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1 **Barriers & enablers to providing community-based occupational therapy to**  
2 **people with Functional Neurological Disorder: An interview study with**  
3 **occupational therapists in the United Kingdom**

4

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25

26 **Research Ethics**

27 Approved by the School of Health Sciences Ethics Committee, City University of  
28 London (REC reference number: MRes/16-17/10) on the 24 February 2017 and  
29 appropriate local authorities.

30 **Declaration of conflicting interests:**

31 None

32 **Contributorship**

33 CN was the primary investigator for this study, researched the literature and applied  
34 for ethical approval. CN, FL, JF contributed to the methodology. CN and FL

1 interpreted the data. CN wrote the first draft of the paper. All authors reviewed and  
2 edited the manuscript.

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10

1 **INTRODUCTION**

2 Functional Neurological Disorder (FND), also known as conversion disorder, is  
3 characterised by altered sensory or motor function that is inconsistent with known  
4 structural neurological disease (Espay et al., 2018). Symptoms are diverse and may  
5 include movement disorder, weakness, sensory symptoms, cognitive dysfunction and  
6 dissociative seizures (Stone, 2009). Functional symptoms are common, with an  
7 estimated 30% of neurology outpatients and 10% of neurology inpatients presenting  
8 with symptoms that are 'somewhat or not at all explained by disease' (Stone et al.,  
9 2010). The symptoms of FND can be highly disabling and are associated with high  
10 levels of distress, reduced quality of life and high health and social care utilisation  
11 (Carson et al., 2011; Gelauff et al., 2019; Merkler et al., 2015).

12 Inpatient multi-disciplinary (MDT) rehabilitation involving occupational therapy is  
13 currently considered the gold standard treatment for FND. There is growing evidence  
14 to support this treatment approach, which includes several large cohort studies  
15 (participant numbers 26–78) (Demartini et al., 2014; Jacob et al., 2018; McCormack  
16 et al., 2014; Saifee et al., 2012) and a small randomised study (n=60)(Jordbru et al.,  
17 2014). Yet, there are limitations to inpatient MDT rehabilitation for FND. Inpatient  
18 settings do not suit all individuals; inpatient rehabilitation is costly, resource intensive  
19 and in the UK, it is oversubscribed resulting in long waiting times. An alternative  
20 treatment model is outpatient and community-based therapy, which has the advantage  
21 of lower costs. However, evidence for occupational therapy in this context is lacking;  
22 anecdotally clinicians can lack experience, confidence and expertise in FND with many  
23 community occupational therapists working in isolation with reduced MDT support.

24 People with FND are commonly seen by neurological occupational therapists in the  
25 UK and abroad in a variety of settings (Gardiner et al., 2018; Paget and Rigby, 1996;  
26 Ranford et al., 2020a), but little is known about the experiences or interventions of  
27 occupational therapists working in community settings. A systematic review of studies  
28 of physical interventions for people with FND from 2013 found no examples of  
29 occupational therapy or home-based treatments (Nielsen et al., 2013). Our search of  
30 the literature for examples of barriers / enablers to community occupational therapy  
31 practice for people with FND found only a single case study from 1996. This paper  
32 described ongoing compensation claims and non-acceptance of the diagnosis as  
33 barriers to therapy engagement in a patient with functional overlay following traumatic

1 brain injury (Paget and Rigby, 1996). Barriers to the treatment of people with FND in  
2 settings other than the home have been identified by other health care professionals,  
3 and have relevance to occupational therapy practice. Perceived barriers include lack  
4 of knowledge of the condition and its treatment; stigma about the diagnosis; reduced  
5 confidence in discussing psychological aspects of the diagnosis; reduced access to  
6 mental health support and lack of continuity of care from hospital to community  
7 environments (Ahern et al., 2009; Barnett et al., 2021; Demartini et al., 2014; Edwards  
8 et al., 2012; Kanaan et al., 2009; Nielsen et al., 2018).

9 Despite the growing recognition of the role of occupational therapy in FND, the lack of  
10 published guidance and limitations of community therapy frameworks leave many  
11 services feeling unable to accept referrals (Barnett et al., 2020; Demartini et al., 2014).  
12 As a result, a large group of patients with high levels of distress and disability are  
13 unable to access treatment. We therefore sought to explore the barriers and enablers  
14 to delivering community-based interventions to people with FND, from the perspective  
15 of neurological occupational therapists in the UK. This exploration was supported by  
16 the application of a behavioural science framework; the Theoretical Domains  
17 Framework (TDF) (Phillips et al., 2015). The TDF provides a framework in which  
18 influences on behaviour (delivery of occupational therapy intervention) and behaviour  
19 change (changes in clinical practice) can be systematically explored. Published  
20 descriptions of occupational therapy interventions for FND are limited (Gardiner et al.,  
21 2018; Nicholson et al., 2020; Ranford et al., 2020a, 2020b) and it remains unclear as  
22 to whether occupational therapy interventions vary between practice settings, hence  
23 this study also sought to review the practice of neurological occupational therapists  
24 working in community settings.

25

## 26 **METHODS**

### 27 **Research Design**

28 This was a UK based, qualitative, semi-structured interview study modelled on the  
29 Theoretical Domains Framework (TDF) (Michie et al., 2005).

30 The TDF is a behavioural and implementation science framework that synthesises  
31 constructs from 33 behaviour change theories into 12 theoretical domains,  
32 representing the range of individual, sociocultural, and environmental influences of

1 behaviour and behaviour change (e.g. knowledge, motivation and goals, beliefs about  
 2 capabilities, social influences, social identity/role) ( See Table 1) (Francis et al., 2012;  
 3 Roberts et al., 2016). Applying the TDF during data collection enables a systematic  
 4 and comprehensive exploration of possible influences on practice and provides a  
 5 systematic method for selecting change strategies to support change to clinical  
 6 practice (Phillips et al., 2015). The TDF has informed data collection and analysis for  
 7 a number of international studies exploring the barriers and enablers to healthcare  
 8 professional behaviour change across a range of clinical contexts, including  
 9 occupational therapy (Francis et al., 2012; Sakzewski et al., 2014).

## 10 **Materials**

11 The interview topic guide was based on the TDF and was informed by the researchers'  
 12 (CN) clinical knowledge and experiences as an occupational therapist working with  
 13 people with FND and was developed in collaboration with behavioural scientists (FL,  
 14 JF). The target behavior of interest was the delivery of community-based neurological  
 15 occupational therapy for people with FND. The topic guide includes at least one  
 16 question per domain of the TDF. Table 1 presents definitions of each domain alongside  
 17 an example question for each domain. Flexible open-ended follow up questions and  
 18 prompts were used to further elaborate upon participant responses. The topic guide  
 19 was piloted with three community neurological occupational therapists and refined to  
 20 ensure question clarity and flow. The final interview guide is available in supplementary  
 21 file 1.

22 **Table 1. Example of interview questions allocated to the TDF domains** (Atkins et al., 2017)

TDF Domain	Domain Content	Exemplar Interview Question
Knowledge	Knowledge (including knowledge about condition / specific rationale) Schemas + mindsets + illness representations Procedural knowledge	Could you tell me what your understanding of the diagnosis of FND is and how you have developed this understanding?
Nature of Behaviours	Routine / automatic / habit Direct experience / past behaviour Representation of tasks	What sort of interventions do you routinely use with people with FND in the community?
Social / Professional Role & Identity	Professional identity, boundaries & roles Group / social identity Alienation / organisation commitment	As a neuro OT to what extent do you consider providing therapy to people with FND to be part of your role?

Social Influences	Interpersonal interactions that can influence clinicians' thoughts, feelings and behaviours. Social support Team working Supervision, learning and modelling Inter-group conflict	Do you feel any pressure from internal or external colleagues to accept or decline referrals for people with FND?
Emotion	Emotions and reactions experienced by the clinician in relation to their working role.	Do you enjoy working with people with a diagnosis of FND?
Memory, Attention & Decision Making	Memory Attention Decision making (factors influencing decision making)	As an OT how do you decide what to work on with the patient? Do you follow a sequence of steps?
Beliefs About Capabilities	Self-efficacy Perceived Competence (of self & others) Optimism / pessimism	As an OT how easy or difficult do you find it to provide OT to people with FND?
Beliefs About Consequences	Outcome expectancies Perceived risk / threat / benefit Attitudes Appraisal / evaluation	If people with FND were not able to access OT services in a timely fashion what do you think might happen?
Skills	Competence / ability, skills assessment Interpersonal skills (including written & verbal communication skills) Coping strategies	What do you think are the specialist skills or knowledge that are important for OTs to have when working with people with FND?
Motivation & Goals	Goal setting / targets Intrinsic motivation Commitment (commitment to achieving set goals)	Compared to other things you have to do as part of your role, where would you rank providing OT to people with FND in terms of priority?
Environmental Context and Resources	Resources / material resources Environmental stressors, constraints or facilitators Organisation culture / climate	What resources do you feel are necessary to provide quality therapy to people with FND in the community?
Behavioural Regulation	Reflexivity Feedback (given or received) Moderators of intention – behaviour gap Guidelines / Evidence Base Barriers / facilitators	How is your approach to treating people with FND different to what you may use when treating someone with another neurological condition e.g. MS or stroke?

1

## 2 Participants and Sampling



1 Participants were neurological occupational therapists experienced in providing  
2 treatment for people with FND in the community in the UK. Those without experience  
3 or who were unable to read and write in English were excluded. Purposive sampling  
4 via the Royal College of Occupational Therapists (RCOT) Specialist Section for  
5 Neurological Practice (SSNP) was undertaken. The RCOT SSNP is a special interest  
6 group which had approximately 950 members at the time of recruitment. The use of  
7 RCOT SSNP UK members aimed to capture occupational therapists with various  
8 levels of experience and ensured that data could be gathered from across the UK.  
9 Based on recommendations for sample sizes in qualitative research (Francis et al.,  
10 2010), an initial sample of 10 participants were recruited. Following analysis of the first  
11 10 participants, the initial data set was assessed for thematic saturation. If no new  
12 themes emerged, data saturation was deemed to be achieved and no additional  
13 participants would be recruited. If saturation was not achieved, a further three  
14 participants would be recruited, saturation re-assessed, in an iterative manner until  
15 saturation was reached.

## 16 **Procedure**

17 An email invitation to participate in a 60-minute telephone or face-to-face interview  
18 and a participant information sheet was sent by the RCOT SSNP to all registered  
19 members. Interested participants were asked to contact the researcher, at which time  
20 their eligibility for participation was confirmed. Those meeting the inclusion criteria  
21 completed a consent form and the interviews were then scheduled at a date, time and  
22 location convenient to participants. The interviews were undertaken by the lead  
23 researcher (CN). Audio-recordings were transcribed verbatim and anonymised so that  
24 no individual or organisation could be identified.

