type 3 Educational needs



Children with type 3 Gaucher disease may need more support than others to reach their full learning potential.

Many children will attend mainstream schools and some will need a more specialist provision.

The information given here mainly relates to England. In the rest of the UK it is slightly different - in Scotland it is called Additional Support for Learning

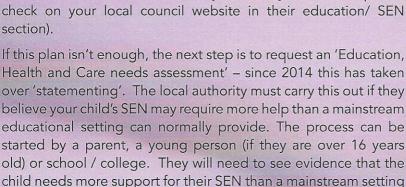
and in Wales and Northern Ireland it is called a Statement of Special Educational Needs. Please check with your local authority for more detailed information.



Before starting school speak to the SENCO (Special Educational Needs Coordinator), show them the main type 3 booklet especially the education section about how type 3 Gaucher disease can effect learning.

Initially support might be provided through the school budget with a School Action / School Action Plus (local authorities call

these different names, such as My Plan / My Plan Plus - please check on your local council website in their education/ SEN section).



Once they have agreed to carry out an assessment they will gather all the information from the parent / child, school/college, other services involved (e.g. Educational Psychologist, specialist

can normally provide. The local authority must tell you in writing

within 6 week whether or not they will assess your child.



teachers, health social care services). The local authority will then decide whether to issue an Education. Health and Care Plan.

The final plan must be issued within 20 weeks from initial application.

If you would like further advice around education please do contact our Patient and Family Support Worker on 01453 549 231