## Caring for people; keeping people well

Shaping Our Future Wellbeing – Developing the UHB's Clinical Services Strategy

Cancer Workshop

5th November 2014



www.cardiffandvaleuhb.wales.nhs.uk/home

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## Agenda

12:00	Welcome Service Users and Carers Introduction to the afternoon and each other
1:00pm	Welcome to Clinical Staff and Partners Director of Planning to Welcome everyone to the workshop
1:20pm	Setting the Scene Assistant Medical Director for Cancer Services
1:40pm	What does it feel like to use the UHB's services? Service User Story
2:00pm	What do the Clinical Services Principles mean to you? Attendees describe and discuss what the Principles mean to them
2:40pm	Coffee
3:00pm	Working together, what could the services of the future look like? Putting the service user at the centre, attendees draw a service model of the future considering the impact of technology and new ways of working
4:00pm	Sharing the service models developed by each table Attendees present and debate their service models of the future
4.45pm	Next Steps



#### Introduction

- It was fantastic to welcome such a variety of people; service users, staff and 3<sup>rd</sup> sector partners, to our recent Shaping Our Future Wellbeing, Cancer Workshop.
- The aim of the afternoon was to begin to describe what, if we work together, cancer services of the future could look like. Ultimately, the output of the session will support the development of a Shaping Our Future Wellbeing – Clinical Services Strategy for Cardiff and Vale University Health Board.
- Having had the scene set by Abigail Harris, Director of Planning and Dr Meriel Jenny,
  Assistant Medical Director for Cancer Services, the session really began to build
  momentum as those in the room who have used our Cancer Services described their
  journeys.
- This report, the presentations from the day and the storyboards generated during participants discussions, will shortly be available on-line. The website will also provide an insight into the breadth of engagement work underway and track progress through to the publishing of Shaping Our Future Wellbeing – Clinical Services Strategy in September 2015.

Rachel Rayment

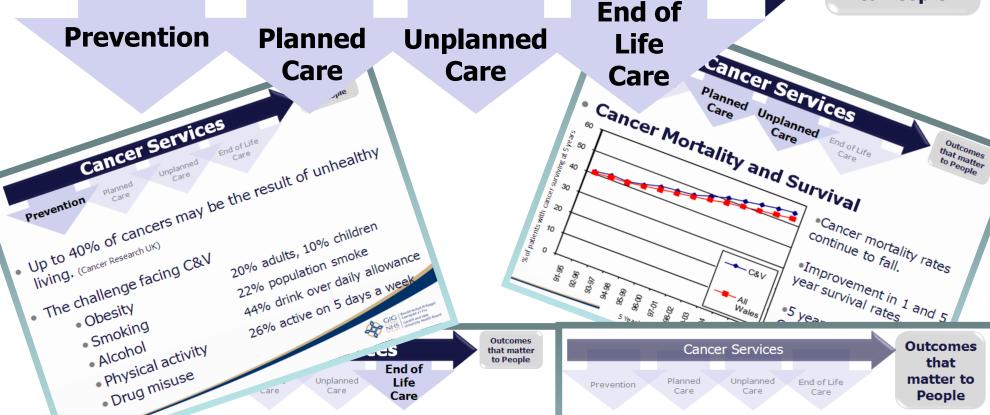
Clinical Lead, Shaping Our Future Wellbeing

Why are we here today?



#### **Cancer Services**

**Outcomes** that matter to People



Page

 Care Priorities are defined for guiding last days of life (Care Priorities Document).

Unplanned

Life

Care

- Use of Rapid Discharge Pathway to speed up discharge and prevent readmission.
- Ongoing education around palliative care and end of life care.

Planned Unplanned End of Life Prevention

matter to People

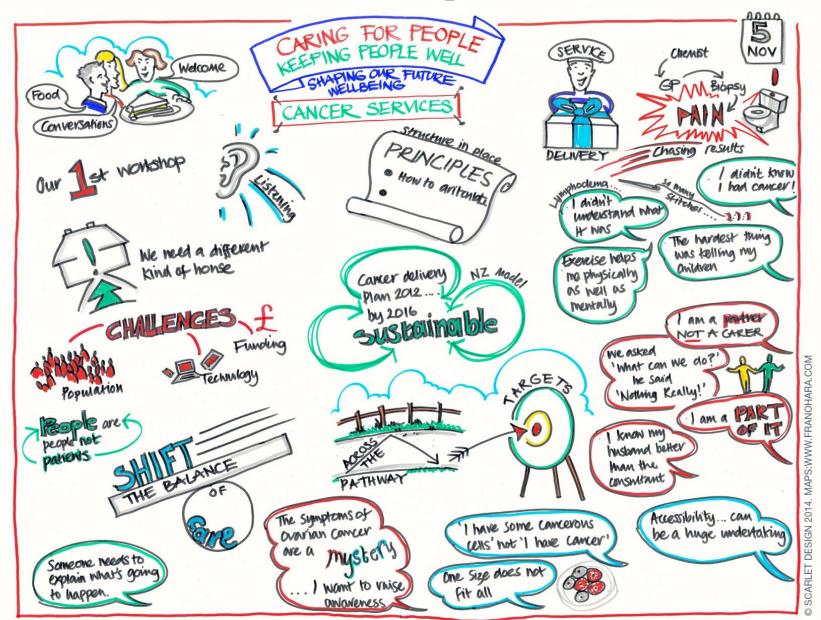
#### Wales Cancer Patient Experience Survey

