

Population based — Individual Screening Protocol for Rural Indian Populations

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January 2011



Introduction

1. Response to Chronicity

The WHO Health Systems framework (WHO, 2007) defines six components of a Health System – service delivery; health workforce; information; medical products, vaccines & technologies; financing; leadership and governance. Within this framework, the evolving 'chronicity' (long duration and slow progression) of the global disease burden requires a highly integrated response to provide continuum of healthcare, within a population, that does not have tertiary care as its focal point (Allotey, 2010) but community based care. In 2000, the World Health Assembly endorsed a three component global strategy (WHO, 2005) for the prevention and control of non-communicable diseases comprising of (i) surveillance to track and monitor major risk factors, (ii) promotion of health to reduce these risk factors and (iii) improved management to promote access to healthcare (Alwan, 2010). Tobacco use and excess weight (body mass index \geq 25) were reported as key risk factors for non-communicable diseases (Alwan, 2010) (Prospective Studies Collaboration, 2009) that needed to be urgently dealt with by the "new" health system.

It is indeed the case that prevention, treatment and management of chronic diseases entails a core set of interventions – primary prevention, proactive high-risk case identification (through routine screening and assessment of risk factors), education, efficient referrals, pharmacological and psychosocial interventions, long term surveillance, and monitoring and assessment of quality of care (Samb, 2010). In a health system designed to address the double burden of infectious and chronic disease and implement this set of interventions, Health Extension Workers (HEWs) are needed to bring care close to the community and play a critical part in early detection and support for long-term self-management and home-based care (Samb, 2010). The behavioural risk factors for chronic diseases closely linked with obesity are diet and physical inactivity. Improving diet and increasing physical activity are cost-effective interventions when dealt with at a policy level e.g. pricing interventions, regulation of food advertising (Cecchini, 2010) (Geneau, 2010). However, strategies targeting individuals e.g. tobacco control, salt reduction, and a multi-drug strategy to treat individuals with high-risk cardiovascular diseases have strong support in terms of cost-effectiveness for scale-up in low and middle income countries (Gaziano, 2007) (Asaria, 2007).

2. Definition of Screening

It is clear from the above discussion that treatment of chronic diseases needs a number of steps to be taken at the community level of which the very first one is screening. The World Health Organization (WHO) defines screening as the "presumptive identification of unrecognized disease or defects by means of tests, examinations or other procedures that can be applied rapidly". The Australians state



that the aim of screening for a disease or a risk factor is to reduce the burden of the disease in a community including incidence of disease, morbidity from the disease or mortality (Population Based Screening Framework, Screening Subcommittee, Australian Population Health Development Principal Committee, 2008).

The USPSTF (U.S. Preventive Services Task Force Grade Definitions, 2008) graded recommendations allow standard evidence based format for preventive/screening protocol development within a nation specific context. As listed by the USPSTF the Grade A&B recommendation across various risk groups are:

- <u>Adults (USPSTF A and B Recommendations, 2010)</u>: Blood pressure screening (aged 18 years and older); cholesterol abnormalities screening (men 35 years and older; women 45 years and older); cervical cancer screening in women who have been sexually active and have a cervix; tobacco use counselling and tobacco cessation interventions for those using tobacco products; colorectal cancer screening (adults beginning at the age of 50 years until age 75 years); HIV and syphilis screening for persons at increased risk; Chlamydia infection screening for women (sexually active, aged 24 years and younger, for older women at increased risk)
- Pregnant Woman (USPSTF A and B Recommendations, 2010): Bacteriuria screening, Hepatitis-B screening at first prenatal visit, Rh (D) blood typing and antibody testing, syphilis screening, tobacco use and counselling (providing augmented pregnancy tailored counselling to those who smoke) Chlamydia infection screening for women (aged 24 years and younger, for older women at increased risk)
- 3. <u>New Born Infants (USPSTF A and B Recommendations, 2010)</u>: Hemoglobinopathies (sickle cell disease) screening; congenital hypothyroidism and Phenylketonuria (PKU) screening

The WHO STEPS surveillance protocol also details the risk factors to be assessed periodically for chronic disease management (WHO, 2005). The WHO STEPS instrument for chronic disease risk-factor surveillance (WHO, 2005) allows for a careful design through screening of non-invasive parameters e.g. demographic information, physical activity assessment, personal medical history, physical non-invasive parameter measurement e.g. height, weight, waist circumference, hip circumference, blood pressure and heart rate by a non-medical provider in a non-clinical setting. Biochemical measurements e.g. blood glucose, blood lipids, triglycerides and HDL Cholesterol as recommended in the STEPS instrument can be performed in a clinical setting.

Despite evidence of their effectiveness, the delivery of preventive services continues to remain relatively low. In a US based study it was estimated that 7.4 hours per working day of physician's time was required to provision USPSTF Grade A&B recommended services to a population of 2,500 individuals defining a typical practice (Yarnall, 2003). Whereas, for only Grade-A recommended services the time implication is estimated at approximately 2 hours per working day (Yarnall, 2003). This study further substantiates the case for task-shifting routine screening activities towards HEWs and establishing effective referral pathways for '*high-risk*' individuals.

3. ICTPH Health Systems Approach

This paper details the age-specific evidence based protocol as assembled by the authors – all of whom are senior researchers at the IKP Centre for Technologies in Public Health (ICTPH). ICTPH is a not-for-profit research organization with its mission to innovate health-systems designs for remote rural populations (ICTPH, 2010) (ICTPH Epidemiology Data Set, 2010) – particularly those that are facing the dual burden of disease mentioned earlier. The Health Systems approach at ICTPH aims to redefine primary healthcare towards an inclusive continuum, accommodating evolving chronicity (Johar, ICTPH Health Systems Approach, 2010). The four elements defining the ICTPH Health Systems approach are human resource, infrastructure, technology, and financing.

Provisioning enhanced diagnostic capability through the village-based Rural Micro Health Centre (RMHC) allows for optimum condition management. Basic haematology, biochemistry, ophthalmology, curative care, intervention management are provisioned through a RMHC for a population of 10,000 people through a village-based nurse and a network of health extension workers (Johar, Financing Health Systems, 2010) (Johar, Provisioning Ophthalmic Care for Remote Rural Indian Populations, 2010) (Mor, 2010). The village-based RMHCs anchor multiple functions, wherein establishing a relationship focused on *'Wellness'* with the household is of utmost importance. Wellness as defined within the ICTPH Health Systems context, not only provisions curative services for management of acute conditions but also *risk-profiles populations* targeting secondary prevention at a population level. Identified *high-risk groups* are then exclusively targeted for provisioning specific interventions e.g. iron/folic acid supplementation through sprinkles for 6 month – 24 month old infants.

4. Plan of the Paper

This paper aims to detail the *individual risk-profiling protocol* as developed and implemented at the population level through a network of health extension workers within the ICTPH Health Systems context as implemented in close collaboration with its Thanjavur, Tamil Nadu based partner SughaVazhvu Healthcare (SughaVazhvu Healthcare, 2010).



