

Long COVID and the role of physical activity: a qualitative study

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BMJ Open Long COVID and the role of physical activity: a qualitative study

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ABSTRACT

Objectives To explore the lived experience of long COVID with particular focus on the role of physical activity.

Design Qualitative study using semistructured interviews.

Participants 18 people living with long COVID (9 men, 9 women; aged between 18–74 years; 10 white British, 3 white Other, 3 Asian, 1 black, 1 mixed ethnicity) recruited via a UK-based research interest database for people with long COVID.

Setting Telephone interviews with 17 participants living in the UK and 1 participant living in the USA.

Results Four themes were generated. Theme 1 describes how participants struggled with drastically reduced physical function, compounded by the cognitive and psychological effects of long COVID. Theme 2 highlights challenges associated with finding and interpreting advice about physical activity that was appropriately tailored. Theme 3 describes individual approaches to managing symptoms including fatigue and ‘brain fog’ while trying to resume and maintain activities of daily living and other forms of exercise. Theme 4 illustrates the battle with self-concept to accept reduced function (even temporarily) and the fear of permanent reduction in physical and cognitive ability.

Conclusions This study provides insight into the challenges of managing physical activity alongside the extended symptoms associated with long COVID. Findings highlight the need for greater clarity and tailoring of physical activity-related advice for people with long COVID and improved support to resume activities important to individual well-being.

INTRODUCTION

As of January 2021, more than 93 million cases of COVID-19 have been reported worldwide.¹ Although the majority of people infected recover within weeks, many people experience persistent symptoms, irrespective of age and underlying health conditions, termed ‘long COVID’.^{2,3} Long COVID appears to be a multisystem disease associated with a complex array of respiratory, neurological, cardiovascular, gastrointestinal, musculoskeletal, rheumatological, dermatological and immunological symptoms ranging in severity, frequency and duration.^{3–6} Symptom surveillance surveys have reported estimates of long COVID (defined as symptoms lasting ≥12 weeks) affecting 10% of people testing positive for COVID-19.⁷ Preliminary findings

Strengths and limitations of this study

- To our knowledge, this article is the first to explore the role of physical activity in the lived experience of long COVID using a qualitative approach.
- The study design enabled in-depth enquiry of lived experiences in a diverse sample.
- Inductive thematic analysis ensured descriptions and interpretations of the lived experience were tested and found to be grounded in the data.
- Participants were recruited from members of a long COVID research interest database who registered via an online form, meaning study findings might not capture the views of digitally excluded populations

from MRI investigations also show that ~70% of ‘low-risk’ individuals testing positive for COVID-19 present signs of impairment in one or more organs 4 months after symptom onset.⁸ In a recent survey, the three most debilitating symptoms of long COVID were identified as fatigue, shortness of breath and cognitive dysfunction. 89% of survey respondents also reported that mental and/or physical exertion triggered the relapse of symptoms (postexertional malaise).⁴

Along with personal accounts,⁹ qualitative studies have provided valuable insight into the lived experience of long COVID.^{2 10 11} People with long COVID describe an illness trajectory and heterogeneous symptomatology that did not conform to initial expectations and were not acknowledged in public health advice.^{2 10 11} This has posed challenges for the medical profession, who have lacked evidence-based guidance to treat and support patients. Individual experiences of medical care have varied from well-meaning but inconclusive to disbelief.^{2 10 11} This lack of understanding about long COVID extends beyond the medical and scientific communities to employers, family and friends, leaving many people feeling frustrated and isolated in their self-management efforts^{2 10 11} and increasingly turning to online communities of people with long COVID for validation and advice.^{2 10 11}

A commonly reported approach to self-management for long COVID is pacing.^{29 11} This involves the strategic use of physical and mental energy reserves, with the aim of preventing or mitigating symptom flareups.^{12 13} Formal guidelines now include the Pace, Plan and Prioritise principles.^{12 13} Nevertheless, the role of physical activity (PA) in the management and rehabilitation of long COVID remains unclear. It is hypothesised that persistent debilitating symptoms are caused by organ dysfunction induced by the virus, but symptoms are likely to be compounded by deconditioning as a result of sedentary behaviour.^{14–16} While it is evident that PA can exacerbate symptoms, it is unknown how PA influences the broader recovery trajectory and how individuals have managed this aspect of their recovery. This study explores the lived experience of people with long COVID—focusing on the role of PA—to inform the design and implementation of rehabilitation support.