- Some good
  - Assigning a clinical nurse specialist/key worker
  - •Giving enough info about psychological support
  - Confidence and trust in ward nurses
- Some areas needing improvement
  - Info on potential long term side effects
  - Providing written care plans
  - Offering help and guidance on financial issues

·Health Board working with Partners to improve patient experience



#### **Service User Experiences**



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#### **Clinical Services Principles**

use people who provide services between the Promote equity

**Empower** the Person

- •Support people in choosing healthy behaviours
- •Encourage self-management of conditions

**Home first** 

•Enable people to maintain or recover their health in or as close to their own home as possible

**Outcomes that** matter to People

•Create value by achieving the outcomes and experience that matter to people at an appropriate cost

Avoid harm, waste and variation

- •Adopt evidence based practice, standardising as appropriate
- •Fully use the limited resources available, living within the total
- Minimise avoidable harm
- •Achieve outcomes through minimum appropriate intervention

# What do the Clinical Services Principles mean to you?



A wordle 'word cloud' generated identifying the words used most frequently on the participants post-its.



#### **Empower** the Person

- Timely information
- Time in clinic for patient to fully understand and ask questions
- Clinicians understand my problems as well as clinical priorities
- Empowerment during survivorship
- Give people options don't decide treatment plan before speaking to service user
- Shared decision-making
- Support from people who have experienced my cancer
- Trying to stop your diagnosis being your life
- What can I do to help myself
- Help to make the person feel in control
- Sharing experiences
- Prevention health education
- Detox alcohol and smoking cessation
- No barriers across working with third sector
- Signpost to support







#### **Home first**

- Online/Telephone appointments with relevant education and tests done pre-appointment.
- Specialist roles within the community to provide support. E.g. Use of patients who have completed treatment.
- Cancer nurse specialist home visit.
- Tele-surveillance.
- Access to community palliative care beds for patients who need admission rather than acute beds.
- Improved access to social services for care packages and access to physiotherapy/occupational therapy support to prevent admission.
- Parking!
- Hub and spoke model for testing services, with results by post.
- Improved communication between hospital and home, especially clear plan at discharge.
- Community phlebotomy and practice nurse service every day, and out of hours weekend cover.







#### **Outcomes that** matter to People

- Health promotion
- Shared outcomes, may be different to family, may be different to professionals.
- Importance of psychological support for patient and family
- Telephone support 24/7
- "Key worker" coordinated care with all services working together
- More clinical trials
- Individualised treatment based on patient values
- Acknowledgement that it is not always a death sentence
- Managing expectations
- Follow-up as needed not always routine.
- Honest discussion of risk with respect for patient choice
- Pain and symptom control as well as survival/cure
- Accurate timely diagnosis, avoid over diagnosis



#### Avoid harm, waste and variation

- Balancing risks and benefits
- Effective communication between all parties
- Increase medical practitioner awareness/training
- Adopting a new attitude
- Access to good evidence base
- Standardising practices
- Innovation to avoid waste
- Feeling in control for both patients and providers
- Focus on the patient and care
- Optimise expected benefits
- Listen to the patient
- Reduce waste around appointments
- Identify the barriers to doing what is intuitively right
- Best use of resources time, staff, services, etcetera





