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This paper is a draft submission to the

WIDER Development Conference

Human capital and growth

6-7 June 2016 Helsinki, Finland

This is a draft version of a conference paper submitted for presentation at UNU-WIDER's conference, held in Helsinki on 6-7 June 2016. This is not a formal publication of UNU-WIDER and may reflect work-in-progress.

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The test effect: Behavioral change and potential biases due to (biomedical) testing in surveys[☆]

February 9, 2016

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Abstract

This paper is the first to rigorously analyze unintended effects of biomedical testing in surveys. Random assignment of blood pressure measurements in a 2013 household survey in Tanzania, and a second survey of the same individuals two years later, allows for the identification of this “test effect” on hypertension (high blood pressure) awareness, health care provider consultations for hypertension, and uptake of voluntary health insurance. As these were the baseline and follow-up surveys of a health insurance impact evaluation, the possible bias in the insurance impact estimates caused by the test effect can also be estimated. Since, complying with ethical standards, respondents who were tested were told their test result, the differential effect of a “good” versus a “bad” test result can be determined. A *bad* test result is found to significantly increase both hypertension awareness, and likelihood of health care provider consultations for hypertension. There is weaker evidence that a *good* test result decreased both the likelihood to self-report hypertension and the likelihood to use health care for hypertension. There is no evidence of a test effect for health insurance enrollment. In particular, no evidence is found of a bias in health insurance impact estimates due to the blood pressure measurements, for any of the outcomes.

Keywords: test effect, survey methodology, biomedical test, blood pressure measurement, health insurance, Tanzania.

JEL: C83, C93, I12, I13, O15.

[☆]I am grateful to the attendees of the 2015 NEUDC conference, Joachim de Weerd, Respichius Mitti, Daniëlla Brals, Marleen Hendriks, Heleen Nelissen, Constance Schultsz, Chris Elbers, Wendy Janssens, Menno Pradhan, Melinda Vigh, Jacques van der Gaag, and Martin Wiegand for useful comments. Many thanks to Economic Development Initiatives Ltd. and the Amsterdam Institute for Global Health and Development for excellent collaboration in survey design and data collection. I would like to thank the Kilimanjaro Native Co-operative Union, MicroEnsure, PharmAccess Foundation, and the Health Insurance Fund for ample assistance with fieldwork preparations. The Health Insurance Fund funded the data collection and has made the data available for this research, for which I am grateful. Finally, I thank the Amsterdam Institute for International Development for providing the funding needed to write this manuscript.

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1. Introduction

As developing countries generally have no population-wide administrative health care data, health economics research requires population representative data to be collected through surveys. These often include self-reported morbidity indicators as measures of health status.

5 However, these subjective health measures suffer from misreporting due to respondents' lack of knowledge of their true health status (Sen, 2002), respondents' tendency to report a socially desirable health status (Latkin and Vlahov, 1998; Adams et al., 2005), or because of recall bias (Das et al., 2012). Instead, researchers can opt to collect objective health measures by including biomedical tests in the survey, such as anthropometric measurements, blood
10 pressure measurements, and blood or saliva tests. While this allows more precise knowledge of a respondent's true health status, biomedical testing in surveys has its own drawbacks. The test may give false positives or false negatives (Banoo et al., 2008), there may be non-response bias due to test refusal (e.g. Reniers and Eaton, 2009; Janssens et al., 2014), and test results can even be faked by interviewers (Janssens et al., 2010).

15 The current paper is about another potential drawback of biomedical testing in surveys, namely its ability to change a respondent's future health care seeking behavior. Such change in behavior could effectively cause a carefully chosen representative sample to cease being representative after the survey. In a panel setting it may bias research outcomes—such as impact estimates of a health care intervention.¹

20 It is well known that being surveyed can change behavior. Knowledge of being observed in the scope of an impact evaluation can cause behavioral change in both the treatment and control group, called the Hawthorne and John Henry effect, respectively (Duflo et al., 2007). The so-called question-behavior effect, which includes self-prophecy and mere-measurement effects, occurs when people are asked questions about future behavior and consequently change
25 their actions in line with their answers (Sherman, 1980; Feldman and Lynch, 1988; Sprott et al., 2006; Dholakia, 2010). The survey effect occurs when asking questions about a certain subject causes a reminder or salience shock that makes people act more responsibly afterwards (Bridge et al., 1977; Zwane et al., 2011; Crossley et al., 2014; Axinn et al., 2015).

This paper concerns itself with unintended behavioral changes due to biomedical testing
30 in a survey setting. Disclosing previously unknown information about one's health status closes an information gap, raising awareness of one's true health status, rather than being

¹Note that there may be a positive side to this behavioral change as well. If receiving biomedical measurements causes a rise in health insurance demand among the poor, which has proven notoriously difficult to achieve in developing countries (Gwatkin et al., 2004; Victora et al., 2004; Gwatkin and Ergo, 2011), large-scale biomedical testing may get us one step closer to universal health coverage.

a reminder of something already known. This effect, which I will call “test effect”, is thus intrinsically different from those previously mentioned. It is expected to occur when health implications of a revealed test result can be mitigated by behavioral change, such as health care use or healthy living.² Non-biomedical tests can fall into this category as well if revealing the test result closes an information gap on a personal health related issue. For example the quality of a household’s drinking water directly affects the health of its members, and receiving information that the water is contaminated may lead to better hygiene and water management practices. *Rapid* biomedical tests—where the interviewer can see the test result—are special, because the interviewer is obliged to report the test result to the respondent for ethical reasons. If notification of the test result changes the respondent’s behavior, and subsequently sample representativeness, such rapid biomedical tests can have unintended (and unwanted) consequences for research.

Several authors have rigorously analyzed how behavior changes when individuals learn their HIV status (Thornton, 2008, 2012; Delavande and Kohler, 2012; Gong, 2015) or their household’s water quality (Jalan and Somanathan, 2008; Davis et al., 2011; Luoto et al., 2011; Hamoudi et al., 2012) in the scope of a survey. These authors generally find heterogeneous behavioral effects of a “good” versus a “bad” test result.³ Furthermore, Gong (2015) observes that the behavioral response occurs when the test result reveals unexpected information, i.e. that which is contrary to prior beliefs. However none of the mentioned experiments use rapid tests—thus notification of the test result was optional in principle—and none look at how the behavioral change due to testing may affect impact estimates of a related intervention.⁴ Evidence is lacking on these unintentional test effects in particular.

Evaluating a health insurance intervention in rural Nigeria, Hendriks et al. (2014) seem to detect a test effect with blood pressure measurement—a rapid test, and the subject of this paper. They note that individuals who had high measured blood pressure at baseline—an indication of hypertension—were more likely to use anti-hypertensive drugs, and had lower average blood pressure two years later. This was observed both in the insurance intervention and in the control area—where the health insurance was not available. However, for lack of a test control group, the authors could not assess whether the baseline measurements caused this behavioral change or whether they biased the health insurance impact estimates.⁵

²Height measurement for example is not expected to change behavior.

³See Appendix A for a short summary of these papers.

⁴Zwane et al. (2011) did explore such unintended *survey* effects. They found that frequent surveying biased the impact estimate of protected water sources on child diarrhea. They however detected no bias in health insurance price sensitivity due to surveying.

⁵Tarozzi et al. (2014) encounter an unintended test effect in rural India, where a rapid diagnostic malaria test was included in the baseline survey. Receiving an offer of insecticide treated bednets approximately four months later, demand was observed to be higher in households where at least one member had tested positive for malaria at baseline. Without a test control group however, the causal effect of the test could not be identified.

In the current paper these effects can be identified, in the scope of an impact evaluation of a voluntary health insurance scheme in the Kilimanjaro region of Tanzania. It uses panel data from two extensive household surveys in 2013 and 2015, where both socioeconomic and health related information were collected. In the first survey 80% of households were randomly
5 assigned to receive blood pressure measurements. Complying with ethical standards, the survey medical officer, upon measuring a respondent’s blood pressure, informed the individual of the result. In case of high blood pressure in two out of three measurements, i.e. a “bad” test result, the respondent was warned of the cardiovascular risk of high blood pressure and advised to seek medical care.⁶ Half of the sample gained access to the health insurance
10 program—which included hypertension treatment— five to nine months after the baseline survey, in July–October 2013.

With these data it is possible to assess the effect of the blood pressure measurements on hypertension awareness and related health care seeking behavior, including take-up of health insurance, in a panel household survey setting. The effect of a good and a bad test result
15 on these outcomes can be differentiated, and the bias in the insurance intervention impact estimates for these outcomes can be estimated. This bias is expected to occur if the blood pressure test alters behavior differently in the presence of the novel insurance program than without. Finally, since behavioral response to the test is expected to differ by subjective beliefs of one’s own hypertension status, these differential effects will be identified as well.
20 To the author’s knowledge, this paper is the first to rigorously analyze unintended effects of biomedical testing in surveys.

The paper will proceed as follows. The next section will provide background information on the research population and the health insurance intervention. The experiment is described in Section 3, together with the data collection. Descriptive statistics are given in Section 4. In
25 Section 5 the model is presented, followed by the econometric analysis in Section 6. Section 7 concludes.

