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**Psychological Factors in Symptom Severity and Quality of Life in Raynaud's
Phenomenon**

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Abstract

Background Despite emotional stress being recognised as a key trigger for Raynaud’s phenomenon episodes, research in the area is still in its infancy.

Aims This study investigated the role of psychological factors relating to symptom severity and quality of life, and differences between Raynaud’s types (primary and secondary) to further inform the development of intervention in this field.

Method A cross-sectional design was used. Two hundred and ten adults with Raynaud’s completed an online questionnaire measuring stress, anxiety, depression, anxiety sensitivity, beliefs about emotions, symptom severity and quality of life.

Results Primary and secondary Raynaud’s groups differed in anxiety ($p < .004$), symptom severity ($p < .001$) and quality of life ($p < .001$). Stepwise multiple regressions indicated anxiety and Raynaud’s type explained 23% variance in hand symptom severity ($p < .001$); anxiety, Raynaud’s type and anxiety sensitivity explained 29% variance in symptom severity (global impact, $p < .001$); depression, Raynaud’s type and anxiety sensitivity explained 32% variance in quality of life ($p < .001$).

Conclusions Results highlight the importance of psychological factors in Raynaud’s phenomenon, indicating possible targets for treatment. Interventions such as cognitive behavioural therapy, which target both physical and psychological wellbeing, bear some promise as an adjuvant therapy for this group.

Keywords: Raynaud’s Phenomenon, quality of life, symptoms, anxiety, depression, psychological distress.

51 Raynaud's phenomenon is an intrusive condition that causes vasospastic episodes in
52 the extremities, usually in response to cold, sudden temperature changes or emotional stress,
53 which can induce pain and paraesthesia, compromising hand function (Shapiro & Wigley,
54 2017). The condition is either primary and idiopathic, or secondary to an underlying
55 condition, such as scleroderma where more than 95% patients have Raynaud's (Meier et al.,
56 2012). Primary Raynaud's affects approximately 5% of the general population, although rates
57 vary by country and population (Garner et al., 2015).

58 Although most Raynaud's episodes are precipitated by cold exposure, studies have
59 shown that emotional stress triggered approximately a third of episodes and that thematically
60 relevant stressors (e.g. losing gloves in a snowstorm) are particularly important (Freedman &
61 Ianni, 1983, 1985; Hughes et al., 2015). Raynaud hypothesised that this response is due to
62 overactivity of the sympathetic nervous system in Raynaud's patients, which exaggerates
63 vasoconstriction via the release of norepinephrine (Fardoun et al., 2016; Freedman & Ianni,
64 1983). An inability to habituate to stressful stimuli has been alternatively suggested to explain
65 repeated excessive vasoconstriction in Raynaud's patients (Edwards et al., 1998). Affective
66 factors can trigger or exacerbate symptoms in other episodic and inflammatory conditions,
67 demonstrating the interaction between psychological and physical functioning (Harth &
68 Nielson, 2019; Marmura, 2018).

69 The National Institute for Health and Care Excellence (NICE) recommend reducing
70 stress and retaining warmth if this is a trigger as a first-line management of the condition but
71 there are currently no interventions in line with these recommendations or other lifestyle
72 recommendations (Daniels et al., 2018; NICE, 2022). A systematic review by Daniels and
73 colleagues (2018) which examined the efficacy of behaviour change interventions concluded
74 that there was not currently enough evidence to support or refute behaviour change
75 interventions in Raynaud's due to low quality studies, but posited that there remained a

76 strong case to further a psychological understanding of the condition, which could provide
77 targets for intervention.

78 Evidence-based non-pharmacological interventions may provide a more acceptable
79 alternative to pharmacological treatments which are commonly ineffective or cause adverse
80 side effects, such as headaches, dizziness and oedema (Choi, & Henkin, 2021).

