

Palliative care for children with complex cardiac conditions: survey results

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ABSTRACT

Objective To explore perspectives of paediatric cardiac and palliative care professionals on providing palliative care to children with complex cardiac conditions.

Design A national survey including closed-ended and open-ended questions as well as clinical scenarios designed to capture referral practices, attitudes towards palliative care, confidence delivering key components of palliative care and perspectives on for whom to provide palliative care. Responses to closed-ended questions and scenarios were analysed using descriptive statistics. Open-ended responses were analysed thematically.

Participants Paediatric cardiac and palliative care professionals caring for children with complex cardiac conditions in the UK.

Results 177 professionals (91 cardiac care and 86 palliative care) responded. Aspects of advance care planning were the most common reasons for referral to palliative care. Palliative care professionals reported greater confidence than cardiac colleagues with such discussions. Clinicians agreed that children with no further surgical management options, comorbid genetic disorders, antenatal diagnosis of a single ventricle, ventricular device in situ, symptomatic heart failure and those awaiting heart transplantation would benefit from palliative care involvement.

Conclusions Components of palliative care, such as advance care planning, can be provided by cardiac care professionals alongside the disease-directed care of children with complex cardiac conditions. Further research and training are needed to address confidence levels in cardiac care professionals in delivering components of palliative care as well as clarification of professional roles and parent preferences in delivery of family-centred care for children with complex cardiac conditions.

INTRODUCTION

Approximately 15%–20% of children with major heart conditions have high-risk complex cardiac conditions.¹ While survival rates for these children have improved,^{2 3} not all have a successful outcome.⁴⁻⁷ Death is more likely to occur in the first year of life, commonly in high-acuity settings following withdrawal of intensive therapies.^{5 8} Parents report significant suffering of their child at their end of life and may only realise their child was dying within their last 24 hours.⁴ Those who survive their first year do so with complex morbidity⁹ and ongoing possibility of early death.^{10 11}

Given these risks, there is an increasing call to provide palliative care alongside disease-directed

What is already known on this topic?

- ▶ Despite improved operative survival rates, children with complex cardiac conditions have a high burden of morbidity and risk of death in childhood.
- ► Palliative care delivered in parallel to diseasedirected care is recognised as beneficial.
- ➤ The specific roles for palliative and cardiac care professionals delivering key components of palliative care to children with complex cardiac conditions are unclear.

What this paper adds?

- Children antenatally diagnosed with single ventricle disease, no surgical options, ventricular device in situ, comorbid disorders, heart failure or awaiting transplantation should have palliative care involvement.
- ► Identification of cardiac care professionals' self-reported lack of confidence in conducting advance care planning discussions with parents.
- Support for development and study of parental views of packages of care integrating principles of palliative care into care for children with complex cardiac conditions.

management. 11–13 By planning disease-directed and palliative care in parallel, clinicians can better address parental concerns, improve parental well-being and broaden the focus of parent–clinician discussions. 14–20 Specifically, palliative care professionals (PCPs) can support complex decision-making, particularly regarding advance care planning, and improve the quality of life for children with complex cardiac conditions and their families throughout the illness, 10 11 13 not just at the end of life.

In the UK, children with complex cardiac conditions receive care in 1 of 16 paediatric cardiac centres, 11 with cardiac surgical facilities, 21 including 2 with cardiothoracic transplant programmes. PCPs, predominantly nurses, largely provide direct care in community and hospice settings. Most medical PCPs are paediatricians or general practitioners with special interests in palliative care. 22

Despite increasing interest in incorporating palliative care into the care of children with complex cardiac conditions, referrals remain low⁴ ²³ and the



specific role of PCPs in this context remains largely unexplored. This study addresses this gap through an examination of current referral practices, attitudes, confidence levels and perspectives of cardiac care professionals (CCPs) and PCPs who provide palliative care to children with complex cardiac conditions.

METHODS

Study design

This national cross-sectional e-survey study was registered at the Great Ormond Street Hospital National Health Service Foundation Trust and was exempt from research ethics committee review. Deidentified data with no institutional-based linkage were collected.

Inclusion criteria

CCPs and PCPs, regardless of discipline, who care for children with complex cardiac conditions in the UK.

Recruitment

An invitation and secure email link were distributed to members of the: Paediatric Intensive Care Society Study Group, British Congenital Cardiac Association, Congenital Cardiac Nurses Association, Association for Paediatric Palliative Medicine and Royal College of Nursing's Children's Palliative Care Special Interest Group. Snowball sampling was encouraged. The survey remained open from 20 June 2018 to 31 August 2018.

The survey

The survey aimed to identify current practices, attitudes, confidence and perspectives of participants on providing palliative care to children with complex cardiac conditions. The survey (online supplemental file 1) was divided into six parts: (1) non-identifiable participant demographics, (2) palliative care referral practices, (3) attitudes towards palliative care and its key components, (4) confidence delivering key components of palliative care, (5) palliative care involvement in clinical scenarios, (6) open-ended questions exploring the appropriate time for referral, barriers to discussing goals of care when a child may have a poor outcome and additional comments regarding palliative care involvement.

Questions and topics included items used in previous surveys, ^{8 24 25} new items, as well as clinical scenarios developed using existing literature ¹⁰ in consultation with PCPs and CCPs. The surveys were hosted on SurveyMonkey. ²⁶

Data analysis

Quantitative analysis

Descriptive analysis was performed using SPSS.²⁷ Data related to current referral practices were discrete response choices that were summarised according to percentage of either PCP or CCP who selected each response option. Data from questions related to attitudes towards palliative care and its key components, confidence delivering the key components and palliative care involvement in clinical scenarios were obtained from a 4-point Likert scale. These data, predominantly presented graphically, were summarised according to the proportion of responses made by PCPs or CCPs. A two-sided Mann-Whitney U test with continuity and tie corrections, with p value set at ≤ 0.05 , was used to test differences between responses of each group—PCPs and CCPs. We then dichotomised these data to analyse level of agreement/disagreement or confidence within each group. When considering the views of participants, we used an a priori level of 75% to indicate overall agreement/disagreement or confidence in each statement or activity. Statistical details