2. Health insurance intervention

The background of the surveys is an impact evaluation of the so-called KNCU Health Plan, a voluntary subsidized health insurance scheme in a rural part of the Kilimanjaro region
30 of Tanzania—on the slopes of mount Kilimanjaro. It was funded by the Health Insurance Fund (Dutch NGO), and implemented by the PharmAccess Foundation (Dutch NGO) and MicroEnsure (an insurance company operating i.a. in Tanzania). This health insurance

⁶Determining whether someone has *hypertension*, i.e. chronically high blood pressure, is possible after high blood pressure is measured multiple times over the span of several days. During the baseline survey individual blood pressure was measured on the same day, which is insufficient for diagnostic purposes.

was offered to small scale coffee farmers who are active members of the Kilimanjaro Native Cooperative Union (KNCU), and their household members.⁷ Insured individuals could go with their insurance card to a designated primary health facility in their vicinity—usually an upgraded faith based dispensary—where they would receive free outpatient treatment. Referrals to a district hospital were covered for pregnancy related complications only. The insurance was available at the household—rather than individual—level, and was on an annual basis.⁸ The annual co-premium had to be paid in advance in cash and was priced on a sliding scale, ranging from 12,000 Tanzanian Shilling (TZS) for one person, to 45,000 TZS for 9–12 people per household.⁹

KNCU is organized into about 90 so-called primary societies (PSs), which are spread throughout the Kilimanjaro region (KNCU). Because the KNCU Health Plan was gradually expanded by PS since April 2011, the insurance intervention and control groups consisted of several PSs in rural districts Hai, Moshi Rural, and Rombo: five PSs in the insurance intervention group, and four in the control group. These were chosen such that they were far enough apart to prevent spill-overs from the intervention to the control group, and such that they were similar at baseline on key characteristics such as access to health facilities, altitude, distance to Moshi town, type of coffee grown, and in which district they were located. A map of the insurance intervention and control area is shown in Figure C1 of the Appendix.¹⁰

Approximately seven months after the baseline survey the KNCU Health Plan was offered to the insurance intervention group in districts Hai and Moshi Rural only. It proved not possible to introduce the insurance in Rombo district, which is why subsequently the insurance intervention and control primary society in this district were excluded from the KNCU Health Plan impact evaluation—and from the follow-up survey.¹¹

There is heterogeneity in the KNCU Health Plan roll-out between the Hai and Moshi Rural PSs (see Table C1 of the Appendix). The two insurance intervention PSs in Hai (Moshi Rural) district received access to the insurance in June–July (October) 2013, and had an upgraded

⁷Based on interviews with local village leaders, approximately two out of five households in the study area are estimated to be such “KNCU households”, ranging from 10% to 89%, depending on the village.

⁸Note however that it was difficult for the insurer to check how many individuals resided in the household at the time of insurance, because people would claim that certain household members had moved out since the census (conducted in November 2012 to January 2013, see Appendix B). In households that enrolled in the KNCU Health Plan, on average 84% of all household members who did not move out since the baseline survey were covered by the insurance at first enrollment. 63% of these households were fully insured, i.e. all household members who did not move away since baseline were insured. There is no clear negative relationship between household size and the percentage of insured individuals per household: the correlation coefficient, after removing one exceptionally large household with 11 individuals, is -0.16 ($p=0.170$).

⁹Since the full premium was 14,000 TZS per person, this implies a subsidy level of 14% up to 73%, depending on household size.

¹⁰For more detailed information on the insurance intervention and control group choice and similarities at baseline, see AIID and AIGHD (2013).

¹¹This loss of sub-population is not expected to have caused sample selection, since both the choice of insurance intervention and control group as well as the sampling was stratified by area (Rombo vs. non-Rombo district), see AIID and AIGHD (2013) and Appendix B.

(non-upgraded) dispensary as primary KNCU Health Plan facility. Furthermore, in Hai district the KNCU Health Plan could be renewed after 12 months, while the two Moshi Rural primary societies received free extension until the end of 2014, after which re-enrollment was not possible. Namely, in that district, the KNCU Health Plan was replaced by a new insurance since the start of 2015, as will be described in this section’s final paragraph. Heterogeneous test effects between the two districts will be explored in Section 6.

The study population in the districts Hai and Moshi Rural is quite poor, with median per capita consumption at 2000 TZS or 1.25 USD (3.13 USD at PPP) at baseline.¹² The full KNCU Health Plan per capita premium thus amounts to one week of median per capita consumption. At baseline, 10.8% of the survey sample had health insurance; 8.5% were insured by the National Health Insurance Fund (NHIF), and 2.1% by the Community Health Fund (CHF). CHF is a community based health insurance, available to the full district population, and managed by the district government. Its co-premium is 10,000 TZS per household per year, and covers up to six household members. Insured individuals are entitled to outpatient treatment in public health facilities in the district. However, treatment is not always free in practice. NHIF is the government health insurance, mandatory for government employees (NHIF). The co-premium is 3% of income, and is half of the full premium. It covers all health care in public, and selected private, health facilities. The spouse and up to four legal dependents are eligible as beneficiaries. NHIF is available to informal sector employees for 987,500 TZS (\approx 617 USD) per family per year, more than the median annual per capita consumption in the research area (730,600 TZS). The KNCU Health Plan was introduced to provide high quality health care at a low price, hypothesizing that this was not available to the target population—outside of the public sector. Note that hypertension treatment is included in all three insurance schemes.

In Moshi Rural district, the KNCU Health Plan joined with CHF as of 1 January 2015, in a public-private partnership with the local governments of these districts. It was re-named *improved* Community Health Fund (*i*CHF), and is now available to the full district population. Note that the insurance control group is located in Hai district. At the time of the follow-up survey the marketing of *i*CHF had not yet reached the research population—only one household in the treatment group had heard of it, and three in the control group—and none of the surveyed individuals were insured by *i*CHF at the time of the follow-up survey in March 2015. In May of that same year—one month after completion of all data collection—*i*CHF became available to the Hai district population as well.¹³

¹²Source: baseline survey. 1 USD \approx 1,600 TZS in February 2013 (Oanda). 1 USD \approx 0.4 USD at Purchasing Power Parity (PPP) in Tanzania in 2013 (World Bank).

¹³Siha was the first Kilimanjaro district where *i*CHF started, in November 2014.

3. Experimental design and data collection

The baseline survey was conducted between 25 January to 6 March 2013 by Economic Development Initiatives (EDI) Ltd., a Tanzanian survey firm. Six teams totaling 24 interviewers and seven health officers conducted household interviews and biomedical tests, respectively, in 1,500 KNCU households; half of which belonged to the insurance intervention group. The random sample was stratified at the sub-village level.¹⁴ Interviewers were extensively trained to introduce themselves as being from an independent survey firm, working together with two Amsterdam research institutes—the Amsterdam Institute for International Development (AIID) and the Amsterdam Institute for Global Health and Development (AIGHD)—performing research on health insurance. The KNCU Health Plan was not mentioned. To check whether respondents still anticipated that the survey was part of KNCU Health Plan related research, one of the open questions posed was—after explaining the concept of health insurance—which health insurances respondents had heard of. Only 6% of households had heard of the KNCU Health Plan—8% in the insurance intervention area, and 4% in the insurance control area—lessening concerns of an anticipation effect.

The household questionnaire was conducted in Swahili, the lingua franca of Tanzania,¹⁵ using computers with the specialized survey software Surveybe. It was very extensive, containing sections on education, work, consumption, household assets, gifts and loans, coffee production, risk and time preferences, self-reported health status, health care expenditures, and health care seeking behavior. All individuals, including those not assigned to receive blood pressure measurements, were asked to give written consent for the biomedical part of the interview.¹⁶ Complying with ethical standards, without this consent biomedical measurements and biomedical questions were fully excluded from the interview.

Furthermore 80% of sampled households were randomly assigned for all adult household members to receive blood pressure measurements, stratified by sub-village.¹⁷ This was done by the survey medical officers using the OMRON M6 Comfort digital blood pressure device. They were extensively trained by medical doctors from AIGHD to be able to perform the measurements according to the highest standards. To signal professionalism of the procedures,

¹⁴See Appendix B for more sampling details.

¹⁵Only for six households translation was needed to Chagga, the area's tribal language.

¹⁶Either the caretaker or the household head was asked to give consent for minors.

¹⁷Individuals between 12 and 59 years of age in the same subset of households were also assigned to receive lung function measurements. However, since it turned out extremely difficult for respondents to correctly perform these lung function measurements—one needed to blow three times into a machine with considerable strength—only 34% successfully completed the procedure. Furthermore, only 5% of those individuals (2% of the total) showed signs of obstructive pulmonary disease, and were advised to seek medical care for this condition. This is why lung function measurements will be excluded from the analyses. Subsequent results are robust to exclusion of individuals with signs of obstructive pulmonary disease at baseline. Furthermore, it should be noted that all consenting individuals in all surveyed households received anthropometric measurements. Because individuals were not warned of health issues concerning anthropometrics, a behavioral response from these measurements is not expected.

the survey medical officers were dressed in doctors' white coats. If an individual had high blood pressure (systolic blood pressure ≥ 140 mmHg or diastolic blood pressure ≥ 90 mmHg) in two out of three measurements, the medical officer would point out the health risks of high blood pressure with the help of a leaflet especially designed for the survey, and would advise the person to visit a health care professional for additional testing and treatment. The leaflet, written in Swahili, was finally handed to the respondent for their information.

