ABSTRACT Routine information systems for mental health in many Eastern Mediterranean Region countries are rudimentary or absent, making it difficult to understand the needs of local populations and to plan accordingly. Key components for mental health surveillance and information systems are: national commitment and leadership to ensure that relevant high quality information is collected and reported; a minimum data set of key mental health indicators; intersectoral collaboration with appropriate data sharing; routine data collection supplemented with periodic surveys; quality control and confidentiality; and technology and skills to support data collection, sharing and dissemination. Priority strategic interventions include: (1) periodically assessing and reporting the mental health resources and capacities available using standardized methodologies; (2) routine collection of information and reporting on service availability, coverage and continuity, for priority mental disorders disaggregated by...
Systèmes d'information et de surveillances de la santé mentale

RÉSUMÉ Les systèmes d'information de routine pour la santé mentale dans de nombreux pays de la Région de la Méditerranée orientale sont rudimentaires ou font défaut, ce qui rend difficile la compréhension des besoins des populations locales et la planification correspondante. Les composantes clés des systèmes d'information et de surveillance de la santé mentale sont les suivantes : un engagement et un rôle de premier plan à l'échelle nationale pour garantir que des données pertinentes et de haute qualité sont recueillies et transmises ; un ensemble de données minimales servant d'indicateurs clés pour la santé mentale ; une collaboration intersectorielle permettant le partage approprié des informations ; le recueil de données systématique complété par des enquêtes périodiques ; un contrôle qualité et la confidentialité ; et de la technologie et des compétences pour appuyer le recueil, le partage et la diffusion des données. Parmi les interventions stratégiques prioritaires, on peut citer : 1) l'évaluation périodique des ressources et des capacités en santé mentale disponibles et la notification de ces informations à l'aide de méthodologies normalisées ; 2) le recueil et la notification de données systématiques sur la disponibilité des services, leur couverture et leur pérennité pour les troubles de santé mentale prioritaires, ventilées par âge, sexe et diagnostic ; et 3) l'enregistrement et la notification obligatoires des suicides à l'échelle nationale (à l'aide des codes CIM pertinents).

1Institute of Brain, Behaviour and Mental Health, University of Manchester, and Lancashire Care NHS Foundation Trust, United Kingdom (Correspondence to R. Gater: richard.gater@manchester.ac.uk). 2Department of Mental Health and Substance Abuse, World Health Organization, Geneva, Switzerland. 3Department of Psychological Sciences, University of Liverpool, Liverpool, United Kingdom.

The case for mental health surveillance & information systems

Surveillance involves the collection, analysis and interpretation of health data and the timely communication of these data to policy-makers and others. The availability of relevant information enables actions to be monitored and improvements in service provision to be detected. Mental health information systems are vital for collecting, processing and analysing information about mental health determinants, needs, system responses and the impact of interventions. But it is also crucial that findings are communicated in a form that is accessible and useful to those who will utilize them. Only then can the mental health information system perform its functions of facilitating effective planning, budgeting, delivery and evaluation of
mental health care. This information loop from data collection, through analysis and reporting to informed implementation of plans (Figure 1), needs to be driven by an infrastructure of training and supervision of all staff involved, of quality assurance and of confidentiality. These activities require clear leadership to oversee and manage the process in its entirety.

Preliminary findings from the World Health Organization (WHO) Mental health atlas survey 2014 [in press] shows that more than one-third of Eastern Mediterranean Region (EMR) countries have not published a specific mental health information report in the past 2 years. Approximately half were unable to provide any financial information and less than one-sixth knew their total expenditure for mental health. About two-thirds of countries in the Region did not know the total number of staff in the mental health workforce. Although more than half could report the number of persons treated at mental hospitals, the great majority of countries were unable to report the number of persons with mental disorders who received care in mental
health outpatients departments or in primary care facilities. About half of EMR countries had data on length of stay and involuntary admissions, but few reported on the proportion of persons discharged from hospital who had a follow-up visit within one month. Only one-quarter of EMR countries were able to report data on numbers of suicides. Suicide rates in EMR countries may be under-reported or even unreported for social, religious and cultural reasons. The difficulty in providing information for the Mental health atlas, particularly on expenditure on mental health care, number of professionals working in different settings, mental health service coverage and suicide data, suggests that many countries are managing with very rudimentary information systems, making it difficult for them to understand the needs of local populations and to plan accordingly.

