

SFHCHS204

Design solutions to meet technical, scientific and/or engineering requirements for healthcare



Overview

This standard relates to the new development and risk assessment of solutions required to support clinical development and the delivery of specific services to meet the clinical needs of the individual.

It may involve designing a prototype to test out a proposal prior to construction or it may relate to the design of specific features of facilities and accommodation to house specialist medical equipment especially those related to safety.

Individuals undertaking this function will liaise closely with key stakeholders and relevant others to confirm accuracy of information and compliance with relevant legislation.

Users of this standard will need to ensure that practice reflects up to date information and policies.

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Performance criteria

You must be able to:

- P1 work within your level of competence, responsibility and accountability
- P2 liaise and work with key stakeholders to ensure that all current influencing factors regarding the design have been taken into consideration
- P3 discuss with key stakeholders any costs involved for developing the preferred solution and factor in any prototypes as required
- P4 propose the relevant and realistic scientific, engineering or technical solutions and identify any feasible options
- P5 carry out risk assessments to identify risk factors involved in the design process and take appropriate steps to minimise these
- P6 review options generated with key stakeholders and reach agreement on optimum solution
- P7 outline the design for the option using the appropriate methodologies at the required level of detail
- P8 produce a design specification which meets the intended outcome/purpose
- P9 ensure the specification has taken full account of any specific design features, requirements or additional accessories to meet its intended purpose and environment
- P10 where appropriate develop a prototype and check it meets the clients needs
- P11 identify defined timescales for the manufacture, adaptation and/or implementation of the approved design solution
- P12 collate the appropriate technical, scientific or engineering information to produce a detailed report
- P13 present approved solutions to key stakeholders, in the appropriate format, and at the required level of detail for the proposed manufacture
- P14 maintain full, accurate and legible records of information collected and store in correct location in line with current legislation, guidelines, local policies and protocols

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Knowledge and understanding

You need to know and understand:

- K1 your own level of competence, authority and specialist knowledge base related to the design process
- K2 the range of stakeholders involved in the design process and their clinical and/or information needs
- K3 the importance of and how to liaise with key stakeholders to select and agree options to meet the clinical need of individuals
- K4 where, when and how to obtain specialist advice and support
- K5 methods of risk analysis, risk benefit analysis and the risk factors associated with the design solution and how to apply them
- K6 the range of design methodologies and how to design solutions or make adaptations to existing resource or facility to address the health condition of an individual or a healthcare group within your work practice
- K7 the range of manual or computer assisted drawing and technical techniques used in the design process
- K8 where applicable, how to apply your speciality knowledge into technical drawings and turn them into a specification for the manufacturing process for a prototype/bespoke model or for a adaptation to an existing resource or facility to meet the individual's clinical requirements
- K9 the importance and role of ergonomics in the design process
- K10 the types, range, properties and limitations of equipment and materials required to meet the intended solution and their interactions with the new design or existing facility
- K11 when and how prototypes and/or adaptations are applicable for verification and validation within the design process
- K12 the framework for intellectual property protection and exploitation related to the design
- K13 the importance of and techniques used in safety test requirements for design prototypes/adaptations and/or products relevant to your work practice
- K14 the current national legislation, guidelines, local policies and protocols which affect your work practice
- K15 the policies and guidance that clarify your scope of practice, accountabilities and the working relationship between yourself and others

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Additional Information

External Links

This standard links with the following dimension within the NHS Knowledge and Skills Framework (October 2004):

Dimension: G2 Development and innovation

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Originating organisation Skills for Health

Original URN CHS 204

Relevant occupations Health, Public Services and Care; Nursing and Subjects and Vocations Allied; Health Professionals; Healthcare and Related Personal Services

Suite Clinical Health Skills

Key words adapt, assistive technology, rehabilitation, calibrate