## 25 **Data Analysis**

26 Interview transcripts were analysed in accordance with published guidance for using  
27 the TDF, using a six-step combined content and framework analysis approach (Atkins  
28 et al., 2017) (see figure 1). This process is:

29 *1. Familiarisation with the data:* The lead researcher transcribed a sub-set of the  
30 interviews, reading and re-reading the interview transcripts and listening to each audio  
31 recording at least once. This process allowed the researcher to familiarize themselves

1 with the data, gain a 'sense' of each participant and record emerging themes and  
2 subjects of interest (Smith et al., 2009).

3 *2. Developing a coding agreement:* Two researchers (CN, FL) independently coded  
4 the first interview, using the TDF as a coding framework by categorising participant  
5 responses according to the domain they were judged to best represent. Results were  
6 compared and coding heuristics were developed (Atkins et al., 2017). Both  
7 researchers coded the second interview separately and compared the results to  
8 ensure coding consistency (Patey et al., 2012). The agreed strategy was then  
9 employed to code the remaining transcripts. This ensured consistency between  
10 researchers and minimised subjective bias (Atkins et al., 2017; Thompson et al.,  
11 2004).

12 *3. Deductive Coding of interview transcripts into theoretical domains:* The lead  
13 researcher coded the remaining interview transcripts independently. Participant  
14 responses were coded into the TDF domains they were judged to best represent.  
15 Responses were deemed to be relevant if they were directly related to the research  
16 question or target behaviour (e.g., *"And I'm also much more comfortable at saying,*  
17 *"Look, I'm here to support you but this is actually your problem and if you don't take*  
18 *ownership of it, I can't help you"* (P03), coded into the domain *Behavioural Regulation*  
19 as the therapist is trying to encourage the person to be an active participant in their  
20 own rehabilitation. Participant responses that were thought to address more than one  
21 domain were allocated to multiple domains (e.g., *"This is why we've had a very big*  
22 *education programme drive going on with the GPs... because they've only got ten*  
23 *minutes, they want that person out of the door"* (P08) coded to both *Skills* and  
24 *Environmental Context and Resources*. The coding was reviewed between CN and  
25 FL regularly in order to cross-check agreement, improve analytical rigour and minimise  
26 researcher bias (Nowell et al., 2017).

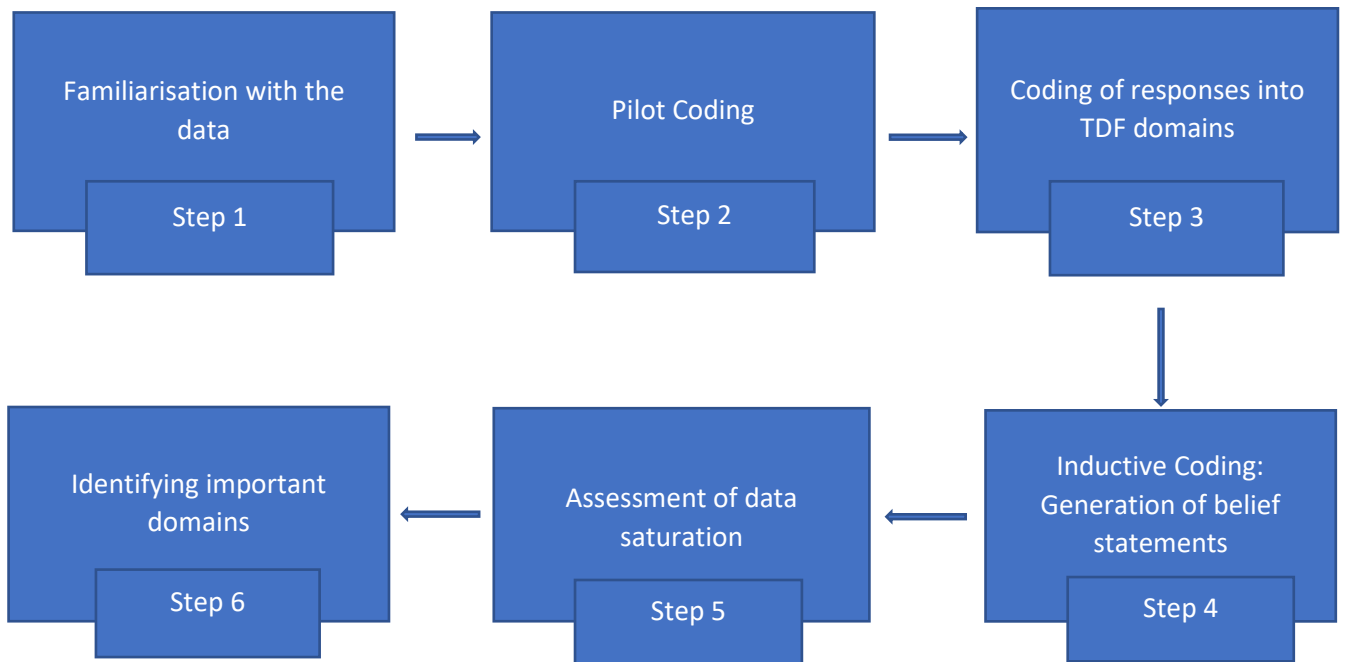
27 *4. Inductive coding: Identifying belief statements within each domain:* Next, similar  
28 responses coded to the same domain were grouped, and a belief statement (theme)  
29 was inductively generated. Each belief statement summarised the shared meaning of  
30 the grouped participants responses and represented a specific belief that one or more  
31 of the participants had shared (See Table 3). Belief statements were then classified

1 as a barrier, enabler, or mixed (i.e., a barrier to some participants but enabler to  
2 others), to delivering occupational therapy.

3 **5. Data Saturation:** Data saturation was assessed in terms of thematic saturation.  
4 Thematic saturation is reached when the data fails to contribute new information that  
5 may have influence over the target behaviour (Atkins et al., 2017).

6 **6. Assessing domain importance:** The domains of greatest importance to influencing  
7 occupational therapy intervention with people with FND in the community were  
8 identified using criteria proposed by Atkins et al., 2017 and Francis et al., 2009: (1)  
9 Frequency of each belief statement across all interviews/participants. Higher  
10 frequency equated higher importance. (2) Presence of discordance / conflicting  
11 beliefs. (3) Expressed importance; domains that had specific beliefs at either high or  
12 low frequency can be considered important if participants reported that they were key  
13 factors that guided their practice or behaviour, as expressed with definitive language.  
14 (See Table 2) (4) Domain Elaboration, number of belief statements identified within  
15 each domain, with more belief statements representing greater elaboration and thus  
16 greater importance.

17 **Figure 1. 6 Stage Data Analysis Process** (Amended from (Atkins et al., 2017; Roberts et al.,  
18 2016)



# 1 RESULTS

## 2 Recruitment and data saturation

3 Sixteen occupational therapists expressed an interest in participating. Ten met the  
4 inclusion criteria and consented to participate. One face-to-face and nine telephone  
5 interviews were undertaken. The mean interview duration was 72 minutes (SD 8,  
6 range 46-83).

## 7 Participant characteristics

8 All participants were female with 9 to 32 years of occupational therapy experience  
9 (mean 19, SD 9) and 1.5 to 25 years of experience working in community neurological  
10 rehabilitation settings (mean 10.7, SD 6.8). Seven participants were employed at band  
11 7-level and three were employed at Band 6. Two participants worked in London, five  
12 elsewhere in England and three in Scotland.

## 13 Data Saturation

14 No new belief statements were generated after participant nine therefore thematic  
15 saturation was deemed to have been achieved and the sample size of ten was  
16 considered sufficient to ensure content validity. (See supplementary file 2: Data  
17 Saturation Table.)

## 18 Assessment of Domain Importance

19 When applying the importance criteria of *frequency of specific beliefs*, *discordant*  
20 *themes* and *expressed importance*, the top five domains deemed to be important in  
21 affecting the delivery of occupational therapy intervention to people with FND in the  
22 community were; (i) Beliefs About Consequences; (ii) Memory, Attention and Decision  
23 Processes; (iii) Nature of Behaviours; (iv) Environmental Context and Resources; and  
24 (v) Social/Professional Role and Identity (Table 2).

25 **Table 2. Assessment of Domain Importance**

TDF Domain	Level of Elaboration (number of themes)	Number of Participants Coded to Domain	Discordant Themes	Expressed Importance	Expressed Importance Quote Examples
Beliefs About Consequences	31 (+ 3 sub-themes)	10	Yes (n=3)	Yes (n=9)	'We'd need support from the MDT as well, because I think that's very key, that

					<i>it needs to be an MDT approach'. (P03, pg. 20)</i>
<b>Memory, Attention and Decision Processes</b>	24	10	Yes (n=1)	Yes (n=1)	<i>'...and it's always, it's always, it's always OT and physio, it always is, it just always is'. (P08, pg. 12) (Professionals who see people with FND in the community).</i>
<b>Nature of Behaviours</b>	17	10	Yes (n=1)	Yes (n=4)	<i>'I think obviously things like pain and fatigue management again, and best routine and structure are fundamental in trying to help'. (P03, p.10)</i>
<b>Environmental Context &amp; Resources</b>	17	10	No	Yes (n=6)	<i>'I think having access to clinical psychology is very helpful, whether it is directly with the individual or whether it is advice and support for the OT and physio colleagues. (P09, p.g 26)</i>
<b>Social / Professional Role &amp; Identity</b>	15	10	Yes (n=2)	Yes (n=2)	<i>'...so I think the bottom line is that this is a disorder that affects the way the nervous system functions and therefore is totally appropriate to see within the context of neuro rehab'. (P05, pg. 23)</i>
<b>Social Influences</b>	14 (+ 1 sub-theme)	10	Yes (n=1)	Yes (n=2)	<i>'I don't know how I'd manage without them'. (P06, pg., 6) (support of Psychologist on the mainland)</i>
<b>Knowledge</b>	14 (+ 4 sub-themes)	10	No	Yes (n=2)	<i>'...the knowledge that's needed, so that's first and foremost'. (P08, pg. 28) (Knowledge of FND)</i>
<b>Skills</b>	12 (+3 sub-themes)	10	No	Yes (n=4)	<i>'There is only so much college can do but I think what sets OT apart from everyone else are functions and activity analysis'. (P06, pg. 35)</i>
<b>Emotion</b>	8	10	Yes (n=1)	Yes (n=1)	<i>'Absolutely love it; absolutely love it'. (P08,</i>

					pg. 25) (Working with people with FND)
<b>Beliefs About Capabilities</b>	7	10	Yes (n=4)	Yes (n=1)	<i>'It's making a demarcation between where I feel competent and able to practice and saying, "No, I can't do that" and feeling the confidence to turn around and say I can't do that, because that can be quite challenging as well'. (P06, pg. 29)</i>
<b>Motivation &amp; Goals</b>	7	10	No	Yes (n=3)	<i>'So, it is very much a client centred approach, and that is the only way to work for me'. (P05, pg. 16) (Client centred goal setting)</i>
<b>Behavioural Regulation</b>	6	9	No	No	

1

## 2 **Coding interview transcripts into theoretical domains**

3 In total, 2646 responses across interviews were coded into the theoretical domains.  
4 The domains with the most responses were Beliefs About Consequences (n=467);  
5 Environmental Context and Resources (n=381); Memory, Attention and Decision  
6 Making (n=363). The domains with the least responses were Emotion (n=84),  
7 Motivation and Goals (n=87), and Behavioural Regulation (n=30). Responses were  
8 synthesised into 174 belief statements (themes). Of these, 30% were deemed to be  
9 barriers, 34% enablers and 36% mixed. Beliefs About Consequences presented the  
10 most barriers (n=8), enablers (n=11) and mixed themes (n=12). See Supplementary  
11 File 3 for a full table of TDF domains and associated belief statements (1-12). See  
12 Table 3 for coding interview transcripts (examples from each domain).