Two years later, in March 2015, EDI Ltd. returned to interview the same households. Because Rombo district was not anymore included in the follow-up survey (see Section 2), the sample was reduced from 1,500 to 1,000 households. The follow-up survey questionnaire was less extensive than that of the baseline, but included questions on self-reported health status, health care utilization, health expenditures, and detailed health insurance questions—including health insurance status one year before the follow-up survey, in March 2014.

Medical ethical clearance for both survey rounds was received from the Tanzania National Institute for Medical Research (NIMR). The Tanzania Commission for Science and Technology (COSTECH) gave general research clearance.

4. Data description

The 1,000 baseline households from districts Hai and Moshi Rural had a total of 4,122 household members, out of which 2,530 were adults (BP test: 2,038; No BP test: 492)¹⁸. Out of all adults 2,159 (85%) consented for the biomedical part of the survey (BP test: 1,738 = 86%; No BP test: 421 = 85%). From all who received the blood pressure measurements—namely all consenting adults, except 43 individuals who did give consent but were not available for the measurements—588 (35%) had high blood pressure in at least two out of three measurements, i.e. a bad test result.

Out of these 1,000 baseline households 34 could not be reached at follow-up (BP test: 29 = 4%; No BP test: 5 = 3%), out of which 19 (56%) were in the insurance intervention group. These households had either moved, were unavailable or had refused an interview. Out of the baseline consenting adults 1,800 (83%) were still household members at follow-up (BP test: 1,444 = 83%; No BP test: 356 = 85%) and 1,536 (85%) consented at follow-up (BP test: 1,243 = 86%; No BP test: 293 = 82%).¹⁹ These 1,536 individuals are used in the analyses.

Individuals who left the household or did not give consent at follow-up were on average 12 years younger, more likely to be male, healthier, better educated, more likely to have worked in the past year, and more likely to have experienced a financial health shock at baseline,

¹⁸“BP test” are the individuals in households that were randomly *assigned* to receive blood pressure measurements, and “No BP test” are those in households that were not assigned to receive the measurements.

¹⁹Only 25 of those had a missing test result at baseline.

compared to those in the research sample.²⁰ The baseline characteristics of those not available for the (biomedical part of the) follow-up survey were however balanced between the blood pressure test and control group.²¹

Table 1: Means of baseline variables, by blood pressure test assignment

	BP test Mean (N=1243)	No BP test Mean (N=293)	ΔMean p-value
<i>Main</i>			
Insurance intervention area	0.48	0.54	0.186
Self-reported HT	0.23	0.26	0.189
BP check - past 12 months	0.34	0.37	0.467
Consult for HT - past 12 months	0.16	0.19	0.229
Any health insurance	0.15	0.13	0.367
<i>Socio-economic characteristics</i>			
Age (years)	54.8	57.7	0.016*
Female	0.61	0.59	0.476
Married ^a	0.69	0.70	0.686
Worked ^b - past 12 months	0.21	0.17	0.073 ⁺
Religion: Christian	0.96	0.96	0.887
Mother tongue: Chagga	0.99	0.97	0.250
Educ ^c : None	0.09	0.13	0.094 ⁺
Educ: Less than primary school	0.31	0.32	0.827
Educ: Primary school	0.54	0.49	0.163
Educ: More than primary school	0.06	0.06	0.853
<i>Self-reported illness/ injury</i>			
Chronic illness	0.41	0.46	0.157
Acute illness / injury - past 12 months	0.50	0.52	0.553
Hospitalization - past 12 months	0.07	0.08	0.546
<i>Household characteristics</i>			
Annual consumption ^d - PC (TZS/1,000)	860	872	0.742
Financial health shock - past 12 months	0.37	0.39	0.632
Household size	4.40	4.09	0.073 ⁺
#Young children in HH (age < 5)	0.20	0.12	0.005**
#Elderly in HH (age ≥ 60)	0.70	0.78	0.199
#Reproductive age women (15–45) in HH	0.51	0.39	0.042*

Note: The table shows statistics for all adults who gave consent for the biomedical part of both surveys (questions and measurements). Means are weighted and p-values clustered at the household level, in accordance with the sampling method. BP=blood pressure; HT= hypertension; HH=household; PC=per capita; ^aIncludes mono- and polygamous marriage; ^bApart from household chores and family farming; ^cHighest completed educational level; ^dOne outlier excluded.

⁺ p<.10, * p<.05, ** p<.01, *** p<.001.

Table 1 compares, for all adults who consented for the biomedical part in both surveys, the baseline means of the outcome variables and other control variables, by blood pressure test assignment. Out of the 22 baseline characteristics, three are not balanced at the 5% level, all age related. Individuals not assigned to receive blood pressure measurements are found to be on average three years older than those assigned to receive the measurements.²² Since hypertension risk increases with age, this must be taken into account in the analyses. Consequently observations will be reweighted according to age in the robustness checks.

²⁰See Table C2 of the Appendix.

²¹See Table C3 of the Appendix.

²²Looking at the mean age separately for the insurance intervention and control group it turns out that the imbalance is small and not statistically significant in the intervention area (BP test: 54.7 years; No BP test: 55.7 years; p=0.479). However the difference in means is quite severe in the insurance control area (BP test: 54.9 years; No BP test: 60.1 years; p=0.008). When regressing blood pressure test assignment on age-group dummies, insurance intervention area, and their interactions, it becomes clear that the oldest group (70 years and older) is overrepresented in the *non-tested* individuals of the *insurance control* area.

Baseline characteristics are balanced between the insurance intervention and control area, except for high blood pressure prevalence, which is almost one and a half times higher in the insurance control area than in the insurance intervention area (Table C4).²³ Since observed test results can be controlled for, this imbalance is not problematic for analyses.

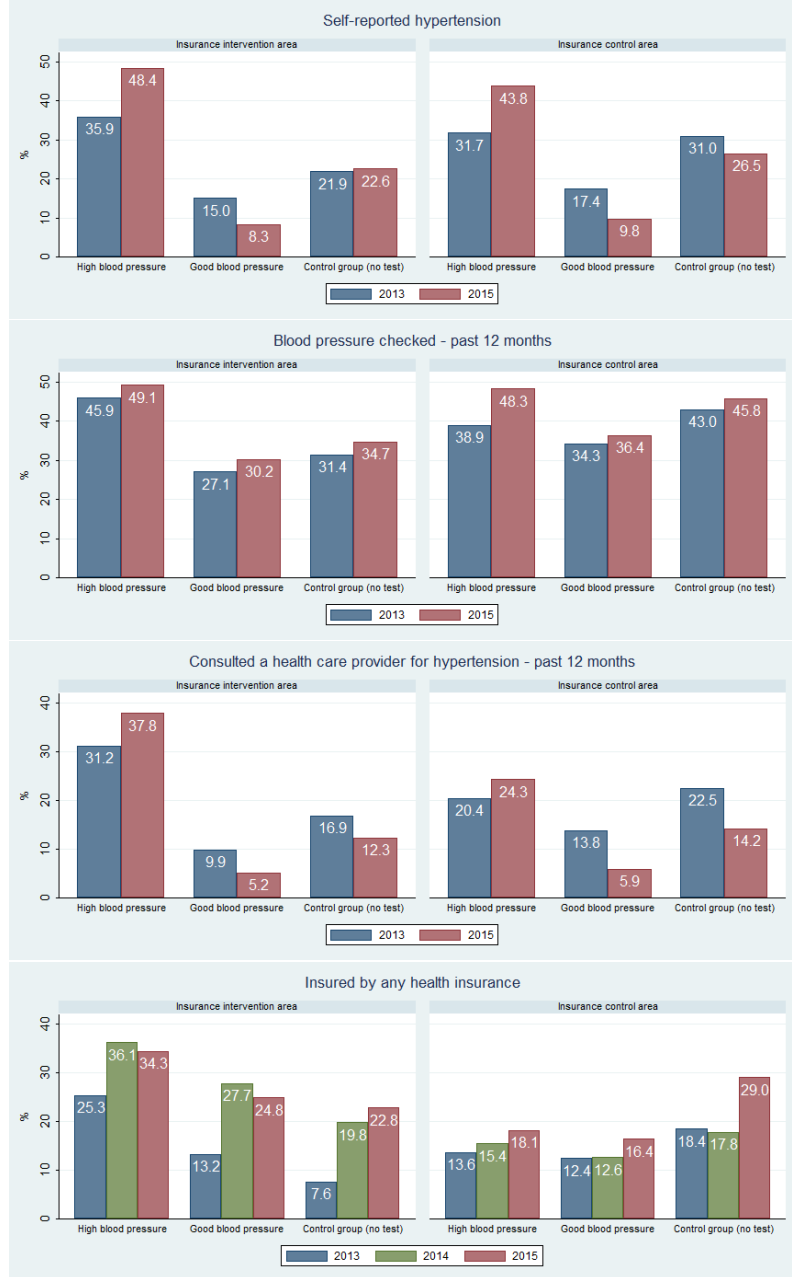


Figure 1: Outcomes by test result and insurance area (N=1511). Observations in the insurance intervention [control] area: Total: 789 [722]; High blood pressure: 209 [258]; Good blood pressure: 425 [326]; Control group (no test): 155 [138].

