 1). ‘Culturally modified/adapted’ interventions, are developed for a majority population and then modified for use in other ethnic groups; the core content, however, is the same. ‘Culturally targeted’ interventions are developed from a bottom-up process that considers the shared characteristics and context of a cultural group before developing an intervention. Finally, bottom-up interventions that assess and are aimed at the unique cultural characteristics and dimensions of individuals within a cultural group, with individualised intervention delivery are known as ‘culturally tailored’. Culturally targeted or tailored interventions are generally suggested to be more effective than culturally modified interventions, though the evidence for this has focussed mainly on children. Studies and clinical practice guidelines often indiscriminately apply findings from a majority population in a South Asian country, as relevant and applicable to South Asian minorities and majorities in other countries, despite differences in time and space of lived experiences and cultural shifts. Not only are the South Asian and Black population heterogeneous groups, but culture is

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Table 1. Definition of terms

| Term | Definition | Examples |
|------|------------|----------|
| Culturally modified/adapted interventions | Pre-existing generic interventions modified for the intention of being relevant to ethnic groups using various strategies, though the content is primarily the same | Language translation, and use of images and bilingual educators from a similar ethnicity as the target population |
| Culturally targeted interventions | A bottom-up process which considers the shared characteristics and dimensions of collective individuals of a culture before developing an intervention, aimed at a group level | Religion |
| Culturally tailored interventions | A bottom-up process which considers the unique cultural characteristics and dimensions of individuals within a cultural group before developing an intervention, aimed at individuals within a group | Level of religious identification or spirituality |
| ‘Majority’ South Asians | Interventions from South Asian countries where the population forms a majority | South Asians in India |
| ‘Minority’ South Asians; ‘Minority’ African Americans | Interventions from countries where the population forms a minority | South Asians in the UK or Canada; African Americans in the USA |

RESULTS

Characteristics of included trials

From a total of 3174 citations, we included 17 papers (reporting 16 trials) (see Fig. 1). The randomised control trials (RCTs) were conducted between 1995 and 2016; four South Asian trials were from India (labelled ‘majority’ South Asian),30–33 four South Asian trials were from the UK34–36 and one from Canada,37 (labelled ‘minority’ South Asian), and nine African-American trials were from the USA (labelled ‘minority’ African American)26–34 (see Table 2). The overall risk of bias within trials was uncertain,30,33,37–41 or high,31,32,36,42–45 Three trials had low risk,34,35,46 (see Table 3).

Participant characteristics: The ‘majority’ population in the South Asian trials comprised of Indians30–33 whereas ‘minority’ South Asian trials included Indians37 and mixed subcultures (e.g., Bangladeshi, Pakistani, Indian or Sri Lankan).34–36 All Black population trials studied the African-American minority population in the USA.38–46 Most trials (fourteen studies) did not define ethnicity; only three ‘minority’ South Asian trials defined ethnicity according to self-identification or language spoken.34,35,37 All trials aimed interventions at asthma patients (whether this was children, adolescents, adults or elders).30–46 In addition, some trials also targeted parents,30,32,38,46 trained African-American coaches and/or residents,38,46 or healthcare professionals (clinicians and nurses).30,32,34–36

Study setting: All ‘majority’ South Asian trials were based in tertiary care hospitals.30–33 In contrast, ‘minority’ South Asian trials were conducted in primary care,35,36 or a combination of community, primary care and hospital (secondary/tertiary) settings.34,37 Similarly, the African-American trials were conducted in various settings: primary or secondary schools,30,32,46 tertiary care hospitals,38–41 emergency department38 and three trials used a combination of settings; community, school and hospital (secondary/tertiary).38,44,46

Geographical area and socioeconomic status: Among the ‘minority’ trials that specified the demographic location of patients, these were described as urban in six trials,34–36,40,41,46 and one African-American trial was conducted in mixed urban and rural areas.43 Eight trials were described as from economically deprived or low-income areas,34,35,38–41,45,46 and two ‘minority’ trials (South Asian and African American) were conducted in low-middle class areas.36,44

Intervention characteristics: Table 2 describes intervention characteristics. All interventions included patient education, though the approach, method of delivery and content varied. Examples included education-sessions or classes,30,32,33,35,36,38–46 training for patients,30,32,33,35,36,45,46 and healthcare professionals, coaches or residents,30,32,34–36,46 education in written,31–33,35,39,44,46 or video format,35,37,42 education in the form of social support,46 or a local education/promotional campaign.38 Twelve out of 17 interventions were delivered by healthcare professionals,30,32,34–36,48 five of whom were specifically trained for the project.30,32,35,42,43 Three interventions from minority countries were delivered in South Asian languages by healthcare professionals or research facilitators,35–37 two ‘majority’ South Asian trials had written materials in Hindi or Tamil30,33 and two USA interventions were delivered by trained African American lay people or university staff who were residents in the community.18,46 Intervention duration ranged from 40 minutes to 1 year and follow-up lengths ranged from 1 month to 3 years (see Table 3 for details on the latter).

Strategies for reinforcing knowledge or self-management behaviours included follow-up classes,30,45 nurse
Eligibility

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for the control groups varied, 30 practice technique,43 asthma diary/workbook,30,32,33,42 peak intervention strategies were based on speci

Intercollegiate Guideline Network (SIGN).33 Institute, Global Initiative for Asthma (GINA) and Scottish National Institutes of Health, National Heart Lung and Blood

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evaluation culturally targeted interventions.31,37 Behera et al. 31 culturally tailored interventions, and only two of seventeen trials

monitoring,30,34,36,37,39,41,42,44,45 medication counselling 33 and
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other intervention characteristics used alongside education,30,32,34,36,39,42,44,45 placebo inhalers to

nation of both), that incorporated cultural beliefs and attitudes,

scienti
educational videos included three intervention possibilities (i.e.,

Fig. 1 PRISMA flow diagram

records identified through database searching

(n = 4772)

Additional records identified through other sources

(n = 231)

Records after duplicates removed

(n = 3174)

Records screened

(n = 3174)

Records excluded

(n = 3064)

Full-text articles assessed for eligibility

(n = 110)

Studies included in quantitative synthesis (narrative analysis)

(n = 17 papers (reporting 16 trials))

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(n = 17 papers (reporting 16 trials))

clinics34,39,44,45 and written materials.42,43 Most trials described other intervention characteristics used alongside education,30,32–46 including the use of written PAAPs in all South Asian trials (majority and minority), and some African-American trials,44,46 provision of emergency oral corticosteroid courses,34 asthma medication/therapy,30,34,36,39,42,44,45 placebo inhalers to practice technique,43 asthma diary/workbook,30,32,33,42 peak flow monitoring,30,34,36,37,39,41,42,44,45 medication counselling33 and access to free asthma organisation helplines.42 In seven trials, intervention strategies were based on specific guidelines, e.g., National Institutes of Health, National Heart Lung and Blood Institute, Global Initiative for Asthma (GINA) and Scottish Intercollegiate Guideline Network (SIGN)33–36,39,44,46 Usual care for the control groups varied30–36,39–44,46 including illustrative leaflets,37 routine education classes,45 and recruiting similar neighbourhood areas to the intervention sites.38

Fig. 1 PRISMA flow diagram

(1) Features of culturally relevant interventions. In line with our definition and that in previous literature,14,15 we did not find any culturally tailored interventions, and only two of seventeen trials evaluated culturally targeted interventions.31,37 Behera et al.31 (‘majority’ South Asian trial at high risk of bias) provided a targeted written self-care booklet in Hindi (including a PAAP) developed collaboratively from patient knowledge, relevant literature and expert advice. Pourselami et al.37 (‘minority’ South Asian trial at unclear risk of bias) developed educational videos in collaboration with community members and healthcare professionals. The educational videos included three intervention possibilities (i.e., scientific knowledge, community opinions/narratives or a combination of both), that incorporated cultural beliefs and attitudes, e.g., cultural gestures, humour, storytelling and social interaction styles appropriate for Punjabi Indians. The aim was to facilitate patients’ trust in the community member and/or clinician who delivered the intervention. Both interventions were piloted in focus groups to improve clarity, relevance and acceptability and were refined before evaluation. These trials were not classified as culturally tailored because they were delivered to the specified cultural group without distinguishing or measuring individual cultural differences within that group.31,37