13

## 14 **Table 3. Coding Interview Transcripts (examples)**

<b>TDF Domain</b>	<b>Example Belief Statement (theme)</b>	<b>Example Utterance (Statement)</b>	<b>Barrier / Enabler / Mixed</b>
Emotion	Patients with FND find the symptoms and resulting disability distressing	<i>'I think that sometimes people, and completely understandably are</i>	Barrier (6 participants)

		<i>frightened and frustrated</i> '. (P02)	
Environmental Context & Resources	We don't have access to the resources that are needed to treat patients with FND effectively.	<i>'It's difficult. It's frustrating. You know you could make a difference with a great many but you don't have the time or the energy or the resources to do it</i> '. (P06)	Barrier (9 participants)
Knowledge	Guidelines would help to develop the knowledge of OT's practice in this area.	<i>'...most people who work with people with functional symptoms aren't specialist OTs like me, they're OTs in social services who've been referred somebody and don't know which end to start with</i> '. (So guidelines would be helpful). (P04)	Barrier (2 participants)
Skills	Working with patients with FND requires specialist skills	<i>'I think neuro OTs are well placed to see them, but I think that the OT has to realise that the techniques and the therapy techniques that they are using with this group are not going to be the same as somebody for example with a stroke, or another brain injury</i> '. (P09)	Mixed (10 participants)
Beliefs About Capabilities	OT's feel that they (do not) have the capabilities to manage distress, anxiety and mental health issues in patients with FND	<i>'...we don't necessarily feel we have the skill to deal with some of the psychological elements of the condition...'</i> (P07)	Barrier (5 participants)
Beliefs About Consequences	Patients require access to community OT in a timely fashion to maximise their potential for	<i>'...increased dependency, possibly greater need for package of care, greater carer burden on family and friends and</i>	Barrier (10 participants)

	recovery (outcomes are worse without it).	<i>social isolation with potentially their mood deteriorating'. (P02)</i>	
Social Influences	Working collaboratively with other healthcare professionals is essential in providing quality care for patients with FND.	<i>'I think in the team that I am working in now it works very well as we tend to all take the same approach'. (P02)</i>	Mixed (10 participants)
Social Professional Role & Identity	Community neurological OT's do (do not feel) that they should see patients with FND.	<i>'I absolutely think it is part of my role, and it is a funny thing because there has been this big debate over should neuro therapists be seeing these cases, and I find it quite difficult to see why they are not all neuro cases'. (P05)</i>	Mixed (9 participants)
Motivation & Goals	Not all OTs want to work with patients with FND.	<i>'... I think it depends on the training and the expertise of the OT, and also the willingness because I can see that this group would not be every OTs preferred group to work with'. (P09)</i>	Barrier (2)
Nature of Behaviours	Intervention strategies for working with patient with FND are (are not) different to those used with patients with organic illness.	<i>'...one of the things I do is I acknowledge that I believe that their symptoms are real, and that I don't think they are making it up, and that they have a genuine problem. So, that is something I would probably never discuss with any other client group'. (P05)</i>	Mixed (9 participants)
Memory, Attention & Decision Making	Community neuro OTs have to clinically reason whether to provide aids and adaptations	<i>'Because there is a kind of, "Oh, we shouldn't give equipment to functional patients",</i>	Mixed (7 participants)



	to patients with FND.	<i>it can be a challenge at home because actually sometimes they need that bit of equipment because they're not actually able to achieve their goal otherwise'.</i> (P01)	
Behavioural Regulation	OT's need to employ reflexive strategies within clinical practice to improve their practice.	<i>'I think we've been quite good as a service at reflecting on things that haven't worked, and just maybe adjusting them, tweaking them a bit, but I think one of the big things that has probably changed is us identifying quite early on either people that we don't think are going to benefit from rehab, or we're going to have very limited results with'.</i> (P09)	Enabler (5 participants)

1

2 A narrative summary of findings within each domain is presented below. Where  
3 participant quotes are provided these are denoted with the participant number e.g.,  
4 P1-10.

5 **Nature of behaviours**

6 All participants indicated that there are routine strategies and interventions that they  
7 use when working in the community with people with FND (e.g., education about the  
8 diagnosis, fatigue and pain management, re-training of activities of daily living, graded  
9 goal setting, graded task practise, anxiety management and relaxation strategies).  
10 Discordant beliefs were reported when the participants were asked if the intervention  
11 strategies and/or skills used with people with FND were different to those used when  
12 treating people with other neurological conditions. *"It's interesting, if I've got a patient  
13 with a stroke or a patient with a functional illness who mimics a stroke, I would still  
14 treat them in the same way,"* (P10) versus; *"So, definitely there are distraction  
15 techniques and the use of rhythm that are quite exclusive to this group."* (P04). Two  
16 other participants also indicated that a point of difference with FND rehabilitation

1 versus standard neurological rehabilitation was the provision of psychoeducation and  
2 assistance to identify symptom triggers.

### 3 **Beliefs About Consequences**

4 Beliefs About Consequences was the most elaborated domain (n=31 beliefs: 8  
5 barriers, 11 enablers and 12 mixed). All ten participants thought that timely access to  
6 community occupational therapy is essential to prevent chronicity of symptoms and  
7 dependency. However, timely access and the quality of interventions provided was  
8 often prevented by resource restrictions; *"It's difficult. It's frustrating. You know you  
9 could make a difference with a great many but you don't have the time or the energy  
10 or the resources to do it"* (P06).

11 Most (n=9) felt that, as a general rule, better therapeutic outcomes are achieved when  
12 treatment occurs in home / community environments with MDT support. However, it  
13 was widely acknowledged (n=10, barrier) that for some, inpatient treatment may be  
14 the best option; *"...somebody who has a very high level of disability and...a high level  
15 of traumatic manifestation just needs a whole team approach..."* (P08).

### 16 **Memory, Attention & Decision Making**

17 Many beliefs in this domain concerned decision making regarding whether to accept  
18 referrals. One participant reported that their team as a whole has FND as an exclusion  
19 criterion, citing lack of resources and specialist skills as the reason. Two participants,  
20 who regularly saw patients with FND, indicated that an important barrier was that  
21 psychologists within their team and the local mental health service would not see  
22 people with FND.

23 The decision of when to discharge people with FND was highlighted as challenging by  
24 nine participants. Flexibility in the intervention period was identified as being beneficial.

### 25 **Environmental Context & Resources**

26 Environmental Context and Resources was a highly elaborated domain (n=17 beliefs:  
27 8 barriers, 4 enablers and 5 mixed). Lack of resources (staff and time) had resulted in  
28 barriers to MDT working secondary to different waiting times for different  
29 professionals. Participants from London and parts of Scotland reported better MDT  
30 access and fewer restrictions on commissioning of services. Lack of funding for key

1 therapies such as physiotherapy, speech and language therapy and psychology were  
2 identified as a barrier in parts of rural England and Scotland. In one region in rural  
3 England, the neurological rehabilitation service and mental health service were unable  
4 to see people with FND citing lack of commissioning. Those working in specialist MDT  
5 FND pathways (n=2) cited better clinical outcomes.

6 Two participants indicated that working in the community limited their access to  
7 supervision and professional support. Whilst staff in under-resourced areas or working  
8 in isolated geographical locations had developed cross professional competencies to  
9 meet the needs of their patients.

## 10 **Social / Professional Role & Identity**

11 All but one participant indicated that they felt community-based neurological  
12 occupational therapists should see people with FND. In teams where there was little  
13 support from psychology, occupational therapists often adopted a dual role as there is  
14 a need for psychological services but no-one to refer onto. This was a concern to some  
15 who worried about working outside of their professional boundaries.

## 16 **Knowledge**

17 Lack of understanding of the condition (n=9) amongst other clinicians and a lack of  
18 evidence base to guide treatment were identified as barriers (n=5). Whilst access to  
19 the website [www.neurosymptoms.org](http://www.neurosymptoms.org) and key papers in the field supported FND  
20 related knowledge (n=10).

## 21 **Skills**

22 Training in FND was identified as beneficial, however, financial constraints and lack of  
23 availability (n=9) made access difficult. Training access was greater in London and  
24 parts of Scotland. Whilst one participant reported the benefits of undertaking a training  
25 program with their local GPs about FND and MDT treatment options; *“This is why  
26 we’ve had a very big education programme drive going on with GPs...because they’ve  
27 only got ten minutes, they want that person out the door”* (P08).

28 Participants recognised that their dual physical and mental health training and core  
29 occupational therapy skills (activity analysis / rehabilitation through function) enhanced  
30 their ability to work with people with FND.

## 1 **Motivation and Goals**

2 The participants (n=8) stressed the importance of establishing client-centred goals in  
3 order to maximise functional outcomes. Goal achievement was seen as motivating  
4 (n=8) for both patients and clinicians whilst professionally, participants identified FND  
5 as an area that they were motivated to improve their skills in.

## 6 **Social Influences**

7 Working collaboratively with other healthcare professionals was seen as essential in  
8 providing quality care for people with FND. One participant indicated that GPs were  
9 their most frequent referrers for all other neurological conditions, yet their team rarely  
10 received referrals from GPs for people with FND. Concern was raised about the  
11 knowledge of some GPs regarding FND and its management. Most participants (n=9)  
12 also reported that they routinely invite significant others to be involved in the  
13 rehabilitation process, most commonly in; education about the diagnosis, symptom  
14 triggers and symptom management strategies.

