

The qualitative DIGNISPACE study: The co-design of a life review, meaning-making and legacy leaving digital intervention for young people with life-limiting conditions

RODRIGUEZ, Alison M., KELLEHEAR, Allan, LANFRANCHI, Vitaveska, MCSHERRY, Wilfred, TATTERTON, Michael, WATTS MBE, Lucy, YAZIJI, Nahel and SMITH, Joanna

Available from Sheffield Hallam University Research Archive (SHURA) at:

http://shura.shu.ac.uk/32652/

This document is the author deposited version. You are advised to consult the publisher's version if you wish to cite from it.

Published version

RODRIGUEZ, Alison M., KELLEHEAR, Allan, LANFRANCHI, Vitaveska, MCSHERRY, Wilfred, TATTERTON, Michael, WATTS MBE, Lucy, YAZIJI, Nahel and SMITH, Joanna (2023). The qualitative DIGNISPACE study: The co-design of a life review, meaning-making and legacy leaving digital intervention for young people with life-limiting conditions. Illness, crisis & loss.

Copyright and re-use policy

See http://shura.shu.ac.uk/information.html



The Qualitative
DIGNISPACE Study: The
Co-Design of a Life Review,
Meaning-Making and Legacy
Leaving Digital Intervention
for Young People with
Life-Limiting Conditions

Illness, Crisis & Loss I-26 © The Author(s) 2023

Article reuse guidelines: sagepub.com/journals-permissions DOI: 10.1177/10541373231202842 journals.sagepub.com/home/icl



Alison M. Rodriguez¹, Allan Kellehear², Vitaveska Lanfranchi³, Wilfred McSherry⁴, Michael Tatterton⁵, Lucy Watts MBE¹, Nahel Yaziji⁶, and Joanna Smith¹

Abstract

Background: Young people with life-limiting conditions have unmet psycho-spiritual needs. Dignity Therapy is a psycho-therapeutic end-of-life intervention that addresses life review, meaning making and legacy leaving. Studies of Dignity Therapy including young people are limited. **Aim:** To co-design a digital Dignity Therapy-based intervention for young people with life-limiting conditions (DIGNISPACE). **Design:** A qualitative study including focus groups (n=5) with hospice-based healthcare professionals (n=23), semi-structured interviews with young people with life-limiting conditions (n=13) and family carers (n=12). Data were analyzed using framework analysis. **Results:** Three main themes were derived; intervention purpose,

Corresponding Author:

Alison M. Rodriguez School of Healthcare, Room 1.29 Baines Wing, University of Leeds, UK, LS2 9JT. Email: a.m.rodriguez@leeds.ac.uk

School of Healthcare, University of Leeds, Leeds, UK

²Dept of Nursing, College of Nursing and Health Sciences, and Dept of Medicine, The Robert Larner College of Medicine, University of Vermont, Burlington

³Devices for Dignity MedTech Co-operative and Dept Computer Science, Sheffield Teaching Hospitals Trust and University of Sheffield, Sheffield, UK

⁴Dept of Nursing, School of Health, Science and Wellbeing, Staffordshire University, Stoke-on-Trent, UK ⁵School of Nursing and Healthcare Leadership, University of Bradford, Bradford, UK

⁶Dept of Health Science and Population Research, Institute of Psychiatry, Psychology and Neuroscience, School of Mental Health and Psychological Sciences, Kings College London, London, UK

amendments to the Dignity Therapy question protocol and content for a digital application. Findings influenced the development of DIGNISPACE and a model of dignity in young people with life-limiting conditions. **Conclusions:** DIGNISPACE addresses the psycho-spiritual needs of young people with life-limiting conditions, including tenets of the model of dignity in young people with life-limiting conditions in its content and delivery.

Keywords

life-limiting conditions, young people, Dignity, qualitative, digital intervention

Background

Worldwide over 21 million children and young people are living with life-limiting conditions (Coombes et al., 2022). These include conditions where treatment may be faesible but can fail (e.g., cancer); conditions where a shortened life is inevitable (e.g., cystic fibrosis); progressive conditions that do not have curative treatment options (e.g., mucopolysaccharidoses) and irreversible conditions that cause severe disability (e.g., multiple disabilities following brain injury) (Together for Short Lives, 2023). Historically, children diagnosed at birth/childhood with a life-limiting condition had little hope for survival into adulthood. However, increasing numbers of children and young people are now living longer than expected and transitioning to adult healthcare. Studies suggest young people (18–24 years) with life-limiting conditions (young people) experience an increased complexity of needs, including unmet psycho-spiritual needs, and a higher healthcare burden as they move into adult services (Booth et al., 2018; Jarvis et al., 2022).

In adult palliative care settings, Dignity Therapy (Chochinov et al., 2005), an evidence-based psychotherapeutic intervention, is offered to support psycho-spiritual wellbeing. This is achieved through a process of life-review, meaning-making and legacy leaving. Dignity Therapists undertake one-to-one semi structured interviews with individuals, usually lasting no more than an hour, drawing on the The Dignity Therapy Question Protocol (Figure 1). These questions encourage a reflective discussion about valued memories, accomplishments, life roles and lessons, and future hopes for significant others. The interview is digitally recorded, transcribed, edited into a storied narrative and returned to the individual, who can suggest changes before the document is finalised. This may involve several meetings. The document is considered a generativity document or written legacy (Chochinov et al., 2005). Many people share their documents with others prior to death, providing opportunities for meaningful conversations that can assist bereavement processes (McClement et al., 2007).

A literature review exploring Dignity Therapy with young people (Rodriguez et al., 2018) found few studies had included young people. Most studies included people middle aged upwards with cancer conditions. More recently, studies have

The Dignity Therapy Question Protocol

1. Tell me a little bit about your life history, particularly the parts that you either remember most, or think are the most important. When did you feel most alive?

- 2. Are there specific things that you would want your family to know about you, and are there specific things you would want them to remember?
- 3. What are the most important roles that you have played in life (family roles, vocational roles, community roles, etc)? Why were they so important to you, and what do you think you accomplished in those roles?
- 4. What are your most important accomplishments, and what do you feel most proud of?
- 5. Are there particular things that you feel still need to be said to your loved ones, or things that you would want to take the time to say once again?
- 6. What are your hopes and dreams for your loved ones?
- 7. What have you learned about life that you would want to pass on to others? What advice or words of guidance would you wish to pass along to your (son, daughter, husband, wife, parents, others)?
- 8. Are there words or perhaps even instructions you would like to offer your family to help prepare them for the future?
- 9. In creating this permanent record, are there other things you would like included?