Table 1 Participant demogra	priics		
Characteristic	PCPs (n=86)	CCPs (n=91)	P value
Female	74 (87.1%)	72 (79.1%)	0.368
Clinical role			0.010
Nurse	37 (43.5%)	59 (64.8%)	
Doctor	46 (54.1%)	30 (33%)	
Other	2 (2.4%)	2 (2.2%)	
Specialty			
Hospital-based team	19 (22%)	-	
Community-based team	8 (9.3%)	_	
Children's hospice	25 (29.1%)	_	
>1 listed affiliation	11 (12.8%)	_	
Cardiology	-	37 (40.7%)	
Intensive care	_	46 (50.5%)	
Cardiac surgery	-	2 (2.2%)	
Other	14 (16.3%)	6 (6.6%)	
No response	9 (10.5%)	0	
Seniority*	63 (76.8%)	58 (65.9%)	0.130
Years in current position			0.124
<1	9 (11.5%)	7 (7.9%)	
1–3	19 (24.4%)	17 (19.1%)	
4–6	14 (17.9%)	20 (22.5%)	
7–9	15 (19.2%)	8 (9.0%)	
>10	21 (26.9%)	37 (41.6%)	
Conducts prenatal consultations	37 (47.4%)	11 (12.4%)	< 0.001
Institution has separate cardiac ICU	27 (34.6%)	61 (68.5%)	< 0.001

^{*}Band 7 (senior clinical nurses, nurse managers) or higher or consultant position. CCPs, cardiac care professionals; ICU, intensive care unit; PCPs, palliative care professionals.

of results not presented graphically are in the text. Supporting numerical data are available in online supplemental file 2.

Qualitative analysis

Responses to open-ended questions were analysed thematically with reference to closed-ended questions and clinical scenarios thereby capturing both a priori themes from the survey and literature as well as newly emerging themes. Qualitative data analysis was managed in NVivo.²⁸

RESULTS

Participants

Two-hundred and one professionals commenced the survey. Thirteen did not go beyond the initial screening questions, eight did not answer any survey questions and three did not identify as PCPs or CCPs. Of the 177 included professionals, 86 identified as PCPs and 91 as CCPs. Demographics are described in table 1.

Current palliative care referral practices

The most commonly reported reasons for referral of children to PCPs were for: assistance with preferred place of death discussions, advance care planning, symptom management, death/dying discussions or to aid parental decision-making (table 2).

There was no overall agreement among either CCPs or PCPs on whether referrals of children with complex cardiac conditions to palliative care were untimely.

Attitudes towards palliative care and its key components

There were significant differences between CCPs and PCPs not only in their views and attitudes towards palliative care, but also

Original research

Table 2 Reasons for referral to palliative care			
Cited reason	Reported by cardiac care providers	Reported by palliative care providers	
Preferred place of death discussions	73/74 (98.6%)	51/72 (70.8%)	
Advance care planning discussions	70/74 (94.6%)	42/72 (58.3%)	
Symptom management	67/74 (90.5%)	45/72 (62.5%)	
To discuss death/dying issues with parents	64/74 (86.5%)	37/72 (51.4%)	
Assist with parental decision-making	58/74 (78.4%)	33/72 (45.8%)	
To discuss anticipated outcomes	54/74 (73%)	32/72 (44.4%)	
Interprofessional conflict	47/74 (63.5%)	21/72 (29.2%)	
Do-not-attempt resuscitation discussions	43/74 (58.1%)	32/72 (44.4%)	

in its key components and impact on parental hope (figure 1). Both CCPs and PCPs reported that palliative care extended beyond the end of life phase (statement 1), the last weeks of life (statement 2) and could be instituted even when management had not yet been decided (statement 5).

While CCPs disagreed among themselves on the acceptability (statement 3) and impact of introducing palliative care on parental hope (statement 4), they agreed on its value as a support to clinicians (statement 9), when there was disagreement among professionals (statement 6) and in managing symptoms (statement 8).

Confidence in delivering key components of palliative care

Both CCPs and PCPs reported confidence in: discussing 'goals of care' (statement 1) and concerns over 'death and dying' (statement 3), 'caring for a deteriorating child over hours to days' (statement 4) and 'providing care during the 'end of life period' (statement 8). PCPs were more confident than CCPs in the first three of these activities and also more confident in discussions around preferred place of death (statement 6). There was, however, no difference in confidence between PCPs and CCPs in providing care during the 'end of life period' (statement 8) (figure 2).

While PCPs reported confidence in discussing resuscitation with 16-year-old children, CCPs did not. Neither PCPs nor CCPs expressed confidence discussing resuscitation with 12-year-old children (p<0.001 for both).

Both CCPs and PCPs reported similarly low levels of confidence in prognostication (p=0.27). For both, this was the area of lowest reported confidence.

Palliative care involvement in clinical scenarios

CCPs and PCPs agreed that children with: no further surgical options (scenario 1), an antenatal diagnosis of single ventricle disease (scenario 2), a ventricular assist device in situ (scenario 3), a comorbid genetic condition (scenario 4), symptomatic cardiac failure from myocardial dysfunction (scenario 5) and those awaiting cardiac transplantation (scenario 11) should be referred to palliative care (figure 3). Of these, PCPs were more likely than CCPs to agree that babies diagnosed antenatally with single ventricle disease (scenario 2) and those with a comorbid genetic condition (scenario 4) should be referred to palliative care.

While there was lack of agreement among CCPs about whether a child with a prolonged intensive care unit stay (scenario 6), neonate with single ventricle disease proceeding to surgery (scenario 10) and those with lack of professional consensus over the treatment plan (scenario 12) should be referred to palliative care, there was no significant overall difference in these responses between PCPs and CCPs.

Although neither group agreed on palliative care involvement for a planned ventricular assist device (scenario 9), implantable cardiac defibrillator (scenario 8) or for a child receiving extracorporeal membrane oxygenation (scenario 7), CCPs were more likely to agree with the statement that palliative care should be involved for a child being planned for a ventricular assist device.

PCPs were ranked by both CCPs and PCPs as the choice for assistance after a colleague in the same team, but before a clinical ethics panel in cases of disagreement about resuscitation status, or in cases where withdrawing artificial nutrition/hydration, further cardiac surgery, or tracheostomy insertion and long-term ventilation were being considered.

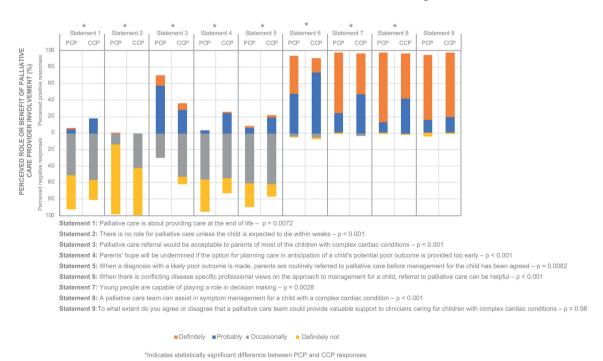


Figure 1 Attitudes towards palliative care. CCP, cardiac care professional; PCP, palliative care professional.

