The health effects of demand side cost sharing in European health insurance

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Abstract

The rationale for demand side cost sharing in health insurance is to deter patients from using low value care. But if agents are cash constrained, demand side cost sharing can lead them to postpone or forgo valuable treatments. We use data on European (NUTS 2) regions to show that the interaction between poverty rate and out-of-pocket (oop) payments leads to unmet medical needs and higher mortality.

JEL codes: I11, I13, I18

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1 Introduction

Most developed economies face rising healthcare expenditures. In many countries the healthcare sector grows faster than the economy as a whole (OECD 2021). One of the instruments that governments have to curb this expenditure growth is demand side cost sharing. The effect of demand side cost sharing on healthcare utilization is well known. As cost sharing increases, healthcare becomes more expensive for the individual and demand for treatments falls. It is less clear whether and to which extent demand side cost sharing induces people to forgo low value care only (Newhouse and the Insurance Experiment Group 1993; Schokkaert and van de Voorde 2011). On the one hand, insurance subsidizes health consumption inducing people to get (expensive) treatments with small health benefits. Economists tend to refer to this as moral hazard. As the social costs (in contrast to an individual's out-of-pocket -oop- expenditure) of such treatments exceed their value (utility gain), an increase in demand side cost-sharing that reduces moral hazard is seen as welfare enhancing. The traditional trade off is between this increase in efficiency (due to reduced moral hazard) and the increased oop risk faced by a risk averse agent. In this paper we focus on the trade off due to behavioral hazard which refers to the case where cost-sharing leads patients to forgo valuable treatments (Baicker, Mullainathan, and Schwartzstein 2015). If a patient decides to skip a treatment where (utility) value exceeds costs because it is too expensive for her, social welfare is reduced.

The goal of this paper is to identify a health effect of demand side cost sharing through treatment becoming too expensive using European data. We start from the following two ideas. First, if demand side cost sharing reduces valuable healthcare by making it (too) expensive, this effect will be stronger for people on low income. Health is a normal good and people with high (enough) income invest in it even if it becomes expensive. Low income can force a patient to postpone or forgo treatment due to liquidity constraints. Second, if there is a substantial demand reduction for high value care, we should be able to detect this in aggregate mortality statistics.

To identify the health effects of cost-sharing we use mortality statistics of Eurostat at the NUTS 2 (Nomenclature of Territorial Units for Statistics) region level. Figure 1 illustrates the NUTS 2 regions used in this paper. Mortality varies by region/year/age/sex. In regions where the percentage of people on low income is high and demand side cost sharing is high, we expect to see high mortality. Since we have panel data, we control for NUTS 2 fixed effects.

Measuring how generous a health insurance system is, is non-trivial. Systems

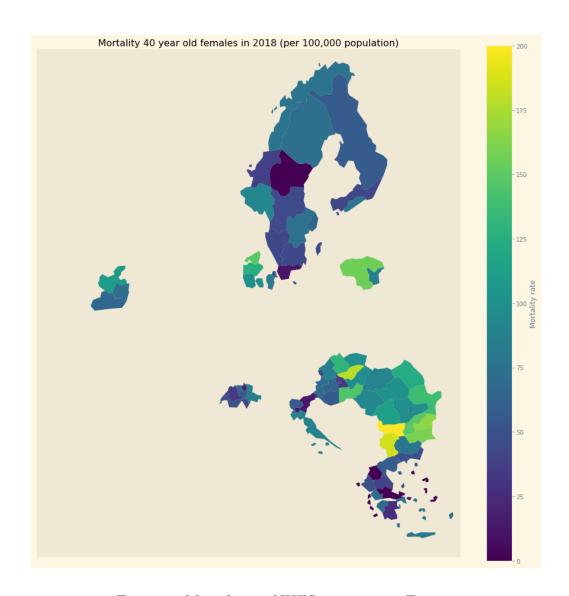


Figure 1: Mortality in NUTS 2 regions in Europe

tend to combine coinsurance with health expenditure caps, like a deductible. Hence, we introduce a model that links observed variables related to mortality, poverty, oop expenditures and people forgoing treatment because it is too expensive. The combination of the model and these variables allows us to identify the mechanism from an increase in oop via poverty to people forgoing treatment thereby raising mortality.

Figure 2 summarizes our main results in the following way. For each country in our data, we consider the NUTS 2 region where poverty is highest and therefore the effect likely to be the strongest at the regional level. Using the estimated model, we simulate the effect of a 500 euro increase in oop on mortality. We report this effect as the increase in deaths (due to the increase in oop) per 1000 dead. The motivation for this measure is two-fold. First, mortality is —thankfully—low and hence the effect of a change in oop on mortality is going to be (very) small. Reporting the increase in mortality per 1000 dead helps to interpret the numbers. Below we also present this measure for diseases that have similar orders of magnitude. Second, in our model this measure (per 1000 dead) is age-independent. That is, the number of people dying due to an increase in oop varies with age (as 25 year olds are less likely to die than 80 year olds). But the fraction of people dying due to the oop increase as a fraction of the total number of deceased is the same across age (and gender).

The blue bars indicate the average simulated effect of the 500 euro increase for this region within each country; the black lines indicate the 95% probability interval of the effect. The three countries with the biggest effects –Bulgaria, Greece and Romania– have the highest poverty levels. For these countries we can easily see that the 95% probability interval of the effect is bounded away from 0. For the Scandinavian countries, Slovenia and Switzerland the effects are close to zero at the region level because poverty is very low (even in the NUTS 2 region with highest poverty per country). Another reason for small simulated effects is a government scheme targeted at the poor helping to finance healthcare expenditures. The poor then face lower oop than our country wide oop variable would suggest. Finally, the dots present the probability that the effect in the region exceeds 0.05 (per 1000 dead). For Bulgaria, Greece and Romania this probability is close to one. For Hungary this probability exceeds 0.7. For the Scandinavian countries and Switzerland the probability that the effect exceeds 0.05 is basically zero.

The results suggest the following policy implications. An increase in oop has a measurable effect on mortality in regions where poverty is high, but the effect on the regional level is small. As we show below, if we believe this effect is (mainly) driven

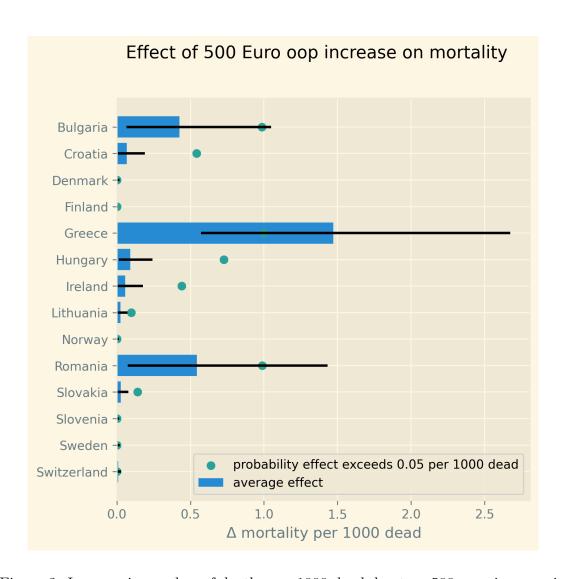


Figure 2: Increase in number of deaths per 1000 dead due to a 500 euro increase in oop for the region in each country where poverty is highest.

by the group of people indicating that they postponed treatment because it was too expensive, the effect among people in this group is an order of magnitude larger. Policies to address this include a scheme that subsidizes healthcare expenditure (on top of health insurance) for poor people; e.g. through means-tested cost-sharing. A downside of such a targeted intervention is a higher marginal tax rate at low income levels contributing to a poverty trap. Alternatively, a government can introduce co-payments that vary with the cost-effectiveness of the treatment. Treatments with high value added would then feature a low co-payment to prevent people from postponing valuable care. This can also help to reduce the mortality associated with cost sharing (Chernew et al. 2008).

This is not the first paper to consider the effects of demand side-cost sharing on mortality. There is a string of recent papers using innovative methods to identify the causal effect of health insurance on health and mortality. There are a number of issues identifying this effect of health insurance on health and mortality using individual level data. First, mortality is a rare event at most ages. Hence, identifying the effect is difficult, especially if the changes in oop are small. Second there is the selection effect that people with low health status tend to buy (generous) health insurance (as they expect high expenditure). This can bias results in the direction that individuals with (generous) insurance have adverse health outcomes (e.g. high mortality). Moreover, cost-sharing tends to be non-linear with e.g. a cap on expenditures that have to be paid oop as with a deductible. In this case, people with high care use tend to face low (marginal) treatment prices. As low health status is likely to increase care use, people facing low marginal prices (suggesting generous insurance) are likely to experience adverse health outcomes.

