Developing a globally applicable evidence-informed competency framework to support capacity strengthening in clinical research

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ABSTRACT
Capacity development for clinical research is held back by a lack of recognition for the skills acquired through involvement in clinical trials and in other varied types of global health research studies. Although some competency frameworks and associated recognised career pathways exist for different clinical research roles, they mostly apply to a single role or study setting. Our experience supports the need for an integrated approach, looking at the many roles in parallel and at all types of clinical research beyond trials. Here, we propose a single, flexible framework which is applicable to the full global health research team, and can be used for recognising staff by highlighting acquired skills and possible progression between various roles. It can also illuminate where capacity needs strengthening and contribute to raising research engagement. Through systematic analysis of existing competency frameworks and current job descriptions covering 11 distinct, broad clinical research roles, we identified and defined 50 key competencies required by the team as a whole and throughout the study life cycle. The competencies are relevant and adaptable to studies that differ in design, geographical location or disease, and fall in five main areas—(1) Ethics, Quality and Risk Management; (2) Study and Site Management; (3) Research Operations; (4) Scientific Thinking; and (5) Professional Skills. A pilot framework and implementation tools are now available online and in paper format. They have the potential to be a new mechanism for enabling research skills development and career progression for all staff engaged in clinical research globally.

INTRODUCTION
In its 2013 World Health Report (WHR), WHO urged the global research and policy communities to take integrated efforts to strengthen the capacity of all countries to generate evidence for the improvement of their health systems. The need for action towards health research capacity development had already been taken up at the regional level by the Pan American Health Organization (PAHO), which first issued a policy document on the matter in 2009. These two documents particularly emphasise...
the need to develop research capacity in low and middle income countries (LMICs), so that these countries can take ownership in addressing local health issues. Research agendas and practices which are set in high-income settings indeed often fail to adequately target the disease burdens that affect the majority of the world’s populations. To make real changes to global health outcomes, LMICs must therefore equip themselves with the skills to carry out high-quality research studies that address locally relevant research questions.

The capacity needed to conduct global health studies is multifaceted, and ranges from the funding and physical facilities (ward space, laboratories), through to managerial support and personnel training. While all building blocks of a well-functioning research system are necessary and interlinked, arguably the key to developing sustainable local capacity comes from having well-trained teams, and individuals who can demonstrate leadership in research, whatever their role. This concept of a capable workforce is central to many capacity development models, which stress the importance of training individuals and ensuring that they are situated within environments of organisational support, mentoring and supervision.

Although on the rise, the availability and accessibility of adequate training programmes remain limited for many research staff, and particularly for those who are not qualified as medical doctors. Technological developments now make it possible to reach new populations through various models of online training. However, with a lack of understanding of the clinical research skills that need to be developed, it is not clear whether existing ‘clinical trials’ curricula (whether face-to-face, or online and distance learning) cover the full range of topics to be mastered by global health research staff. In addition, many core roles within clinical research are little understood and still relatively unrecognised as viable career paths. This is the feedback we received during discussions with local partners and when we evaluated training exercises we regularly conduct in LMICs, especially on the role of research nurses, trial managers and data staff.

Furthermore, sustainable capacity building must enable research groups to independently plan, obtain funding for and operate high-quality, locally led studies that address most pressing issues. Tackling those require a whole range of clinical research; from epidemiology, to observational studies and cross-sectional surveys, through to interventional trials. When defining a set of competencies for clinical research, we therefore consider that an integrative approach which aims to capture information about a spectrum of clinical research will be most successful and practicable, as it will best help staff determine how to transfer their advanced research skills to different types of project.

Several groups have worked towards clarifying the competencies for specific clinical research roles, and the ‘Multi Regional Clinical Trials Center (MRCT) Harmonized Core Competency Framework’ makes an excellent move towards combining the information about several roles (principal investigator, clinical research coordinator and clinical research associate) into one common set of knowledge domains. Here, we further this integration effort to many more roles, including essential though less recognised ones in more operational or technical positions (eg, laboratory technician, data entry clerk). We also take particular account of LMIC settings.