Figure 1. The dignity therapy question protocol (Chochinov et al., 2005).

explored the utility of Dignity Therapy for children and young people. The Dignity Therapy question protocol has been adapted for adolescents with cancer in Portugal (Julião et al., 2020), and a digital story board technique for children with cancer (Akard et al., 2020) has been created in the USA. In addition, Dignity Therapy has been used effectively in hospital settings in the USA, involving children, young people, and families (Schuelke & Rubenstein, 2020). Consultation activities undertaken in the UK, however, suggest that although Dignity Therapy could be a supportive intervention for young people, the approach would benefit from adaptations to better reflect their needs (Watts et al., 2020a & 2020b). As life-limiting conditions advance there are especially high burdens placed on families to support care (Ammann-Schnell et al., 2021; Veronese et al., 2015). The option of family engagement in Dignity Therapy for young people would acknowledge their diversity.

In exploring how and in what ways a Dignity Therapy-based intervention could meet young people's needs, the theoretical bases of Dignity Therapy need to be considered. Dignity Therapy is based upon the Dignity Conserving Model of Care (Chochinov et al., 2002). This model constitutes what dignity means to adults, drawn from the end-of-life experiences of middle aged and older adults with advanced cancer conditions. Dignity Therapy relates especially to several tenets of this model including generativity, the continuity of self, the maintenance of pride, hopefulness, role preservation, aftermath concerns, care tenor and social support.

Cai et al. (2023) explored the dignity model of conserving care for terminally ill children, from the perspectives of parents (n=11) of children with cancer conditions and healthcare professionals (n=14). Although the tenets of the adult dignity model (Chochinov et al., 2002) were deemed relevant for children, additional tenets were recognised. It was suggested that Dignity Therapy for children should further consider how stigma can cause psychological distress, and how acknowledging regrets and supporting the child's sense of security can assist coping. In addition, dignity in children can be supported by them having the company of loved ones, showing them 'fairness', exploring any unfinished wishes, resolving any family disputes, and having a decent and dignified death. It is plausible that the tenets of dignity found in both adult (Chochinov et al., 2002) and child (Cai et al., 2023) models of dignity may be relevant to young people.

The current study aims to co-design a life review, meaning-making and legacy leaving intervention for young people. DIGNISPACE will be a Dignity Therapy-based digital intervention; a way of delivering supported psycho-spiritual care acceptable to young people with a range of life-limiting conditions and their families. Through the co-design process dignity in young people will be explored. The sections that follow outline the study's methodology, a thematic discussion of findings, detail of a proposed model of dignity in young people and detail of the content of the Dignity Therapy-based digital app for young people 'DIGNISPACE', with a discussion and concluding section that draws together key points.

Methods

Design

A descriptive qualitative design, drawing on naturalistic inquiry (Erlandson et al., 1993) facilitated the co-design of DIGNISPACE.

The study includes two phases:

- 1. A qualitative exploration of dignity and Dignity Therapy for young people.
- 2. A small-scale qualitative user evaluation of the DIGNISPACE digital app prototype and the related question protocol.

An advisory group of eight young people and eight family carers supported the study. Reporting follows the COREQ Checklist (Tong et al., 2007).

Participants and Setting

The study was undertaken in the North of England, in a mixed urban-rural region. Participants were recruited from child and adult hospice settings due to the palliative and psycho-spiritual focus of the proposed intervention.

Participant inclusion criteria was as follows:

 Young people: 18–24 years; living with a life-limiting condition, can provide informed consent; access to internet/smartphone technology; no evidence of confusion/delirium based on clinical consensus.

- Family carers: family member of a young person with a life-limiting condition; directly involved in care; access to internet/smart phone technology.
- *Healthcare professionals*: Employed by a hospice; directly involved in care; access to internet/smartphone technology.

Sampling

Sampling was purposive as appropriate to the aims and qualitative design of the study.

Recruitment

Recruitment was facilitated by six hospices with young people caseloads. Hospice healthcare professionals were research gatekeepers, and shared study information documents with eligible participants. Recruitment was undertaken between April 2021 and December 2022.

Data Collection

In phase 1, participants were involved in semi-structured virtual interviews (young people) or virtual focus group discussions (healthcare professionals) via the ZOOM video conferencing digital platform. Participants were invited to share their views on dignity in young people and the acceptability of Dignity Therapy for young people. The Dignity Therapy question protocol (Chochinov et al., 2005) with some suggested amendments for young people (Watts et al., 2020a & 2020b) along with a basic mock-up of DIGNISPACE were shown to participants. Participants shared recommendations for question amendments and digital development.

In phase two, participants were invited to participate in semi-structured virtual interviews (young people and family carers) or face-to-face focus group discussions (healthcare professionals) at a regional hospice setting. Participants were invited to evaluate the co-designed DIGNISPACE question protocol and DIGNISPACE digital app prototype and suggest any necessary amendments.

Data Analysis

All data were digitally recorded, transcribed, and anonymised. Framework analysis (Smith & Firth, 2011) was undertaken on all data and included:

- Developing an initial coding frame from study aims, refined following review of the first focus group and two young person interview transcripts.
- The coding frame was applied to the whole data set, with categories reviewed following team and advisory group discussions.
- Categories were condensed to form overarching themes and sub-themes.

Following analysis, the model of dignity in young people was developed and the DIGNISPACE question protocol and digital app prototype were finalised.

Ethical Considerations

The study received ethical review and approval from the School of Healthcare Research Ethics Committee, University of Leeds, UK. The study was approved 28/01/2021 (reference: HREC 20–005). Before any participant was recruited, governance approvals from participating organisations (hospice settings) were achieved. All team members are experienced in research with young people. AR (psychologist, Dignity Therapy trained) undertook interviews and chaired focus group discussions, supported by MT and JS. Processes were in place to support participant withdrawal and distress/safeguarding need.

Results

Participant Details

The study included 48 participants: 18 semi-structured interviews with young people (mean age 21 yrs.), 12 semi-structured interviews with family carers (8 siblings, 4 parents) and five focus groups with 23 healthcare professionals (13 nurses, 8 therapists, 2 physicians). Data collection was from April 2021 to January 2023 with interviews 40–90 min and focus groups 50–70 min in duration. Tables 1–3 detail participant characteristics.

Table	I.	Participant	Characteristics	Young People.
-------	----	-------------	-----------------	---------------

	Participant (Pseudonym)	Study phase	Age	Gender	Ethnicity	Religious orientation
ī	Alishba	I	19	Female	South Asian	Muslim
2	Charlotte	I	22	Female	White British	Christian
3	Shane	I	23	Male	White British	Christian
4	Simon	1&2	22	Male	White British	Christian
5	Nicholas	1&2	22	Male	White British	Christian
6	Luke	1&2	23	Male	Black British	Christian
7	Joshua	1&2	24	Male	White British	Christian
8	Julia	I	24	Female	White British	Christian
9	Darius	I	19	Male	Black European	Muslim
10	Abdul	1&2	24	Male	South Asian	Muslim
П	Violet	2	18	Female	Black British	Christian
12	Рорру	2	23	Female	White British	Christian
13	Eve	2	21	Female	White Eastern European	Christian

Table 2. Participant Characteristics Healthcare Professionals.

