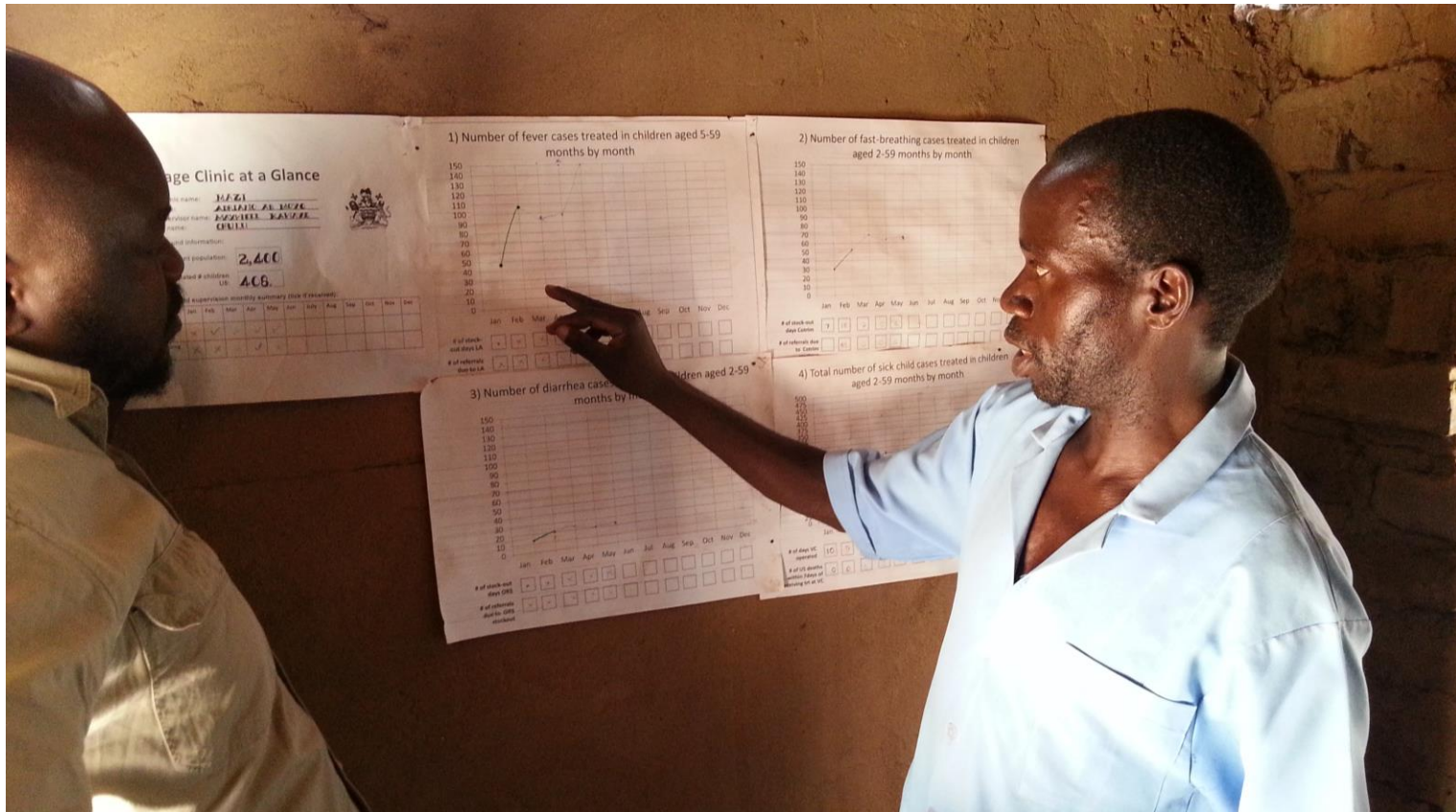


Special considerations for community level data



Improving Routine Data for Child Health in National Health Information Systems: Africa Regional Workshop

September 21, 2017

Jeanne Koepsell, Tanya Guenther, Gail Snetro-Plewman

Overview

- Importance of routine community level data
- Characteristics of community data
- Design principles for CHIS (#1-4)
- Community level data quality & use

Routine monitoring systems for integrated community case management programs: Lessons from 18 countries in sub-Saharan Africa

Tanya Guenther¹, Yolanda Barberi Lainez², Nicholas P. Olliphant³, Martin Dale⁴, Serge Baharison⁵, Laura Miller⁶, Geoffrey Namara⁷, Theresa Diaz⁸

dhis2

DHIS2 Community Health Information System Guidelines



empowered to use data for decision-making [8]. Monitoring systems for many of the same shortcomings of the health information systems (HIS), but at the community level at scale present a challenge. While the literature of ICCM programs has expanded, little that explores the monitoring system's successful implementation.

