



Proveca

Medicines for Children

Sialanar®

**ADMINISTRATION VIA
FEEDING TUBES**

GUIDE FOR CARERS

This leaflet has been given to you by your child's doctor because
Sialanar® has been prescribed for your child



1



Always wash hands
before giving medication

2

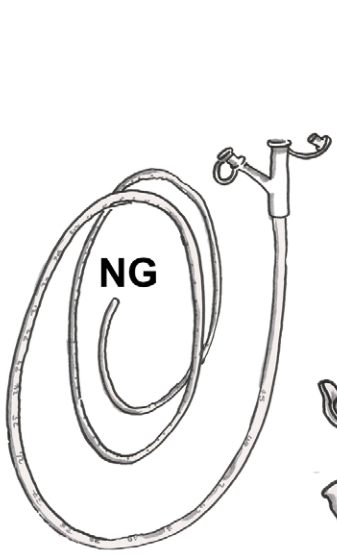
Check the dose & read the
patient leaflet supplied with
the pack



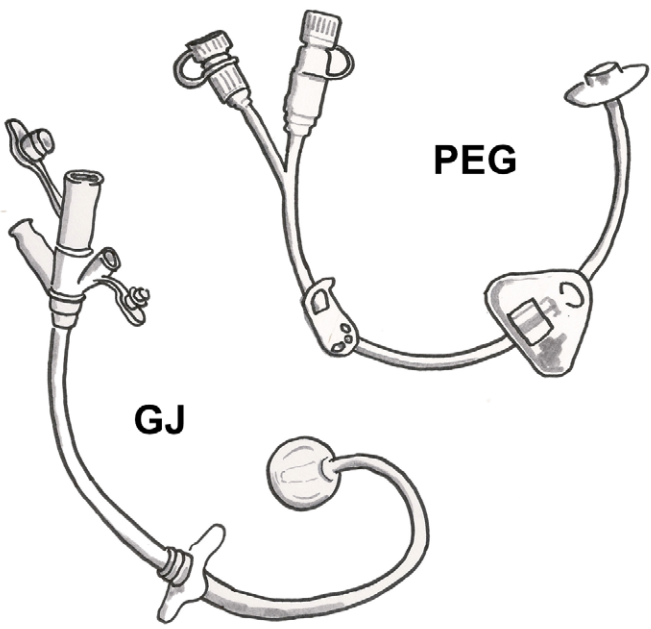
Unscrew child
resistant cap

**NB DO NOT USE THE PRE-SUPPLIED ADAPTOR WHEN USING A
SYRINGE OTHER THAN THE ONE SUPPLIED IN THE PACK**

4 Use a syringe which fits the valve of the feeding tube to draw up the correct dose of Sialanar. Replace cap on bottle



5 Check the dose and then insert the syringe into valve



6 Sialanar will easily flow down the tube



7 After use, always flush with 10mls of water to make sure all the medication has gone down the tube

IMPORTANT INFORMATION ON THE ADMINISTRATION OF SIALANAR

Always give Sialanar® exactly as the doctor has told you. You should not increase the dose without the doctor's permission. Check with your doctor if you are not sure.

Further information about taking Sialanar® can be found in the Patient Information Leaflet and can be obtained from your child's doctor.

How to store Sialanar®

Do not use Sialanar® after the expiry date stated on the label after EXP: The expiry date refers to the last day of that month.

Do not store above 25° C.

This medicine must be used within 2 months of first opening the bottle.

Sialanar® should not be used if the packaging has been opened or damaged.

Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help protect the environment.

Reporting of side effects

If your child gets any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in the package leaflet. You can also report side effects directly via the Yellow Card Scheme at www.mhra.gov.uk/yellowcard

By reporting side effects you can help provide more information on the safety of this medicine.

Side effects should also be reported to Proveca Limited.

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