	Participant (Pseudonym)	Study phase	Focus group	Gender	Ethnicity	Role
I	Melanie	I	ı	Female	White British	Physician
2	Rebecca	I	I	Female	White British	Nurse ^a
3	Sarah	I	1	Female	White British	Physician
4	Jenny	I	2	Female	White British	Therapist ^b
5	Donna	I	2	Female	White British	Therapist ^b
6	Sally	I	3	Female	White British	Therapist ^b
7	Mandy	I	3	Female	White British	Therapist ^b
8	James	I	3	Male	White British	Therapist ^b
9	Helen	I	3	Female	White British	Therapist ^b
10	Dehlia	I	3	Female	White British	Therapist ^b
П	Annalise	1	3	Female	White British	Therapist ^b
12	Gemma	3	4	Female	White British	Nurse ^a
13	Louise	3	4	Female	White British	Nurse ^a
14	Beryl	3	4	Female	White British	Nurse ^a
15	Petra	3	4	Female	White British	Nurse ^a
16	Sue	3	4	Female	White British	Nurse ^a
17	Pearl	3	4	Female	White British	Nurse ^a
18	Chloe	3	5	Female	White British	Nurse ^a
19	Linda	3	5	Female	White British	Nurse ^a
20	Phoebe	3	5	Female	White British	Nurse ^a
21	Terri	3	5	Female	White British	Nurse ^a
22	Emma	3	5	Female	White British	Nurse ^a
23	Cara	3	5	Female	White British	Healthcare assistant

^aNurse, child, adult or learning disability trained.

Framework Analysis

Findings from phases 1 and 2 were analysed, deriving three overarching themes influencing the development of DIGNISPACE: 'intervention purpose', 'intervention components, content and delivery' and 'networks and support systems' (Table 4).

THEME 1: Intervention Purpose

Young people emphasised the potential of DIGNISPACE to help them make sense of their lives. Whereas healthcare professionals highlighted the value of an evidence-based tool in their practice to support young people's psycho-spiritual wellbeing:

^bTherapist including occupational therapy, clinical psychology, counselling, music therapy and chaplaincy.

	Participant (Pseudonym)	Age	Gender	Ethnicity	Religious orientation	Relationship
ı	Graham	27	Male	White British	Christian	Sibling
2	Finlay	30	Male	White British	Christian	Sibling
3	Maryam	26	Female	South Asian	Muslim	Sibling
4	Shafqut	53	Male	South Asian	Muslim	Parent
5	Ayman	48	Male	Black European	Muslim	Parent
6	Marie	57	Female	White British	Christian	Parent
7	Joyce	54	Female	White British	Christian	Parent
8	Mark	28	Male	White British	Christian	Sibling
9	Eleanor	23	Female	White British	Christian	Sibling
10	Michael	29	Male	White British	Christian	Sibling
П	Esmae	27	Female	White British	Christian	Sibling
12	Bob	30	Male	White British	Christian	Sibling

Table 3. Participant Characteristics Family Carers.

Table 4. Themes and Subthemes.

Themes	Subthemes
I. Intervention purpose	 Structured psycho-spiritual support Incorporating a DIGNISPACE question protocol Facilitating young peoples' agency
2. Intervention components, content, and delivery	 Practical considerations App design, functionality, and functions Inclusivity
3. Networks and support systems	Family supportPeer supportHealthcare professional support

'I think this would be a lovely tool for them... and I think in the adult hospice world there's a lot of holistic and spiritual support need as well and which I think they will see the value in this as well... brilliant' (Sarah, doctor).

Family carers perceived DIGNISPACE could meet the young person's needs, facilitating their agency. Participants in general, perceived ongoing psycho-spiritual support was essential to help young people see they have purpose, to make sense of life events and for them to have some control over how their legacy will be shaped.

Several young people talked about how they did not have an outlet for their troublesome thoughts and feelings. They had struggled to speak about their death anxiety and related wishes. Some had desires to talk about challenging topics in

depth but did not feel they had anyone they could easily access to discuss these topics with. They did not want to place any more upset on their families and to uphold their dignity they wanted to be able to have control over what they spoke about, when and to whom, for example:

'Talking about my life is something I've found difficult as I don't want to upset people around me. Having a space where I could share moments from my life good or bad would be a real weight off my shoulders mentally.' (Simon, young person).

Healthcare professionals suggested they needed tools/guidance to support conversations around challenging topics, acknowledging that young people live with the anguish of having things unsaid. A framework of questions would assist healthcare professionals to access young people's views and give them confidence to initiate and support discussions on challenging topics. Healthcare professionals considered the terminology and tenor of Dignity Therapy-based questions and how they would feel in delivering these questions to young people:

'Just thinking about working with a young person... when do you feel most happy and what inspires you... that feels very current and that feels comfortable to ask somebody that. '(Dehlia, therapist)

Young people suggested there could be pages within the proposed app to reflect the nature of associated questions. This would maintain user interest and understanding. Suggestions included: 'my identity', 'my top memories so far', 'remember me', 'my bucket list', 'making life better', 'my future messages' and 'final wishes'. Broad topic focused pages would provide an overarching framework for young people to structure their DIGNISPACE content. It was recognised however, that some of the questions would need to be more reflective of what dignity means to young people. Having a means to acknowledge challenges, reflect on growth, help others, share experiences, and offer words of wisdom to benefit others, were important to young people's dignity alongside having a sense of feeling that they have mattered and that their presence, memories and wishes will transcend death. The potential emotionality involved in working on the content of DIGNISPACE and considering legacies was recognised strongly by family carers. One parent highlighted how working with DIGNISPACE meant addressing sensitive topics, so there needed to be appropriate support available for young people:

'As soon as they start touching on them topics... the flood gates are going to open so... I think the phrasing needs to be sensitive and the support ready' (Shafqut, parent).

However, all participants agreed that DIGNISPACE would be a safe space for young people to record their perceptions and experiences and enable 'a working through' of all kinds of emotions. Participants suggested that DIGNISPACE would be an

important, motivational, and supportive intervention, identifiable as an activity that belonged to the young person. The proposal of DIGNISPACE belonging to the young person was deemed important for all young people regardless of the amount of support they may need from healthcare professionals or family carers to engage with the intervention, for example:

'I think it is important the activity is theirs... for some with neuro problems ownership might be difficult but the family can share that ownership... what is produced even if minimal, or if drawing on shared experiences would be extremely valuable' (Mark, sibling).

Where significant neurodegeneration was expected, healthcare professionals suggested that facilitating DIGNISPACE earlier would be beneficial. This would enable others to later get a better sense of how that person has lived, how they want to be remembered, their perceived identity, their interests and wishes.