METHODS

Participants and recruitment

Interviews were conducted with English-speaking adults aged >18 years who self-identified as recovering from COVID-19, were not hospitalised (did not receive inpatient treatment) and had experienced a recovery period lasting 3 weeks or more, a time frame consistent with definitions of ‘postacute COVID-19’ at the time of study initiation.¹⁷ Ethical approval was granted by Sheffield Hallam University. Participants were recruited from a COVID-19 research interest database (the RICOVR database) established by the Advanced Wellbeing Research Centre (AWRC) at Sheffield Hallam University.¹⁸ At the time of the study, 2022 people were registered on the AWRC database (see [table 1](#) for characteristics). Consistent with other long COVID research, a positive COVID-19 test was not an inclusion criterion.^{17 19}

Purposive selection was used to ensure that the sample reflected a range of ages, genders and ethnicities. In accordance with UK demographic estimates,²⁰ we aimed to recruit 15% of participants from black, Asian and minority ethnic groups and 85% from the white British population, with an even sex ratio. To achieve a widespread of ages, two to three participants were selected from each age category (18–24 years; 25–34 years; 35–44 years; 45–54 years; 55–64 years; ≥65 years). Database members were stratified by age group, gender and ethnicity, and then selected chronologically according to the date that they registered with the RICOVR database. Two rounds of invites were sent by email; if invitees did not respond after 2 weeks or decided not to participate, we sent a new invite to the next person registered on the database (demographically matched). A total of 35 people were invited to participate; 21 responded to indicate interest. Respondents were provided with full details about the research and invited to an informal telephone discussion with the interviewer (HH) to discuss the research aims and procedures. Eighteen people took

Table 1 Characteristics of people registered on the RICOVR database at study commencement

	n	%
Total: 2022		
Sex		
Male	271	13.4
Female	1655	81.8
Undisclosed	96	4.7
Age		
18–24	28	1.4
25–34	219	10.8
35–44	575	28.4
45–54	686	33.9
55–64	337	16.7
>64	86	4.3
Undisclosed	91	4.5
Ethnic background		
White	1794	88.7
BAME	120	5.9
Undisclosed	108	5.3

BAME, black, Asian and minority ethnic.

up this opportunity; all subsequently decided to proceed. Participants completed a written consent form returned via email before interview. Seventeen participants were residing in the UK and one was living in the USA. After interviewing 18 participants, the research team were satisfied that thematic saturation was reached²¹ and recruitment ceased.

Patient and public involvement

During the study design, the AWRC Public Involvement in Research Group²² reviewed study aims and all materials. The group provided feedback to refine documents including clarification of language in the participant information sheet, rewording of interview questions and the addition of information about support for carers in the postinterview support document.

Interview procedures

A semistructured interview guide (online supplemental file) was developed to elicit participants’ stories about their lived experience of long COVID and the role of PA within that experience. Open questions explored four broad topic areas: (1) illness and recovery trajectory, (2) sources of support, (3) experiences of PA and (4) future priorities and concerns. All interviews were conducted during September and October 2020 via telephone with the exception of two interviews carried out using Zoom video conferencing to suit participants’ needs. All interviews were conducted by HH, an experienced female qualitative researcher in public health and exercise psychology. Interviews were audio-recorded and limited

to a maximum of 45 min to limit any cognitive burden for participants. Participants were not reimbursed but were signposted to information detailing sources of support within and beyond the University should any distress have been caused by the interview.

Data analysis

All recordings were transcribed verbatim by a professional transcription service. Transcripts were sent to participants for review; one participant responded with clarifications which were included in our analysis. Reflexive thematic analysis with inductive, semantic coding²³ was used to interpret the data. Consistent with recommendations, we did not set out to achieve intercoder reliability.²⁴ Instead, multiple researchers coded the transcripts to encourage reflexivity and ensure our analysis considered different possible interpretations. Two researchers (HH and LK, both with postgraduate psychology qualifications, training and experience in qualitative interviewing) reviewed 50% of the transcripts each. HH and LK independently developed preliminary coding frameworks presenting initial themes, which they compared and refined with input from a third researcher (NK) who had read a cross section of the transcripts. Following discussion with a fourth researcher (RC), HH, LK and NK returned to the transcripts to sense-check candidate themes and ensure that they offered an appropriate representation of the data, at which point themes were defined and named.

RESULTS

Participants

Eighteen people took part in an interview (see table 2 for sample characteristics).

