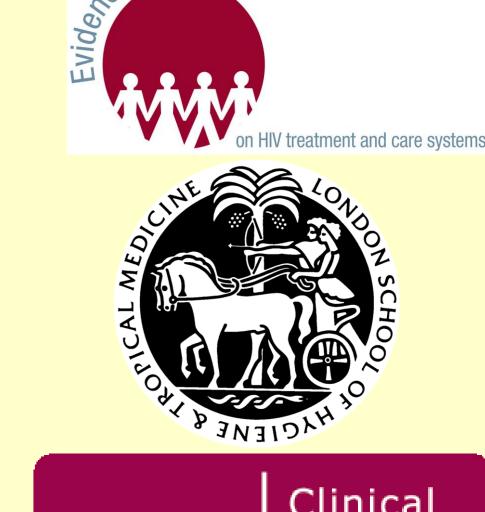
Poster number: MOPE0999

Getting research into policy for Cotrimoxazole prophylaxis for HIV related infection: a comparative policy analysis in Malawi, Uganda, and Zambia

E. Hutchinson¹, J. Parkhurst¹, N. Chishinga², B. Droti³, S. Phiri⁴, S. Hoskins⁵, D.M. Gibb⁵

1 London School of Hygiene and Tropical Medicine, 2 Zambia AIDS Related TB Project, Zambia 3 Medical Research Council - Uganda Virus Research Institute, 4 Lighthouse Trust, Malawi, 5 Medical Research Council Clinical Trials Unit, UK.





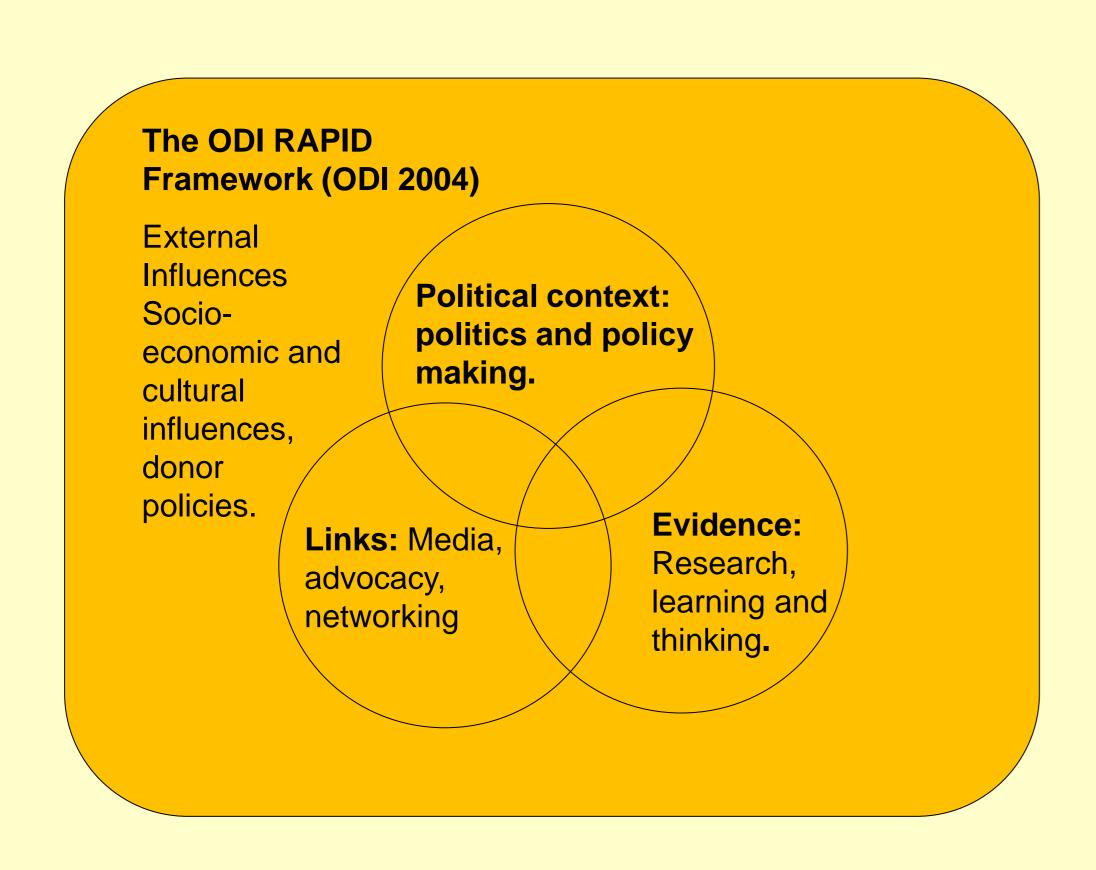
Background

- Cotrimoxazole prophylaxis (CPT) prevents opportunistic infections, saves lives, and is cost-effective.
- 3 medical journals have questioned why CPT has not been rapidly scaled up across Africa, expressing frustration that it is not more widely available.
- Policy analysis approaches are regularly used to understand local processes and structures to explain the take up of evidence into policy.

