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A Qualitative Exploration of the Impact of COVID-19 on Individuals with Eating Disorders in the UK

Running Title: Impact of COVID-19 on people with eating disorders

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Abstract

COVID-19 may have substantial impact on the mental health at a population level, but also has the potential to significantly affect those with pre-existing mental health difficulties such as eating disorders. This qualitative study explores the impact of COVID-19 and associated public health measures on adults with eating disorders within the UK. We conducted 10 in depth interviews with adults (24-38 years) with a self-reported eating disorder during lockdown. Data were analysed using an inductive thematic analysis approach. We identified core themes related to social restrictions (social isolation, changes in accountability to others, and increased responsibility for self and others), functional restrictions (lack of routine and structure, a need to intentionally plan activity, a desire for secrecy particularly around food shopping) and restrictions in access to mental health services. Overall, the impact of the lockdown was experienced as a catalyst for either increased disordered eating behaviours or for a drive for recovery, depending on individual circumstances going into these restrictions. This study is the first in depth interview approach with adults with mixed eating disorder presentations in the UK. Findings have important implications for post lockdown intervention care and practice.

Keywords

Coronavirus, COVID-19, eating disorders, qualitative, thematic analysis, lockdown
Coronavirus disease 2019 (COVID-19) is a global pandemic with far-reaching consequences for the physical and mental health of the population, leading to the World Health Organisation (WHO) declaring it a Public Health Emergency of International Concern in January 2020. The significant morbidity and rapid spread of the virus has led to the activation of various levels of public health measures, including “lockdown” in a high proportion of countries and physical distancing measures to prevent transmission, resulting in unprecedented impacts on social interactions, employment and the world economy. While the impact of the pandemic, and associated management, on mental health is not yet fully known, the potential for psychological distress is significant, as a result of the effects of social isolation, the economic fallout, grief and trauma for survivors (Reger, Stanley, & Joiner, 2020). Initial reports indicate increased levels of distress and anxiety among the general population (Wang et al., 2020) and specific concern has been noted for the potential impact on vulnerable populations, including those with pre-existing mental health difficulties (Holmes et al., 2020).

One particularly vulnerable group in this context may be those with eating disorders. The effects of the COVID-19 lockdown on individuals with an eating disorder could be broad ranging. While there may be potential for some protective consequences – e.g., reduced interpersonal triggers such as face-to-face body-based social comparisons (Cooper et al., 2020), increased opportunity of support from loved ones (Murphy, Calugi, Cooper, & Dalle Grave, 2020), or services embracing new technology in the delivery of psychological therapies (Murphy et al., 2020) – the overwhelming concern within the field is the potential for severe, adverse impacts (Weissman, Bauer, & Thomas, 2020). Initial small scale pilot data from Spain indicates worsening of the mental health of individuals with an eating disorder including a deterioration in eating disorder symptomology for one third of respondents (Fernández-Aranda et al., 2020) and in a large scale survey in Australia conducted within the first few weeks of the pandemic, a significant proportion of individuals who self-identified as having an eating disorder (n=180) reported an
exacerbation of restricting, binge eating, purging and exercise behaviours, relative to before COVID-19 (Phillipou et al., 2020).

Rodgers and colleagues (2020) outline three pathways that may either precipitate the development of disordered eating during the pandemic, or exacerbate existing difficulties. The first involves the impact of disruption and restrictions to daily activities as a result of public health interventions aimed at reducing transmission of COVID-19. This includes public restrictions on exercise, grocery shopping and concerns around scarcity of specific foods. These restrictions may be highly provoking for individuals with rigid and inflexible exercise or eating patterns, for example, by increasing the perceived need to stockpile food, and associated risk of binge episodes (Touyz, Lacey, & Hay, 2020). This pathway also highlights the potential for reduced social support, including restrictions in access to treatment as a consequence of limitations placed on traditional face-to-face treatment as a result of social distancing (Touyz et al., 2020) and restructuring and reorientation of health services to prioritise the management of COVID-19 (Davis et al., 2020).

The second pathway relates to the effects of media (Rodgers et al., 2020). The authors propose that social distancing may lead to an increased use of social media and consequently enhanced exposure to harmful eating and appearance-related content (e.g. review by Holland & Tiggemann, 2016), and to stressful and traumatic world events which has been shown to negatively impact eating behaviours (Rodgers, Franko, Brunet, Herbert, & Bui, 2012). In addition, the increased necessity to use video-conferencing may be distressing for individuals with body avoidance.