²³This is not due to age imbalance, since regressing the insurance intervention area on age-group dummies for those who received blood pressure measurements yields a p-value of 0.499 for the joint significance F-test.

Figure 1 shows the means of the outcomes of interest—self-reported hypertension, blood pressure checks in the past 12 months, health care provider consultations for hypertension in the past 12 months,²⁴ and health insurance enrollment—over time, disaggregated by test result and by the insurance intervention and control area.

5 The general picture from Figure 1 is that—as expected—already at baseline, individuals with a bad test result were more likely to self-report hypertension, more likely to have utilized health care for hypertension in the past year, and more likely to have health insurance, compared to those with a good test result at baseline. Baseline means of individuals from the test control group—residing in households not assigned to receive blood pressure measurements—
10 are a priori expected to be in between those with a good and a bad test result. This can indeed be seen in the insurance intervention group, for all outcomes except for the insurance prevalence, which is lower than expected for those in the test control group. However, in the insurance control group, the baseline means are as high or higher than those of individuals with high blood pressure at baseline. This is an indication that hypertension prevalence in
15 the test control group may very well be higher than that of the population, which is in line with the earlier observation that non-tested individuals in the control group are on average older than those assigned to be tested (footnote 22). Reweighting according to age should control for this imbalance.

From the first set of graphs in Figure 1, it can be seen that after two years, as expected, the
20 percentage of self-reported hypertension increased (decreased) in both the insurance intervention and control area among those who had a bad (good) test result at baseline. Among those who were not assigned to receive blood pressure measurements the percentage of self-reported hypertension surprisingly decreased in the insurance control area (statistically insignificant at a 10% level). As can be seen from the second set of graphs, there is a small increase
25 between baseline and follow-up in the percentage of individuals who had their blood pressure checked in the past 12 months, for all subgroups.²⁵ The third set of graphs, showing the percentage of individuals who consulted a health care provider for hypertension in the past 12 months, displays a pattern similar to self-reported hypertension: an increase over time for those with high blood pressure at baseline, and a decrease for those with good blood pressure.
30 For the test control group there is an unexpected decrease in likelihood of consultations for hypertension over time, for both the insurance intervention and control group.

The final set of graphs in Figure 1 shows the percentage of individuals enrolled in any health insurance scheme. As mentioned in Section 2, already at baseline almost 11% of the

²⁴Note that the follow-up survey occurred 25–26 months after the baseline survey. Thus if during the follow-up survey someone reports to have utilized health care in the past 12 months, this visit occurred at least one year after the baseline survey.

²⁵The household survey blood pressure test was explicitly excluded when phrasing this survey question.

surveyed individuals had health insurance (2% CHF, 9% NHIF). In the insurance intervention area there is a rise in health insurance in 2014 and 2015, irrespective of the blood pressure measurement. In the insurance control group there is hardly any change in health insurance enrollment between 2013 the baseline survey and 2014, but there is a rise in 2015, for all three groups. The rise in insurance take up in the insurance intervention group is mainly due to uptake of the KNCU Health Plan,²⁶ while in the control group it is CHF insurance take up that increased between baseline and follow-up (Figure C2).²⁷ There is no indication that those with a bad test result are more likely to take up health insurance than those with a good test result. This is to be expected with household level insurance, such as the KNCU Health Plan and CHF: healthy household members become insured along with the sick.

Finally, the previously mentioned age imbalance between the test and no test group is problematic if the test effect is heterogeneous by age. There is no indication of this however (Figure C3).

5. Model

For the k th outcome of interest, Y_k , the test effect—irrespective of its result—is captured by parameter β_k in the following difference-in-differences individual fixed effects model:²⁸

$$y_{kit} = \beta_k(M_i \times T_t) + \gamma_k T_t + \delta_{ki} + \epsilon_{kit}, \quad (1)$$

where y_{kit} is the k th outcome value for individual i at time t , which is equal to 0 at baseline (January–February 2013), and 1 at follow-up (March 2015)—except for health insurance take-up. Namely, when considering health insurance take-up t will be equal to 1 in March 2014 (one year before the follow-up survey), because this reflects the *first-time* KNCU Health Plan choice.²⁹ M_i is a dummy equal to 1 if individual i was assigned to receive blood pressure measurements at baseline,³⁰ and 0 otherwise. The time dummy T_t is 0 at baseline and 1

²⁶ Note that the KNCU Health Plan was introduced there in July–October 2013. Because one KNCU Health Plan insurance package lasts for one year—except in the Moshi Rural insurance intervention area, where it was extended to 15 months (Table C1)—this means that the (March) 2014 insurance status reflects the first-time KNCU Health Plan insurance choice. The (March) 2015 insurance status does *not* reflect the KNCU Health Plan second-time insurance choice well, because there was no second-time enrollment possible in Moshi Rural due to the introduction of the new *i*CHF insurance (Section 2).

²⁷ NHIF insurance shows no substantial changes over time, which is unsurprising for a government employee insurance that is extremely expensive for the informal sector (see Section 2).

²⁸ Equation 1 is a linear probability model, since all outcomes of interest are binary. A non-linear model is in theory preferred, e.g. the “changes-in-changes” model by Athey and Imbens (2006), as suggested by Blundell and Dias (2009). However, as shown by Angrist and Pischke (2009, p. 197-205) and Wooldridge (2002, p. 472), estimated marginal effects and standard errors from a non-linear model are in general similar to those of its linear counterpart.

²⁹ See footnote 26. The insurance status in March 2015 unfortunately does not reflect the second-time KNCU Health Plan choice well, which is why it will not be used in the analyses.

³⁰ This is thus excluding individuals who did not give consent for the biomedical part of the survey, and including the 25 individuals who gave consent, but did not receive the test.

when $t = 1$, such that parameter γ_k captures the common time trend. The individual time invariant characteristics are captured in the fixed effect δ_{ki} , and ϵ_{kit} is the error term.

Because the blood pressure measurement is combined with notification of the result—it is expected is necessary to obtain ethical clearance for rapid biomedical testing in surveys—it is expected
 5 that the test *result* will drive behavior, rather than the act of testing.³¹ To differentiate between the effects of a good and a bad test result M_i is split up in $M_i = G_i + B_i$, such that G_i and B_i are dummy variables for a good and bad test result, respectively.³² The parameters of interest are then β_{kg} and β_{kb} in the following equation:³³

$$y_{kit} = \beta_{kg}(G_i \times T_t) + \beta_{kb}(B_i \times T_t) + \gamma'_k T_t + \delta'_{ki} + \epsilon'_{kit}. \quad (2)$$

The test effect is heterogeneous by test result iff $\beta_{kg} \neq \beta_{kb}$. A priori it is expected that
 10 $\beta_{kb} \geq 0$ for all outcomes: individuals hearing that they have high blood pressure will become more aware of this, will utilize health care more often, and are more likely to take up health insurance afterward. Symmetrically, it is expected that $\beta_{kg} \leq 0$ for all outcomes, except health insurance uptake, which could still be attractive, since it protects against many types of health risks—not only hypertension. Moreover, in case that health insurance is at the
 15 household level, such as with the KNCU Health Plan and CHF, healthy household members are expected to take up health insurance along with the sick.

Adding to equation 1 interaction terms with dummy D_i , denoting the *insurance* intervention area, i.e. the insurance program intent to treat (ITT), gives:

$$y_{kit} = \tilde{\beta}_k(M_i \times T_t) + \tilde{\eta}_k(M_i \times D_i \times T_t) + \tilde{\theta}_k(D_i \times T_t) + \tilde{\gamma}_k T_t + \tilde{\delta}_{ki} + \tilde{\epsilon}_{kit}. \quad (3)$$

Parameter $\tilde{\theta}_k$ captures the insurance intervention ITT on the outcome, expected to be
 20 non-negative for all outcomes, and distinctly positive for insurance uptake. The parameter of interest in equation 3 is $\tilde{\eta}_k$, since it represents the bias from the blood pressure test

³¹Note, since there is no intermediate control group which receives the test, but does not learn its outcome, the effect of administering the test only—without letting the respondent know its outcome—cannot be determined.

³²This split is impossible for the 25 individuals in the treatment group who gave consent but were not tested, thus their G_i and B_i value is set to be missing.

