Adherence to limiting weight-bearing activity in patients

2

with diabetic foot ulcers: A qualitative study

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17

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- 25

26 ABSTRACT

27

28 Aims: Patients with diabetic foot ulcers are advised to limit weight-bearing activity for 29 ulcers to heal. Patients often disregard this advice although the reasons are not yet fully 30 understood. This study explored 1) patients' experiences of receiving the advice and 2) 31 factors influencing adherence to the advice. 32 Methods: Semi-structured interviews were conducted with 14 patients with diabetic foot ulcers. Interviews were transcribed and analysed using inductive thematic analysis. 33 34 **Results:** Advice regarding limiting weight-bearing activity was described by patients as 35 directive, generic and conflicting with other priorities. Rapport, empathy and rationale supported receptivity to the advice. Barriers and facilitators to limiting weight-bearing 36 37 activity included: demands of daily living, enjoyment of exercise, sick/disabled identity 38 and burden, depression, neuropathy/pain, health benefits, fear of negative 39 consequences, positive feedback, practical support, weather, and active/passive role in 40 recovery. Conclusions: It is important that healthcare professionals pay attention to how limiting 41 42 weight-bearing activity advice is communicated. We propose a more person-centred 43 approach in which advice is tailored to individuals' specific needs with discussion around 44 patient priorities and constraints.

45

46 Keywords: patient adherence, motivation, diabetic foot ulcer, communication,

47 physician-patient relations

48 Key Messages

- Patients with diabetic foot ulcers find adhering to advice to limit weight-bearing
 activity challenging
- This study explored patients' experiences of being advised to limit weight-bearing
 activity and factors influencing adherence to this advice.
- Semi-structured interviews were conducted with 14 patients with diabetic foot
 ulcers
- Footrest was generally viewed as impractical due to the constraints of everyday
 life and conflicting health priorities.
- Patients require tailored support to manage physical activity and ulceration and
- are more receptive to advice when delivered with rapport, empathy and the
- 59 rationale behind recommendations explained.

60 1. INTRODUCTION

Individuals with diabetes are at increased risk of developing lower limb complications 61 62 such as foot ulcers. Diabetic foot ulcers (DFUs) are estimated to affect up to 25% of 63 patients with the disease, with a high rate of recurrence (40% within 1 year).(1,2) Some 64 ulcers never heal and can lead to severe complications including infection, amputation, 65 hospitalisation, and premature mortality.(3) Both ulcers and amputations are associated 66 with reduced quality of life with nearly half of patients with DFUs experiencing symptoms of depression.(4,5) Moreover, the financial burden of DFUs is significant, costing the NHS 67 68 an estimated £837-£962 million a year.(6) The likelihood of developing a DFU, and the 69 subsequent healing process, is affected by patients' self-management strategies and 70 adherence to clinical advice. Thus, research exploring factors influencing DFU patient 71 adherence to treatment recommendations is needed.

The International Working Group of the Diabetic Foot recommend patients with preulcerative signs or active ulceration limit weight-bearing activity (WBA) to aid healing.(7,8) DFUs are caused by a combination of factors, the most common pathway being excessive mechanical stress on insensate neuropathic plantar tissue.(2,9) Weightbearing is thought to increase the cumulative plantar tissue stress, which in turn, can increase the risk for foot ulceration.(9) Consequently, current practice is for healthcare professionals to recommend patients reduce WBA for an ulcer to heal.(7)

Adherence to treatment advice for patients living with DFUs has been consistently low.(10-12) Qualitative research exploring the experiences of patients living with DFUs has reported patients' lack of engagement with advice to limit WBA on the ulcerated foot.(13-18) However, the reasons for patients' non-adherence to limiting WBA advice are not yet fully understood.

Previous research on adherence to treatment for patients living with DFUs have focused on barriers and facilitators to footcare advice in general (e.g., inspection of feet, appropriate footwear, attendance at foot screening appointments), rather than WBA

specifically. A qualitative meta-synthesis reported factors that influence adherence to foot care behaviours include knowledge of appropriate foot care, perceived importance, practical support from family and friends, prior experience of consequences (e.g., amputation), and communication style of healthcare providers.(19) Some of these factors may play a role in patients' adherence to limiting WBA but there may also be other factors specific to this behaviour that have not yet been explored.

