

NEUROFIBROMATOSIS TYPE 1 (Nf1) PASSPORT

Many parents feel that they are constantly repeating themselves with schools, GP's, consultants at hospital etc. about their child/ren's condition and how it affects them.

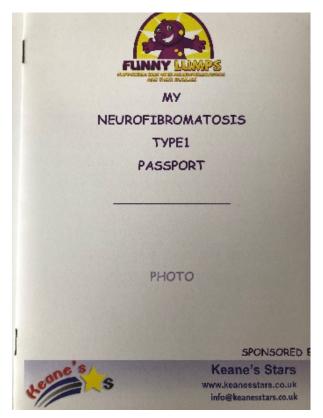
Every time your child changes teacher, sees a different GP or a new consultant it can feel like telling the same story repeatedly. It's not always easy to remember everything and many times you leave a consultation thinking "I wish I had told them about ...".

Taking this into consideration, Funny Lumps developed the Nf1 Passport.

It gives a short explanation of Nf1 and allows you to individualise it for your child.

This means you have separate information for each child in the family who has Nf1 contained in a handy booklet. Whilst there are tick boxes for some of the more common effects of Nf1, there is also ample space to personalise this document with information about your child.

It's also somewhere you can keep a note of the different consultants etc that your child sees and their contact details.



If you would like a copy please email with your full name and address to:

info@funnylumps.org or you can download a copy on our website.