15 Regarding the provision of mentorship and professional supervision, eight participants  
16 reported that they do not have direct access to a senior occupational therapist, but  
17 rather draw support from peers or other AHPs such as psychologists.

## 18 **Beliefs About Capabilities**

19 Concerns were raised (n=6) about abilities to effectively treat people with FND; *“Well,*  
20 *I don’t feel competent or skilled. I just look and I think, “Oh, god, how am I going to*  
21 *get through this one?”* (P10). Participants (n=5) also highlighted that they lacked the  
22 necessary mental health skills to effectively work with some people with FND (in the  
23 absence of psychology support). Whilst concerns were also raised about the abilities  
24 of some medical staff (GPs and neurologists) to recognise, diagnose and manage the  
25 condition in a sensitive and comprehensive manner.

## 26 **Emotion**

27 When asked whether they enjoyed working with people with FND there were mixed  
28 responses. Only one participant stated definitively that they did not enjoy it, citing  
29 patient complexity and lack of resources as reasons.

1 Other emotions expressed by participants were feeling frustrated at having to offer  
2 sub-optimal care due to limited resources, being scared to treat people with the  
3 condition and feeling overwhelmed from managing the complex physical and  
4 psychological aspects of the condition.

## 5 **Behavioural Regulation**

6 This was the least cited domain. Three participants indicated that guidelines would  
7 help to facilitate occupational therapy practice with people with FND. Others (n=3)  
8 highlighted the benefits of joint working with junior staff to enhance their knowledge  
9 and skills about FND.

10 Guiding people with FND in the use of self-management strategies was also identified  
11 by three participants as integral to their practice.

12

## 13 **Discussion**

14 The results of this study have highlighted individual, professional and service-level  
15 barriers and enablers to the delivery of community-based neurological occupational  
16 therapy to people with FND in the UK. Lack of resources, reduced access to timely  
17 interventions, lack of professional role clarity, and reduced understanding of the  
18 condition and treatment modalities were deemed to have the greatest influence on the  
19 delivery of occupational therapy interventions.

20 Within clinical practice there is often conjecture about which professionals are best  
21 placed to treat people with FND; neurological or mental health specialist clinicians.  
22 The results of this study have indicated that participants were enthusiastic about the  
23 potential benefits of community-based neurological occupational therapy interventions  
24 for FND with the majority of those interviewed citing that it was appropriate for  
25 neurological occupational therapists to see people with FND in the community.  
26 Reasons identified for this belief were: that it is a condition that affects the way that  
27 the neurological system works and the symptoms impede a patient's occupational  
28 performance. Similarly, Edwards et al., 2012 reported in their survey of neuro-  
29 physiotherapists working with people with functional movement disorder that 82% of  
30 respondents felt that physiotherapy was an appropriate treatment with more than 10%  
31 of their time being spent with this patient group. In contrast, Ahern et al., 2009 found

1 in their study of neuro-sciences nurses that 34% of those surveyed felt that neurology  
2 wards are not best placed to manage people with FND, whilst Ricciardi and Edwards,  
3 2014 highlighted long-standing ambiguity about which physicians (neurologists or  
4 psychiatrists) should treat people with FND.

5 Resource availability was unsurprisingly identified as a major barrier to the provision  
6 of effective and timely treatment. Others have previously reported barriers to  
7 accessing community therapy teams for people with FND in the UK (Demartini et al.,  
8 2014). Lack of funding is usually cited as the main issue, as well as prioritisation of  
9 people with other neurological conditions. Similar themes have also been  
10 highlighted in the US where a recent economic evaluation reported that despite  
11 acute healthcare costs for people with FND being comparable to those of other  
12 neurological conditions, funding for rehabilitation and psychiatry input for FND  
13 remains low (Stephen et al., 2020). In a review of commissioning for  
14 neuropsychiatric conditions (including FND) in London, services were found to be  
15 disjointed and variable (Bhattacharya et al., 2015). In 2019, the national  
16 neurosciences advisory group highlighted that people with the most complex  
17 needs often get the most fragmented care from the NHS (National Neurosciences  
18 Advisory Group, 2019). Fragmented care was reported in the current study.

19 The importance of close MDT working and harnessing the skills of a mix of allied  
20 health professionals (AHPs) within neurological rehabilitation settings is widely  
21 acknowledged (Chard, 2006; Health Improvement Scotland, 2012; Turner-Stokes,  
22 2008; Wade, 2015). However, participants noted that MDT working with people with  
23 FND was often prevented by lack of resources (staff, time and funding), resulting in  
24 varying waiting times for different professions, meaning that many occupational  
25 therapists often saw people with FND in isolation, before or after their physiotherapy  
26 and psychology colleagues. This problem was amplified in the teams who had limited  
27 access to psychological services or who were working in rural and geographically  
28 isolated areas where resources were diluted. Insufficient resources to provide timely  
29 treatment may lead to worse outcomes. Chronicity of symptoms has been linked with  
30 worse functional outcomes, greater rates of unemployment and uptake of disability  
31 related benefits (Carson et al., 2011; Sharpe et al., 2010). Participants also highlighted  
32 the benefits of joint working with GPs to improve continuity of care across the treatment  
33 pathway. This included offering training to GPs about FND management which has

1 been shown to result in a reduction in GP visits for people with somatoform disorders  
2 (Rief et al., 2006).

3 Participants working in services with limited psychological support raised concerns  
4 regarding lack of professional role clarity between occupational therapy and  
5 psychology. It was common for occupational therapists to inadvertently take up  
6 aspects of the psychology role (e.g., strategies for managing anxiety and low mood),  
7 in order to move forward with their own treatment plan. Edwards et al. (2012) also  
8 highlighted this as an issue for neuro-physiotherapists, indicating that they needed to  
9 take a greater supportive role for patients when psychology was unavailable within  
10 their service. However, it was widely recognised by the occupational therapy  
11 participants that they felt unable to manage people with significant psychiatric illness,  
12 such as post-traumatic stress disorder. This stance is supported by Nicholson et al.,  
13 2020 who highlighted that although occupational therapists are dually trained in  
14 physical and mental health rehabilitation, not all are well equipped to manage the  
15 sequelae of serious mental health problems and psychological trauma. Additional  
16 training, accreditation and onward referrals to mental health services may be required.

17 This study highlighted significant variability in the provision, quality and access to  
18 community-based neurological occupational therapy for people with FND depending  
19 on geographical location. Similar problems have also been found in the provision of  
20 community services for people with long-term neurological conditions in London, with  
21 service provision between boroughs being inconsistent and inequitable (Siegert et al.,  
22 2014). Occupational therapists working within specialist FND pathways reported the  
23 least number of barriers to effective service delivery. Within these services, referrals  
24 were received directly from a neurologist post diagnosis and after having received a  
25 thorough explanation. The teams had ongoing access to multidisciplinary support  
26 including neurologists, psychiatrists and psychologists. Intervention periods were  
27 flexible with discharge determined by the therapists and patients, based on patient  
28 identified goals and goal achievement. Participants from these services reported  
29 feeling more confident and competent in their abilities to provide quality care to people  
30 with FND, citing a high degree of job satisfaction. Neuro-science and mental health  
31 experts in the UK agree that it is important for community services to have access to  
32 specialists in neurology and mental health as it helps to facilitate the development of  
33 integrated and holistic treatment pathways whilst 'operating and delivering services in

1 silos results in poor patient experience' (National Neurosciences Advisory Group,  
2 2019).

3 Although occupational therapists reported high motivation to work with people with  
4 FND, they found the work challenging due to limitations in knowledge and a lack of  
5 published information to guide practise. The absence of occupational therapy  
6 treatment guidelines and limited access to affordable specialist training for FND was  
7 highlighted as a barrier. Participants reported that their knowledge of other  
8 neurological conditions is greater as there are established guidelines to direct their  
9 practice e.g. National Clinical Guideline for Stroke, 2016 (ISWP, 2016). To support  
10 FND specific knowledge and learning occupational therapy consensus  
11 recommendations for FND have recently been published (Nicholson et al., 2020) and  
12 The Functional Neurological Disorders Society (<https://www.fndsociety.org/>) has been  
13 established. Despite having a lack of documented guidance at the time of this study,  
14 practice descriptions as described by participants are largely in line with those  
15 suggested in recent publications specific to occupational therapy for FND (Gardiner et  
16 al., 2018; Nicholson et al., 2020). Similarities were also found between the  
17 interventions undertaken by the community neurological occupational therapists in this  
18 study and occupational therapy interventions for FND described in in-patient settings  
19 (Demartini et al., 2014; Jacob et al., 2018; Jordbru et al., 2014; McCormack et al.,  
20 2014; Saifee et al., 2012).

21

## 22 **Implications for Policy and Practice**

23 The study has highlighted barriers and enablers to effective community-based  
24 neurological occupational therapy, including reduced MDT support, lack of guidelines  
25 and evidence base to guide interventions and delays to treatment which may lead to  
26 worse patient outcomes. Despite these challenges, community-based neurological  
27 occupational therapists remain largely positive about the benefits of occupational  
28 therapy interventions for people with FND. The findings suggest that there is a need  
29 for better access to education about FND and /or specialist community treatment  
30 pathways for FND which include occupational therapy. Integral to this model of care  
31 would be timely and co-ordinated MDT treatment by clinicians who have developed  
32 an interest, specialist knowledge and skills in FND. A more coordinated approach to



1 commissioning physical and mental health services is required to adequately address  
2 the bio-psychosocial components of this condition and to improve the disjointed  
3 access to services that has been highlighted in the findings. Finally, there is a need  
4 for improved channels of communication between neurology, psychiatry, general  
5 practice and allied health professionals for patients with complex needs in order to  
6 improve continuity of care throughout the treatment pathway.

7

## 8 **Strengths and Limitations**

9 The structured and systematic approach of the TDF has uncovered individual and  
10 sociocultural barriers that may not have been otherwise identified with other methods  
11 (e.g., blurring of roles between occupational therapy and psychology, reduced  
12 opportunity for MDT working secondary to the holding of separate waiting lists).

13 A limitation of this study is a small sample size that was confined to UK-based  
14 clinicians. However, thematic saturation was achieved. Recruiting participants via the  
15 OT special interest group SSNP UK, enabled targeted purposive sampling of key  
16 stakeholders. This, along with inclusion of occupational therapists from different  
17 geographical locations increases the generalisability of the findings, ensuring that  
18 variations in knowledge, scope of service and funding provision within the UK are  
19 captured.

20 The nature of interventions described in the study are largely in line with recently  
21 published occupational therapy consensus recommendations for FND which was a  
22 collaboration of international clinicians, representing England, Scotland and the US.  
23 This suggests that findings may have some international relevance.