**What information is needed for mental health policy, planning and evaluation?**

Collecting a small number of carefully selected indicators thoroughly and consistently over time (both within and across countries) is more effective than collecting a large number of indicators that are never implemented. The collected indicators should be meaningful to health planners, acceptable to stakeholders, valid, reliable, comparable over time and sensitive to change. They need to be disaggregated by sex and age and by other variables, in order to capture the diverse needs of subpopulations, including individuals from geographically diverse communities (for instance, urban versus rural) and vulnerable populations.

Three WHO sources—the Comprehensive Mental Health Action Plan 2013–20 (1); the EMR Regional Framework (2); and the WHO Mental Health Gap Action Programme (mhGAP) monitoring and evaluation tool kit (3)—can be used to identify a minimum data set for mental health. A set of indicators assembled from the Comprehensive Mental Health Action Plan and the EMR Regional Framework are included in Table 1.

**How can information for mental health be generated?**

Data relating to internationally agreed as well as locally determined mental health indicators can be collected routinely or periodically. Ideally, most of the data requirements should be generated on a routine basis via local information systems; for example, deaths attributable to suicide and self-harm should be recorded in vital registration systems, while cases of mental disorder receiving care and treatment should be identifiable through facility-based recording systems (see Box 1 for an example from Saudi Arabia).

In situations where routine health information systems may not yet be in place or functioning well, or where more periodic assessment may be sufficient (e.g. the compliance of local mental
health legislation with international or regional human rights instruments), periodic but regular surveys can be used to monitor developments. For example, in order to measure current and increased service coverage for severe mental disorders—a core mental health indicator of the global Action Plan—many countries may consider carrying out a baseline and repeat survey of provider facilities in one or more defined geographical areas of the country.

Table 2 provides examples of expected data collection strategies and sources for a number of key mental health indicators.

**How can information be used?**

The information loop is completed when the information is presented in a meaningful way and it is used to inform service planning. In 2010 the regional report based on the WHO Assessment Instrument for Mental Health Systems (AIMS) found that 71% of countries had a formally defined list of mental health data items to collect, and 65% of countries had published the data; however, only 30% published the data with comments (5). In other words, although information was being collected, it was seldom analysed so that it could be used as a tool for action.

Traditionally, reporting has been in the form of printed statistical tables with a commentary, but Internet-based technology now offers the opportunity for information integrated from different sources to be disseminated rapidly to end-users in a relevant and interactive format at a local, national, regional or global level. The WHO Regional Office for the Eastern Mediterranean (EMRO) is currently engaged in developing a regional National Health Information Systems (NHIS) strategy and a set of core health indicators which countries should report to EMRO on a regular basis. The strategy anticipates that NHIS will be required to move to systems that are deployed on the Internet, and by design are integrated based on principles of data warehousing. This will enable the access of data from different sources, and facilitate circulation of accessible data.

Whatever reporting system is in place, it is important that policy-makers and service planners have the skills to interpret and apply the evidence from information systems; and that reporting is part of an ongoing dialogue with policy-makers and service planners, to ensure that information is relevant and presented in a useful format.

**Key recommendations**

1. Periodically assess and report the mental health resources and capacities available using
standardized methodologies.

Establish a national focus of expertise and leadership to implement the development, reporting and use of mental health surveillance and information involving collaboration between relevant service providers working in the field of mental health: ministries of health, university hospitals, psychiatric associations, private hospitals, nongovernmental organizations and key opinion leaders.

Develop procedures, regulations and training to ensure that the processes of collecting, analysing, reporting and using data meet standards of quality and confidentiality.

2. Routinely collect information and report on service availability, coverage and continuity, for priority mental disorders disaggregated by age, sex and diagnosis.

Develop or strengthen national mental health information systems incorporating the indicators in Table 1.

Collaborate with the EMR regional strategy for NHIS strengthening to develop information systems that utilize web-based technologies and data warehousing to facilitate the integration of information. These will enable the access of data from different sectors and settings, and facilitate circulation of accessible data.

3. Record and report on deaths as a result of suicide at the national level (using the relevant International classification of diseases (ICD) X-codes).

References