A number of papers use the Medicaid eligibility expansion of the Affordable Care Act which was introduced in different US states at different times. This allows for a diff-in-diffs identification strategy. Using individual level data, a number of papers have shown that the Medicaid expansion (more generous health insurance coverage) reduced mortality (Borgschulte and Vogler 2020; Miller, Johnson, and Wherry 2021). Other papers, focusing on particular causes of death, find similar results: the Medicaid expansion was associated with lower cardiovascular mortality in middle-aged adults (Khatana et al. 2019) and lower 1-year mortality among patients with ESRD initiating dialysis (Swaminathan et al. 2018).

Others analyze Medicare part D prescription drug coverage where the end-ofyear price is non-linear in expenditure. One paper uses enrollment month (related to birth month) to get exogenous variation in end-of-year expenditure for people aged 65 (Chandra, Flack, and Obermeyer 2021). The main finding is that increases in the oop costs of drugs, reduce drug use including use of high value treatments. This, in turn, raises mortality. Another approach is to show that the implementation of Medicare Part D increased the use of drug treatments for cardiovascular disease which reduced mortality (Huh and Reif 2017). By using exogenous exit of plans in the Medicare Advantage market to control for endogeneity problems, it is possible to show that more generous prescription drug coverage leads to lower mortality (Abaluck et al. 2020).

Finally, Goldin and coauthors use an experiment where a subset of people who should buy health insurance under the Affordable Care Act were reminded that they would face a financial penalty if they did not comply. This reminder tended to induce people to buy insurance instead of remaining uninsured (Goldin, Lurie, and McCubbin 2020). Mortality turns out to be lower among the people who received the reminder compared to the control group who were not reminded in this way.

Compared to these papers on health insurance (generosity) and mortality, our paper differs along the following lines. First, we use European instead of US data. The advantage is that within a European country health insurance is more homogeneous than in the US. Within a US state or county, people may have generous employer sponsored insurance, benefit from Medicaid or Medicare or have no insurance at all. Hence, a change in Medicaid coverage may have no detectable effects at the aggregate level (while an effect can be found with individual level data). In European countries a number of health insurance features are determined nationally. Consider the first two rows of the OECD Health Systems Characteristics Survey (https://qdd.oecd.org/data/HSC) showing the share of the population obtaining basic primary health care coverage through automatic or compulsory insurance coverage. For all European countries this is above 90% and for most 99% or 100%. For the US this is less than one third. Hence, country or region wide statistics in Europe give a better picture of the situation applying to most citizens in that region than in the US. This does not imply that the aggregate statistics perfectly represent everyone's insurance situation (some people may buy complementary insurance where others do not), but it may be representative enough to identify the interaction effect of poverty and oop payments we are interested in.

Moreover, individual level data sets tend to be within a country not across countries. But the variation in oop across countries is far bigger than within a country. Hence, across country data—although aggregated at the region level— helps us to identify the effect of oop on health and mortality.

Second, we show that mortality is high in regions where both oop and poverty are high. This follows the literature showing that healthcare consumption is liquidity sensitive (Gross, Layton, and Prinz 2020; Nyman 2003). People on low income tend to postpone or forgo valuable treatments if these are expensive. This focus on low incomes can imply that we under-estimate the mortality effect of cost-sharing if higher incomes also forgo valuable treatments due to oop (Brot-Goldberg et al. 2017; Chandra, Flack, and Obermeyer 2021). This is then not so much caused by liquidity problems but by behavioral hazard. In this sense, the results below are a lower bound on the mortality effects of cost-sharing.

Third, we use the regional structure of the Eurostat data. We analyze the effects of the OOP variable times Poverty interaction on mortality per age-gender class at the NUTS 2 regional level. This helps to solve the following potential endogeneity issue. A country with a population that has low health status (across ages), decides to have, say, generous health insurance. This causal effect is in the opposite direction from the one we are interested in –from health insurance generosity to health status and mortality. We avoid this problem by considering within a country how health per region varies with oop and poverty, while using NUTS 2 fixed effects to correct for other factors affecting health. By analysing health/mortality per age cohort, our results are not affected by a country's or region's age distribution. By filtering out these other effects we mitigate power issues associated with the use of mortality data (Black et al. 2021).

Fourth, Eurostat variables based on the EU-SILC survey allow us to zoom in on the relevant causal mechanism. This survey asks people whether they had unmet medical needs in the past months and if so the reason for the unmet needs. One of the answers is that treatment was postponed or skipped because it was too expensive. This allows us to simultaneously estimate the fraction of people in a NUTS 2 region that forgo treatment because it is too expensive and the effect of unmet medical needs on mortality. In this way, we capture that in regions where the OOP × Poverty interaction is high, more people postpone treatment because it is too expensive and these unmet medical needs raise mortality in the region.

Finally, our focus on the OOP × Poverty interaction distinguishes our paper from the literature on the effect of income and wealth on health (Chetty et al. 2016; Mackenbach et al. 2008; Semyonov, Lewin-Epstein, and Maskileyson 2013) where papers use cross country data. This literature typically finds that lower income and wealth is associated with lower health status, although the causal mechanism is not clear (Cutler, Lleras-Muney, and Vogl 2011). Two possible mechanisms are

that higher income leads to more expenditure on treatments (normal good) and better health. Alternatively, healthier people are more productive and earn higher incomes. The combination of fixed effects and the use of the survey question on unmet medical needs allows us to zoom in on the mechanism where high $00P \times Poverty$ interaction leads to unmet medical needs and hence to low health status and high mortality.

Summarizing, compared to papers using individual level data our approach is more broad brush and less precise in estimating the size of the effect of insurance generosity on mortality. To illustrate, we do not determine the mortality effect of a 1% change in a deductible. But we estimate the mortality effect of a 500 euro increase in oop. We do not have data on the oop details of each country's health insurance system, like what is the coinsurance rate for different types of treatments, which treatments are exempt from oop etc. Even if we had such detailed institutional data, it is not obvious how one would summarize the different systems in a way that makes them comparable across countries. Instead we use the fraction of oop payments in total healthcare expenditure, OOP, as a summary measure of a health insurance system's generosity. The theory section derives that OOP and the fraction of people postponing treatment because it is too expensive are parametric functions of the underlying exogenous parameters coinsurance rate and deductible level. This allows us to interpret the relation between OOP and mortality.

Although results based on aggregate data are less precise than those based on individual level data, our approach is more robust in the sense that it applies across a number of countries instead of a particular sub-population (like 65 year old Medicare users in the US). Although we do interpret our results using the size of the effect, our main goal is to establish that an increase in OOP in a poor region increases mortality. In particular, we quantify how sure we are that this effect is positive (bigger than 0.05 per 1000 dead in Figure 2).

We estimate our model to explain mortality for each age-gender category per NUTS 2 region per year. Figure 1 shows mortality for 40 year old women in 2018 across Europe. For each region/year/age/gender combination we observe population size and the number of deaths. We model the number of deaths as a binomial distribution where the probability of death depends on the fraction of people with unmet medical needs in a region and control variables like age, poverty and fixed effects. Simultaneously, we model the fraction of people with unmet medical needs due to financial reasons as a function of poverty and the interaction of oop and poverty. The next section presents a model explaining the relationship between

the variables mortality, poverty, OOP and the fraction of people forgoing treatment because it is too expensive.

The rest of the paper is organized as follows. After the theory section, we describe the Eurostat data that we use. We explain the empirical model that we estimate in Section 4. Estimation results are presented for the baseline model and we show that these are robust with respect to a number of our modeling choices. We conclude with a discussion of the policy implications. The appendix contains the proofs of our results and more details on our data and robustness analyses.

2 Theory

As described in the next section, the relevant variables in our data are mortality per region/year/age/sex category, OOP measuring the percentage of healthcare expenditure paid out-of-pocket (oop), the poverty rate and the fraction of people per region postponing or forgoing treatment because it is too expensive. We introduce a model to explain how these variables are related.

Consider a population in an EU region where a fraction $\alpha \in \langle 0, 1 \rangle$ has low income l and fraction $1-\alpha$ high income h. Let π^j denote the probability that someone with income j=l,h falls ill. As is well known, low income people tend to have a lower health status (Cutler, Lleras-Muney, and Vogl 2011). We capture this by assuming $\pi^l > \pi^h$. People on low income may have a less healthy diet, exercise less etc. due to either the cost of or knowledge about healthy lifestyle choices. This makes it more likely that they fall ill.

