With the widespread use of highly active antiretroviral therapy (HAART), the epidemic of paediatric HIV has evolved into a chronic disease of childhood. The difference between this disease and most other chronic diseases of childhood, however, is that it is not only a life-threatening diagnosis but also an extremely stigmatising one, resulting in highly emotionally charged responses to disclosure of such a diagnosis to the patient. The disclosure process is made much more difficult when the person being disclosed to is a child. One of the major difficulties is that one is dealing with many layers of disclosure: disclosure of HIV status to the child; the concomitant disclosure of HIV status of the parent/s and other siblings or other family members; and having to anticipate the child’s own disclosure to his/her friends, extended family and community. Approaches to disclosing the diagnosis may vary between parents and between healthcare providers, but one thing that has been established in many studies worldwide is that the child needs to be informed, many suggesting sooner rather than later during the course of the illness. It has been suggested that disclosure of the diagnosis to the child is an integral part of providing comprehensive medical care to a child infected by HIV and may impact positively on adherence to medication. During adolescence, disclosure of the diagnosis may also aid in preventing high-risk behaviour, thereby curbing spread of the disease. Disclosure of HIV to a child should be seen as an ongoing process that may last several years depending on the cognitive development of the child. Health care providers who deal with HIV infection in children need to develop a plan that will enable them to support parents in disclosure.

FACTORS INFLUENCING THE DISCLOSURE OF HIV STATUS BY PARENTS TO THEIR CHILDREN

- Parents feel responsible for shielding their children from potentially harmful knowledge.
- Discrimination against infected persons by peers, at schools, in day-care centres and in the community as a whole.
- Parents who are coping with their children’s illness must at the same time cope with their own illness, declining health and imminent death.
- Parental feelings of guilt, blame and responsibility for their children’s HIV infection.
- Fear that the diagnosis will negatively affect the child’s will to live.
- Fear of anger from the child related to perinatal transmission.
- Fear of inadvertent disclosure by the child to the broader community.

WHEN TO WITHHOLD/DELAY DISCLOSURE

- Very young child (usually under the age of 5 years) or a child with developmental delay (as defined by IQ < 70).
- Children with severe emotional disturbances.

These children should have their situations assessed periodically should their circumstances change.

FACTORS ASSOCIATED WITH EARLY DISCLOSURE IN HIV-INFECTED CHILDREN

Lester et al. reported their findings in a cross-sectional study of 51 HIV-infected children. They concluded that the following factors play an important role in determining whether disclosure to a child takes place earlier rather than later:
A child with a higher IQ. Higher IQs relate to the child's developmental readiness to understand the complex and changing nature of HIV diagnosis and prognosis.

Urban family residence. Although families from both rural and urban settings expressed fear about negative reactions from the community, it was found that the more negative attitudes were prevalent in communities that lacked HIV/AIDS education and services. Disclosure to children in these communities was therefore found to be delayed.

Families with higher expressiveness. Assessment of family communication style is important in determining whether a child should be disclosed to earlier rather than later.

They also found that at the time of study initiation, subjects who had already been disclosed to had:

- Higher mean ages (10.5 years).
- Increased major life events within the family in the preceding year.

**PROCESS OF DISCLOSURE**

Disclosure is a process that should take place over a period of time. It requires ongoing and effective communication within families. It requires that health professionals respect family needs, wishes and expectations as well as community norms and pressures.

Disclosure must take into account:

- The needs of the child AND the needs of the parent/family. Pushing the process before the family is ready may disrupt the therapeutic alliance and have a negative impact on the ongoing care of the child.
- Evolving developmental and psychological status of the child.
- Special adherence challenges for children.
- The evolving nature of the illness.
- Time for discussion with parents during each visit and time to prepare them for difficult questions that may come later.
- The need to decrease social isolation – this can be achieved by encouraging discussions between parents and other caregivers during clinic visits.