81 However, prior to the development of such an intervention there is a need to
82 operationally define the terms used to describe triggers of an episode, “emotional stress”
83 being the most commonly used in the literature, and suggests a combination of anxiety and
84 stress, which are also used interchangeably in the field. Stress can be characterised as a
85 ‘response’ to pressure caused by an *external* trigger, whereas anxiety is more of a persistent,
86 excessive worry that remains even without the stressor being present (American
87 Psychological Association, 2019). Brown et al. (2001) investigated both constructs and
88 concluded that anxiety, rather than stress, predicted frequency and severity of attacks. The
89 term “emotional distress” is also used, which more broadly includes depression (Evers et al.,
90 2011; Newton et al., 2012). Depression is common across many physical health conditions
91 and often associated with poorer outcomes (Daré et al., 2019); depressed patients may have
92 lower treatment adherence and less likely to take care of themselves by keeping warm or less
93 willing to seek help for symptoms, thereby increasing symptom severity (DiMatteo et al.,
94 2000). The non-specificity in the literature creates confusion and resolving this is likely to be
95 pivotal in the development of appropriate non-pharmacological interventions for this group,
96 where there are currently none.

97 As found in other medical conditions, Raynaud’s symptoms have a detrimental
98 influence on quality of life, with impact on everyday activities and the requirement of
99 adjustment and adaptation (Murphy et al., 2021; Pauling et al., 2018). However, current
100 research has not gone beyond simply showing that Raynaud’s patients have poorer quality of

101 life than healthy individuals, and that quality of life is lower in secondary Raynaud's
102 compared to primary (De Angelis et al., 2008; Fábíán et al., 2019). The latter of which may
103 be attributed to the absence of underlying pathology and lesser severity in primary Raynaud's
104 (Shapiro & Wigley, 2017). Further research is needed to address this knowledge gap;
105 improving quality of life is considered at least as important as treating symptoms (Shapiro &
106 Wigley, 2017).

107 There has been a growing interest in the psychophysiology literature about the role of
108 anxiety-related constructs in health conditions. Anxiety sensitivity, the fear of anxiety
109 symptoms (physical and emotional) and believing they may cause illness, harm or
110 embarrassment, is one such construct (Horenstein et al., 2018). Anxiety sensitivity may be
111 particularly relevant to Raynaud's patients due to the symptomatic presentation (tingling,
112 numbness) and visible nature of the symptoms (triphasic colours). An overlapping yet distinct
113 construct with anxiety, anxiety sensitivity has been independently associated with
114 symptomology and quality of life in other conditions (Asmundson et al., 2000; Smitherman et
115 al., 2014). Anxiety sensitivity has also been reported to affect quality of life by perpetuating
116 anxiety, depression and avoidance of physical and mental health-promoting activities
117 (Bernstein et al., 2019; Ouimet et al., 2016); anxiety sensitivity may inadvertently increase
118 symptom severity and worsen quality of life through increased fear and avoidance.

119 Understandable fear of anxiety may give rise to negative beliefs about experiencing
120 and expressing emotions, a factor reported to be associated with adverse health outcomes
121 (Bowers & Wroe, 2016; Brooks et al., 2017). Consistent with this hypothesis, a qualitative
122 study reported that patients with scleroderma, most of whom have Raynaud's, reported
123 coping with distress by actively suppressing upsetting thoughts and feelings and were
124 reluctant to seek support (Newton et al., 2012). These avoidant strategies are likely to result

125 in reduced social support and helpful coping strategies, serving to induce or maintain low
126 mood (Bowers & Wroe, 2016; Ouimet et al., 2016).

127 Biopsychosocial models, which explain conditions as a complex interplay between
128 biological, psychological and social factors as seen in Raynaud's, have been increasingly
129 used to explain symptomology in physical health problems and promote a multidisciplinary
130 approach to treatment (Greenen & Dures, 2019; Miaskowski et al., 2020). Efficacious and
131 acceptable interventions in behavioural medicine which draw on this model, such as
132 cognitive behavioural therapy, are well placed to be adapted for use in this group. Based on
133 the notion that thoughts, feelings, behaviour and physiology are interlinked, evidence
134 supports use in similar conditions such as inflammatory arthritis (Marques et al., 2021). Such
135 an integrated approach has the potential to improve care by broadening intervention options
136 and optimising efficacy of treatment (Daniels & Turner-Cobb, 2017).