The final pathway relates to fear of contagion, which may in turn lead to an increase in restrictive eating patterns and orthorexia-based cognitions, alongside increased levels of general stress and emotional distress, increasing the risk of disordered eating patterns (Rodgers et al., 2020). Other authors also highlight the potential financial impact of the pandemic, inclusive of reduced capacity for carers to support individuals with eating disorders, due to the need to increase working hours (Davis
et al., 2020) or the impact on food insecurity on financial ability to purchase “safe”
or binge foods (Touyz et al., 2020; Weissman et al., 2020).

These theoretical pathways provide a useful framework to explore the potential
impact of COVID-19 and its management on those with eating disorders. Given the
rapidly changing environment, to date these mechanisms and pathways are largely
hypothesised, with relatively little literature based on those with lived experience of
an eating disorder. There are increasing calls for researchers to understand the
psychological, social and neuro-scientific effects of the COVID-19 pandemic on
mental health in collaboration with those with lived experience (Holmes et al.,
2020). In this study we therefore aim to contribute to this growing body of work by
exploring the experience of adults affected by eating disorders during the COVID-19
pandemic in the UK using in depth interviews.

2. Methods
This study is reported in line with the COREQ guidance (COnsolidated criteria
for REporting Qualitative research) (Tong, Sainsbury, & Craig, 2007). Ethical approval
was provided by the University of Edinburgh (Ref: STAFF181, 05/06/20).

2.1 Participants and Sampling
Purposive sampling was used to recruit adults living in the UK who self-identified
as experiencing an eating disorder. Due to international differences in how
governments approached the pandemic, our study only included UK residents.
Advertisements for the study were posted on social media, mainly Twitter and
Facebook. All advertisements provided a study link to a participant information
sheet, to inform potential participants about the aim of the study, to verify eligibility
criteria and to obtain informed consent to be contacted. Participants were asked
to provide an email address to be contacted to schedule a one-time interview
alongside brief demographic information.

In total, 44 individuals noted an interest in the study of whom 15 consented to take
part, met the eligibility criteria and provided a valid email address to be contacted.
Of these, two opted out of the study and three did not respond to email communication. Therefore, in-depth interviews were conducted with ten adults, which was the minimum sample size striven for to enable data saturation. Interviewers had no prior relationship to interviewees, except for one participant who knew the interviewers from a previous eating disorder awareness event. Nine participants identified as female; one identified as non-binary. The mean age of participants was 29.6 years, ranging from 24 to 38 years, and all participants identified as White. Five participants lived in England, five in Scotland. Five participants lived alone at the time of the interview, two participants lived with family, one with a roommate, one with a partner, and one with a partner and family. All participants identified with disordered eating behaviours for more than two years. Six participants mainly identified with Anorexia Nervosa, two with Eating Disorders Not Otherwise Specified and one with Bulimia Nervosa.

2.2 Data Collection
Participants were contacted via email to arrange a one-on-one Skype interview with one of two female interviewers (SMB (MSc) and MCO (MSc), both PhD students in the field of eating disorders at the time of the interviews). Skype interviews were audio-recorded and transcribed verbatim by three researchers (SMB, MCO and IP). No field notes were taken during the interviews. The purpose of the study was fully disclosed to all participants prior to the study and interviewees had the opportunity to enquire about the researchers’ motivations and interests in this research topic subsequent to the interview. A semi-structured interview schedule was used, which was provided to participants beforehand if they requested it to reduce the likelihood of being triggered by any of the questions. The schedule was pilot tested by each interviewer and focused on the general impact of COVID-19 and resulting lockdown measures, the specific impact on eating behaviours, food purchasing and exercise behaviours. In addition, participants were asked about how the media had impacted their mental well-being during lockdown and how their support systems were impacted by the restrictions. In the UK, “lockdown” was enforced on March 26th, 2020 where all UK residents were asked to stay at home unless purchasing basic necessities, for medical need, essential key worker travel to work or one form of
exercise a day. Restrictions on outdoor activities (e.g., exercise) started to ease in mid-May 2020, and non-essential businesses remained closed until mid-end June (specific dates vary across the UK). Interviews were conducted from 14\textsuperscript{th} May to 4\textsuperscript{th} June 2020 and lasted between 45 and 120 minutes. All participants were debriefed after completion of the interview and provided with external support resources. Transcripts were not returned to participants for comments or corrections.