93 The recommendation to limit WBA directly opposes the general advice consistently given for patients with diabetes to be physically active.(20) Engaging in regular activity is 94 95 associated with many benefits for patients with diabetes including, but not limited to, 96 weight control, reduction in blood pressure, improved glucose control, reduced risk of 97 cardiovascular disease, and improvements in mental health and health-related quality of 98 life.(21) Thus, long-term adherence to limiting WBA may be particularly problematic for 99 those who experience high recurrence of ulceration. To our knowledge, no research has 100 explored DFU patients' experience of receiving advice to limit WBA and perceptions of 101 managing seemingly conflicting advice. Exploring patients' views and experiences may 102 inform our understanding of the factors that influence adherence. Findings could be used 103 to assist healthcare professionals in supporting patients to limit WBA.

Therefore, the aim of the study was to 1) explore patients' experiences of receivingadvice to limit WBA and 2) factors that may influence adherence to this advice.

106

107 **2. METHODS**

108 2.1. Study Design

In depth qualitative semi-structured interviews were conducted with patients with DFUs.
The study was approved by the East Midlands – Derby Research Ethics Committee (REC
Number 18/EM/0162) and written informed consent gained from all participants.

112 2.2. Recruitment

113 Participants were an opportunistic sample recruited from a specialist Diabetes Foot Clinic 114 in the East Midlands, UK. To be eligible individuals needed to be a patient at the diabetes 115 clinic, aged 18 years and over, and be able to communicate in English. Potential 116 participants were initially approached by a member of the patient's usual care team who 117 provided patients with detailed written information pertaining to participation in the study. 118 It was explained that participation was entirely voluntary and that their treatment and 119 care would not be affected by their decision. On consenting to take part the patients' 120 contact details were shared with the researchers to organise the interview.

121 2.3. Data collection

Interviews were conducted by two authors (JEH and CEH) both trained and skilled in qualitative interviewing. Most interviews took place face-to-face in a private room within the Diabetes Foot Clinic. One interview was conducted via phone as the patient was unable to attend in-person. All interviews were audio-recorded and lasted no more than 1-hour.

127 Interviewers utilised a semi-structured guide which was formulated from best practice 128 guidance.(22) The guide included questions covering patients' views on and experience 129 of being advised to limit WBA (e.g., how the advice was communicated, how relevant 130 they felt the advice was for them, anything that was said or done that made them more 131 or less likely to limit their activity, how confident they feel in limiting their WBA, the 132 things that help them to reduce their WBA, and things that make it difficult for them to 133 limit their WBA).

134 2.4. Data analysis

Interviews were transcribed verbatim and anonymised. Data were analysed in NVivo (version 12) using inductive thematic analysis following the recommendations of Braun and Clarke.(23) Analysis was iterative, occurring concurrently with interviews so that sampling continued until saturation was reached with no new information emerging.(24) Authors familiarised themselves with the data through 'active reading' of the transcripts

and noted initial analytic observations. Due to the explorative nature of the study, initial
codes were developed using an inductive approach. Coded data was reviewed and
collated into potential themes. Themes discussed between authors and refined until
consensus was reached. Detailed field notes and a clear audit trail of analytic decisions

144 were kept to maximise transparency and ensure credibility and quality.

145

146 **3. RESULTS**

147 Fourteen participants with DFUs (13 male, 1 female) were interviewed in 2019. The 148 gender split broadly reflects the clinical population, with DFUs affecting men more than 149 women.(25) All participants were White British with ages ranging from 28-81 (mean 150 age=60.36, s.d.=15.07). Two participants were diagnosed with type 1 diabetes and 12 151 with type 2 diabetes mellitus. Mean BMI was 32.42 (range 22.53-48.70). Marital status 152 varied: 3 reported as single, 2 partnered, 6 married, 1 divorced, and 2 widowed. 153 Patients had been receiving treatment at the specialist Diabetes Foot Clinic for an 154 average of 25 months (range 3-72 months). Interviews lasted on average 37 minutes 155 (range 23-59).

