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Abstract

Background: Research-conducive environments are mandatory for planning, implementing and translating research findings into evidence-informed health policies.

Aim: This study aimed at comprehensive situation analysis of health research institutions in the Region.

Methods: We collected data on: institutional characteristics, research scope, capacity building, ethics, governance and resources.

Results: We contacted 575 institutions, of which, 223 (38.8%) responded, indicating that they conducted population research (82%). Reported studies were mostly in medicine, public health and epidemiology, while reported capacity building mainly focused on scientific writing (20.6%), research proposal writing (18%) and quantitative research methods (17%). Most institutions reported having collaborating partners (82%) – predominantly national (77%). Sixty-four percent of institutions received their own funding, with 48% reporting always having access to national databases.

Conclusion: Governments in the Eastern Mediterranean Region and international funding agencies are called upon to support health research production through increasing allocated

support and capacity building in health research.

Keywords: mapping; health research; institutions; Eastern Mediterranean Region

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Introduction

Health research is an integral component in developing health systems, understanding the roots and consequences of poor health, as well as anticipating and mitigating the effects of various factors on health. Promoting and fostering an environment conducive for health research is mandatory for planning, designing and implementing research and for sharing, using and translating its findings into evidence-informed health policies and cost-effective interventions (1).

Mapping studies have been attempted in Africa (2), Canada (3), Europe (4), Latin America and the Caribbean (5). Recent studies in the World Health Organization (WHO) Eastern Mediterranean Region have revealed that the performance of health systems research is weak across all sectors, including governance, finance, workforce, medical and other technology, health information and service delivery. Such studies have referred to critical deficits in stewardship and translation of research into policy and practice, and often absence of an identified research agenda based on emerging priorities (6–8). It should be emphasized that there were prior attempts for health research mapping in the Region, but such studies were not comprehensive and only involved some member states (9–11), or were of institutional (rather than national) scope (10). Thus, to plan effectively for supporting health research to address current priorities and challenges in the Region, comprehensive mapping was deemed essential.

This mapping study aimed to review the scope of existing health research in the Region and identify the challenges that need to be addressed to promote health research further. More specifically, the study aimed to synthesize prior mapping attempts for health research in the Region; to provide a rationale for carrying out comprehensive situation analysis of the range of health research institutions; and to identify possible gaps in the research cycle (research question, objectives, methods, data management, report writing and knowledge translation).

Methods

The study followed a cross-sectional approach over 2 phases: the first was to synthesize prior attempts for mapping health research in the Eastern Mediterranean Region; and the second was to conduct a mapping survey of health research institutions in the Region.

Synthesis of prior mapping exercises in the Region

For synthesis of prior attempts, a literature search was carried out to identify prior mapping attempts for health research in the Region, using databases (Medline, PubMed, Academic Search Complete, WHO Index Medicus for the Eastern Mediterranean Region and Google Scholar); websites [WHO, Council on Health Research and Development (COHRED), Alliance for Health Policy and Systems Research]; and reference lists of relevant studies. Search terms used to search the different databases are found in [Appendix 1](#) ; Medline search strategies for mapping exercises in the Region and globally are found in

[Appendix 2](#) ; and identified studies are listed in Appendix 3.

Health research institutions mapping survey

For the health research institutional mapping survey, a tool was developed to provide a more comprehensive situation analysis of the landscape of health research in the Region at the country level. The tool was a modified version of tools used in previous regional/global mapping exercises (3,6,10–15) and consisted of 6 sections, each comprising several questions on: background information; institutional characteristics; scope of research; training and capacity building; ethics, leadership and governance; and resources (human, financial and technical). In addition, participants were asked to provide narrative remarks reflecting challenges (national, financial, human resources and technical) faced when conducting research within their respective institutions. The tool was pilot-tested to ensure validity and reliability and estimate the time of completion using a guiding protocol for pilot testing. Pilot testing resulted in refinement of some questions.

Focal persons were identified in each of the study countries by the WHO Regional Office for the Eastern Mediterranean to assist data collection. Upon their approval to collaborate, detailed information was provided to them including the overall approach and data collection tool. The responsibilities of the focal person were to assist with: identifying institutions and corresponding focal points in each country; and encouraging institutions to complete the survey tool and submitting online within a specific deadline. The focal persons also validated the final list of institutions after elimination of duplicates and nonresearch centres.