Generally speaking, oop payments tend to take two forms that we want to capture: a coinsurance rate, which we denote $\xi \in [0, 1]$, and a maximum expenditure, which we denote D (for deductible). Some systems have a combination of the two.

Conditional on falling ill, there is a probability $\zeta_i \in [0,1]$ that the patient is advised to get treatment i at cost x_i for i in the set of "illnesses" I. We define I_{ξ} as the subset of I where $\xi x_i < D$ and $oop_i = \xi x_i$ and I_D where $\xi x_i \geq D$ and $oop_i = D$. To keep things simple, we assume that ζ_i is exogenous for the patient. We model the treatment decision on the extensive margin only: accept or reject the treatment proposed by a physician. A pure coinsurance system has $\xi < 1$ and $I_{\xi} = I$. A pure deductible system $\xi = 1$ and I_D non-empty. A combination of the two has $\xi < 1$ and there is a maximum on the oop payment. Health insurance systems in Europe tend to have such maximum oop expenditure. An increase in either ξ or D is interpreted

 $^{^1\}mathrm{See}$ question 12 in https://qdd.oecd.org/data/HSC specifying for most European countries

as making health insurance less generous.

Whereas with individual level data one can determine whether an individual faces a positive treatment price at the margin (E.g. using the end-of-year price as in Keeler, Newhouse, and Phelps 1977; Ellis 1986), this is not possible with the aggregate data that we use here. Hence, we rely on an aggregate summary variable, denoted 00P, measured as oop payments over total healthcare expenditure. That is, the fraction of healthcare expenditure paid by patients oop. We interpret this variable as capturing the generosity of the health insurance system. To illustrate, if healthcare is free at point of service, 00P equals zero; if there is no health insurance at all, 00P equals 1. The challenge is to capture changes in ξ and D although we do not directly observe these variables in the data. This is what the model sets out to do.

If a patient receives treatment $i \in I$, we denote her (expected) health σ_i , while without treatment (expected) health equals σ_0 with $0 \le \sigma_0 < \sigma_i \le 1$. Health is normalized at value one for a patient who does not fall ill. We simply assume that utility is multiplicative in health and consumption and we model the patient's treatment decision as:

$$\nu \sigma_i u(y^j - oop_i) > \sigma_0 u(y^j) \tag{1}$$

where utility u(.) is determined by how much money can be spent on other goods: y^j minus oop in case of treatment and y^j if no treatment is chosen. The utility function u(.) is increasing and concave in consumption: u(.), u'(.) > 0 and u''(.) < 0. Further, parameter ν captures other factors than pure financial ones affecting a patient's treatment choice.

In our data, we have a variable "unmet medical needs" based on a number of motivations: treatment is too far away to travel to, there is a long waiting list, the patient is scared to undergo treatment etc. To make our point, it is enough to assume that such factors affect utility in a multiplicative way. To illustrate, if the patient has to travel far for treatment, utility is reduced by multiplying it with $\nu < 1$. The cumulative distribution function of ν is given by $G(\nu)$ and its density function by $g(\nu)$. Other factors can include travel time to treatment, belief that the condition will resolve itself without intervention, poor decision making and focus on the short term undervaluing the benefit of treatment. If inequality (1) holds, the agent accepts the treatment. In the proofs in the appendix we assume that G is a Pareto distribution.

a spending cap.

²To ease notation we do not let σ_0 vary with i.

Hence, the probability that a patient with income y^j accepts treatment i offered by a physician equals

 $\delta_i^j = 1 - G\left(\frac{\sigma_0}{\sigma_i} \frac{u(y^j)}{u(y^j - oop_i)}\right)$

that is, ν is big enough that inequality (1) holds. With probability $G\left(\frac{\sigma_0}{\sigma_i}\frac{u(y^j)}{u(y^j-oop_i)}\right)$ the patient decides to postpone or forgo treatment i.

The probability that a patient postpones or skips a treatment because it is too expensive is given by

$$G\left(\frac{\sigma_0}{\sigma_i} \frac{u(y^j)}{u(y^j - oop_i)}\right) - G\left(\frac{\sigma_0}{\sigma_i}\right) \tag{2}$$

These are agents ν that would have chosen treatment i if it were free $(oop_i = 0$ and $u(y^j)/u(y^j - oop_i) = 1)$ but who forgo treatment now that it costs $oop_i > 0$. The probability $G(\sigma_0/\sigma_i)$ captures factors like waiting lists or the patient hoping that the health problems resolve themselves without treatment. That is, reasons for postponing treatment not related to oop payments. In the proof of the lemma at the end of this section, we show that the probability of treatment is increasing in income y^j and decreasing in oop_i , as one would expect.

An agent's health is affected by the probability of falling ill and then getting treatment (or not). We assume that agents' mortality is affected by health in the following way, where we define mortality m as the prophability of dying in a given period.

$$\ln(m_{agt}) = \ln(\tilde{\eta}_{ag}) + \gamma \ln\left(\frac{m_{a-1,g,t-1}}{\bar{m}_{a-1,g}}\right) - (\alpha(1-\pi^l) + (1-\alpha)(1-\pi^h))$$
(3)

$$-\alpha \pi^l \sum_{i \in I} \zeta_i (\delta_i^l \sigma_i + (1 - \delta_i^l) \sigma_0) - (1 - \alpha) \pi^h \sum_{i \in I} \zeta_i (\delta_i^h \sigma_i + (1 - \delta_i^h) \sigma_0)$$

where we use the following subscripts: age a, gender $g \in \{f, m\}$, year t. In words, log mortality in a region depends on the biology of age and gender, $\tilde{\eta}_{ag}$. As people get older, they tend to become less healthy and more likely to die. We define this effect as independent of country or year (in the period that we analyze). Then there are a number of effects that increase or decrease mortality compared to $\tilde{\eta}_{ag}$. The health of the age-gender cohort in the previous period: if in a NUTS 2 region there was a shock in t-1—when this cohort was aged a-1—that increased mortality above the average (across years and regions) mortality for this cohort, we interpret this as a negative health shock. For the people that survived in this cohort, this health

shock can affect their mortality in period t. This is captured by the coefficient γ .³ People who do not fall ill, have the highest health level (normalized to 1) and hence reduce mortality to the biggest extend. People who do fall ill and get treatment, get health $\sigma_i \leq 1$ and reduce mortality to a smaller extent. Finally, people falling ill but forgoing treatment lead to the smallest reduction σ_0 in mortality.

As we show in the proof of the lemma below, we can write the expression for log mortality as:

$$\ln(m_{ag2t}) = \ln(\tilde{\eta}_{ag}) + \mu_2 + \gamma \ln\left(\frac{m_{a-1,2,g,t-1}}{\bar{m}_{a-1,g}}\right) + \beta_{poverty}\alpha_{2t} + \beta_{unmet} \text{Unmet}_{2t} \quad (4)$$

where subscript 2 indicates that the variable varies with NUTS 2 region, μ_2 denote NUTS 2 fixed effects, poverty α varies with NUTS 2 region and year and Unmet denotes the fraction of people indicating unmet medical needs in a region in year t.

In our data, the variable Unmet varies with NUTS 2 region and year and not by age or gender. Hence, in terms of our model, we define this variable as follows:

$$Unmet_{2t} = \sum_{i \in I} \zeta_i (\alpha_{2t} \pi^l (1 - \delta_{ict}^l) + (1 - \alpha_{2t}) \pi^h (1 - \delta_{ict}^h))$$
 (5)

with δ_i^j varying with country c and year t because oop varies with countries over time.

Further, in our data we have the variable OOP defined as oop payments as a percentage of healthcare expenditure. In terms of our model, we write this as

$$OOP = \frac{\sum_{i \in I} \zeta_i oop_i(\alpha \pi^l \delta_i^l + (1 - \alpha) \pi^h \delta_i^h)}{\sum_{i \in I} \zeta_i x_i (\alpha \pi^l \delta_i^l + (1 - \alpha) \pi^h \delta_i^h)}$$

The numerator of 00P contains the oop payments oop_i and the denominator expenditures x_i . If $I_{\xi} = I$, it is clear that OOP = ξ . Because I_D is non-empty (European countries have a maximum oop payment), the expression for 00P is actually non-trivial. We can also write 00P as the ratio of average oop per head and average healthcare expenditure per head:

$$OOP_{ct} = \frac{\overline{oop}_{ct}}{\bar{x}_{ct}} \tag{6}$$

In our data these variables vary by country and year.