Population based Individual Screening Protocol (PISP)

The Population based Individual Screening Protocol (PISP) as assembled by ICTPH aims to screen all individuals of a household using a special tool which is implemented by hamlet-based HEWs (referred to as the ICTPH Guide (Chockalingam, 2010). In concordance with several international screening guidelines which are primarily age-specific the PISP approach also targets four age groups within the intervened population – Adult (all individuals above 18 years of age); Adolescent (all individuals between 10 years – 18 years of age); Child (all individuals between 2 years to 10 years of age); and Infant (all individuals below 24 months of age).

Effective implementation of a high quality screening programme requires the development of a household roster and obtaining consent from the individuals that are sought to be screened.

- 1. <u>Household Roster:</u> Within the Karambayam, Thanjavur, Tamil Nadu context wherein PISP was implemented in close collaboration with SughaVazhvu (SughaVazhvu Healthcare, 2010), basic demographic details of a household entailing the name, age, gender, relationship to a predefined family head, address including GPS coordinates were obtained from Kshetriya Gramin Financial Services (KGFS) a local financial institution (IFMR Trust, 2008), which they shared with us after obtaining the consent of the households served by them no financial or economic information about the household was obtained from KGFS. Using the geo-tagged data and GIS technology, each one of the twelve HEWs were allocated a set of households that they would serve (ICTPH Guide; Lakshmanan, 2010) within the overall catchment population of 10,000 people for the RMHC, allowing proximate household allocation of approximately 200 households for each Guide. Pre-populated household specific rosters accompanied the age-specific PISP Optical Mark Recognition (OMR) sheets, along with the ICTPH Guide Kit (Das, Optical Mark Recognition Technology for Rural Health Data Collection, 2011), providing her with relevant demographic details and equipment for acquiring PISP related information.
- 2. <u>Consent Form</u>: Consent using a Standardised Form was obtained from all the PISP assessed members of a household, though either a signature or a thumb impression. For all PISP assessed infants, children and adolescents consent was obtained from their respective caregivers.

The following sections briefly describe the Tool used for each age group mentioned earlier. The Appendices have more detailed descriptions.

- 1. <u>Population based Individual Screening Protocol (PISP) ADULT (Appendix 1)</u>
- a. <u>Additional Demographic</u>: The code for the health extension worker, and individual ID as indicated in the household roster, date of birth, age completed in years, education status (illiterate, matriculate,



higher secondary, graduate, post graduate), currently enrolled in school, marital status, occupation in the last one year (daily wages on other people's land, cultivation on own land, manual labor, selfemployed non-farm work, government employee, non-government employee, student, homemaker, retired, unemployed (able to work), unemployed (unable to work) are cited as independent fields on the PISP OMR sheet.

b. <u>Anthropometric Measurement</u>: Height and weight for all individuals computing the Body Mass Index (BMI) (only exception for BMI computation – pregnant female) along with waist circumference (Dobbelsteyn, 2001) (Prospective Studies Collaboration, 2009) were measured using a measuring tape, weighing scale, cardboard slab and a marking pencil. The normal cut-off for waist circumference was < 80 cm for women and < 90 cm for men (WHO, 2000). The normal BMI range for adults was defined as 18.5 – 25 (with an additional cut-off at 23 for Asian populations). BMI of < 18.5 was categorized as underweight, BMI range of 26 – 29 as overweight, and BMI of ≥ 30 as obese (WHO, 2000).</p>



Figure 1: A snapshot of the PISP Adult tool in Tamil. This PISP tool has been implemented in Thanjavur, Tamil Nadu by one of the health extension workers in her routine PISP beat in her catchment area.

- c. <u>Blood Pressure:</u> Two blood pressure measurements for all individuals using a mercury sphygmomanometer separated by a minimum of 30 minutes were obtained (USPSTF Screening for High Blood Pressure, Topic Page, 2007).
- d. <u>Acute Conditions</u>: Through a home-based assessment of common acute conditions, the PISP protocol aims to identify individuals requiring immediate medical attention. Other than an extensive listing of common acute conditions, their onset in the last month, debilitating effect, treatment sought were also recorded. If the individual reports current existence of the reported symptoms immediate referral to a healthcare provider by the health extension worker is recommended. A listing of acute conditions persistent cough, fever, diarrhoea, weakness, fatigue, vomiting, worms



in stool, difficulty breathing, pain in abdomen, genital ulcers, painful urination, swelling ankles, difficulty hearing, skin problems/irritation, chest pain, paralysis, night sweats, weight loss, other. *Additional female*: menstrual problem (pain, irregularity, heavy bleeding), abnormal discharge (bad odour, change in colour). *Additional male*: Abnormal penile discharge (bad odour, change in colour) (Banerjee, 2010).

- e. <u>Personal Illness</u>: A recall based approach to record personal illness allows appropriate risk profiling of an individual for chronic conditions. Related surgical/non-surgical intervention, along with current treatment sought is also recorded. A listing of conditions assessed heart disease, high blood pressure, diabetes/high blood sugar, high cholesterol, liver disease, respiratory disease, tuberculosis, cancer, HIV/AIDS, STD, trauma, seizure disorder.
- f. <u>Family History</u>: A recall based approach to record family (mother, father and/or siblings) illness allows close assessment of acquired risk. A listing of conditions assessed heart disease, high blood pressure, diabetes/high blood sugar, high cholesterol, liver disease, respiratory disease, tuberculosis, cancer, HIV/AIDS, STD, trauma, seizure disorder.
- g. <u>Sexual and Reproductive Health</u>: Other than a query assessing the awareness and usage of contraceptive methods, this section entails to assess the reproductive health of a female. If the female reports a pregnancy an immediate referral to a healthcare facility for ante-natal care (ANC) is recommended. Other queries aim to assess if ANC is being currently sought, previous birth history of the female, including miscarriage, abortions and still-births.
- h. <u>Smoking Habits</u>: The PISP nicotine dependence assessment is an adaptation of the Fagerstrom Tolerance Questionnaire (Heatherton, 1991) incorporating tobacco consumption through chewing. The nicotine dependence through smoking, as assessed through a score follows the standard Fagerstrom protocol (Heatherton, 1991).
- i. <u>Alcohol Consumption</u>: The PISP alcohol consumption assessment is an adaption of the FAST Alcohol Screening Test (Hodgson, 2002). A clear scoring mechanism as illustrated in the FAST tool allows careful assessment of hazardous alcohol consumption within the population (Hodgson, 2002).
- j. <u>Visual Acuity</u>: The PISP protocol aims to assess myopia related refractive error leading to vision impairment using a Snellen Chart, and a 6mt long string, with markings at 1mt each, so as to move the patient closer to the Snellen chart in units of a meter each on reporting compromised reading capability beginning at 6mt. Upon reporting a compromised visual acuity of < 6/6 a referral to a healthcare provider is recommended by the health extension worker i.e. ICTPH Guide.



