This is me My Care Passport

It should be kept with me and brought with me into any care setting, including hospital.

	My name is:
Click here to add your photo	
from your computer.	I like to be known as:
P	

Please return my passport to me when I go home.

This is essential reading for all staff working with me. It gives important information about **me.** This passport should be kept visible and used when you talk to me or think about me.



My last three hospital visits/admissions

Please give details of these visits/admissions below: The dates, hospital, consultant, reason and outcomes.



More basic information about me

This passport needs to be updated if my needs change.

Where I currently live:

For example - supported living or my family home.

Hours of support I get each day:

Any 1:1 hours I have at home should be provoided in hospital if I am there

Who to contact for more information about me:

Please say name, role and contact phone number.

Other key professionals involved in my care:

Please say name, role and contact phone number.

Learning Disability Acute Liaisons nurses can give extra support to people with a learning disability in hospital. See the back page for contact details of your local team

Key person / people to liaise with about my admission and discharge:

This passport was filled in by:



Date:

Things you must know about me

1. Adverse drug reactions, allergies or intolerances.



Please give details including what my reactions would be.

2. Communication - How well I use and understand speech



Other ways I communicate - signing, pictures or other languages ? How I show how I feel. How I communicate yes and no.

3. Food and drink - Food allergies / intolerances and help choosing



Do I need help filling in menus? How I make food and drink choices. See also the likes and dislikes section.

4. Eating and drinking - What help I need



Does my food need to be cut up or liquidised? Do I use dentures to eat? Do I use special equipment?

If there is a risk I may choke please give details of my management plan and seating & posture.



5. Pain - How I show I'm in pain and how to support me



6. Other medical conditions - Such as diabetes, epilepsy, asthma and depression

See separate medication list.



7. How I take medication - One tablet at a time, on a spoon or via a syringe

Do I need help to make sure I have swallowed?



8. How to support me with medical interventions



Things like taking my temperature, blood pressure, blood test and having injections.

9. How I usually am - for example do I sleep a lot, am I usually very quiet?



10. How do I react to strange places?



11. Keeping me	safe - Do I wander?	Could I fall out of bed?	Do I fall?

12. Things that may worry or upset me - How I may show this.

13. How to support me if I'm anxious or upset - Also see the likes and dislikes section.



14. Behaviours I have that may be challenging or cause risk

What you can do to support me with my behaviours - things that help me relax.

15. My sight - Any problems I have, aids I use like glasses or magnifying glass.

Can I clean my glasses myself?



16. My hearing - Any problems I have, aids I use like a hearing aid?

Can I put my hearing aid in myself? Do I know how to turn it on?



17. Other vital information - Such as advance care decision.

If I have a 'Lasting Power of Attorney' please specify whether it covers 'Health and Welfare' and/or 'Finance and Property'. Please also say if I have an 'End of Life Care Plan'.

Please also say who holds these documents and how to contact them.



Things that are important to me

Important people

Family, friends & staff who support me.

Level of support I need when well

Who needs to stay and how often.





Personal care - support I need with things like dressing, washing and teeth cleaning.



Moving around - for example posture in bed, walking aids and wheelchair.

Do I need help with moving around?



Sleeping - my sleep pattern / routine / time of waking.



My likes and dislikes

Things I like

Could include:

Music, TV, foods, activities and how I relax.



Things I don't like

Could include:

Things that worry me, foods, activities and ways I don't like being treated.

My history - What is important that you know about my life (past and present)

Please also use this space for any further information.

Based on a previous 'Hospital Passport' this version is designed to be used for all people within a variety of care settings.

The content was developed together by Surrey and Borders Partnership NHS Foundation Trust Acute Liaison, Specialist Therapies and Older Adults services, Royal Surrey County Hospital and the Surrey Alzheimer's Association. This passport was designed by The Clear Communication People Ltd and funded by The Learning Disability Partnership Board in Surrey.

Learning Disability Acute Liaison Team contact details:

These are my local Acute Learning Disability Nursing Team contact derails.

You can contact them regarding any care I am having in hospital.