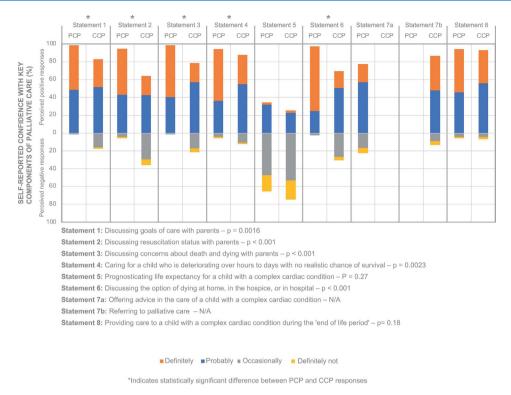


Figure 2 Self-reported confidence in delivering key components of palliative care. CCP, cardiac care professional; N/A, not applicable; PCP, palliative care professional.

Qualitative analysis

One-hundred and thirty-seven participants (69 of 91 (75.8%) CCPs; 68 of 86 (79%) PCPs) responded to one or more of the open-ended questions; elaborating on their responses to closed-ended questions regarding attitudes to palliative care, confidence in delivering palliative care and when to refer to PCPs.

Additional information not directly touched on in closed-ended questions included: (1) foregrounding quality of life as an indicator for referral; (2) nature of uncertainty and (3) management of parental hope.

While the term 'quality of life' did not appear in the survey (as meaning is variously interpreted²⁹), it was raised by both

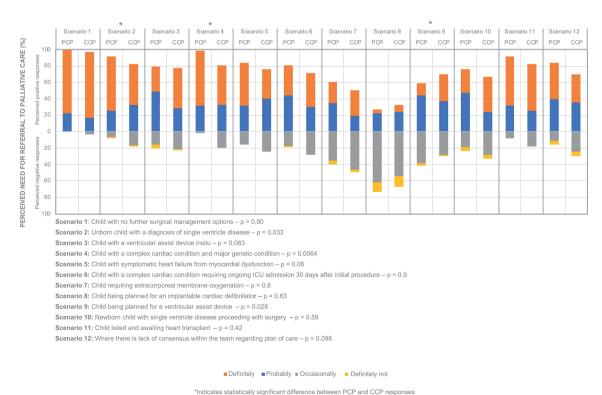


Figure 3 Referral practices related to clinical scenarios. CCP, cardiac care professional; ICU, intensive care unit; PCP, palliative care professional.

Original research

CCPs and PCPs as an indicator for referral. For example, referring a child 'ideally when [the child] becomes unstable/quality of life is in question' (CCP, <1-year experience in current role) or 'at the point of diagnosis ... selling all the positive benefits about focusing on having the best life they can rather than approaching it from the end of life care perspective' (PCP, 7–9 years of experience in current role).

The nature of uncertainty, particularly around surgery, as an indicator for referral to PCPs, was made more explicit in some responses. For example, in CCPs' statements, while there was support for earlier referral, this was defined as when there were 'no further surgical options' (CCP, 4–6 years of experience in current role) or 'at the point where treatment has 'failed'' (CCP, >10 years of experience in current role). PCPs remarked on surgery, but more broadly, including risk and the future as in 'when there is uncertainty about the future or [the child is] facing potentially risky surgery' (PCP, 1–3 years of experience in current role).

Management of hope also figured in CCPs' comments on referral, which should occur 'as early as possible PROVIDING it is managed carefully with the families and that they are helped to see that this is not saying that we are giving up on their child' (CCP, 4–6 years of experience in current role).

DISCUSSION

This study revealed significant agreement between CCPs and PCPs on when children with complex cardiac conditions should be referred to PCPs. There was also strong agreement on the roles of PCPs and their contributions to enhancing care. Given this strong support for referral to PCPs and specific indicators for doing so both in this study and others, ⁴⁸ ^{10–13} ²⁴ ^{30–32} why do we repeatedly see calls for earlier and greater referral to palliative care not only for children with complex cardiac conditions, but also for other life-threatening conditions? ¹⁴ ¹⁹ ²⁴ ³¹ ^{33–35}

We would suggest the solution to this conundrum lies in reframing the question from: How can we achieve earlier and greater referral to palliative care? to: How can we achieve the benefits of palliative care in the face of conscious and unconscious individual and institutional barriers? These barriers often establish a dichotomy between 'fixing' the child versus supporting best quality of life. Previous studies indicate that while parents prefer palliative care principles and practices integrated into their child's treatment, ^{36–39} they do not necessarily want to receive those services from PCPs. They prefer to receive them from their treating teams.

Building on CCPs and PCPs' shared vision of which children would benefit, we would recommend integrated models of care for children with complex cardiac conditions such as that described by Moynihan *et al*⁴⁰ enhanced by PCPs providing an educative and supportive role to CCPs in such aspects of care as advance care planning. We hypothesise that with such an approach, CCPs will gain greater confidence in delivering essential components of palliative care as well as better management in face of deterioration and death. Moreover, parents will be able to have advance care planning conversations with those they want to—the treating team. ^{36 37} Not inconsequentially such an approach may also enhance clinicians' skills in other areas of practice including talking with parents and children about less charged issues, as well as reducing the risk of overwhelming the availability of palliative care resources.

Strengths and weaknesses/limitations of the study

This study was robust, involving senior multidisciplinary professionals matched in seniority in both groups, capturing

perspectives of those with valuable experience. Reflecting the field, the survey also included all potential providers of palliative care. As a national survey, the findings extend beyond practices and perspectives within a single institution.

To reach the greatest number of professionals, participants were recruited through national organisations and snowball sampling was encouraged. However, given this strategy, we cannot report the response rate. We recognise the potential biases within the sample, based on gender, given significant differences in medical and nursing respondents, limited number of cardiac surgical respondents and those from non-medical/nursing backgrounds. We agree that professionals who are interested in this area may have been more likely to participate, and note that those with different attitudes towards palliative care may not be represented in our sample. We acknowledge the implications that this lack of inclusion might have on planning service model innovations, and as such, hope future research will capture the variety of perspectives of individuals within institutions. In addition, we encourage further study of interpretations of quality of life, content of advance care planning discussions and how these might fit within cardiac treatment plans.

Aware of the inherent limitations in exploring perspectives through a survey, we used 4-point Likert scales, removing the neutral option, ensuring a more definitive response. The disparity in overall number of responses to closed-ended questions with multiple options and no requirement to rank responses was addressed through open-ended questions which provided opportunities for elaboration and clarification of closed responses. Finally, the survey was unique in inclusion of clinical scenarios which identified how CCPs applied their perspectives in practice and provided opportunities for PCPs to specify which children they could offer the most help with.