³³Controlling for a bad test result in both the treatment and the test control group, as in Jalan and Somanathan (2008), is not possible due to lack of a control group that does not learn the test result (footnote 31). Because the testing treatment was randomly assigned, the a priori expectation is that the bad test result would have been balanced between the treatment and control group, if the latter would have received the test. From the descriptive results in Table 1 it seems however that the high blood pressure prevalence in the test control group is likely higher than that of the treatment group, since the latter individuals are on average younger. Fixed effects regression should control for this imbalance as long as the treatment effect is not heterogeneous by age. As mentioned at the end of Section 4, there is no indication of heterogeneity by age in the outcomes of interest. Any remaining age imbalance will be corrected for once the regressions are reweighted by age in the robustness analyses.

(disregarding its result) in the insurance intervention ITT estimate.³⁴ Note finally that $M_i \perp D_i$, because the randomization of the testing treatment (M_i) was equally divided between the insurance intervention and control area by design.

The bias may heterogeneous for those with a good versus a bad test result. Splitting M_i into $G_i + B_i$ in equation 3 gives:

$$y_{kit} = \tilde{\beta}_{kg}(G_i \times T_t) + \tilde{\beta}_{kb}(B_i \times T_t) + \tilde{\eta}_{kg}(G_i \times D_i \times T_t) + \tilde{\eta}_{kb}(B_i \times D_i \times T_t) + \tilde{\theta}'_k(D_i \times T_t) + \tilde{\gamma}'_k T_t + \tilde{\delta}'_{ki} + \tilde{\epsilon}'_{kit}, \quad (4)$$

where the parameters of interest are $\tilde{\eta}_{kg}$ and $\tilde{\eta}_{kb}$, giving the bias in the insurance intervention ITT estimate from a good and bad test result, respectively. Here $\tilde{\eta}_{kb}$ is expected to be non-negative for the two health care utilization outcomes, as well as for insurance uptake. Namely for individuals with a bad test result at baseline, access to the KNCU Health Plan is a priori expected to further increase health care use and insurance uptake, i.e. it is expected to amplify the effect of a bad test result in absence of the KNCU Health Plan. However, $\tilde{\eta}_{kb}$ may be (close to) zero because (i) household level insurance reduces adverse selection, (ii) because insured individuals do not increase health care use (no moral hazard), or (iii) because when the KNCU Health Plan became available in the insurance intervention area (five to nine months after the baseline survey) people had already forgotten the results of the baseline survey measurements. In case of self-reported hypertension the sign of $\tilde{\eta}_{kb}$ is unclear, since the baseline high blood pressure may have been acute instead of chronic (see footnote 6). No additional effect of the insurance intervention is expected in case of a good test result, thus $\tilde{\eta}_{kb}$ is expected to be zero.

Finally, the behavioral response to the test is expected to differ conditional on prior beliefs of hypertension status, i.e. self-reported hypertension at baseline. In particular, as in Gong (2015), it is expected that the behavioral response will be strongest in those “surprised” with a good/bad test result, than in those who are “unsurprised”. These heterogeneous effects will be explored in the next section as well.

³⁴This can be seen as follows. Defining $\Delta y_{ki} := y_{ki1} - y_{ki0}$, omitting the subscript k , and writing M_i (D_i) instead of $M_i \times T_t$ ($D_i \times T_t$) for ease of notation, gives:

$$\begin{aligned} E(\Delta y_i | M_i, D_i) &= C + E(\Delta \epsilon_i | M_i, D_i), \\ &= C + M_i D_i E(\Delta \epsilon_i | M_i = 1, D_i = 1) + (1 - M_i) D_i E(\Delta \epsilon_i | M_i = 0, D_i = 1) \\ &\quad + M_i (1 - D_i) E(\Delta \epsilon_i | M_i = 1, D_i = 0) + (1 - M_i) (1 - D_i) E(\Delta \epsilon_i | M_i = 0, D_i = 0), \\ &=: C + M_i D_i E_{11} + (1 - M_i) D_i E_{01} + M_i (1 - D_i) E_{10} + (1 - M_i) (1 - D_i) E_{00}, \\ &= C + E_{00} + M_i (E_{10} - E_{00}) + D_i (E_{01} - E_{00}) + M_i D_i [(E_{11} - E_{10}) - (E_{01} - E_{00})], \end{aligned}$$

where C is a constant. The term $(E_{01} - E_{00})$ is the ITT of the insurance program in absence of blood pressure measurements; $(E_{10} - E_{00})$ is the ITT of the blood pressure measurements without the availability of the insurance program; and $[(E_{11} - E_{10}) - (E_{01} - E_{00})]$ is the ITT of the insurance program in the presence of blood pressure measurements, minus the ITT in absence of the blood pressure measurements, i.e. the bias in the insurance program ITT due to the blood pressure measurements. Note finally that the within estimator is equivalent to the first difference estimator up to a constant.

6. Analysis

Because of the random assignment of households into receiving the blood pressure tests at baseline a priori it could reasonably be assumed that the time trend of the outcome of interest in absence of the baseline test would be the same between the test and no-test group. Then, using a two period balanced panel, the above equations could be consistently estimated with the fixed effects estimator (Cameron and Trivedi, 2009).

However, from Section 4 it was clear that the randomization was unlucky in terms of age—a risk factor for hypertension—since the test group is on average younger than the no-test group. The individual fixed effects estimator is still consistent if the test effect is not heterogeneous by age—of which there is no indication, as also noted at the end of Section 4. In case there is some age heterogeneity, reweighting observations by baseline age-group would solve the issue. This will be done as robustness check.

All subsequent regressions are performed using the command “xtreg, fe” in Stata 11.2 software (StataCorp, 2009). According to the sampling frame, observations are weighted by their sampling probabilities, and errors are clustered at the household level. Only adults who gave consent for the biomedical part of both surveys are included in the regressions.

6.1. Main results

Since the test effect is expected to be heterogeneous by test outcome, and even with opposite signs for three out of four outcome variables, it seems prudent to first look at the results corresponding to equation 2 and 4.

Table 2: Main results, corresponding to equation 2

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
Good BP	-0.053 (0.032)	-0.003 (0.042)	0.001 (0.031)	0.016 (0.034)
High BP	0.140*** (0.039)	0.039 (0.045)	0.113** (0.038)	-0.009 (0.035)
Constant	-0.018 (0.028)	0.029 (0.034)	-0.064* (0.027)	0.063* (0.028)
Observations	3014	3008	3006	3022
$P(\beta_{kg} = \beta_{kb})$	<.001***	0.260	<.001***	0.289

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation “× Time” omitted). BP=blood pressure; HT=hypertension; 12m=12 months;
⁺ $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.

Table 2 shows the estimation results for equation 2. Here evidence is seen of a test effect for self-reported hypertension and consultations for hypertension in the past 12 months. There is a 14 pp increase (significant at the 0.1% level) in self-reported hypertension for those individuals who had a bad test result at baseline, compared to those who were not tested. There is also a 5 pp decrease in self-reported hypertension for those with a good test result.

However, this is not significantly different from zero at a 10% level. Having had a bad test result at baseline increases the probability to consult a health care provider for hypertension significantly (at the 1% level), by 11 pp. A good test result is found to have no influence on consultations for hypertension. Heterogeneous test effects between a good and a bad test result for these two outcomes are confirmed by the small p-value corresponding to the Wald test of $\beta_{kg} = \beta_{kb}$.

There is however no significant effect of a good or bad test result on visiting a health care provider for blood pressure checks (column 2 of Table 2) or on insurance take up (column 4 of Table 2). While the signs in column 2 are as expected, the test effects in case of health insurance take-up have the opposite sign. Thus Table 2 gives no evidence of adverse selection into health insurance due to the test.³⁵

There is no evidence of any test effect when not differentiating by test result, for any outcome (Table C5). The heterogeneous test effects of a good and bad test result thus seem to cancel each other out.

Table 3: Main results, corresponding to equation 4

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
Good BP	-0.029 (0.048)	-0.010 (0.056)	0.001 (0.043)	0.008 (0.046)
High BP	0.169** (0.056)	0.064 (0.058)	0.121* (0.051)	0.024 (0.046)
Good BP \times Ins.area	-0.044 (0.065)	0.013 (0.083)	-0.000 (0.061)	0.016 (0.067)
High BP \times Ins.area	-0.053 (0.079)	-0.061 (0.089)	-0.006 (0.077)	-0.037 (0.070)
Ins.area	0.055 (0.056)	0.001 (0.067)	0.034 (0.054)	0.128* (0.055)
Constant	-0.048 (0.042)	0.028 (0.044)	-0.082* (0.038)	-0.006 (0.039)
Observations	3014	3008	3006	3022
$P(\tilde{\beta}_{kg} = \tilde{\beta}_{kb})$	<.001***	0.143	0.003**	0.512
$P(\tilde{\eta}_{kg} = \tilde{\eta}_{kb})$	0.891	0.327	0.921	0.281

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation " \times Time" omitted). BP=blood pressure; HT=hypertension; 12m=12 months; Ins.area= Insurance intervention area; ⁺ $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.

Table 3 shows the estimation results corresponding to equation 4. Even more pronounced than in Table 2, a significant positive effect can be seen of a bad test result on self-reported hypertension (17 pp increase, significant at the 1% level) and health care provider consultations for hypertension (12 pp increase, significant at the 5% level). Also similarly to Table 2, there is no significant effect of a good or bad test result on the other two outcomes (at the

³⁵When running regression 4 of Table 2 at the household level (replacing the individual bad test result with a dummy for a bad test result in the household, and similarly replacing the individual good test result with a dummy for only good test results in the household) the signs and significance of the parameter estimates remain the same.