3.1. Patients' experiences of receiving advice to limit weight-bearing activity

157 Two themes and five sub-themes were identified. See Table 1 for a summary.

158

159 [TABLE 1]

160

161 **3.1.1. Advice**

162

163 *3.1.1.1. Directive*

Patients described repeatedly being advised to rest and limit their WBA. Such advice was
often viewed as 'nagging', particularly when delivered in a directive/paternalistic
manner.

167	
168	"The podiatrists, the doctors, everyone that's involved with my care, they all say
169	the same to me. You must, must limit your activitySo from the word go all the
170	advice I received at each consultation, it's always the same: are you limiting your
171	activity? Yeah, yeah doctor, as much as I can, blah-blah-blah."
172	
173	When advice was delivered in a controlling way (e.g., using phrases such as "You
174	must") patients reported feeling like they were being told off which made them less
175	likely to want to adhere.
176	
177	"Not putting my feet up. Always get told offIf people tell me to do something, I
178	just do the oppositethe more they tell me to do something I'm less likely to do
179	it."
180	
181	Patients preferred it when healthcare professionals explained the reasoning behind their
181 182	Patients preferred it when healthcare professionals explained the reasoning behind their recommendations.
182	
182 183	recommendations.
182 183 184	"They're not telling you; they're explaining it to you, which is a subtle difference,
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195	
196	"Yes, she described it as little things are trying to mesh and knit together, and
197	every time you put your foot down it's like smashing them apart again. But we try
198	to not smash them too much because they're trying to knit togetherBut also, she
199	asks the question, shall I go on telling you these things?"
200	
201	3.1.1.2. Generic
202	Treatment advice was described by many as generic and 'one-size-fits-all' with a lack of
203	discussion around individual needs and lifestyles.
204	
205	"The advice at the clinic is general knowledge really. Keep off your footthey
206	haven't got time to think about what's going on in my lifeThey seem to tar
207	everybody with the same brush."
200	
208	
208	Patients felt they would benefit from advice being tailored to their specific needs via a
	Patients felt they would benefit from advice being tailored to their specific needs via a process of collaborative problem-solving.
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209 210 211 212 213 214 215 216 217 218 219	 process of collaborative problem-solving. "I think certainly the more individual you can tailor the advice, the more it works for individual peopleIt's a case of me sort of saying well hang on, what about this? What you're looking for is a suggestive process rather than an instructive process. What works for me almost certainly doesn't work for someone else." 3.1.1.3. <i>Conflicting</i> advice Patients reported receiving conflicting advice, with doctors recommending they exercise to manage their diabetes or other health conditions (e.g., osteoarthritis, depression) and

223 "Because I've got osteoarthritis my doctor tells me to keep active because 224 it stops me from ceasing up. But then the hospital are telling me to 225 offload." 226 227 Some patients expressed being active was more important to them than the ulcer 228 healing. 229 230 "The dilemma is that the podiatry want me to rest, my doctor wants me to exercise, what do I do? It seems to me that that's because they'll be some 231 232 weight loss with that [exercise] as well as lots of peripheral gains. If the 233 other side of the coin is that it's not helping my feet, well I'll probably 234 have to live with it." 235 236 237 3.1.2. Relationships with health care providers 238 239 3.1.2.1. Rapport 240 Participants explained that when healthcare professionals showed interest in them 241 and their life it made them feel supported and understood. 242 "We've been coming here for such a long time now, she [the podiatrist] shows a 243 regular interest...they're the little things which make us smile, make us feel 244 wanted and cared for. Understanding you as a person rather than just caring 245 about what is going on with your foot." 246 247 Participants reported being more willing to open-up about their challenges with 248 limiting WBA when there was a sense of connection with healthcare professionals. 249 "I think just knowing that someone's there as well and you can talk to them. They're 250 easily approachable."