CREATING THE FRAMEWORK

To create the Special Programme for Research and Training in Tropical Diseases (TDR) Global Competency Framework for Clinical Research, we looked at the complete set of activities needed for a study to take place, regardless of which individual may perform them. We then examined this information independently of the initial job role it pertained to, to determine the shared competencies (tasks, skills and knowledge) required to support the research endeavour as a whole. The rationale for this approach is that each research project is unique, and the exact distribution of roles and responsibilities within the team are likely to vary depending on the size and demands of the study, as well as on the resources and team available to conduct the trial on the ground. Yet, in any clinical research project there are core, invariant aspects that need to be covered: a protocol must be written, ethics approval must be obtained, data must be collected, verified and analysed, etc. Therefore, the research process and the team undertaking it can be envisioned as a continuum for the definition of cross-cutting ‘competency areas’.

This systematic examination and categorisation of clinical research activities involved qualitative content analysis of 28 past competency frameworks and 116 current job descriptions from partners around the world, altogether covering the following 11 broad research roles: data staff (from entry to managerial roles), laboratory scientist (from technician to head of laboratory), trial pharmacist, community engagement staff, research nurse, study physician, principal investigator, trial manager or project coordinator, quality control monitor, ethics review board member and sponsor. The data-driven, evidence-based framework resulting from this work was then appraised by a panel of 20 expert trialists, and accordingly refined—detailed methods are provided in the TDR Global Competency Framework for Clinical Research development report. In the remainder of this article, we describe the framework (accessible at https://globalhealthtrials.tghn.org/global-competency-framework-clinical-research/), as well as its associated implementation tools; we share the lessons learnt while developing this global competency framework, and we highlight its potential as a mechanism to strengthen clinical research capacity.

CLINICAL RESEARCH ACTIVITIES REVOLVE AROUND 50 COMPETENCIES

After several rounds of iterative, thematic categorisation of the textual data that we gathered from multiple
sources and partners around the world, we found that information from the records included in the analysis crystallised to around 50 distinct competencies, which can be grouped into five balanced categories (figure 1): Ethics, Quality and Risk Management; Study and Site(s) Management; Research Operations; Scientific Thinking; and Professional Skills. The ‘Ethics, Quality and Risk Management’ area encompasses all the competencies related to putting safeguards into place, so as to ensure the risks to participants and of not reaching a scientific conclusion from the research work are kept as low as possible. The ‘Study and Site(s) Management’ area focuses on research management, from project coordination through to finances and human resources management, and also includes the ability to communicate to relevant parties; an essential part of large, multinational clinical research. The ‘Research Operations’ area contains more action-based competencies, and covers most of the practical requirements of interacting with participants, delivering interventions or analysing biological samples to collect data on a daily basis during the study. The ‘Scientific Thinking’ area concentrates on the planning and interpretation aspects of research, which may require various levels of competencies in key disciplines (eg, medicine, statistics, pharmacology, public health), as well as an understanding of the ‘science’ of doing research itself and the methods that are applicable to different research questions. Finally, the ‘Professional Skills’ are generalisable, essential competencies, such as leadership, interpersonal skills or organisational skills.

While the day-to-day tasks and behaviours suggesting competency in the corresponding area will differ between team members and vary according to their specific role, the thorough examination of employers’ requirements alongside relevant literature suggests that the underlying competencies are remarkably similar and cross-cutting.

A competency is defined as the knowledge or skill required to carry an activity out, not the activity itself.20 21 Our framework distinguishes between the following three types of competencies: (1) the theory, or knowledge-based competency, which reflects a level of theoretical understanding of a particular topic, and which can be acquired through learning about a field; (2) the capability, or task-based competency, which necessitates the application of technical or practical knowledge to the performance of a specific task, and which is better learnt by doing and (3) the trait, or skill-based competency, which corresponds to the demonstration of appropriate behaviours in various situations, and which is better developed by experiencing and reflecting on the experience.

SIMILARITY BETWEEN COMPETENCIES ALLOWS FOR A UNIFYING FRAMEWORK

The finalised TDR Global Competency Framework for Clinical Research consists primarily of an encompassing ‘Competency Wheel’ (figure 1), which is applicable to all research roles. It visually presents the framework with its 50 competencies, distributed into the five categories. The core ‘Professional Skills’ category is surrounded by the four thematic, clinical research-specific categories (Ethics, Quality and Risk Management; Study and Site(s) Management; Research Operations; Scientific Thinking); and the latter are each further divided into 3–4 areas of competencies. While figure 1 provides a paper version of the framework, an interactive version allows navigating it more easily and is available online (https://globalhealthtrials.tghn.org/competencywheel/).