10% level). The insignificant parameter estimates of “Good BP \times Ins. area” ($\tilde{\eta}_{kg}$) and “High BP \times Ins. area” ($\tilde{\eta}_{kb}$), as well as the insignificant Wald test of $\tilde{\eta}_{kg} = \tilde{\eta}_{kb}$ for all outcomes, give no evidence that the test biases the KNCU Health Plan impact estimates. Possibly this is because the KNCU Health Plan is offered at the household level only, thus limiting adverse selection.³⁶ Unsurprisingly, there is a 13 pp impact (significant at the 5% level) of offering the KNCU Health Plan on health insurance enrollment in the insurance intervention area. There is again no evidence of a test effect when not differentiating by test result, for any outcome (Table C6).

6.2. Test effect conditional on prior beliefs

Table 4 shows results corresponding to equation 2, when additionally controlling for self-reported hypertension at baseline (in the levels only). The test effects for the first three outcome variables are now more pronounced than in Table 2. The effect of a good test result on self-reported hypertension is now much larger at -11 pp (significant at the 0.1% level). Additionally, a good test result now has a negative effect on health provider consultation for hypertension, as a priori expected. However this is not significant at the 10% level.

Table 4: Controlling for self-reported hypertension at baseline, corresponding to eq. 2

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
Good BP	-0.116*** (0.027)	-0.039 (0.042)	-0.040 (0.027)	0.018 (0.034)
High BP	0.187*** (0.035)	0.065 (0.044)	0.144*** (0.036)	-0.011 (0.035)
Self-reported HT	-0.635*** (0.031)	-0.359*** (0.038)	-0.419*** (0.039)	0.018 (0.027)
Constant	0.147*** (0.024)	0.122*** (0.035)	0.046* (0.023)	0.059* (0.028)
Observations	3014	3008	3006	3022
P($\beta_{kg} = \beta_{kb}$)	<.001***	0.004**	<.001***	0.236

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation “ \times Time” omitted). BP=blood pressure; HT=hypertension; 12m=12 months; + $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.

Adding additional interaction terms “Good BP \times Self-reported HT” and “Bad BP \times Self-reported HT” gives information on whether the test effect differs in case the result revealed unexpected information, instead of expected information. An interaction term of zero means that there is no significant difference between the two. Table 5 shows that all the interaction terms are not significantly different from zero, except “Good BP \times Self-reported HT” when the outcome is self-reported hypertension—in which case the interaction term is -21 pp (significant at the 5% level). Thus, someone who reported to have hypertension at baseline but

³⁶Running regression (4) of Table 3 at the household level, analogously to footnote 35, gives results similar to those of Table 3.

had a good test result is much more likely to report not having hypertension at follow-up than someone unsurprised with a good test result.

From Table C7 of the Appendix it can be seen that there is again no evidence of bias caused by the test, even when adding self-reported hypertension interaction terms to equation 4.

Table 5: Heterogeneity by self-reported hypertension at baseline, corresponding to equation 2

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
Good BP	-0.074** (0.026)	-0.016 (0.045)	-0.042* (0.020)	0.035 (0.034)
Good BP \times Self-reported HT	-0.208* (0.081)	-0.103 (0.098)	-0.019 (0.098)	-0.068 (0.066)
High BP	0.185*** (0.038)	0.077 (0.050)	0.105*** (0.031)	0.017 (0.034)
High BP \times Self-reported HT	-0.012 (0.081)	-0.049 (0.093)	0.127 (0.099)	-0.098 (0.064)
Self-reported HT	-0.551*** (0.065)	-0.300*** (0.075)	-0.463*** (0.077)	0.082 ⁺ (0.046)
Constant	0.125*** (0.024)	0.107** (0.037)	0.057** (0.019)	0.042 (0.027)
Observations	3014	3008	3006	3022

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation “ \times Time” omitted). BP=blood pressure; HT=hypertension; 12m=12 months; ⁺ $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.

6.3. Robustness checks

Results are robust to sub-village level clustering of standard errors and age-group reweighting. Furthermore, no heterogeneous test effect is found between the Hai and Moshi Rural treatment areas. Robustness check results are not shown here, but are available upon request.

7. Conclusion

This paper provided strong evidence that individual behavior can be substantially influenced by the inclusion of rapid biomedical tests in surveys (“the test effect”). The test—in this case blood pressure measurement—was embedded in an impact evaluation of a voluntary health insurance intervention in the Kilimanjaro region of Tanzania. This allowed to assess whether the test effect differed between the insurance intervention and control areas, something that would bias the insurance (intention to treat) impact estimates.

It was found that measured high blood pressure (a “bad test result”) significantly increased hypertension awareness, as well as the likelihood to consult a health care provider for hypertension. There is also evidence, albeit weaker, that a good test result reduced both the likelihood to self-report having hypertension, as well as the likelihood to consult a health care provider for hypertension. The test effect did not differ between those surprised and those unsurprised with their blood pressure measurement, as in Gong (2015). Unexpectedly, there

was no test effect on health insurance enrollment. Finally, no evidence was found that the blood pressure measurements biased health insurance impact estimates.

Medical clearance procedures mandate that results of rapid biomedical tests are revealed to the individuals who are tested during a survey. Considering the above evidence of behavioral
5 change due to blood pressure measurements, as well as that of earlier papers with HIV and water quality testing (e.g. Thornton, 2008; Gong, 2015; Jalan and Somanathan, 2008; Hamoudi et al., 2012) it is recommended that the relatively easy and cheap method introduced in this paper, i.e. the random exclusion of a small percentage of interviewed individuals from participating in the test, should be routinely adopted in such surveys, to facilitate rigorous
10 testing of whether these tests have biased the outcomes of interest.

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Appendix A Behavioral response to household survey testing in the literature

A.1 HIV testing

Thornton (2008) conducted an experiment in rural Malawi where people were randomly assigned monetary incentives to learn their HIV status after having been tested during a household survey. She finds that sexually active HIV positive individuals who learn their status are three times more likely to buy condoms two months later, compared to those who did not learn their status. She finds no test effect on condom purchases for HIV negative individuals. Based on the same experiment, Thornton (2012) finds no effect of learning one's HIV status on economic behavior two years later. Delavande and Kohler (2012), also using data from this experiment, observe surprisingly that becoming aware of an HIV negative status increases one's subjective beliefs of being HIV positive after two years. Those who learned they were HIV positive self-reported less risky sexual behavior—more condom use and less sexual partners—than HIV positive individuals who did not learn their result.

Using data from experiments in urban Kenya and Tanzania where HIV tests were randomly assigned during a survey, Gong (2015) finds that individuals *surprised* by their HIV positive (negative) result had higher (lower) likelihood to contract gonorrhea or chlamydia—sexually transmitted infections (STIs), and proxies of risky sexual behavior—six months after the experiment, compared to untested individuals with similar prior beliefs about their HIV infection probability. However, no test effect on STI infection was observed for those unsurprised by their test result.

A.2 Water quality testing

Jalan and Somanathan (2008) conducted a household survey in urban India, where households' own drinking water was tested for the presence of fecal bacteria, and a random sample learned their test result. Households initially not purifying their water that received a bad test result—denoting presence of fecal bacteria in their drinking water—were more likely to start purifying their water within two months of learning the result, compared to those not learning their test result. Initially purifying households that received a good test result did not reduce purification.

Davis et al. (2011) use data from a household survey experiment in peri-urban Tanzania where all households received general information on water quality and treatment, and randomly chosen households were notified of their own stored drinking water and/or hand-rinse water quality. Households that received the test results—all of which were bad, i.e. contaminated—were more likely to self-report improved water management and hygiene behaviors, but less likely to improve their actual drinking water or hand-rinse water quality two months later, compared to the control group—which received general information only.

Luoto et al. (2011) tested both source water and households' own water quality in rural Kenya in the scope of a household survey. All households received water purification kits, and a random sub-sample learned that their source water was contaminated. Furthermore, a random sub-sample of the latter additionally received the test result of their household's own
5 stored water—most times (87%) contaminated. The authors find that information provision of source water quality significantly increases water purification efforts two months later. These do not increase further once the household's own water quality is revealed.

In a household survey in rural India a random sample of households learned the quality of their own drinking water and received information on water management practices, while the
10 control group received neither test nor information. Using data from this experiment Hamoudi et al. (2012) find, in a situation where almost 90% of tested households had contaminated drinking water, that water quality testing increased demand for commercial water sources, but had no effect on time-intensive water management practices.

