

This guide has been provided to give you easy-to-follow explanations of the day-to-day aspects of tube feeding, as well as handy tips from other parents and caregivers. To begin with, all this information may seem overwhelming but with practice and support, all of these things will become second nature to you and your family.

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If the pump stops working or you have any questions please contact Nutricia Homeward:

UK 0800 093 3672 NI 0800 169 5173 ROI 1800 221 800

What's inside this guide?

Even once the decision has been made to go ahead with tube feeding for your child, you are still bound to have concerns and questions. Asking yourself questions such as 'what happens next?' and 'can I manage this?' is normal at this stage, as everything about this is new and not what you are used to.



Adjusting to life with a feeding tube may take a bit of time, and it's likely that more practical questions will arise as you and your child get into a routine. With the right information and support, over time you will become experienced in tube feeding your child, and begin to experience the many advantages that tube feeding brings.

Do not feel nervous about discussing any concerns and questions you have with the healthcare professional (HCP) involved in your child's care. There are also a number of specialist organisations and parent associations (details at the end of the guide) that can provide you with advice and support at this stage and in the future.

Many parents want to learn more at this stage, and express the need for more general tube feeding information and practical tips so that they feel more confident when handling their child's feeding tube.

Choice of feeding tube

You will choose which type of feeding tube is best for your child after a discussion with your child's doctor/dietitian, taking into consideration his or her specific needs.

There are three types of feeding tubes, which are described below. Each of these feeding tubes differ slightly in terms of placement; however, their overall aim is the same; to provide your child with all the nutrients they need to help them grow, stay healthy and be happy, supporting them to get stronger to cope with their underlying conditions.

1. Nasogastric tube (NG tube)

This type of tube goes directly to the stomach, but the insertion point is through the nostril, not the stomach. It is typically considered for short term use (i.e. usually 6–8 weeks, although the time of tube feeding varies depending on the child's condition, and the consultant's advice).

For this type of feeding tube no surgery is required for its placement.

These type of tubes may accidentally move out of place; tube position needs to be checked before each feed. If the tube is needed for longer than 6-8 weeks a replacement tube is needed, a small dressing may be required on the nose or cheek.



2. Nasojejunal tube (NJ tube

These are less commonly used than other types of feeding tubes. They are also inserted through the nostril, but – unlike the NG tube – reach into the small intestine, not the stomach.

This type of feeding tube is suitable if a child is having problems with digestion or absorption.

These types of tubes always have to be replaced in the hospital. More easily dislodged, daily position check needed, short term tube replaced regularly, a small dressing may be required on the nose or cheek.



3. Gastrostomy tubes

A gastrostomy tube is placed directly into the stomach through an opening in the abdomen, known as a stoma, allowing enteral feed to be fed directly into the stomach. When a child needs a tube for a longer period of time, gastrostomy is in most situations recommended as the preferred option. Gastrostomy tubes include initial placement tubes and replacement tubes.

Initial placement tube:

Percutaneous endoscopic gastrostomy (PEG tube)

The tube is placed by means of an endoscope, typically under a general anesthetic in children. Once the stoma has healed, the tube can be replaced, if required, by a gastrostomy balloon replacement tube. This can take 12 weeks or more depending on waiting times

Less visible than the NG tube and can last up to two years or as indicated by specific manufacturer's guidelines.

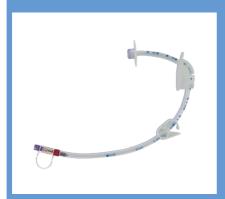
- Requires surgery for placement.
- Stoma infections can occur.

Replacement tube:

Gastrostomy balloon tube (G tube)

Low profile gastrostomy balloon tube (button)

This has a much shorter tube than the PEG tube and sits close to the skin. There are different types of buttons available, for example the MIC-KEY and the CuBBy button. This option is generally preferred by healthcare professionals and parents alike as it is more practical for use in children.





Your child's healthcare professional will provide advice on how to care for the tube



Equipment you may need

Once, after discussion with your child's doctor/dietitian, it has been established what the best feeding tube is for your child, you are probably asking yourself certain questions about what happens next:

Additional equipment required to start tube feeding

- Pump
- Giving set
- Extension set (only applicable when button devices placed)
- Syringe

Type of enteral nutrition product*

- Ready to feed nutrition, i.e. tube feeds only
- A combination of oral feed and ready to feed nutrition, i.e. tube feed

Feeding routine*

- How do I give my child enteral nutrition through the tube?
 - Pump method
 - Gravity/bolus method
- Feeding regimen
 - Continuous feeding
 - Overnight feeding
 - Bolus feeding

^{*}Type of enteral nutrition products and feeding routines should be decided by the dietitian or doctor in partnership with the parents/carers.