³Although we think of $\gamma > 0$, we allow for $\gamma < 0$. The interpretation in the latter case would be that some people with low health status in cohort a-1 passed away early, increasing average health for people remaining in this cohort.

Finally, using equation (2) our model allows us to formalize the fraction of people that forgo treatment because it is too expensive.

TooExp =
$$\alpha \pi^l \left(\sum_{i \in I} \zeta_i \left(G \left(\frac{\sigma_0}{\sigma_i} \frac{u(y^l)}{u(y^l - oop_i)} \right) - G \left(\frac{\sigma_0}{\sigma_i} \right) \right)$$
 (7)

$$+(1-\alpha)\pi^{h}\left(\sum_{i\in I}\zeta_{i}\left(G\left(\frac{\sigma_{0}}{\sigma_{i}}\frac{u(y^{h})}{u(y^{h}-oop_{i})}\right)-G\left(\frac{\sigma_{0}}{\sigma_{i}}\right)\right)$$

In our data, TooExp varies with Nuts 2 region and year. The following lemma summarizes the main results from the model and presents the equations that we estimate below. The innovation is to view equations (6) and (7) as being parametrized by ξ and D which are not observed in our data. We show that this leads to an equation where TooExp is a function of OOP and poverty.

Lemma 1 Healthcare demand $\delta = 1 - G(.)$ is increasing in income y^j and decreasing in oop_i (ξ or D). We write the expression for female/male mortality of age cohort a in Nuts 2 region 2 at time t as:

$$m_{ga2t} = \frac{e^{\beta_{ag}}}{1 + e^{\beta_{ag}}} e^{\left(\mu_2 + \gamma \ln\left(\frac{m_{a-1,g,2,t-1}}{\bar{m}_{a-1,g}}\right) + \beta_{poverty}Poverty_{2t} + \beta_{unmet}Unmet_{2t}\right)}$$

where $\beta_{poverty}, \beta_{unmet} > 0$. The linear expansion of TooExp with respect to OOP can be written as

$$TooExp_{2t} = b_{0,2} + b_{0,t} + OOP_{ct}\bar{x}_{ct} \left(b_{oop,c} + b_{interaction,c}Poverty_{2t}\right)$$

We model $\tilde{\eta}_{ag}$ as a sigmoid of age and gender fixed effects, β_{ag} . This makes sure this part of the probability of death is between 0 and 1. We multiply this baseline probability with a multiplier taking care of other effects. In particular, NUTS 2 region fixed effects which capture regional variation in π^h . Whether this age cohort experienced a health shock in the previous period (compared to the average mortality of this cohort). Poverty level and the fraction of people with unmet medical needs in the region in year t. If the sum of these terms is negative, the multiplier is less than 1 and mortality for this gender/age/region/year combination is reduced compared to the baseline probability given by the sigmoid. If the sum of the terms is positive, mortality for this observation is higher than the baseline probability.

We use a linear expansion of TooExp in terms of OOP. The appendix shows how we derive this relation using the exogenous variables ξ and D which affect OOP and TooExp simultaneously. It turns out that there is a direct effect of OOP on TooExp and an interaction effect with the fraction of people below the poverty line in a region. We show that b_{oop} , $b_{interaction} > 0$: a region that lies in a country with high OOP tends to have high unmet needs and especially so if the region features a high poverty rate. In our estimation we want to ensure that TooExp is between 0 and 1. For this we assume that TooExp has a logit-normal distribution. That is, the log-odds of TooExp is normally distributed.

Figure 3 illustrates this approximation of the relation between (log-odds) TooExp and 00P for simulated values in the model above. We simulate data for two countries which differ in poverty rate (see web appendix for details). As we vary ξ and D, both 00P and expenditure per head vary leading to the graph in the left panel of Figure 3. As explained in the proof of the lemma, the linear expansion of TooExp in 00P and 00P × Poverty interaction does not determine the intercept b_0 . Therefore, we allow b_0 to vary by region and year: $b_{0,2} + b_{0,t}$. For both sets of simulated data, the approximation where the (log odds of) fraction of people forgoing treatment because it is too expensive depends linearly on 00P × Poverty seems reasonable. As shown in the appendix, we need to multiply 00P and 00P × Poverty by healthcare expenditure per head because the underlying changing variable is not the endogenous 00P but the policy parameters ξ and D. As illustrated in equation (6), the relation between changes in D and 00P is multiplied by expenditure per head: dOOP/ $dD \propto 1/\bar{x}_{ct}$.

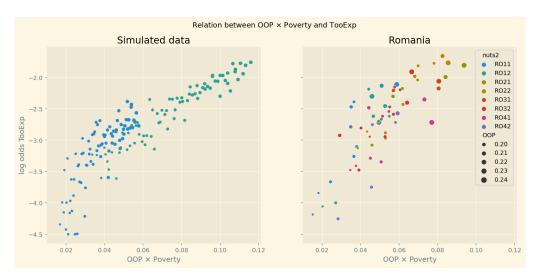


Figure 3: The simulated relation between fraction of people who forgo treatment because it is too expensive and 00P measure for different values of ξ , D and α (left panel) and this relation for NUTS 2 regions and years in Romania (right panel).

The right panel of Figure 3 illustrates this relation for regional data from Roma-

nia. Again a linear approximation looks reasonable.

3 Data

The data that we use is from Eurostat's regional database and provides for NUTS 2 regions population size and number of deaths per age-gender category. In principle, we have data on 14 countries and 78 NUTS 2 regions for the years 2009-2019, ages 35-85 for women and men. The years 2009-2019 were chosen because, at the time of the analysis, data on poverty was available from 2009 onward and data on the number of deaths ran till 2019. We start at age 35 because at ages below 35, mortality is so low that there is hardly a difference between mortality in regions with different poverty levels (Figure 4 below). For ages above 85 population numbers get very low. We drop NUTS 2 region-year combinations where for an age-gender category—due to reporting issues or people moving—the number of deaths in a year exceeds the population size at the start of the year. We focus on observations where we have complete records on mortality, the fraction of people indicating they postponed treatment because it was too expensive and oop expenditure.

Table 1 shows the summary statistics for our variables. We have more than 50k observations. The average population size per region-age-gender category is about 7500 and the average number of deaths 100. Median population size per category equals 6500 and median number of deaths 56. Percentage of people dying in a NUTS 2/year/age/gender category (mortality) equals 2% on average with a maximum of 20%.

Table 1: Summary statistics main variables

	count	mean	std	min	median	max
population	52612.00	7491.28	4805.28	440.00	6477.00	36117.00
deaths	52612.00	103.19	126.49	0.00	56.00	1033.00
mortality	52612.00	2.12	2.94	0.00	0.81	20.72
poverty	52612.00	16.47	6.50	2.60	15.30	36.10
deprivation	52612.00	11.23	12.78	0.00	3.40	52.30
too exp.	52612.00	2.00	3.09	0.00	0.60	16.00
unmet	52612.00	4.93	3.73	0.00	4.00	20.00
out-of-pocket	52612.00	22.03	8.88	8.83	19.46	47.74
voluntary	52612.00	3.12	3.07	0.33	1.59	15.20
expend. per head	52612.00	3379.56	2688.57	307.69	3559.49	8484.88

We use two measures for poverty; each of these measures comes from the EU statistics on income and living conditions (EU-SILC) survey. The first is "at-risk-of-poverty rate" that we refer to as poverty. This is a relative poverty measure: the share of people with disposable income after social transfers below a threshold based on the national median disposable income. The material deprivation measure (denoted deprivation) refers to the enforced inability to pay unexpected expenses, afford adequate heating of the home, durable goods like a washing machine etc. See the appendix for details.

In our data, the (unweighted) average (across regions and years) percentage of people at risk of poverty equals 16% with a maximum of 36%. For material deprivation these numbers are 11% and 52%. These measures vary by NUTS 2 region and year but not by age or gender. We use deprivation in our baseline analysis because it captures more closely the idea of postponing treatment due to financial constraints. The poverty variable is used in a robustness check.

Also from the EU-SILC survey, we use the variable capturing unmet healthcare needs because the forgone treatment was too expensive (too exp). The variable unmet measures percentage of people that postpone or forgo treatment because it is either too expensive, the hospital is too far away, there is a waiting list for the treatment, the patient hopes that symptoms will disappear without treatment or because the patient is afraid of treatment. As explained in the model above, our analysis uses both too exp and unmet (which includes too exp as reason for unmet medical needs) as variables.

