

Fluids and the use of artificial hydration

This factsheet is for guidance only and has been produced with assistance from healthcare professionals

We hope to answer your concerns, but if you have any more questions please ask a member of staff

This factsheet answers some frequently asked questions about fluids and the use of artificial hydration (fluid given by a drip) in advanced illness. It is aimed mainly at carers, but some patients may also find this information helpful.

If someone doesn't drink, won't they die of dehydration? Isn't fluid even more important than food?

It is normal for people who are dying from advanced cancer and other illnesses to eventually stop drinking and to stop feeling thirsty. As the body weakens and the systems start to work less well, there is less and less need for fluid.

It is important to remember that it is the illness which is making the body systems fail, not a lack of fluid. If someone is very weak and is given fluid by mouth it may go down the 'wrong' way, and make them cough and splutter.

What can I do if they complain that they have a dry mouth?

A dry mouth can be a very common problem at any stage of the illness. Ask the medical or nursing staff about it. This feeling is quite different from feeling thirsty. Medicines such as special saliva sprays and gels may be helpful.

What can I do to help?

If staff feel it is safe to do so, you can carry on offering drinks (as the staff will do when you are not there) but don't be surprised if your relative only wants a few sips at a time. The staff will help you to make sure that the drinks are not causing coughing and spluttering.

Some people like to have their favourite drink frozen as an ice-lolly or ice chips – this can be easier to suck on than trying to drink.

When someone is dying and no longer taking drinks from a cup, you can use a sponge or soft toothbrush dipped in cold water (or the person's favourite drink) to help to stop the mouth getting dry. A member of staff will be happy to show you how to do this.

Do you ever use drips?

Yes. A drip is the name for fluid, which is usually sterile salt water, given to the patient through a tube that goes in either under the skin or through a vein. Sometimes the doctors and nurses may feel that giving a drip might help, particularly if there is a suggestion that the person is thirsty. If a drip is started, it will be reviewed by staff over the next 24 to 48 hours to see if it helps or if there are any side effects from it.

However, for most people with only hours or days to live, their body systems are shutting down and thirst is usually not a problem. Drips may not help and good mouth care is the most important comfort measure.

Can you give drips at home if they might help?

The pros and cons of a drip at home will be weighed up carefully in every situation. These decisions can be difficult and are always tailored to the individual patient and in accordance with national guidance. It is sometimes possible to give drips at home, though a nurse needs to be present regularly to monitor them.

Are there disadvantages to drips?

Yes, in the last few hours or days of life the body cannot handle fluid as effectively as before – giving drips can sometimes make things worse by overloading the delicate fluid balancing mechanisms of the body. If this happens the person may experience ‘chestiness’ or noisy breathing, and swelling of the arms or legs, as their body cannot process the fluid from the drip.

Once a decision about a drip has been made is it final?

No. The doctors and nurses will always monitor the situation and discuss it on a regular basis. The patient always has the right to say that they want to have a drip removed. If the patient is too ill to make that decision the doctors and nurses will make a careful assessment and have a discussion with the patient’s family about the right thing to do.

The ultimate responsibility for decisions about starting and stopping a drip rest with the senior doctor or clinical nurse specialist caring for the patient. Any decision that is made can always be reviewed.

This leaflet is updated regularly. For the most up to date information please visit <https://loros.co.uk/our-care>.

In cases of comments or complaints, please contact: Chief Executive, or Director of Care Services, LOROS, Groby Road, Leicester LE3 9QE or, Care Quality Commission, East Midlands Office, Citygate, Gallowgate, Newcastle Upon Tyne NE1 4PA

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LOROS

Hospice Care for Leicester, Leicestershire & Rutland

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and your family

LOROS Groby Road, Leicester LE3 9QE

☎ (0116) 231 3771

✉ info@loros.co.uk

✕ LOROSHospice

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