2.3 Theoretical Position and Analysis

This study was informed by Houston’s (2001) illustration of critical realism, which recognises human subjectivity, while acknowledging that personal meaning is shaped by social structures. In the present context, the COVID-19 pandemic and resulting lockdown measures can be seen as structures and powers that actuate specific psychological mechanisms. These mechanisms in turn cause so-called tendencies (e.g. behaviours, thoughts, feelings). Our main analytical goal was to understand and explain these tendencies, considering underlying psychological mechanisms and structures (Houston, 2001).

Positioning us as researchers within the framework of critical realism involves questioning our own assumptions, to better understand how those participating in our study interpret their own actions, thoughts and feelings (e.g. Manicas, 2009). Firstly, all researchers involved in this project experienced the lockdown measures in the UK first-hand which may have had an influence on how we expected participants to feel during this time. All authors are mental health researchers in the field of eating disorders, which constitutes an ‘insider conflict’ (Aguinis & Henle, 2002; Holian & Coghlan, 2013). One of the authors has lived experience with disordered eating behaviours. Therefore, we must acknowledge the impact of assumed knowledge, use of vernacular and assumed shared beliefs on our research.

All transcripts were coded line-by-line using NVivo (QSR International, Melbourne, Australia) and a thematic analysis was conducted in accordance with the steps outlined by Braun and Clarke (2006), using an inductive approach. Three researchers (SMB, MCO, IP) coded four transcripts each to identify preliminary themes, while
allowing for partial cross-validation between coders. During two meetings, the researchers discussed identified themes in the context of critical realism. Subsequently, preliminary themes were grouped and, if necessary, adapted. This process was followed by a second coding phase, which focused on the identification of common underlying structures, psychological mechanisms and resulting tendencies. Prevalent “patterned responses” (Braun & Clarke, 2006, p. 10) were identified to investigate meaningful structures across all data sets. Initial codes were reviewed among the coders and three overarching themes were determined as coherently representing the complexity of the data. Eventually, all coders were familiar with all transcripts and two further meetings were used to finalise the thematic analysis by clearly defining all themes and subthemes. Participants did not provide feedback on the findings.

3. Results

Across all interviews, the impact of the lockdown could be described as a catalyst for either disordered eating behaviours or the effort to recover. Participants who were managing better during lockdown attributed their coping skills to comparatively better personal circumstances at the onset of lockdown and expressed concern about the possibility of being in lockdown during a severe phase of disordered eating.

Our study identified three main themes of underlying lockdown structures: social restrictions (changes in how people were socialising), functional restrictions (changes in daily routines around work, shopping etc.) and restrictions in access to professional support.

3.1 Social Restrictions

Social distancing measures were introduced during lockdown to contain the spread of the virus. Restrictions in social interactions were therefore the most decisive overarching structure influencing participants’ mental well-being. Under this theme we identified tendencies related to participants experiencing social isolation, changes in accountability to others meaning increases or decreases in disordered
eating behaviours, and participants needing to take on more responsibility for themselves and others.

3.1.1 Social Isolation

The COVID-19 public health restrictions had a significant impact on most participants’ social interactions, especially for those living alone. Loneliness was a prevalent theme in all interviews as illustrated by participant 2696:

“Times when I would normally kind of be doing something potentially social or something like that over the weekend...Obviously with more free time, I might have gone back to see my parents--that [...] feeling, of like, existential loneliness felt incredibly desperate and really quite painful. But it was...It came in bursts to begin with, and I think as lockdown has gone on, it’s that feeling of real painful loneliness.” (R2696)

Being socially isolated while struggling with an eating disorder was linked with the tendency to become even more focused on food and disordered eating behaviours:

“Whereas, since lockdown, because I live alone...I’m on my own in the house because there’s nobody else around and I’ve got my house full of food, I have more and more preoccupied thoughts about food.” (R4880)