251				
252	3.1.2.2. Empathy			
253	Some patients reported a lack of empathy in discussions regarding limiting WBA. They			
254	explained the challenges of putting the advice into practice in their everyday life was not			
255	always acknowledged.			
256				
257	"Obviously they know I live alone, and I've got to work but sometimes I think they			
258	don't live in the real worldJust go home and put your foot up, doesn't fit with my			
259	lifestyle."			
260				
261	However, others felt that practitioners did take time to recognise the difficulties patients			
262	face in adhering to treatment recommendations.			
263	"There's empathy there, and they do understand what I'm going through. They			
264	don't go over the top, it is what it is, but they do understand that there is a lifestyle			
265	change in the fact that you've got something like this."			
266				
267	3.2. Factors influencing adherence to advice to limit weight-bearing activity			
268	Themes were organised into barriers and facilitators to limiting WBA (see Table 2).			
269				
270	[TABLE 2]			
271				
272	3.2.1. Barriers to limiting weight-bearing activity			
273	3.2.1.1. Demands of daily living			
274	Many patients understood that limiting WBA would likely aid healing but expressed that			
275	it was just not practical. Participants described needing to engage in various activities of			

276 daily living which involve WBA (e.g., work, shopping, driving, cleaning, gardening,

attending hospital appointments and caring for family members and pets).

278 "I know I've got to keep off it...I try and keep off it as much as I can, but I've got
279 stuff to do: cook, clean the home, you know."

Financial hardship was an insurmountable barrier for some. Participants working in jobs that require WBA (e.g., manual labour, shop assistant) described not being able to afford to change job or reduce hours worked.

283 "I mean they're [healthcare professionals] always telling me to stop, to not go to
284 work, but obviously I need the money. I've got to earn... I'm in a job that doesn't

pay fantastically. I don't get paid if I'm off sick, apart from statutory sick pay.

And the last time I was off, I had a shock, I nearly lost my house because I

287 couldn't pay the mortgage."

288

289 *3.2.1.2. Enjoyment of exercise*

For those who being active was part of their identity and considered essential for health and well-being, being told to rest, and limit WBA conflicted with their core values.

292

293 "You've got to be active haven't you; otherwise rigor mortis sets in."

294

A few patients acknowledged that non-WBA could provide a means for remaining active but these types of activities (e.g., chair-based exercise) did not appeal. The preference was for physical activity which is enjoyable and/or purposeful.

298

299 "I suppose there are exercises you can do lying down. But we tend to do
300 activities like taking the dog out, so that means quite a bit of walking. So
301 that's contrary to what I should be doing."

302

303 *3.2.1.3. Sick/disabled identity and burden*

Being viewed as someone who was chronically ill and limited in their ability to get around was not an identity that patients wanted to associate with. Resting was often attempted in the short-term but not viewed as a viable long-term option.

307 "Well, they just tell me to sit down and rest it. You know, sit at home and
308 watch whatever's on TV if you like. Which I did for a couple of weeks, but
309 that's not me."

Family relationships and caring responsibilities changed because of patients developing
 foot ulcers. Patients reported loss of independence and concern regarding overburdening

312 others.

313 "It's just not that easy dragging folk in, people do help us, don't get me

314 wrong. But other people have got busy lifestyles as well haven't they,

315 can't just down tools and help us all the time."

316

317 3.2.1.4. Depression

318 One patient, when asked about the importance of limiting WBA, explained that it 319 depended on how they were feeling emotionally, with their depression sometimes 320 leading them to want to weight-bear to make the ulcer worse.

321 "That's down to my moods and the way that I feel in myself and not

322 wanting to feel well really...The depression is manifesting itself in ways

323 that I want to punish myself I suppose."

324 Another patient, diagnosed with clinical depression, described a conflict between

knowing that they need to rest for their ulcer to heal and remaining active for

326 their physical and psychological well-being.

- 327 "From the first moment I saw a podiatrist I've been told to rest and put
- 328 my feet up and from day one I've ignored it. I heard it and it makes sense
- but looking at my big picture it makes less sense...I'm suffering from a
- clinical depression, being so weak and feeble doesn't do much for my
- 331 spirit."
- 332

333 *3.2.1.5. Neuropathy*

334 Some patients explained that because they do not feel pain from the ulcer it does335 not seem as high a priority to limit WBA.