Both trials significantly improved knowledge. Pourselami et al.37 improved adherence to physician instructions on medication and inhaler use, and Behera et al.31 reported reduced symptoms, hospital admissions and use of breathing exercises during acute attacks. Although, the former trial achieved significant findings on all outcomes for Punjabi Indians, the Chinese population (who were studied as a parallel group with their own culturally targeted intervention) performed even better. The authors considered that this may be related to participant demographics; the Punjabi Indians were older and less educated than the Chinese community.37

In contrast, 15 out of 17 interventions were found to be culturally modified30,32–36,38–46 They used strategies such as adapting existing interventions or materials for the target ethnic group,32,35,39,44 e.g., an African-American training video was re-recorded with South Asian actors.35 and ethnically relevant images were used such as African-American celebrities.34,35,42 Other studies applied interventions to several ethnic groups without considering cultural differences; thus, providing written or oral education (e.g., classes, PAAPs and workbooks) translated from English to the target participant language or using bilingual educators, without adjusting intervention content.33–38 However,
| Study, Country | Population characteristics | Intervention characteristics |
|----------------|-----------------------------|-----------------------------|
| **Majority’ South Asian trials** | | |
| Agrawal30 India | Evaluated efficacy of PAAPs for asthma control | Indian; Patients; Parents; 2–12; 60 (32/28) Tertiary (university clinic) | Education; sessions, training including on asthma symptom diary and peak flow measurements No PAAP, standard asthma therapy and education Trained physician; social scientist (-) Individual; Written material Modified |
| Behera31 India | Assessed patient knowledge of self-care needs and develop/evaluate a self-care manual | Indian Tertiary (university clinic) | Education-booklet in Hindi (included a PAAP) Booklet evaluation/not stated No specific instructions/pilot study used to develop booklet in Hindi (n = 45) Not stated (Hindi) Written material Targeted |
| Ghosh32 India | Assessed the impact of self-management education and training on health status and resource use | Indian Tertiary (university clinic) | Education; sessions, training, written instructions, audio-visual aids, role models, group/scenario discussions Regular care e.g. drug administration Trained social scientist (-) Group; Modified |
| | Patients: Parents; 10–45; 276 (140/136) | | Daily diary (included symptom assessment and financial workbook) Asthma therapy PAAPs/four 2 hour sessions Written material |
| Shanmugam33 India | Provided pharmaceutical care through partnership of pharmacists and patients for good asthma control | Indian Tertiary (university hospital) | Education; sessions, asthma care diary in English and Tamil (including leaflet), PAAP and symptom log sheet No pharmaceutical care Not stated (English and Tamil) Written material; Modified |
| | Patients Age; –; 66 (33/33) | | Medication counselling/not stated Other methods not stated |
| **Minority’ South Asian trials** | | |
| Griffiths34 UK | Tested whether specialist nurses across ethnically diverse and deprived areas reduce unscheduled care | South Asians (mostly Bangladeshi White Caucasians, Other (Black/African Caribbean/Other) Primary/secondary (out-of-hours GP service/hospital) | Education; training based on guidelines, nurse review with advice PAAP explained in English and Sylheti Usual care; single nurse visits to discuss asthma guidelines and check inhaler technique Trained nurse specialists (partially; PAAPs explained in Sylheti) Individual; Modified |
| | | | Written material; |
| Study, Country | Population characteristics | Intervention characteristics |
|---------------|--------------------------|-----------------------------|
| **Aim** | | Ongoing clinical support for professionals on computer prompts |
| | | Peak flow meters provided |
| | | Oral corticosteroids/2 one hour visits for GP practices; 194 days |
| | Patients; 4-60; 164 (95/69) | Deprived/urban |
| **Griffiths** UK | Tested whether culturally specific education programmes adapted from USA interventions reduce unscheduled care | Primary (GP) |
| | South Asians (Bangladeshi, Pakistani, Indian, Sri Lankan) | Patient; Primary/secondary care clinicians; 3 and above; 375 (183/192) |
| | Deprived/urban | Usual care; nurse follow-ups to book appointments (CDSMP), research training with video based on guidelines; South Asian actors and manualised programme (PACE)/PACE; two seminars; CDSMP; 2-hour session |
| | | Trained GP (South Asian) |
| **Moudgil** UK | Tested whether bilingual education of treatment optimisation and follow-up reduce urgent healthcare and improve quality of life | Primary (GP) |
| | South Asian (mainly Indian and Pakistani), White European | Patients; GP; 11-59; 344 (171/173) |
| | Low or medium deprivation/urban | Booklet including PAAP (based on BTS guidelines) and peak flow measurements |
| | | GP trained on prescribing, optimal treatment, knowledge and medication |
| | | Peak flow meter provided |
| | | Asthma therapy/40 minutes |
| | | Written material |
| **Poureslami** Canada | Explored the effectiveness of different culturally relevant information | South Asians (Indian Punjabi), Chinese Other/tertiary (home, university clinic)- |
| | Education; videos (physician-led, community and physician-led/ | Pictorial pamphlet in either Mandarin, Cantonese or Punjabi |
| | | Research facilitators (South Asian) |
| | | Group/video |
| | | Targeted |
| Study, Country | Population characteristics | Intervention characteristics |
|----------------|-----------------------------|-----------------------------|
| **Aim** | | formats and impact on self-management |
| Study setting; SES/area | | community combination) |
| Intervention description/length | | Peak flow meter |
| Control /other group | | PAAPs/1 month |
| Delivery (ethnicity; language) | | /Co-development of intervention (n = 35); focus group sessions (n = 40) |
| Mode of delivery | | Video/DVD; |
| Modified; Targeted; Tailored | | Written material |

**'Minority' African American trials**

**Blixen**

USA: Tested feasibility of a culturally appropriate in-patient education programme for hospitalisation

- **African Americans**
- **Tertiary** (hospital)
- **Patients**: 21 and above; 45 (33/12)

- **Intervention**: Education; sessions and video, asthma workbook using African-American images, references to famous celebrities, written education posted as follow-up
- **Control/other group**: Received usual care
- **Delivery**: Trained nurse (Not stated)
- **Mode of delivery**: Individual; Modified

**Fisher**

USA: Tested community-based intervention to improve asthma awareness, attitudes, management practices and reduce acute care rehospitalisation

- **African Americans, White Caucasians, Others**
- **Other (community, school)**
- **Patients**: 8–50; 28 (14/14)

- **Intervention**: Education; promotion campaigns, sessions, training residents to support patients in school and community/12 months
- **Control/other group**: Four areas in the same location with similar SES characteristics
- **Delivery**: Trained university staff/residents (African American)
- **Mode of delivery**: Group; Modified

**Fisher**

USA: Tested whether community health workers can reach low-income parents of hospitalised children and to reduce rehospitalisation

- **African American**
- **Other/secondary (community, hospital)**
- **Low income/ urban**
- **Patients**: 18–70; 241 (119/122)

- **Intervention**: Education; sessions by asthma coach based on guidelines and parental support contacts/meetings for readiness to change, training for asthma coaches (including PAAPs)/2 years
- **Control/other group**: Usual care; inpatient education and discharge planning with PAAP, a suggested follow-up primary care within 1 week of discharge
- **Delivery**: Nurse, Individual; Modified

**Ford**

USA: Reanalysed an education programme that assessed the effects on asthma outcomes

- **African Americans**
- **Secondary (emergency department)**
- **Urban and rural**

- **Intervention**: Education; sessions and follow-ups, handout, mailed sessions for non-attenders
- **Control/other group**: Received no intervention
- **Delivery**: Trained healthcare professionals and nurses (not stated)
- **Mode of delivery**: Group; Modified