137 The study seeks to address gaps in the literature that could inform future treatment
138 development. Specifically, the relative impact of psychological factors on symptom severity
139 and quality of life in Raynaud's phenomenon, with a view to identify possible targets for
140 intervention.

141 **Method**

142 **Participants and Procedure**

143 Cross-sectional online questionnaire data was collected from adults with Raynaud's
144 using Qualtrics software, recruited via snowballing techniques on social media and two
145 associated charities (Scleroderma & Raynaud's UK and Raynaud's Association). Inclusion
146 criteria stipulated only adults (18+) who self-identified as having either primary or secondary
147 Raynaud's be included in the study sample. After reading the information sheet, participants
148 completed an informed consent form before moving on to the questionnaires battery.
149 Participants could withdraw by exiting the survey before the end. Data was collected between

150 9th June and 7th July 2020, early on in the Coronavirus infectious disease (CovID-19)
151 pandemic. Sampling took place over a limited four-week period to ensure stability of the
152 relative temperature and weather. Ethical approval was granted by the Psychology Research
153 Ethics Committee at University of XXXX (ref: 20-114).

154 Of the 269 who participated, 59 participants were removed due to incomplete data or
155 failing to meet age inclusion criteria, leaving a final sample of 210 ($n = 92$ primary; $n = 101$
156 secondary Raynaud's). Average time since diagnosis was 18.35 years ($SD = 14.60$) and mean
157 age was 47 years old ($SD = 13.63$). The sample was mostly female (94.3%), white (94.8%)
158 and either married/partnered (71.4%), with 55.2% having an education level of Bachelor's
159 degree or higher and only 6.2% were current smokers.

160 **Measures**

161 **Independent Variables.** The 21-item Depression, Anxiety and Stress Scales (DASS-
162 21; Lovibond & Lovibond, 1995) contains three 7-item subscales measuring depression,
163 anxiety and stress. Participants rate how much each statement (e.g. *I found it hard to wind*
164 *down*) applied to them over the previous week and relevant item scores (0-3) are summed and
165 multiplied by two to calculate subscale scores. The developers have recommended cut-off
166 scores for "normal", "mild", "moderate" "severe" and "very severe" that correspond to each
167 subscale (Lovibond & Lovibond, 1995). Internal consistency was good or acceptable for
168 stress ($\alpha = .88$), anxiety ($\alpha = .71$) and depression ($\alpha = .92$) subscales in the current study. The
169 total scale and subscales have been validated (Antony et al., 1998).

170 The 16-item Anxiety Sensitivity Index (Reiss et al., 1986) measures anxiety
171 sensitivity. Participants responded to items such as "*Unusual body sensations scare me*"
172 using a 5-point Likert scale (0 = "very little" to 4 = "very much"). Item scores can be
173 summed to produce a total score. Scale items were internally consistent here ($\alpha = .91$) and
174 validity has been established (Peterson & Plehn, 1999).

175 The 12-item Beliefs about Emotions Scale (Rimes & Chalder, 2010) measures beliefs
176 about the unacceptability of experiencing and expressing emotions, with items such as “*I*
177 *should be able to control my emotions*”. Participants respond using a 7-point Likert scale (6 =
178 “totally agree” to 0 = “totally disagree”). The scale showed strong internal consistency within
179 this sample ($\alpha = .93$) and has good validity (Rimes & Chalder, 2010).

180 **Dependent Variables.** Due to the lack of suitable outcome measures for this group
181 (Daniels et al. 2018), it was necessary to use two symptom severity measures to assess
182 specific and global aspects, a method used in other measures, such as the EQ-5D (EuroQol,
183 2017). The questionnaires battery consisted of measures with low overall item totals, making
184 it convenient for participants who may tire from completing larger, more time-consuming
185 batteries, especially within clinical samples (Waltz et al., 1991).