A comprehensive process was developed to establish and validate a database for targeted health research institutions and contact persons. The process went through the following steps: (1) identifying WHO contact lists of health institutions; (2) designing a comprehensive database schema with a unified structure; (3) converting all lists to the unified structure and synthesizing them into the contacts database with a unique identifier and a link to its origin (1415 contacts from 14 sources); (4) reviewing to exclude nonresearch institutions, which resulted in selection of the 575 targeted institutions; (5) validating to exclude duplicates; and (6) sharing the contact list with focal persons for final review and amendments. Data were collected between July 2015 and March 2016.

An automated tool was developed by WHO to send a cover message to each targeted participant/institution, explaining the requirement and communicating the authentication credentials for accessing the online form. Ethical clearance to conduct the study was obtained from the Regional Office. Tool users received continuous communication to ensure clarity of the tasks and to facilitate data collection and reporting to the core team. Survey tools were developed on an interactive web interface using WHO DataCol version 4.4 software.

Data analysis

Data from the survey were analysed using SPSS version 24 for quantitative data. Descriptive analysis including means and standard deviations were used as appropriate. Testing for significant differences across groups was done using the χ^2 test.

Results

Synthesis of prior mapping exercises in the Eastern Mediterranean Region

The search identified 67 health research mapping exercises, of which, 35 were conducted in the Region ([Appendix 3](#)). Upon screening this subgroup, the following gaps were identified. (1) Scope of mapping: mapping addressed some institutional categories (e.g., academic institutions, nonacademic research centres, governmental bodies, international organizations and nongovernmental organizations), focusing on certain parts of the research cycle and support systems. (2) Methodology: among the 35 screened articles, 29 used bibliometric

analyses to assess health research production. Only 6 studies used a cross-sectional method with surveys to map the situation of health research in the Region. (3) Coverage of countries in the Region: the cited studies focused on a subset of Member States in the Region (maximum of 10). No single study has assessed all 22 countries, to gain a comprehensive understanding of the state of health research in the Region. (4) Type of health research: most mapping exercises were topic specific such as mental health research, noncommunicable diseases, nutrition and dietetics. Several articles focused on assessment of biomedical research. Only a few focused on health policy and systems research or public health research.

Outcomes of the synthesis of prior mapping exercises in the Region paved the way and provided a rationale for carrying out comprehensive situation analysis of health research institutions in the Region, identifying possible gaps in the research cycle.

Health research institutions mapping survey

A total of 575 facilities were contacted across 22 countries in the Region, of which, 223 (38.8%) responded to the survey ([Table 1](#)). Respondents' positions were diverse, including directors (27%), deans (15.3%) and chairpersons (12.6%); most of them reported holding a PhD or MD (84.7%), while only 10.5% reported holding an MS or MA.

Institutional characteristics

About half of the respondent centres were academic research institutions (44.7%), mostly (59.8%) established after 2001. Organizational sectors were mostly public (64.2%); noting that 40% of nonacademic research institutions were nongovernmental.

Scope of research

The majority of responding institutions reported conducting population/public health research (84.2%); 76.9% reported conducting social/behavioural research; and 74.6% reported conducting clinical/experimental research. Academic research centres/institutes conducted the majority of research in the areas shown in [Table 2](#) , but the differences from schools and nonacademic centres were only significant for biomedical/basic science research, clinical experimental research and population/public health research.

Training and capacity building

The most commonly reported topics of capacity building sessions were scientific writing/publishing (20.1%), health research proposal writing/funding (18.8%) and quantitative research methods (15.9%). Institutions hosted in high-income countries in the Region reported

issuing calls for proposals annually (37.8%), while institutions hosted within high-to-middle- and low-to-middle-income countries did not report issuing such calls (28.8% and 28.4%, respectively). We used the World Bank classification for countries (<https://datahelpdesk.worldbank.org/knowledgebase/articles/906519-world-bank-country-and-lending-groups>).

Incentives

Institutions were asked about the type of incentives offered to staff to engage in health research. The most reported incentive (50/109) was financial, which included providing grants to researchers, scholarships, awards, monetary incentives and supporting publishing in peer-reviewed journals. The second most reported incentive was linking research activities and publications to academic promotion (25/109). Other incentives reported by respondents included participation in local and regional conferences, collaboration with local and international research centres and organizations, technical support, paid research leave, and training.

