Intelligent Systems

CLINICAL PRACTICE GUIDELINES - INTRODUCTION



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Summary

Clinical Practice Guidelines

- Definition
- Clinical Specialty and Category
- Types of Clinical Practice Guidelines
- Development Process
- Important organisations
- Benefits
- Shortcomings
- The role of Artificial Intelligence in guideline application

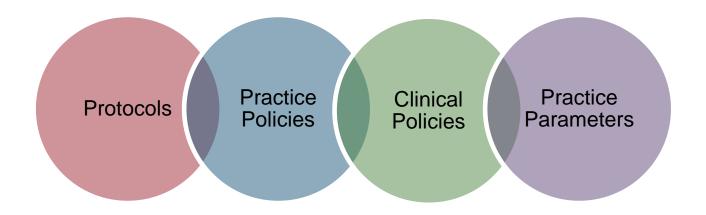
Objectives

- Understand the function and scope of Clinical Practice Guidelines (CPGs);
- Identify the key phases in the development process of CPGs;
- Identify important organisations in the creation and standardisation of CPGs;
- Identify benefits and shortcomings of these documents;
- Determine which aspects Artificial Intelligence may help to improve.

Definition

CPGs are systematically developed statements containing **medical recommendations** to assist healthcare professionals and patients about appropriate healthcare in specific clinical circumstances (Miller and Kearney 2004).

Other designations include:



Clinical Specialty and Category

Medical fields touched by the CPG

Clinical Specialty

- Family practice
- Pediatrics
- Cardiology
- Surgery

• . . .

Determined by the types of recommendations provided

Category

- Diagnosis
- Evaluation
- Treatment
- Management
- Prognosis

• . . .

 According to the type of information CPGs are built upon, there are two types of documents:

Consensus-based guidelines

Evidence-based guidelines

Which one is more appropriate



Consensus-based Guidelines

A common form of guidelines resulting from the consensus of expert opinions. A process that seeks the agreement of all parts.

The opinions of the **minority** must be considered.





Consensus-based Guidelines

Nominal Group

Delphi Technique

Team members write down their ideas of the problem;	It utilizes a series of questionnaires administered by a central individual of experts who never meet together;
Each idea is summarized so that all members can see them. No ideas are discussed until all are presented and recorded;	As the respondents reply, their questionnaires are summarized;
An open discussion of ideas follows to clarify ideas that members do not understand. No attack or defence is allowed;	A new questionnaire based on their responses to the first, is developed;
Next, members vote (in secret) on the top ideas in order of priority. The eventual decision of the nominal group is the vote outcome.	This repeating process continues until a team consensus on the problem is reached.

Evidence-based Guidelines

- Guidelines developed after the extraction and revision of scientific information from bibliography, clearly distinguishing what is proof from what is opinion;
- Besides advising which treatment is better, the document quantifies in absolute terms the benefits and costs of adopting this or other procedure.

Evidence-based Guidelines

Features of evidence-based guidelines in the COGS (Conference on Guideline Standardisation) checklist (2003)

Clear definition of the scope and objectives

Involvement of all the stakeholders

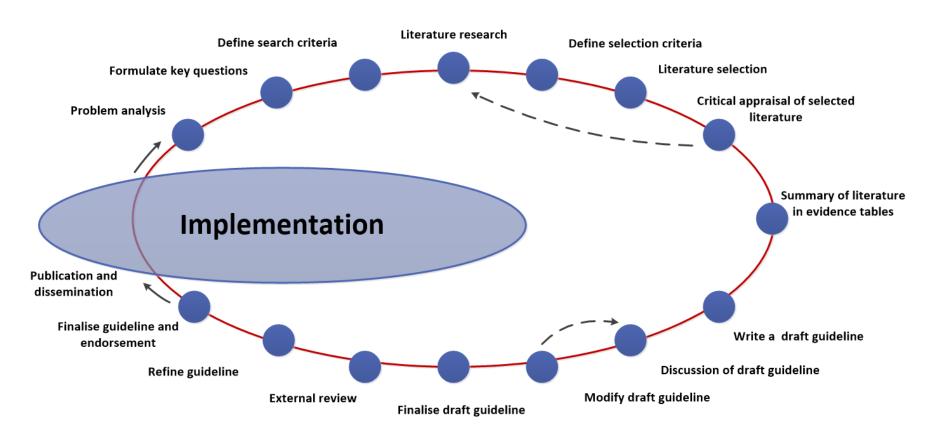
Rigor in the development process

Clarity in the presentation

Applicability

Editorial independency

 There is no standard development process for CPGs. However organisations follow a set of similar phases.



Problem analysis

Formulate key questions Define search criteria Literature research Define selection criteria

Literature selection Critical appraisal

Discussion of the subjects touched by the guideline.

Interviews and questionnaires to stakeholders.



Formulation of key questions to be used as criteria for the bibliographic research.



Search in the literature of the available scientific evidence to answer the key questions.

Definition of selection constraints.