A ‘Competency Dictionary’ has been created as a direct companion to the Competency Wheel, and provides detail on what is meant by each of the 50 competencies (figure 2). The Competency Dictionary thus explains the overall, generic framework by suggesting a definition, further clarified with keywords, for each competency. Essential skills and knowledge related to mastering the competency are also listed along with the competencies that are related to it and that appear elsewhere in the framework. Usual abilities associated with the competency are given and directly derived from analysed documents; without those illustrative examples being job specific. The full Competency Dictionary is downloadable at https://globalhealthtrials.tghn.org/site-media/media/medialibrary/2016/11/TDR_Framework_Competency_Dictionary.pdf.

THE FRAMEWORK CAN FACILITATE TRAINING AND CAREER DEVELOPMENT

The core, unifying wheel and the dictionary are supported by practical implementation tools, particularly meant to support assessment and follow-up of an individual’s competencies.

Drawing on previously developed guidance on scoring and evaluating one’s skill level,22 a ‘Competency Radar’ (figure 3) has been developed, and can be used to grade individuals on each of the areas of competency defined within the TDR Global Competency Framework for Clinical Research. Consistent grading should be used as much as possible, in line with the scale developed for the Professional Membership Scheme (PMS),22 and to facilitate comparisons of competency levels, over time and between individuals. In the future, this consistent grading process will be further facilitated through the release of role-specific frameworks mapped to the generic one, to provide more detailed, targeted examples of tasks and expected levels for each role and competency.

Once filled in, the Competency Radar (figure 3) readily shows the areas of strengths of the concerned individual based on their past experience, and outlines which of their skills could be transferred to a different study or setting. It is indeed important to consider one role’s activities within the global picture of competencies, so as to highlight the significance of their...
Design & planning of research
- Health-related knowledge
- Research methodology
- Developing a protocol
- Attracting funding

Protocol operationalization
- Developing study plans and documents
- Developing the QMS and SOPs
- Developing the CRF and DMS

Interpretation of study results
- Analysing data
- Disseminating research findings

Ethics & Quality & Risk management
- Professional skills
  - Cognitive skills
  - Strategic leadership
  - Interpersonal skills
  - Language & communication
  - Organisational skills
  - Record-keeping
  - Computer & IT skills
  - Work ethic

Safeguards
- Ethics and human subject protection
- Risk and safety management
- Determining liability and insurance needs

Quality assurance
- Good Clinical (or other) Practice
- Working as per the QMS
- Controlling quality of research

Regulations & governance
- Securing or maintaining approvals
- Securing or maintaining contracts
- Governance and organisational context
- Research regulations

Study & Site(s) management
- Oversight
  - Initiating study
  - Closing study
  - Project management
  - Tracking study progress
- Study communications
  - Reporting
  - Liaising or acting as a link
  - Facilitating or attending meetings
- Staff management
  - Human resources
  - Creating or delivering training
  - Supervising or mentoring
- Resources management
  - Overseeing essential documents
  - Logistics and facilities management
  - Finances management

Research Operations
- Data flow
  - Creating and maintaining a database
  - Collecting accurate data
  - Data management

Clinical & laboratory operations
- Providing clinical care
  - Ensuring appropriate use of IMPs
  - Handling biomedical products
  - Performing laboratory assays

Interaction with public & participants
- Engaging with the community
- Enrolling and retaining participants
- Supporting and advising participants throughout the informed consent process

Figure 1  The Competency Wheel or one-page overview of the framework. All 50 competencies are visible in this overall representation of the framework. CRF, Case Record Form; DMS, Data Management System; IMPs, Investigational Medical Products; IT, Information Technology (computing); QMS, Quality Management System; SOPs, Standard Operating Procedures.
Figure 2 Annotated sample ‘Competency Dictionary’ entry. The ‘Supervising or mentoring’ competency from the Study and Site(s) Management area of competency is used as an example to illustrate how competencies have been defined. Each of the 50 competencies listed above have its specific entry in the companion dictionary, which can be downloaded at https://globalhealthtrials.tghn.org/site_media/media/medialibrary/2016/11/tdr_framework_competency_dictionary.pdf.