**THE CHILD**

- Information appropriate to the age and development (including emotional development) of the child in response to questions, especially with regard to clinical visits, drawing of blood, and medication.
- Parents should never knowingly tell untruths, as this will decrease the trust between parent and child.
- Ascertain what the child knows or suspects about his/her diagnosis.
- Determine the child's state of mind – depressed, worried, anxious?

- Determine whether the child is sexually active or not.
- Children who ask direct questions are ready to hear about their diagnosis and will seek that information elsewhere if the parents and health care providers are not forthcoming in providing it.

**THE PARENT**

Factors to consider:

- Parental HIV status.
- Disease stage.
- Who is the child's primary caregiver?
- Who is emotionally closest to the child?
- Is the parent living at home?
- Communication style within the family.
- Impact of the disease on other family members.
- Fears, concerns and attitudes with regard to disclosure.

The book by Mary Tasker, *How Can I Tell You?*, outlines the four common phases of disclosure by a parent, namely:

- **Secrecy/privacy phase.** This occurs immediately after learning the child's diagnosis. The diagnosis is kept secret or private from anyone outside the medical centre.
- **Exploratory phase.** This occurs later, when the parent is willing to disclose the diagnosis to a close friend or family member. The parent will often also offer the child a plausible though not entirely thorough explanation for the numerous medical visits.
- **Readiness phase.** This occurs when the parent begins to disclose to a larger network and starts entertaining the idea of fully disclosing to the child.
- **Disclosure phase.** This is when full disclosure to the child occurs. Lee et al. found that disclosure is best done over a period of time at different phases of the child's life and involving a multidisciplinary approach that includes the parents/caregivers.

Parents can be helped through these phases taking into account what has been ascertained about the child, the parent, the family and the community as a whole.

**REACTIONS TO DISCLOSURE**

**DENIAL**

Children may deny having the disease, or may claim that they are cured of the disease. This becomes a problem when the child starts to refuse medical care, misses appointments or sabotages treatment. These children may need the intervention of a psychologist to help them cope with coming to terms with the diagnosis. Support groups with similar children may be helpful.

**SADNESS AND DEPRESSION**

This is a more common response but is usually short-lived. The child requires reassurance and assistance in exploring and coming to terms with his or her feelings, fears and anxieties. Should a more chronic period of depression set in, a professional should be consulted to help deal with the depression.
SUICIDE
This is extremely rare, and should there be an expression of suicidal ideation a mental health professional needs to be consulted.

GOOD WAYS TO DISCLOSE

- Disclosure should occur between the primary caregiver, parent and child with or without health care providers.
- It should be:
  - driven by the child’s agenda
  - planned
  - private
  - loving and reassuring
  - unhurried.

WAYS NOT TO DISCLOSE

Disclosure should not be done:

- accidentally
- in the heat of anger
- AFTER the child has figured it out for him/herself
- when the child puts a provider ‘on the spot’.

WHAT TO SAY

‘You know you have been coming to this clinic for a long time? You come because you have a long-standing illness/infection that requires regular check-ups and medication. The medication you take helps the illness, and helps to keep you well.’

When the parents and health care provider are sure that the child understands these concepts, the disclosure can go a bit further. ‘That illness or infection we spoke about – well, it is caused by a virus/germ. Do you know what a virus/germ is? The medication that you take to keep you well works by killing this virus/germ.’ After a period of time, the parents or health care provider may then name the virus.

The parent should practise the disclosure with the health care provider. They should also explore answers to some of the difficult questions, like ‘How did I get it?’ or ‘How did you get it?’

CONCLUSIONS

We believe that every health care provider should develop a plan with the HIV-infected child’s parents or guardians that will outline the process of HIV disclosure. Disclosure is a process that should occur over a period of time, and should occur before the child discovers his or her own diagnosis. Role-playing with the child’s parents or guardians will enable them to prepare answers for difficult questions and facilitate support for adverse outcomes following disclosure.

REFERENCES

2. Lipson M. Disclosure within families. AIDS Clinical Care 1993; 5: 43-44.