24 The results could be strengthened through exploration of the barriers and enablers to  
25 community occupational therapy from the perspectives of people with FND and their  
26 significant others. Future work should explore the content and evidence for  
27 occupational therapy in different settings, including inpatient rehabilitation, acute and  
28 community care services.

## 29 **Conclusion:**

1 Functional Neurological Disorders are common, disabling and associated with high  
2 levels of health care utilisation, yet funding for rehabilitation remains relatively low.  
3 This study has identified multifactorial, inter-dependent barriers and enablers to the  
4 delivery of neurological occupational therapy to people with FND in the community.  
5 There was significant variability in the provision, quality and access to occupational  
6 therapy depending on geographical location. Although the participating occupational  
7 therapists reported that they enjoyed working with people with FND, resource  
8 limitations limited the quality of the service that they were able to provide. The findings  
9 suggest that many existing community service structures are inadequate and  
10 additional resources are required to support occupational therapists delivering  
11 treatment to people with FND in the community.

12

### 13 **KEY FINDINGS:**

14 *Key Enablers to community occupational therapy for people with FND were:*

- 15 - Access to training to increase knowledge and skills
- 16 - Access to the website [www.neurosymptoms.org](http://www.neurosymptoms.org) and key papers
- 17 - Teaching patients self-management strategies to decrease reliance on  
18 therapists
- 19 - Dual mental and physical health training
- 20 - Core occupational therapy skills and values

21 *Key Barriers were:*

- 22 - Lack of available resources (staff and funding)
- 23 - Varying waiting times for different professions limiting access to MDT working
- 24 - Lack of access to psychological services leading to role blurring between  
25 occupational therapy and psychology
- 26 - Lack of guidelines and evidence base for occupational therapy assessment  
27 and treatment

### 28 **What the study has added:**

- 29 - Key barriers and enablers to the delivery of occupational therapy interventions  
30 for people with FND are identified together with areas of need for future  
31 service provision and commissioning.
- 32 - An identified need for the development of guidelines/recommendations to  
33 support occupational therapy assessment and treatment of people with FND.
- 34 - Illustrates a theory-based approach to exploring influences on clinical practice  
35 as a basis for designing healthcare quality improvement interventions.

36

37

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1 **Supplementary Files**

2 **Supplementary File 1:**

3 **Final Interview Guide**

4 **Topic Guide**

5 *Reported barriers and enablers to providing community-based occupational therapy*  
6 *to patients with functional neurological symptoms: an interview study with occupational*  
7 *therapists in the UK.*

8

9 **Knowledge and guidelines:**

10 1/ Could you tell me what your understanding of the diagnosis of FNS is and  
11 how you have developed this understanding?

12

13 2/ Have you come across any resources that you have used for your own  
14 knowledge or for use with patients regarding FNS? *(Can you tell me about*  
15 *those and whether you found them helpful? What information would have been*  
16 *more helpful?)*

17

18 3/ Are you aware of any guidelines or evidence base about Occupational  
19 Therapy intervention for patients with FNS? *(If so, what do you think of them /*  
20 *are they helpful? If not, what do you think they should recommend?)*

21

22 **Referrals:**

23 4/ Approximately how often would you say your team receive referrals for  
24 patients with this diagnosis?

25

26 5/ As an OT how frequently do you see patients with this diagnosis in the  
27 community?

28

29 6/ What professionals in your team see patients that are referred with a  
30 diagnosis of FNS?

31

32 7/ Who do you routinely receive referrals from for patients with FNS?

33

34 8/ When you receive a referral to see a patient with FNS, how do you and your  
35 team decide whether to accept the referral or not? *(Are there any factors that*  
36 *influence this decision? Do you make that decision alone or is it a team*  
37 *decision?)*

38

39 9/ Are there other professionals or teams that you think should be involved that  
40 currently are not? *(Can you tell me a bit more about that?)*

41

42 10/ Do you feel any pressure from internal or external colleagues to accept or  
43 decline referrals for patients with FNS? *(If yes, can you elaborate on that a*  
44 *little?)*

45

1 11/ Within your team are referrals to see a patient with FNS well received?  
2 *(Please elaborate.)*  
3

4 **Interventions:**

5 12/ When treating patients with FNS do you generally do this in isolation or as  
6 part of an MDT? *(Why is this?) If part of an MDT is there a team process or*  
7 *pathway in place for patients with FNS? Can you tell me a bit more about that?)*  
8

9 13/ Before seeing a patient with FNS how do you know if they are involved with  
10 any other health care professionals or if they have been in the past? *(Can you*  
11 *tell me a bit more about how you come to that knowledge? If so, does it change*  
12 *your interventions at all?)*  
13

14 14/ As an OT how do you decide what to work on with the patient? Do you follow  
15 a sequence of steps?  
16

17 15/ What sort of interventions do you routinely use with patients with FNS in the  
18 community? *(How did you come to the decision as to what type of interventions*  
19 *to undertake?)*  
20

21 16/ When delivering therapy do you tend to include the patients' significant  
22 others in the process? *(If so, how so? If not, why not?)*  
23

24 17/ In your experience how do you think the patients social and physical  
25 environment impacts on their ability to participate in therapy? *(Can you tell me*  
26 *a bit more about that?) Does this change your intervention approach at all?)*  
27

28 18/ What challenges have you encountered when delivering occupational  
29 therapy to patients with FNS? *(Can you talk me through these....how did you*  
30 *overcome these?)*  
31

32 19/ As an OT how easy or difficult do you find it to provide occupational  
33 therapy to patients with FNS? *(What in particular do you find easy/difficult? Is*  
34 *this for all patients with FNS or does it vary across patients?)*  
35

36 20/ Do you enjoy working with patients with a diagnosis of FNS? *(If so why, if*  
37 *not, why not? If not do you think that it influences your therapeutic relationship*  
38 *or ability to develop rapport with your patient?)*  
39

40 21/ As a neuro OT to what extent do you consider providing therapy to patients  
41 with FNS to be part of your role?  
42

43 22/ Compared to other things you have to do as part of your role, where would  
44 you rank providing occupational therapy to patients with FNS in terms of  
45 priority?  
46

47 23/ Is providing care for this patient group something you would like to do more  
48 or less of? *(Can you please elaborate on that...)*  
49



1 24/ Do you think your intervention methods with this patient group have  
2 changed over time? *(If so, can you tell me a bit more about why you think this*  
3 *is the case?)*

4  
5 25/ What do you think helps or hinders patients with FNS to carry out their set  
6 OT goals in the community?  
7

8 26/ How is your approach to treating patients with FNS different to what you  
9 may use when treating someone with another neurological condition e.g MS or  
10 stroke?  
11

12 27/ What do you think are the specialist skills or knowledge that are important  
13 for OT's to have when working with patients with FNS? *(To what extent do you*  
14 *feel you are equipped with such specialist skills/ knowledge?)*  
15

16 28/ What do you think would be the best way to help OT's build their skills and  
17 knowledge around working with patients with FNS?  
18  
19

### 20 **Community Therapy:**

21 29/ What do you think are the advantages of delivering occupational therapy to  
22 patients with FNS in the community setting?  
23

24 30/ If patients with FNS were not able to access occupational therapy services  
25 in a timely fashion what do you think might happen?  
26

27 31/ What resources do you feel are necessary to provide quality therapy to  
28 patients with FNS in the community? *(Do you feel that you have access to these*  
29 *resources?)*  
30

31 32/ Can you tell me your thoughts about the benefits and disadvantages of the  
32 provision of in-patient vs community therapy for patients with FNS?  
33  
34

### 35 **Support:**

36 33/ If a patient with FNS is distressed about their condition, how does this  
37 impact your intervention?  
38

39 34/ Can you tell me who you would go to if you were experiencing difficulties  
40 managing a patient with FNS?  
41  
42

### 43 **Supplementary File 2:**

#### 44 **Data Saturation Table**

45  
46  
47 **Table showing theme presence in each transcript with data saturation**  
48 **achieved by participant 9**  
49

	P1	P2	P3	P4	P5	P6	P7	P8	P9	P10
<b>Domain: Emotion</b>										
<b>T denotes theme (belief statement)</b>										
<b>T1</b> Therapists find lack of access to care for patients frustrating and upsetting	X	X						X		
<b>T2</b> Seeing patients with FNS is anxiety provoking for OTs	X								X	
<b>T3</b> Patients and families fear the diagnosis of FNS		X		X						X
<b>T4</b> I do (not) enjoy working with patients with FNS	X	X	X	X	X	X	X	X	X	X
<b>T5</b> Patients with FNS find the symptoms and resulting disability distressing	X	X				X		X		X
<b>T6</b> Working as an OT in the community can be stressful						X				X
<b>T7</b> Working with patients with FNS is tiring				X				X		
<b>T8</b> OT's enjoy working in the community setting with patients with FNS								X		
<b>Domain: Environmental context &amp; resources</b>										
<b>T1</b> We have access to the resources that are needed to treat patients with FNS	X	X	X	X	X	X	X	X	X	X
<b>T2</b> We don't have access to the resources that are needed to treat patients with FNS effectively.	X	X	X	X		X	X	X	X	X
<b>T3</b> Community neuro OT's in under-resourced teams have to be multi-skilled to meet the needs of the patient.						X		X		
<b>T4</b> The patient's physical environment impacts on their ability to participate in therapy	X	X	X	X	X	X	X	X	X	X
<b>T5</b> The patient's social environment can prevent engagement in therapy or encourage it.				X	X	X	X	X	X	X
<b>T6</b> Working in the community limits access to professional support	X					X				
<b>T7</b> Staffing levels vary from service to service		X		X	X	X	X	X		
<b>T8</b> Patient information is shared amongst services via an electronic notes system		X	X		X	X	X	X	X	
<b>T9</b> GP's are influential in funding community services		X								
<b>T10</b> The duration of the intervention period varies across services	X	X	X		X	X		X	X	X
<b>T11</b> There are no OT guidelines for treating patients with FNS				X	X	X	X	X		X
<b>T12</b> Resource pressures hinder MDT working in the community		X		X	X					X
<b>T13</b> Different waiting times for different services hinder MDT working in the community			X	X				X	X	X
<b>T14</b> Delivering OT to patients with FNS in the community is resource intensive				X						
<b>T15</b> Therapy delivered within the community is more embedded in the patient's real life					X	X	X	X	X	
<b>T16</b> OTs would like effective standardised assessment tools and outcome measures to use with patient with FNS				X	X					
<b>T17</b> Psychologists in some services will not see patients with FNS due to funding, resources, inadequate intervention periods.		X								X
<b>Domain: Knowledge</b>										
<b>T1</b> OTs are able to explain the diagnosis of FNS	X	X	X	X	X	X	X	X	X	
<b>T2</b> I am not aware of OT guidelines	X	X	X		X	X	X	X		X
<b>T3</b> There is a lack of evidence to inform practice	X			X	X	X			X	