CONCLUSION

This study provides evidence-based guidance for integration and delivery of palliative care to children with no further surgical management options, comorbid genetic disorders, an antenatal diagnosis of a single ventricle, ventricular device in situ, symptomatic heart failure from myocardial dysfunction and those listed and awaiting heart transplantation. Further research and training are needed to address confidence levels in cardiac care professionals in delivering components of palliative care as well as clarification of professional roles and parent preferences in delivery of family-centred care for children with complex cardiac conditions.

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Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions

Introduction

Thank you for taking part in this national survey.

We are seeking your views to better understand and improve services for children with complex cardiac conditions, their parents and professionals caring for them.

We estimate that the survey will take approximately 10 minutes to complete. Your responses cannot be saved so for your views to be shared, you must finish the survey after opening it. The deadline for survey completion is 30 June 2018.

We greatly appreciate your time and input

If you have any questions about the survey or would like more information, please contact Sid Vemuri (s.vemuri@ucl.ac.uk).

By continuing with this survey you acknowledge that:

- Submission of the survey indicates your consent to participate.
- All responses are anonymous and neither you nor your institution will be identifiable.
- Participation is entirely voluntary and you may cease answering questions at any time you wish.

Once you have submitted the survey your data cannot be withdrawn.
* 1. Do you agree to continue with this survey?
Yes
○ No

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions	
Screening for inclusion	
* 2. Does your practice include clinical care for children with complex cardiac conditions?	
Yes No	

	atric palliative care providers' perspectives and practices in the care of children omplex cardiac conditions	
emog	graphics	
3. G	ender	
	Male	
	Female	
	Prefer not to say	
* 4. Pl	lease select your profession/occupation	
	Doctor	
	Nurse	
	Other (please specify)	

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions
Demographics
5. Please select your role
Consultant
Specialty or Associate Grade doctor
Trainee

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions
6. Please select your current band
Band 5
Band 6
Band 7 or above
* 7. Please select your primary professional affiliation
Hospital-based palliative care team
Children's hospice
Community-based palliative care team
Other (please specify)
* 8. How long have you worked in your current role?
Less than 1 year
1-3 years
4-6 years
7-9 years
10 years or more
* 9. Which of the following is your primary field ?
General practice
Community children's nursing
Children's hospice care
Children's community palliative care
General paediatrics
Specialist paediatric palliative care
* 10. Do you participate in any prenatal consultations?
Yes
○ No

	Does your main referring institution have a separate cardiac intensive care unit?
	Yes
	No
	On average, what percentage of children who you have provided end of life care to in a given year had a complex cording condition?
	e had a complex cardiac condition?
	<1%
	1-5%
	5-25%
	25-50%
	>50%
\bigcirc	Unsure

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions
Clinical practice and experience
* 13. When a child may have a poor outcome, do you think there are barriers to discussing goals of
care? Yes
○ No

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions	
Clinical practice and experience	
14. Please describe the barriers	
	8

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions
Clinical practice and experience

	Strongly disagree	Disagree	Agree	Strongly agre
Palliative care is primarily about providing care at the end of life				
There is no role for palliative care involvement unless the child is expected to die within weeks		\circ	\circ	\circ
Palliative care referral would be acceptable to the parents of most of the children with complex cardiac conditions				
Parents' hope will be undermined if the option for planning care and treatment in anticipation of a child's potential poor outcome (including death) is provided too early				
When a diagnosis with a likely poor outcome is made, parents are routinely offered referral to the palliative care team before management for the child has been agreed				
When there are conflicting disease-specific professional views on the approach to management of a child, a referral to the palliative care team can be helpful				
Young people are capable of playing a role in decision-making				
A palliative care team can assist in symptom management for a child with complex cardiac conditions	0	\circ	\circ	0

	Very confident	Confident	Not confident	Not at all conf
Discussing the goals of care with parents				
Discussing resuscitation status with parents	\bigcirc			
Discussing concerns about death and dying with parents				
Caring for a child who is deteriorating over hours-to-days with no realistic chance for survival	\circ	0	0	0
Prognosticating life expectancy for a child with a complex cardiac condition				
Discussing the option of dying at home, in the hospice or hospital	0	0	0	0
Offering advice in the care of a child with a complex cardiac condition				
Providing care to a child with a complex cardiac condition during the 'end-of-life care period' (defined as the time after which you realise that the child has no realistic chance for survival)				0
L7. Please select if you a			-	
Aged 12 years	Goals of care	Resuscitat	tion status Conce	rns about death ar
		L	_	
Aged 16 years				

Paediatric palliative care p with complex cardiac cond		s and practices in the ca	are of children
Palliative care involvemen	t in the care of childre	en with complex cardiac	conditions
* 18. To what extent do you to clinicians caring for child	= =	·	provide valuable support
Strongly disagree	Disagree	Agree	Strongly agree
			0
* 19. In the last year, how of complex cardiac conditions	-	eferrals from clinicians car	ing for children with
Never	Rarely	Occasionally	Regularly
* 20. Why have children with apply) To support parents in decis	ion making		? (Please select all that
To assist in discussions wit	h parents around expected o	utcomes and expectations	
To allow parents to plan ca	re in the event of a potential p	oor outcome for the child	
To provide symptom mana	gement to the child or young p	person	
To discuss DNAR status wi	th parents		
To discuss with the parents	the option of the child or you	ng person dying at home or in a	hospice
To discuss the issues of illn	ess and dying with the paren	ts and/or child or young person	
To assist in finding a resolu	tion of differences between th	ne team and parents wishes for c	are
Other (please specify)			

* 21. In general, do you feel At the appropriate time		referred to palliative care	Too late
At the appropriate time	,		

Paediatric palliative care providers' perspectives and practices in the care of children with complex cardiac conditions
Palliative care involvement
22. When would be the more appropriate time to refer to palliative care?

	ediatric palliative care th complex cardiac cor		spectives and praction	ces in the care of c	hildren
Sc	enarios				
*	23. Do you think the child				
	Child where there are no further surgical management options technically possible	Definitely	Probably	Occasionally	Definitely not
	Unborn child with an antenatal diagnosis of a single ventricle	\circ	\circ	\circ	0
	Child with a ventricular assist device in-situ				
	Child with a complex cardiac condition and a major genetic disorder (excluding trisomy 21)	0	\circ	\circ	0
	Child with symptomatic heart failure from myocardial dysfunction				
	Child with a complex cardiac condition who requires ongoing ICU/HDU admission 30-days after their initial procedure	0			
	Child requiring ECMO				
	Child being planned for an implantable cardiac defibrillator		\circ	\bigcirc	\bigcirc
	Child being planned for a ventricular assist device				
	Newborn child with a single ventricle proceeding with surgery	0	0		0
	Child listed and awaiting heart transplantation				
	Where there is lack of professional consensus within the treating team regarding the plan of care	0			