THEME 2: Components, Content, and Delivery

Central to the DIGNISPACE app development were participant views about content, and practice delivery. There was some initial confusion amongst healthcare professionals with respect to how DIGNISPACE would be different to advance care planning. In sharing the Dignity Therapy question protocol and DIGNISPACE mockup, healthcare professionals could see how DIGNISPACE would be a supportive intervention separate to, but also complementing, advance care planning. Several of the healthcare professionals identified how DIGNISPACE would support young people and healthcare professionals to constructively discuss topics that may have therapeutic benefit. In practice there can be a struggle for young people to initiate talk and for healthcare professionals an absence of tools to assist timing and engagement, for example:

'It's not to replace the children and young people's advance care plan; it's the idea of those therapeutic conversations, that's what this is meeting, and other tools don't meet currently' (Petra, nurse)

Although advance care planning processes are individualised, young people perceived that meetings were not fully tailored to meet all their psycho-spiritual needs. As suggested by Violet, young people thought that engaging with DIGNISPACE would be very different to their experience of advance care planning:

'This is different to advance care planning, and I think more motivating... it's about wishes but it's about how life has gone...who am I... and what do I want to say... and pass on about myself, my experiences' (Violet, young person).

Family carers highlighted how advance care planning had given them reassurance that end-of-life care wishes would be put into place. However, there was recognition that there were still issues not being addressed. Family carers identified how DIGNISPACE would include questions that may address the young persons worries and to some extent their mental health difficulties. Joyce suggested that addressing worries and issues linking to young people's mental health was something needed and missing in current practice:

'Having someone who's trained to help young people talk though their worries and concerns is something I feel has been missing from mine and my son's life... we have a lot of medical support but no real support for his mental health' (Joyce, parent).

It was suggested that the introduction of DIGNISPACE would help people to recognise that young people are living with unmet psycho-spiritual needs. DIGNISPACE may enable a greater openness towards psycho-spiritual issues amongst families. However, to promote young person engagement, DIGNISPACE would need to have relevant content, functionality, and good aesthetics. The DIGNISPACE app 'mockup' was received positively by the young people in general, for example:

'To be honest... this is going to sound rude... but it is better than what I thought... Yes, I mean, it's really... it's got a really appealing interface' (Poppy, young person).

While participants perceived the DIGNISPACE question protocol would encourage deep thoughts and a range of emotional responses, they highlighted how linking to their social media would enable them to access and upload content, giving them a variety of memories to link to and reflect upon. They suggested a section of the app be dedicated to linking to 'their socials'. Julia was quite specific in how the app could be organised:

'Obviously you'd need a good navigation menu so you'd have the time capsule here (the time capsule incorporates all the DIGNISPACE questions and topic pages) with the key questions, you'd have your socials listed separate that you could lift content from' (Julia, young person).

The application could facilitate the uploading of different types of media. Participants welcomed the creativity of the app and discussed how some young people would enjoy working on and uploading artwork, music, and vlogs. The phrasing and language content of the app would need to be considered to maximise engagement. However, opinions differed around some proposed terminology. Healthcare professionals were more protective with their views, for example:

'These you know some of my top memories, and these are the things I want you all to remember, and these are all the things I want you to learn, and you know my words of wisdom kind of thing would be so wonderful because I do think it's a beautiful thing to do... I'm a little nervous about the idea of the bucket list, calling it that' (Melanie, doctor).

Whereas young people wanted directness and clarity. They suggested this was needed to help them to consider their answers and content for the sections that mattered to them, for example:

'Maybe you could put when did you feel at your worst and how did you bounce back or something like that... It sort of normalises real life emotions. It's not always incredible, you have your rubbish times... and everyone has a bucket list' (Nicholas, young person).

Family carers also outlined the importance of capturing the 'essence of the young person' through the questioning route, for example:

'It would be good for all families living this, when they lose some capacity, it can get harder and harder but working on this together can keep us thinking about who that person is, they are still there inside what is happening to them and it could become a focus for us all and help us through' (Michael, sibling).

Young people made further question suggestions, including acknowledging the importance of helping others and recording end-of-life and after death wishes. Participants suggested different DIGNISPACE formats be developed or for the app to have multiple functions, to meet the needs of all regardless of physical ability. Family carers stated how young people lived with the support of different types of assistive technology and that such would need to be considered for the app to be inclusive. Marie considered her daughter's condition, highlighting her current needs which included needing alt text:

'We would need alt text to be able to help my daughter with the digital app, but I think the digital app allows you and them the space to get on with it or think about it at least' (Marie, parent).

For young people with difficulties communicating orally, digital flexibility allowing typing and the uploading of various multimedia content was welcomed. To offer further inclusivity allowing the use of assistive technology such as eye gaze technology, and including emoticons and phrase banks would be enabling. In addition, incorporating a translate engine would support users where English is not their first language.

Participants identified how some questions would be more emotionally difficult than others, suggesting that there could be 'locked' sections/pages, and facilities to alert supporters to help young people to manage the more thought-provoking content from the outset (while still ensuring agency). Young people assumed

support would be available for the more 'difficult' questions and suggested ways that support could be triggered. Shane suggested an informal approach:

'(On alerting a supporter) Could it be a text? Yeah, like hello, are you okay, but very colloquial: are you okay, do you want to talk about ...? Yeah, is all good, are you happy with what you've written, if not call us or give us a shout' (Shane, 23 yrs., young person).

As a supported therapeutic intervention, the tenor of care being empathic, non-judgmental, encouraging, and respectful would be important to establish the therapeutic alliance between young person and supporter. The young person would need support to work through both the positive and negative revisiting of events and in thinking about their shortened life trajectories and wishes. Participants agreed the DIGNISPACE supporter would need training to be equipped to deliver support with appropriate tenor, offer guidance for content, and support pathways for referral, if needed, to engage in other care processes such as advance care planning, will writing or more specialist mental health support.

THEME 3: Networks and support systems

Participants' suggested support networks be embedded into DIGNISPACE. Across participants the role of family was highlighted as fundamental in supporting young people and therefore, in the broadest sense, families could be involved in aspects of DIGNISPACE and legacy leaving activities. Donna further highlighted the diversity of young people and families in her care and the implications of such for engagement with DIGNISPACE:

'I think some might need more help and support from families than others,

some just no communication, but they haven't got the cognitive ability to understand any of this process whatsoever, and that it might be useful for the parents to do it on their behalf, and then be useful for the parents', (Donna, therapist).

Healthcare professionals and family carers highlighted how DIGNISPACE could also meet parent's needs, in supporting memories of their child. There was an appreciation of how introducing DIGNISPACE to some families may be unsettling but would provide an opportunity to discuss with families the benefits of 'openness' and how enabling the recording of memories, meaning making and legacy leaving could ease worries. For example, some healthcare professionals discussed how young people may not just have concerns for how they may die, how they may be remembered, what has been their purpose, but also what may become of their families once they have died. Rebecca reflected on the needs of one young person historically in her care:

'Lot of his (young person) questions were all about who was going to look after his family, they weren't about him at all, or they were about how his family was going to be, who was going to look after the family. What were they going to remember', (Rebecca, Nurse).

Healthcare professionals also highlighted that DIGNISPACE could provide young people with a strategy for managing emotions following the death of a peer, or as a way of remembering them though shared memories. Jenny highlighted how young people need focused support and a means of sharing memories when one of their peers dies:

'They are now reflecting on the fact that one of their friends has died... they weren't able to say goodbye... they approached me to say, could we set something up' (Jenny, therapist).