**Interventions to enhance the adoption of asthma self-management**

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| Study, Country | Population characteristics | Intervention characteristics |
|----------------|---------------------------|----------------------------|
| **Aim**        |                           | Intervention description/length |
| **Ethnicity; Participants; Sample age; Sample size (I/C)** |                           | Control /other group descriptions |
| **Study setting; SES/area** |                           | Delivery (ethnicity; language) |
| **Mode of delivery** |                           | Modified; Targeted; Tailored |
| **Keslo**<sup>39</sup> USA | Provided major long-term therapeutic intervention and intensive education | Wallet sized card (with medication list, dose, frequency); Placebo inhaler to practice/3 sessions |
| African Americans | Secondary /tertiary (emergency department/ university clinic) | Education; sessions based on NIH guidelines, Follow-up clinics |
| Patients: 18 and above; 52 (30/22) | Low; deprived | Usual care |
| **Keslo**<sup>44</sup> USA | Tested if a long-term management programme (emphasising ICS and patient education), would improve outcomes | Education booklet (including diary card for measurements and 1-page summary of asthma prevention, medications, triggers and peak flow meter product literature) |
| African Americans | Tertiary (university based clinic) | Asthma therapy for ICS Peak flow meter (colour-coded stickers), inhaled b-agonist and aero chamber provided/1-hour session |
| Patients: 18 and above; 39 (21/18) | Low; working and middle-class college students | Follow-up clinics (including diary)/2 years |
| **Velsor-Friedrich**<sup>40</sup> USA | Tested the effect of a school-based education programme (Open Airways) on the psychosocial outcomes | Educational booklet Written instructions on asthma crisis management |
| African Americans | Other (Eight public primary school) | Asthma therapy and peak flow meter (colour-coded stickers), MDI and other medications Follow-up clinics (including diary)/2 years |
| Patients: 8-13; 102 (40/62) | | Education; sessions/2 weeks, six 45 minute sessions per week |

*Interventions to enhance the adoption of asthma therapy for ICS*
| Study, Country | Population characteristics | Intervention characteristics |
|----------------|-----------------------------|-----------------------------|
|                | Aim                         | Intervention description/length | Control /other group descriptions | Delivery (ethnicity; language) | Mode of delivery | Modified; Targeted; Tailored |
|                | Ethnicity; Participants; Sample age; Sample size (I/ C) | Study setting; SES/area | Education/sessions (as above) | Patient education | Academic professor, academic nurse (-) | Individual; Written material | Modified |
| Velsor-Friedrich | An extension of the study above (Velsor-Friedrich 2004): tested a two-part school-based education programme | African Americans; Other (eight public primary schools with nurse clinics) | Low/Urban | As above and all students received a PAAP | | |
| USA | Patients; 8-13; 52 (28/24) | | | | | |
| | An extension of the study above (Velsor-Friedrich 2004): tested a two-part school-based education programme | Education/sessions, coping skills training including role-playing & technology use (with a booster session as follow-up) | Routine education | Clinician, Individual; Group | | Modified |
| Velsor-Friedrich | Evaluated efficacy of a school-based asthma education program on psychosocial & health outcomes | African Americans; Other (5 secondary schools) | Low | Nurse practitioner reinforcement & clinic visit | | |
| USA | Patients; 13–19; 137 (74/63) | | | Provided MDI, hydrofluoroalkane & static free chamber | | |
| | | | | Peak flow diary | | |
| | | | | PAAP/Six 45 minute sessions over 6 weeks | | |

Note: Missing data obtained from authors is noted in italic in the table
| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher’s interpretation for the harvest plot |
|---------------------------------------------------------|------------------------|------------------------------------------------------------------------------------------------|-----------------------------------------------|
| Agrawal30 n = 60 children FU: 4 m | Clinical-unscheduled care, 4 m | Compared to controls, children in the intervention group had: | Illustrated as a consistent significant positive effect |
| Overall risk of bias: Unclear | Clinical-asthma control, 4 m | Fewer acute asthma events: I: 0.50 (SD 0.71) vs. 1.0 (SD 0.61); p = 0.02 | Illustrated as a consistent significant positive effect |
| | | Compared to controls, children in the intervention group had: | |
| | | Improved symptom score (from the symptom diary): I: 21.9 (SD 14.4) vs. C: 33.7 (SD 10.9); p = 0.0006 | |
| | | Fewer nocturnal awakenings: I: 1.75 nights/month (SD 1.30) vs. C: 3.25 (SD 1.201); p = 0.001 | |
| | | Reduced school absenteeism: I: 1.5 days/month (SD 1.4) vs. C: 2.54 (SD 1.79); p = 0.015 | |
| Behera31 CCT n = 523 adults | Clinical-unscheduled care, 1 yr | A reduction in hospital admissions is illustrated graphically (the authors state that there was a significant decrease in hospital admissions in the intervention group at FU compared to the control group) | Illustrated as a consistent significant positive effect |
| FU: 2 wks, 6 m, 1 yr Overall risk of bias: High | Clinical-asthma control, 2 wks, 6 m, 1 yr | Symptom scores decreased in both groups | Illustrated as a consistent significant positive effect |
| | | I: Baseline: 18.14 (SD 41.23) vs. FU 1 yr: 12.61 (SD 28.66) | |
| | | C: Baseline: 18.76 (SD 42.64) vs. FU 1 yr: 10.69 (SD 24.30) | |
| | | Logistic regression: compared to the control group, more intervention group patients showed a significant improvement in symptom scores at 2 w, 6 m and 1 yr (p < 0.001) | |
| | Process, 2 wks, 6 m, 1 yr | Knowledge scores increased significantly in the intervention group and fell in the control group; | Illustrated as a consistent significant positive effect |
| | | I: Baseline: 13.04 (SD 4.06) vs. FU 1 yr: 28.13 (SD 15.70); p = < 0.001 | |
| | | C: Baseline: 11.44 (SD 4.0) vs. FU 1 yr: 9.47 (SD 2.89); p = < 0.001 | |
| | | Logistic regression: Compared to the control group, more intervention group patients showed a significant increase in knowledge scores at 2 wks, 6 m and 1 yr (p < 0.001) | |
| | Behavioural, 2 wks, 6 m, 1 yr | Reported self-care in acute attacks showed no change in attitudes in either group, but significantly more patients in the intervention group adopted the recommended position (sitting, leaning forward) and practiced breathing exercises during an acute attack as compared to control patients | Illustrated as a significant positive effect but hatched to show inconsistency |
| Ghosh32 n = 276 adult, adolescent, children/parent | Clinical-unscheduled care, 1 yr (assessed by diary in months 4, 8 and 12) | Fewer total number of ED visits, but no between group difference in proportion with ED visit | Illustrated as positive but hatched to indicate inconsistency |
| | | Number of ED visits in the 3-month diary: I: 11.6 (SD 16.2) vs. C: 21.8 (SD 25.0); p = 0.002 | |
| | | Proportion with ED visits in the 3-month diary: I: 42.9% vs. 50.0% (p = 0.117) | |
| | | Number and duration of hospitalisations were both significantly reduced | |
| | | Hospital days in the three diary months: I: 5.8 (SD 10.7) vs. 12.5 (SD 19.8); p = 0.016 | |
| | | Proportion hospitalised in the three diary months: I: 27.1% vs. C: 36.8%; p = 0.043 | |
| FU: 4 m, 8 m, 1 yr Overall risk of bias: High | | Fewer productive days lost in the intervention group during the three diary months | Illustrated as a consistent significant positive effect |
| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher’s interpretation for the harvest plot |
|---------------------------------------------------------|------------------------|-------------------------------------------------------------------------------------------------|-----------------------------------------------|
| Clinical-asthma control, 1 yr (assessed by diary in months 4, 8 and 12) | | | |
| Process | Not assessed | Not assessed | |
| Clinical-unscheduled care | | | |
| FU: 29 days Overall risk of bias: unclear | | | |
| Process | Not assessed | Not assessed | |
| Griffiths 34 n = 44 practices/324 – (South Asians I: 95 C: 69 n = 164), adults, adolescents, children | Clinical-unscheduled care, 1 yr | Asthma control improved in the intervention group compared to the control group | Illustrated as a consistent significant positive effect |
| | | Mean ACT score for each question was greater in the intervention group at FU: p < 0.05 | |
| | | (Overall mean ACT scores are not reported) | |
| | | Lung function showed a greater increase in the intervention group compared with control | |
| | | PEFR (L/min): Baseline: I: 282 (SD 95) vs. C: 265 (SD 93); FU: I: 336 (SD 88) vs. C: 268 (SD 85); p < 0.05 | |
| | | Process | Not assessed | |
| | | Behavioural | Not assessed | |
| FU: 2 m, 9 m, 1 yr Overall risk of bias: low | Clinical-asthma control, 2 m, 1 yr | Symptoms: no between group differences in whole population. No data for South Asian sub-group, but authors state that ‘intervention effect was not significant for other sub-group analysis’ | |
| | | Process | Not assessed | |
| | | Behavioural, 2 m, 1 yr | Not assessed | |
| Griffiths 35 n = 84 practices/375 elders, adults, adolescents, children, primary and secondary care clinicians | Clinical-unscheduled care: 171 days/ I: 189 days; 72 days/ C: 339 days 1 yr Unscheduled care: there was no between group difference in healthcare use | Illustrated as a consistent no effect | |
| | | *Time to first unscheduled contact FU: HR = 1.19 (0.92 to 1.53); p = 0.185 | |
| | | Proportion without unscheduled care FU: OR = 0.72 (0.45 to 1.16); p = 0.175 | |
| | | Time to first unscheduled primary care contact FU: HR = 1.20, 0.92 to 1.57 p = 0.177 | |
| | | Time to first routine review in primary care FU: HR = 2.22, 1.67 to 2.95 p < 0.001 | |