* 24. In the following clinic support for the treating of		t and rank the most useful :	sources of additional
	Colleague in their team	Clinical ethics panel	Palliative Care team
Scenario: Child with a single ventricle, post first-stage palliative surgery, who has recurrent cardiac arrests Issue: The family want full resuscitation but the treating clinician wants to limit ongoing resuscitation interventions			
Scenario: Extremely premature child, now two months corrected age, with chronic lung disease, who has required multiple dilatations for pulmonary vein stenosis affecting all pulmonary veins			
Issue: The family want to pursue surgical repair but the treating clinician feels this should not be offered			
Scenario: Child who has remained in the intensive care for six months with CPAP dependence, following palliative cardiac surgery Issue: The family want a tracheostomy inserted but the treating clinician does not feel it should be offered			
Scenario: Child with trisomy 21, failure to thrive and corrected congenital heart disease Issue: The family want to remove the nasogastric tube and continue with oral feeding only, but the treating clinician thinks the child would benefit from ongoing nasogastric feeds and/or gastrostomy insertion			

Pae with	diatric palliative care providers' perspectives and practices in the care of children complex cardiac conditions
	5. Any other comments regarding palliative care involvement in children with complex cardiac
C	onditions
Tha	ank you for your participation.

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions

Introduction

Thank you for taking part in this national survey.

We are seeking your views to better understand and improve services for children with complex cardiac conditions, their parents and professionals caring for them.

We estimate that the survey will take approximately 10 minutes to complete. Your responses cannot be saved so for your views to be shared, you must finish the survey after opening it. The deadline for survey completion is 31 August 2018.

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- All responses are anonymous and neither you nor your institution will be identifiable.
- Participation is entirely voluntary and you may cease answering questions at any time you wish.

• However, once you have submitted the survey your data cannot be withdrawn.
* 1. Do you agree to continue with this survey?
Yes
○ No

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Screening for inclusion
* 2. Dono your practice include clinical care for shildren with complex conditions?
* 2. Does your practice include clinical care for children with complex cardiac conditions ? Yes
○ No

emon	graphics				
Jillog	партнос				
	ender				
	Male				
	Female				
	Prefer not to say				
4. Pl	ease select you	r profession	occupation/		
	Doctor				
	Nurse				
	Other (please speci	fy)			

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Demographics
5. Please select your current role
Consultant
Specialty or Associate Grade doctor
Trainee

Vemuri S, et al. Arch Dis Child 2021;0:1–6. doi: 10.1136/archdischild-2020-320866

6. Pleas	e select your cur	rent band		
Band	d 5			
Band	d 6			
Band	d 7 or above			

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Demographics
* 7. Specialty
Cardiology
Intensive care
Cardiac surgery
General paediatrics
6

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions

Demographics	
* 8. Which of the following is your primary field?	
General cardiology	Heart failure/transplant
Fetal cardiology	Electrophysiology
Single ventricle service	Cardiac imaging
Interventional cardiology	
* 9. How long have you worked in your current role?	
Less than 1 year	7-9 years
1-3 years	10 years or more
4-6 years	
* 10. Do you conduct any prenatal consultations?	
Yes	
○ No	
	industrial and the second of t
* 11. Does your institution have a separate cardiac	intensive care unit?
Yes No	
) NO	
* 12. Does your institution have access to a palliativ	e care team?
Yes	
○ No	
Unsure	
* 13. Have you had any professional experience wit	h a (please select all that apply)
Hospital-based palliative care team	
Children's hospice	
Community-based palliative care team	

* 14. On average, what percer	tage of children under your care have died in a given year?	
<1%	25-50%	
1-5%	>50%	
5-25%	Unsure	
		8

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Clinical practice and experience
* 15. When faced with a child who may have a poor outcome, do you think there arebarriers to discussing goals of care?
Yes
○ No

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Palliative care involvement
16. Please describe the barriers

	Strongly disagree	Disagree	Agree	Strongly agree
Palliative care is primarily about providing care at the end of life				
There is no role for palliative care involvement unless the child is expected to die within weeks		0	0	0
Palliative care referral would be acceptable to the parents of most of the children I look after				
Parents' hope will be undermined if the option for planning care and treatment in anticipation of a child's potential poor outcome (including death) is provided too early				
When a diagnosis with a likely poor outcome is made, parents are routinely offered referral to the palliative care team before management for the child has been agreed				
When there are conflicting professional views on the approach to management, a referral to the palliative care team can be helpful			0	
Young people are capable of playing a role in decision-making				
A palliative care team can assist in symptom management in the children I look after	0	0	0	0

Discussing goals of care with parents Discussing resuscitation status with parents Discussing concerns about death and dying with parents Caring for a child who is deteriorating over hours-to-days with no realistic chance for surrivial Prognosticating life expectancy Discussing the option of dying at home, in the hospited or hospited I hospited		Very confident	Confident	Not confident	Not at all confic
resuscitation status with parents Discussing concerns about death and dying with parents Caring for a child who is deteriorating over hours-to-days with no realistic chance for survival Prognosticating life expectancy Discussing the option of dying at home, in the hospice or hospital Referral to a palliative care team Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years In a young person (please check all that apply)					
about death and dying with parents Caring for a child who is deteriorating over hours-to-days with no realistic chance for survival Prognosticating life expectancy Discussing the option of dying at home, in the hospice or hospital Referral to a palliative care team Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years	resuscitation status	\bigcirc		\circ	\bigcirc
deteriorating over hours-to-days with no realistic chance for survival Prognosticating life expectancy Discussing the option of dying at home, in the hospice or hospital Referral to a palliative care team Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realiste that the child has no realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) and the dead of the confidence of the confi	about death and dying				
Discussing the option of dying at home, in the hospice or hospital Referral to a palliative care team Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realise that the child has no realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years	deteriorating over hours-to-days with no realistic chance for	\circ	\circ	\circ	\circ
dying at home, in the hospice or hospital Referral to a palliative care team Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realise that the child has no realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years					
Providing care to a child with heart disease during the 'end-of-life care period' (defined as the time after which you realise that the child has no realistic chance for survival) L9. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years	dying at home, in the	0	\circ	0	\circ
with heart disease during the 'end-of-life care period' (defined as the time after which you realise that the child has no realistic chance for survival) 19. Please select if you are confident discussing with a young person (please check all that apply) Goals of care Resuscitation status Concerns about death and Aged 12 years					
Aged 12 years Goals of care Resuscitation status Concerns about death and	with heart disease during the 'end-of-life	\bigcirc	\circ	\circ	
Aged 12 years	realise that the child has no realistic chance				
	realise that the child has no realistic chance for survival)	are confident discus	sing with a youn g	ı person (please ched	ck all that apply)
Aged 16 years	realise that the child has no realistic chance for survival)				
	realise that the child has no realistic chance for survival) 19. Please select if you a				
	realise that the child has no realistic chance for survival) 19. Please select if you a Aged 12 years				
	realise that the child has no realistic chance for survival) 19. Please select if you a Aged 12 years				