Promote equity between the people who use and provide the service

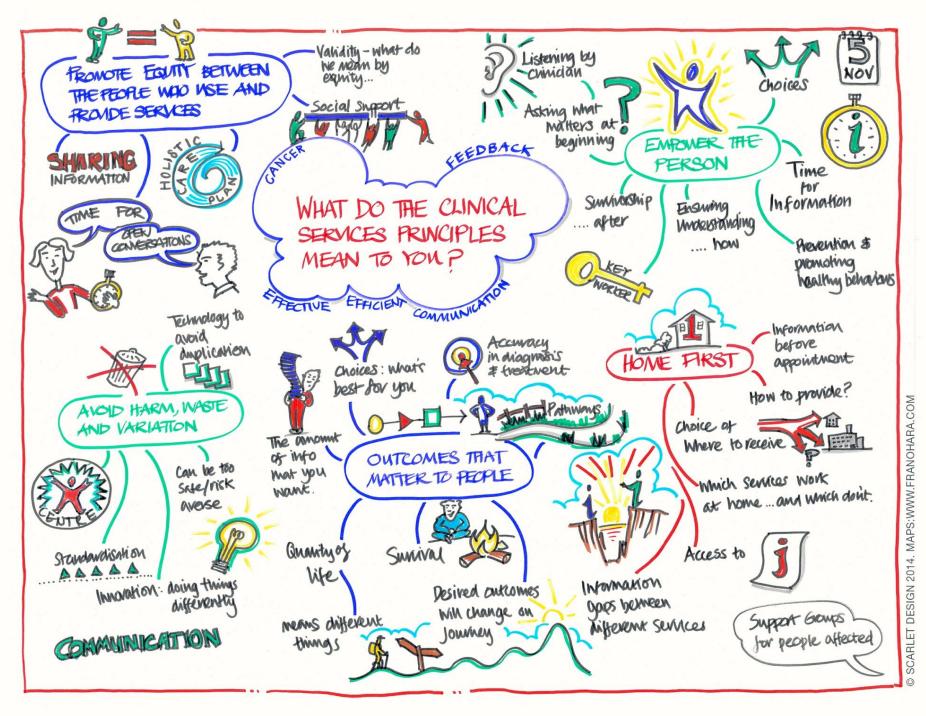
- Right information for the right person at the right time
- People empowered to identify what care they want
- Honesty and openness
- Listening
- More emphasis on partnership in care
- Supported self-care and management
- Coproduction of services
- Opportunity for the person to discuss their needs and be listened to-not just a treatment plan-but a care plan and to have a written copy of the agreed plan to take away
- Able to be responsive to change
- Support for professionals to change practice/perspectives
- Wherever you are diagnosed with cancer that you get the same psychological support
- Accountability and responsibility for outcomes shared
- Family should be the unit of care
- Equity of access involves doing more in areas where people have the poorest health