* indicates the primary outcome (if stated)
| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher's interpretation for the harvest plot |
|----------------------------------------------------------|------------------------|-------------------------------------------------------------------------------------------------|-----------------------------------------------|
| FU: 3 m, 1 yr Overall risk of bias: low                  | Clinical-asthma control, 3 m, 1 yr Process, 3 m, 1 yr                                      | Corticosteroid prescriptions: There was no between group difference in steroid prescriptions Steroids FU: I: 1.16 vs. 0.98 Adjusted incidence rate ratio: 1.14 (0.87–1.49) | Illustrated as a consistent no effect |
| Moudgil36 n = 689 (White Europeans 345, Indian subcontinent 344); adults, adolescents, children | Asthma control: there was no between group difference in symptom score                       |                                           |                                |
|                                                          |                                      | Researcher's interpretation for the harvest plot                                               |                                |
|                                                          |                                      | Symptom score FU 1 yr: 9.9 (SD 5.0) vs. C: 10.1 (SD 4.2) AHR: −0.04 (−1.16 to 1.09); p = 0.949 |                                |
|                                                          |                                      | Self-efficacy was improved at 3 m but not at 1 yr follow-up; At 3 months: I: 6.7 (2.1) vs. C: 6.3 (1.9) AHR: 0.44 (0.05 to 0.82); p = 0.027 At 12 months: I: 6.4 (1.8) vs. C: 6.3 (1.6) AHR: 0.25 (−0.13 to 0.63); p = 0.188 |                                |
|                                                          |                                      | Not assessed                                                                                    |                                |
|                                                          |                                      | [Note: these data are an a priori sub-group analysis]                                            |                                |
|                                                          |                                      | Number of asthma events/episodes for South Asians: no between group differences                |                                |
|                                                          |                                      | *Proportion with an admission. I: 5.3 vs. C: 6.3% OR 0.83 (0.28 to 2.44); p = 0.9081          |                                |
|                                                          |                                      | Proportion with an A&E attendance. I: 1.4 vs. C: 4.0% OR 2.92 (0.52 to 21.2); p = 0.3184     |                                |
|                                                          |                                      | Proportion with out-of-hours primary care. I: 2.8 vs. C: 2.6% FU: OR 0.95 (0.19 to 4.60); p = 1 |                                |
|                                                          |                                      | Proportion with a GP consultation. I: 55.9 vs. 50.3% OR 0.80 (0.49 to −1.30); p = 0.3971    |                                |
|                                                          |                                      | Proportion with a steroid course. I: 20.3% vs. 19.9% OR 0.97 (0.53 to 1.79); p = 1           |                                |
| FU: 4 m, 8 m, 1 yr Overall risk of bias: High            | Clinical-asthma control, 1 yr ISC n = 280                                                   | Quality of life in South Asians was significantly better in the intervention group             | Illustrated as a consistent significant positive effect |
|                                                          |                                      | Change in AQLQ FU: I: 0.11 vs. −0.15. Between group mean difference 0.26 (0.17–0.36); p = < 0.001 |                                |
|                                                          |                                      | Not assessed                                                                                    |                                |
|                                                          |                                      | [Note: these data are an a priori sub-group analysis]                                            |                                |
|                                                          |                                      | Number of asthma events/episodes for South Asians: no between group differences                |                                |
|                                                          |                                      | *Proportion with an admission. I: 5.3 vs. C: 6.3% OR 0.83 (0.28 to 2.44); p = 0.9081          |                                |
|                                                          |                                      | Proportion with a GP consultation. I: 55.9 vs. 50.3% OR 0.80 (0.49 to −1.30); p = 0.3971    |                                |
|                                                          |                                      | Proportion with an A&E attendance. I: 1.4 vs. C: 4.0% OR 2.92 (0.52 to 21.2); p = 0.3184     |                                |
|                                                          |                                      | Proportion with out-of-hours primary care. I: 2.8 vs. C: 2.6% FU: OR 0.95 (0.19 to 4.60); p = 1 |                                |
|                                                          |                                      | Proportion with a steroid course. I: 20.3% vs. 19.9% OR 0.97 (0.53 to 1.79); p = 1           |                                |
|                                                          |                                      | Insufficient data                                                                               |                                |
| Poursalmi37 n = 92 (47 Chinese, 45 Punjabi); Adults      | Process, 3 m, 6 m Punjabi n = 43                                                           | Knowledge: no comparison data for intervention and control groups                              | Insufficient data |
| FU: 3 m, 6 m; 1 telephone survey interview Overall risk of bias: unclear | Behavioural, 3 m, 6 m Punjabi n = 43                                                       |                                                        |                                |
|                                                          | Understanding physician instructions; on medication and proper inhaler use skills: no comparison data for intervention and control groups |                                                        |                                |
| Blixen42 n = 28, Adults                                   | Clinical-unscheduled care, 3 m, 6 m                                                        | Healthcare use: no data provided, though stated as no significant between group differences    |                                |
|                                                          | Clinical-asthma control, 3 m, 6 m                                                        | Quality of life: There was no significant between group differences                            |                                |
|                                                          | Overall AQOL score. FU 6 m: I: 4.59 (SD 1.48) vs. C: 4.43 (SD 1.52); p = 0.12                |                                                        |                                |
|                                                          | Process                                                                                  |                                                        |                                |
|                                                          | Not assessed                                                                               |                                                        |                                |
|                                                          | [Note: these data are an a priori sub-group analysis]                                      |                                                        |                                |
|                                                          | Number of asthma events/episodes for South Asians: no between group differences            |                                                        |                                |
|                                                          | Proportion with a GP consultation. I: 55.9 vs. 50.3% OR 0.80 (0.49 to −1.30); p = 0.3971    |                                                        |                                |
|                                                          |Proportion with an A&E attendance. I: 1.4 vs. C: 4.0% OR 2.92 (0.52 to 21.2); p = 0.3184     |                                                        |                                |
|                                                          | Proportion with out-of-hours primary care. I: 2.8 vs. C: 2.6% FU: OR 0.95 (0.19 to 4.60); p = 1 |                                                        |                                |
|                                                          |Proportion with a steroid course. I: 20.3% vs. 19.9% OR 0.97 (0.53 to 1.79); p = 1           |                                                        |                                |
Table 3 continued

| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher’s interpretation for the harvest plot |
|----------------------------------------------------------|------------------------|--------------------------------------------------------------------------------------------------|-------------------------------------------------|
| Fisher³⁸ \(n = 249\) Adolescents, children, parents     | Behavioural, 3 m, 6 m   | Self-management behaviours: no data, though stated as no-significant between group differences     | Illustrated as a consistent no effect            |
|                                                          | Clinical-unscheduled care, quarterly for 3 yrs | *Acute care: no data given (results illustrated graphically), though authors stated no significant between group differences in acute care (hospitalisations and ED attendances \(p = 0.35\)) | Illustrated as a consistent no effect            |
|                                                          | Clinical-asthma control Process | Not assessed                                                                                     | –                                               |
|                                                          | Behavioural, every quarterly until 3 yrs | *Asthma management: no significant between group differences in the non-validated assessment of parent’s reported attitude about asthma and asthma management Attitudes about asthma FU: I: 2.34 vs. C: 2.24 (\(p = 0.35\)) Appropriate thresholds for seeking help Baseline: I: 30 vs. C: 47%; FU: I: 51 vs. C: 53% \(p = 0.77\) | Illustrated as a consistent no effect            |
| FU: 3, 6, 9, 12, 16, 20, 24, 28, 32, 36 m Overall risk of bias: unclear |                              |                                                                                                  |                                                  |
| Fisher³⁹ \(n = 191\) parents, coaches                   | Clinical-unscheduled care, 1 yr, 2 yr | *Hospitalisation Compared to controls, the intervention group had fewer hospitalisations; Hospitalised at least once FU I: \(n = 35/96\) (36.5%), \(55\) vs. C: \(55/93\) (59.1%); 95% CI (0.11–0.34); \(p = .002\) | Illustrated as a consistent significant positive effect |
| FU: 6, 12, 18, 24 m Overall risk of bias: low           | Clinical-asthma control Process | Not assessed                                                                                     | –                                               |
|                                                          | Behavioural              | Not assessed                                                                                     | –                                               |
| Ford⁴³ \(n = 241\) (African American = 163, Caucasian = 78) | Clinical-unscheduled care, 4 m, 8 m, 1 yr | *ED visits No impact [Note: these data are an \(a\ priori\) sub-group analysis] ED visits/year I: Baseline: 5.0 (SD 3.6) vs. FU 2.7 (SD 3.3); C: Baseline: 6.7 (SD 8.4) vs. FU: 4.8 (SD 6.8) No between group comparisons reported | Illustrated as a consistent no effect            |
|                                                          | Clinical-asthma control, 4 m, 8 m, 1 yr | Limited days of activity No impact [Note: these data are an \(a\ priori\) sub-group analysis] Days/person: I: Baseline: 20.6 (SD 25.4); FU: 18.7 (SD 36.8) C: Baseline: 27.8 (SD 33.4); FU: 27.9 (SD 55.7), no between group differences reported |                                                  |
| FU: 4 m, 8 m, 1 yr Overall risk of bias: high            | Process, 1 yr            | *Knowledge and beliefs: no effect [Note: these data are an \(a\ priori\) sub-group analysis] Mean scores I: Baseline: 14.1 (SD 2.9); FU: 14.6 (SD 3.2) C: Baseline: 14.3 (SD 2.3); FU: 14.7 (SD 2.3) No between group differences reported | Illustrated consistently no effect              |
| Keslo⁴⁰ \(n = 52\) adults                               | Behavioural              | Not assessed                                                                                     | –                                               |
|                                                          | Clinical-unscheduled care, 1 yr | Unscheduled care: compared to controls, the intervention reduced ED visits but not hospitalisations *Change in ED visits Baseline: I: \(4.4\) (SD 2.7) vs. C: \(3.4\) (SD 2.6); FU: I: \(2.6\) (SD 2.6 vs. C: \(3.5\) (SD 2.7) Between group difference \(p < 0.01\) Change in hospitalisations Baseline: I: \(1.3\) (SD 1.3) vs. C: \(1.0\) (SD 1.2); FU: I: \(0.5\) (SD 0.8) vs. C: \(0.5\) (SD 0.9) Between group difference \(p = 0.37\) | Illustrated as a significant positive effect but hatched to show inconsistency |
| FU: 1 yr, telephone every 2 wks to every 6 m Overall risk of bias: unclear | Clinical-asthma control Process, After intervention | Not assessed                                                                                     | –                                               |
|                                                          | Not assessed                                                                                     | No data reported for knowledge Insufficient data |                                                  |
| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher’s interpretation for the harvest plot |
|--------------------------------------------------------|------------------------|-------------------------------------------------------------------------------------------------|-----------------------------------------------|
| Keslo*44 n = 39, adults                                  | Behavioural            | Not assessed                                                                                   | Illustrated as a consistent significant positive effect |
| FU: every month then every 2-3 m                        | Clinical-unscheduled care, 1 yr, 2 yr | Unscheduled care: Intervention group had a greater reduction in hospitalisations and ED visits |
| Overall risk of bias: High                             | Process, before and after intervention | *Change in ED visits. Median (IQR) visits 2 years, I: 0 (0,0) vs. C: 2 (1.5, 2); p = < 0.05 |
|                                                        | Behavioural            | *Change in hospitalisations. Median (IQR) hospitalisations, I: 0 (0,0) vs. C: 0.5 (0, 1); p = < 0.05 |
| Velsor-Friedrich*40 CCT n = 102, children               | Clinical-unscheduled care, 2 wks, 5 m, 1 yr | Unscheduled care: the intervention group had significantly more unscheduled visits at 5 m and 1 yr |
| Overall risk of bias: unclear                          | Process, 2 wks, 5 m, 1 yr | Medical visits at 5 m. Mean (SE) I: 0.12 (0.05) vs. C: 0.02 (0.04) |
|                                                        | Process, 2 wks, 5 m, 1 yr | Medical visits at 1 yr. Mean (SE) I: 0.07 (0.03) vs. C: 0.00 (SD 0.02); p = 0.01 |
|                                                        | Behavioural, 2 wks, 5 m s | Symptom days: greater reduction in days with symptoms in intervention compared to control |
|                                                        | Behavioural, 2 wks, 5 m s | Symptom days at 5 m. Mean (SE) I: 2.15 (SE 0.30) vs. C: 1.42 (SE 0.21) |
|                                                        | Behavioural, 2 wks, 5 m s | Symptom days at 1 yr. Mean (SE) I: 1.26 (SE 0.33) vs. C: 1.49 (SE 0.23); p = 0.047 |
|                                                        | Behavioural, 2 wks, 5 m s | PEFR: intervention group had greater increase in PEFR at both FU time-points |
|                                                        | Behavioural, 2 wks, 5 m s | % increase in PEFR at 5 m. I: 2.9% (SE 2.0%) vs. C: 2.9% (SE 1.0%) |
|                                                        | Behavioural, 2 wks, 5 m s | % increase in PEFR at 1 yr. I: 7.5% (2.0%) vs. C: 2.9% (SE 1.2%); p = 0.046 |
|                                                        | Behavioural, 2 wks, 5 m s | School absences: no between group difference in days absent from school |
|                                                        | Behavioural, 2 wks, 5 m s | Days absent at 1 yr. I: 9.03 vs. C: 14.4 days |
|                                                        | Asthma knowledge test at 5 m. I: 14.05 (SE 0.55) vs. C: 13.35 (SE 0.38) |
|                                                        | Asthma belief survey at 5 m. I: 4.23 (SE 0.10) vs. C: 4.15 (SE 0.08) |
|                                                        | Self-perception inventory at 5 m. I: 2.80 (SE 0.08) vs. C: 2.85 (SE 0.05) |
|                                                        | Self-practice/asthma self-care: No significant between group differences |
|                                                        | Denyes self-care agency instrument at 5 m: 72.03 (SE 2.46) vs. 70.57 (SE 1.68) |
|                                                        | Asthma self-care instrument at 5 m: 68.87 (SE 2.89) vs. C:70.41 (SE 2.00) |
| Velsor-Friedrich*41 CCT n = 52, children                | Clinical-unscheduled care, 2 wks, 5 m, 1 yr | Urgent medical visits (and medications): no significant between group differences at any time point |
| Overall risk of bias: unclear                          | Clinical-asthma control, 2 wks, 5 m, 1 yr | Urgent doctor visits at 12 m. I: n=4 (14%) vs. C: n=5 (20%) |
|                                                        | Clinical-asthma control, 2 wks, 5 m, 1 yr | No data; some data on medicine use was provided |
|                                                        | Clinical-asthma control, 2 wks, 5 m, 1 yr | Symptoms, PEFR and school absences: no significant between group differences at any time point |