186 The Symptom Burden Index–Hands (Kallen et al., 2010) was used to measure
187 symptom severity (hand function). It is a 5-item subscale of a 40-item measure of symptom
188 burden in systemic sclerosis, a closely related condition. For this study, participants were
189 asked to consider symptoms relating to Raynaud’s (Pauling et al., 2018) over the last two
190 weeks and responded to items (*e.g. How often were hands a problem?*) using a rating scale
191 (0-10). The subscale showed excellent internal consistency in this study ($\alpha = .98$), while the
192 complete index has been validated in systemic sclerosis patients (Kallen et al., 2010).

193 The Bath Ankylosing Spondylitis Patient Global Score (Jones et al., 1996) was used
194 to measure symptom severity (global impact). Two VAS items (0-10) that ask participants to
195 indicate the effect their disease has had on their wellbeing over the last week and last six
196 months are averaged to provide the global score. The two items were highly correlated ($r =$
197 $.77, p < .001$) and the measure has been previously validated (Jones et al., 1996).

198 The ONS4–Life Satisfaction (Tinkler & Hicks, 2011) is a validated single-item
199 measure of personal wellbeing asking “*Overall, how satisfied are you with your life*

200 *nowadays?”* (0-10). As wellbeing is comparable to quality of life, the measure was
201 considered suitable given the lack of relevant measures for this group (Camfield &
202 Skevington, 2008). It is included in the Office for National Statistics (ONS) Annual
203 Population Survey to estimate personal wellbeing in the UK, demonstrating its utility as a
204 wellbeing measure (ONS, 2018). The single item measure also allows for direct measurement
205 of personal wellbeing, reflecting good face validity (Wanous et al., 1997).

206 **Analytic Strategy**

207 Total (sub)scale scores were computed in SPSS, v.26. Missing data was replaced with
208 the series mean, as was suitable given the sizable sample and low rate of missing data (1.5%)
209 that were missing completely at random, as determined using Little’s MCAR test (Parent,
210 2012). Cronbach’s alpha coefficient was calculated to assess the internal consistency of each
211 scale.

212 Descriptive statistics were calculated for the total sample, as well as primary and
213 secondary Raynaud’s separately, for stress, anxiety, depression, anxiety sensitivity, beliefs
214 about emotions, symptom severity (hand function), symptom severity (global impact) and
215 quality of life, as were the proportion within each DASS-21 subscale severity label. Summary
216 data *t*-tests were calculated to make comparisons with previous normative/nonclinical data.

217 Bivariate correlations (*Pearson’s R*) assessed the relationship between stress, anxiety,
218 depression, anxiety sensitivity, beliefs about emotions, symptom severity (hand function),
219 symptom severity (global impact) and quality of life. Concern for multicollinearity was
220 considered using a threshold of $r > .8$ (Field, 2013).

221 Two-tailed independent samples *t*-tests were performed to assess group differences
222 between primary and secondary Raynaud’s in stress, anxiety, depression, anxiety sensitivity,
223 beliefs about emotions, symptom severity (hand function), symptom severity (global impact)
224 and quality of life. Welch’s *t*-test was reported where appropriate as indicated by Levern’s

225 test for violations in equality of variances assumption. To account for nonnormal distribution
226 of (sub)scale scores, 95% percentile bootstrapped confidence intervals (2,000 resamples)
227 were calculated for *t*-tests (Field, 2013). As Raynaud's type could not be inferred from
228 participants who did not specify this, 17 cases were excluded from these analyses.

229 Stepwise multiple regressions were conducted separately on symptom severity (hand
230 function), symptom severity (global impact) and quality of life to assess the R^2 variance
231 accounted for by stress, anxiety, depression, anxiety sensitivity and beliefs about emotions.
232 Based on prior research, Raynaud's type, condition duration, age, gender and smoking history
233 were controlled for in each regression to account for confounding (Garner et al., 2015).
234 Violations of linearity, normality and homoscedasticity were judged through visual
235 inspection of histograms and scatterplots of the residuals and models were checked for
236 influential outliers (standardized residuals ± 3 and Cook's distance > 1) and multicollinearity
237 (Tolerance $< .01$). An alpha level of .05 was used for analyses.