2. <u>Population based Individual Screening Protocol (PISP) – ADOLESCENT (Appendix 2)</u>

The Adolescent PISP Tool (for individuals of 10 years – 18 years of age) is primarily an adaption of the Adult PISP Tool. Within the demographic details an additional field to capture the Caregiver's ID was incorporated to ensure the consent of an adult/caregiver was obtained while the Adolescent PISP tool is implemented by the Health Extension Worker. The Sexual and Reproductive Health section is a slight adaptation of the Adult Tool.

Following the USPSTF recommendations for screening blood pressure (BP) at the population level (USPSTF Screening for High Blood Pressure, Topic Page, 2007) BP was excluded from the PISP Adolescent Tool. The Fagerstrom Tolerance Questionnaire (Heatherton, 1991) and FAST Alcohol Screening Test (Hodgson, 2002) are both recommended for adult populations (18 years and above of age) hence, a simplified adaption without any scoring capability was incorporated in the PISP Adolescent Tool.

For the adolescent population z-scores for BMI for age as defined by WHO were used to define standard PISP cut-off range (WHO, 2011). Waist Circumference was not measured for adolescents, in accordance for recommended guidelines. All the other sections of the Adolescent PISP Tool follow the Adult PISP Guidelines.

3. <u>Population based Individual Screening Protocol (PISP) – CHILD (Appendix 3)</u>

The PISP Child Tool is a further adaptation of the PISP Adolescent Tool accommodating an additional section detailing episodic occurrence of common childhood illness i.e. measles, chicken pox, mumps and polio. The sexual & reproductive health section, the smoking and alcohol history section as detailed in the PISP Adolescent Tool were deleted from the PISP Child Tool. The acute conditions section was also simplified further excluding menstrual and penile discharge sections as explained previously.

For child population ranging from 2 years – 10 years of age the z-scores for weight for age, and height for age, as defined for boys and girls by WHO were used to define standard PISP cut-off range (WHO, 2011).

4. <u>Population based Individual Screening Protocol (PISP) – INFANT (Appendix 4)</u>

The PISP Infant Tool was assembled to primarily focus on birth history, feeding practices, immunization, along with other parameters listed in the PISP Child Tool.

Detailed birth history is sought including:

a. Place of delivery - at home, government hospital, government health centre, private hospital/clinic



- b. Antenatal Care (ANC): *provider*: health professional/doctor, nurse/midwife, traditional birth attendant, no care sought; *Health-check-up*: at home, government hospital, government health centre, private hospital clinic
- c. Birth weight if possible from a standard infant delivery card, or else recall
- d. Immunization history as per the WHO India guidelines for Polio, DPT, BCG, Measles, MMR (WHO India, 2011)

The acute conditions and the childhood illness section remain the same as detailed in the PISP Child Tool. For infant population < 2 years of age the z-scores for weight for age and height for age, as defined for boys and girls by WHO were used to define standard PISP cut-off range (WHO, 2011).

The breast feeding practice, exclusivity, and frequency; complementary feeding – initiation, pattern, and feeding practice is assessed using the *WHO tool for assessing infant and young child feeding practice* (WHO, 2008). The WHO tool is recommended as an indicator for population level assessment of infant and young child feeding practices (WHO, 2008). A clear recommendation towards usage of the WHO tool for large-scale surveys and national programs, and against indicator definition translation into caregiver messaging is noted (WHO, 2008). This limits PISP towards making individual caregiver specific diet recommendation upon assessment.

Human Resource facilitation for PISP

Within the ICTPH Health Systems context primarily two levels of human-resource are necessary at the village level. Each village-based RMHC is managed by a nurse, servicing a population of 10,000 people and for every 1,000 people (primarily a hamlet within the village context) a local health extension worker is required (Lakshmanan, 2010). The RMHC is supervised by an assigned medical practitioner overseeing all provisioned curative care, along with anchoring other activities. The health extension worker referred to as *Guide* within the ICTPH Health Systems context is the primary anchor point of the population to the Health System.

The tasks of a *Guide* primarily entail individual screening at the population level, intervention implementation, clinical rotation at the RMHC and follow-up care. Amongst all of these, individual screening establishes her status within the community, through methodically approaching households within her service area. This allows her an opportunity to carefully assess the profile of all screened individuals and categorize them within assigned risk-groups.

The integrated data-collection platform allows a smooth flow of accumulated information to the RMHC based individual electronic health record (hosting personal, demographic (geo-tag), family history and health information) allowing automated analysis and risk-profiling of all individuals within a household.

The analyzed individual health assessment reports are then delivered back to the household with a turnaround time of a week, with automated individual risk-factor specific recommendations. The recommendations are further detailed by the *Guide* to various members of the household reiterating the message to prioritized '*high*' risk individuals. The individualized recommendations allows an opportunity to the *Guide* to establish a trusting rapport across all age groups within the household allowing her to stress the urgency to follow-up based on individualized risk exposures.

Within the ICTPH Health Systems approach, a through community understanding is attained through methodically interacting and assessing *all* members of the community. This not only allows a deeper understanding of the serviced population to the healthcare provider but also empowers resident individuals with pertinent health information.

Technology facilitation for PISP

Among various data collection technologies, Optical Character Recognition (OCR), Optical Mark Recognition (OMR) and conventional manual data entry methods were evaluated. For PISP implementation considering the cost, turnaround time, error rates and responses from other user group, OMR emerged as the preferred choice in the short-run (Das, Optical Mark Recognition Technology for Rural Health Data Collection, 2011) (Das, Data collection and technology, 2010). Going forward the intent is to switch to mobile phone based data entry technologies which will enable the reduction in the error rates and the level of latency (delay between data entry and printing of final analysis).

Figure 2 shows the process-flow protocol for completing a PISP cycle, i.e., assessing an individual in a household to risk-assessed report delivery as a multi-step process with a current turn-around time of two weeks. The PISP-OMR forms are collected weekly by the Guide supervisor and are sent for scanning, and error checking to the technology team. The scanned results of the PISP-OMR are obtained in a tabular, excel sheet format as shown in *Table-1*. The excel sheets are then integrated with the Electronic Health Record (EHR) portal, wherein household specific individual information is mapped to the demographic information as obtained from KGFS, and detailed in the Household Roster section of this paper.



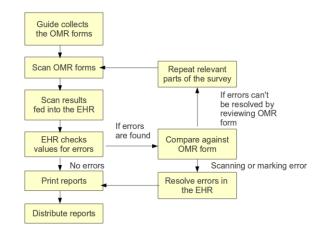


Figure 2: The process flow protocol for PISP. OMR sheets collected weekly, by the Health Extension Worker (ICTPH – Guide) supervisor, forwarded for scanning and error checking to the technology vertical, back-checks conducted if deemed necessary, validated PISP-OMR processed for automated reporting as generated through the ICTPH Electronic Health Record (EHR) portal and delivered to the relevant household through the Guide network.