The measure 00P that we use in the baseline model, is based on household oop payments (out-of-pocket). In particular, this measures the percentage of healthcare expenditures paid oop. This varies by country and year. The higher 00P, the less generous the healthcare system is (in terms of higher coinsurance ξ or deductible D in the model above). We expect that high 00P is especially problematic in regions with a high percentage of people in poverty.

In a robustness analysis we consider the sum of oop and payments to voluntary health insurance as a percentage of health expenditures (voluntary) as our OOP measure. The reason why we also consider the sum of expenditure on voluntary insurance and oop payments is that basic or mandatory insurance packages can differ between countries. If people are willing to spend money on voluntary insurance, it can be the case that this voluntary insurance covers treatments that people deem to be important. Put differently, a country that finances all expenditure ("free at point of service") for a very narrow set of treatments would appear generous if we

only used oop payments. The narrowness of this insurance would then be signalled by people buying voluntary insurance to cover more treatments.

As can be seen in Table 1, out-of-pocket is the most important component of the two OOP inputs. Percentage of healthcare expenditure paid oop is a multiple of the percentage financed via voluntary insurance (both in terms of the mean and of the minimum, median and maximum reported in the table). Therefore, the baseline model works with oop payments (only).

Finally, as shown in Lemma 1, healthcare expenditure per head (expend per head) affects how 00P influences the fraction of people forgoing treatment because it is too expensive. Expenditure per head is on average 3300 euro for the countries in our data. But the variation is big: minimum value of 308 euro per year and maximum value of 8500 euro.

Figure 4 (left panel) shows average mortality as a function of age for women and men. This is the pattern that one would expect: clearly increasing with age from age 40 onward and higher for men than for women (as women tend to live longer than men). Figure 4 (middle panel) shows the effect we are interested in: mortality is higher in regions where the interaction OOP × Poverty is high than where it is low and this difference increases with age. Both for women and for men, we plot per age category the difference between average mortality in regions that are at least one standard deviation above the mean for $OOP \times Poverty$ and regions that are one standard deviation below the mean. Around age 80, this mortality difference equals approximately 1.75 percentage points. In the raw data, for 100 women, there are 1.75 additional deaths in regions with high $OOP \times Poverty$ compared to regions with low interaction. Note that this plot of the raw data does not correct for other factors, like the poverty level itself, and thus over-estimates the size of the effect of $OOP \times$ Poverty on mortality. The right panel in this figure does a similar exercise with the fraction of people reporting unmet medical needs. Mortality is higher in regions where unmet needs are at least one standard deviation above the mean compared to regions where it is one standard deviation below the mean.

The observation from the figure that the difference between the two sets of regions is approximately zero for people below 35, is our motivation to include ages above 35 only in our data. Further, the difference in mortality between the regions increases with the mortality level in the left panel. This is in line with our specification in Lemma 1 where unmet needs has a multiplicative effect on the underlying mortality rate modeled by $e^{\beta_{ag}}/(1 + e^{\beta_{ag}})$.

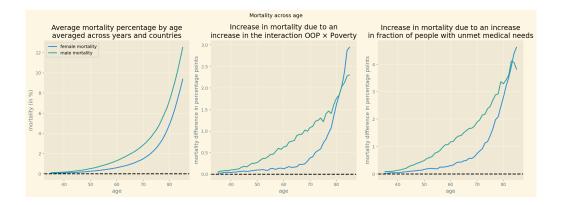


Figure 4: Mortality and difference in mortality between regions with high and low interaction $OOP \times Poverty$ and high and low unmet medical needs.

4 ESTIMATION

In this section, we explain the model that we estimate.

4.1 Empirical model

We estimate a binomial model with population size as the number of draws and deaths as the number of events. We do this for every combination of NUTS 2 region, calendar year, age and gender in our data. The probability of $k \leq n$ deaths out of a population n is then given by

$$\binom{n}{k}m^k(1-m)^{n-k}$$

where m denotes mortality, the probability of death. The advantage of modeling k as a binomial distribution is that it automatically captures that the variance in the proportion of deaths will be bigger if population size n is smaller. The equation that we estimate for m_{2atg} is given in the Lemma 1. The coefficient we are interested in is β_{unmet} . Once we know how OOP increases the fraction of people postponing treatment because it is too expensive, this increase affects Unmet and β_{unmet} then gives us the increase in mortality.

Hence, we also estimate the equation determining the fraction of people forgoing treatment because it is too expensive as specified in Lemma 1.

4.2 Bayesian estimation

We use Markov Chain Monte Carlo (MCMC), in particular the NUTS sampler to explore the posterior distributions of our parameters. We use regularized priors on the coefficients: centered at 0 with a small standard deviation. This reduces the risk of over-fitting: a parameter only deviates from 0 if there is strong evidence for this in the data. Details on the prior can be found in the online appendix.

5 Results

In this section we present the results of the estimation of the baseline model. Before presenting the outcome of our estimation, we present graphically two checks of our model.

5.1 model fit

First, Figure 5 gives an idea of the fit of the model in terms of predicting deaths per gender/age/region/year category and the fraction of people postponing treatment because it is too expensive.

The left panel shows observed number of deaths per category on the horizontal axis and the posterior predictive for this on the vertical axis. For each row in our data, we have observed number of deaths and a prediction of this number. In particular, the Bayesian model generates 8000 posterior samples (4 chains with 2000 samples each). In the figure, we show the average prediction of deaths across the samples. The predictions are not perfect but do follow the 45-degree line closely.

The right panel shows the (log odds of the) fraction of people per region/year indicating they went without treatment (for a while) because it was too expensive. Two things are different in this panel compared to the left. First, this fraction does not vary by gender and age. Hence, we do not have a prediction for each row in our data. Second, this fraction TooExp is based on (EU-SILC) survey data where we do not know the number of people interviewed. Hence, we cannot model this as a binomial distribution where we predict the number of people indicating unmet medical needs because of financial constraints.

Therefore, the right panel shows the observed and predicted fraction for TooExp per region/year. The dots indicate the average posterior prediction of this log-odds ratio and the vertical black lines indicate the 95% prediction interval.

A final observation is that **TooExp** equals 0 for a number of region/year combinations. To handle this numerically, we use of lower bound of -5 for the log-odds. This corresponds to a probability of 0.007 which is close enough to zero for our purposes. The right panel shows this bunching for a number of observations at -5.

Compared to the observed number of deaths, the predictions for TooExp seem less accurate. This is to be expected as there are a lot fewer observations for this variable compared to mortality. But all in all the fit does not seem unreasonable as the points cluster around the 45-degree line.

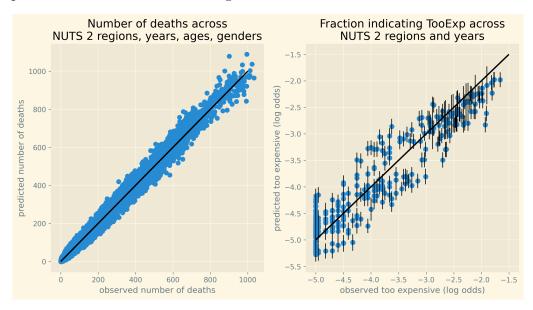


Figure 5: Fit of estimated and observed mortality across all observations and observed and predicted fraction of people indicating TooExp across NUTS 2 regions.

Another way to check how well the model fits, is to see how well it captures the age profile of mortality. This we present in Figure 6. The left panel shows the age profile $\tilde{\eta}_{ag} = e^{\beta_{ag}}/(1+e^{\beta_{ag}})$. If the other terms in equation (4) equal 0, $\tilde{\eta}_{ag}$ gives the probability of death for category ag. The right panel includes for every region and calendar year the correction on $\tilde{\eta}_{ag}$ to yield mortality for that combination of gender/age/region/year. On average, the model captures the age profile perfectly.

The appendix presents two further checks of the model. Figure 7 shows the trace plots for the parameters of interest. The figures in the left panel show the posterior distribution of the parameters in the figure. The coefficients b_oop, b_interaction vary by country and hence we have different colors for the distributions in these graphs. The beta parameters do not vary with country (or another index) and hence there is one color only. In the beta figures it is easy to see that there are four distributions per parameter. These correspond to the four chains that are sampled by the NUTS algorithm.

The right panels show the same samples but now ordered across the horizontal axis as they were drawn. We check these plots for the following three features. First, the plot should be stationary; that is, not trending upward or downward.