Four themes were identified: *the interconnection of physical and psychological symptoms*; *lack of clear and consistent PA-related advice*; *learning to balance symptoms and activity*; and *adapting to an altered life*. Themes are described below with illustrative participant quotes.

Interconnection of physical and psychological symptoms

Participants recounted their experiences of long COVID in detail, describing diverse individual illness trajectories and varied patterns of symptoms presenting throughout the body. All participants described significant debilitation, with their physical function drastically reduced. This meant several weeks, and in many cases months, of being virtually housebound. During this period, basic activities of daily living (ADLs) including self-care and housework became challenging or impossible, and formal exercise was unconceivable:

The slightest thing was an effort in a way I've never ever conceived before, it's the most fatigued I have ever been ... things like changing my bedding, I did in stages like one pillowcase and then later in the day I'd do another pillowcase, it was that sort of level of difficulty with day-to-day tasks. (IV2)

Table 2 Participant characteristics

ID	Sex	Age group	Ethnic group	Place of residence
1	Male	55–64	White—Irish	UK
2	Female	45–54	Asian or Asian British—Indian	UK
3	Male	35–44	White—British	UK
4	Male	18–24	Black or Black British—Caribbean	UK
5	Male	45–54	White—British	UK
6	Male	65–74	White—British	US
7	Female	25–34	Mixed—White and Asian	UK
8	Female	65–74	White—British	UK
9	Male	35–44	Asian or Asian British—Indian	UK
10	Male	65–74	White—British	UK
11	Female	18–24	White—British	UK
12	Male	18–24	White—British	UK
13	Female	65–74	White—British	UK
14	Female	55–64	White—British	UK
15	Male	45–54	White—Other	UK
16	Female	35–44	White—Other	UK
17	Female	35–44	White—British	UK
18	Female	45–54	Asian or Asian British—Indian	UK

It has affected me to the point of like debilitating ... I've had to contact the school and say I can't do the one-way system to drop off. So literally I drop my son off right outside the school so I don't have to walk the 10 minutes' walk around the one-way system. I can't do that. (IV7)

The combination of physical and cognitive symptoms added to an overall sense of debilitation:

So if I do something physical I suffer. If I walk I suffer in my legs. If I do something with my hands I suffer with my hands. If I start to think too much I then get a foggy head. If I type an email on the computer and it goes on too long, I then can't think enough to shut the computer down. (IV5)

Physical and psychological impacts of long COVID were interlinked. For some participants, prolonged and unexpected physical incapacitation had emotional implications, including lowered self-esteem, frustration and guilt about not being able to fulfil everyday responsibilities. Some participants believed that stress was a specific trigger for physical symptoms:

I'm the wrong generation to just not go to work. So I know I couldn't go to work but I felt guilty about not

going to work ... once I knew every week that I had to be in touch I'd get really anxious because I knew I couldn't string a sentence together or even send a proper email. (IV14)

You see, stress brings it on and that makes it really bad. (IV13)

Lack of clear and consistent PA-related advice

Participants had varied expectations of how the health-care system could support them with PA. Some were deterred from help-seeking by unhelpful responses or a perception that they did not meet the threshold for help. Others acknowledged the constraints facing clinicians, yet felt compelled to seek advice about unnerving symptoms. Questions about how to manage PA challenged the expertise of medical professionals, who lacked an evidence base for this novel condition and were unable to provide conclusive advice. Consequently, participants sought information and validation online. Social media forums provided a community that normalised the long COVID experience and provided coping strategies. In addition, most participants adopted a 'researcher' role, reading scientific articles and health resources to better understand their condition:

I think the GP that I've managed to speak to has been trying his best, but he is a little bit unsure himself you see the most things that have been useful are the support groups on Facebook, I hate to say it but that's where I've found most of my information. And then obviously I've been reading a lot of evidence papers as well. So I've been ... trying to form my own opinion and diagnosing myself. (IV18)

As a consequence, professional advice about PA often arrived after information had already been accessed online:

They [physiotherapist] offered loads of advice just about pacing really. But I think at that point because of all the communities that have sprung up everywhere online people had already been sharing this information. (IV3)

While information accessed online was broadly helpful, it lacked specificity for long COVID:

There's obviously people that have had ... different types of viruses, and they're all claiming that doing exercise and whatnot is harmful for your recovery. I'm not sure if you've heard of something called ME, a lot of people are stating ... don't do graded exercise ... don't do none of that ... (IV4)

Some participants also struggled with digesting online information, finding long periods of concentration difficult. One participant highlighted the challenges associated with critically appraising evidence:

... I don't really have any knowledge of just medical sciences or have the ability to just go and start reading

the journals of the latest publications ... the common person can't do that. (IV12).