- 336 "If it hurt more, I would think it was more important, but it doesn't. And
- that is probably the biggest problem. If it hurt, your motivation on
- 338 accelerating the healing process would go up."
- 339

340 *3.2.1.6. Passivity*

- 341 Many patients described adopting a passive role in management of their
- 342 condition, relying on healthcare professionals to fix their ulcer. This seemed to be
- the case particularly for those who had had ulcers for a long-time and given uphope.
- 345 "I just come, get treated, go home. I've been doing it for so long that I don't
 346 think I attend to it. I don't think I'm that bothered...so from that point of
 347 view I keep walking."

348

349 **3.2.2. Facilitators to limiting weight-bearing activity**

350 *3.2.2.1. Health benefits*

351 Belief that limiting WBA will lead to ulcer healing appeared to be related to

352 patients' willingness to adhere.

353 "Knowing that I'll get fixed if I do as I'm told... Because less weight bearing on my
354 foot, quicker healing recovery. That's it really."

355

356 *3.2.2.2. Fear and/or experience of negative consequences*

357 Seeing the negative implications from not limiting WBA made some re-assess

358 their activity.

359 "Just because I feel it's getting worse. So, like I say you just look at it, reality

360 check, and you think well, it's going to get even worse if you're going to keep doing

361 it. So, you think right OK enough is enough, you need to stay off your feet."

362 Fear of consequences (e.g., amputation) appeared to be a strong motivating

363 factor.

364 "There's the thought in the back of my mind, well if I don't get this right then, you
365 know, in my worst-case scenario I might have to have my toe amputated. And I
366 don't particularly want that, so."

Patients also talked about the long-term implications of not limiting WBA to include notbeing around to care for loved ones.

369 "My partner says well I want you here for at least another 20, so I need to see if I
370 can achieve that. And for my grandson's sake as well!"

371

372 *3.2.2.3. Lifestyle adjustments*

373 Participants described a process of adjustment using trial and error. Awareness of

374 which activities aggravate the ulcer was described as key.

375 "The biggest thing is awareness. If you know what you're doing is wrong, then
376 you can limit and adjust. Give you an example, one of the things I used to do is
377 merchandising, which involves hoarding cages of stock...So you look at it and
378 think well, probably best if I don't do that...It's a constant balance."

This participant went on to describe how living with an ulcer involves planning ahead,taking time to consider activities and the impact they may have.

381 "It's all about risk assessment...It's just a case of taking what I'm offered and
382 analysing it far more than I ever used to. Now it's more a case of when somebody
383 says to me, well we want you to do this, you ask them what's involved and then
384 make a balanced decision. Rather than just turn up and do it, no big deal."

385 Identifying ways of adapting their activities to minimise the time spent on their feet was386 described as requiring conscious effort and problem-solving.

387 "So not just go for a walk for the sake of it. So yeah, I try to be mindful. If I go to
388 a shop, rather than using the steps, use the elevator or the escalator. So just trying
389 to think ahead how to minimise what I will need to do."

390 The process of adjusting behaviours to reduce activity was described as time consuming391 and requiring patience.

392 "Yeah, it can be difficult, you have to think about a lot of different things. Can you

do this, can you do that? So, putting it into practice, it is harder said than done.

394 It can be done, but it just takes a lot of time and patience. And I think that's the395 key thing, patience."

396 Some participants found identifying activities they could do while resting, that

they find enjoyable and purposeful, helped with adapting their lifestyle.

398 "What I've done is I've filled up my life with learning and education whilst I'm
399 waiting for my foot to heal"

400

401 *3.2.2.4. Positive feedback*

402 Confirmation from healthcare professionals that what patients are doing to limit pressure403 on their ulcer is working was described as a key facilitator.

404 "When they do the camera, take the photograph, and give you feedback, it is useful 405 because if it is positive, even if it's telling you like half a mil that it's reduced, that 406 does give you a bit of a, like hey great, that's good then. I need that confirmation 407 that I'm getting better."

408

409 *3.2.2.5. Practical support and reminders*

410 Practical support (e.g., cooking, shopping, driving) from family and friends was411 described as vital.

412 "Well, my sister lives locally, and I've got friends and family that can help.

413 But it's a case of mainly deliveries, or I can get people to drive me places."

414

415 A key component of the support from family involved provision of verbal

416 reminders and recognition of the importance of patients limiting their WBA.