One participant compared the first weeks of confinement during lockdown with being back in hospital, while two other participants described their realisation that the lockdown only emphasised how socially isolated they had been before. Becoming aware of this loneliness was seen as painful, but experiencing this social isolation due to external circumstances made it more apparent that enhanced social support would be helpful:

“Sometimes I think I found I couldn’t manage the intensity of what I was going through [with the eating disorder]...I’ve lost relationships because of it. Now I feel like I’ve got a [...] a very small [support system] compared to how I feel like I would need. But I feel like I could do with a lot more.” (R7375)

3.1.2 Changes in Accountability
Depending on their living situation, participants in this study either experienced an increase in or decreased accountability to others for their behaviours during lockdown. Even though feelings towards accountability were ambivalent, participants had the tendency to associate an increase in accountability with improved eating pathology. Working from home without face-to-face social contacts led one of the participants to actively engage in her eating disorder:

“And there were more days of not eating the week before lockdown. I of course wasn't allowed in work and it was before they’d realise that, "Oh! You can do your job from home!" And...I just...I didn’t eat for that entire week because it was like, ‘Hey! I’m not accountable! No one else is here! This is the dream!’...Which is very dysfunctional! But it was absolutely...‘This is all I’ve ever wanted!’ with my ‘eating disorder brain’. [...]” (R5082)

Another participant felt more accountable at home, where she lives with her partner:

“I’m very busy at work. And no one pays any attention to what I am eating, in my job. Whereas when I am at home with my partner, erm... and we eat together, it’s much more difficult for me to not eat. Because he will ask me to eat a meal with him, or a snack with him [...]. I don’t think that he’s – he’s perhaps as aware of that happening as I am. So, he just wants to eat lunch together.” (R1443)

All participants showed a high level of self-awareness for their disordered eating behaviours and how they previously or currently engage in them. An ambivalence towards accountability was experienced by most participants, which reflected a tension between both distress and relief associated with disordered eating behaviours. Even though accountability was seen as helpful, not being accountable (due to being alone) was also associated with feelings of safety by one of the participants:

“Like – if I was going to my work every day, I wouldn’t be able to like exercise this much in the morning or I could, but I would have to get up super early. And I’ll be out of the house longer and people will expect me to eat lunch and they’ll
expect me to not just eat salads and my mum will expect me to go out with her
more [...] like there’s lots of more expectations of...I suppose less opportunity to
hide and be quite like safe and withdrawn.” (R7260)

The social restrictions of the lockdown are therefore an opportunity to evade
expectations for recovery. This, however, was associated with anxiety regarding a
future post-lockdown, when it wouldn’t be possible to actively engage in certain
behaviours anymore.

“[...] I used to slightly be anxious about coronavirus, but now I’m just anxious
about where it’s going to go from here. I’ve kind of gotten used to those
thoughts. Now, I’ve got future worries about how I’m gonna go back out into the
world” (R9143)

3.1.3 Increased Responsibility

Due to the social distancing measures, participants had less or modified professional
support and communicated with their friends and family primarily via phone and
online. Having had experiences of continued recovery was therefore an opportunity
to claim responsibility for certain accomplishments.

“I hope that I won’t slip back into that habit [not eating lunch], because I think
actually, I’m doing quite well now. [...] and it will be nice if I could take
responsibility for that myself as well, really.” (R1443)

Again, the conditions appeared to influence how participants coped with the
increased responsibility for themselves and others. Participant 1443 was working
towards recovery before the lockdown was introduced and received additional
support from her partner. Other participants were living on their own or had to take
on additional responsibilities due to the pandemic. Participant 4110, who was taking
care of her two younger brothers due to her mother being part of the high-risk
population, described how buying foods for others increased her preoccupation with
food and compensated for not eating the food herself:

“[...] I am buying a lot of food just like for my brothers, because I am doing all
the shopping because my mum is not here. I am buying so much food. [...] – and
a lot of it I am doing because I know that I can’t eat it. Like I am – I just buy
everything, [...] it’s not even pleasure, I just don’t know, I am obsessed with it.
Like I hate food shopping I absolutely hate it. But I spend - I have never spent so
much money on food shopping in my life.” (R4110)

Heightened responsibility was experienced by most participants, but resulting
behaviours and cognitions differed depending on their living situation, eating
disorder progression and how readily accessible additional support was during the
first weeks of lockdown.