Appendix B Sampling

In November 2012 to January 2013 a census was conducted of all households belonging to active KNCU members of the insurance intervention and control areas in districts Hai, Moshi Rural, and Rombo of the Kilimanjaro region. A random sample of 1,500 households was then
5 selected from the census, stratified by geographic area and insurance intervention.³⁷ Namely 500 households were drawn from the Rombo district and 1,000 from the Hai and Moshi Rural districts (the study population in Moshi Rural district is near the border with Hai district), such that half of the sample in each of these areas was drawn from the insurance intervention area, and the other half from the insurance control area. Because of logistical reasons an
10 additional stratification was made at the smallest administrative unit, the sub-village, such that from each sub-village, in each stratum, approximately the same number of households was randomly drawn. Thus, sampling probability weights are necessary in the analyses.

Additionally, per sub-village, 40% extra households were randomly drawn, to serve as replacement in case a household in the original sample could not be interviewed. Ultimately
15 84 households (5.6%) of the original sample were replaced (equally distributed between the insurance treatment and control group), mostly due to households' unavailability for the interview (39%), followed by: household moved (17%), household listed twice (17%), household refused to be interviewed (16%), household unknown (6%), household no longer exists (6%).

³⁷The census data were furthermore used by the KNCU Health Plan as administrative base for the insurance.

Appendix C Additional tables and figures



Figure C1: Insurance intervention and control KNCU primary societies (PSs) and their villages. *Hai* district is located on the far west side of the map (containing PSs Isuki, Lemira Mroma, Masama Saawe, Machame Nkuu, and Narumu). *Moshi Rural* is the district east of Hai (containing PSs Umbwe Ndoo and Kombo), while *Rombo* is the most eastern district (containing PSs Shimbi and Mkuu Masaseni). For completion, Moshi Urban district is located to the south of Moshi Rural, and is not included in the research. The (control) treatment villages in this figure denote the areas where—at the time of the baseline survey—the health insurance intervention, i.e. the KNCU Health Plan, was *planned* (not) to be offered. Because it proved not possible to introduce the KNCU Health Plan in the Rombo district, the two PSs in Rombo were excluded from the follow-up survey and subsequent analyses. Note that the research population resides on the slopes of mount Kilimanjaro, increasing in altitude towards the north-northwest of the map. Substantial travel time is needed between treatment and control group villages, sometimes even between villages of the same primary society. Lemira Lutheran dispensary is located within the cluster of Isuki and Lemira Mroma villages (north side). Umbwe Parish dispensary is located in the Umbwe Ndoo and Kombo village cluster (southeast side). PSs Masama Saawe, Machame Nkuu, and Narumu have a popular dispensary within their village clusters as well. Source: GPS coordinates collected by EDI Ltd. in November 2012. Map adapted from Google maps.

Table C1: KNCU Health Plan details per primary society (insurance intervention group)

Primary society	District	Primary health facility	Start date	End date
Isuki	Hai	Lemira Lutheran dispensary ⁺	1-Jul-2013	30-Jun-2014*
Lemira Mroma	Hai	Lemira Lutheran dispensary ⁺	1-Aug-2013	31-Jul-2014*
Umbwe Ndoo	Moshi Rural	Umbwe Parish dispensary	1-Oct-2013	31-Dec-2014**
Kombo	Moshi Rural	Umbwe Parish dispensary	1-Oct-2013	31-Dec-2014**
Shimbi	Rombo	n/a	n/a	n/a

Note: ⁺Lemira Lutheran dispensary was upgraded in the scope of the KNCU Health Plan in May–August 2013. *In Hai district the KNCU Health Plan could be renewed for another year after the shown end date. **In Moshi Rural the first insurance period was extended until the end of 2014 (three additional months for free). As of 1 January 2015 the KNCU Health Plan was terminated in this district. Instead, all households in Moshi Rural could then enroll in the new *iCHF* insurance (see Section 2).

Table C2: Attrition: baseline means of sample vs. individuals lost to follow-up

	Sample Mean (N=1536)	Lost to follow-up Mean (N=623)	Δ Mean p-value
<i>Main</i>			
BP test	0.81	0.80	0.841
High BP ^a	0.31	0.21	<.001***
Insurance program area	0.50	0.41	<.001***
Self-reported HT	0.23	0.14	<.001***
BP check - past 12 months	0.35	0.26	<.001***
Consult for HT - past 12 months	0.17	0.12	<.001***
Any health insurance	0.15	0.14	0.575
<i>Socio-economic characteristics</i>			
Age (years)	55.4	43.1	<.001***
Female	0.60	0.48	<.001***
Married ^b	0.69	0.42	<.001***
Worked ^c - past 12 months	0.20	0.25	0.008**
Religion: Christian	0.96	0.97	0.475
Mother tongue: Chagga	0.99	0.99	0.160
Educ ^d : None	0.10	0.10	0.997
Educ: Less than primary school	0.31	0.18	<.001***
Educ: Primary school	0.53	0.57	0.148
Educ: More than primary school	0.06	0.16	<.001***
<i>Self-reported illness/ injury</i>			
Chronic illness	0.42	0.28	<.001***
Acute illness / injury - past 12 months	0.50	0.42	<.001***
Hospitalization - past 12 months	0.07	0.07	0.931
<i>Household characteristics</i>			
Annual consumption ^e - PC (TZS / 1,000)	862	862	0.998
Financial health shock - past 12 months	0.37	0.41	0.087 ⁺
Household size	4.34	4.86	<.001***
#Young children in HH (age < 5)	0.19	0.20	0.758
#Elderly in HH (age \geq 60)	0.73	0.61	<.001***
#Reproductive age women (15-45) in HH	0.50	0.62	<.001***

Note: The table shows statistics for all adults who gave consent for the biomedical part of the baseline survey (questions and measurements). Means weighted and p-values clustered at the household level, in accordance with the sampling method BP=blood pressure; HT= hypertension; HH=household; PC= per capita; ^a43 individuals with missing test result excluded (Sample: 25; Lost to follow-up: 18);

^bIncludes mono- and polygamous marriage; ^cApart from household chores and family farming; ^dHighest completed educational level; ^eTwo outliers excluded;

⁺ p<.10, * p<.05, ** p<.01, *** p<.001.

Table C3: Baseline means of individuals lost to follow-up, by test assignment

	BP Test Mean (N=495)	No BP test Mean (N=128)	Δ Mean p-value
<i>Main</i>			
Insurance program area	0.41	0.44	0.613
Self-reported HT	0.14	0.15	0.813
BP check - past 12 months	0.27	0.21	0.214
Consult for HT - past 12 months	0.11	0.13	0.738
Any health insurance	0.13	0.17	0.392
<i>Socio-economic characteristics</i>			
Age (years)	43.1	42.8	0.892
Female	0.50	0.40	0.108
Married ^a	0.42	0.40	0.667
Worked ^b - past 12 months	0.24	0.28	0.439
Religion: Christian	0.97	0.97	0.989
Mother tongue: Chagga	0.99	0.99	0.686
Educ ^c : None	0.10	0.09	0.710
Educ: Less than primary school	0.17	0.21	0.399
Educ: Primary school	0.58	0.49	0.090 ⁺
Educ: More than primary school	0.14	0.21	0.150
<i>Self-reported illness/ injury</i>			
Chronic illness	0.27	0.31	0.485
Acute illness / injury - past 12 months	0.42	0.43	0.894
Hospitalization - past 12 months	0.07	0.09	0.497
<i>Household characteristics</i>			
Annual consumption ^d - PC (TZS / 1,000)	857	883	0.629
Financial health shock - past 12 months	0.42	0.38	0.578
Household size	4.90	4.70	0.385
#Young children in HH (age < 5)	0.21	0.15	0.299
#Elderly in HH (age \geq 60)	0.60	0.64	0.645
#Reproductive age women (15–45) in HH	0.64	0.55	0.349

Note: The table shows statistics for all adults who gave consent for the biomedical part of the baseline survey (questions and measurements), but who were lost to follow-up. Means are weighted and p-values clustered at the household level, in accordance with the sampling method. BP=blood pressure; HT= hypertension; HH= household; PC=per capita; ^aIncludes mono- and polygamous marriage; ^bApart from household chores and family farming; ^cHighest completed educational level; ^dOne outlier excluded; ⁺ p<.10.

Table C4: Baseline characteristics, by insurance intervention/control area

	Ins. area Mean (N=797)	Not Ins. area Mean (N=739)	Δ Mean p-value
<i>Main</i>			
BP test	0.79	0.82	0.188
High BP ^a	0.25	0.37	<.001***
Self-reported HT	0.22	0.25	0.154
BP check - past 12 months	0.33	0.37	0.060 ⁺
Consult for HT - past 12 months	0.17	0.17	0.685
Any health insurance	0.15	0.14	0.743
<i>Socio-economic characteristics</i>			
Age (years)	54.9	55.8	0.307
Female	0.60	0.61	0.712
Married ^b	0.70	0.69	0.666
Worked ^c - past 12 months	0.20	0.20	0.808
Religion: Christian	0.94	0.98	0.003**
Mother tongue: Chagga	1.00	0.97	0.002**
Educ ^d : None	0.11	0.09	0.176
Educ: Less than primary school	0.31	0.32	0.711
Educ: Primary school	0.54	0.52	0.438
Educ: More than primary school	0.04	0.07	0.003**
<i>Self-reported illness/ injury</i>			
Chronic illness	0.41	0.44	0.252
Acute illness / injury - past 12 months	0.49	0.51	0.463
Hospitalization - past 12 months	0.08	0.07	0.337
<i>Household characteristics</i>			
Annual consumption ^e - PC (TZS / 1,000)	851	873	0.382
Financial health shock - past 12 months	0.39	0.36	0.432
Household size	4.27	4.40	0.340
#Young children in HH (age < 5)	0.18	0.16	0.486
#Elderly in HH (age \geq 60)	0.72	0.72	0.893
#Reproductive age women (15–45) in HH	0.50	0.49	0.841

Note: The table shows statistics for all adults who gave consent for the biomedical part of both surveys (questions and measurements). Means are weighted and p-values clustered at the household level, in accordance with the sampling method. BP=blood pressure; HT= hypertension; HH=household; PC=per capita; ^a25 individuals with missing test result excluded (Ins. Treat: 8; Ins. Control: 17); ^bIncludes mono- and polygamous marriage; ^cApart from household chores and family farming; ^dHighest completed educational level; ^eOne outlier excluded; Ins.=Insurance intervention; ⁺ p<.10, * p<.05, ** p<.01, *** p<.001.