<b>T4</b> Guidelines would help to develop the knowledge of OTs practice in this area	X			X						
<b>T5</b> Knowledge of the way that neurological symptoms arise and present is essential when working with patients with FNS to aid diagnosis and treatment	X		X	X	X	X	X	X		X
<b>T6</b> A patient's lack of understanding of the diagnosis and symptoms can impede recovery.	X		X		X	X	X	X	X	X
<b>T7</b> Understanding of the condition is important for OTs to deliver effective interventions (OT's are able to explain the diagnosis)	X	X	X	X	X	X	X	X	X	
<b>T8</b> There is an identified need to increase knowledge about FNS amongst OT staff	X				X					
<b>T9</b> FNS is a difficult diagnosis to understand for both clinicians and patients	X		X	X	X			X		
<b>T10</b> Community OTs must have knowledge of other services to link patients in with to provide holistic care		X				X				
<b>T11</b> Some neurologists do not have the knowledge to provide a diagnosis of FNS		X			X					
<b>T12</b> Some GPs do not have good knowledge of the condition						X		X	X	
<b>T13</b> Available guidelines are not always helpful when treating patients with FNS								X		
<b>T14</b> Community neuro OTs have developed their understanding of FNS via a number of avenues	X	X	X	X	X	X	X	X	X	X
<b>T15</b> The mind and body are linked in FNS				X	X	X			X	X
<b>Domain: Skills</b>										
<b>T1</b> Training has helped OTs to increase their knowledge and skills around FNS	X	X	X	X	X	X	X	X	X	X
<b>T2</b> Working with patients with FNS requires specialist skills	X	X	X	X	X	X	X	X		X
<b>T3</b> The identification of FNS requires specialist skills	X			X		X	X	X		X
<b>T4</b> OTs have identified training needs regarding working with patients with FNS	X	X	X	X	X	X	X	X	X	X
<b>T5</b> OTs do not feel they have the psychology skills to see patient's with FNS without psychology support						X	X			
<b>T6</b> Core OT skills mean that OTs are well placed to treat patients with FNS					X	X			X	X
<b>T7</b> Some OTs feel that they have to work out of their skill set to treat patients with FNS									X	X
<b>T8</b> Training opportunities are limited	X									X
<b>T9</b> There is an overlap of some skills between professionals when working with patients with FNS					X	X				
<b>T10</b> Specialist skills are required to work with patients with functional overlay			X				X			
<b>T11</b> Explaining the diagnosis requires specialist skills					X		X			X
<b>T12</b> Senior staff try to dispel negative beliefs and attitudes about patients with FNS through providing education to less experienced staff					X					
<b>Domain: Beliefs About Capabilities</b>										
<b>T1</b> OTs do (not) feel confident to question if a patient with FNS is suitable for the team to treat	X						X			
<b>T2</b> Community neuro OTs (do not) feel confident in their abilities to work with patients with FNS	X	X	X	X		X	X	X	X	X
<b>T3</b> Community neuro OTs (do not) feel confident in their teams' abilities to work with patients with FNS.	X	X	X	X				X	X	X

<b>T4</b> OTs feel that they (do not) have the capabilities to manage distress, anxiety and mental health issues in patients with FNS	X			X	X	X	X			
<b>T5</b> OTs find it difficult to see patients when they do not have an official diagnosis	X	X								
<b>T6</b> Community neuro OTs (do not) feel confident in the abilities of medical staff to diagnose and discuss the diagnosis of FNS				X		X		X	X	
<b>T7</b> OTs and teams with more experience of working with patients with FNS are more confident in their abilities to manage them successfully				X	X	X			X	X
<b>Domain: Social, Professional Role &amp; Identity</b>										
<b>T1</b> There is a lack of professional role clarity between OT and psychology in teams where there is little or no psychology support				X		X	X	X		X
<b>T2</b> OTs are (not) clear about when they need to seek the support of other health care professionals	X	X	X	X	X	X	X		X	X
<b>T3</b> Community neurological OTs do (do not feel) that they should see patients with FNS	X	X	X		X	X	X	X	X	X
<b>T4</b> Community psychologists do (do not) see it as their role to see patients with FNS		X				X	X			
<b>T5</b> There are certain interventions that are recognised as being in the realm of OTs		X	X	X	X	X	X		X	
<b>T6</b> Community neuro OTs take on an advocate role for patients with FNS	X	X	X			X	X	X	X	X
<b>T7</b> Community neuro OTs are aware of the roles that other professionals play in treating patients with FNS			X	X		X	X	X	X	X
<b>T8</b> Community neuro OTs feel that it is important to maintain professional identity when working with patients with FNS					X					
<b>T9</b> OTs and Physiotherapists are most often involved with patients with FNS in the community		X			X	X	X	X	X	X
<b>T10</b> Community neuro OTs feel that it is a neurologist's role to provide and discuss the diagnosis		X	X	X	X	X				X
<b>T11</b> There is a blurring of roles between professionals working with patients with FNS in the community						X		X		
<b>T12</b> Community neuro OTs that work in isolated areas feel pressure to take on more professional roles than those in more well serviced areas						X				
<b>T13</b> Patients with FNS are often seen as complex cases by community neuro OTs					X	X	X			X
<b>T14</b> In some services, junior staff will not see patients with FNS	X				X					
<b>T15</b> GPs see patients with FNS	X	X				X				
<b>Domain: Beliefs About Consequences</b>										
<b>T1</b> Patients require access to community OT in a timely fashion to maximise their potential for recovery (Outcomes are worse without it)	X	X	X	X	X	X	X	X	X	X
<b>T2</b> There are benefits of in-patient treatment for some patients with FNS	X	X	X	X	X	X	X	X	X	X
<b>T3</b> Better patient outcomes are achieved when delivering therapy in the community	X	X		X	X	X	X	X	X	X
<b>T4</b> When OTs run a mixed caseload it is difficult to develop expertise in any one area	X	X								
<b>T5</b> Patient's mental health issues and FNS are often related and may / may not be related to past physical or emotional trauma	X				X	X		X		X
<b>T6</b> Providing a diagnosis in a sensitive manner and way that patients can understand will lessen the chance of relapse and improve outcomes	X			X	X	X		X	X	X
<b>T7</b> Psychological input can be very beneficial for some patients with FNS	X	X								X

<b>T8</b> There are more patients in the community with FNS than the number that are referred for therapy	X									X
<b>T9</b> Patients without a diagnosis of FNS may not fit the criteria required for community therapy / some pts with FNS do not meet some team's criteria	X		X				X			X
<b>T10</b> It is difficult to deliver the right care to patients with FNS when teams or team members are generalists not specialists	X			X	X	X		X	X	
<b>T11</b> It is easier to deliver care to patients with FNS when teams are specialists not generalists	X				X			X	X	X
<b>T12</b> Having FNS will mean that the likelihood of recovery is poor (good)	X					X			X	X
<b>T13</b> Patients with FNS who have good insight and motivation are likely to have better outcomes	X		X	X		X	X			X
<b>T14</b> Some referrals lack adequate information / finding out additional information from other professionals about a patient that has been referred for therapy can be time consuming and unproductive (productive)	X		X			X		X		X
<b>T15</b> It is easier (more difficult) to work with patients with FNS if they and their significant others understand and are (not) on board with the diagnosis / supportive of treatment	X	X	X	X	X	X	X	X	X	
<b>T16</b> Adapting a patients' home (who has FNS) or providing equipment / care services / benefits can be beneficial (unhelpful)	X		X	X	X	X		X		X
<b>T17</b> It is important for OTs to be able to recognise when a patient is (is not) ready for therapy	X	X		X	X					X
<b>T18</b> Some patients with FNS will not do well (will do well) with a short intervention period	X	X		X				X	X	X
<b>T19</b> It is important for Neuro OTs to learn about FNS so that they can effectively treat patient's and share their knowledge with other staff	X			X	X	X		X	X	X
<b>T20</b> It is difficult for OTs to work with patients with FNS when they have not received an official diagnosis / easier to work with them when they have had a diagnosis		X	X	X	X					
<b>T21</b> OTs need to be careful with the language that they use when working with patients with FNS	X				X	X				X
<b>T22</b> Close MDT working and having MDT support is important when treating patients with FNS (organisational issues sometimes prevent this)		X	X		X	X	X	X	X	X
<b>T23</b> OTs find it difficult to treat patients with FNS without the support of psychology / mental health professionals		X	X			X	X	X		X
<b>T24</b> A lack of understanding of FNS can result in a negative view of patients with the condition and result in misinformation to the patient		X	X	X	X	X	X	X	X	
<b>T25</b> Some patient's mental health symptoms can make it difficult to engage in daily activities and community OT		X		X	X					X
<b>T26</b> Group interventions can be useful for patients with FNS		X		X						
<b>T27</b> Engaging in therapy can be very stressful for some patients with FNS and compliance can be affected				X						
<b>T28</b> Adjunct therapies can be helpful for some patients with FNS				X		X				
<b>T29</b> Outcomes for patients with FNS participating in community neuro OT vary from patient to patient				X	X	X	X	X	X	X
<b>T30</b> Therapeutic use of self and taking time to build the therapeutic relationship is important when working with patients with FNS					X	X		X		X
<b>T31</b> Treating patients with FNS can be taxing on the mind and body for community OTs					X			X		