	ic cardiology care provid ent in the care of childre			palliative care		
Palliative	Palliative care involvement					
* 20. To your te	what extent do you agree o	or disagree that a pa	alliative care team can pro	ovide valuable support to		
:	Strongly disagree	Disagree	Agree	Strongly agree		
palliati Ne	the last year, how often hav ve care team? ver rely	e you made referra	ls or encouraged others to	o make a referral to the		
O 0	casionally					
Re	gularly					
To ou To ou To pe Ott	nen would you consider mal support parents in decision makir assist in discussions with parents toomes and expectations allow parents to plan care in the etcome for the child provide symptom management to rson her (please specify)	around expected event of a potential poo	To discuss DNAR status w To discuss with the parent: person dying at home or ir To discuss the issues of illi and/or child or young person To assist in finding a resoluteam and parents wishes for	ith parents s the option of the child or young a hospice ness and dying with the parents on ution of differences between the or care		
At t	the appropriate time	Too early	Too late	Not applicable, as no children I look after are referred to palliative care		

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Palliative care involvement
24. When would be the more appropriate time to refer to palliative care?

Paediatric cardiology care providers' perspectives and practices towards palliative care involvement in the care of children with complex cardiac conditions
Scenarios

25. Do you think the children in the following clinical scenarios should be referred to palliative care				
	Definitely	Probably	Occasionally	Definitely not
Child where there are no further surgical management options technically possible				
Unborn child with an antenatal diagnosis of a single-ventricle				\bigcirc
Child with a ventricular assist device in-situ				
Child with a complex cardiac condition and a major genetic disorder (excluding trisomy 21)	\circ		\circ	0
Child with symptomatic heart failure from myocardial dysfunction				
Child with a complex cardiac condition who requires ongoing ICU/HDU admission 30-days after their initial procedure				
Child requiring ECMO				
Child being planned for an implantable cardiac defibrillator	\circ	\bigcirc	\bigcirc	\circ
Child being planned for a ventricular assist device				
Newborn child with a single-ventricle proceeding with surgery	\bigcirc	\bigcirc	\circ	0
Child listed and awaiting heart transplantation				
Where there is lack of professional consensus within the team regarding the plan for care				0
ioi care				

*	26. In the following clinica	ıl scenarios, please selec	ct and rank who you would g	go to for assistance
		Colleague in my unit	Clinical ethics panel	Palliative care team
	Scenario: Child with a single ventricle, post first-stage palliative surgery, who has recurrent cardiac arrests Issue: The family want full resuscitation but you want to limit ongoing resuscitation interventions			
	premature child, now two months corrected age, with chronic lung disease, who has required multiple dilatations for pulmonary vein stenosis affecting all pulmonary veins Issue: The family want to pursue surgical repair but you feel this should not be offered			
	Scenario: Child who has remained in the intensive care unit for six months with CPAP dependence following palliative cardiac surgery Issue: The family want a tracheostomy inserted but you do not feel it should be offered			
	Scenario: Child with trisomy 21, failure to thrive and repaired congenital heart disease Issue: The family want to remove the nasogastric tube and continue with oral feeding only, but you think the child would benefit from ongoing nasogastric feeds or gastrostomy insertion			

aediatric cardi volvement in t	ology care prov the care of child	viders' perspe dren with com	ctives and p plex cardiac	ractices towa conditions	rds palliative	care
	comments regard	ding palliative c	are involveme	nt in children w	ith complex ca	ırdiac
conditions						
hank vou	for your p	articipatic	n.			
nam you	.o. you. p	a. c.o.pac.e				

Palliative care for children with complex cardiac conditions: survey results

Raw data

Q13. Do you think there are barriers to discussing goals of care?

	Yes	No
Palliative Care Provider (PCP)	58/78 (74.4%)	20/78 (25.6%)
Cardiac Care Provider (CCP)	73/89 (82%)	16/89 (18%)

Q15. To what extent do you agree or disagree with the following?

a) Palliative care is primarily about providing care at the end of life

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	30/72 (41.7%)	37/72 (51.4%)	4/72 (5.6%)	1/72 (1.4%)
ССР	18/75 (24%)	43/75 (57.3%)	14/75 (18.7%)	0

b) There is no role for palliative care involvement unless the child is expected to die within weeks

	Strongly	Disagree	Agree	Strongly agree
	disagree			
PCP	61/72 (84.7%)	10/72 (13.9%)	0	1/72 (1.4%)
ССР	43/75 (57.3%)	32/75 (42.7%)	0	0

c) Palliative care referral would be acceptable to the parents of most of the children with complex cardiac conditions.

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	0	21/72 (29.2%)	42/72 (58.3%)	9/72 (12.5%)
ССР	7/75 (9.3%)	40/75 (53.3%)	22/75 (29.3%)	6/75 (8%)

d) Parents' hope will be undermined if the option for planning care and treatment in anticipation of a child's potential poor outcome is provided too early

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	28/72 (38.9%)	41/72 (56.9%)	3/72 (4.2%)	0
ССР	14/75 (18.7%)	41/75 (54.7%)	19/75 (25.3%)	1/75 (1.3%)

e) When a diagnosis with a likely poor outcome is made, parents are routinely referred to palliative care before management for the child has been agreed

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	21/72 (29.2%)	44/72 (61.1%)	5/72 (6.9%)	2/72 (2.8%)
ССР	11/75 (14.5%)	47/75 (62.7%)	15/75 (20%)	2/75 (2.7%)

f) When there are conflicting disease specific professional views on the approach to management for a child, a referral to palliative care can be helpful

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	1/72 (1.4%)	3/72 (4.2%)	35/72 (48.6%)	33/72 (45.8%)
ССР	2/75 (2.7%)	4/75 (5.3%)	56/75 (74.7%)	13/75 (17.3%)

g) Young people are capable of playing a role in decision making

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	1/72 (1.4%)	0	18/72 (25%)	53/72 (73.6%)
ССР	0	2/75 (2.7%)	36/75 (48%)	37/75 (49.3%)

h) A palliative care team can assist in symptom management for a child with a complex cardiac condition.