238 **Results**

239 Descriptive statistics for variables and participant proportions within each DASS-21
240 severity label category are presented in Table 1. In comparison to normative data (Crawford
241 & Henry, 2003; Peterson & Plehn, 1999), participants had significantly higher stress
242 ($t(251.93) = 7.73, p < .001$), anxiety ($t(236.78) = 10.62, p < .001$), depression ($t(243.10) =$
243 $7.46, p < .001$) but not anxiety sensitivity ($t(220.01) = -0.14, p = .89$). Participants also had
244 significantly higher negative beliefs about emotions than a previous non-clinical sample
245 (Rimes & Chalder, 2010; $t(175.36) = 5.01, p < .001$).

246 Bivariate correlations showed that measures of stress, anxiety, depression, anxiety
247 sensitivity, beliefs about emotions, symptom severity and quality of life were all significantly
248 correlated in the expected directions (Table 2). A strong correlation was found between hand
249 function and global impact ($r = .78, p < .001$) due to the convergence around symptom

250 severity measurement. Strong correlations were also found between stress and depression ($r =$
251 $.65, p < .001$) and anxiety and anxiety sensitivity ($r = .63, p < .001$) indicating the variables
252 were related but below the threshold for possible multicollinearity, i.e. they are distinct
253 constructs.

254 Independent samples t -tests indicated that participants with primary Raynaud's had
255 significantly lower anxiety ($t(172.10) = -2.89, p = .004, 95\%$ bootstrapped CI $[-4.69, -0.97], d$
256 $= .41$), symptom severity in the domain of hand function ($t(191) = -5.11, p < .001, 95\%$
257 bootstrapped CI $[-2.64, -1.22], d = .74$) and domain of global impact ($t(191) = -5.30, p <$
258 $.001, 95\%$ bootstrapped CI $[-2.36, -1.12], d = .76$) but were higher in relation to quality of life
259 ($t(188.52) = 3.46, p = .001, 95\%$ bootstrapped CI $[0.44, 1.60], d = .49$) than participants with
260 secondary Raynaud's. The two diagnostic groups did not significantly differ on measures of
261 stress ($t(191) = -0.23, p = .822, 95\%$ bootstrapped CI $[-2.82, 2.07], d = .03$), depression
262 ($t(191) = -1.97, p = .051, 95\%$ bootstrapped CI $[-4.99, -0.97], d = .28$), anxiety sensitivity
263 ($t(184.57) = -1.69, p = .092, 95\%$ bootstrapped CI $[-6.13, 0.37], d = .24$), or beliefs about
264 emotions ($t(191) = -0.55, p = .582, 95\%$ bootstrapped CI $[-5.85, 3.22], d = .08$).

265 Stepwise regression analyses indicated that anxiety and Raynaud's type (primary or
266 secondary) explained 23% of the variance in symptom severity (hand function; $R^2 = .23, F(2,$
267 $207) = 30.50, p < .001$), see Table 3. Anxiety accounted for 16% of the variance ($\beta = .40, p <$
268 $.001$), while Raynaud's type accounted for an additional R^2 change of 7% ($\beta = .27, p < .001$).
269 All other entered variables were excluded.

270 A three-predictor model containing anxiety, Raynaud's type and anxiety sensitivity
271 accounted for 29% of the variance in symptom severity (global impact; $R^2 = .29, F(3, 206) =$
272 $27.34, p < .001$), see Table 4. Anxiety explained 20% of the variance ($\beta = .44, p < .001$).
273 Raynaud's type contributed an additional R^2 change of 7% ($\beta = .26, p < .001$) and anxiety

274 sensitivity explained a further R^2 change of 2% ($\beta = .19, p = .016$). All other entered variables
275 were excluded.

276 Three significant predictors explained 32% of the variance in quality of life ($R^2 = .32,$
277 $F(3, 206) = 32.68, p < .001$), see Table 5. Depression accounted for 29% of the variance ($\beta =$
278 $-.54, p < .001$), Raynaud's type explained an additional R^2 change of 2% ($\beta = -.15, p = .011$),
279 anxiety sensitivity contributed a further R^2 change of 1% ($\beta = -.13, p = .046$) to the model. All
280 other entered variables were excluded.