*Change in ED visits. Median (IQR) visits 2 years, I: 0 (0,0) vs. C: 2 (1.5, 2); p = < 0.05

*Change in hospitalisations. Median (IQR) hospitalisations, I: 0 (0,0) vs. C: 0.5 (0, 1); p = < 0.05
Table 3 continued

| Citation design, sample group/size and risk of bias score | Outcome categories, FU | Reported outcomes-values for intervention (I)/control (C) *indicates the primary outcome (if stated) | Researcher’s interpretation for the harvest plot risk of bias score |
|----------------------------------------------------------|------------------------|------------------------------------------------------------------------------------------------|-------------------------------------------------------------------|
| Process, 2 wk, 5 m, 12 m                                   | Proportion with > 1 day with symptoms/2 wks at 1 yr. : I 14 (50%) vs. C 13 (54%) % increase in PEFR from baseline at 1 yr. : I 26.21% (SD 0.22) vs. C 27.80% (SD 0.31) Average days absent from school. : I 9.03 vs. C 14.4 | Illustrated as a consistent positive effect but hatched to show inconsistency |
| Behavioural, 2 wks, 5 m, 1 yr                             | Knowledge and self-efficacy: Intervention group had higher scores at all time-points, but neither group improved over time Asthma Knowledge: test at 1 yr. Adjusted mean : I 14.28 (SE 0.80) vs. C 11.88 (SE 0.87); p = 0.03 Asthma belief scale at 1 yr. Adjusted mean : I 4.09 (SE 0.14) vs. C 3.82 (SE 0.15); p = 0.01 Self-esteem: no significance between group differences at any time point Self-perception inventory at 1 yr. Adjusted mean : I 2.71 (SE 0.08) vs. C 2.78 (SE 0.10) | Illustrated as a consistent positive effect |
| Velsor-Friedrich RCT n = 137, adolescents                  | Hospital visits: no significance between group differences p > 0.05 (no other data provided) Symptoms reduced in both groups; no significant between group differences PEFR: no significance between group differences School absences reduced in both groups; no significant between group differences Knowledge, self-efficacy improved in both groups; no significant between group differences Coping frequency/efficacy, no significance between group differences | Illustrated as a consistent no effect Symptom takes priority. Illustrated as a consistent no effect Symptom takes priority. Illustrated as a consistent no effect Illustrated as a consistent no effect |
| Clinical-Unscheduled care, 6 m, 12 m                     | Asthma self-care practice/general self-care: intervention group had higher scores at all time-points, but neither group improved over time Denyes self-care agency instrument. : I 75.55 (SE 2.60) vs. 67.41 (SE 2.82); p = 0.01 General self-care. : adjusted mean : I 72.99 (SE 3.26) vs. C 63.75 (SE 3.53); p = 0.2 | Illustrated as a consistent positive effect |
| Clinical-asthma control, 6 m, 1 yr                        | Hospital visits: no significance between group differences p > 0.05 (no other data provided) Symptoms reduced in both groups; no significant between group differences | Illustrated as a consistent no effect |
| FU: 2 m, 6 m, 1 yr Overall risk of bias: high             | Process, 6 m, 1 yr Knowledge, self-efficacy improved in both groups; no significant between group differences Coping frequency/efficacy, no significance between group differences | Illustrated as a consistent no effect |
| Behavioural, 6 m, 1 yr                                    | Self-care practice, no significance between group differences | Illustrated as a consistent no effect |

For conflicting outcomes within a category, the decision process was dependent upon priority of evidence including:
- Defined primary outcomes in an adequately powered sample/sub-group analysis (for the latter we will consider a prior sub-group analysis)
- Outcomes measured using a validated instrument (as opposed to non-validated instruments)
- Outcomes that were clinically and statistically significant (e.g., achieved significance defined minimum clinically important difference)
- If doubts remain, the author’s interpretation was considered to provide context for the final decision

Note:
- For quality of life outcomes, we will use the overall score, if no overall score is stated the outcome will not be plotted
- Asthma related quality of life scales will be given priority (e.g., AQLQ) over generic quality of life scales (e.g., ED5D)
- For the clinical-asthma control category, symptoms will be a priority over other outcomes in the same category as it is a better indicator of asthma control

Abbreviations: FU follow-up, wks weeks, m month, yr year, RCT randomised control trial, CCT clinical control trial, ED emergency department visits, I intervention, C control, CI confidence interval, AQLQ quality of life questionnaire, AQ20 the Airways questionnaire 20, ACT asthma control test, F F statistics, AHR adjusted hazard ratio, HR hazard ratio, OR odds ratio, EES estimated effect size, PEFR peak expiratory flow rate, SD standard deviation, SE standard error, DF degree of freedom, p p-values
the distinction between modified, tailored and targeted interventions is not clear-cut. Both culturally targeted interventions also incorporated some modified components. \(^{31,37}\) e.g., adaptation of language in PAAPs to meet the target population needs. \(^{31}\)

(2) Effectiveness of interventions in different sociocultural contexts.

In the harvest plot (Fig. 2 and Table 3), the four outcome categories (i.e., unscheduled care, asthma control, process and behavioural), are plotted for the three ethnic groups, ‘majority’ South Asian, ‘minority’ South Asian and ‘minority’ African American. \(^{37}\) The harvest plots show that the interventions in the ‘majority’ South Asian trials were effective, though notably they were all based in tertiary care settings potentially serving a relatively severe asthma population (thus with greater potential for improvement). \(^{30–33}\) In addition, risk of bias, was either high, \(^{31,32}\) or unclear \(^{30,33}\), and two of these trials had short follow-up periods (1 and 4 months). \(^{30,33}\)

In contrast, trial outcomes from studies involving both ‘minority’ communities were inconsistent, though more trials were at a low risk of bias, \(^{34–36}\) in contrast to ‘majority’ trials. In the ‘minority’ South Asian trials, most of the outcomes did not show significant benefit. \(^{34–36}\) The exceptions were improved quality of life in a trial at high risk of bias, \(^{36}\) and in another study improved self-efficacy at 3 months, which was not sustained at 12 months. \(^{35}\) Similarly, in ‘minority’ African-American trials (all but one were at high or unclear risk of bias), \(^{36}\) most interventions were ineffective, \(^{43,45}\) or inconsistent. \(^{39–41}\) In addition, one trial at unclear risk of bias had a negative impact on unscheduled care. \(^{46}\) Three trials had positive outcomes (unscheduled care and behavioural), \(^{41,44,46}\) of which one trial was at a low risk of bias. \(^{46}\)

(3) Identified barriers and facilitators to self-management in included trials.