Young people can feel isolated at home. However, they can make strong bonds with others from attending hospices and networking via social media. They can suffer multiple bereavements as their peers die. DIGNISPACE could facilitate memories with peers and engage the support of healthcare professionals to talk through their emotions and reflections as the immediacy of the death of others impacts on their own lived realities. This sentiment was also expressed by family carers, for example:

'We are a small but tight knit community, when someone's child passes, we all feel and need to talk about it' (Ayeman, parent).

The key role of healthcare professionals would be to provide support to young people and families with DIGNISPACE. Healthcare professionals suggested training was needed to ensure DIGNISPACE implementation was maximized. Helen suggested there would be training needed in how and when to introduce DIGNISPACE and managing support demands:

'I think that comes down to the therapists, well trained, well supervised in, knowing when's a good time and how to manage that conversation' (Helen, therapist)

The unpredictable nature of lives lived, and of times ahead, means young people are living amidst much uncertainty, working through lots of emotional challenges in silence. DIGNISPACE could enable an emotional outlet and satisfy many current unmet psycho-spiritual needs. It is important that support is accessible to work through what could be difficult thoughts and emotions and encourage a positive framing of events to be able to live in the moment and to be more comfortable with what may lie ahead.

Table 5. The Model of Dignity for Young People.

Illness related concerns - issues emanating from the health condition that impact the persons sense of dignity	Dignity conserving repertoire - the persons psycho-spi outlook, personality traits and internal resources that impact their perception of dignity	- the persons psycho-spiritual internal resources that nity	Dignity conserving repertoire - the persons psycho-spiritual Social dignity inventory - social issues, outlook, personality traits and internal resources that relationships and networks that influence the persons sense of dignity
Symptom distress: • Physical distress • Psychological distress • Medical uncertainty • Death anxiety • Stigma • Isolation Level of independence • Cognitive acuity • Functional capacity	Dignity conserving perspectives ways of coping with lived experiences Role preservation Generativity, legacy and remembrance Feeling they have mattered Maintenance of pride and voice (agency) Hopefulness Helping and remembering peers Informed and autonomy Acceptance of dying and death Resilience or fighting spirit Acknowledging regret Acknowledging regret Sense of security	Dignity conserving practices - resources that person can draw upon to strengthen their sense of dignity - Living in the moment - Normalcy - Seeking spiritual comfort - Seeking spiritual comfort - Seeking spiritual comfort - Seeking spiritual comfort - Explaining disease and death - Guiding to find the meaning - of life - Expressing love - Expressing lo	 Privacy boundaries Social support Care tenor Burden to others Aftermath concerns and wishes Aftermath concerns Addressing challenging conversations

The Model of Dignity for Young People with Life-Limiting Conditions

The model of dignity for young people with life-limiting conditions has been developed from the findings of this study, as highlighted in Table 5. Within this model, tenets of dignity relevant to young people with life-limiting conditions are highlighted in bold alongside the already identified tenets of dignity provided within the adult (Chochinov et al., 2002) and child (Cai et al., 2023) (provided in italics) models of dignity. The model of dignity in young people is considered within the DIGNISPACE question protocol and digital app prototype.

DIGNISPACE

The DIGNISPACE app prototype is developed as a React Native app accessible via Android and Apple devices, and web browsers. The DIGNISPACE question protocol is mapped to defined sections, broken down into questions and prompts to assist user reflection and meaning-making. To ensure flexibility, users can insert multi-media content when completing questions. The app is built around the idea of bubbles: bubbles are groups of people the user wants to share information with. The user can define different bubbles with any number of people and share sections of the app with them. Users can request support from a healthcare professional (supporter) from within the app, generating a support request alert. The same mechanism applies for sensitive questions; they can be unlocked; a message is then sent to the supporter and content then completed with the supporter.

The app has been developed using:

- 1. Database: Secure and encrypted data repository in the cloud; implemented using mongo DB installed on a university server.
- 2. API: Core element in the software that allows interaction between the database, the web app, and the mobile app implemented in the form of a RESTFUL API, directly linked with the back-end (server and database) and the front-end (mobile and web visualisation).
- 3. Web app: Application for users, accessible via web browser, to complete and review the questions.
- 4. Mobile app: Smartphone application for users to complete questions with text, voice messages, photos and/or videos.

The DIGNISPACE app prototype content incorporating the DIGNISPACE question protocol is summarised in Table 6.

Discussion

The study has co-designed a Dignity Therapy-based psychotherapeutic intervention for young people. DIGNISPACE (name suggested by advisory group and agreed by

Table 6. DIGNISPACE

App page

Content

My Time Capsule

Profile detail and access to all tabs.

My Bubbles My Social My Settings

My identity

Select bubbles.

Screen message: multimedia content? Directions

My top memories

so far...

Select bubbles.

Screen message: multimedia content? Directions

Remember me

My bucket list

Directions

Bubbles

Select bubbles.

Screen message: multimedia content? Directions

Screen message: multimedia content?

Drop down boxes:

 In the event of my death, I would like my supporter to access my DIGNISPACE and release content to hubbles

Selection of individuals for legacy content Links to other social media platforms

Inc links: emoticons, stock sentences, language translation engine.

Tell me a little about you/ your story...

 How do you see yourself? What are your likes/dislikes?
 When have you felt, or do you feel the happiest? What or who inspires you? Do you want to reflect on your condition?

Screen message: need help? Yes/No. message to supporter.

What are your most memorable moments?

 Are there any events/activities you can remember that you really enjoyed? What made that event/activity special?

Screen message: need help? Yes/No. message to supporter.

How do you think you will be remembered?

 How do you think your parents/family/friends will remember you? How would you like to be remembered?
 Would you like to be remembered differently by the different people in your life?

What has challenged you?

 Would you like to reflect on getting through that/ those challenge(s)? What was difficult? What do you feel most proud of?

What/who has made you happy in your life? Is there anything you want to tell people close to you about these things?

Screen message: need help? Yes/No. message to supporter.

Are there things/activities you would like to do?

Can we create a special time for you?

A special day/activity with family/friends? Art/music activity? **Screen message:** need help? Yes/No. message to supporter.

(continued)

Table 6. (continued)

My future messages

App page

Content

Making life better: sharing with and supporting others

Bubbles

Bubbles

Directions

Screen message: multimedia content? Directions

Screen message: multimedia content?

Is there anything you would like to share with others who may have your condition/a life-limiting condition?

To help them to make their life better?

- Services? transition, respite care, PAs,
- Leisure? holidays, sports/hobbies.
- Social support? peer groups, games, social media, blogs.
- Tips for others to make life better?

Screen message: need help? Yes/No. message to supporter.

Screen message:

Your supporter knows you have entered this area of DIGNISPACE. Please record your thoughts, your supporter will contact you to arrange a time to discuss/create content.

Message to supporter.

Is there anything you want to say or repeat to the people you care about?

• family? friends? PA? carer(s)? others?

Do you have any specific hopes or wishes for the people close to you?

family? friends? PA? carer(s)? others?