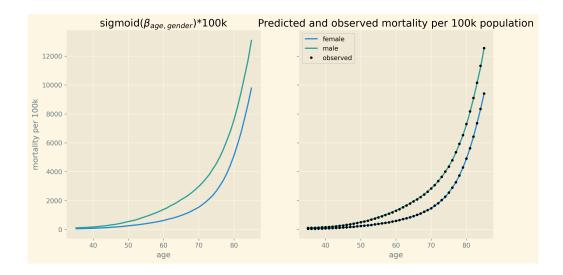


Figure 6: Fit of average mortality by age

This implies that the posterior mean of the coefficient is (more or less) constant as we sample. Second, there should be good mixing which translates in condensed zig-zagging. In other words, the algorithm manages to draw values across the whole domain of the posterior quickly after each other. Finally, the four chains cover the same regions. All three features are satisfied in the right panel of the figure.

Another check on the convergence of the algorithm are the r-hat values in Table 5. This table summarizes the posterior distribution for the slopes that we arter interested in. It provides the mean and standard deviation for each of these parameters, the 95% probability/credibility intervals and the number of effective samples for each parameter. As the number of these samples is clearly above 1000 for each parameter, this looks fine. The final column presents the values for r-hat for each parameter. Since these are all equal (close) to 1.00, we can be confident that the NUTS algorithm converged for these parameters.

5.2 size of effects

Table 5 presents the values for each of the parameters. Here we focus on the effect we are interested in: what is the increase in mortality due to an increase in oop? As we show in the appendix, it is routine to verify that 500 euro increase in oop leads to the following increase in mortality:

$$\frac{dm_{ga2t}}{m_{ga2t}} = \beta_{unmet} \text{TooExp}_{2t} (1 - \text{TooExp}_{2t}) 500 (b_{oop,c} + b_{interaction,c} Poverty_{2t})$$

Note that this increase in the number of deaths dm_{ga2t} per the number of deaths m_{ga2t} is independent of age. This is due to our formulation of mortality in equation

(4) where we have a baseline mortality $\tilde{\eta}_{ag}$ and a deviation from this baseline based on poverty and unmet medical needs etc. Figure 2 reports the expression in the equation above multiplied by 1000. That is, we report the increase in deaths due to the oop increase per 1000 deaths.

As the expression for dm/m varies with country, year and NUTS 2 region, Figure 2 summarizes our main findings in the following way. For each country we focus on the region where deprivation is highest. This is the region where we expect the mortality effect to be highest as many people could have problems paying medical bills. Table 2 presents this region for each country in our data together with the value of deprivation and the fraction of people with unmet medical needs due to financial constraints.

Table 2: Region per country with highest fraction of material deprivation

region	country	deprivation	too expensive
BG33	Bulgaria	0.40	0.08
HR04	Croatia	0.13	0.01
DK02	Denmark	0.04	0.00
FI1C	Finland	0.03	0.00
EL63	Greece	0.28	0.07
HU31	Hungary	0.32	0.02
IE06	Ireland	0.07	0.02
LT02	Lithuania	0.12	0.01
NO01	Norway	0.02	0.00
RO22	Romania	0.32	0.11
SK04	Slovakia	0.11	0.01
SI03	Slovenia	0.05	0.00
SE22	Sweden	0.02	0.00
CH01	Switzerland	0.02	0.02

Substituting these values from the table into the expression for dm/m we get the numbers in Figure 2. As mentioned, the blue bars give the average effect of the 500 euro increase in oop on mortality. As we have the posterior distributions for each of the parameters, we also have the posterior distribution for the mortality effects per country. The black horizontal lines present the 95% intervals around the mean effect. Finally, the dots present the probability that the effect is at least 0.05 increase in deaths per 1000 dead.

The first observation is that for Bulgaria, Greece, Romania and –to a lesser extent– Hungary the 95% probability interval is clearly bounded away from zero. For these countries we can clearly see that an increase in oop negatively affects health and increases mortality. The probability that the effect is at least 0.05 (per 1000) is close to 1.0 for the former three countries and almost 0.75 for Hungary. In this sense, for our European data we do not suffer from power issues in identifying mortality effects in aggregate (regional) data (Black et al. 2021).

Why are the effects smaller for the other countries? The effects are basically zero for the Scandinavian countries, Slovenia and Switzerland. As shown in Table 2, for these countries both deprivation and the fraction of people indicating unmet medical needs because treatment is too expensive are small. For the Scandinavian countries in the region with highest deprivation, TooExp is basically zero. For low income families there is enough financial support to ensure that treatments are not postponed for financial reasons.

Another reason why the effects are small for some countries is that the underlying parameters b_oop, b_interaction are small for these countries. This can be seen in Table 5 in the appendix. If countries have policies to subsidize healthcare for poor families, the effects of country wide OOP on these families is small as they actually pay a lower fraction of their treatments' costs oop. Indeed, these coefficients for the Scandinavian countries are relatively low. Greece in the years of the financial crisis had less scope to support low income families in this way.

Summarizing, we can identify in our data the effect that an oop increase, raises the number of people with unmet medical needs due to financial constraints and hence increases mortality. This was the main objective of this paper.

A follow up question is: how big is this effect? In order to interpret the size of the oop effect, Table 3 presents the number of people dying from a particular cause per 1000 dead.⁴ If we would consider all causes and add them up, the sum would equal 1000. The table focuses on causes of death with relatively small numbers involved. The table is based on EU wide data in 2017 for ages 35-85.

Note that the comparison of the numbers in Figure 2 with the numbers in the table is just to get an idea of the order of magnitude. But –strictly speaking– the causes are not comparable. Nobody dies of an increase in oop in the way people die from influenza. Due to an increase in oop, people may have gone without treatment which can then lead to death from, say, hepatitis. Hence, one should be careful in

⁴We use the icd-10 classification here.

interpreting the simulation results with the numbers in Table 3. But the table does provide some context in interpreting the size of the simulated effects.

Table 3: Number of people dying by cause (per 1000 dead) for ages 35-85 (EU average).

icd-10	per 1000
Tuberculosis	0.93
Chronic viral hepatitis B and C	0.28
HIV	0.72
Malignant melanoma of skin	4.20
Malignant neoplasm of thyroid gland	0.96
Malignant neoplasm of ovary	8.06
Malignant neoplasm of stomach	14.03
Diseases of the blood and blood-forming organs	2.69
Leukaemia	10.15
Mental and behavioural disorders due to use of alcohol	4.90
Influenza (including swine flu)	1.18
Asthma and status asthmaticus	1.22
Rheumatoid arthritis and arthrosis	1.06
Alzheimer disease	13.08
Drug dependence, toxicomania	0.38
Accidental drowning and submersion	1.28
Intentional self-harm	12.82
Assault	0.67

The average mortality effect due to a 500 euro increase in oop in Greece is approximately 1.5 (per 1000 dead). This exceeds the effect due to accidental drowning. The effects in Bulgaria and Romania are around 0.5 which places them between deaths due to drug dependence and due to HIV.

However, these are effects aggregated at the regional level. If we are willing to assume that the incidence of the increase in mortality due to increase of oop by 500 euro falls in the group of people that indicate that they postponed or did without a treatment because it was too expensive, we can divide the mortality effect that we have by the size of this group. To get a rough idea of the size of the effect in this case, consider again Table 2. For the countries where we find sizable effects—Greece, Romania, Bulgaria—the fraction of people indicating they found treatment

too expensive equals approximately 0.1 (or less) which we use to facilitate the backof-the-envelop calculations here.

Hence, roughly speaking, in the NUTS 2 regions (which are worst affected in terms of deprivation per country), to find the mean overall effects on mortality as indicated in the figure, the incidence within the group too expensive would vary between 0.5/0.1 = 5 and almost 1.5/0.1 = 15.

Then within the group of people indicating they sometimes postpone or forgo treatment because it is too expensive, the increase in deaths due to an increase in oop is of an order comparable to deaths due to Leukaemia or Alzheimer disease.

In countries where poverty is high, a 500 euro increase in oop does not lead to a big increase in mortality. But the order of magnitude of deaths among 1000 dead are comparable to a number of well known conditions varying from HIV (at the regional level) to Leukaemia and Alzheimer disease if we assume the incidence is among the group of people indicating they forgo treatment because of financial reasons.

6 Robustness Checks

6.0.1 **TODO** In this section we discuss three robustness checks.