A range of barriers and facilitators to asthma self-management were identified and differentiated according to ethnicity and sociocultural context (Illustrated in Fig. 3). Key findings were that:

- Across both ethnic groups and all social contexts, barriers included insufficient knowledge and understanding of asthma and related factors, \(^{31,36,37,43}\); facilitators included providing self-management education, \(^{31,32,37,39,44,45}\) and support from healthcare professionals (with continuity of care), \(^{31,32,37,41,44}\)

- In ‘minority’ trials, even though language barriers were accounted for, \(^{36,37}\) a barrier identified for South Asians, was insufficient consideration of individual learning styles related to age, \(^{36,37}\) gender \(^{36,37}\) and level of education. \(^{36}\) In a minority’ African-American trial, culturally/age specific self-management strategies (e.g., gaming) were identified as a facilitator. \(^{45}\)

- A facilitator that occurred frequently in studies involving South Asians across both majority and minority settings was providing culturally and linguistically appropriate educational materials. Language barriers were not an issue for ‘minority’ African-Americans. \(^{31,36,37}\)

- Some barriers and facilitators were specific to one of the two ethnic groups or social context. For instance, facilitators for ‘majority’ South Asian trials included generic self-management strategies, \(^{30–32}\) e.g., use of PAAPs; \(^{50}\) written reinforcement, \(^{31}\)
and practising preventative behaviour. One African-American trial observed that stressors (e.g., neighbourhood violence), interfered with generic self-management strategies such as relaxation and breathing exercises in adolescents. Similarly, three African American trials incorporated discussions of managing common stressors in daily African American lives as a facilitator, because this allowed individuals to focus on asthma. Another African-American trial identified social support as a facilitator.

**DISCUSSION**

Main findings

We identified seventeen RCTs, most at unclear or high risk of bias, which tested asthma self-management interventions for South Asian or African-American communities. Education was a component of all interventions, but content, mode of delivery and additional strategies varied. Only two interventions were culturally targeted, in contrast to 15 culturally modified interventions and no culturally tailored interventions. Trials based in South Asian countries, appeared to be more effective than those delivered to minority populations (for both South Asians and African Americans), though with the caveat that none of the ‘majority’ population trials were at low risk of bias and targeted populations were from tertiary care hospitals (in whom it may have been easier to demonstrate health benefits due to more severe asthma). Hence, it is unclear whether culture or minority-status of an ethnic group influences the variance in self-management outcomes. Education with on-going professional support was identified as a facilitator to asthma self-management in all groups. Other facilitators included focusing on individual learning styles in minority communities, culturally and linguistically appropriate education for minority and indigenous South Asians, generic self-management strategies in ‘majority’ South Asian communities, and strategies for dealing with stress and social support in African-American populations.

Interpretation of findings in relation to previously published literature

A previous systematic review concluded that a culturally targeted intervention (in line with the definitions of this review) was more effective than generic programmes in improving asthma outcomes, and revealed that most interventions were culturally modified. We found only two culturally targeted interventions, suggesting that this recommendation has not been adopted, hence progress in this area of research has not advanced. This may be due to the expensive and lengthy nature of developing targeted or tailored interventions compared to the ease of adapting or re-testing modified interventions, however, in the long-term culturally targeted or tailored interventions may be more cost-effective. Trials have typically considered ethnic groups as homogenous, e.g., they do not consider variation among smaller subcultural groups of South Asians or African Americans, or the influence of acculturation in minority communities, potentially important for designing interventions.
The two culturally targeted trials also included some modified characteristics, e.g., language adaptation for PAAPs, so the distinctions between culturally relevant interventions is not absolute. This is supported by a previous systematic review, which found interventions labelled as targeted or tailored also incorporated modified features, e.g., community/participatory approach to smoking cessation. It may be that modification of certain proven asthma self-management strategies, e.g., PAAPs, together with customising by culturally specific elements is an optimal approach.

Targeted trials customise the development of interventions to a cultural group rather than just adjusting the content. For instance, interventions developed collaboratively with target groups helped existing self-management strategies to be linguistically and culturally relevant. This can be further understood as aiming at deep structures, e.g., cultural beliefs, norms, lifestyles, environmental and social contexts, which aid receptivity of information and behaviour change. The Person-Based Approach to intervention development suggests that comprehension of user perspectives and contexts based on qualitative studies at every stage of development is central to customisation. In contrast, modifying surface structures to observable traits, e.g., language, ethnicity, food and clothing, may influence information processing but not behaviour change (a common characteristic of modified interventions). For instance, two ‘minority’ South Asian trials modified interventions according to language with mostly ineffective outcomes, suggesting merely focussing on language modifications is insufficient for their needs. However, more rigorous trials are needed, as both targeted interventions had either high or unclear risk of bias.

Similarly, some ‘majority’ South Asian interventions were modified from generic programmes rather than developed for their own community. For example, Ghosh et al. a trial from India, adapted self-management strategies from an intervention from Colorado, USA. Trials from diverse sociocultural contexts and different cultural groups demonstrate the potential pitfalls of extrapolating findings from one context and applying it to another. A possible explanation for ‘majority’ South Asian trials incorporating culturally modified strategies may be that international clinical guidelines for respiratory diseases, promote a generic model of self-management interventions with evidence and examples from high-income populations and recommendation of adaption to low or middle-income countries (LMICs). While remaining true to the core evidence-based features of supported self-management presented in guidelines, intervention developers also need to deliberate on the principles of cultural relevance to the targeted local community, rather than depending on translation. For LMICs, this may be challenging due to the lack of resources, training and manpower, as well as public health priorities and models of care focusing on communicable rather than long-term conditions. GINA guidelines acknowledge these difficulties, but do not offer specific guidance on providing targeted or tailored self-management, in contrast to the advice about cost-effective options for diagnosis and treatment in LMICs.

Conceptualising culture with its interaction with context offers new avenues of comprehending the role of culture in health. Apart from better outcomes in ‘majority’ South Asian trials based in tertiary care settings compared to ‘minority’ communities, poor reporting with limited descriptions of SES and diversity of trial settings meant we were unable to draw conclusions about associations between outcomes and contextual data. This is an important point as variations in SES within a culture has been suggested to determine health outcomes, e.g., restrictions in accessing services. In LMICs such as India, tertiary care may currently be the only practical setting for delivering asthma self-management interventions due to lack of community-based clinical and research expertise, as well as social and financial barriers that result in under-diagnosis, under-treatment and limited treatment availability. In the absence of adequately resourced primary care, it is common for individuals in these populations (particularly for children) to only access healthcare during exacerbations, rather than receiving preventative care.

Strengths and limitations of this study

To our knowledge, this review is one of few studies analysing the effectiveness of South Asian or African-American asthma self-management interventions. By identifying barriers and facilitators across two different ethnic groups and sociocultural contexts, our review can inform the customisation of interventions. We included seventeen trials, though the exclusion criteria of requiring separate outcome data for the specific groups of interest may have restricted the number of articles included in the final analysis; identification of more culturally targeted and even some tailored trials would have been informative. Limited resources precluded duplicate selection of papers, but we undertook a ten percent reliability check of the selection process. Risk of bias assessment was duplicated and data extraction was fully checked by a second reviewer.

Further, limited descriptions of the studies made it difficult to know how the interventions were developed or on what they were based on, particularly in the ‘majority’ South Asian trials in addition, few authors responded to our request for further information. This meant that one of the targeted trials was excluded from the harvest plot analysis because data on between group differences were missing. Additionally, some harvest plot decisions relied upon sub-group analyses, which reduce study power and thereby could have increased the potential for null findings. However, primary outcomes were prioritised and, for clarity, inconsistent findings were indicated by hatched bars to limit over interpretation. Subjectivity in assessing the outcomes for the harvest plot was minimised by specifying predefined criteria that were replicable, and all the judgements were checked by at least two reviewers. Additionally, even though harvest plots are a good technique of illustrating heterogeneous findings and can be personalised to the requirements of the review, they may neglect some important outcomes that cannot be reported in the plots and overemphasise others.