	Α	В	С	D	E	F	G	Н	1	J	К	L	М	N	0	—
1	Form_No	F101	F102	F103	F201D	F201M	F201Y	F301D	F301M	F301Y	F401	F501	F601	F701	F801	F90
2	0000001122	03	3748203	3748202	13	12	2010	16	12	1995	14	6	3	Y	N	Y
3	0000001123	03	3748204	3748202	13	12	2010	02	03	2000	10	6	1	Y	Y	Y
4	0000001124	07	205905	205901	06	11	2010	02	03	1996	14	6	3	N	N	Ν
5	0000001125	07	183004	183001	09	11	2010	24	04	1993	17	6	3			
6	0000001126	07	183005	183001	09	11	2010	11	06	1997	13	6	2	Y	Y	Y
7	0000001127	07	203903	203901	09	11	2010	01	05	1998	12	6	2	N	N	Ν
8	0000001128	07	176405	176401	07	11	2010	06	12	1994	16	6	3	N	N	Ν
9	0000001129	07	2920*	29201	30	10	2010	21	19	1998	12	6	2	Y	Y	Y
10	0000001130	01	790307	790301	12	11	2010	23	03	2001	08	6	1	N	N	Ν
11	0000001131	01	796603	796601	26	11	2010	09	01	1998	11	6	2	N	N	Ν
12	0000001132	08	191804	191801	27	11	2010	12	02	1993		6	3	N	N	Ν
13	0000001133	08	184103	184102	01	12	2010	23	01	1996	15	6	3	N	N	Ν

<u>**Table 1**</u>: A snapshot of the excel data dump of the scanned PISP – OMR sheets. The excel sheets are then merged with the back-end household specific individual database in the Electronic Health Record (EHR) portal for further analysis and processing.

Post the OMR-PISP and demographic data mapping within the EHR, the hamlet specific Guide information is also mapped to the data. Automated error checking of the data within cut-off limits as defined in *Table-2* for BMI, blood pressure, waist circumference, nicotine and alcohol dependence are applied. For most other fields within PISP e.g. acute conditions, family history etc. are primarily recall



based binary fields with a yes/no reporting. These are checked for skip-logics, empty fields and gender consistency e.g. female reporting penile discharge, male pregnancies are flagged as errors.

The error checking stage is followed by automated risk analysis through the EHR. Based on the cut-offs defined in *Table-2* pre-determined messaging for various levels of identified risk along with specific recommendations are tagged to the individual report. For example, relating to blood pressure three levels are defined, normal, high and very high. Individuals with very-high blood pressure are prioritized for follow-up and are strongly recommended to seek medical attention given the associated probability of hypertension. Similarly for pregnant women, a strong ANC recommendation is made through the PISP report. Existing acute-conditions are also recommended for attention from a medical practitioner.

BMI		Blood Pressure	
Range	Assessment	Range	Assessment
< 10 or > 40	ERROR	<60/30 or >300/150	ERROR
10 - 14.9	Very Low	120/80	Normal
15 - 18.4	Low	>120/80 and <140/90	High
18.5 - 22.9	Normal	>140/90	Very High
23 - 24.9	Above Normal	Nicotine Dependence	
25 - 29.9	High	Score	Assessment
30 - 40	Very High	≤3	Low Risk
Waist Circumfere	nce	3 - 7	High Risk
Range	Assessment	7 - 10	Very High Risk
> 80 cm (Female)	High	Alcohol Dependence	
> 90 cm (Male)	High	Score	Assessment
		> 3	High Risk

<u>**Table 2**</u>: Risk definition for five conditions as protocolized in the Electronic Health Record (EHR) portal for automated risk analysis for PISP assessed individuals. Height & weight (computing the BMI), waist circumference, blood pressure, nicotine dependence as implemented through the Fagerstrom Score (Heatherton, 1991), and alcohol dependence as implemented through the FAST Score (Hodgson, 2002)

As shown in *Fig. 3*, the mapped OMR-PISP and demographic information in the EHR post error checking and risk-profiling are assembled together as individual reports. The household specific individual reports are consolidated together in the household folder which is delivered back to the respective household through their hamlet specific guide.



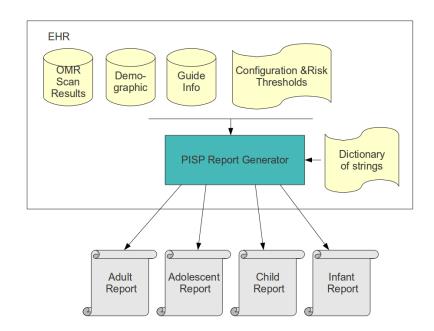


Figure 3: Automated report processing protocol for PISP screened individuals. The scanned PISP-OMR sheets are mapped to respective demographic and Guide details, as obtained from the household specific individual database in the Electronic Health Record (EHR) portal. Pre-configured risk analysis of all PISP assessed individuals yielding age-specific PISP risk assessment report generation.

PISP – An Early Snapshot from Karambayam, Tamil Nadu

The PISP tools, as developed by ICTPH for four different age groups, were implemented in close collaboration with SughaVazhvu Healthcare, Thanjavur, Tamil Nadu (SughaVazhvu Healthcare, 2010). The Karambayam Rural Micro Health Centre, servicing a population of 10,000 people along with a network of 12 Health Extension Workers implemented the PISP tool, following a pre-specified beat in October 2010.

Fig. 4 illustrates the CVD risk-profiles of 547 adults based on their PISP assessment, as collected through the first-batch of PISP-OMR in the first two weeks of PISP implementation in Karambayam through Health Extension Workers. The three risk parameters indicated are BMI, blood pressure and waist circumference. 16% of adult population was assessed with high blood pressure, with an additional 5% of the adult population with very high blood pressure indicating the need of an immediate therapeutic intervention (risk classification categories for high and very-high blood pressure as listed in *Table-2*),



27% with high BMI (joint representation of high and very-high BMI categories as listed in *Table-2*), 30% with low BMI (joint representation of low and very-low BMI categories as listed in *Table-2*), and 24% with high waist circumference (joint representation of male and female WC categories as listed in *Table-2*). Other PISP assessment risk indicators that also contribute towards an individual's CVD risk score are family history, personal history, age and sex.