*I'd say family and friends, they know how important it is for me to stay off my
feet. They realise if I don't stay off my feet, I could end up losing my feet. They
just say how much walking have you done today? They ask me little questions like
that so that I think well, yeah, I have been on my feet quite a bit today, for the
rest of the afternoon I need to stay off my feet."

422

423 *3.2.2.6.* Cold weather

The weather was described to influence WBA. Participants explained they are more likelyto stay indoors and rest in winter months when it is cold and dark outside.

*Recently, we've had some days of solid rain. Well, those two days I couldn't go
outside, I did more sitting down. But you've got to organise the weather to get me
to do that."

429 In the summer months individuals expressed a desire to go out and about more.

430 "I think I've been more motivated this month just because it's cold weather and
431 you want to try and stay as warm as possible. This summer just gone; I think I
432 was less motivated than ever before, just because it's warm weather. You just want
433 to go out and do your day-to-day stuff."

434

435 *3.2.2.7. Pain*

436 A participant without neuropathy described how pain from his ulcer acted as a437 reminder not to put pressure on it.

438 "If I get a bit of pain, I will stop. Yeah, well I just put my foot up for half439 an hour, not very long. As soon as the pain's gone I start walking about."

440

441 *3.2.2.8. Depression*

442 Depression acted as a facilitator for one individual who explained they are more likely to 443 rest and limit WBA when they feel low.

444 "Although because I've been depressed, I do go to bed an awful lot, lie in bed 445 watching the telly or go to sleep, I don't have to think about what's going on."

446

447 3.2.2.9. Active role in recovery

448 Participants who expressed feelings of empowerment and self-determination in their

449 motivation were more willing to limit their WBA.

- 450 "At the end of the day all I want to do is get my foot fixed, but there's only me451 that can do it. It's me that's got to make the change."
- 452

453 **4. DISCUSSION**

454 This study aimed to explore 1) DFU patients' experiences of receiving advice to limit 455 WBA and 2) factors influencing adherence to this treatment advice. Interviews with 456 patients revealed advice to limit WBA to be perceived to be delivered in a directive 457 manner via the use of controlling language (such as, 'you must', 'you need to'). Use of 458 controlling language by significant others is understood to be an antecedent of poor-459 quality (or non-self-determined) motivation driven by internal or external contingencies 460 such as pressure or guilt.(26) Thus, the language used to convey the treatment 461 recommendation may influence patients' experiences of receiving the advice and, in 462 turn, their motivation for adhering. The directive approach also included regular 463 repetition of the recommendation. Previous research exploring podiatrists' views on 464 patient adherence to footcare behaviours, revealed podiatrists to believe that regular 465 reiteration of advice increased the likelihood of patient adherence.(17) The present 466 findings suggest that may not be the case, with regular repetition of generic advice 467 viewed as 'nagging' and not considered conducive to engagement.

468 Findings revealed patients viewed advice to limit WBA as generic and 'one-size fits all' 469 rather than that which is specifically tailored to their individual circumstances. This is 470 aligned with findings by Searle who reported advice given by podiatrists to be generic 471 rather than tailored specifically to patients' needs.(17) The quality of the relationship 472 with healthcare professionals was also found to be important factor within patient 473 experiences of being advised to limit WBA. Patients appeared to view advice to limit WBA 474 more favourably when a rapport was already established with healthcare professionals. 475 Similar experiences were described by Coffey et al. in their qualitative meta-analysis, 476 which reported development of rapport and empathy as important factors in DFU patient

477 engagement and willingness to follow treatment advice.(19) Therefore, the findings from
478 previous research and the current study suggests healthcare professionals may need
479 support to develop skills in person-centred care.

480 The main barrier identified in relation to limiting WBA was the practicalities of daily life. 481 Patients reported various activities of daily living (e.g., work, shopping, caring 482 responsibilities) that affected their ability to limit WBA. Searle also reported patients to 483 view footrest as impractical due to the constraints of everyday life.(17) Patients were 484 often left to work out for themselves how to implement the recommendation into their 485 everyday life. Patients described making lifestyle adjustments as a process of trial and 486 error and requiring forward planning. This is a strong indicator that support from 487 healthcare professionals in the form of collaborative problem-solving may be beneficial 488 to aid the behaviour change process.