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	1/72 (1.4%)	0	10/72 (13.9%)	61/72 (84.7%)
ССР	1/75 (1.3%)	1/75 (1.3%)	32/75 (42.7%)	41/75 (54.7%)

Q16. In general, how confident do you feel with the following clinical activities?

a) Discussing GoC with parents

	Very confident	Confident	Not confident	Not at all confident
PCP	36/72 (50%)	35/72 (48.6%)	1/72 (1.4%)	0
ССР	23/75 (30.7%)	39/75 (52%)	12/75 (16%)	1/75 (1.5%)

b) Discussing resuscitation status with parents

	Very confident	Confident	Not confident	Not at all confident
PCP	37/72 (51.4%)	31/72 (43.1%)	3/72 (4.2%)	1/72 (1.4%)
ССР	16/75 (21.3%)	32/75 (42.7%)	22/75 (29.3%)	5/75 (6.7%)

c) Discussing concerns about death and dying with parents

	Very confident	Confident	Not confident	Not at all confident
PCP	42/72 (58.3%)	29/72 (40.3%)	1/72 (1.4%)	0
ССР	16/75 (21.3%)	43/75 (57.3%)	13/75 (17.3%)	3/75 (4%)

d) Caring for a child who is deteriorating over hours to days with no realistic chance of survival

	Very confident	Confident	Not confident	Not at all
				confident
PCP	42/72 (58.3%)	26/72 (36.1%)	3/72 (4.2%)	1/72 (1.4%)
ССР	25/75 (33.3%)	41/75 (54.7%)	8/75 (10.7%)	1/75 (1.3%)

e) Prognosticating life expectancy for a child with a complex cardiac condition

regressive and the expectation for a crima with a complex caraliae container				
	Very confident	Confident	Not confident	Not at all confident
PCP	2/72 (2.8%)	23/72 (31.9%)	34/72 (47.2%)	13/72 (18.1%)
ССР	2/75 (2.7%)	17/75 (22.7%)	40/75 (53.3%)	16/75 (21.3%)

f) Discussing the option of dying at home, in the hospice, or at hospital

	Very confident	Confident	Not confident	Not at all confident
PCP	52/72 (72.2%)	18/72 (25%)	2/72 (2.8%)	0
ССР	14/75 (18.7%)	38/75 (50.7%)	20/75 (26.7%)	3/75 (4%)

g) Offering advice in the care of a child with a complex cardiac condition

	Very confident	Confident	Not confident	Not at all confident
PCP	15/72 (20.8%)	41/72 (56.9%)	12/72 (16.7%)	4/72 (5.6%)
ССР				

Referring to PCT

	Very confident	Confident	Not confident	Not at all confident
PCP				
ССР	29/75 (38.7%)	36/75 (48%)	7/75 (9.3%)	3/75 (4%)

h) Providing care to a child with a complex cardiac condition during the 'end of life care period'

Very confident	Confident	Not confident	Not at all
			confident

PCP	35/72 (48.6%)	33/72 (45.8%)	3/72 (4.2%)	1/72 (1.4%)
ССР	28/75 (37.3%)	42/75 (56%)	3/75 (4%)	2/75 (2.7%)

Q17. Please select if you are confident discussing the following with a young person:

a) Aged 12 years

GOC

	Yes	No
PCP	64/69	5/69
ССР	52/57	5/57

Resus

	Yes	No
PCP	42/68	27/69
ССР	18/57	39/57

Death and dying

	Yes	No
PCP	57/69	12/69
ССР	33/57	24/57

b) Aged 16 years

GOC

	Yes	No
PCP	68/71	3/71
ССР	61/63	2/63

Resus

	Yes	No	
PCP	57/71	14/71	
ССР	36/63	27/63	

Death and dying

	Yes	No
PCP	63/71	8/71
ССР	45/63	18/63

Q18. To what extent do you agree or disagree that a palliative care team could provide valuable support to clinicians caring for children with complex cardiac conditions.

Page **4** of **9**

	Strongly disagree	Disagree	Agree	Strongly agree
PCP	3/72 (4.2%)	0	12/72 (16.7%)	57/72 (79.2%)
ССР	1/74 (1.4%)	0	15/74 (20.3%)	58/74 (78.4%)

Q19. In the last year, how often have you received referrals from clinicians for children with complex cardiac conditions (PC) or how often have you made referrals to the PC team (cardiologists)?

	Never	Rarely	Occasionally	Regularly
PCP	5/72 (6.9%)	11/72 (15.3%)	39/72 (54.2%)	17/72 (23.6%)
ССР	9/74 (12.2%)	20/74 (27%)	27/74 (36.5%)	18/75 (24.3%)

Q20. Why have children with complex cardiac conditions been referred to you?

PCP		ССР		
PPoD Discussions	51/72 (70.8%)	PPoD Discussions	73/74 (98.6%)	
Symptom management	45/72 (62.5%)	Parental care planning	70/74 (94.6%)	
Parental care planning	42/72 (58.3%)	Symptom management	67/74 (90.5%)	
Discuss death/dying issues	37/72 (51.4%)	Discuss death/dying issues	64/74 (86.5%)	
Parental decision making	33/72 (45.8%)	Parental decision making	58/74 (78.4%)	
Outcome discussions	32/72 (44.4%)	Outcome discussions	54/74 (73%)	
DNAR discussions	32/72 (44.4%)	Interprofessional conflict	47/74 (63.5%)	

Q21. In general, do you feel that children were referred to palliative care

	At the appropriate time	Too early	Too late	N/A
PCP	30/72 (41.7%)	0	42/72 (58.3%)	0
ССР	21/73 (28.8%)	0	46/73 (63%)	6/73 (8.2%)

Q23. Do you think children in the following clinical scenarios should be referred to palliative care?

a) Child where there are no further surgical management options possible

	Definitely	Probably	Occasionally	Definitely not
PCP	49/63 (77.8%)	14/63 (22.2%)	0	0
ССР	53/66 (80.3%)	11/66 (16.7%)	2/66 (3%)	0

b) Unborn child with an antenatal diagnosis of a single ventricle

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		Definitely	Probably	Occasionally	Definitely not		