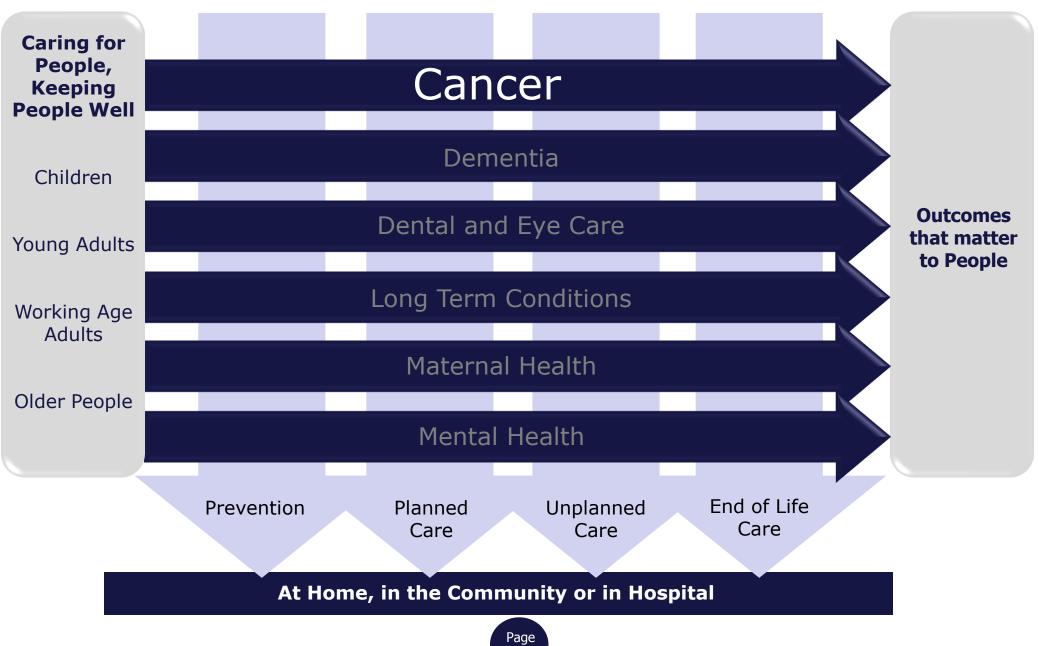


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### **Future Services**

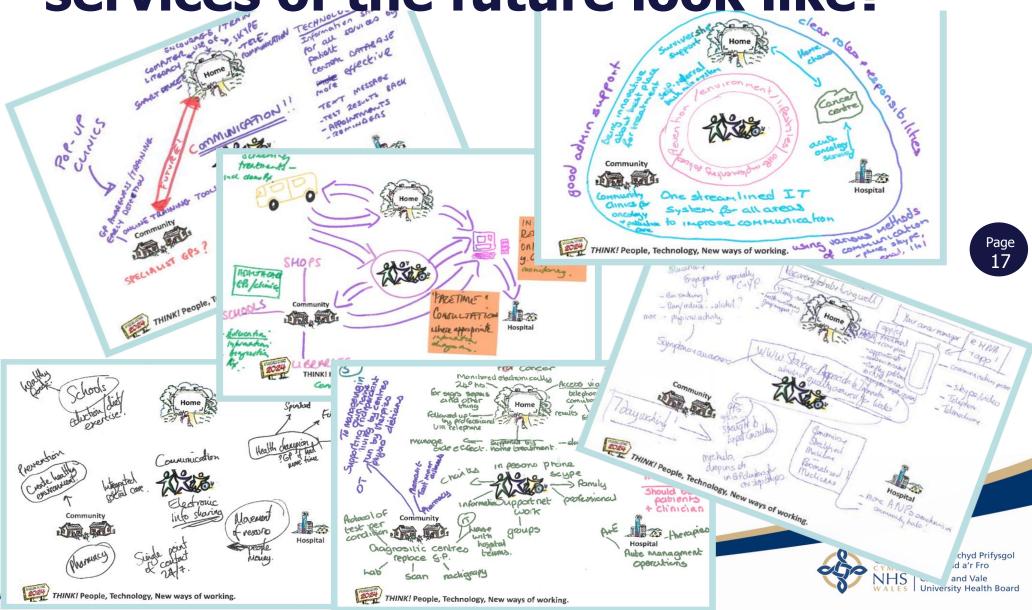


#### **Clinical Services Framework**

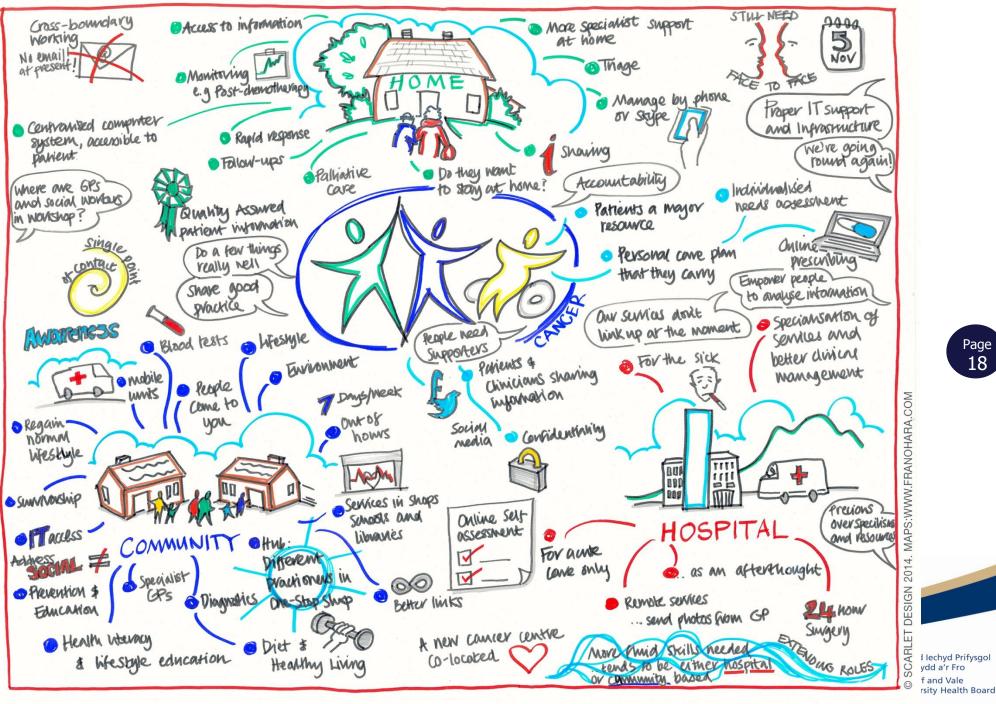


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Working together, what could the services of the future look like?







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#### **Next Steps**

 Within the month, provide you with the output of today's workshop



- Refine the workshop output through key interest groups
- At the Feb Clinical Senate, and at Feedback
   Session 13<sup>th</sup> March provided combined feedback on all the workshops
- Engage on a draft Clinical Service Strategy document
- UHB Board approve Clinical Service Strategy September 2015