EQUIPMENT THAT MAY BE REQUIRED TO START TUBE FEEDING

The feeding tube that has been selected for your child comes as part of a package of equipment designed to make feeding your child as straightforward as possible. All of the additional equipment you need to start tube feeding at home will be given to you by your child's care team, who will also explain to you how to use it.

The kit includes:



A pump to deliver the feed through the giving set at a controlled rate set by you.



A plastic **extension tube** that attaches the giving set to the gastrostomy tube (needed only if low profile/button is in place).



A **feeding tube** may have two ports; one for balloon inflation with water (Luer slip syringe required), and one for administering the enteral nutrition product. However, a lot of PEGs and extension sets will only have a male Luer port.



A **giving set** – a plastic tube attached to the bag containing the feed at one end and the feeding tube or extension set at the other with the feeding pump in the middle to deliver the enteral nutrition product at a set rate and volume.



Syringes* to administer feed, water, medication, or aspirate (remove liquid from the feeding tube). Syringes come in different types and different sizes (e.g. 10 or 60 ml).

^{*}Syringes used for balloon checks are different to enteral flushing and feeding syringes.

TYPES OF TUBE FEEDING

When it comes to deciding what to put into the tube, your child's dietitian or doctor will work with you to find what type of enteral nutrition product is best. The choice will depend on your child's nutritional needs, age, growth rate and any allergies for example.

Ready to feed nutrition

Most dietitians/doctors recommend the use of ready to feed nutrition. Its advantages are that it is:

- Specifically made for children who need a feeding tube
- Specifically designed to meet children's nutritional requirements
- Sterile
- Convenient

- Flows easily through the feeding tube so less risk of blockages
- Can be used alone or in combination with oral feeding



Combination feeding

If your child's doctor/speech and language therapist agrees, and your child can still consume food orally, you may prefer to combine feeding him or her by mouth in addition to tube feeding. Always discuss the possibilities with your healthcare professional before commencing combination feeding.

Some children like to eat normally during the day and are tube fed only at night, others are tube fed only after meals if they cannot eat enough food by mouth. In some cases, children may not want to eat by mouth at all, and there are a few reasons why children are reluctant with oral intake – this can be because they associate it with feeling ill or their doctor has recommended them not to

Did you know?

If needed the tube can also be used to administer medication. This can be an important benefit if your child has difficulties swallowing or dislikes the taste of the medicine. Your pharmacist will be able to give you advice on when your child's medication should be taken (e.g. on an empty or a full stomach), and how best to administer different forms of medication (i.e. liquids, tablets, or capsules) through the tube.

How do I put the feed in the tube?

Getting feed into the tube can be achieved using a pump or syringe.

With the pump, you can programme the rate at which the enteral nutrition product flows through the tube so that you know exactly how much your child has been given. Your choice of pump will depend on your circumstances and your child's needs. Specific details on how to use different types of pump are provided by your pump supplier.

Gravity feeding is the method of feeding using an open syringe/bolus where you slowly pour an enteral feed into the syringe or an extension set. The speed of delivery can be varied depending on the height of the syringe. If required, your dietitian/doctor will be able to provide more information on the differences between these feeding methods. In some cases a syringe can be used with a plunger to administer the feed.

For how long should I feed my child?

Depending on your child's nutritional requirements, medical needs and his/her preferences, your child's doctor or dietitian may recommend either continuous or bolus feeding.



With continuous feeding, the enteral nutrition product is given slowly over a number of hours using a pump (see below). It is possible for this to happen overnight whilst your child is sleeping.



Bolus/portion feeding is when the enteral nutrition product is given by pump or syringe over short periods of time several times a day.

Visit *nutriciaflocare.com* to learn more about pumps, feeding tubes and giving sets. The website also gives access to interactive training tools, such as an online pump simulator.

^{*}Feeding routines should be decided by the dietitian or doctor in partnership with the parents/carers.



POTENTIAL COMPLICATIONS

No matter how experienced you are, occasionally things can go wrong and accidents may happen when tube feeding your child.

If you are in any doubt as to what to do when tube feeding your child, or if any problems arise, there are always places where you can seek help.

- If the pump stops working please contact Nutricia Homeward on 0800 093 3672
- You should contact your local hospital if, for example:
 - The feeding tube comes out
 - The tube is still blocked
 - Your child is unwell (diarrhoea, constipation, upset stomach, etc)
 - Your child has pain on feeding.
 In this instance please stop feeding
- If your child has redness, swelling or discharge around the tube site, please contact your doctor immediately.