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PCP	42/63 (66.7%)	16/63 (25.4%)	4/63 (6.3%)	1/63 (1.6%)
ССР	33/67 (49.3%)	22/67 (32.8%)	11/67 (16.4%)	1/67 (1.5%)

c) Child with a VAD insitu

	Definitely	Probably	Occasionally	Definitely not
PCP	19/63 (30.2%)	31/63 (49.2%)	10/63 (15.9%)	3/63 (4.8%)
ССР	33/67 (49.3%)	19/67 (28.4%)	14/67 (20.9%)	1/67 (1.5%)

d) Child with a complex cardiac disorder and a major genetic condition

	Definitely	Probably	Occasionally	Definitely not
PCP	42/63 (66.7%)	20/63 (31.7%)	1/63 (1.6%)	0
ССР	32/67 (47.8%)	22/67 (32.8%)	13/67 (19.4%)	0

e) Child with symptomatic heart failure from myocardial dysfunction

	Definitely	Probably	Occasionally	Definitely not
PCP	33/63 (52.4%)	20/63 (31.7%)	10/63 (15.9%)	0
ССР	24/67 (35.8%)	27/67 (40.3%)	16/67 (23.9%)	0

f) Child with a complex cardiac condition who requires ongoing ICU admission 30 days after their initial procedure

	Definitely	Probably	Occasionally	Definitely not
PCP	23/63 (36.5%)	28/63 (44.4%)	11/63 (17.5%)	1/63 (1.6%)
ССР	28/67 (41.8%)	20/67 (29.9%)	19/67 (28.4%)	0

g) Child requiring ECMO

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	Definitely	Probably	Occasionally	Definitely not		
PCP	16/63 (25.4%)	22/63 (34.9%)	22/63 (34.9%)	3/63 (4.8%)		
ССР	21/67 (31.3%)	13/67 (19.4%)	31/67 (46.3%)	2/67 (3%)		

h) Child being planned for ICD

	Definitely	Probably	Occasionally	Definitely not
PCP	3/63 (4.8%)	14/63 (22.2%)	39/63 (61.9%)	7/63 (11.1%)
ССР	6/67 (9%)	16/67 (23.9%)	36/67 (53.7%)	9/67 (13.4%)

i) Child being planned for VAD

	Definitely	Probably	Occasionally	Definitely not
PCP	9/63 (14.3%)	28/24 (44.4%)	24/63 (38.1%)	2/63 (3.2%)
ССР	22/67 (32.8%)	25/67 (37.3%)	19/37 (28.4%)	1/67 (1.5%)

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j) Newborn child with a single ventricle preceding with surgery

	Definitely	Probably	Occasionally	Definitely not
PCP	18/63 (28.6%)	30/63 (47.6%)	12/63 (19%)	3/63 (4.8%)
ССР	29/67 (43.3%)	16/67 (23.9%)	19/67 (28.4%)	3/67 (4.5%)

k) Child listed and awaiting heart transplant

	Definitely	Probably	Occasionally	Definitely not
PCP	38/63 (60.3%)	20/63 (31.7%)	5/63 (7.9%)	0
ССР	38/67 (56.7%)	17/67 (25.4%)	12/67 (17.9%)	0

I) Where there is a lack of professional consensus within the treating team regarding the plan of care

	Definitely	Probably	Occasionally	Definitely not
PCP	28/63 (44.4%)	25/63 (39.7%)	7/63 (11.1%)	3/63 (4.8%)
ССР	23/67 (34.3%)	24/67 (35.8%)	16/67 (23.9%)	4/67 (6%)

Q24. In the following clinical scenarios, please select and rank the most useful sources of additional support for the treating clinician

a) Resuscitation

COLLEAGUE

	1st	2nd	3rd	N/A
PCP	34/63 (54%)	17/63 (27%)	11/63 (17.5%)	1/63 (1.6%)
ССР	62/67 (92.5%)	1/67 (1.5%)	2/67 (3%)	2/67 (3%)

CLINICAL ETHICS PANEL

	1st	2nd	3rd	N/A
PCP	10/63 (15.9%)	14/63 (22.2%)	37/63 (58.7%)	2/63 (3.2%)
ССР	2/67 (3%)	10/67 (14.9%)	47/67 (70.1%)	8/67 (11.9%)

PCT

	1st	2nd	3rd	N/A
PCP	22/63 (34.9%)	31/63 (49.2%)	10/63 (15.9%)	0
ССР	5/67 (7.5%)	51/67 (76.1%)	4/67 (6%)	7/67 (10.4%)

b) Surgery

COLLEAGUE

	1st	2nd	3rd	N/A
PCP	44/63 (69.8%)	7/63 (11.1%)	12/63 (19%)	0
ССР	61/67 (91%)	2/67 (3%)	2/67 (3%)	2/67 (3%)

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CLINICAL ETHICS PANEL

	1st	2nd	3rd	N/A
PCP	9/63 (14.3%)	12/63 (19%)	40/63 (63.5%)	2/63 (3.2%)
ССР	3/67 (4.5%)	12/67 (17.9%)	44/67 (65.7%)	8/67 (11.9%)

PCT

	1st	2nd	3rd	N/A
PCP	14/63 (22.2%)	40/63 (63.5%)	8/63 (12.7%)	1/63 (1.6%)
ССР	7/67 (10.4%)	46/67 (68.7%)	9/67 (13.4%)	5/67 (7.5%)

c) Tracheostomy

COLLEAGUE

	1st	2nd	3rd	N/A
PCP	36/63 (57.1%)	11/63 (17.5%)	16/63 (25.4%)	0
ССР	55/67 (82.1%)	9/67 (13.4%)	1/67 (1.5%)	2/67 (3%)

CLINICAL ETHICS PANEL

	1st	2nd	3rd	N/A
PCP	10/63 (15.9%)	19/63 (30.2%)	28/63 (44.4%)	6/63 (9.5%)
ССР	6/67 (9%)	8/67 (11.9%)	45/67 (67.2%)	8/67 (11.9%)

PCT

	1st	2nd	3rd	N/A
PCP	19/63 (30.2%)	33/63 (52.4%)	10/63 (15.9%)	1/63 (1.6%)
ССР	9/67 (13.4%)	43/67 (64.2%)	7/67 (10.4%)	8/37 (11.9%)

d) Nutrition/Hydration challenge

COLLEAGUE

	1st	2nd	3rd	N/A
PCP	36/63 (57.1%)	16/63 (25.4%)	10/63 (15.9%)	1/63 (1.6%)
ССР	61/67 (91%)	3/67 (4.5%)	0	3/67 (4.5%)

CLINICAL ETHICS PANEL

	1st	2nd	3rd	N/A
PCP	12/63 (19%)	22/63 (34.9%)	25/63 (39.7%)	4/63 (6.3%)
ССР	3/37 (4.5%)	18/67 (26.9%)	25/67 (37.3%)	21/67 (31.3%)

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PCT

	1st	2nd	3rd	N/A
PCP	18/63 (28.6%)	23/63 (36.5%)	16/63 (25.4%)	6/63 (9.5%)
ССР	5/67 (7.5%)	28/67 (41.8%)	11/67 (16.4%)	23/67 (34.3%)