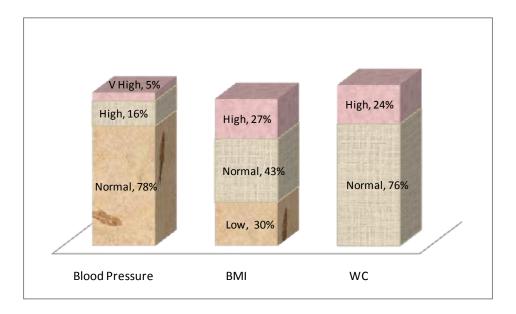


Figure 4: CVD risk profiling of 547 Adults based on three risk factors as obtained through the PISP Adult Tool. The three reported *CVD Risk Factors* are *Body Mass Index (BMI) for 473 adults* as computed from height and weight, *blood pressure for 494 adults*, and *waist circumference for 518 adults*. All adult individuals screened through the PISP Adult Tool in the year 2010, were residents of Karambayam Village, Thanjavur District in Tamil Nadu, being serviced by the ICTPH/SughaVazhvu Karambayam Rural Micro Health Centre.

All the other parameters relating to acute illness, visual acuity, immunization for infants, ANC for pregnant mothers are also individually assessed and consolidated as household specific individual reports.

A multi-factorial individual risk-factor analysis for all the variables gathered through the PISP tool along with village-level 'Wellness Index' computation will be presented in a separate working paper.



Conclusion

This paper demonstrates the feasibility of population-based individual screening for age-specific risk factors.

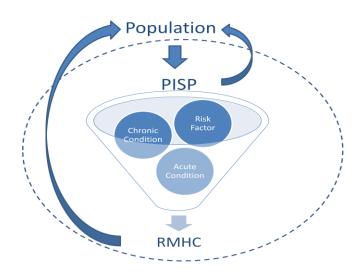


Figure 5: The Annual PISP cycle as implemented in rural Indian geographies utilizing Optical Mark Recognition (OMR) technology through trained Health Extension Workers (ICTPH Guide). Healthy populations are tracked annually for risk factor development. Individuals with elevated risk factors, chronic disease condition and/or acutely-illness are referred to the RMHC and/or other healthcare provider for a follow-up.

As shown in *Fig. 5*, population screening for risk profiling, and detection of existing acute and chronic conditions allow appropriate individual prioritization and long-term condition management. The effectiveness of the PISP intervention as monitored by population outcome through risk factor indicators and enhanced disease-condition management is yet to be established.

The PISP individualized assessment provides an opportunity for customized risk profiling and focused recommendations. The population level interpretation of the *WHO tool for assessing infant and young child feeding practice* (WHO, 2008) integrated in the PISP Infant tool limits making individual caregiver specific diet recommendations.

Appropriate technology intervention through OMR, facilitated the turn-round time but the latency and error rate can be further reduced through the use of mobile-based technological solutions. PISP implementation by Health Extension Workers (ICTPH Guide) was proven feasible through the training and implementation phase.

The enormity of the data fields associated with the age-specific PISP tools, allowed minimum interaction time between the ICTPH Guide and the household individual. One of the targets of using specific tools like PISP was to establish a dialogue on '*Wellness*' within the household. Mechanical and elaborate data fields minimize free interaction, limiting the outcome of the PISP exercise. As a recommendation, a more methodological approach, wherein the PISP interaction is accomplished through multiple-visits to a household, not only allows an opportunity to carefully assess an individual but also simultaneously acton already acquired information yielding specific recommendations of either a follow-up, or an intervention e.g. iron-supplementation for a 6 month old infant.

Within the current PISP protocol, a strong emphasis has been laid on accumulating risk factors relating to cardiovascular diseases and diabetes. As a recommendation, other chronic conditions, most importantly mental illness (Alexander, 2008), cataract, and presbyopia may also be included within the PISP Adult Tool framework.

Individual prioritization based on risk-factor profiling is only a one-dimensional analysis. Ensuring accessible and equitable healthcare provisioning to serviced populations, poverty identifying markers other than the usual BPL (Below Poverty Line) card need to be integrated in the PISP framework. Given the simplicity of Grameen Foundation's *Progress out of Poverty Index* (Schreiner, 2008) to assess the poverty index of a household may be evaluated for integration within the PISP framework. These poverty indicators coupled with PISP risk-factor profile will then classify the populations on a multi-dimensional high-risk matrix including both individual's health and ability to afford/seek healthcare.

The PISP protocol as illustrated in this paper has been implemented in a population based setting through Health Extension Workers. Given the protocolized nature of PISP, this can also be integrated into the SOAP (Subjective Objective Assessment Plan) methodology as implemented in a primary-care clinic based setting, facilitating opportunistic screening.



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608	How many miscarriages, a	bortions or s	tillbirth	s have voi	u had?							567		
701	Have you ever chewed toba	cco? Yes 🔿) N	o (Skip to 70	3) 🔿	702	Do you o	currently che	ew tobacco	o?	١	res 🔿		No C
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ADOLESCENT QUESTIONNAIRE OMR SHEET

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FLAG: ONLY ADMINISTER THE QUESTIONNAIRE IF THE ADOLESCENT IS PRESENT; ADOLESCENT SHOULD BE BETWEEN THE AGES OF 10 AND 18; ONLY ASK THESE QUESTIONS AFTER THE CONSENT HAS BEEN SIGNED. ADOLESCENT SHOULD ANSWER QUESTIONS WITHOUT THE INFLUENCE OF A CAREGIVER IF ABLE TO DO SO.

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INFORMATION PANEL

SVG Code		A	dolesc	ent ID		_			Caregiv	er ID						f Interview	
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DISEASE SYMPTOMS	- Flag	: Only	ask Q.	-302-3	04 if th	e ansv	ver is `										
301. Have you experienced conditions in the last month?		owing	302. A	bility			303 M					301 AS		•*2		304. S	Still
Condition:	Yes	No	affecte Yes	ed? No						-	locatio					exists/ Yes	Pe
1. Cold symptoms (runny nose, congestion, sneezing)	Y		(Y)		1	2	3	4	5		7	8	9	10	(11)	(Y)	+
2. Persistent Cough	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	-
3. Fever	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
4. Diarrhea	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
5. Weakness / Fatigue	Y	\mathbb{N}	Y	\mathbb{N}	1	2	3	4	5	6	7	8	9	10	11	Y	T
6. Vomiting	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
7. Worms in stool	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
8. Difficulty in breathing	Y	\mathbb{N}	Y	\mathbb{N}	1	2	3	4	5	6	7	8	9	10	11	Y	
9. Pain in abdomen	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
10. Genital ulcers	Y	\mathbb{N}	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
11. Painful urination	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
12. Swelling of ankles	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
13. Difficult in hearing	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
14. Skin problems / irritation	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
15. Chest pain	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
16. Paralysis	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
17. Night sweats	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
18. Weight loss	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
19. Other pain	Y	\mathbb{N}	Y	\mathbb{N}	1	2	3	4	5	6	7	8	9	10	11	Y	
IF [ADOLESCENT] IS FEM	IALE :																
20. Menstrual problems (pain / irregularity / heavy bleeding)	Y	N	Y	N	1	2	3	4	5	6	7	8	9	10	11	Y	
21. Abnormal discharge (bad odour / change in colour)	Y	\mathbb{N}	Y	\mathbb{N}	1	2	3	4	5	6	7	8	9	10	11	Y	
IF [ADOLESCENT] IS MAL	E :																
22. Abnormal penile discharge (bad odour / change in colour)	Y	N	Y	\mathbb{N}	1	2	3	4	5	6	7	8	9	10	(11)	Y	