3.2 Functional Restrictions
The lockdown not only limited people’s social interactions, but also the way they
could organise their daily life. Many activities such as the commute to and from
work, meal time routines and food shopping had to be altered, meaning participants
had to build up new routines. Related to these functional restrictions, we identified
tendencies associated with this lack of structure, becoming increasingly ‘intentional’
in planning social activities and exercise, and managing a desire for
anonymity/secrecy in the context of food purchasing.

3.2.1 Lack of Routine and Structure
All participants referred to rigid behaviour when describing their disordered eating
behaviour. Routine and structure were not only seen as important, but also essential
to being able to cope with dysfunctional thoughts and behaviours. The lockdown
disrupted established routines and heightened participants’ need for introducing
new structures and routines into their lives:

"So…So, my eating routines have changed probably for the better. Because I’m
with my partner more. Erm…Yeah. I think…[pause] Erm… [pause] I-I think I have
struggled a lot with…Worry about not getting food that I feel comfortable
eating." (R1443)

Maintaining both daily routine and structure functions to both perpetuate and
mitigate disordered eating psychopathologies. Most respondents referred to these
behaviours, especially in relation to times and environments associated with eating. Participant 7260 referred to this in terms of social cues and expectations from colleagues:

"Yeah because I see what was keeping me in my routine was having people around me, so like some people at my work knew so they’d be like ‘It’s lunch time!’ and like we would all be all over so we might not eat together but like because -and we would eat at our desks and stuff as we were working, but because people said like it’s lunchtime like it was easier to do things in a routine when you’ve got more of a routine. Whereas like, the whole day just seems the same even if you’re working, even if you’ve got meetings or whatever, it’s not, the day is split so like yeah." (R7260)

Similarly, participants referred to a lack of routines making disordered eating habits less severe. Without pre-lockdown routines and structures, Participant 4110 did not feel compelled to mitigate maladaptive eating behaviours:

"I don’t know, like I think – there’s just always like just that thing that like I go to in my life that as soon as things like kind of change and go a bit crazy like it’s my kind of go to and the – and my brain is automatically like, 'Ok, well, like, let’s just stop it then forever, d’you know- let’s just cut down, or let’s do this or...' There was definitely change or it was quite slow at first it wasn’t like let’s stop eating altogether it was just like let’s cut back a bit and see how that works." (R4110)

As established routines and structures were inevitably impacted due to the lockdown, participants reported being disconcerted by having to deconstruct rigid regimens to adjust to the current situation. For Participant 2696, the lack of physical boundaries distinguishing work and leisure has been unpleasant:

"But, it has been kind of strange, and I personally have actually hated working from my flat. I really, really like to implement those physical boundaries around saying, "Okay, I'm going to work now"--treating study like work, going to the library, getting to my lectures and then I come back to the flat. "Fine. This is where you don't work. This is where I chill out. I rest." So, that's been very difficult." (R2696)
3.2.2 A Need for Intentionality

Prior to the lockdown, participants’ daily routines were to some extent externally regulated and offered diverse opportunities to socialize, without actively choosing to do so. Having to compensate for this new type of confinement and a more sedentary lifestyle led participants to introduce more intentional, consciously-planned activities or to intensify their exercise routines, which were in some instances perceived as compulsive.

“Yeah, because now it is like exercise for exercise sake, whereas like before it was like a social thing. I was doing that with people, and I was going to the gym to see people and then I was going running with people and now it’s just like I need to exercise because I am in the house sitting still being lazy all day.” (R7260)

Having to schedule all social interactions further meant that participants had to actively reach out for support if needed. Participant 9143 described how her problems with communication are part of her eating disorder and part of why she misses the casualness of social interactions before lockdown:

“[..] It's harder to bring things up if today I'm struggling...Before, it was a lot easier in-person to pass it by in conversation rather than make such a big deal out of it. That's what I feel like it is now—just a lot of emphasis rather than just notice I'm not very well. [...] But, when I'm with some people I know, I find it hard to open up, and they usually can tell a lot by my body language and behaviours. That's probably a big reason why I have an eating disorder and still do...It's a way of communicating, isn't it? If I'm not okay and people can't see that, I find it hard then to communicate how I am, or if I need help or something without being in person.” (R9143)