AT A GLANCE CHECKLISTS



Use the following checklist as a reminder before and during feeding time. Additional information is available on tube care from your healthcare professional.

Thi	ings to check at feeding time
	Check the label of the tube feed to make sure it is correct.
	Follow the hygiene rules as outlined on the label.
	Check that the feeding tube is in the right position before feeding (your child's care team will show you how to do this).
	Flush the tube through before and after adding the feed (or medication) to avoid blockages, or follow the instructions provided by your dietitian/doctor.
	It is important to position your child in an upright position whilst feeding and for one hour after feeding as advised by your nurse.
Thi	ings to check every day
	Clean around the insertion site of the feeding tube every day.
	Look after the skin around the tube to avoid irritation (your child's care team will advise you about what this entails).
	Make sure your child stays hydrated by giving them water through the tube, or orally if safe. Speak to your dietitian to discuss how much water you can use.

In the beginning, it may seem that there is lot to remember, but with time all of these things will become second nature to you, and you and your child will be able to enjoy feeding time.

AWAY FROM HOME



Use this checklist if you want or need to travel away from home, either for a special occasion, holiday or to visit the hospital for a check-up or test, to ensure that you have everything you need.

Enteral nutrition product
Medication, if applicable
Water for flushing
Syringes
Container
Extension tube and clamp if needed
Feeding pump (if needed) and pump charger
Spare giving sets and replacement feeding tubes, syringes and sterile water, i you are comfortable doing this yourself
Towels, wipes, tape
Soap for skin care
A way to hang the feeding bag while away from home (usually in a carry system)
Plastic containers/bags to store feeding supplies
Phone numbers for the healthcare provider
Your child's care plan (see next page)
This booklet

Nutricia Homeward Holiday Service

At least six weeks in advance of a holiday abroad, contact Nutricia Homeward for information on the Holiday Service

YOUR CHILD'S CARE PLAN

This page is designed for you to fill in and refer to, or to give to a caregiver, for example, so they have all the information they need to feed your child for the day.

CHILD'S PHOTO

Type of tube	
Size of tube	
Date placed	
Feeding method	
Feeding time	
Name of enteral nutrition product	
Amount of product per day:	
The flow rate set on the pump for your child is	mls/hour
The dose or volume on the pump for your child is	mls/hour
Flushing	
Amount before and after feeding	mls
Amount during continuous feeding (everyhours)	mls
Amount before and after medication	mls
Doctor's name and number	
Dietitian's name and number	
Patients & carers helpline	
In case of an emergency, ring	



Frequently asked questions

Do I need to keep the enteral nutrition product refrigerated?

Store unopened product at room temperature (between 5–25°C) in a cool, dry place. You do not need to keep unopened product in the fridge. Once opened, it can be kept for a maximum of 24 hours.

The nasogastric tube has been accidentally removed or has changed position. What should I do?

If the tube has been accidentally removed or changed position it is important to contact a healthcare professional for advice — it is important the tube is not used until it has been confirmed that it is in the correct position — the stomach. Otherwise contact the Nutricia Homeward Enteral Nurse Specialist, district nurse or GP. They will arrange to have another tube inserted. Always re-check its position before starting the feeding process.

My child's PEG tube has accidentally been removed. What do I do?

Place a dressing over the site and go to hospital immediately for reinsertion of the tube. A replacement tube should be inserted as quickly as possible. Contact the hospital beforehand to avoid waiting in Accident & Emergency.

What should I do with giving sets and reservoirs after I have used them?

You can dispose of giving sets, reservoirs, containers, bolus adapters, 2-pack connectors and used product packs with household waste.

If my child needs to take a break during feeding what should I do?

If your child is taking a short break that is less than an hour, you should:

- Disconnect the tube from the giving set;
- Replace the end cap on the giving set; and
- Flush the feeding tube.

If your child is having a longer break of more than one hour you should do as above and then put the entral nutrition product into the fridge, leaving the giving set attached. When you take it out of the fridge, leave it at room temperature for 30 minutes before you start feeding again.

FREOUENTLY ASKED OUESTIONS CONTINUED

Can my child have baths, showers or swim?

Yes, once the stoma tract has been fully formed after they have had the PEG or JEJ tube in place your child can bath, shower or swim as normal. When your child is allowed to take baths and swim, make sure that the tube is closed before immersing it in water. Be sure that you dry the area thoroughly afterwards.