Do you have any advice that you would like to pass along to your friends and family?

family? friends? PA? carer(s)? others?

Are there any words or instructions you would like to offer your friends and family to help prepare them for life without you?

family? friends? PA? carer(s)? others?

Final wishes

Bubbles

Screen message: multimedia content? Directions

Screen message: Your supporter knows you have entered this area of DIGNISPACE.

Please record your thoughts, your supporter will contact you to arrange a time to discuss/create content. Message to supporter.

What are your final needs & wishes? Screen message:

You may decide to release your legacy content before you die. Your supporter can help you to consider conversations and any additional support you think you may need.

(continued)

Table 6. (continued)

App page

Content

Do you have any after death/funeral wishes?

Do you want to add detail about your will/legacy?

Do you want to add detail about where you would like to be after death?

Do you have any funeral wishes? Type of service? Music? Messages to share?

Is there anything that needs to be included in your DIGNISPACE that we have not talked about?

participants), considers what dignity means to young people and offers a solution to meeting some of the psycho-spiritual needs young people face in living with medical uncertainty and a shortened life prognosis.

DIGNISPACE is:

An activity to promote dignity, psychological and spiritual well-being. Young people with therapeutic support are encouraged to reflect upon life experiences, poignant memories and related meaning making. They are supported to share identity forming elements of 'their story', to voice 'their views', detail 'their wishes' and create content of sentimental value and guidance for significant others including family members, friends, and peers. The activity content and processes enable young people to engage in challenging conversations and explore the many ways in which they have had purpose and impact across their lifetimes.

DIGNISPACE and Psycho-Spiritual Needs

A systematic review of Dignity Therapy highlights its international appeal with studies undertaken in North America, China, Japan, Taiwan, New Zealand, Australia and Western Europe (Martínez et al., 2017). Several randomised controlled trials have explored its effectiveness compared to standard palliative care, with improved outcomes in anxiety, depression, quality of life, sadness, dignity and a sense of purpose (Chochinov et al., 2011; Hall et al., 2011; Juliao et al., 2014; Rudilla et al., 2016; Silva et al., 2019). It is possible that DIGNISPACE could reduce psycho-spiritual burden in young people to a greater effect than standard palliative care.

With increasing medical advancements young people are living longer sicker or with increased frailty (Cook et al., 2016). This impacts the complexity of their psychospiritual needs, particularly as they move to adult services with differing support available (Barker et al., 2019). A recent review of adult palliative care for young people

discussed themes linked to psycho-spiritual burden and adjustment difficulties, suggesting many young people (irrespective of location) receive no or very little specialist support (Abdelaal et al., 2023). We found emotional distress was mostly linked to feeling isolated, a loss of privacy and the fear of dying. Many participants coped by comparing their own condition to others or by drawing on the deeper meanings of their 'suffering' (Silva et al., 2019). To some degree humour and engagement with peers is helpful but there remains a need for tailored support (Jamalimoghadam et al., 2019; Mohammadi et al., 2023).

In addition to the considerations of Chochinov and colleagues (2002) and Cai and colleagues (2023) in their models of dignity conserving care and dignity in children respectively, the dignity of young people is impacted by several additional tenets. In considering illness related concerns leading to psychological distress, being or feeling isolated is relevant to this group. Dignity conserving practices including the need to preserve identity, feeling they have mattered, maintaining their voice and agency, helping, and remembering others and acknowledging the challenges they both have faced and overcome are important. In addition, dignity conserving repertoire practices for young people include the desire to not have things left unsaid. In terms of the social dignity inventory, it was important for several participants that not only would aftermath concerns be addressed but also their post death wishes be acknowledged alongside other challenging conversational needs.

It was important to participants that DIGNISPACE would be inclusive and flexible to the needs and abilities of all its potential users. The use of storytelling, narrative, and multimedia can enable creative DIGNISPACE content and help people even with profound and multiple disabilities to have their preferences acknowledged about their lives and wishes (Watson et al., 2019). It is in this way that DIGNISPACE offers a mechanism to give voice to young people who are seldom "heard" especially those with difficulties communicating their needs with words or whom may not have a shared first language with their formal carers (Liddiard et al., 2019; Spencer et al., 2020).

Young people enjoy arts-based activities for memory-making (Foster et al., 2012), but there are no structured tools for healthcare professionals to use to support allied meaning-making and legacy leaving. The Children and Young People's Advance Care Plan Collaborative (2023) supports healthcare professionals, providing resources, to plan and record end of life directives. However, the focus is on care choices. DIGNISPACE could enhance advance care planning, offering young people, families, and healthcare professionals a structured approach to preserving dignity, offering broader psycho-spiritual support, and incorporating much creativity.

Implications for Practice

Studies suggest end-of-life conversations are best introduced by palliative care teams (Sansom-Daly et al., 2020). Healthcare professionals with experience of working with young people could be trained to become DIGNISPACE supporters. DIGNISPACE

supporters would need to introduce and guide young people to use DIGNISPACE, help them to develop content, work with them through emotions and address adjunct challenging conversations. The digital app is designed to have inbuilt flexibility with functions to support physical needs and to work with family units if preferred. DIGNISPACE could strengthen partnerships between child and adult palliative care services and offer a new approach to community palliative care through transition.

Implications for Research

Through this study some of the challenges that healthcare professionals and families can face in supporting young people to address their psycho-spiritual needs have been identified and these warrant further investigation. The tenets of previously derived models of adult and child dignity are relevant to the participants of this study. However, further important tenets of dignity for this group have been identified and considered within a model of dignity in young people. This model of dignity could be used for the development of other dignity focused interventions for young people.

Further research is needed to explore the role of Dignity Therapy-based interventions in supporting young people at home and especially through transition to adult services. DIGNISPACE will be feasibility tested and the fidelity of training across child and adult hospice settings explored, examining the effectiveness of different models of healthcare professional support. This work will consider young people's behaviour change and engagement and how and in what ways communities of DIGNISPACE practice are formed.

Study Strengths and Limitations

To our knowledge this is the first study that has co-designed a Dignity Therapy-based intervention to be delivered digitally. As a research team we were guided by an expert by experience team member and an advisory group of young people and family carers. However, because hospices offer fewer short breaks compared to pre-pandemic service levels, our ability to promote the study to families, and to support recruitment was impacted. Sadly, several young people were unable to participate due to their health deterioration or death. However, a depth of data was still achieved.

Conclusion

Evidence based interventions to support young people's psycho-spiritual needs are limited. Dignity Therapy is a validated approach that young people view as acceptable, however, the question protocol and delivery of DIGNISPACE is more reflective of their lives lived and accessibility needs. DIGNISPACE is the first co-designed Dignity-Therapy-based intervention for young people that can be delivered digitally. It is proposed that DIGNISPACE will complement the excellent work already undertaken in hospices and palliative care teams by providing a supported, structured

intervention for young people (and where appropriate families) to engage in life reflection, meaning making and legacy leaving.