Leadership and governance

The majority of institutions reported having an advisory board (65%), especially public and nongovernmental organizations. The types of advisory board members were: general public or service recipients (e.g., citizens, patients and clients) (35/314, 11.2%; 314 exceeds the total number of respondent institutions, as categories are not mutually exclusive); other academic faculties/schools/research institutes (86, 27.4%); policy-makers in government (43, 13.7%); representatives of health care institutions (55, 17.5%); representatives of donor agencies (17, 5.4%); representatives of health professional associations (45, 14.3%); and representatives of nongovernmental organizations (33, 10.5%).

The majority of institutions reported having collaborating partners (81.4%), with national collaborators comprising a majority (70.7%) (regional 66.4%, international, 55.4%). Nonacademic research centres/institutes, nongovernmental organizations and high-income countries were all more likely to have collaborating partners.

Research ethics boards

The majority of institutions reported having an ethics review board (73.5%); mostly reported by high-income countries. The areas represented in the ethics board included medicine (58.8%), health systems (44.4%) and public health (41.3%), in addition to statistics, epidemiology and sociology. The majority of institutions (70.8%) reported having a policy that outlined the structure and function of the ethics review committee, but low-to-middle-income countries were significantly less likely to have such policies (58%). About two thirds (63.7%) of institutions

reported providing training or continuing education to scientists/researchers/new members on research on health ethics. Verification and auditing procedures are outlined in [Table 3](#). Only half (50.9%) the sampled institutions reported having policies for conducting onsite audits of their research ethics committee rules. Only 35% conducted internal/external assessment of the function of the research ethics committee as part of quality improvement.

Human, technical and financial resources

Gender distribution of researchers was reported to be almost the same across participating institutions (i.e., almost equal numbers of men and women), but the average number of PhD holders was higher than those with an MS, BS/BA or MD. International institutions were found to have a significantly higher number of MS degree holders than public institutions had.

Health research funding sources were: own institution (145/458, 31.7%; 458 exceeds the total number of respondent institutions, as categories are not mutually exclusive); local public institution (89, 19.4%); regional public institution (49, 10.6%); WHO (47, 10.3%); international aid agency (36, 7.9%); local private institution (35, 7.6%); pharmaceutical company (34, 7.4%); other United Nations agency (32, 7.0%); and regional private institution (25, 5.5%).

The majority of institutions reported always having computers (84.5%), telephones (78.8%), internet connections (76.1%), printers (75.7%) and scanners (67.7%) at the disposal of their researchers. Only 45.6% reported always having access to national databases, while 32.3% reported always having access to international databases. Participant institutions reported providing IT support (73.5%), staff for research on health (65.9%), research management (57.1%), and health proposal writing services (52.2%) ([Table 4](#)). Respondent institutions reported that their research laboratories were predominantly clinical (41.0%) or biotechnology (36.2%) laboratories.

Challenges facing institutions conducting health research (qualitative feedback)

At the national level, some respondents (12/51) indicated that lack of awareness of policy-makers of the importance of research is one of the challenges facing health research conduct and utilization of its findings. Respondents also reported a lack of national strategy for health research, and a lack of national policies and regulations that govern the conduct of health research. The need to set national research priorities and raise awareness about the importance of research and evidence-informed policy-making was also reported. The limited availability and access to data, mainly governmental data, was another challenge raised by respondents. Other challenges reported included limited national funding and mismatch between funding and national priorities.

At the financial level, most respondents (61/89) reported limited funds available for health research, including limited internal funding and lack of grant management systems. Other reported financial challenges included mismatch between funding opportunities and priorities, sustainability of funding, and competitive aspects of international funding.

At the human resources level, almost half the respondents (33/69) reported a shortage of and challenges in recruitment of health researchers, as well as difficulty in retaining qualified researchers, research assistants and associates (i.e., brain drain). Respondents raised the need to build capacities and skills of health researchers, especially on research methods, writing proposals, complementary and integrative medicine research, and health policy and systems research.

At the technical level, the challenges mostly reported

by respondents included: limited availability of laboratories facilities, equipment and devices; limited skills and capacities of the staff; and lack of collaboration between the research institutions and other national, regional and international institutions. Sanctions were also reported to challenge some institutions from adopting new technologies, buying equipment and accessing websites.

Discussion

This mapping exercise provides an overview of the scope of existing health research cycles and support systems in the Eastern Mediterranean Region. We found that most institutions reported undertaking public health research followed by clinical research, and fewer institutions reported undertaking health policy and systems research. This is in line with a previous mapping exercise that showed the low production of health policy and systems research in the Region (8). Most institutions reported having collaborating partners (82%), predominantly national (77%), and to a lesser extent international collaborators (55.4%). Most participating institutions were public and received internal funding (64%).