Conclusions and implications for future research, policy and practice

Asthma self-management interventions delivered in South Asian and African-American minority communities were less effective than interventions delivered in indigenous populations in South Asia, though the design/conduct of the latter studies meant that they were at greater risk of bias. Additionally, most trials from India are not designing interventions to their community, instead they are following guideline recommendations from studies in high-income countries. Studies that improve understanding of sociocultural contexts, allow a deeper appreciation of customising interventions and how to prevent inequalities in self-management behaviour, both are needed to inform international asthma guidelines. Targeted or tailored intervention development does not exclusively include collaboratively developed components customised to beliefs and needs of the target ethnic group, but may also include adoption of existing resources. Intergroup subcultural heterogeneities, cultural changes over generations (due to acculturation) and individual learning styles, add to the complexity of self-management behaviour and all need to be explored further. Rigorous trials of culturally targeted or tailored interventions are needed. Moreover, there needs to be standard recommendations on how trials verify participant ethnicity/culture, as only three ‘minority’ South Asian trials defined ethnicity according to self-identification or language spoken and culture

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was not considered and/or perceived to be synonymous to ethnicity.34,35,37

METHODS

The review protocol is registered with the PROSPERO database (registration number CRD42015020174). We followed the procedures described in the Cochrane handbook for systematic review of interventions.56

Search strategy

Our key search terms were ‘asthma’ ‘AND’ ‘self-management’ ‘AND’ ‘population’ (including terms for South Asian and Black communities as summarised in Table 4 (detailed in Supplementary Appendix 1)). We searched for RCTs on eight electronic databases (Medline, EMBASE, Web of Science, PsycINFO, Scopus, Elsevier Science Direct, Cochrane Library including Cochrane Airways Group Register of Trials and Google Scholar), three research registers in [February 2015] (PROSPERO, The University of York's Centre for Reviews and Dissemination, and the Clinical Trials Database), manually searched relevant journals (Patient Education and Counselling, Health Psychology and Ethnicity and Health), and searched reference lists of identified systematic reviews. The search was not confined by publication year or language.

Inclusion and exclusion criteria

We included RCTs evaluating self-management interventions delivered to South Asian or Black asthma patients, the parents/carers of children with asthma, lay or healthcare professionals who care for people with asthma from these communities. The search included populations of all ages and in any country. Black African Americans, were included because they are from another well-studied minority population, with experience of socioeconomic deprivation, and our scope of literature suggested there was a relatively large evidence base. Outcomes of interest were clinical (e.g., unscheduled care and asthma control),57 process, behavioural (e.g., knowledge and medicine adherence). We excluded studies that did not specify their population (e.g., trials using broad terms when describing their population such as 'West Indians' or 'Asians'), and trials of multiple ethnic populations that did not provide separate asthma outcome data for the ethnic groups of interest (see Fig. 1; The PICO strategy is summarised in Table 5).

Study selection

A PRISMA diagram was used to report the number of studies identified, the screening process and the final list of included studies (see Fig. 1). All titles, abstracts and full texts were screened by one reviewer (S.A.), and a random 10% by two other reviewers (L.S., H.P.). Disagreements were resolved by discussion and the inclusion/exclusion criteria clarified as necessary.

Data extraction and risk of bias

A standardised Cochrane data extraction sheet was modified for this study.58 All data extraction was completed by one reviewer (S.A.) and independently checked by a second reviewer (K.H.). Discrepancies were resolved by discussions between reviewers and the wider team (L.S., H.P.), until consensus was achieved. Trial authors were contacted by email to clarify any missing, unclear or additional data required. If contact with the author failed, the uncertainty was noted on the data extraction form. The Cochrane

### Table 4. Search strategy terms

| Asthma | Self-management | Population search |
|--------|-----------------|-------------------|
|        | Self management OR asthma control OR self care | South Asians |
|        | Barriers OR facilitators | Bengali OR Bangladeshi OR Bangladesh |
|        | Beliefs OR attitudes | Indian OR India |
|        | Knowledge OR asthma education | Pakistani OR Pakistan |
|        |                                            | Black OR African OR Afro Caribbean |
|        |                                            | Ethnic OR ethnicity |

### Table 5. PICO search strategy

| PICO | Criteria |
|------|----------|
| Population | South Asian communities (Indian, Pakistani, Bangladeshi etc.), or Black populations (African, Caribbean or Other) asthma patients, their parents/carers, healthcare or lay professionals. The search considered all population ages and countries |
| Intervention | Asthma self-management interventions in any healthcare, community or remote settings. We used the self-management definition of the US Institute of Medicine: ‘The tasks that individuals must undertake to live with one or more chronic conditions. These tasks include having the confidence to deal with medical management, role management and emotional management of their conditions.460 |
| Comparator | Asthma patients, parents/carers of children with asthma, healthcare or lay professionals supporting asthma patients, who did not receive asthma self-management intervention |
| Outcomes | Outcomes of interest were: |
|          | 1. Clinical outcomes: (i) current asthma control was defined as the degree to which different asthma manifestations were reduced/eliminated by treatment. Here, main categories include clinical-asthma control level (ii) future risk of adverse events and unscheduled healthcare utilisation. All clinical outcomes are aligned with the American Thoracic Society/European Respiratory Society Task Force standardised definitions57 |
|          | 2. Process outcomes: any outcome that occurred because of certain steps in a process, e.g., knowledge and self-efficacy |
|          | 3. Behavioural outcomes: outcomes related to behaviour, e.g., medicine adherence and inhaler technique |
| Exclusion | 1. All studies that did not explicitly specify population were excluded e.g., trials that did not provide details on which ethnic group they are referring to when they used broad terms such as 'West Indians' or 'Asians' |
|          | 2. Studies of multiple ethnic populations that did not provide outcome data separately for the South Asian and the Black ethnic groups or subgroups were excluded |
|          | 3. Trials studying multiple illnesses but did not provide separate outcome data for asthma were excluded |
EPOC Risk of Bias Assessment Checklist,\(^{39}\) was used to evaluate bias in included studies. This was independently coded by two researchers (S.A., K.H.), and any discrepancies were resolved by another researcher (L.S.).

Analysis
We anticipated that studies would be too heterogeneous for meta-analysis, and, therefore, used a narrative synthesis, illustrating key findings on trial effectiveness with a harvest plot.\(^{55}\) Harvest plots allow visual representation of the findings of a narrative synthesis (comparable to Forrest plots in a meta-analysis), facilitating comparison across studies.\(^{55}\) They enable identification of interesting patterns among varying outcomes, and may highlight the strongest or most inconsistent evidence, areas of possible concern, and gaps in the research. If there were various outcomes in one category (e.g., the asthma control category might include symptom scores, symptom-free days, or days off work/school with a range of significant and non-significant results), the overarching outcome was determined according to pre-defined criteria (see note to Table 3), applied and agreed by three researchers (S.A., H.P. and/or L.S.).\(^{55}\) Sizes of lines and colour hatchings were used to illustrate features of the trial according to predefined convention (see summary in footnote to Fig. 2 and detailed description in Table 3). Barriers and facilitators were identified from data and/or interpretations of study authors.

Data availability
All included papers are published; no further data are available. Requests for further information should be addressed to the corresponding author.

Disclaimer
The views expressed are those of the author(s) and not necessarily those of the NHS, the NIHR or the Department of Health.

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AUTHOR CONTRIBUTIONS
Development of concept and design of the work (S.A., L.S., H.P., S.J.C.T.), data collection (S.A., L.S.), screening and second reviewing titles, abstracts and full texts (S.A., L.S., H.P.), data extraction (S.A., K.H.), data analysis and interpretation (S.A., L.S., H.P.), second review of harvest plots analysis table (H.P.), third review of harvest plots analysis table (L.S.), initial draft of the manuscript (S.A.), critical revision of the article (S.A., H.P., L.S., S.J.C.T., K.H.), Salima Ahmed (S.A.); Hilary Pinnock (H.P.); Liz Steed (L.S.); Stephanie JC Taylor (S.J.C.T.); Katherine Harris (K.H.).

ADDITIONAL INFORMATION
Supplementary Information accompanies the paper on the npj Primary Care Respiratory Medicine website (https://doi.org/10.1038/s41533-017-0070-6).

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