

REVIEW

When is the right time? Complex issues around withdrawing life-sustaining treatment in children

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When should one withdraw treatment in children? The challenge is to recognise when a decision needs to be made. Parents may be in denial, and deciding which questions to ask may be difficult. Ethically, the guiding principle should be the child's best interests. May the parents or primary caregiver decide what the child's best interests are? Legislation in South Africa prevents a parent or caregiver from refusing treatment that medical professionals deem to be in the child's best interests. This article discusses the ethical and legal aspects around the decision to palliate in children.

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Healthcare practitioners are often faced with difficult decisions, but the real challenge is to recognise when a decision needs to be made. In a busy general practice, managing a chronically ill child you are not attending to daily, the most straightforward approach may be to continue with a treatment plan for the child and family without reflecting on the course of events over time. The parents may be in denial about the condition of the child and it may be particularly difficult to step back and identify which questions need to be asked.

Case study

Let me introduce Sara and Jade. Sara is a 10-year-old girl with quadripastic cerebral palsy. Her mother, Jade, has been bringing her to you since they moved to the area many years ago. Sara is profoundly impaired – physically and mentally. Both her sisters are much older and have left home already. Sara is the centre of Jade's life. Sara's father and Jade are divorced. He could not cope with Jade's 'obsession' with Sara.

Sara does not make eye contact and is unable to communicate. Although she barely moves and has severe spasticity, she has generally been well – until recently. During the past 9 months Sara has refused to eat, vomited frequently, and has been admitted to the local hospital five times for pneumonia. She does not sleep well and cries much of the time. She has undergone numerous invasive investigations,

but her condition is not improving. Gastrostomy feeding has been discussed, but the surgical team did not feel it was necessary as neither parent was in favour of Sara undergoing surgery.

Jade is not coping and feels that Sara is misbehaving. Sara's father doesn't see her often as he travels for work and finds it easier not to interfere with Jade's management of Sara.

Your heart sinks as you see they have another appointment. When Jade and Sara enter, you immediately realise that Sara has pneumonia. She is short of breath and moaning. Her muscle spasms seem worse. Jade is exhausted. Sara has lost more weight and refuses her normal feeds.

You call their paediatrician and again Sara is admitted to the ward, with referrals to the physiotherapist and dietician. As Jade leaves your rooms she asks, 'What are we going to do next?'

Jade's question is very important. In such a case it is very easy to practise reactive medicine, responding to each acute episode individually without asking difficult questions. Perhaps we need to ask, 'What *should* we do next?'

How do we approach a case such as this one?

There are three key considerations when regarding difficult ethical decisions in caring for children:

- *What* is the ethical issue in this case?
- *Who* should make this decision?
- *How* is the decision made?

What is the ethical issue in this case?

Before you can make decisions, you must define the questions. Sara is a chronically ill, neurologically impaired child whose clinical condition is deteriorating. She is uncomfortable and does not tolerate food. The burden of caring for her is enormous and taking its toll on her mother.

We need to consider whether managing Sara in this reactive manner is in her best interests. What is the benefit versus harm of recurrent hospitalisation? Do we have alternative options with regard to her management? What should we be doing about feeding her?

Who should make this decision?

Very often, healthcare decisions are left to the parents or primary caregiver as it is assumed that they will understand the child's best interests. Parents and healthcare workers should be working as a team, ensuring open and honest communication at all times. The healthcare professional must identify the key decision makers in a family. However, it is important to note that legislation in South Africa prevents a parent or caregiver refusing treatment that medical professionals deem to be in the child's best interests.^[1] Dealing with children is complex – one must be able to manage the parents and the child.

What about the child?

In our case study, the child lacks the capacity to be autonomous as she is severely mentally impaired. There is, however, increasing evidence that children should be involved in their own healthcare decisions.^[2-5] The Children's Act 38 of 2005 (amended by Act 41 of 2007), promulgated in 2010, emphasises child participation. 'Every child that is of such an age, maturity and stage of development as to be able to participate in any matter concerning that child has the right to participate in an appropriate way and views expressed by the child must be given due consideration.'^[6]

The act also states, 'Patients, including children, have a right to have sufficient information about their health to enable them to make an informed decision about treatment. The information must be "relevant and must be in a format accessible to children, giving due consideration to the needs of disabled children"^[6] This is echoed in the Health Professions Council of South Africa's *Guidelines for the Withholding and Withdrawing of Treatment*.^[7]

Deciding whether a child has the capacity to make decisions may be difficult, but should take into account the child's ability to reason, their understanding (which may be based on previous experience), their willingness to consent and the seriousness of the situation.^[4] There are no hard-and-fast rules, as children vary greatly depending on age, rate of development, social circumstances and educational level. Each situation needs to be assessed individually. In South Africa, children are very seldom involved in their healthcare decisions and a shift in attitude needs to occur if we are to comply with the Children's Act.^[6]

Many healthcare professionals have been involved in Sara's care over a long period of time. It is imperative that everyone be included in the discussions and decision making to prevent conflicts arising in the team. Both her parents, and possibly even the adult siblings who are also affected by the decisions, should be included in the discussion.

How do decisions get made?

Key ethical principles

- Autonomy
- Beneficence
- Non-maleficence
- Justice

When considering children, the standard of 'best interests' applies. Best interests usually refer to the 'highest net benefit among the available options that apply to any situation in which a decision has to be made regarding the health of the child'.^[8]

The Children's Act^[6] states that 'The best interests of a child are of paramount importance in every matter concerning the child'. This has to take into account every aspect of the child's life, including age, maturity, family dynamics, illness, educational needs and disability.

It is not always easy to know what is in the child's best interests, but Kopelman^[9] argues that taking a child's best interests into account does not require what is ideal, but what is reasonable. In deciding what is in the child's best interests, most healthcare workers try to calculate the consequences of the various treatment options available and choose the one likely to do the least harm (non-maleficence) and the most good (beneficence) to the child.

Although the best interests of the child must carry the most weight, 'any decision made in respect of the child must carefully consider the interests of all potentially affected persons, most usually other family members, old or young, who will live with the child or are dependent upon the immediate family in other ways'.^[10]

To apply the standard of best interests we need the following:

- a good understanding of the ethical principles
- all the available information relevant to the case
- an understanding of the child's quality of life
- a clear idea of treatment goals
- the ability to examine our own motives.

Quality of life

Quality of life is difficult to define but has been described as an individual's satisfaction or happiness with life in domains he or she considers important.^[11] It is a subjective measure, but in the case of the disabled child it is important to consider the primary caregiver's opinion on the quality of the child's life.

The Royal College of Paediatrics and Child Health (RCPCH), UK, has defined the following five situations associated with poor