Findings from the current study support previous research detailing barriers and motivators to foot self-care behaviours.(19,27) For example, the present study also found patient understanding of the importance, prior experience of consequences (e.g., amputation), practical support from family and friends and a sense of personal responsibility/empowerment to be key influencing factors. This provides further support for targeting these aspects within routine clinical care.

495 This study advances our understanding of DFU patient adherence by identifying aspects 496 which are unique to the behaviour of limiting WBA. For instance, patients reported 497 enjoyment and/or importance of exercise as a barrier to limiting WBA. Regular physical 498 activity is essential for the management of diabetes and other health conditions(21) and 499 patients described the advice to limit WBA as conflicting with other health priorities. 500 Although the advice given to patients is to limit WBA, this does not necessarily mean 501 stopping exercise altogether.(7) There is scope therefore for further research to identify 502 how to most effectively support patients' engagement in non-weightbearing alternatives 503 as part of the management and treatment plan.

504 Findings revealed depression to have a potentially complex relationship with adherence 505 to limiting WBA. A recent meta-analysis showed nearly half of DFU patients report 506 symptoms of depression.(5) Individuals with depression have been found to be less 507 physically active and engage in higher levels of sedentary behaviour compared to aged-508 matched controls.(28) Our findings partially concur with this, with some patients 509 reporting feelings of depression to make it more likely for them to rest and limit WBA. 510 However, other patients described their depression manifesting in a way which led to 511 them engaging in WBA to either to improve their psychological well-being or purposefully 512 cause themselves injury. Emotional support for such patients is essential to reduce the 513 risk of further self-harm.

514

515 4.1. Practical implications

516 The findings of the present study highlight the need for improved patient support with 517 managing physical activity and ulceration. Patients have reported dissatisfaction with the 518 way in which treatment advice to limit WBA is communicated. Future research is needed 519 to identify how to most effectively support health professionals to communicate in a way 520 which empowers patients' to actively engage with treatment advice and adhere to 521 recommendations to limit WBA. This may involve taking time to develop a rapport and 522 connection with patients and delivery of advice using non-controlling language and 523 rationales. Such communication strategies have been found to promote autonomous 524 motivation for health behaviours.(29,30)

Patients expressed a preference for advice which is person-centred and tailored to individuals' specific needs. For many patients being told to rest and limit WBA conflicted with either their lifestyle or identity which created a sense of resistance and lack of adherence. Rather than the generic message being to 'rest and limit WBA' it may be that a shift to a more personalised approach involving discussion around 'safe physical activity levels' for individual patients would be beneficial.(31) Such an approach could be

- 531 consistently communicated by all healthcare professionals, reducing patient confusion
- 532 resulting from conflicting advice.
- 533

534 **5. Conclusion**

- 535 To our knowledge, this is the first study to explore DFU patients' experience of receiving
- advice to limit WBA and perceptions of managing such conflicting advice. Various
- 537 barriers and facilitators to limiting WBA have been identified and can be used to inform
- 538 practice. An understanding of patient priorities and constraints may aid healthcare
- 539 professionals in developing person-centred treatment plans.

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Table 1. Themes and subthemes

Theme	Subtheme
3.1.1. Advice	3.1.1.1. Directive
	3.1.1.2. Generic
	3.1.1.3. Conflicting
3.1.2. Relationship	3.1.2.1. Rapport
	3.1.2.2. Empathy

Table 2. Themes rel	lating to barriers	and facilitators to	limiting WBA
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Barriers		Facilitators
5.1.1.1.	Demands of daily living	3.2.2.1. Health benefits
5.1.1.2.	Enjoyment of exercise	3.2.2.2. Fear and/or experience of
5.1.1.3.	Sick/disabled identity and	consequences
	burden	3.2.2.3. Lifestyle adjustments
5.1.1.4.	Depression	3.2.2.4. Positive feedback
5.1.1.5.	Neuropathy	3.2.2.5. Practical support and reminders
5.1.1.6.	Passivity	3.2.2.6. Cold weather
		3.2.2.7. Pain
		3.2.2.8. Depression
		3.2.2.9. Active role in recovery