Table C5: Results corresponding to equation 1

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
BP test	0.022 (0.032)	0.014 (0.039)	0.044 (0.030)	0.009 (0.032)
Constant	-0.018 (0.028)	0.029 (0.034)	-0.064* (0.027)	0.063* (0.028)
Observations	3064	3056	3056	3072

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation “ \times Time” omitted). BP=blood pressure; HT=hypertension; 12m=12 months;

⁺ p < .10, * p < .05, ** p < .01, *** p < .001.



Figure C2: Detailed health insurance enrollment, by test result and insurance area (N=1511). No one in the insurance control area took up the KNCU Health Plan, as envisaged in the research design. Observations in the insurance intervention [control] area: Total: 789 [722]; High blood pressure: 209 [258]; Good blood pressure: 425 [326]; Control group (no test): 155 [138].

Table C6: Results corresponding to equation 3

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
BP test	0.064 (0.047)	0.025 (0.051)	0.056 (0.043)	0.018 (0.044)
BP test × Ins.area	-0.079 (0.064)	-0.023 (0.077)	-0.021 (0.061)	-0.002 (0.064)
Ins.area	0.055 (0.056)	0.001 (0.067)	0.034 (0.054)	0.128* (0.055)
Constant	-0.048 (0.041)	0.028 (0.044)	-0.082* (0.038)	-0.006 (0.039)
Observations	3064	3056	3056	3072

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation "× Time" omitted). BP=blood pressure; HT=hypertension; 12m=12 months; Ins.area= Insurance intervention area; + $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.

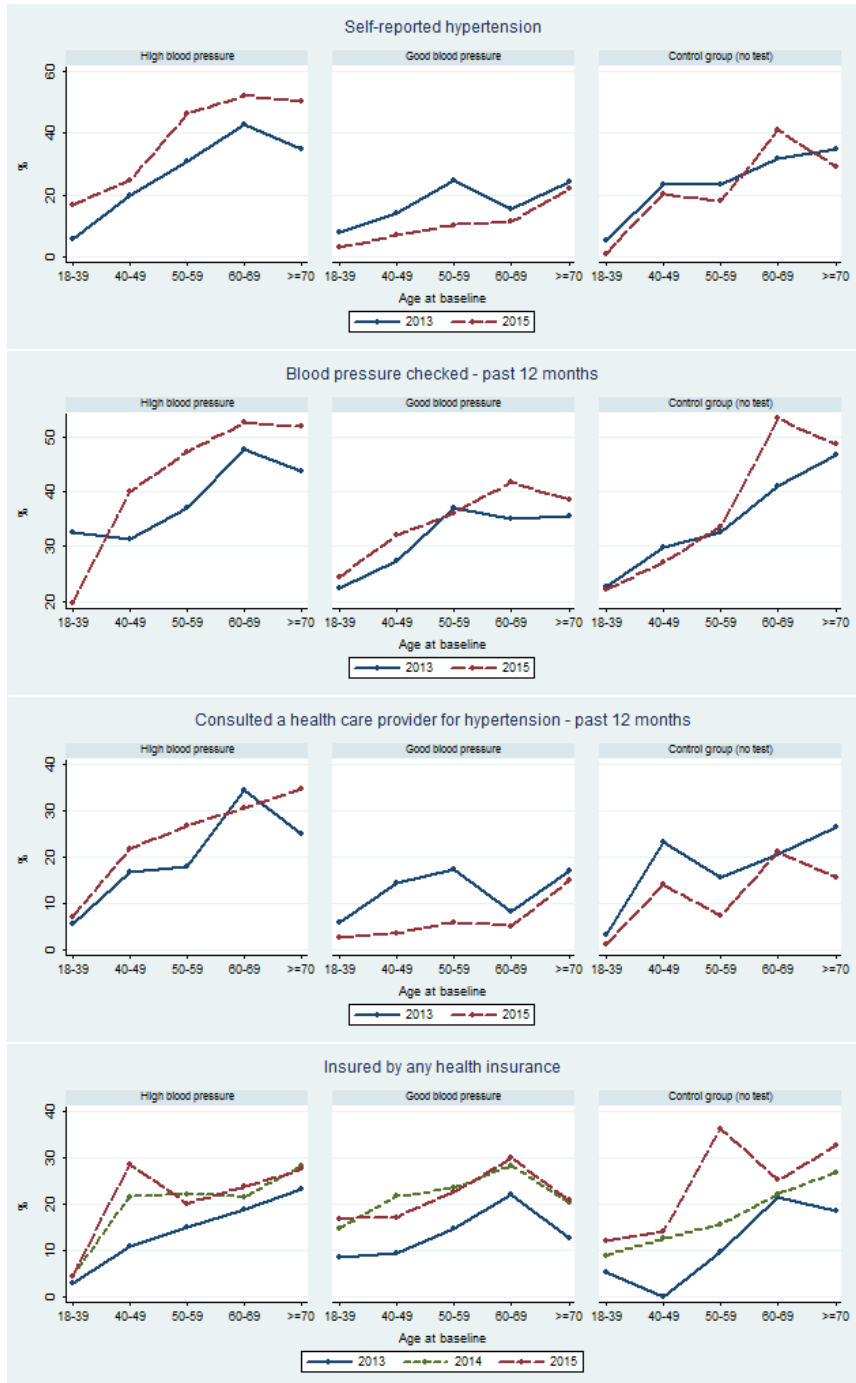


Figure C3: Outcomes by age group and test result (N=1511). The baseline and follow-up lines roughly have parallel trends, indicating no heterogeneity in the test effect by age. Observations per test result: High blood pressure: 467; Good blood pressure: 751; Control group (no test): 293. High blood pressure age-cohort 18-39 has the smallest number of observations, namely 16. The second smallest observation number is 43, in the same age-cohort of the control (no test) group.

Table C7: Heterogeneity by self-reported hypertension at baseline, corresponding to eq. 4

	(1) Self-repor- ted HT	(2) BP check: 12m	(3) Consult for HT: 12m	(4) Insured
Good BP	-0.081* (0.040)	-0.012 (0.066)	-0.064* (0.029)	0.009 (0.051)
Good BP × Self-reported HT	-0.155 (0.116)	-0.131 (0.120)	0.018 (0.130)	0.007 (0.088)
High BP	0.177*** (0.053)	0.100 (0.069)	0.067+ (0.040)	0.036 (0.049)
High BP × Self-reported HT	-0.011 (0.113)	-0.109 (0.116)	0.182 (0.125)	-0.040 (0.086)
Good BP × Ins.area	0.013 (0.053)	-0.007 (0.090)	0.040 (0.040)	0.052 (0.068)
Good BP × Ins.area × Self-reported HT	-0.108 (0.163)	0.071 (0.199)	-0.069 (0.199)	-0.171 (0.131)
High BP × Ins.area	0.012 (0.078)	-0.041 (0.102)	0.092 (0.064)	-0.004 (0.073)
High BP × Ins.area × Self-reported HT	0.008 (0.162)	0.115 (0.188)	-0.134 (0.202)	-0.158 (0.131)
Ins.area	-0.019 (0.049)	0.038 (0.074)	-0.010 (0.037)	0.094+ (0.055)
Ins.area × Self-reported HT	0.089 (0.132)	-0.237 (0.153)	0.017 (0.156)	0.162+ (0.094)
Self-reported HT	-0.594*** (0.092)	-0.187* (0.086)	-0.472*** (0.095)	0.019 (0.062)
Constant	0.136*** (0.037)	0.085 (0.054)	0.063* (0.026)	-0.012 (0.045)
Observations	3014	3008	3006	3022

Note: Fixed effects regression on balanced two-period panel; weighted and standard errors clustered at household level, according to the sampling frame. Standard errors in parentheses. All independent variables are interacted with time dummy (notation “× Time” omitted). BP=blood pressure; HT=hypertension; 12m=12 months; Ins.area= Insurance intervention area; + $p < .10$, * $p < .05$, ** $p < .01$, *** $p < .001$.