- other definition poverty: material deprivation
- other definition oop: include voluntary insurance
- separate unmet and TooExp effects in mortality equation

7 Discussion and Policy implications

The Introduction discusses a recent literature using individual level data analyzing whether demand side cost sharing reduces expenditure on low value treatments (usually referred to as moral hazard) or whether it leads patients to postpone or forgo valuable treatments thereby negatively affecting their health. This literature is focused on US data and argues that it is hard to find negative health effects of cost sharing in aggregate data. We use European data at the (NUTS 2) regional level to show that a high share of out-of-pocket expenditures (in total healthcare expenditures) has a clear effect on mortality in regions where the fraction of low income households is high.

Healthcare costs keep increasing in most, if not all, developed countries. Demand side cost sharing is a well known instrument to curb the growth in expenditure. This paper shows that there is a upper bound on oop beyond which regions with high poverty levels start to show increased mortality rates. To avoid this mortality effect, policy makers need to search for alternative instruments.

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A Proof of results

Proof of Lemma 1 First, we show that the probability of treatment, $1-G(\sigma_0/\sigma_i u(y^j)/u(y^j-oop))$, is increasing in y^j and decreasing in oop. Taking the derivative

$$\frac{d\left(\frac{u(y^{j})}{u(y^{j}-oop)}\right)}{dy^{j}} = \frac{u'(y^{j})u(y^{j}-oop) - u(y^{j})u'(y^{j}-oop)}{u(y^{j}-oop)^{2}} < 0$$

because u is positive and increasing in y, u' > 0 is decreasing in y and oop > 0. Hence, the probability of treatment is increasing in y^j . Similarly, the treatment probability falls with oop.

The expression for mortality follows from equation (3) which we can write as:

$$\ln(m_{agt}) = \ln(\tilde{\eta}_{ag}) + \gamma \ln\left(\frac{h_{a-1,g,t-1}}{\bar{g}_{a-1,g}}\right) - (1 - \pi^h) - \pi^h \sum_{i \in I} \zeta_i + \alpha(\pi^l - \pi^h)(1 - \sum_{i \in I} \zeta_i)\sigma_0 + \sum_{i \in I} \zeta_i(\sigma_i - \sigma_0)(\alpha \pi^l (1 - \delta_i^l) + (1 - \alpha)\pi^h (1 - \delta_i^h))$$

We capture $\tilde{\eta}_{ag}$ with a sigmoid of age and gender fixed effects, β_{ag} . The NUTS 2 fixed effects capture $(1-\pi^h)+\pi^h \sum_{i \in I} \zeta_i$. As α denotes poverty, we have

$$\beta_{poverty} = (\pi^l - \pi^h)(1 - \sum_{i \in I} \zeta_i)\sigma_0 > 0$$

With expression for Unmet in equation (5), we find that

$$\beta_{unmet} = \frac{\sum_{i \in I} \zeta_i (\sigma_i - \sigma_0) (\alpha \pi^l (1 - \delta_i^l) + (1 - \alpha) \pi^h (1 - \delta_i^h))}{\sum_{i \in I} \zeta_i (\alpha \pi^l (1 - \delta_i^l) + (1 - \alpha) \pi^h (1 - \delta_i^h))}$$

which is a weighted average of the $\sigma_i - \sigma_0 > 0$ terms. Hence, $\beta_{unmet} = E(\sigma_i - \sigma_0) > 0$.

Finally, we derive how the fraction of people that forgo treatment because it is too expensive depends on OOP. We first derive this for an increase in D. We start from

$$\frac{d\text{TooExp}}{d\text{OOP}} = \frac{d\text{TooExp}}{dD} \left(\frac{d\text{OOP}}{dD}\right)^{-1} = \frac{d\text{TooExp}}{dD} \frac{\sum_{i \in I} \zeta_i x_i}{\sum_{i \in I_{\xi}} \zeta_i \xi x_i + \sum_{i \in I_D} \zeta_i D}$$
(8)

where we use that in Europe oop payments are small relative to yearly income and hence we use the approximation that δ_i^j is constant across $i \in I$, $\delta_i^j \approx \delta_i$:

$$OOP = \frac{\sum_{i \in I_{\xi}} \zeta_{i} \xi x_{i} \delta_{i} + \sum_{i \in I_{D}} \zeta_{i} D \delta_{i}}{\sum_{i \in I} \zeta_{i} x_{i} \delta_{i}}$$

Further, equation (7) implies we can approximate the slope of TooExp with respect to D around 0 as:

$$\frac{d\text{TooExp}}{dD} = \sum_{i \in I_D} \zeta_i \left(\alpha \pi^l g_i^l \frac{\sigma_0}{\sigma_i} \frac{u'(y^l)}{u(y^l)} + (1 - \alpha) \pi^h g_i^h \frac{\sigma_0}{\sigma_i} \frac{u'(y^h)}{u(y^h)} \right)$$

where we use notation $g_i^j = g(\sigma_0/\sigma_i * u(y^j)/u(y^j - D))$ for $i \in I_D$. Further, for y^j big compared to D, we have that $g_i^l = g_i^h = g_i$:

$$\frac{d\text{TooExp}}{dD} = \sum_{i \in I_D} \zeta_i g_i \frac{\sigma_0}{\sigma_i} \left[\pi^h \frac{u'(y^h)}{u(y^h)} + \alpha \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) \right]$$

where $\left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)}\right) > 0$ because $\pi^l > \pi^h, u'(y^l) > u'(y^h)$ and $u(y^l) < u(y^h)$. A similar derivation shows

$$\frac{d\text{TooExp}}{d\xi} = \sum_{i \in I_{\varepsilon}} \zeta_i x_i g_i \frac{\sigma_0}{\sigma_i} \left[\pi^h \frac{u'(y^h)}{u(y^h)} + \alpha \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) \right]$$

Using equation (8), we find that

$$\frac{d \text{TooExp}}{d \text{OOP}} = \left[\pi^h \frac{u'(y^h)}{u(y^h)} + \alpha \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) \right] \sum_{i \in I_D} \zeta_i g_i \frac{\sigma_0}{\sigma_i} \frac{\sum_{i \in I} \zeta_i x_i \delta_i}{\sum_{i \in I_D} \zeta_i \delta_i}$$

And, similarly, for ξ :

$$\frac{d\text{TooExp}}{d\text{OOP}} = \left[\pi^h \frac{u'(y^h)}{u(y^h)} + \alpha \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) \right] \sum_{i \in I_{\varepsilon}} \zeta_i x_i g_i \frac{\sigma_0}{\sigma_i} \frac{\sum_{i \in I} \zeta_i x_i \delta_i}{\sum_{i \in I_{\varepsilon}} \zeta_i \delta_i x_i}$$

Note that we can write

$$\sum_{i \in I_D} \zeta_i g_i \frac{\sigma_0}{\sigma_i} \frac{\sum_{i \in I} \zeta_i x_i \delta_i}{\sum_{i \in I_D} \zeta_i \delta_i} = \left(\sum_{i \in I} \zeta_i x_i \delta_i\right) \sum_{i \in I_D} \frac{\zeta_i \delta_i}{\sum_{i \in I_D} \zeta_i \delta_i} \frac{g_i \frac{\sigma_0}{\sigma_i}}{1 - G_i} = \left(\sum_{i \in I} \zeta_i x_i \delta_i\right) \kappa$$

where $\kappa > 0$ denotes the parameter of the Pareto distribution:

$$1 - G(\nu) = (\underline{\nu}/\nu)^{\kappa}$$

if $\nu \geq \underline{\nu}$ and 1 other wise. With this distribution we have $g(\nu)\nu/(1-G(\nu)) = \kappa$ for each $\nu > \underline{\nu}$. It is routine to verify that we get the same expression for the expansion with respect to ξ : a weighted average of the parameter κ .

Hence, irrespective of whether we expand with respect to D or ξ , we find that

$$\frac{d\text{TooExp}}{d\text{OOP}} = \kappa \left[\pi^h \frac{u'(y^h)}{u(y^h)} + \alpha \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) \right] \sum_{i \in I} \zeta_i x_i \delta_i$$

Using this, we estimate the following linear expansion TooExp = $b_0 + \frac{d\text{TooExp}}{d\text{OOP}}$ OOP:

$$TooExp_{2t} = b_{0,2} + b_{0,t} + OOP_{ct}\bar{x}_{ct} \left(b_{oop,c} + b_{interaction,c} Poverty_{2t}\right)$$
(9)

where

$$b_{oop,c} = \kappa \pi^h u'(y^h) / u(y^h) > 0$$

and

$$b_{interaction,c} = \kappa \left(\pi^l \frac{u'(y^l)}{u(y^l)} - \pi^h \frac{u'(y^h)}{u(y^h)} \right) > 0$$

As it is hard to know what determines the intercept for this linear expansion, we allow it to vary with NUTS 2 region and calendar year: $b_0 = b_{0,2} + b_{0,t}$. Finally, to facilitate the estimation of this equation we assume that TooExp has a logit-normal distribution. That is, the log-odd of TooExp is normally distributed with the mean given by equation (9). This ensures that TooExp in the estimation always lies between 0 and 1.