Acknowledgements

We would like to acknowledge Dr Mari-Cruz Villa-Uriol and Mr Jiaqi Fu, University of Sheffield, for their work assisting VL with digital design and coding. We would also like to thank all participants for sharing their experiences and perceptions and to members of the advisory group 'Striding Forward' for their consistent support and guidance. We would also like to thank Prof Harvey Max Chochinov for his support at the time of study design and set up, to use the Dignity Therapy materials. Finally, we would like to remember Miss Lucy Watts MBE, research team member and youth ambassador, whose ideas and strength of commitment to this work was invaluable.

Author's Note

Joanna Smith is currently affiliated with College of Health, Wellbeing and Life Sciences, Sheffield Hallam University, Sheffield and Sheffield Children's Hospital, Sheffield.

Declaration of Conflicting Interests

The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The authors disclosed receipt of the following financial support for the research, authorship, and/ or publication of this article: The research reported in this publication was supported by the National Institute for Health Research (NIHR) Devices for Dignity MedTech Co-operative. The project was funded by the National Institute for Health and Care Research (NIHR) under its Research for Patient Benefit (RfPB) Programme (Grant reference number NIHR201597). The views expressed are those of the authors and not necessarily those of the NIHR or the Department of Health and Social Care.

ORCID iDs

Alison M. Rodriguez https://orcid.org/0000-0001-9104-1999 Michael Tatterton https://orcid.org/0000-0002-7933-5182

References

Abdelaal, M., Avery, J., Chow, R., Saleem, N., Fazelzad, R., Mosher, P., Hannon, B., Zimmermann, C., & Al-Awamer, A. (2023). Palliative care for adolescents and young adults with advanced illness: A scoping review. *Palliative Medicine*, *37*(1), 88–107. https://doi.org/10.1177/02692163221136160

Akard, T. F., Wray, S., Friedman, D. L., Dietrich, M. S., Hendricks-Ferguson, V., Given, B., Gerhardt, C. A., Hinds, P. S., & Gilmer, M. J. (2020). Transforming a face-to-face legacy intervention to a web-based legacy intervention for children with advanced cancer.

 ${\it Journal~of~Hospice~\&~Palliative~Nursing,~22(1),~49-60.~https://doi.org/10.1097/NJH.~000000000000614}$

- Ammann-Schnell, L., Groeschel, S., Kehrer, C., Frölich, S., & Krägeloh-Mann, I. (2021). The impact of severe rare chronic neurological disease in childhood on the quality of life of families—a study on MLD and PCH₂. Orphanet Journal of Rare Diseases, 16(1), 211. https://doi.org/10.1186/s13023-021-01828-y
- Barker, M. M., Beresford, B., Bland, M., & Fraser, L. K. (2019). Prevalence and incidence of anxiety and depression among children, adolescents, and young adults with life-limiting conditions: A systematic review and meta-analysis. *JAMA Pediatrics*, 173(9), 835–844. https:// doi.org/10.1001/jamapediatrics.2019.1712
- Booth, A., Maddison, J., Wright, K., Fraser, L., & Beresford, B. (2018). Research prioritisation exercises related to the care of children and young people with life-limiting conditions, their parents and all those who care for them: A systematic scoping review. *Palliative Medicine*, 32(10), 1552–1566. https://doi.org/10.1177/0269216318800172
- Cai, S., Guo, Q., Lin, J., Deng, C., Li, H., & Zhou, X. (2023). The dignity of terminally ill children in pediatric palliative care: Perspectives of parents and healthcare providers. BMC Palliative Care, 22(1), 1–11. https://doi.org/10.1186/s12904-022-01116-x
- Child and Young Persons Advance Care Plan Collaborative. (2023). http://cypacp.uk/ (Last accessed 14.03.2023).
- Chochinov, H. M., Hack, T., Hassard, T., Kristjanson, L. J., McClement, S., & Harlos, M. (2005).
 Dignity therapy: A novel psychotherapeutic intervention for patients near the end of life.
 Journal of Clinical Oncology, 23(24), 5520–5525. https://doi.org/10.1200/JCO.2005.08.391
- Chochinov, H. M., Hack, T., McClement, S., Kristjanson, L., & Harlos, M. (2002). Dignity in the terminally ill: A developing empirical model. *Social Science & Medicine*, 54(3), 433–443. https://doi.org/10.1016/S0277-9536(01)00084-3
- Chochinov, H. M., Kristjanson, L. J., Breitbart, W., McClement, S., Hack, T. F., Hassard, T., & Harlos, M. (2011). Effect of dignity therapy on distress and end-of-life experience in terminally ill patients: A randomised controlled trial. *The Lancet Oncology*, 12(8), 753–762. https://doi.org/10.1016/S1470-2045(11)70153-X
- Cook, K. A., Jack, S. M., Siden, H., Thabane, L., & Browne, G. (2016). Investing in uncertainty: Young adults with life-limiting conditions achieving their developmental goals. *Journal of Palliative Medicine*, 19(8), 830–835. https://doi.org/10.1089/jpm.2015.0241
- Coombes, L., Braybrook, D., Roach, A., Scott, H., Haroardottir, D., Bristowe, K., Ellis-Smith, C., Bluebond-Langner, M., Fraser, L. K., Downing, J., Farsides, B., & Murtagh, F. E. M. (2022). Achieving child-centred care for children and young people with life-limiting and life-threatening conditions—a qualitative interview study. *European Journal of Pediatrics*, 181(10), 3739–3752. https://doi.org/10.1007/s00431-022-04566-w
- Erlandson, D. A., Harris, E. L., Skipper, B. L., & Allen, S. D. (1993). *Doing naturalistic inquiry:* A guide to methods. Sage.
- Foster, T. L., Dietrich, M. S., Friedman, D. L., Gordon, J. E., & Gilmer, M. J. (2012). National survey of children's hospitals on legacy-making activities. *Journal of Palliative Medicine*, 15(5), 573–578. https://doi.org/10.1089/jpm.2011.0447
- Hall, S., Goddard, C., Opio, D., Speck, P. W., Martin, P., & Higginson, I. J. (2011). A novel approach to enhancing hope in patients with advanced cancer: A randomised phase II trial of dignity therapy. *BMJ supportive & Palliative Care*, 1(3), 315–321. https://doi.org/10.1136/bmjspcare-2011-000054