281 The regression models met the necessary assumptions of linearity, normality, and
282 homoscedasticity of the residuals. A single outlier was identified but retained as Cook's
283 Distance indicated that it was not influential. Tolerance values confirmed absence of
284 multicollinearity, meaning the regression models were statistically stable and regression
285 coefficients were reliable (Field, 2013). As prior power analysis indicated a sample of 98 was
286 needed to detect a medium effect size observed in prior related work (Ryan & McGuire,
287 2016; Wan et al., 2014) using $\alpha = .05, 1 - \beta = .8$, we can confidently report these results.

288 Discussion

289 Stress, anxiety and depression were found to be higher in those with Raynaud's when
290 compared to normative data, consistent with a body of research showing that mental health is
291 poorer in people with physical conditions (Crawford & Henry, 2003; Daré et al., 2019).

292 Those with primary and secondary Raynaud's did not significantly differ in terms of stress,
293 depression, anxiety sensitivity or beliefs about emotions, suggesting overall mental health is
294 similar between Raynaud's types, however anxiety was higher in those who experience
295 Raynaud's secondary to another health problem.

296 Group differences in relation to symptom severity and quality of life reflect a more
297 significant detrimental impact in secondary Raynaud's in comparison to primary Raynaud's,

298 in keeping with prior research (Fábián et al., 2019; Shapiro & Wigley, 2017). This may be
299 partly attributable to more systemic health problems in those with secondary Raynaud's.

300 Taken together, these findings suggest that psychological factors and quality of life
301 are integral to functioning and physical health and should be routinely assessed in Raynaud's
302 alongside a primary focus on symptom severity and health status.

303 Advancement of our understanding regarding the role of psychological factors is
304 reflected in the finding that anxiety, not stress, was independently associated with symptom
305 severity. This suggests that the term anxiety may more accurately describe the "emotional
306 stress" commonly purported to trigger episodes. This result agrees with previous findings by
307 Brown et al. (2001), however the sample here consisted of both primary and secondary
308 Raynaud's participants rather than just primary Raynaud's. It also provides further support
309 for Raynaud's original sympathetic overactivity hypothesis, which describes a hyperactivity
310 of internal fear-response systems that are associated with anxiety.

311 Anxiety, known to be amenable to evidence-based therapies such as cognitive
312 behavioural therapy, may provide a target for intervention in Raynaud's. As an intervention
313 which targets emotional wellbeing, quality of life and promotes effective-self management in
314 Raynaud's, cognitive behavioural therapy would be suitably aligned as a potential treatment
315 option, with further modification for this clinical group. It is particularly relevant given the
316 prolific evidence-base for cognitive behavioural therapy as a treatment for anxiety (NICE,
317 2020), which inherently aims to reduce hyperactive sympathetic responses associated with
318 anxiety that are thought to facilitate vasoconstriction in these patients. Given the
319 neurobiological basis behind Raynaud's, it is important that any psychological intervention
320 emphasises these aspects alongside targeting illness specific beliefs and behaviours which are
321 serving to maintain an overactive sympathetic nervous system (Mosely & Butler, 2015).
322 Employing a multidisciplinary approach which incorporates input from specialist

323 physiotherapy alongside cognitive behavioural therapy might be especially beneficial to
324 people with Raynaud's.

325 Anxiety sensitivity accounted for some of the variance (albeit marginal) in quality of
326 life and global impact, but not hand function. This is in line with associations found in related
327 rheumatological conditions (Bernstein et al., 2019; Mehta et al., 2016), of which many will
328 feature Raynaud's. This finding indicates a sensitivity in the physiological response to
329 anxiety and the physiology of their condition in Raynaud's patients, which may impact
330 symptom experience and quality of life. Although highly correlated with anxiety, anxiety
331 sensitivity was independently associated with symptom severity, demonstrating that they are
332 indeed distinct constructs and worthy of consideration separately as targets for intervention.