Q.E.D.

B DATA

The data on population and deaths come from Eurostat's regional demographic statistics. Table 4 shows the dimensions over which our variables vary: country, NUTS 2, calendar time, age and sex. We also present a clickable link to the variable on the Eurostat website for ease of reference. The file ./getting_data.org presents the code to download the Eurostat data.⁵

Table 4: Variables and the dimensions over which they vary.

variable	country	NUTS 2	time	age	sex	missing	reference
population		X	X	X	X		link
deaths		X	X	X	X		link
at-risk-of-poverty		X	X			X	link
material deprivation		X	X			X	link
fraction too expensive		X	X			X	link
unmet		X	X			X	link
out-of-pocket	X		X			X	link
voluntary	X		X			X	link
expenditure per head	X		X			X	link

The variables on poverty, deprivation and access to care (unmet and too expensive) come from the EU statistics on income and living conditions (EU-SILC) survey.

⁵This file can be found on: https://github.com/janboone/out_of_pocket_payments_and_health.

From the Eurostat Glossary: "The at-risk-of-poverty rate is the share of people with an equivalised disposable income (after social transfers) below the at-risk-of-poverty threshold, which is set at 60 % of the national median equivalised disposable income after social transfers. This indicator does not measure wealth or poverty, but low income in comparison to other residents in that country, which does not necessarily imply a low standard of living. The equivalised disposable income is the total income of a household, after tax and other deductions, that is available for spending or saving, divided by the number of household members converted into equalised adults; household members are equalised or made equivalent by weighting each according to their age, using the so-called modified OECD equivalence scale."

"Material deprivation refers to a state of economic strain and durables, defined as the enforced inability (rather than the choice not to do so) to pay unexpected expenses, afford a one-week annual holiday away from home, a meal involving meat, chicken or fish every second day, the adequate heating of a dwelling, durable goods like a washing machine, colour television, telephone or car, being confronted with payment arrears (mortgage or rent, utility bills, hire purchase instalments or other loan payments)." Our variable "material deprivation" equals the share of people in a NUTS 2 region in material deprivation.

Fraction of people with self-reported unmet needs for medical examination is based on the same survey. In particular, the definition of this item is "Self-reported unmet needs for health care: Proportion of people in need of health care reporting to have experienced delay in getting health care in the previous 12 months for reasons of financial barriers, long waiting lists, distance or transportation problems.". We use both the general definition of unmet needs and the specific reason that treatment was too expensive.

We characterize how generous a health insurance system is using the variable OOP in our analysis. This variable is derived from data on health care expenditure by financing scheme. For our OOP measure we focus on voluntary healthcare payment schemes (voluntary) and household out-of-pocket payment (out-of-pocket). Both measured as share of total current health expenditure.

Expenditure per head refers to healthcare expenditure per head at the country level.

C ESTIMATION

This section presents the trace plots for the baseline model, the table with the relevant coefficients and the derivation of the change in mortality as a function of the change in oop.

C.1 trace plots

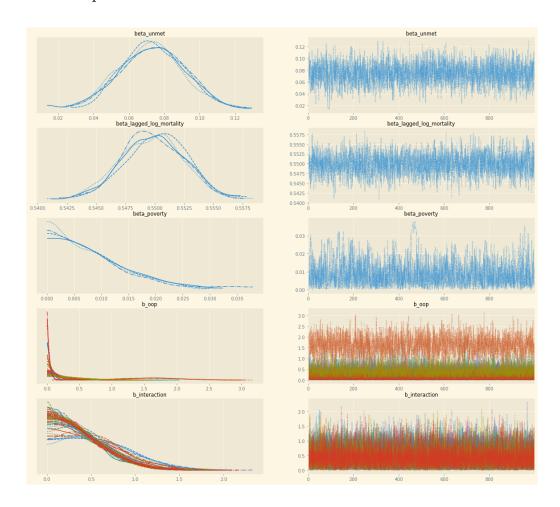


Figure 7: Trace plots of the coefficients of interest

C.2 table of coefficients baseline model

C.3 derivation of the effect of oop on mortality

As we assume that TooExp has a logit-normal distribution, the derivative of the expression in Lemma 1 with respect to $OOP\bar{x}$ is given by

$$\frac{dTooExp}{d(OOP\bar{x})} = TooExp(1 - TooExp)(b_{oop,c} + b_{interaction,c}Poverty_{2t})$$

Table 5: Summary statictics for estimated coefficients

	mean	sd	$hdi_3\%$	hdi_97%	ess_bulk	r_hat
beta_unmet	0.07	0.02	0.04	0.11	1773.00	1.00
beta_lagged_log_mortality	0.55	0.00	0.55	0.56	1815.00	1.00
beta_poverty	0.01	0.01	0.00	0.02	922.00	1.00
$b_oop[Bulgaria]$	0.20	0.18	0.00	0.53	2458.00	1.00
$b_oop[Croatia]$	0.36	0.28	0.00	0.86	2715.00	1.00
$b_oop[Denmark]$	0.12	0.12	0.00	0.34	1755.00	1.00
$b_oop[Finland]$	0.20	0.17	0.00	0.53	1594.00	1.00
$b_oop[Greece]$	1.60	0.44	0.81	2.45	1737.00	1.00
$b_oop[Hungary]$	0.18	0.17	0.00	0.50	2719.00	1.00
$b_oop[Ireland]$	0.24	0.20	0.00	0.61	2227.00	1.00
$b_oop[Lithuania]$	0.30	0.25	0.00	0.76	2645.00	1.00
$b_oop[Norway]$	0.05	0.05	0.00	0.15	2644.00	1.00
$b_oop[Romania]$	0.26	0.21	0.00	0.63	2982.00	1.00
$b_oop[Slovakia]$	0.26	0.22	0.00	0.67	2158.00	1.00
$b_oop[Slovenia]$	0.33	0.27	0.00	0.82	3299.00	1.00
$b_oop[Sweden]$	0.09	0.08	0.00	0.24	1549.00	1.00
$b_oop[Switzerland]$	0.03	0.03	0.00	0.09	1848.00	1.00
$b_interaction[Bulgaria]$	0.57	0.38	0.00	1.23	2575.00	1.00
$b_interaction[Croatia]$	0.40	0.29	0.00	0.92	3113.00	1.00
$b_interaction[Denmark]$	0.39	0.29	0.00	0.91	2655.00	1.00
$b_interaction[Finland]$	0.39	0.29	0.00	0.91	3291.00	1.00
$b_interaction[Greece]$	0.52	0.35	0.00	1.15	2936.00	1.00
$b_interaction[Hungary]$	0.44	0.32	0.00	1.00	2706.00	1.00
$b_interaction[Ireland]$	0.39	0.30	0.00	0.91	2733.00	1.00
$b_interaction[Lithuania]$	0.39	0.29	0.00	0.91	2526.00	1.00
$b_interaction[Norway]$	0.38	0.29	0.00	0.93	2961.00	1.00
$b_interaction[Romania]$	0.44	0.31	0.00	0.99	2758.00	1.00
$b_interaction[Slovakia]$	0.38	0.29	0.00	0.90	2650.00	1.00
$b_interaction[Slovenia]$	0.40	0.30	0.00	0.93	3301.00	1.00
$b_interaction[Sweden]$	0.38	0.29	0.00	0.90	2216.00	1.00
$b_interaction[Switzerland]$	0.39	0.29	0.00	0.92	2928.00	1.00

In the simulation we work with a 500 euro increase in oop: $d(OOP\bar{x}) = 500$. We assume that the increase in TooExp translates one-for-one in an increase in Unmet. Hence, the change in mortality is given by:

$$\frac{dm_{ga2t}}{m_{ga2t}} = \beta_{unmet} TooExp(1 - TooExp)500(b_{oop,c} + b_{interaction,c} Poverty_{2t})$$

This is the increase in deaths per one dead. In Figure 2 we multiply this expression by 1000: number of deaths per 1000 dead.

D ROBUSTNESS ANALYSIS

- different definition oop: include vol. ins.
- different defition poverty: at risk of poverty
- separate TooExp effect from Unmet in mortality equation