To understand the variation in timing and policy content in the development of CPT national policy processes for CPT across Malawi, Uganda and Zambia, with a particular focus on the ways in which national context impacts on health policy making.

Study methods:

- Comparative policy analysis of research to policy processes in Malawi, Uganda and Zambia.
- Case study approach using the Overseas Development Institute's RAPID research framework.
- Published and unpublished documentation supplemented by interviews with 47 key informants across all three countries.



CPT key research and policy timeline

Cote d'Ivoire RCTs CPT effective in HIV-1 infected adults in area of low resistance to cotrimoxazole (CTX) (Anglaret et al 1999,

S

Wiktor et al 1999).

Malawi observational study. CPT is feasible, safe and reduces mortality among HIV positive TB patients in area of high resistance to

CTX (Zacharia et al).

Zambia RCT CPT reduces mortality and morbidity among HIV infected children in area of high resistance to CTX (Chintu et al)

2004

WHO/UNAIDS/

UNICEF joint

advocates use of

exposed and HIV

infected children.

CPT for all HIV

statement

Malawi cohort

patients is

study. HTC and CPT

associated with good

for HIV positive TB

well accepted and

treatment outcome

Malawi descriptive

routinely implement

VCT and CTX for TB

patients (Chimzizi et

observational study

positive TB patients

bacterial resistance

(Mwaungulu et al)

observational study.

morbidity among HIV

infected adults in

resistance to CTX

Malawi

Uganda

CPT reduces

mortality and

area of high

(Mermin et al)

CPT reduces

mortality in HIV

in area of high

study. It is feasible to

(Chimzizi et al).

Malawi RCT. No significant differences in terms of efficacy between two different doses of CPT in HIV positive TB patients (Boeree et al).

Malawi observational study showing Incidence of Trimethoprim-Sulfamethoxazole-Preventable Infections in Malawian Adults Living With HIV (van Oosterhout et al)

Uganda observational study. CPT reduces mortality and morbidity among HIV infected adults in area of high resistance to CTX (Watera et al)

1999

2000

WHO, UNAIDS

2001

provisional recommendations that CPT be scaled up across Africa. Malawi, Uganda and Zambia reject scale-up.

2002

Malawi policy to provide CPT for HIV+ tuberculosis (TB)

patients is agreed.

tuberculosis patients is published.

2003

Malawi policy to

provide CPT for HIV+

ANECCA statement advocates use of **CPT** in infants and children.

2005

Malawi policy to provide CPT for all eligible HIV + patients is agreed and published.

Ugandan policy to provide CPT for all HIV + patients is agreed and published.

Zambian policy to provide CPT for all

2006

eligible HIV + patients is agreed WHO guidelines recommend CPT

among children,

adolescents and

limited settings.

adults in resource

provide CPT for all eligible HIV + is published.

2007

Zambian policy to

Discussion

- A favourable policy context is central to the adoption of research into policy. Even if a sound evidence base upon which policy can be constructed is in existence, an unfavourable national context makes it difficult for policy to develop.
 - Government structures and focus, donor interest and involvement, healthcare infrastructure and other clinical uses of cotrimoxazole shaped the development of CPT Policy in the three countries.
- The evidence base is an important aspect of a policy development process, but that it does not in itself drive policy-while poor evidence may stall a policy process, even when strong clinical research evidence is available to policy makers, it does not necessarily lead to policy change.
 - In our three countries, the only country to conduct a randomized clinical trial took the longest amount of time to create national policy on CPT. Policy makers in Malawi and Uganda valued the local evidence on the feasibility of an intervention (as in Malawi 2002 and 2005 and Uganda 2005) as well as its clinical efficacy.
- Policy entrepreneurs are central in driving policy processes, and can emerge from different types of healthcare institutions (NGOs, government or donor agencies).
 - In each of the three countries, the policy entrepreneurs held senior positions in well funded organisations concerned with both policy making and implementation. In Malawi when the research evidence was contested (2002) a powerful, well positioned policy entrepreneur, supported by policy champions played a particularly important role in driving policy forward.