<b>Domain: Motivation &amp; Goals</b>										
<b>T1</b> Not all OTs want to work with patients with FNS					X				X	
<b>T2</b> Some patients are very motivated to participate in therapy and make gains and some are not	X		X	X			X		X	X
<b>T3</b> Some staff members are keen to learn more about FNS and develop their skills	X	X	X							X
<b>T4</b> Seeing patients make functional improvements is highly motivating for OTs	X			X	X	X	X	X	X	X
<b>T5</b> OTs prefer to see patient's with FNS as part of a mixed caseload				X	X				X	
<b>T6</b> Flexible client driven OT goals are important for successful outcomes			X	X	X	X	X			
<b>T7</b> Goal setting is routinely undertaken by community neuro OTs			X	X	X	X	X	X	X	X
<b>Domain: Memory, Attention &amp; Decision Making</b>										
<b>T1</b> Only adequately funded services will accept referrals for patients with FNS	X	X	X	X		X				
<b>T2</b> Some services will see patients with functional overlay but not pure FNS			X							
<b>T3</b> Some mental health teams and psychology services will not see patients with FNS			X				X			
<b>T4</b> Community neuro OTs have to be able to explain their clinical reasoning when making clinical decisions	X								X	
<b>T5</b> Community neuro teams feel (do not feel) pressure to take on patients with FNS	X	X			X	X			X	
<b>T6</b> Community teams sometimes have to make decisions about whether to accept a patient or not based on little referral information or without a diagnosis	X	X		X		X		X		X
<b>T7</b> Undiagnosed patients with suspected FNS will be referred by community OTs for further consultations and tests	X		X		X	X			X	X
<b>T8</b> The decision of when to discharge a patient with FNS is multi-factorial	X	X	X	X	X	X	X		X	X
<b>T9</b> There are various sources of referrals of patients with FNS to community OTs	X	X	X	X	X	X	X	X	X	X
<b>T10</b> The number of referrals of patients with FNS varies across community OT services				X	X	X	X	X	X	X
<b>T11</b> Referrals are more likely to be accepted from trusted sources	X			X	X				X	
<b>T12</b> Team members on rota triage referrals and decide which referrals to accept and which waiting list to put them on	X				X	X	X			X
<b>T13</b> Those triaging the referrals may seek a second opinion from a colleague before accepting the patient					X					
<b>T14</b> When reviewing a referral, the decision is made as to whether that patient should be seen independently or jointly with another colleague	X	X	X	X	X	X	X	X	X	X
<b>T15</b> Consultations between community OTs and other HCPs may (may not) alter the treatment plan			X	X			X		X	X
<b>T16</b> Community neuro OTs have to clinically reason whether to provide aids and adaptations to patients with FNS	X		X	X	X	X		X		X
<b>T17</b> Treatment plans are often developed after discussions with the MDT	X			X		X	X			X
<b>T18</b> Community neuro OTs make decisions around prioritisation of patients	X	X	X		X		X	X	X	X
<b>T19</b> Deciding if a patient with FNS requires specialist in-patient care or specialist community services (e.g. psychiatry) requires complex consideration	X	X			X	X		X	X	X
<b>T20</b> When a patient with FNS is very distressed, community neuro OTs need to make a clinical decision as how that may be best managed	X	X			X	X		X	X	X
<b>T21</b> The patient's clinical need determines which clinicians in the team will review them		X	X		X		X	X	X	X

T22 Variation in treatment duration across community services affects decision making around treatment plans			X	X			X	X	X	X
T23 FNS can be identified through recognising incongruity between physical symptoms and function						X	X			X
T24 FNS may initially be diagnosed as something else						X				X
<b>Domain: Social Influences</b>										
T1 Working with other health care professionals regarding patients with FNS can be difficult		X	X			X				
T2 Mentorship and supervision from fellow OTs is valued but not always available	X	X	X	X			X	X	X	X
T3 Community neuro teams do not have the MDT support required to see patients with FNS			X				X	X	X	X
T4 Community neuro OTs draw support from other team members if OT supervision is not available				X	X	X		X		X
T5 Participation of patients' significant others in rehab process is (not) beneficial	X	X	X	X	X	X	X		X	X
T6 Working collaboratively with other healthcare professionals is essential in providing quality care for patients with FNS	X	X	X	X	X	X	X	X	X	X
T7 Family and clinicians with a lack of understanding of FNS may inadvertently hinder patient recovery	X				X	X			X	
T8 The way in which the diagnosis is delivered by the neurologist can be a positive or negative experience for patients with FNS		X	X					X	X	
T9 Community neuro OTs would like to have some influence over funding decisions and inclusion criteria			X							X
T10 Neuro services feel obligated / pressured to take on patients with FNS or suspected FNS as no other services will see them	X					X	X			X
T11 Community neuro OTs sometimes consult with other health care professionals involved with the patient before meeting	X	X	X	X	X		X	X	X	X
T12 Family members are sometimes involved in the therapy programme	X	X	X	X	X	X		X	X	X
T13 OTs can find it challenging to guide the clinical practice of others who are less experienced in the field		X	X		X	X				
T14 The use of open and consistent communication methods with patients with FNS and fellow team members is seen as an important intervention strategy with patients with FNS		X				X		X	X	X
<b>Domain: Behavioural Regulation</b>										
T1 Formal guidelines would be helpful to guide OT's new to the area of FNS	X		X					X		
T2 Supervision of junior OT staff by more senior OTs helps skill development	X		X		X					
T3 Current training and published papers are useful in guiding clinical practice for OTs working with patients with FNS		X			X					
T4 OTs teach patient's with FNS self-management principles to reduce reliance on their therapists			X					X	X	
T5 OTs need to employ reflexive strategies within clinical practice to improve their practice				X	X		X	X	X	
T6 OTs use a number of strategies to guide the behaviour of patients with FNS					X	X				
<b>Domain: Nature of Behaviours</b>										

T1 Intervention strategies for working with patient with FNS are (are not) different to those used with patients with organic illness	X	X		X	X	X	X	X	X	X
T2 OTs explain their intervention methods to patients in different ways					X					X
T3 The intervention strategies used by OTs with patients with FNS have changed as they have gained experience		X	X	X	X	X		X	X	X
T4 Education about the diagnosis and management strategies is a large component of OT intervention with patients with FNS	X		X	X	X	X	X	X	X	X
T5 The way that OTs explain the diagnosis may be different if it is to a patient or a clinician					X					
T6 There are some routine OT practices and techniques that are undertaken in the community with patients with FNS	X	X	X	X	X	X	X	X	X	X
T7 Community neuro OTs see patients with a variety of neurological and health conditions, as well as FNS			X	X	X					
T8 FNS can be intertwined with other organic symptoms	X							X		
T9 The presentation of FNS and recovery rates are different from patient to patient	X			X		X		X	X	X
T10 OTs routinely seek the advice and expertise of their colleagues when working with patients with FNS	X			X		X		X	X	X
T11 Community work means that community neuro OTs have to be able to work autonomously and creatively	X			X		X			X	
T12 Community neuro OTs are regularly seeing patients with FNS but also routinely run mixed caseloads		X	X	X						
T13 Community neuro OTs involve significant others in the rehab process in various ways		X		X	X	X	X		X	X
T14 The community environment influences the practice methods and approaches of community neuro OTs		X		X					X	
T15 Patients often have had input from a number of different specialities before coming in contact with their community neuro rehab team				X	X			X	X	X
T16 Some OTs see patients with FNS for maintenance sessions over time						X			X	X
T17 FNS are common and a huge problem in the community						X		X		

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3 **Supplementary File 3:**

4 **TDF domains and associated belief statements (1-12). Three examples from**  
5 **each domain are provided only.**

6 NOTE: For ease of representation sub-themes are highlighted in green

7 **1/ Knowledge**

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ Understanding of the condition is important for OTs to be able to deliver effective interventions.</b>	<b>OTs lack understanding of the condition</b>	E B (sub-theme 1)	9	<b>I think that kind of... often with people that have these conditions, have lots of physical symptoms as well, and I am not sure that mental health teams always have the necessary experience and</b>	...the knowledge that's needed, so that's first and foremost. (P08, pg. 28)



				<p>knowledge to be able to treat these people, which I think is a bit of a shame, really. (P03, pg. 5)</p> <p>I think having a good understanding of other factors, other than the physical symptoms, and having a good understanding of the influence of psychological and emotional factors, and social factors. (P09, pg. 22)</p>	
<b>2/ Guidelines would help to develop the knowledge of OT's practice in this area.</b>	There are currently no OT guidelines to inform OT practice with patients with FNS in any setting.	B	2	...most people who work with people with functional symptoms aren't specialist OTs like me, they're OTs in social services who've been referred somebody and don't know which end to start with. (So guidelines would be helpful) (P04, pg. 28)	
<b>3/Community neuro OTs have developed their own and their patients understanding of FNS via a number of avenues.</b>		E	10	My colleagues and I use the Neuro Symptoms website. We've used it for our own training purposes, and originally gathering knowledge, and we would use it with clients. (P09, pg. 3)	

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## 2 2/ Skills

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ Training has helped OTs to increase their knowledge and skills around FNS</b>		E	10	...we as a team have done loads of training to make sure that we always share in our understanding, so it's a common district wide understanding. (P08, pg. 12)	I think that having that training available so that you know that you are doing the best thing for that patient group. (P02, pg. 18)
<b>2/ Working with patients with FNS requires specialist skills</b>	Some staff members do not have those specialist skills	M B (sub-theme)	10	<p>I think many of the staff in the unit don't feel that they have the necessary skills to manage, so I think lots of people come, the junior staff that come to the service to work here, haven't come across it before and don't know anything about it.(P03, pg. 9)</p> <p>I think neuro OTs are well placed to see them, but I think that the OT has to realise that the techniques and the therapy techniques that they are using with this group are not going to be the same as somebody for example</p>	I think often the diagnosis is the make or break place, how it's given, and can be done wonderfully and supportively, or could actually make people fearful of professionals for life. (P08, pg. 4)

				with a stroke, or another brain injury. (P09, pg. 21)	
<b>3/ Core OT skills mean that OT's are well placed to treat patients with FNS</b>		E	4	But I think it draws on our mental health skills, I think it definitely draws on our neuro skills as well. (P10, pg. 15)	There is only so much college can do but I think what sets OT apart from everyone else are functions and activity analysis. (P06, pg. 35)

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## 2 3/ Social / Professional Role & Identity

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ There is a lack of professional role clarity between occupational therapy and psychology in teams where there is little or no psychology support.</b>		B	5	...so as OTs we've always got this tension between how much psychological intervention do we provide within the context of our profession and you know, I'd like that quite clarified as well, especially when working with the people who have got more trauma issues, how far do us OTs go with that. (P04, pg. 6)	
<b>2/Community neurological OTs do (do not feel) that they should see patients with FNS</b>		E B (do not feel)	9	<i>If I could I would hand it on to a mental health team. (P06, pg. 32)</i>  I absolutely think it is part of my role, and it is a funny thing because there has been this big debate over should neuro therapists be seeing these cases, and I find it quite difficult to see why they are not all neuro cases. (P05, pg. 24)	...so I think the bottom line is that this is a disorder that affects the way the nervous system functions and therefore is totally appropriate to see within the context of neuro rehab. (P05, pg. 23)
<b>3/ Community neuro OTs feel that it is important to maintain professional identity when working with patients with FNS</b>		E	2	I think ultimately, we must remember that we are actually still trying to enable people to achieve the occupations that they want and need, and are expected to do. (P05, pg. 5)	I think we would have to go back ... what it is we are trying to do as Occupational Therapists and not to lose sight of that (P05, pg. 5)

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## 4 4/ Beliefs About Capabilities