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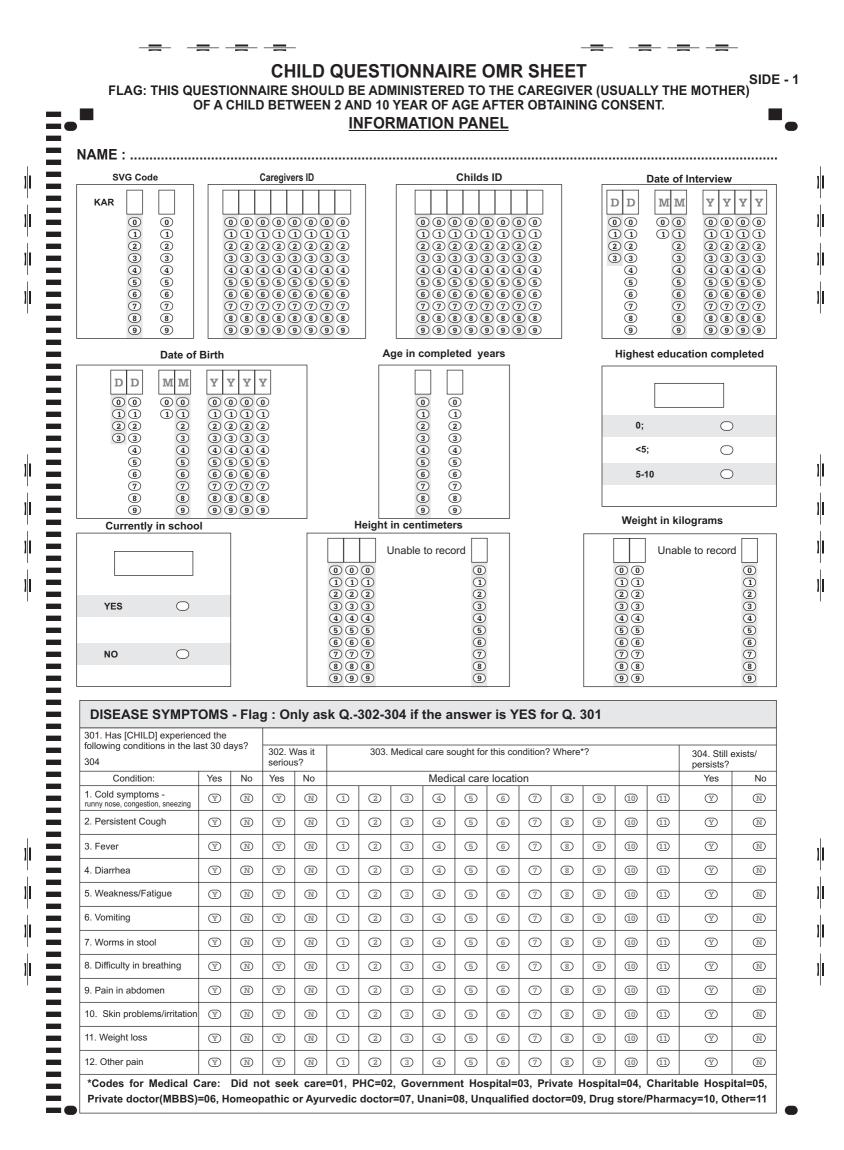
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305				<u> </u>					questio					s (Refer		HC for	check-u	ıp) (\supset		No	\supset
					-	•			2.403 if								4hiaQ, 4	02 4 m		ainal tua		a a ui a a d
401-	401.		as a nea	inncare p	provider	ever ton	u you ina	402.	ESCENT.	403.	02. Di	401			ve surg	Jery IOI	uns?, 4	US. Ally	402.	gical trea	403.	eceiveu?
403		dition :			Yes	No	DK	Yes	No	Yes	No	-	onditi	ion :		Yes	No	DK	Yes	No	Yes	No
	1. H	leart d	isease		Y	\mathbb{N}	ØK	Y	N	Y	\mathbb{N}	7. T	uber	culosis		Y	\mathbb{N}	DK	Y		Y	N
	2.1	ligh Bl	Þ		Y	\mathbb{N}	DK	Y	\mathbb{N}	Y	\mathbb{N}	8. C	ance	er		Y	N	DK	Y	\mathbb{N}	Y	\mathbb{N}
	3. [Diabete	s		Y	\mathbb{N}	DK	Y	\mathbb{N}	Y	\mathbb{N}	9. S	STD			Y	N	DK	Y	\mathbb{N}	Y	\mathbb{N}
	4. H	ligh ch	olester	ol	Y	\mathbb{N}	ØK	Y	\mathbb{N}	Y	\mathbb{N}	9. H	IIV / A	AIDS		Y	N	DK	Y	N	Y	N
	5. L	iver di	sease		Y	\mathbb{N}	ØK	Y	\mathbb{N}	Y	\mathbb{N}	10.	Trau	ma		Y	N	DK	Y	N	Y	N
	6. A	Asthma	I		Y	\mathbb{N}	ØK	Y	\mathbb{N}	Y	\mathbb{N}	11.	Seizı	ure diso	rder	Y	N	DK	Y	N	Y	\mathbb{N}
501 502	Are	e you a you u	aware a se any	about co	ontrace ception	eption r metho	nethod ods?	ls?	ept strid			Y	/es /es has	ever b	een n	narrie	No No ed.			DK DK	0K	
503	Are v	ou curr	ently pre	egnant ?	?					refer t	/es)	N Skip to	0 505	\mathbf{O}		D	ON'T KI kip to Q.	NOW	DK	
504	How	many nt pre	times ł	nave yo , if app	ou beer				clude at didn'		2	3			5	6	7	8	9	10	11	12
505	Have	you e	ver sm	oked to	obacco	?						Y	/es	\bigcirc			No	\bigcirc				
506	Numl	per of	times t	obacco	consu	med in	past n	nonth		0 () 1.	-2 () 3	8-5 🔿	6-9	\bigcirc	10-19	⊃ 20-	29 🔿	30 (DON'T KM	IOW DK
507	Have	you e	ver had	d any a	lcohol,	like be	er, win	e, or li	quor?			Ye	s	\bigcirc				No (Sk	ip to Se	ection 6	5) 🔿	
508	How	nany a	lcoholic	drinks o	did you l	nave in	the last	month?)	0 () 1	-2 (3	8-5 🔿	6-9	\bigcirc	10-19) 20-	29 🔿	30 (DON'T KM	IOW DE
6. Bl	МІ									<u> </u>												
	н	eight	in cent	timeter	rs				Wei	ght in I	kilog	rams	;									
601 602				012346678								6 9 8 9 6										
603		70	9 9 9 9	() () () () () () () () () () () () () (0 7 8 9										
7. VI	SUAL A	CUITY																				
			1	1	1	I	1	1	Ţ,	-												
		6/6	5/6	4/6	3/6	2/6	1/6	CF	НМ	Blind												
701	Right Eye	6	5	4	3	2	1	10	11	0												
	Left																					
	Eye	6	5	4	3	2	1	10	11	0	_											
702	1	lf	less f	then (6/6 re	fer to	opti	cian														

