

# Social Preferences for Health Care Interventions: The SoPHI (“Social Preferences for Health Care Interventions”) Study

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## Introduction & Background

### Empirical Evidence on “Social Value Drivers”:

#### Attributes of the Health Condition

- individual valuation of health conditions
- severity of the condition
- unmet medical need
- urgency of an intervention
- capacity to benefit from an intervention

#### Attributes of the Persons Afflicted

- non-discrimination (and claims-based approaches)
- age (and fair innings)
- other patient attributes
- fairness objectives

Based on a comprehensive literature review, the Swiss HTA project, and an expert consensus on valuation principles for ultra-rare disorders, we developed the protocol of the “Social Preferences for Health Care Interventions” or **SoPHI Study** as a discrete choice experiment (DCE).

### Limitations of the Relevant Literature:

- limited in size and / or scope
- impaired by framing effects and unstable preferences
- sometimes not choice-based experiments
- sometimes of questionable methodology
- often imposed a “zero sum” assumption
- ex ante* severity of health state probably best documented attribute (“contextual variable”), but distinct difficulties to quantify effects

#### Cost Attribute (payment vehicle in most studies)

Typically reflecting an individual health state valuation (/WTP) perspective, whereas citizens’ “social WTP” for coverage of health care programs under a collectively financed health scheme might be more relevant

## Governance

### ESPM (European Social Preference Measurement)

#### Project Group: Scientific Steering Committee

- Silvio Garattini** (Mario Negri Institute, Milan / Italy)
- Sören Holm** (U of Manchester / England)
- Peter Kolominsky** (U of Erlangen / Germany)
- Deborah Marshall** (U of Calgary / Canada)
- Erik Nord** (U of Oslo / Norway)
- Ulf Persson** (IHE, Lund / Sweden)
- Maarten Postma** (U of Groningen / The Netherlands)
- Jeffrey Richardson** (Monash U, Melbourne / Victoria)
- Michael Schlander** (DKFZ & U of Heidelberg / Germany)
- Steven Simoons** (U of Leuven / Belgium)
- Oriol de Sola-Morales** (IISPV, Barcelona / Spain)
- Harry Telser** (Polynomics / Switzerland)
- Keith Tolley** (Tolley HE, Buxton / England)
- Mondher Toumi** (U Aix-Marseille / France)

## Objectives & Methods

### Primary Study Objective(s):

- To investigate how the **general public values selected attributes of health care interventions** (incl. their interaction), with special emphasis on prevalence (“rarity”), and
- to assess the **sensitivity of preferences to potential framing effects** (the level reflection and information offered to respondents).

### Design Elements:

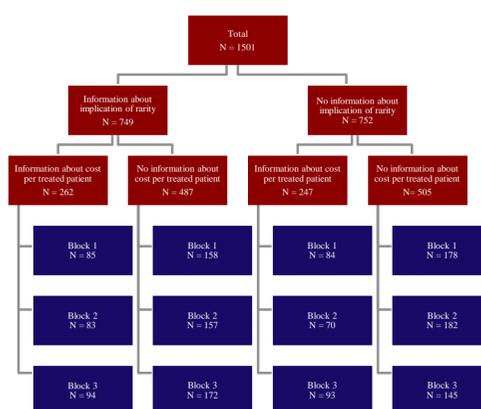
- Representative Swiss population sample** (n=1501)
- Discrete Choice Experiment (DCE) design**
- Perspective on costs:** WTP<sub>public</sub> as payment vehicle
- Utility comparator:** health states derived from EQ-5D-5L
- Initial “**Preference Formation Phase**”
- Additional **socioeconomic questions**
- Testing for potential cognitive overload**
- Pre-tests: qualitative (n=10, “think aloud”) and quantitative (n=201)
- Econometric evaluation**

### Study Population

#### and Subsamples (to control for framing effects):

Respondents (n=1,501) were randomized into 2 x 2 groups, which differed

- by one additional item to reflect on the implications of prevalence (*rarity*; 1:1), and
- by information on implied extra cost *per patient* of new treatment (1:2):