- Jamalimoghadam, N., Yektatalab, S., Momennasab, M., Ebadi, A., & Zare, N. (2019). Hospitalized adolescents' perception of dignity: A qualitative study. *Nursing Ethics*, 26(3), 728–737. https://doi.org/10.1177/0969733017720828
- Jarvis, S., Flemming, K., Richardson, G., & Fraser, L. (2022). Adult healthcare is associated with more emergency healthcare for young people with life-limiting conditions. *Pediatric Research*, 92(5), 1458–1469. https://doi.org/10.1038/s41390-022-01975-3
- Juliao, M., Oliveira, F., Nunes, B., Vaz Carneiro, A., & Barbosa, A. (2014). Efficacy of dignity therapy on depression and anxiety in Portuguese terminally ill patients: A phase II randomized controlled trial. *Journal of Palliative Medicine*, 17(6), 688–695. https://doi.org/10.1089/ jpm.2013.0567
- Julião, M., Santos, A., Albuquerque, S., Antunes, B., Crujo, M., Sobral, M. A., Cancelinha, C., da Silva Soares, D., Fradique, E., Palaré, M. J., & Chochinov, H. M. (2020). Operationalizing dignity therapy for adolescents. *Palliative and Supportive Care*, 18(5), 626–631. https://doi.org/10.1017/S1478951520000589
- Liddiard, K., Runswick-Cole, K., Goodley, D., Whitney, S., Vogelmann, E., & Watts, M. B. E. (2019). "I was excited by the idea of a project that focuses on those unasked questions" co-producing disability research with disabled young people. *Children & Society*, 33(2), 154–167. https://doi.org/10.1111/chso.12308
- Martínez, M., Arantzamendi, M., Belar, A., Carrasco, J. M., Carvajal, A., Rullán, M., & Centeno, C. (2017). 'Dignity therapy', a promising intervention in palliative care: A comprehensive systematic literature review. *Palliative Medicine*, 31(6), 492–509. https://doi.org/10.1177/0269216316665562
- McClement, S., Chochinov, H. M., Hack, T., Hassard, T., Kristjanson, L. J., & Harlos, M. (2007). Dignity therapy: Family member perspectives. *Journal of Palliative Medicine*, 10(5), 1076–1082. https://doi.org/10.1089/jpm.2007.0002
- Mohammadi, F., Oshvandi, K., Khodaveisi, M., Cheraghi, F., Tehrani, T. H., Khalili, A., & Kyle, H. (2023). Caregivers' perception of teenagers' dignity in end of life stages: A phenomenological study. *Nursing Ethics*, 30(1), 121–132. https://doi.org/10.1177/09697330221085776
- Rodriguez, A., Smith, J., & McDermid, K. (2018). Dignity therapy interventions for young people in palliative care: A rapid structured evidence review. *International Journal of Palliative Nursing*, 24(7), 339–349. https://doi.org/10.12968/ijpn.2018.24.7.339
- Rudilla, D., Galiana, L., Oliver, A., & Barreto, P. (2016). Comparing counseling and dignity therapies in home care patients: A pilot study. *Palliative & Supportive Care*, 14(4), 321–329. https://doi.org/10.1017/S1478951515001182
- Sansom-Daly, U. M., Wakefield, C. E., Patterson, P., Cohn, R. J., Rosenberg, A. R., Wiener, L., & Fardell, J. E. (2020). End-of-life communication needs for adolescents and young adults with cancer: Recommendations for research and practice. *Journal of Adolescent and Young Adult Oncology*, 9(2), 157–165. https://doi.org/10.1089/jayao.2019.0084
- Schuelke, T., & Rubenstein, J. (2020). Dignity therapy in pediatrics: A case series. *Palliative Medicine Reports*, 1(1), 156–160. https://doi.org/10.1089/pmr.2020.0015
- Silva, G. C., Reis, D. C., Miranda, T. P., Melo, R. N., Coutinho, M. A., Paschoal, G. D., & Chaves, É. D. (2019). Religious/spiritual coping and spiritual distress in people with cancer. Revista Brasileira de Enfermagem, 72(6), 1534–1540. https://doi.org/10.1590/0034-7167-2018-0585
- Smith, J., & Firth, J. (2011). Qualitative data analysis: The framework approach. *Nurse Researcher*, 18(2), 52–62. https://doi.org/10.7748/nr2011.01.18.2.52.c8284

Spencer, G., Fairbrother, H., & Thompson, J. (2020). Privileges of power: Authenticity, representation and the "problem" of children's voices in qualitative health research. *International Journal of Qualitative Methods*, 19, 160940692095859. https://doi.org/10.1177/1609406920958597

- Together for Short Lives (2023). Key information about children who may need palliative care (togetherforshortlives.org.uk) last accessed 27.07.2023.
- Tong, A., Sainsbury, P., & Craig, J. (2007). Consolidated criteria for reporting qualitative research (COREQ): A 32-item checklist for interviews and focus groups. *International Journal for Quality in Health Care*, 19(6), 349–357. https://doi.org/10.1093/intqhc/mzm042
- Veronese, S., Gallo, G., Valle, A., Cugno, C., Chiò, A., Calvo, A., Rivoiro, C., & Oliver, D. J. (2015). The palliative care needs of people severely affected by neurodegenerative disorders: A qualitative study. *Progress in Palliative Care*, 23(6), 331–342. https://doi.org/10.1179/1743291X15Y.0000000007
- Watson, J., Voss, H., & Bloomer, M. J. (2019). Placing the preferences of people with profound intellectual and multiple disabilities at the center of end-of-life decision making through storytelling. *Research and Practice for Persons with Severe Disabilities*, 44(4), 267–279. https://doi.org/10.1177/1540796919879701
- Watts, L., Smith, J., McSherry, W., Tatterton, M., & Rodriguez, A. (2020a). Stakeholder perceptions of dignity therapy for children and young people with life-limiting and life-threatening conditions in the UK. OBM Integrative and Complementary Medicine, 5(1), 1–3. https://doi.org/10.21926/obm.icm.2001012
- Watts, L., Smith, J., McSherry, W., Tatterton, M., Rodriguez, A., et al. (2020b). Correction: Stakeholder perceptions of dignity therapy for children and young people with life-limiting and life-threatening conditions in the UK. *OBM Integrative and Complementary Medicine*, 5(1), 1–3. https://doi.org/10.21926/obm.icm.2001012

Author Biographies

Alison M. Rodriguez is an associate professor in Child and Family Health (Psychology) within the School of Healthcare at The University of Leeds.

Allan Kellehear is a clinical professor within the Departments of Nursing and Medicine at the University of Vermont. Allan is a Medical and Public Health Sociologist.

Vitaveska Lanfranchi is a senior lecturer in Medical Computing and head of the Pervasive Computing Research Group at the University of Sheffield.

Wilfred McSherry is a professor in Nursing within the School of Health, Science and Wellbeing at Staffordshire University. Wilfred is an adult nurse.

Michael Tatterton is a clinical associate professor within the School of Nursing and Healthcare Leadership at the University of Bradford. Michael is a children's nurse.

Lucy Watts MBE was a Youth Ambassador for young people with life-limiting conditions. Lucy unfortunately became increasingly ill through the course of this work and died shortly after the close of our study.

Nahel Yaziji is a research associate based in the Institute of Psychiatry, Psychology and Neuroscience at Kings College, London. Nahel is a pharmacist with an MSc in Public Health. He has undertaken training in economic evaluation, decision modelling and financial mathematics and works on the economic evaluation components of projects.

Joanna Smith is a professor of Nursing in Child Health at Sheffield Hallam University and Sheffield Children's Hospital. Joanna is a child and adult nurse.