333 Depression did not predict symptom severity but accounted for a large proportion of
334 the variance in quality of life, which corresponds with associations found in prior related
335 work (Hudson et al., 2008; Wan et al., 2016). Surprisingly, anxiety was not independently
336 associated with quality of life, contrasting with previous research in related conditions
337 (Anyfanti et al., 2016; Sierakowska et al., 2019). This does suggest that anxiety does not have
338 as large an impact on quality of life in Raynaud's, as compared to depression, which may be
339 partly attributable to the functional and emotional limitations often associated with
340 depression. As such, treating comorbid depression should also be at the forefront of any
341 intervention as it may work towards improving quality of life in people with Raynaud's.

342 Beliefs about emotions was not significantly predictive of symptom severity or
343 quality of life in the regression models. This is contrary to findings in other stress-related
344 conditions (Bowers & Wroe, 2016), and inconsistent with theories of emotion in Raynaud's,
345 suggesting that cognitions surrounding the experience of emotion and physiology in
346 Raynaud's may be more complex than in other conditions. A considerable association was
347 exhibited between beliefs about emotions and anxiety sensitivity. Indeed, anxiety sensitivity

348 is a belief about emotion itself and its associated physiology, based in the belief that anxiety
349 symptoms are harmful. Therefore, it is possible the beliefs about emotions that impact
350 symptom severity and quality of life in Raynaud's relate specifically to the experience of
351 anxiety and the knowledge that it can trigger episodes. As such, beliefs about anxiety
352 specifically (i.e., anxiety sensitivity) may be more pertinent to address in this group than
353 beliefs about emotions more generally.

354 By looking at beliefs about emotions more generally, we may be missing other
355 important condition-related cognitions and belief systems which indirectly maintain
356 symptoms in Raynaud's and give rise to avoidant coping strategies. In their development of
357 the beliefs about sharing illness experiences scale (BASIE), Wroe and Bowers (2019)
358 reported that beliefs regarding the unacceptability of sharing illness experiences maintained
359 cycles of symptoms and distress in fibromyalgia patients. This may be similarly relevant in
360 people with Raynaud's given the common feelings of fear and embarrassment related to the
361 visibility and impact of symptoms which may further serve to trigger or maintain a
362 Raynaud's episode.s.. Raynaud's sufferers with alexithymia who find it difficult identifying
363 and describing feelings may be particularly vulnerable in this regard as having alexithymia
364 may further reduce support-seeking behaviours and increase suppressive emotion regulation
365 strategies (Fabien et al., 2020). Further research is needed to understand the complex
366 relationship between beliefs about emotions, expression of emotion and coping strategies in
367 Raynaud's. This may support the development and adaptation of a CBT based treatment
368 model

369 The overall findings support a biopsychosocial model for use in Raynaud's;
370 psychological factors have been found to be related to the fear-based activation of the
371 sympathetic nervous system that inhibits blood flow to the extremities, which is likely to be
372 moderated by beliefs that emotional factors are closely related to the activity of their

373 Raynaud's (Newton et al., 2012; Pauling et al., 2018). Adopting a biopsychosocial lens when
374 assessing, formulating and treating anxiety, anxiety sensitivity and depression in Raynaud's
375 patients is vital due to the interaction between these dimensions in Raynauds, particularly the
376 autonomic arousal, role of cognition and the social discomfort commonly seen in Raynaud's.
377 . While the common approach to the treatment and management of long-term conditions is
378 CBT (Daniels Psychologist mag CBT trickles down), it would be imperative that the
379 biological components and autonomic system are adequately taken account within the
380 physiological aspect of this approach.

381

382 Study findings also provide empirical support for NICE first-line recommendations
383 that Raynaud's patients minimise their emotional stress to help manage the condition (NICE,
384 2022). Qualitative research reported that those with scleroderma, of whom almost all will
385 have Raynaud's, were reluctant to seek psychosocial support specifically for their distress,
386 therefore a stepped care integrated approach may be most suitable to accommodate the
387 different levels of care desired by individuals (Newton et al., 2012). Low-intensity care might
388 include patient education and self-management strategies based on the principles of cognitive
389 behavioural therapy (e.g., non-avoidance of temperate changes/stress), as recommended by
390 NICE for anxiety (NICE, 2020).