Once the evidence is selected. The methodologies used for producing the evidence are assessed according to checklists (e.g., size of the sample, characteristics of the population)..

Summary of literature in evidence tables

Production of a draft for the quideline

Discussion,
modification
and finalisation
of the draft
guideline

External Review Refinement

Production of evidence tables with scores for the evidence.

More on evidence grading later.



Based on the evidence, produce the document containing: keyquestions, evidence tables, recommendations, implementation costs and references



An iterative process in which the development group evaluates the draft guideline and proposes changes until the final version of the document is reached.



The development group submits the guideline for external revision by an independent organisation.

Finalise guideline and endorsement

Using the suggestions obtained in the external review, the development group produces the final version of the guideline. The guideline is approved by the responsible organisations.



Publication and dissemination

The guideline is published in online repositories and disseminated through newsletters and conferences.

- Evidence Grading
 - Each organisation implements its ad-hoc grading system;
 - However, efforts haven made towards a unified grading system:

Grading of
Recommendations
Assessment,
Development and
Evaluation
(GRADE)

High
Moderate
Low
Very Low

Features

Study limitations
Inconsistency of
results
Lack of precision
Bias

Important Organisations

National Institute of Health (NHI)

http://www.nih.gov/

- Dutch Institute for Healthcare Improvement CBO http://www.cbo.nl/en/
- Scottish Intercollegiate Guidelines Network (SIGN)
 http://www.sign.ac.uk/
- New Zealand Guidelines Group (NZGG)
 http://www.health.govt.nz/about-ministry/ministry-health-websites/new-zealand-guidelines-group

Important Organisations

In Portugal:

Direção Geral de Saúde

http://www.dgs.pt/

For the standardisation of CPGs:

- Guidelines International Network(G-I-N)
 - Comprises 92 organisations and 127 individual members representing 48 countries from all continents.

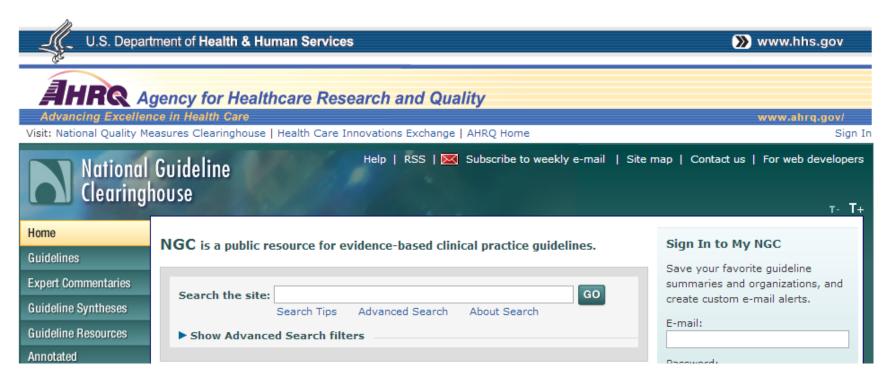
http://www.g-i-n.net/

Important Organisations

Centralised guideline repository:

National Guideline Clearinghouse (NGC)

http://guideline.gov/



Reduce Medical Error

A medical error is defined as a mistake or error committed by a healthcare professional, which results in harm to the patient. It includes errors of execution and errors of planning. It usually results in an adverse event.

Kalra (2004)

5-80 times per 100 000 consultations

11% of all prescriptions

Sanders and Esmail (2003)

39% of adverse event incidence with 18% preventability

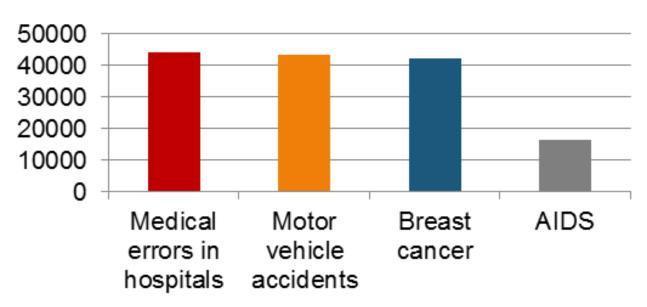
Nicolleta et al. (2003)

108 errors in 64 patients

Proctor et al. (2003)

Reduce Medical Error

Number of Annual Deaths in the US



Institute of Medicine (2000)

Reduce Defensive Medicine

Ordering of treatments, tests and procedures for the purpose of protecting the physician from criticism rather than diagnosing or treating the patient.