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305	Interviewer checkpo	int:Wa	as the a	answer	Yes to a	any que	stion ι	under Q.3	04? Ye	es (Refer to R	MHC for	check-ı	ıp) (qı	\supset	No DK 08 19 10 10 10		
A. CHILDHOOD ILLNESSES 401 401. Has a healthcare provider ever told you that [CHILD] has: Yes No DK 1. Measles 「 ⑤ ⑤ ⑥ ⑥ ⑥ Ø 2. Chicken pox 「 ⑦ ⑧ Ø																	
401	401. Has a healtho	are pr	ovide	r ever	told ye	ou tha	t [CH	ILD] has	:	Yes			No			DK	
	1. Measles									Y			\mathbb{N}			DK	
	2. Chicken pox									Y			\mathbb{N}			DK	
	3. Mumps									Y			\mathbb{N}			DK	
	4. Polio									Y			\mathbb{N}				
5. PE	RSONAL ILLNESSES	6 - Flag	: Only	ask Q	.502 an	nd Q.50)3 if tl	ne answe	er is YES	for Q.501							
501	Flag:501. Has a healt	hcare p	orovide	r ever te	old you	that yo	u have	e?; 502. E)id you hav	ve surgery fo	or this?;	503. Ar	ny non-	surgica	l treatm	nent rec	eived?
									-				-	-		1	
	Condition :	Yes	No	DK	Yes	No	Yes	No	Conditio	on :	Yes	No	DK	Yes	No	Yes	No
	1. Heart disease	Y	\mathbb{N}	ØK	Y	N	Y	N	7. Tuber	culosis	Y	N	ØK	Y	N	Y	\mathbb{N}
	2. High Blood Pressure	• (Y)	\mathbb{N}	ØK	Y	N	Y	N	8. Cance	er	Y	N	ØK	Y	N	Y	\mathbb{N}
	3. High blood sugar			DK	Y	N	_	N	9. HIV / /	AIDS	Y	N	DK	Y	N		
	4. High cholesterol				Y	N	_	N	10. Trau	ma	Y		DK	Y	N		\mathbb{N}
	5. Liver disease					N	_	N	11. Seiz	ure disorder	Y	N	DK	Y	N	Y	\mathbb{N}
	6. Respiratory Disease		N	DK	Y	N	Y										
6. VIS	SUAL ACUITY																
		6/6	5	/6	4/6	3/	/6	2/6	1/6	CF	НМ	BI	ind				
														lf la		en 6/	-
601	Right Eye	6		5	4		D	2	1	10	11					optici	
				_						_				TOTO		puon	
	Left Eye	6		5	4	0	Ð	2	1	10	11						
602	-									1 1							

501	Flag:501. Has a health	hcare p	rovide	r ever t	old you	that you	u have	e?; 502. E	Did you hav	e surgery fo	or this?;	503. Ar	ny non-	surgical	treatm	ent rec	eived
503	501				502		503		501					502		503	
	Condition :	Yes	No	DK	Yes	No	Yes	No	Conditio	n :	Yes	No	DK	Yes	No	Yes	No
	1. Heart disease	Y	\mathbb{N}	DK	Y	\mathbb{N}	Y	\mathbb{N}	7. Tubero	culosis	Y	\mathbb{N}	DK	Y	\mathbb{N}	Y	
	2. High Blood Pressure	Y	\mathbb{N}	DK	Y	N	Y	N	8. Cance	r	Y	\mathbb{N}	DK	Y	N	Y	N
	3. High blood sugar	Y	\mathbb{N}	DK	Y	N	Y	N	9. HIV / A	NDS	Y	\mathbb{N}	DK	Y	N	Y	N
	4. High cholesterol	Y	\mathbb{N}	DK	Y	N	Y	N	10. Traur	na	Y	\mathbb{N}	DK	Y	N	Y	N
	5. Liver disease	Y	\mathbb{N}	DK	Y	N	Y	N	11. Seizu	ıre disorder	Y	\mathbb{N}	DK	Y	N	Y	N
	6. Respiratory Disease	Y	N	DK	Y	\mathbb{N}	Y	N									
6. VIS	SUAL ACUITY	• •					•								•	•	
		6/6	5	/6	4/6	3/	6	2/6	1/6	CF	НМ	Bli	ind				
601	Right Eye	6		5	4	3		2	1	10	11			lf le	ss th		3
														refe	r to o	pticia	an