Key lessons learned from monitoring and implementing systems for routine scale ICCM programs. These lessons from the primary partners supporting us across 18 countries in sub-Saharan Africa with monitoring local persons at documents and tools and informed regulatory routine health information by [1-5]. We first outline the rationale for the challenges ICCM programs face in monitoring systems to generate well-timed the current state of routine monitoring, summarize lessons learned and forward.

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Why is community-level data important?

- Children require health services on an on-going basis
- Surveys occur periodically and do not provide community-level estimates
- Routine data at community level:
 - ❖ Facilitates community ownership and increases demand for quality, responsive health care
 - ❖ Allows tailoring of services to greatest needs
 - ❖ Strengthens linkages to facility

What is community-level data?*

- data on health, health services, and determinants of health
- derived from and relevant to:
 - community admin/geo units
 - community members
 - community level of health systems

*Source: DHIS2 community health information system guidelines, 2017.

https://www.healthdatacollaborative.org/fileadmin/uploads/hdc/Documents/Working_Groups/CHISGuidelines_version_August29.pdf



Unique characteristics of community data

Area	Examples
Scale	<ul style="list-style-type: none">• Thousands of CHWs and communities compared to typically hundreds of health facilities
Complexity	<ul style="list-style-type: none">• Wider range of programs (health promotion, etc)• Different services and definitions• Variable linkages with formal health system
Diversity	<ul style="list-style-type: none">• CHWs range from well educated to illiterate• Communities may have different CHWs for different programs; programs vary by area
Stakeholders	<ul style="list-style-type: none">• Wider group of stakeholders interested in community-level data – has implications for data management, protection, etc
Infrastructure	<ul style="list-style-type: none">• Tends to be weaker the further out you go – less electricity, limited to no internet, poorer roads – all affecting ability to collect and report data

Design principles for CHIS

1. Design the CHIS to strengthen **government ownership** and sustainability.
2. Enable and **strengthen community engagement**.
3. Build a **balance** between reporting burden and provision of care.
4. Strengthen **capacity** of CHWs and other stakeholders as a team.
5. *Follow **incremental** and evolutionary principles of system design and development.*



CHIS Design Principle 1: Government ownership

- Government leadership and coordination essential to promote:

- Rationalization
- Harmonization
- Sustainability
- Integration with HMIS
- Pooling of resources

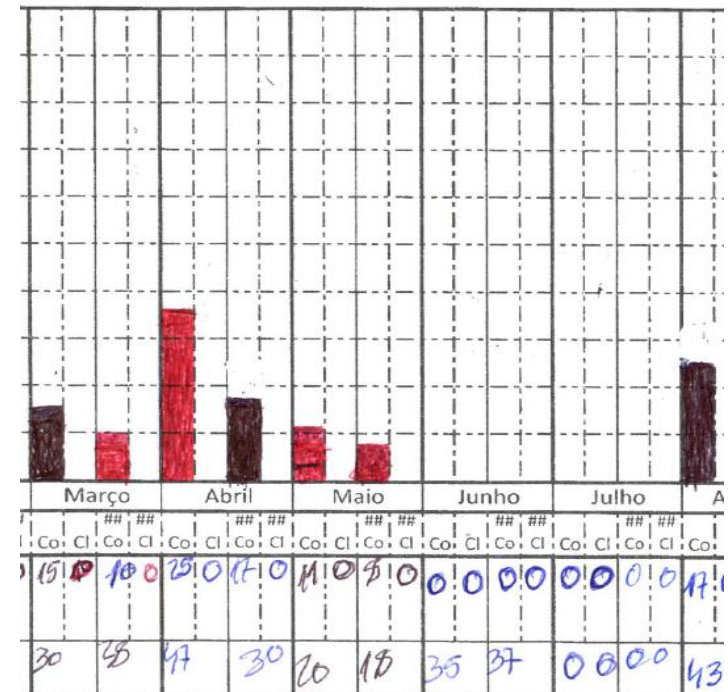
Example: Mozambique data use training

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CHIS Design Principle 2: Community engagement