The intention to socialise across distinct environments within participants’ daily routines became increasingly apparent once interaction frequencies changed and environments became more static. For participant 4880, unintentional social interactions grew more apparent after their day-to-day schedules were disrupted:
“I don't really have a 'social life' so to speak, before lockdown anyway. But the thing that has changed is I'm not having the little interactions I would have been having before with other people at the swimming pool or with my yoga teacher, or with colleagues in the office at work. So, I'm not having any of these interactions.” (R4880)

Overcompensating for both a lack in activity and a perceived inability to purposefully ask for support are the result of fewer opportunities to engage in daily rituals. Participants discussed replacing typical daily lower intensity exercise with more moderate and vigorous exercise once lockdown restrictions were implemented:

“Erm...Yeah, I think I – I think I started running properly, erm... at the beginning of lockdown. Because I think, for me, I couldn’t...Not being able to go out for long walks in [the national park] or wherever was really difficult. So, I think I just felt like I needed something to replace that—to try and keep myself...stable.” (R1443)

Participant 5082 reported utilizing the ‘one form of exercise per day’ mandate by attempting to exercise as much as possible in the allotted opportunity for physical activity:

“[...] I can only get out once a day. I'll have to make the most of it. I'll have to run. There can't be any, ‘Fuck it. I'm not doing it. I feel like shit.’ You have to get out. You have to do it. And then...Thinking of days when I felt I really had to compensate. It would be walking a long way to the shops. And then...Yeah. It was a bit of a grey area, in terms of, "Should I really be out for three, four hours running on the fell? Erm...Probably not.” (R5082)

For most participants, interrupted routines and structures impacted rigid behaviours that served various functions regarding eating pathology. Intentionality across social interactions and exercise regimes became increasingly evident as the lockdown prevented access to work environments, altered social interactions, and increased perceived sedentary behaviours. However, situations differed based on individual
motivations behind established routines, which either mitigated or exacerbated disordered eating behaviours, perception of sociality, and exercise habits.

3.2.3. Secrecy

Concern around being recognised in shops and whether or not disordered eating behaviours were noticeable was raised by some participants. Anxiety surrounding others’ assumed options on deemed ‘non-essential’ food purchases, frequency going to supermarkets, and detectability of disordered eating symptoms contributed to a want to maintain secrecy.

For two respondents who identified with binge-eating disorder and OSFED respectively, heightened awareness of food purchasing behaviours impacted food purchasing behaviours. Participant 2445 referred to frequenting different shops to possibly prevent shop staff from noting perceived inappropriate purchases during lockdown:

"The fear of being recognised is what has made me feel very anxious about going to the shops, so I tend to switch stores every two or three days just to make sure that people don't recognise me and they don't know who I am and I can be free to purchase whatever I want to purchase! It's a bit of an awkward concept." (R2445)

Shame was also evident in discussions around food purchasing and possible staff recognition. With limits on store occupancy and designated for essential workers, shopping behaviours that perhaps once seemed more nondescript now seemed more conspicuous:

"That guilt going to the shops. The 'beige food trolley', which...And it's even better because at eight o'clock, it's yellow sticker shopping. And NHS key workers get straight in. So, you get a full range of the entire binge foods that you could want! Or, what I would use. And then, because of where I live...Well, it's a tourist town. And the shop is normally really, really busy. Loads of tourists struggle to even buy milk. Now, eight o'clock I can go and be the only person in. And if I'm on full-blown 'binge mode', 'This is what is happening!'...The shop
assistants know me. And, if I bump into people and they recognise me, and then it's this whole thing, ‘Oh...I was in here two days ago doing this...’ So, that's been hard...” (R5082)

Depending on participants’ living situation, hiding certain behaviours became a way to avoid friends’ and relatives’ concerns or help:

“I don’t [talk about my eating disorder now], because I don’t want, I don’t want anyone to stop me either, like I kind of do – but I also don’t.” (R7260)

While for those sharing the household with a partner, not being able to keep certain behaviours hidden was a cause of anxiety, because they were forced to accept how disordered their behaviours had been.