Figure 3 Example of completed ‘Competency Radar’. The grading scheme accompanying the Special Programme for Research and Training in Tropical Diseases (TDR) Global Competency Framework was carefully applied to assess the competency level of a junior researcher in global health. The radar shape readily highlights areas of strengths for this young investigator (who has been involved in designing, planning and interpreting study results), as opposed to major areas where further training and experience should be sought, in line with career plans (eg, the present researcher still lacks confidence in the performance of managerial tasks). The image was generated using the online App for the Competency Radar.
contribution to the study, and their possible career pathways. Past experience of TDR and The Global Health Network suggests there is demand for such a flexible and endorsed system, so as to assist individuals with their career progression. The PMS on which the present grading scheme is derived was initially built by The Global Health Network and TDR as a means of providing a free, standardised system for trialists of all roles to track and demonstrate their progression as they build their careers in research; and it currently counts 2500 members as of February 2017.

As a complementary supporting tool, a ‘Piloting Protocol’ (available at https://globalhealthtrials.tghn.org/site_media/media/medialibrary/2016/11/TDR_Framework_User_Guide.pdf) has been developed to provide further guidance about how to pilot test and use the framework in practice. This protocol also contains a feedback questionnaire, which the early adopters are encouraged to use to let the authors know of their experience using this beta version.

FURTHER DEVELOPMENT TO REINFORCE THE FRAMEWORK

While the framework is designed to evolve as we learn, it is currently being released in a beta version, a necessary step to facilitate its further development and testing. Continuous evaluation is indeed envisioned, so as to iteratively improve and validate the work and tools in line with users’ needs. This learning process should offset the limitations discussed below.

First, job descriptions have been used as an easily accessible means of determining the roles, tasks and ideal attributes of applicants. These provide a list of the activities each individual undertakes, the training basis required for that role, and which key skills or personal attributes are desirable or essential. However, we recognise there could be a discrepancy in what a job description describes and the actual day-to-day tasks undertaken by an individual. This problem is likely to have limited impact on the overall framework’s picture, but we recommend additional research and different methods to further the development of role-specific frameworks. This will be particularly needed for roles where we had limited access to job descriptions, such as the role of the study sponsor, which is critical to clarify to enable locally led research in LMICs.

Second, our framework aims to cover all types of clinical research and thus job descriptions from many diverse locations and trial types were taken into consideration. While this provides the widest possible range of competencies and highlights the connections between apparently different roles, it also means that some of the tasks and roles will not be applicable to every situation. Similarly, we strived to derive broadly applicable and adaptable competency definitions; these may, however, appear too generic in some cases. The generic picture of competencies is nonetheless beneficial to promote career development, as it enables to emphasise bridges between roles and to assess the skills and experience research staff have acquired in a comparable manner. The adoption of this whole-encompassing view is anticipated to lead to better career recognition and capacity building, provided that there is institutional buy-in with creation of adequate training and job opportunities on the ground.

In line with Lucia and Lepsinger’s recommendations for the validation of competency frameworks, the pilot testing will seek to: (1) show face validity, by ensuring that individuals who are currently working in each role understand the competencies suggested, and how they apply to their role; (2) determine whether key tasks have been identified and are clearly covered within one competency or another; (3) determine what the ‘predictors of success’ are for each role, that is, which behaviour demonstrates that the individual has mastered a competency.

Data from this testing phase will first serve to clarify how users interpret the proposed definitions, to address the face-validity question. A feedback questionnaire has been developed, allowing for in-depth comments on the easiness of understanding of different competencies. Where necessary, changes will be made to the framework or the wording of competencies and/or their definition, to improve the comprehensiveness of the framework.

Data from different users of the framework will also enlighten the key skills and tasks that all job roles require for the successful conduct of a study, including but beyond the investigator. We therefore encourage research teams to adopt and pilot test this framework to capture whether competencies are present or missing at the individual and at the larger, site level. Research teams should benefit from using the framework, as their progress can be tracked and demonstrated over time using the proposed tools; hence their efforts can be better directed towards addressing most pressing training or recruitment needs.

CONCLUSION

The Global Health Network and TDR have created a unifying framework which makes use of pre-existing, more role-specific or setting-specific frameworks, as well as of real-life data. The TDR Global Competency Framework for clinical research is broadly applicable and has been successfully appraised by multiple experts; in addition, stakeholders based in LMICs have particularly emphasised the need for such a scheme. In light of this support, the TDR framework has been launched along with its tools to enable refining this resource in the future, based on users’ experience of applying it in context. This framework is laying the ground for future work in the field of clinical research capacity strengthening; and we encourage teams to get involved and use this scheme. Their feedback will be very valuable in refining and widely implementing this valuable new tool for research capacity development.
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