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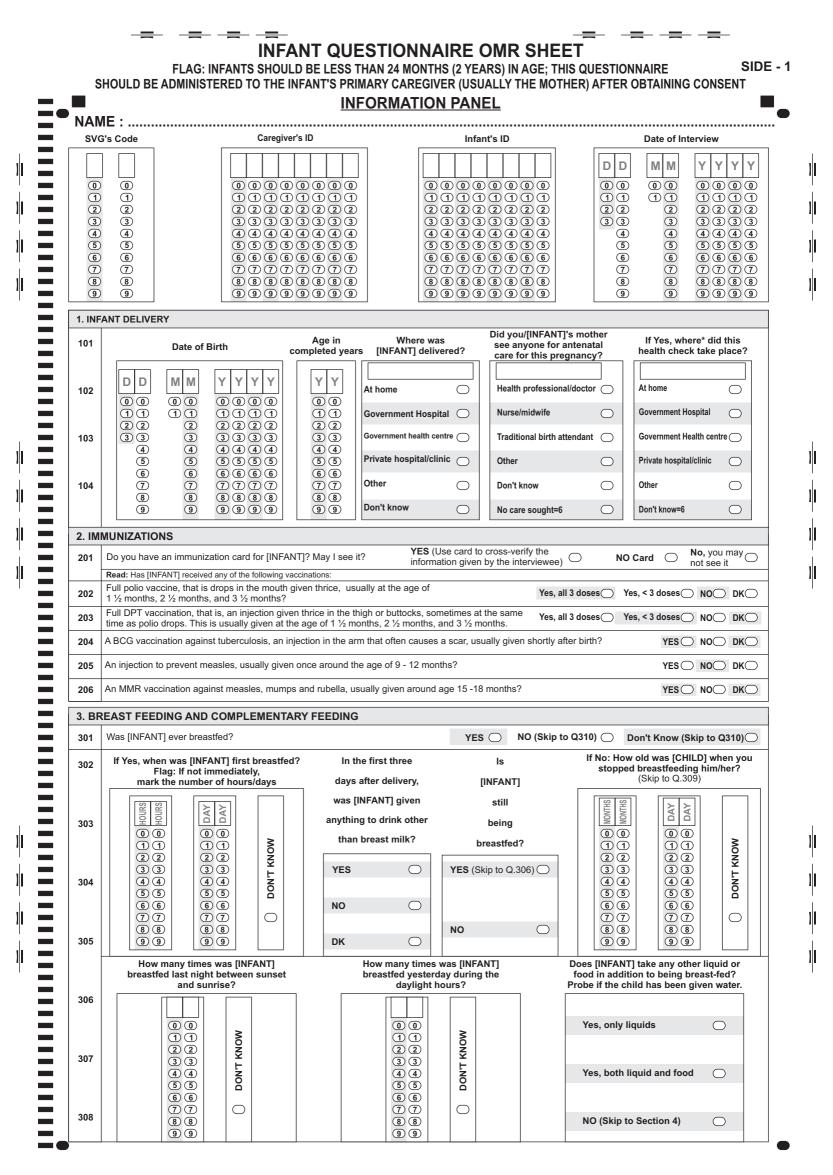
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	0 0 d or fo	>						Plea	ase take				out wha		Is [INFA	NT]			
	If Yes, how old was [INFANT when he/she was first given other liquid or food? Image: Second	DON'T KNOW												Y	ES		NO		DK
	th was []h (c) (c) (c) (c) (c) (c) (c) (c) (c) (c)	ONT			in water?		l infant fr	ormula fo	d as a liqu	uid (for o		octogon	Protoin _(X)						DK DK
	w old			3. An	y other m							•	FIOLEIIIX)	C	Ď		$\overline{\mathbb{N}}$		DK
	Image: Construction	$ \circ $			iit juice? a or coffe	e?									D D				OK OK
					y other lie										\mathbb{D}				DK
311	Please take a m until going to s																ie time o	ot wakin	ıg up
		OOD ITI	M		YES	NO			FOOD IT				YES	NO	412	2			
	 Any porridge or gru Any commercially f 		/ food suc	h as Cerelac	1	2		Any chick Any other	en, duck o meat?	r other bi	rds?		1	2		j –	_≆ ∏		
	or Farex? 3. Any bread, roti, ch	apati. rice. I	noodles, b	iscuits. idli.		2	12.	Any eggs	?				1	2		ni-soli	IFANT at nig		
	or any other foods	made from	grains?						or dried fi	sh or sho	llfich?					id, ser	did []N	11	KNOW
	4. Any pumpkin, carr yellow or orange in	nside?				2							1	2		of soli	g the (33	Т KN
	Any white potatoe other foods made f	rom roots?		a, or any	1	2	14.	Any toods	s made from	m beans,	peas, or l	entils?	1	2		vings	or soft foods (no liquids) did [NFANT] have yesterday during the day or at night?	4 4 5 5	DON'T
	 6. Any dark green, le 7. Any ripe mangoes 			e, or jackfruit?		2		Any nuts? Any chee	? se or yogh	urt?			1	2		ny ser	foods terday	66 77	
	8. Any other fruit or ve 9. Any liver, kidney, h	getables?				2	17.	Any food	made with solid or se	oil, fat, g		tter?	(1) (1)	2		ow ma	r soft	88	
				edis?	0	2	10.	Any other		enn-soliu	1000 ?		0			Ť	o la L	99	
		MEASU	IRES									4.B.	HEIGH		VEIGHT	г			
	Do you				opy the				NFANT]'					Ца	i a la t		14/		
404	birth weig	ght writt	en	i	n kilogra	ims -		g: If bir	st of yo th weigh	t is unl	known,			i	ight in meters			eight in	
401	down? M	card with [INFANT]'s weight o							the blan		0.	404						grams	
					•				•										
402				0		00		0							1				
	YES	C	~	2		$\overline{2}$ $\overline{2}$ $\overline{3}$ $\overline{3}$		2		2 2 3 3		405		223	2		$\overline{2}$ $\overline{2}$ $\overline{2}$ $\overline{3}$ $\overline{3}$ $\overline{3}$	2	record
				4	$\overline{\underline{4}}$	<u> </u>		4	4	4				4 4	4	to re	44		
403		_		5 6	6	5 5 6 6		(5) (6)	6	5 6 6				550	6	Unable to	550	9 6	Unable to
	NO (Skip to	Q.403) ((7) (8)		7 7 8 8		(7) (8)	8	7) (7) 8) (8)					7) 8) :	5	770 880	7) 8	5
				9	90	99		9	90	99				99	9 (\supset	99	9	0
5. DIS	EASE SYMPTON	S - Flag	Only	ask Q50	2-504 if t	he ansv	ver is Y	ES for	Q.501										
	las [INFANT] expe ng conditions in th			500	Was it	1	500			-			501 AS						
504				seric			503.	. Medica	al care so	ugnt io	r this co	natuon	where	?			504.	Still ex	ists?
1 Col	Condition:	Ye		lo Yes			1	1	Medica	al care l	ocation		1			1)	Yes	No
runny no	sistent Cough	ng O				1	2	3	(4) (4)	5	6	7	8	9	10	(1) (1)		Y Y	
3. Fev 4. Dia	er						2	3	(4) (4)	55	6	7	8	99	10		(
5. Wea	akness/Fatigue	0		\mathbb{V}		1	2	3	4	5	6	7	8	9	10	(1)	(\mathbb{Y}	N
	ms in stool	0				1	2	3	(4) (4)	5	6 6	7 7	8	9	10	(1) (1)	(Y Y	
	culty in breathing in abdomen	0					2	3	(4) (4)	5	6	(7) (7)	8	9	10	(1) (1)		Ý Y	
10. SI	kin problems/irrita	ion 🤇				1	2	3	4	5	6	7	8	9	10	1	($\overline{\mathbb{Y}}$	\mathbb{N}
	her pain		-			1	2	3	(4) (4)	5	6	7	8	9	10	(1) (1)		Y Y	
*Code	s for Medical Ca	re: Did r	iot seel	care=01,	PHC=02	, Gover	nment l	lospital	=03, Priv	ate Hos	spital=04	l, Chari	table Ho	spital=0	5, Priva	ite doc	tor(MBB	S)=06,	
Home	opathic or Ayurved		,				,	0			,		MHC for	check-	ar (ar	$\overline{}$		No	0
	ILDHOOD ILLNI	<u> </u>					1		20011										
601	601. Has a hea	Ithcare	provide	er ever to	d you th	at [INFA	ANT] ha	as:					YES		NO		DON'	Γ ΚΝΟΥ	v
	1. Measles												Y		\mathbb{N}		(N	
	2. Chicken p	x											Y		\mathbb{N}		(N	
	3. Mumps												Y			+	(\mathbb{N}	

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