“I used to do nearly all of our food shopping. And... my partner would just let me get on with it, because I did the food shopping. Erm... whereas – because he’s wanted to make sure we’ve had enough of things, and he knows that I’m not likely to judge that very well, erm... he’s started doing the online shopping, or checking it before... before we place the order and adding loads of things... which makes me really anxious. Because [...] I just - I hate it. [...] but also, it kind of is making me realize how... perhaps... disordered some of my habits were. [...] a lot of the... stuff around food during lockdown has made me really anxious. Erm... but I also do think it's-it's teaching me... erm... where I’m still really maintaining quite rigid control. Erm... perhaps without realising it.” (R1443)

3.3 Restrictions in Accessing Professional Support

As support services had to adapt in relation to health and safety concerns, some participants highlighted new and continued barriers to support, while one participant viewed increased online communication as beneficial.

3.3.1. Accessibility of support

All participants mentioned comparisons between personal health concerns and overall health concerns surrounding the COVID-19 pandemic when discussing their thoughts on available supportive resources. For Participant 2445, receiving medical
assistance during lockdown was a mixed experience. Compared to others who
needed medical attention and resources during this time, they believed their
situation was not as critical, but nevertheless required more attention than was
offered:

“A couple of days ago, I was in hospital and they offered me psychiatric help,
and they told me that I was technically allowed to receive it, but I wasn't 'bad
enough' to be in the psychiatric ward or be followed-up by a psychiatrist or a
psychologist. I do understand that there are much bigger problems going on, but
I felt like I wasn't 'sick enough'...Nobody should feel like they're not 'sick enough'
to be taken care of. I feel like I've kind of been let down by the whole system at
this point, and I haven't been able to talk to a therapist. I wasn't able to start a
new round of therapy because, at this point, I finished my journey with the
whole clinic...[...] And it's not very clear how I should approach my GP or how I
should try to find a new counsellor.” (R2445)

Many participants mentioned not necessarily wanting more help, as they believed
others were in greater need for support at the moment, due to COVID-19. Most
mentioned their belief that receiving more support may take away health providers
from perceived more important cases:

“To be honest, I feel lucky to have the support that I do, and, like you said, I am
also really, really aware that there are people all over the country who are
struggling with lockdown who don't have any support whatsoever. I think my
main feeling around it is, to be honest, I feel undeserving of weekly hour
long mental health support when everybody's struggling with their mental
health at the moment...I certainly don't think that I would want more.” (R4880)

Similarly, Participant 1096 referred to the resources and support others affected by
COVID-19 required, and stated that anxiety and age restrictions prevented them
from seeking additional support:

“So, yeah, it’s like the only support that I get, because of COVID-19 I daren’t ask
my GP for more support...GP, the access is there but it is much, much, much
more difficult because it has to be telephone call, there’s no text-based service. I
can’t text, I can’t email, I can’t book appointments online...It makes it much more inaccessible for me. So it exists, but it is not one I can use. I also do the fun thing of being ever so slightly too old for some of the support offered by BEAT because a lot of that is 18-25 and I am 26...” (R1096)

Conversely, some participants had different experiences regarding streamlined medical care and support services, and online options during lockdown. Compared to previous practical contact with service providers, Participant 7260 expressed preference for adapted and restructured eating disorder support services:

“I think it has been really good that support services have had to adapt to using digital technology. Before, everyone had been really quite resistant to like online communication I think, like how long has it taken for any medical records to even be digital...I think that actually there’s a few benefits in it, like no one would choose for this to happen but at the same time I hope that things don’t go back to the way they were, where everyone had to physically turn up to buildings to access a service.” (R7260)

4. Discussion

This study is the first using an in-depth interview approach with adults with mixed eating disorder presentations in the UK. Our results suggest the impact of COVID-19 lockdown in the UK can be described as a catalyst for either the exacerbation of disordered eating behaviours, or for eating disorder recovery. The findings highlighted the structures of social and functional restrictions, as well as restrictions in accessibility to professional support, to be crucial determinants of mental well-being in this group. Personal experiences of disordered eating during lockdown were seen as either facilitated or limited by these restrictions, depending on participants’ living and work situation, as well as their eating disorder progression. Predominant feelings of ambivalence towards lockdown measures were in line with participants’ feelings towards recovery. Ambivalence in eating disorder recovery has previously been described as “a state of dynamic stability” (Bell, 2013) due to conflicting motivations in long-term eating disorder pathology. Participants in this study described feeling safe or proactive while engaging in disordered eating or excessive
exercise, even if they were working towards recovery and recognised that their mental health was affected by their behaviours. Being externally restricted through the lockdown measures might have reinforced the ambivalent perception of agency in the context of disordered eating behaviours (Shohet, 2007).