- Central to data quality and supporting community action based on data
- Requires involvement of range of community-level stakeholders
- Must occur early and continue across the design → implementation phase
- Design process should aim for **simplicity, efficiency** and **scalability** and be suitable for the lowest capacity levels
- Identify and promote local innovations



CHIS Design Principle 3: Balance

- CHWs often bear the heaviest burden for data collection & reporting
- Overly detailed documentation requirements counterproductive
- Prioritize data elements that are:
 - Linked to program performance (e.g. treatment coverage)
 - Tied to specific actions
 - Feasible to collect well



Photo: Courtesy of Yolanda Barbera Lainez, International Rescue Committee



CHIS Design Principle 4: Strengthen team capacity

- Allocate resources to strengthen capacity of teams (CHWs, facility staff, other stakeholders)
- Build in basic concepts of data elements, indicators, coverage, etc
- Provide simple tools and resources
- Promote appropriate use of mobile and other technologies
- Ensure sustained support and supervision



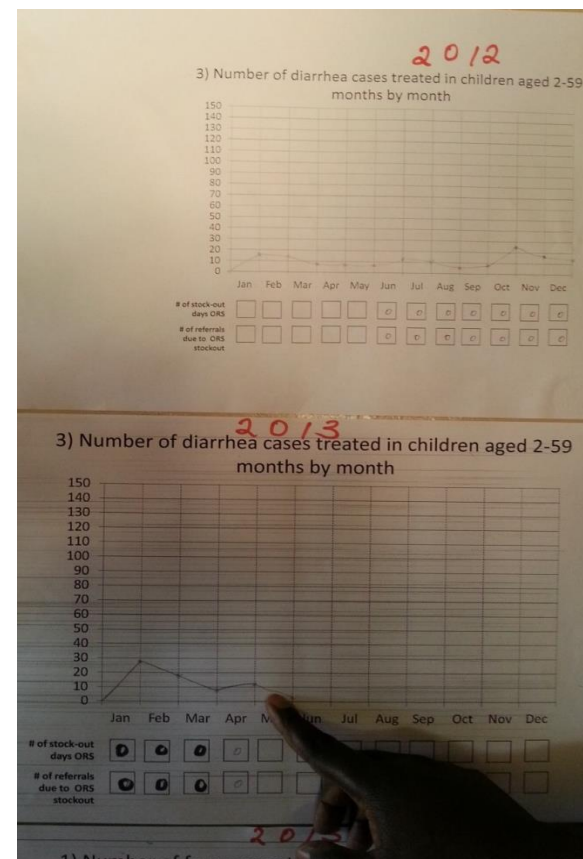
Improving data quality and use by CHWs

Data Quality and Use Intervention Package for iCCM in Malawi*

- Challenge: iCCM data were being compiled but not used at the health facility or HSA levels
- Intervention: Flexible data quality and use improvement package for HSAs and facility staff including: basic data management, analysis and use training; provision of wall charts for data display with action thresholds
- Evaluation results: Provision of wall charts to organize and view monthly iCCM reporting data, along with additional training, leads to more data-based decision making
- Package scaled up across Malawi through MOH and partners

Paper available soon at: <https://doi.org/10.9745/GHSP-D-17-00103>

"Now everyone can see the data"



Community *Use of Data* for Decision-Making

- Communities are the source of data and have a fundamental citizen right to use data for decision-making
- Engaging communities in data collection & use helps:
 - Improve health indicators
 - Leverage local resources to support health programs
 - Increase women's and family participation in community decision-making and action.
 - Strengthen community capacity to act and monitor their progress
 - Increases self-reliance

CH BULLETIN BOARD



Community *Use of Data* for Decision-Making

- Captures key community health data before being *extracted up* the health system
- Makes data understandable to all through use of picture cards and Community Health Bulletin Boards
- Provides feedback back to the community on local health status
- Provides a tool for existing community health groups, such as Community Health Committees; women's groups to engage the broader community in action for health

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2		37	18	41	24	120
3		3	1	2	2	8
4		37	18	41	24	120

Community *Use* of Data: How Does It Work?

- Health promoters collect monthly data on key indicators from families
- Service providers collect service utilization data
- Together they consolidate data at the end of the month.
- Teams use simple tools to share the data with the community.
- Community members review and analyze the information.
- Participants then set priorities and develop plans to improve their priority health indicators.



THANK YOU



Save the Children