

Research Process

- International standardised questionnaire translated into Tokpisin
- Selection and training of PLHIV interview team selective bias to persons of diverse sexualities
 and ensuring that the team had MSM, sex
 worker, transgender, married men, married
 women, single women, single man
- Basic research methods training, basic ethics in research, basic data recording, basic field work responsibilities – strong on confidentiality and responsibility expectations of team.

Research Process

- Selection of sites Mt Hagen and Chimbu based on epidemic patterns, service patterns and preparedness of local PLHIV group/s to be involved
- Prior on-ground consultations with relevant organisations and groups
- Negotiating entry to clinical spaces and sites where PLHIV gathered
- Eighty interviews conducted in late 2011 and data checking with interviewers conducted again in Port Moresby early in 2012

Another way is through language

- What is stigma?
- 'bagarapim ol narapela manmeri'
- 'pasin blo rabisim ol narapela manmeri'
- 'tok basait long ol narapela manmeri'

- What is discrimination?
- 'wokim pasin nogut long ol narapela manmeri'
- 'mekim nogut long manmeri'

- 80 x People Living with HIV were in interviewed.
- Research was conducted in Chimbu & Western Highlands Provinces.

- More then half of the respondents were 15 -29 interviewed.
- High percentage of respondents interviewed live with HIV for 1 – 4 years then followed those between 5 – 9 years.
- More then half the respondents lived in the village around the town settings
- More the half the respondents said they earned more then K201 to over K1,00.00 annual.

- Nearly all respondents indicated that they are sexually active although less than a quarter of the respondents say they are married or cohabiting with a partner.
- Over 10% identified as belonging to the category of sex worker and a small percentage said that they belong to the group MSM or gay.
- A high percentage said that they had been physically assaulted in the previous twelve months and that it was because of their HIV status.

- Over 50% said they had not attended social gatherings because of their HIV status.
- Nearly half the respondents said they had feelings of being ashamed, guilty or blamed themselves because of their HIV status.
- Being gossiped about was identified by respondents as the most persistent, commonly occurring, disturbing and upsetting way in which they experienced stigma and discrimination

- 70% of the respondents have heard of the HAMP Act., however, they have never read, discussed or very much on the Act.
- Half the respondents said their rights have been abused as a Person Living with HIV.
- Nearly 75% said they have never seek redress in court due to poor support and lack knowledge in the process and protocols.

- Over half the respondents said they had confronted, challenged or educated someone who was stigmatised and discriminated them.
- 60% of respondents said they had never seeked assistance with Legal Issues
- 70% of respondents said they have had supported their peers in dealing the stigma and discrimination (emotional and physical support)

- Over 60% respondents said they trusted their own peers.
- A quarter of respondents had disclosed to adult members of their families.
- Health Workers, teachers, social workers and Employers were least to be trusted to disclose.

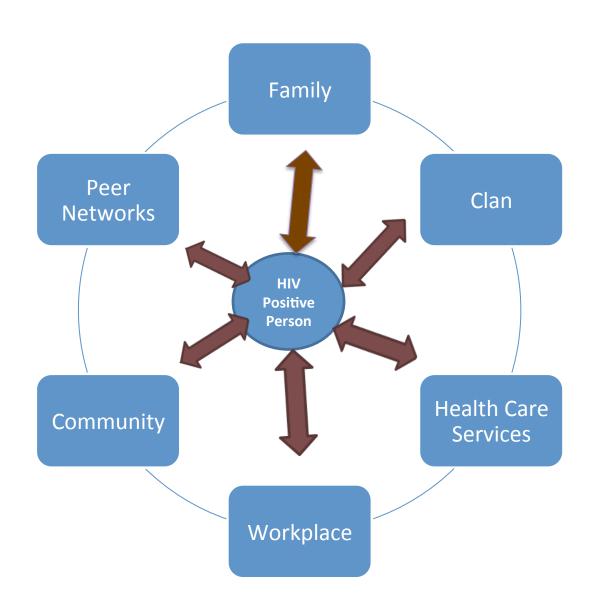
- 95% said they will have access to ART even if they were not on it.
- ART was also a problem due to distance and financial problem.
- Low respondent to the question of having constructive discussions on sex and sexual reproductive health, sexual relationship, emotional well-being including drug use.

- 60% of respondents said they had never been given PPTCT services.
- 5 women said they were coersed in termination of their pregnancy.
- Nearly 90% said they were on ART. There was concern on side effects & 50% said service satisfaction.
- Health Care Workers, Media, Govt employees Social Workers were least if the participants list to disclose

- 2 quarter said they found their disclosure very empowering.
- Shame & Fear including living in denial related to Testing & Diagnosing.
- &70% said they were confronted with internal stigma.

•

One way of showing how it is happening



Another way.....

Action

Effect

Impact

Rejection by family

No support to attend the clinic

Does not take ART

Gossip amongst the clan

Has to leave and go to another place

Loss of income and financial support

Community isolation

Denial of HIV status

Risk of on-going transmission

With acknowledgement of the considerable work of the research team, NACS funding support, UNAIDS funding and technical support.



Thank you