WHO Informal Consultation on fever management in peripheral health care settings: a global review of evidence and practice

Global Malaria Programme, WHO



Section I - Review on etiologies and management of febrile illness

What are we trying to do?

- Intended aim need to be clear:
 - Reduce antibiotic prescription / drug resistance
 - Increase appropriate treatment
 - Reduce severe disease
 - Save money

Etiologies

• Common findings of studies on etiologies so far:

Children<5 years: ¹/₂ ARI, 1/10 to ¹/₄ diarrhoea, rest unspecific fever, UTI always low, typhoid low in Africa, high in Asia

Adults: driven by HIV (40% even in low prev area) more vector-born, live-stock, outdoor (lepto, rickettsia, typhus...)

- Low specificity of RR for pneumonia in underfive confirmed \rightarrow viral etiology
- As we go away from `gold standard diagnosis' towards clinical outcome, `Treatment failure' need to defined (e.g pneumonia)



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Epidemiology

- Good estimates of incidence or prevalence <u>only if</u> clinical data or asympt. group associated to 'crude' laboratory data (biased pop.)
- Serology lack specificity and PCR is too sensitive \rightarrow high pos. rate in asympt group
- Severe disease is very rare at peripheral level, especially community (true?)

The way forward

- No need to repeat extensive etiology studies
 - \rightarrow use simplified design
 - \rightarrow build on existing networks (GEMS, PERCH, TSAP...)
 - \rightarrow at different levels: community / outpatients / admissions
 - \rightarrow in different age groups: underfive, <u>5-15y children</u>, adults

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- Methodology:
 - \rightarrow target <u>unspecific</u> fevers in different areas
 - \rightarrow is asymptomatic control group always necessary?
 - \rightarrow common definitions for diseases

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The way forward

Analytical Considerations

Possible/useful to develop a 'standard' framework for data analysis

- Descriptive epidemiology
- Risk factors for disease progression, severe illness, drug resistance
- Risk factors for treatment with an antibiotic
- Effects of recommending specific treatment (eg doxycycline)
- Modelling to inform target product profiles of new diagnostics
- Disease severity vs pathogen-specific
- Respiratory rate counters, pulse oximetry
- Target sens/spec
- Algorithm design (eg ALMANACH)
- Formulating algorithms
 - \rightarrow etiologies

 \rightarrow other factors (distance to HF, economical stautus, ease of referral..)

- \rightarrow continuum of care
- \rightarrow potential of electronic guides for compliance and data collection

Tools available

Children Adults

Hospital Blue book District manual

Health facility IMCI IMAI

Community (& informal private) iCCM ?

- No guidelines for adults in community
- No guidelines for children 5 to 15 years
- Algorithm for malaria diagnosis&treatment well integrated in most of guidelines
- Home Based Malaria (2002-2005) should be put in archives
- Several points in need for update:
 - Criteria for high and low malaria risk area
 - Testing of anemic children in high malaria risk
 - Testing before referral/pre-referral treatment
 - Time interval new malaria infection (>14 days)
- IMCI & IMAI should be widely disseminated

 \rightarrow no more malaria diagnosis&treatment without IMCI/iCCM

• Adherence to iCCM OK, to IMCI problematic \rightarrow find new strategies for HFs

Algorithms available

- Up to which degree of place and time should algorithms be refined?
 - → need to go below → possible to have them different algorithms according to season?
 - → Probably rather by level of health system (keep it simple for the community level)
- To keep in mind: HWs are trained and leave, trained and leave again...
- Algorithms for typhoid (and Dengue) in high endemic areas are urgently needed
- Carefully evaluate each new test for cost/benefit before adding it (e.g Dengue)

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- IMCI booklets have already become too heavy
- IMAI: How to cope with long list of diseases in the fever branch?

New diagnostic tests

- More specific, more expensive (clinical \rightarrow epidemio \rightarrow severity test \rightarrow pathogen test)
- POCTs already available, some usable as they are (Dengue) other not (Typhoid)
- New POCTs in development
 - ightarrow to specifically detect one pathogen
 - → to 'generically' identify: patients at risk for progression to severe dis.
 patients in need for antibiotic
- Electronic tools to measure essential clinical parameters (RR, O₂ Sat, temp.)