We identified several challenges that undermine health research in the Region. Both quantitative and qualitative data reflected limited national, regional and international funding to institutions conducting health research. This indicates that investment in health research is still low in the Region. Limitations in national funding can be attributed to multiple reasons. National governments and policy-makers, especially in high- and upper-middle-income countries, might still be allocating limited budgets for health research (6). Policy-makers in these countries may be also unaware of the importance of health research in improving health systems and public

health, as reported by some respondents. These countries might also be lacking national strategies and policies that govern health research and define national health research priorities as reported by several respondents. One can also speculate that policy-makers are rarely engaged in health research as only 29.3% of institutions reported having policy-makers on their advisory board. This finding is in line with another mapping exercise conducted in the Region that showed that only 3 of 10 surveyed countries reported setting national health research priorities, and only 2 countries had a dedicated national health research policy (11). Another study from the Region also revealed the poor engagement of policy-makers in health research, with only 16% of researchers interacting with policy-makers and stakeholders in priority setting, and only 20% involved them in their research (16). Another reason for limited national funding might be the financial inability of the country to provide a high budget for health research, especially in low- and middle-income countries. The limited international funding can be explained by the weak institutional capacity to attract funds, in terms of grant management, writing proposals, access to international databases and shortage of skilled human resources, as the findings of the present study show. Some of the reported reasons for this shortage were the difficulty in recruiting and retaining qualified researchers, mainly research assistants and associates, and the brain drain caused by emigration.

The present study reflected the findings from the first comprehensive survey involving all Member States of the Region. The survey also targeted all the health research institutional categories (public, private, academic and nonacademic) and all types of health research (from biomedical to health systems research). It utilized a thorough pilot-tested survey that was based on research evidence and comprised different sections that covered different aspects of health research cycle and support systems.

Although the study had a high target for sampling facilities, the response rate was 38.8%. Despite this, the results provide insight into different aspects of health research. In some countries, focal people were more successful in terms of access to respondents and responsiveness to complete the survey itself. In some instances, questions required respondents to report on their personal perceptions; these questions may have been subject to social desirability responses. Additionally, we were unable to consider the context for understanding the situation of health research institutions, due to the cross-sectional nature of the study and random response of respondent institutions. Hence, the outcomes of the survey should be assessed with caution, in spite of the wide spectrum of respondent institutions from different countries in the Region. We believe that after sharing results with Member States, more health research institutions will come forward and respond to the survey, providing a better spectrum of the health research cycle in the Region.

Conclusion and recommendations

Governments in the Region and international funding agencies are called upon to support

health research through increasing support and capacity building in health research. Health research institutions in the Region should work on developing strategies to retain qualified researchers and building capacities of existing ones. The institutions are also called upon to strengthen their capacities, including grant management systems and access to international databases. This will increase their ability to attract international funding and increase research productivity. As the study showed limited international collaboration, institutions should recognize the importance of such collaboration in advancing and exchanging knowledge and strengthening research capacity and expertise (17).

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Cartographie institutionnelle de la recherche en santé : perspective régionale pour la Méditerranée orientale

Résumé

Contexte : Des environnements favorables à la recherche sont nécessaires pour planifier, mettre en œuvre et traduire les résultats de recherche en politiques de santé reposant sur des bases factuelles.

Objectif : La présente étude visait à mener une analyse de situation des établissements de recherche en santé dans la Région.

Méthodes : Nous avons recueilli des données sur les caractéristiques institutionnelles, le champ des recherches, le renforcement des capacités, l'éthique, la gouvernance et les ressources.

Résultats : Nous avons contacté 575 établissements, sur lesquels 223 (38,8 %) ont répondu qu'ils avaient effectué des recherches en population (82 %). Les études traitées concernaient principalement la médecine, la santé publique et l'épidémiologie ; le renforcement des capacités signalé portait principalement sur l'écriture scientifique (20,6 %), la rédaction des propositions de recherche (18 %) et les méthodes de recherche quantitative (17 %). La plupart des établissements signalaient avoir des partenaires de collaboration (82 %), principalement au niveau national (77 %). Soixante-quatre pour cent des établissements recevaient leur propre financement, et 48 % mentionnaient avoir toujours accès aux bases de données nationales.

Conclusion : Les gouvernements des pays de la Région de la Méditerranée orientale et les bailleurs de fonds internationaux sont appelés à soutenir la production de la recherche en santé en augmentant le soutien et le renforcement des capacités alloués en matière de recherche en santé.

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