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
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<b>1/ Community neuro OTs (do not) feel confident in their abilities to work with patients with FNS</b>		E B (do not)	9	I don't find it any more challenging than any other patient group. (P02, pg. 12)  Well, I don't feel competent or skilled. I just look and I think, "Oh, god, how am I going to get through this one?" but it doesn't matter. (P06, pg. 30)	
<b>2/ OTs feel that they (do not) have the capabilities to manage distress, anxiety and mental health issues in patients with FNS</b>		E B (do not)	5	I think we're all quite equipped with the ability to manage the initial distress, in a sense, and how to question it and pick it out but I think you don't want to take it all on your own. (P01, pg. 31)  ...we don't necessarily feel we have the skill to deal with some of the psychological elements of the condition...(P07, pg. 15)	It's making a demarcation between where I feel competent and able to practice and saying, "No, I can't do that" and feeling the confidence to turn around and say I can't do that, because that can be quite challenging as well. (P06, pg. 29)
<b>3/ OTs find it difficult to see patients when they do not have an official diagnosis.</b>		B	2	...it's when patient's don't have a clear diagnosis is when there can be problems...(P02, p.g 12)	

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## 2 5/ Environmental Context & Resources

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ We don't have access to the resources that are needed to treat patients with FND effectively.</b>		B	9	So, sometimes people have to wait until the other waiting list has come down a bit before we can see them jointly, or try and see them individually which sometimes creates a bit of a barrier really, and not always the best for that person. P03, p.g 20)	So I think there needs to be a bit more funding, and there needs to be a bit more training and I think, and we'd definitely need a consultant for MDT really, to support these people. (P03, p.g 12)
<b>2/ Working in the community limits access to professional support and can be isolating</b>		B	2	Whereas in the community, you haven't got that luxury; you're on your own, you're in the moment and you might have to wait a week before you see the person again and you might not see a senior for two days because you're at one office hub and in and out. (P01, p.g 29)	

3/ Different waiting times for different services and resource pressures hinder MDT working in the community		B	7	"It always end up that physio has to opt out for three months so why don't we just put you in first" dependent on resources. (P08, p.g 17)	Because we have separate waiting lists for OT and physio and speech, it doesn't always coincide but with complex people we definitely try and do joint visits when we can and joint assessments because that's more beneficial. (P10, p.g 21)
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## 2 6/ Beliefs About Consequences

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
1/ Patients require access to community OT in a timely fashion to maximise their potential for recovery.	Outcomes are worse without it.	M B (sub-theme)	10	I suppose, increased dependency, possibly greater need for package of care, greater carer burden on family and friends and social isolation with potentially their mood deteriorating. (P02, pg. 18)  ...people would return to work more quickly, they could return to a full and active life more quickly, they'd manage their families, they'd manage their responsibilities, they'd maintain their roles much more effectively and they wouldn't end up just kind of going round and round and round hospital appointments looking for an answer that they're not getting. (P04, pg. 30)	
2/ It is important for Neuro OTs to learn about FNS so that they can effectively treat patient's and share their knowledge with other staff.		E	7	If neuro OTs are trying to apply the same techniques in a "one size fits all", I think that that is why people that come to us who have been to other services in the past haven't benefited. (P09, pg. 21)	I think it's really important to be quite knowledgeable about this condition and understand some of these strategies that we might use, because sometimes the strategies just unlock the movement of the action you are really looking for. (P05, pg.12)
3/ Close MDT working and having MDT support is important when treating patients with FNS	(organisational issues sometimes prevent this)	E B (sub-theme)	7	I think it's because like with lots of conditions, people have very variable needs and I think that not one professional is more, has all the skills to meet all of those needs, so I think that there is lots of evidence for all	We'd need support from the MDT as well, because I think that's very key, that it needs to be an MDT approach. (P03, pg. 20)

				<p>conditions an MDT approach can be more helpful. (P03, pg. 20)</p> <p>So, sometimes people have to wait until the other waiting list has come down a bit before we can see them jointly, or try and see them individually which sometimes creates a bit of a barrier really, and not always the best for that person. (P03, pg. 20)</p>	
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## 2 7/ Motivation and Goals

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ Not all OTs want to work with patients with FNS.</b>		B	2	So, yes, I think it depends on the training and the expertise of the OT, and also the willingness because I can see that this group would not be every OTs preferred group to work with. (P09, pg. 21)	
<b>2/ Some staff members are keen to learn more about FNS and develop their skills.</b>		E	4	So, it's not a patient group that you think, "Oh, yeah, we're going to see really good gains for me"; it's a patient group that it's more that I just find interesting and want to develop my skills in. (P01, pg. 11)	So, it's quite good MDT availability and everyone is quite keen to learn, is the key. P01, pg. 7)
<b>3/ Flexible client driven OT goals are important for successful outcomes.</b>		E	5	...well I think with any patient group you've got to really get it right with goal setting because you know, if people don't set the right goals for themselves it can just put them off and make them even more miserable and unhappy and feel even bigger failures and I don't want that, I want people to have really positive experiences and to feel that they're moving forwards. (P04, pg. 18)	I think that's really fundamental to goal setting because if it is clinician driven then obviously it's less likely to be as successful. (P03, pg. 10)

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## 4 8/ Memory, Attention & Decision Processes

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ Some services will see patients with functional overlay but not pure FNS.</b>		B	1	...our service has decided that we don't commission people with that diagnosis alone, so it is only the overlay. (P03, pg. 5)	

<b>2/ Some mental health teams and psychology services will not see patients with FNS</b>		B	2	So, it's almost like, sometimes they don't want to take people with this diagnosis either and then there is a bit of a gap, really. (P03, pg. 5)	
<b>3/ Variation in treatment duration across community services affects decision making around treatment plans</b>		M	6	We do have some people that we can have very good results within eight weeks, but with others I think you would have to question whether getting involved at all is just a waste of time and money, if you can only provide six sessions, or eight sessions. (P09, pg. 12)	

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## 2 9/ Social Influences

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/Community neuro OTs draw support from other team members if OT supervision is not available.</b>		M	5	I've got very good mentors who are psychologists on the mainland who I can phone up and say, "I don't know what to do" and they're very, very supportive. (P06, pg. 6)	I don't know how I'd manage without them. (P06, pg., 6) (support of Psychologist on the mainland)
<b>5/Participation of patients' significant others in rehab process is (not) beneficial</b>		E B (not beneficial)	9	Yeah, I think when someone has got a social environment that is very supportive, it's a very positive thing but you want to make sure they're not too supportive and they're allowing the person freedom to engage and participate in things that may be considered risky or difficult but doing it in a safe way. (P01, pg. 19)  ...sometimes I have found that family members can be quite negative in their feedback to the person, so when you are saying, "Look, let's try and do this, you want to be able to do this, so we'll take these steps and do this, this and this" and a family member is always, "Oh, but ...". There is always a negative involved in it, that they are only seeing the negatives rather than the positives. (P09, pg. 17)	
<b>3/Working collaboratively with other healthcare professionals is essential in providing quality care for patients with FNS</b>		M	10	I think in the team that I am working in now it works very well as we tend to all take the same approach and I think that that works well with any patient if everyone is being consistent but I think that that has worked really well with the team that I am working with now. (P02, pg. 9)	

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2 **10/ Emotion**

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/ Seeing patients with FNS is anxiety provoking for OTs</b>		B	2	I think that I know I felt anxious initially because in your head you're having this battle with, "They've got this weakness but it's not neurological", there's not an organic cause for it but you want the person to feel that you're there with them, you're on their journey and you're not trying to discredit what they're experiencing and what they're saying. (P01, pg. 12)	
<b>2/ I do (not) enjoy working with patients with FNS</b>		E B (not)	10	I worked with a lady recently that when we met her had been wheelchair bound for two years and she had intensive Physio and OT and within six weeks she was walking and I mean that is amazing and incredibly rewarding to be able to facilitate someone to be able to do that. (P02, p.13)  No, my heart sinks. (P06, pg. 30)	Absolutely love it; absolutely love it. (P08, pg. 25)
<b>8/ OTs enjoy working in the community setting with patients with FNS</b>		E	1	To me it's the obvious place to treat them, but then I am completely biased, and I'll admit to this in this research, I am biased toward community... because it rocks my boat really. (P08, pg. 29)	

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4 **11/ Behavioural Regulation**

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
<b>1/Formal guidelines would be helpful to guide OTs new to the area of FNS</b>		E	3	I think a guideline would be useful just to have a brief overview of FNS and might also be useful to have a not a step by step but an idea of how you might work through as an OT working with someone who is newly diagnosed FNS or longer standing to actually help you with your initial assessments and the pathway you might take with that client group. (P01, pg. 3)	
<b>2/ Supervision of junior OT staff by more senior</b>		E	3	So, myself and a colleague, the other seven, are just really keen for our juniors to have to access us, "Put something in my diary, book it in,	

OTs helps skill development				reserve it, we'll do joints, do doubles". (P01, pg. 29)	
3/OTs teach patient's with FNS self-management principles to reduce reliance on their therapists		E	3	So, we very much want it to be things that they can carry on with their family members, rather than having the image that you are going to be working on this goal, but you need a health professional to do it with you in your home. (P09, pg. 22)	

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## 2 12/ Nature of Behaviours

Theme (global)	Sub-themes	B/E/M	Freq	Quotes	Exp Imp
1/Intervention strategies for working with patient with FNS are (are not) different to those used with patients with organic illness.		M Are different  Are not different	9	I think also, I am very keen to acknowledge with them, one of the things I do is I acknowledge that I believe that their symptoms are real, and that I don't think they are making it up, and that they have a genuine problem. So, that is something I would probably never discuss with any other client group. (P05, pg. 11)  ...once we've worked out what it is then it becomes a case of restoring movement, restoring action, restoring confidence, restoring normal sleep patterns, reducing anxiety, whatever, and that can come through a fairly standard programme. (P06, pg. 32)	So, I'm very conscious of the language that I use around them. (P05, pg. 11)
2/Education about the diagnosis and management strategies is a large component of OT intervention with patients with FNS		E	9	I think we always continue to go back to discussing why this strategy might work, and why it might look like an unusual strategy to use, but why it will work, and always going back to the diagnosis and what keeps some of these symptoms going, and helping them to understand the whole way along why the intervention might work. (P05, pg. 11)	
3/There are some routine OT practices and techniques that are undertaken in the community with patients with FNS.		E	10	Um, the same that I would with any other patient group so it would depend on what their goals were, we might look at functional task practice, cognitive rehab, upper limb programs, community access so it would just depend on what their goal was. (P02, pg. 10)	I think obviously things like pain and fatigue management again, and best routine and structure are fundamental in trying to help. (P03, p.10)

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