Essential medicines

• High level of bacterial resistance to first line treatments:

- \rightarrow How to quickly adapt guidelines to these changes?
- \rightarrow How to replace cotrimoxazole by amoxicillin for ARI (dispersible)
- Should also think in terms of 'class of antibiotics' (not only yes/no)
- No evidence to split the list by level of health care \rightarrow responsibility of countries
- No injectables in the list for community level (pre-referral antibiotic???)



Section III - Agencies and NGOs experience with iCCM

iCCM task force

- Specific tasks: develop tools (training packages, job aids...)
 - set up supply chain management
 - M&E
 - operational research
 - policy & advocacy
 - country support (difficult)
 - \rightarrow based on lessons learned, new manual to guide countries
- Extension to newborns considered, but not to school-children or adults

Challenges to the scale-up (multi-countries review):

- Retention of CHWs in the context of limited HR:
- Supervision of CHWs: more experience peer rather than clinician of HF
- Severe drug shortages: necessity of introducing parallel system \rightarrow not sustainable
- Care seeking behaviour: communities need to know what care they can expect
- Weak M&E: innovative technologies (basic phones are enough)

Section III - Agencies and NGOs experience with iCCM

Results of operational research

- \downarrow mortality with AM (ongoing studies will tell us for AB)
- High compliance with lab-test, low compliance with clinical-test (RR)
- CHWs not good to pick up danger signs (rarely seen)
- Do not refer (Why? Know that patients will not comply?)
- \uparrow utilisation of CHWs, but still below expected incidence of diseases
- Very difficult for CHWs to identify danger signs in newborns
- How to measure quality of care: DO without reexam, registers, scenarios not enough for RR and danger signs
- Access should take into account other factors than geograph. Distance
- More and more salaries \rightarrow helps for retention of HWs
- Feeling of managers: should remain a limited mandate (regulatory problems)

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Costs: much cheaper to manage sev. pneumonia at community level

Section IV - Experiences of community case management of fevers

Public sector

- Rollout of iCCM in different countries with different adaptation of algorithm, training, supervision, data collection/reporting and remuneration/motivation approaches
- Quantification challenges for RDTs and for different medicines (antimalarials, antibiotics and ORS) due to different prevalence of the 3 diseases in different parts of the country
- Supply chain challenges addressed in different ways: in the future need to integrate the current parallel distribution system with the main drug supply system managed by central medical store
- Issues with services at community level outperforming health facilities, and need to review package of services at referral level
- Need to clarify role of amoxicillin for pre-referral treatment of severe pneumonia at community level
- Simplified algorithm required, focusing on malaria, pneumonia and diarrhoea, with focus on danger signs requiring referral

IMCI - Caring for the sick child in the community



Section IV - Experiences of community case management of fevers

Private sector

- Need to be addressed (important source of care in many, not all, settings)
- Factors: source of care, skills levels, disease etiology, coverage with public sector facilities/agents (CHWs etc)
- Not uniform, needs to be segmented (e.g. drug peddlers, retail shops, non registered drug shops, registered drug shops, private clinics (by level), not-for-profits etc)

 \rightarrow for strategizing research, review and interventions

• Different approaches for different segments

→ e.g: positive incentives (knowledge/training, profits, social marketing, organization into societies etc)

 \rightarrow to come up with an appropriate "mix" (in each context, segment)

• Do not introduce malaria RDTs alone (e.g. blanket advise for referring RDT-

→TOGETHER WITH (diagnostics &) treatment for common conditions in that context (e.g. RR timers and (prepackaged) antibiotics; ORS+Zinc)

• Supervision, Quality Measurement and Quality Assurance of care and products:

 \rightarrow Methods and mechanisms need to be elaborated and evaluated

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Private sector

Surveillance methods need to be elaborated

 e.g. for RDT positivity rate, conditions treated, drug use
 → to integrate information into health management systems

Empower Demand side (knowledge and purchasing/entitlement-enabled consumers)

→ important factor in improving care-seeking and quality of care
 → e.g: "branding" or social franchising drug shops/clinics/individuals
 but ALSO criteria of good quality care (e.g. child examined,
 diagnostic test applied, treatment upon result).

- Need for futhur understanding of:
 - microeconomics of running private sector outlets,
 - construction of (financial) incentive mechanisms that promote desired behaviours (such as profit margins from testing, different treatment combinations etc)