Learning how to balance symptoms and activity

Participants described a loss of freedom to engage with routine activities; any physical or cognitive activity resulted in the onset of fatigue. Some participants considered these relapses worthwhile, either because with each relapse followed a small perceived improvement in baseline function or because it was considered a price worth paying for the sense of normality, control and positive affect the activity provided:

So as much as I'm enjoying [walking the dog], it has the knock-on effect. But that is getting less and less, so the more I'm doing the better I'm feeling afterwards. I think [relapses are] all part of it, just got to get on with it and push myself a little bit harder and then hopefully I'll get better quicker. It doesn't put me off. (IV17)

Some participants were deterred from PA by seeing others with long COVID experience a symptom relapse. The severity and unpredictability of their own symptoms caused others to fear potential adverse events, along with the absence of any guarantee that permanent damage would not be caused by PA:

I read about pacing and I haven't tested myself, so I'm not exercising and I think that's a big thing for me ... So when I'm reading other people saying they get tired when they exercise, maybe that's put me off from trying that. Maybe it's the fear of not knowing. (IV9)

... I've always been one of those people that thinks well, you know, you push through it But this you just can't. And this is something that I'm becoming more afraid of that I think maybe I need to properly back off from as much daily activity as I can to recover from this because I'm scared that I will eventually end up as probably a 50% to a 60% of what I was previously, permanently, or for a longer term. (IV5)

Participants were especially discouraged from PA by symptoms such as breathlessness and heart palpitations. Where medical advice was sought or provided, this tended to err on the side of caution and thus reinforced decisions to avoid PA:

My heart rate being high is a worry. I do experience chest pain and constant tightness on my chest, which again is a worry. And I've never felt as if I could really push myself ... And my doctor has said ... well don't push it, you may do more harm than good by pushing too much. Which again doesn't help me an awful lot ... (IV1)

Most participants established personal strategies for managing PA based on trial and error. All described a limited energy reserve that must be used wisely, planning

their days according to personal schedule and priorities. Activities tended to be selected based on their benefit to well-being and/or responsibilities, for example, walking, getting outdoors and fulfilling caring duties:

I have to try and think well if I need to be around to look after my kids or something in particular in a day I need to be very, very careful of what I do the day before. (IV5)

I do the physical things that look after my mental health. So going outside and getting some fresh air looks after my mental health, and it in doing so helps the other symptoms, if that makes sense. So I focus on those a lot. So on a sunny day I'll go outside, because blue skies do me the world of good. (IV7)

This process of pacing and energy conservation was not without challenges. Many participants expressed a desire for better monitoring and support to manage PA, and bespoke long COVID guidance for establishing a safe, consistent baseline for activity:

Everything that you read is pace yourself, pace yourself. But that's really hard to do, because until you've overdone it you don't know how much you can do without overdoing it, if you see what I mean, so learning by default. Which isn't the best way, but I guess what's enough for me might not be enough for somebody else. (IV8)

... what I found with the advice for chronic fatigue syndrome is there seems to be, I don't know, I'm kind of just assuming this and making it up but it seems that there's more of a standard pattern; whereas with this it seems that no two days are alike so it's really hard to find a baseline because it's so erratic. (IV3)

Adapting to an altered life

There is a clear sense that long COVID is experienced as life-altering. Many participants described a loss of 'self' and a substantial impact on their identity. Some participants made sense of this as a 'pre-COVID-19' and 'post-COVID-19' life, and others described it as a journey. While the prospect of permanent disablement was distressing for all, many older participants drew on life events such as previous illness to make sense of their current experience:

... I'm self-aware when it comes to my health and my mood and I think I'm just accepting of it rather than wanting to change it or be negative about it. (IV13)

In contrast, younger participants (eg, those aged 18–24 years) struggled with their reduced function:

I just want my life back, it's getting a bit tedious. I see myself just becoming a burden, I don't want to live my life like that. (IV4)

The majority of participants had been unable to resume activities that were previously central to their core identity (eg, a parent, an employee). Any activity that provided a sense of normality thus helped to refute the idea that this

changed identity was permanent (a prognosis that was both feared and resisted). Participants for whom PA was a core feature of their self-concept previously referred to this as a strong motivator in their desire to improve functional capacity and resume PA:

I've been a runner and then a cyclist for many years so I had the intent of getting back in the walking. And then as soon as I could I got back in the cycling a little bit. (IV6)

I've always been physically active ... So it was somewhat testing myself, somewhat being hard headed and driven and also hoping that I'm actually fine, it's all over now and I can get back to playing basketball soon ... (IV15)

Some participants had reached a point of 'reluctant acceptance', not necessarily arrived at peacefully but through exasperation and for some, a degree of self-defeat. Participants described needing to give themselves permission (or seeking it from others) to rest, and some described a shift in priorities:

I'm just desperate to get back to exercise but I've had to learn that I can't, and a lot of mindfulness and patience and... the supportive people around me just helping basically, just listening. (IV3)

... I feel there's an opportunity for change. I might reduce my hours going forward. It's difficult but I might try and balance my work-life balance a bit more and pace myself. (IV9)

Family and friends played a significant role in participants' 'journeys'. While participants' symptoms were severe and physical function was most disrupted, friends and family acted as informal carers, supporting self-care activities such as cooking. They also provided emotional support, although as with medical professionals, this was sometimes contingent on an understanding and awareness of long COVID that developed gradually. Over time, family and friends became important PA companions, often adjusting their own PA behaviour to match the level of the person with long COVID and providing a reassurance that fostered confidence in activities such as walking:

I started going out every day and trying to do a bit more of a walk at home where it was relatively safe, my mum was there, she would walk with me and look after me. (IV11)

DISCUSSION

Resuming and maintaining 'normal' activity

Our findings illustrate the impact of long COVID on the energy reserves of individuals, which is consistent with previous reports.^{10 11} Our study highlights the importance of activities of daily living (ADLs (eg, housework, gardening) and outdoor activity to the normality and mental health of people with long COVID interviewed.

ADLs have been defined as fundamental skills required for independent living²⁵ and are often used to assess physical function. Participants prioritised and selected these activities based on their individual meaning, balancing what was physically possible against what provided the biggest return on investment in terms of feeling normal, engaged in family life or what boosted their mood.

Our findings indicate that people experiencing long COVID need better support to manage their symptoms, especially fatigue, while also helping them safely pursue the potential benefits of PA. As in previous research,¹¹ participants reported that guidance about PA was not always tailored to the complexity of long COVID. In particular, they lacked support on how to establish a baseline and upper threshold for activity which accommodated the erratic relapse–recovery cycle common to our participants' experience. Our research suggests that pacing needs to be individualised, considering biographical factors such as life-stage as well as current physical function and activity history. Many people will also require monitoring (either via professional or self-monitoring tools) to provide reassurance about the safety of PA while experiencing other symptoms like rapid heartbeat or breathlessness.

Our findings reflect concerns regarding the potential risk of long-term damage associated with postexertion malaise and PA. Participants differed in their attitudes towards relapse, some believing they were constructive to recovery, whereas others feared danger to long-term health. This paradoxical role of PA in relation to relapse and recovery reflects previous qualitative studies involving people with physically limiting conditions such as multiple sclerosis.²⁶ Parallels have been drawn between long COVID and myalgic encephalomyelitis (ME) and/or chronic fatigue syndrome (CFS).²⁷ Recently, the National Institute for Health and Care Excellence withdrew a recommendation to prescribe graded exercise therapy for patients with ME/CFS following concerns it could cause harm.²⁸ It is imperative to establish consensus, adding to what is already known¹⁷ regarding PA-related advice specifically for people with long COVID, including the identification of individual phenotypes for whom PA might or might not add value to their recovery.

Access to information and the role of the internet

For the majority of our participants, online long COVID communities provided opportunities to relate and empathise with others.^{2 10 11} Online communities pool collective knowledge derived from the lay expertise of members.²⁹ Information being shared is thus vetted and validated by the online community itself,^{30 31} and such groups can provide access to rapidly changing information that is inaccessible or unavailable offline.^{32 33} Advice about PA was accessed online much quicker than it was cascaded via medical professionals. The novelty of long COVID, however, meant that the lay expertise of members was sometimes dependent on learning from patients with other apparently similar conditions. In

the case of graded exercise therapy, this had potential to cause confusion where advice was controversial or disputed by some patients. Most participants also experimented with pacing and graded exercise without support from a health professional, risking potential damage to their health.