Chawla et al. (2008)

8% of all diagnostic testing in the US is defensive

\$2 trillion attributable to defensive medicine

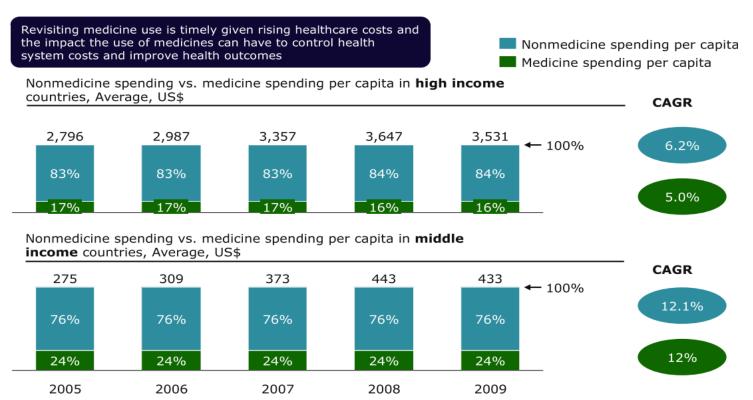
50% of women who will undergo mammography testing will receive a false positive

Chawla et al. (2008)

93 % physicians reported practicing defensive medicine

Studdert et al. (2005)

Reduce Defensive Medicine



^{*}Nonmedicine spending is calculated by subtracting pharmaceutical expenditure from total health expenditure per capita

Sources: IMS Institute for Healthcare Informatics, 2012; World Bank; WHO (latest available data for a subset of countries representing over 50% Of each income group based on World Bank income groupings)

Shortcomings

- Long textual documents that are difficult to consult at the moment of care;
- Difficult maintenance: updating and modifying;
- Healthcare professionals argue that they stifle change and innovation. Moreover, they consider that CPGs are strict rules that do not take into account the social, economical and cultural contexts where the medical practice is developed.

Shortcomings

 Sometimes the documents show some form of ambiguity

Syntactic

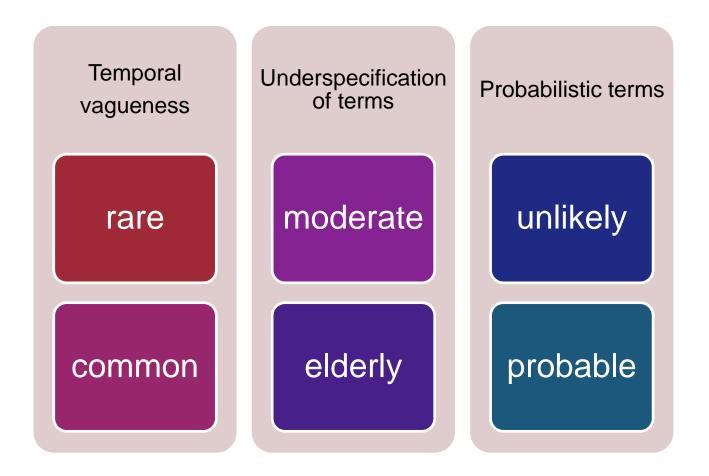
Semantic

Pragmatic

- Syntactic ambiguity occurs when the structure of a statement is not clear, thus impeding its correct interpretation.
- Misplaced (or lack of) punctuation and wrongfully applied Boolean connectors are some of the causes of syntactic ambiguity.
- Characterized by situations where terms can be interpreted in more than one way.
- Misuse of abbreviations, such as the case of the word "cold", which in the context of a guideline can mean "common cold", "cold sensation" or "Chronic Obstructive Lung Disease".
- It happens when the recommendations of CPGs are not consistent or are conflicting with each other.

Shortcomings

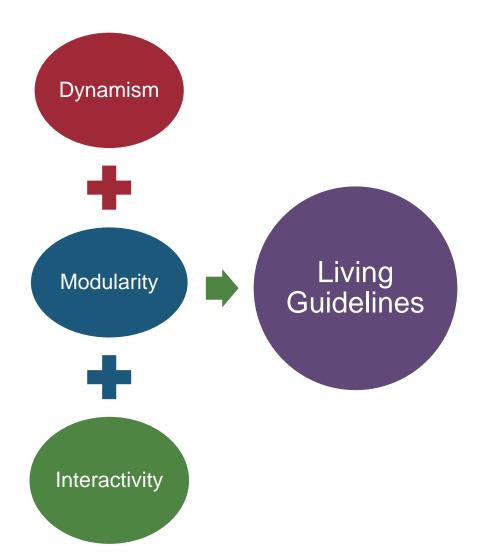
The vocabulary used may also denote vagueness



The role of AI in guideline application

- Al is a field of study that aims to explain and emulate intelligent behaviour in computational processes. It is the branch of computer science that is concerned with the automation of intelligence;
- Al in Medicine is more focused on giving support to healthcare workers rather than trying to replace them.
- Using knowledge representation formalisms, it is possible to create structured representations of CPGs for Clinical Decision Support Systems.
- This way, guidelines acquire a set of desirable features, namely...

The role of AI in guideline application



CPGs in **constant change**, that are easy to update and modify.

These guidelines are modeled as **modules of knowledge**, easily reusable inside other guidelines.

By being integrated in Clinical Decision Support Systems, they provide an **interactive clinical process**. They are fed with the information provided by healthcare professionals and give real-time recommendations.

In the next session...

- Clinical Decision Support Systems
- Computer-Interpretable Guidelines

Intelligent Systems

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