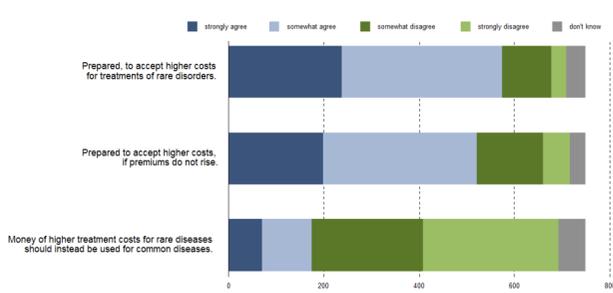


### Attributes included in SoPHI Study:

- Severity** of the initial health state: life expectancy (i.e., *ex ante*, before / without an intervention)
- Severity** of the initial health state: health-related quality of life (i.e., *ex ante*, before / without an intervention)
- Effectiveness** of intervention (“new treatment”): life expectancy gained
- Effectiveness** of intervention (“new treatment”): health-related quality of life gained
- Age** of patients (or “fair innings”)
- Rarity** of disorder (i.e., prevalence or number of persons benefitting)
- Incremental Cost** of intervention (“new treatment”): perspective of a compulsory health scheme (OKP), i.e., payment vehicle = social willingness-to-pay

## Primary Results & Key Observations

### Stated Preferences regarding Rare Disorders:



Stated Preferences during Preference Formation Phase [regarding acceptance of higher cost for rare disorder treatments; subsample randomized to reflect on the statements depicted above]

### The “Rarity” Attribute and Framing Effects:

The valuation of an intervention decreased with decreasing prevalence of the disorder. This effect was smaller than the decrease of prevalence, so that **by implication the accepted cost per patient increased with rarity.**

In order to increase awareness of respondents of the high cost per patient in the rare and ultra-rare cases, we enhanced the Main Survey by a subgroup with additional information on implied cost per patient – which, however, in the main survey had a small impact on valuation only.

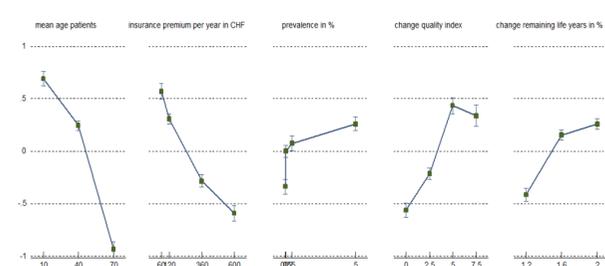
Reflection on the implications of “rarity” influenced the valuation of the attribute, i.e., respondents who were exposed to more information showed a smaller decrease of utility with decreasing prevalence.

### Model Selection:

We estimated a separate model for each attribute, investigating how well a linear model specification approximates the flexible function of the dummy model:

#### Flexible Functional Form (with dummy variables):

The figure below illustrates the point estimates with 95% confidence intervals of the change in utility for each attribute level.



The variables *mean age of patients* and *prevalence [%]* required a nonlinear variable specification.

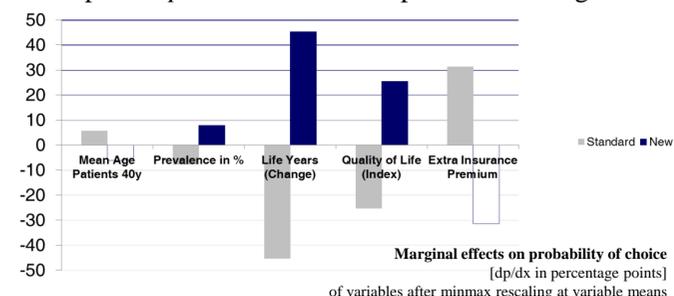
According to the Main Model that we specified, the marginal utility for an additional year of life is decreasing with the total number of years.

#### Interaction Effects:

Inclusion of interaction effects did not improve model fit based on BIC. Therefore, we did not include interactions in our Main Model.

### Importance of Attributes:

- the marginal effect of each variable depends on the overall utility level and is not constant
- variables with the highest **impact on choice probability** were *change in remaining life years*, the *quality of life index*, and *insurance premium* per year
- the negative marginal effect for *older people* was three times larger compared to middle-aged people
- impact of *prevalence* was comparable to the age effect



## Conclusions

Our discrete choice experiment (DCE) – using a payment vehicle from the citizen’s perspective – provides empirical support for the contribution to social value by all attributes tested, **implying that the accepted cost per patient may indeed increase with decreasing prevalence.** Further studies seem warranted to confirm the observed effect in other settings.