Influence of psychological characteristics on PA behaviour

Previous research involving people with long COVID has reported a substantial impact on identity.^{10 11} Our study extends these findings; people for whom PA was a core aspect of their identity pre-COVID-19 were more inclined to push themselves physically, willing to risk relapses or view them as constructive. Their mental health was also more contingent on being able to undertake PA. PA identity has previously been associated with commitment, ability and self-regulation, with research suggesting that identity–behaviour discrepant situations can create negative affect.³⁴ Further research could explore the relationship between PA identity and physiological and psychological responses to exertion and relapses in people with long COVID.

The common-sense model (CSM) of self-regulation³⁵ could provide a useful theoretical framework for understanding PA behaviours in people with long COVID. The model illustrates how deviations from 'normal self' (eg, onset of symptoms and disruption of function) interact with illness beliefs (eg, drawing on past illness experiences) to appraise the threat level of long COVID symptoms. This results in the activation of treatment action plans (in this case, PA choices and behaviours) that are mediated by social influences (eg, family/friends, mass media).³⁶ The CSM highlights a number of attributes that influence appraisal of health threats and subsequent behaviours, including expectations about the timeline for a given illness and perceived efficacy of self-management behaviours. In our study, these factors affected participants' likelihood of continuing to engage in PA; participants who felt that PA had a positive effect were more committed to maintaining PA. In contrast, some participants re-evaluated their PA behaviour over time, reducing PA when symptoms continued longer than expected or if they perceived PA to be harmful.

While the aim of our study was not to undertake a comparative analysis, we were alerted to potentially important age-related differences. Specifically, we noted that among younger participants the debilitating physical symptoms of long COVID appeared to have a particularly negative impact on participants' sense of self, and for these participants, adjusting to an uncertain future presented significant psychological struggle. Future work could examine the particular psychological-related and identity-related challenges of long COVID experienced by differing age cohorts. Sociological theories of illness including 'biographical disruption'³⁷ might offer a potentially useful explanatory framework for such work.

Implications for long COVID rehabilitation

The range of symptoms, severity, frequency and duration associated with long COVID³⁸ presents significant challenges for the publication of precise recommendations for PA in people with long COVID. Recommendations need to be tailored to individuals' clinical status and in some cases will require medical assessment by a specialist.

Our findings endorse ongoing efforts to establish holistic models of care that address both the physical and psychological impact of the condition.³⁹ They also highlight the limited capacity of primary care to provide PA-specific support. We agree with calls for multidisciplinary work to support people with long COVID,⁴⁰ highlighting the potential role of sports medicine and the need for collaboration between centres of well-being, behavioural science and professionals from a range of medical specialties.

Limitations

In our study, 17 out of 35 people invited chose not to participate. We did not ask non-responders to provide a reason for declining participation, and thus cannot speculate on their personal choices, but the scope of our study indicates a potential for response bias towards people with a particular interest in PA. Qualitative research of this kind is typically based on small samples and is not intended to be generalisable, seeking instead to provide trustworthiness and sufficient context to allow readers to make their own transferability judgement.⁴¹ To date, inconsistent diagnoses of long COVID have made it difficult to ascertain the demographic profile of people with long COVID at a population level, with women disproportionately represented in symptom studies⁴² and qualitative research^{10 11} along with people of higher educational background.¹⁰ We took steps to achieve a diverse sample in terms of age, gender and ethnicity, but did not collect information on socioeconomic status or geographical characteristics of participants, so are unable to draw firm conclusions about how these factors may influence our findings. Nevertheless, our participants were recruited from a research interest database indicating a level of engagement and access to online research, and our sample might not be representative of the long COVID population as a whole. Inclusion of people from digitally excluded populations⁴³ and a wide range of demographic profiles is important for future research and a key factor in providing equity of access to long COVID support.

CONCLUSIONS

This study provides insight into the challenges of managing PA alongside the extended symptoms associated with long COVID. Findings highlight the need for greater clarity and tailoring of PA-related advice for people with long COVID and improved support to resume activities important to individual well-being.

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Contributors RC initiated the idea for the study. HH designed the protocol and secured ethics for the study with support from RC. NK undertook stratification of database participants. HH undertook recruitment including invitations to take part, provision of participant information and collecting written informed consent. HH carried out all interviews. HH and LK undertook initial data analysis and developed preliminary coding frameworks. HH, LK and NK reviewed and refined themes with oversight from RC. HH prepared the manuscript, and all authors edited and revised the final version.

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