Our findings partially aligned with the pathways proposed by Rodgers et al. (2020). The disruption to usual life and the resulting influence on meal patterns, routines, and physical activities had a considerable impact on the lives and eating disorder symptoms of participants. Social isolation and removal of social support led to an increased sense of loneliness and resulted in impact on accountability (Akey & Rintamaki, 2014), their routines and sense of responsibility. Participants reported having to redesign and restructure the usual aspects of their lives which heightened participants’ awareness of their behavioural intentions, seeking to reintroduce the incidental aspects of day to day life into the new mode of living during the COVID-19 pandemic. Participants’ responsibility for themselves and intentionality in planning their own actions were highlighted as being key mechanisms influencing their eating disorder behaviours.

In contrast, Rodgers et al.’s (2020) second and third pathways – concerning increasing in detrimental media exposure and health concerns were less apparent in our findings. Regarding health anxiety, participants voiced concern primarily regarding the threat of the virus for others such as the elderly or vulnerable loved ones rather than towards themselves, often feeling undeserving of professional support as others were viewed as needing it more in the pandemic context.

One key, and novel, finding of this study was the lockdown associated with COVID-19 being experienced as a catalyst for recovery from disordered eating behaviours for some, whilst be related to increased difficulties for others. Previous authors have highlighted that COVID-19 may precipitate or exacerbate disordered eating behaviours (e.g. Weissman et al., 2020). However, reported experiences in our study reveal the potential of the pandemic to improve eating disorder symptomatology through its focus on self-efficacy and risk management. Motivations for continuous
recovery were risk avoidance in the context of COVID-19, and a sense of achievement of managing without professional support. That said, not all participants were in a position to experience this; severe eating disorder pathology before and during lockdown was not associated with reported improvements in eating disorder management. Individual perceptions of negative impact were highly dependent on how participants conceptualised current eating disorder symptoms, which was generally ambivalent and differed between expected short-term and long-term impact.

A further important finding of the present study was participants’ concept of the perceived future post-lockdown. The introduction of the lockdown was followed by multiple amendments which will eventually allow UK residents to return to their workplaces. The easing of restrictions is thereby associated with as much, if not more, uncertainty as the introduction of the social distancing measures. Established lockdown routines continuously need to be adapted, which was highly anxiety inducing for most participants in our study. Future studies will have to assess the long-term impact of this uncertainty on eating behaviours and exercise routines.

4.1. Strengths and Limitations

Throughout the pandemic, recommendations and guidance rapidly changed; thus, capturing the experience of a particular period during the pandemic was challenging. The interviews were performed from 14th May until 4th June, with four interviews being conducted after the first amendment to lockdown restrictions was implemented in Scotland and England. However, lockdown restrictions were still seen as significantly impacting participants’ lives and capturing this change might have enabled us to identify additional feelings of uncertainty due to changing circumstances.

In addition, only White, predominantly female participants volunteered to take part in this study, limiting the generalizability of our research findings. Regarding pathology, we were able to interview participants with a wide range of eating
disorder behaviours. Even though most participants identified with Anorexia Nervosa, our findings are based on various clinical pictures, which enriched our analysis.

Finally, all interviews were guided by a semi-structured interview schedule to ensure consistency among interviewers and interviews, while allowing for flexibility in the data collection, depending on participants’ experiences. This, however, meant that interviews varied in length and focused on slightly different topics, if participants chose to elaborate on certain experiences more than on others.

5. Conclusion

Social, functional, and professional support-related restrictions were three main themes of underlying lockdown structures inductively identified from 10 interviews with individuals with an eating disorder. The tendencies identified related to social isolation, accountability, increased responsibility, lack of routine and structure, intentionality, and secrecy all reflect Rodgers et al.’s (2020) first pathway of the COVID-19 pandemic’s impact of disruption and restrictions to established daily routines and interactions. Exploring how existing professional support services can best adapt to help those with eating disorders manage these difficulties would be valuable going forward.
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S.B. and M.O. conducted the interviews. S.B., M.O., and I.P. transcribed the interviews and analysed the results. All authors contributed to the overall design of the study and writing the manuscript. All authors have approved the final article.

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