391 There are several services available in the UK National Health Service (NHS), as part
392 of an initiative to integrate physical and mental healthcare for people with long-term physical
393 conditions (NHS, 2018). Talking Therapies in the NHS currently only offer interventions
394 which focus on low mood and anxiety, addressing only part of the care pathway for people
395 with Raynaud's. It would be optimal to offer a more holistic model and approach from which
396 these services can work from, focusing on the nervous system through education, formulation
397 to identify relevant beliefs and behaviours, and intervention using cognitive behavioural

398 therapy, as part of a multidisciplinary approach. Improving Access to Psychological
399 Therapies (IAPT) services co-located in physical health services allow patients to access
400 NICE-recommended therapies alongside physical treatment (NHS, 2018). This approach
401 promotes greater coordination between healthcare providers to comprehensively address the
402 needs of Raynaud's patients and improve overall care but may not be available everywhere.

403 The findings here could relate to other conditions which are underpinned by the
404 similar physiological mechanisms (e.g. a sensitised autonomic nervous system) and may also
405 benefit from a multidisciplinary approach to treatment. Future research in this area is needed
406 to consider the benefits of having a holistic approach to managing conditions such as this
407 where the relationship between physiological and psychological experience are closely bound
408 by the cognitions and neurobiological mechanisms that trigger and maintain them.

409 **Limitations and Future Research**

410 Due to the cross-sectional design of the study, causal direction cannot be inferred
411 from these results. Prospective longitudinal research is needed to establish a greater
412 understanding of the direction of influence. Nevertheless, the identification of psychological
413 factors that predict variance in symptom severity and quality of life in Raynaud's is important
414 to inform the direction of such research.



415 It was evident that there is a lack of suitable outcome measures for this group, as
416 reported in Daniels et al. (2018), and the online cross-sectional design prohibited the use of
417 the traditional Raynaud's Condition Score diary (Daniels et al., 2018). However, the
418 measures adapted for this study showed good reliability and produced meaningful results that
419 corresponded to measures used in prior Raynaud's studies (Brown et al., 2001; Fábíán et al.,
420 2019; Hughes et al., 2015). These should be further tested and considered for use in future
421 research.

422 Self-selection via online recruitment and self-reporting of Raynaud's type potentially
423 undermined the credibility of the sample. This could have resulted in a biased sample that
424 may not be fully representative, a common challenge in online studies (Gosling & Mason,
425 2015). Future research would benefit from a clinically confirmed representative sample rather
426 than a self-selected online sample.

427 Data was collected during a short period in early summertime in the UK, which
428 limited the confounding role of temperature, given its importance in Raynaud's. It would be
429 useful to repeat this study or use a longitudinal design to observe whether the influence of
430 these factors change with the seasons. As data collection took place early on in the
431 Coronavirus infectious disease (CovID-19) pandemic, it is worth noting the potential impact
432 that elevated anxiety and depression experienced during this time may have had in the
433 context of this study, as anxiety is a known trigger for Raynaud's (Gigante et al., 2020; Rettie
434 & Daniels, 2021). Although elevated levels of anxiety are unlikely to have altered the nature
435 of the relationships between the key variables.

436 **Conclusion**

437 These findings provide pivotal insight into the psychological factors associated with
438 symptom severity and quality of life in people with Raynaud's phenomenon; an area that has
439 previously been relatively under researched despite having a strong theoretical and practical
440 basis for study in this condition. Study results suggest a multidisciplinary biopsychosocial
441 approach which address psychological factors in addition to physical needs may be most
442 appropriate for the treatment of Raynaud's and provide empirical support for NICE first-line
443 recommendations that patients minimise emotional stress to help manage the condition.
444 Cognitive behavioural therapy is suitably aligned as a potential treatment option, considering
445 its recommendations for use with anxiety and robust evidence base supporting its delivery in
446 rheumatological and other medical conditions. This paper presents initial findings that may

447 underpin the adaption of cognitive behavioural therapy for this common, debilitating
448 problem.

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