NOTE: The PEG Clinical Nurse will advise you on exactly when your child can hathe.

Will my child need the tube forever?

Some children need tube feeding for short periods of time, while others may need them long term. Your child's doctor/dietitian will discuss the expected length of time your child will require a feeding tube. Your child's progress will be reviewed regularly by their consultant and dietitian.

Can my child eat normal foods while being tube fed?

You will be informed by your doctor/dietitian if your child can eat while on enteral nutrition products. If they are allowed to eat, you will be given information about what types of food and amounts your child can eat. If your child has swallowing difficulties, a speech & language therapist will assess your child's swallow and will advise you if it is safe for your child to eat by mouth.

I no longer require the feeding pump. What should I do?

It is important to return the pump to Nutricia Homeward when you have finished using it.

You can arrange to return the pump by contacting Nutricia Homeward on 0800 093 3672.

We will arrange to collect all the equipment and we will pay any charges related to the return of equipment.

INFORMATION, ADVICE AND SUPPORT ABOUT TUBE FEEDING

www.pinnt.com www.nutricia.co.uk www.tubefeeding.co.uk For more information please contact your healthcare professional.

READING MATERIALS

The following are additional sources of information about tube feeding which you may find helpful:

How do you feel about tube feeding

Practical advice on the signs that may indicate that your child requires a tube feed.



Making the decision to start tube feeding

For information about the range of enteral nutrition products available.



To request a copy of these booklets please contact the Nutricia Resource Centre on *03457 623653* or *resourcecentre@nutricia.com.*

GLOSSARY OF TERMS

Bolus feeding	The feed is given in smaller volumes (e.g. 200ml) one at a time, several times throughout the day (e.g. up to 7 feeds) with the use of a pump, gravity or a syringe		
Button	A low profile gastrostomy at skin level, and is retained by a small balloon inside the stomach		
Clamp	A small device on the feeding tube which, when pressed closed, prevents leakage from the feeding tube		
Continuous feeding	The feed (e.g. 1000ml or 1500ml) is given slowly over a number of hours using a pump that controls the flow rate of the feed		
Endoscope	A slender tube used to examine the inner part of the body with an attached instrument for biopsy or surgery		
Enteral feeding	A liquid nutrition product passes directly into the stomach or intestine		
Extension set	A tube that can be attached between the giving set and the feeding tube		
Flushes	Water given via the tube to avoid any tube blockage and maintain hydration		
Gastric	Anything related to the stomach		
Gastrostomy	An opening through the abdominal wall into the stomach		
Gastrostomy tube, G tube	A feeding device through which enteral nutrition, fluids and/or medication can be given. A G tube is placed directly into the stomach through an opening in the abdominal wall (stoma) and has a small balloon end present to hold the button in place		
Giving set	A flexible tube that helps to deliver the feed. It is attached to the tube feed bag at one end, and the feeding tube with the pump in the middle		

Gravity feeding	Enteral nutrition that enters the stomach by gravity (i.e. placing the feed above the height of the stomach)
Nasogastric tube, NG tube	A nasogastric tube is designed for tube feeding and placed through the nose, down the oesophagus and into the stomach. The position of the NG tube must be confirmed before each feed can commence
Oesophagus	Often called food pipe, it is a part of the gastrointestinal system where food and drink are moved from the mouth to the stomach
Oral	Anything related to the mouth
Overnight feeding	Typically given during night-time, often through a pump
Pump	A device that delivers the enteral nutrition product through a tube at a set rate and volume
Pump feeding	Enteral nutrition that is given using a pump
Stoma site	An opening from the outside of the body through the skin where the feeding tube enters into the stomach or the small intestine (jejunum)
Syringe	A medical device designed to administer fluid, feed or medications, or remove fluid
Trachea	The 'wind-pipe' that carries air from the mouth to the lungs
Tube feed, formula	Specifically formulated enteral nutrition products designed to meet the patient's specific needs
Tube feeding	A way of getting your body the nutrition it needs. Tube feed is a liquid form of nourishment that's delivered to your body (stomach or small intestine) through a flexible tube
NJ tube	A nasojejunal tube is designed for tube feeding, usually placed in hospital and can't be replaced in the community. The tube is placed through the nose, down the oesophagus, through the stomach and into the small intestine (jejunum)
PEG	A Percutaneous Endoscopic Gastrostomy tube is a gastrostomy tube inserted using an endoscope. The tube has an internal retention disc inside the stomach to prevent it from falling out

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UK 0800 093 3672 NI 0800 169 5173 ROI 1800 221 800

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