The concept of General Consent in Switzerland and the implementation at the University Hospital Zurich, a cross-sectional study

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Summary

BACKGROUND: General Consent (GC) allows the further use of health-related data/samples for multiple, unspecified research projects and/or for the collection in databases and biobanks in Switzerland. The application of General Consent in the context of human research is regulated within the scope of the Human Research Act. At the University Hospital Zurich patients are informed about General Consent to which they can agree (GC = yes) or disagree (GC = no) to the use of their routinely collected data/samples in research. In this paper, we investigated the association of demographic and medical factors on a patient’s General Consent choice.

METHODS: In this cross-sectional study, we investigated the association of age, gender, number of visits and number of diagnoses on General Consent choice. The study population was stratified by General Consent status group (GC choice: Yes, No, Not issued) and examined by means of descriptive statistics, comparative statistics and a multinomial and logistic regression model. A p-value of 0.001 was determined as significant.

RESULTS: The female gender was found to associate with decreased odds in positive General Consent choice (<0.001) whereas age (<0.001) and number of diagnoses (<0.001) were associated with increased odds in positive General Consent choice (reference “GC = no” group). The number of visits (<0.001) as well as the number of diagnoses associated (<0.001) with increased General Consent collection (increase in positive as well as negative General Consent status).

CONCLUSION: General Consent is an innovative concept that simultaneously informs patients about human research in accordance with Swiss regulations and promotes research with routinely collected data and biological samples in an era with large information repositories. Our results show that medical and demographic factors may influence a patient’s choice. Therefore, approaching these populations and taking additional care to adequately inform and ensure ethical conformity and behaviour is essential. Flexible communication channels may help us reach this goal.

Background

Informed consent (IC) is one of the core elements of clinical research which is reflected in national legislations around the world, in addition to international guidelines such as the International Conference on Harmonization - Good Clinical Practice (ICH-GCP) [1]. A patient’s participation in human research projects requires sufficient or full information on all relevant aspects of the specific project followed by the voluntary, verbal and written confirmation of the subject’s willingness [2]. As the gold standard of consent, IC is used in the majority of human research projects around the world [3].

The concept of General Consent (GC) was developed as a consequence of technological advancement and the growth of large data/sample repositories [4]. With databases and biobanks at the disposal of researchers, the collection of specific Informed Consent, for every patient and every project, is extremely resource intensive and time consuming [5, 6]. In Switzerland, General Consent is considered a comprehensive “one-time” consent which allows the use of a patient’s routinely collected health-related data/samples for future unspecified research projects and for its storage in databases and biobanks [7–9].

At the University Hospital Zurich (USZ), General Consent was strenuously implemented as a consequence of the Federal Act on Research involving Human Beings (HRA, Human Research Act) which came into force in 2014 [7]. It has become a standard element of the admissions process for in- and outpatients in every clinic. All patients at the USZ are informed about General Consent by their clinic and can autonomously and without consequence to medical care, agree (GC = yes), or disagree (GC = no) to the further use of their data/samples in human research pro-
projects. Both Informed Consent and General Consent include documentation by means of written, signed, and dated consent forms followed by approval of the specific research project by the local cantonal ethics committee (EC).

The advantages of General Consent include uncomplicated collection and use of routinely collected data and samples. This promotes development of new approaches and technologies, the expansion of new information sources, and the establishment of data and sample networks—a big step forward for cooperative research [6]. Nevertheless, the implementation of such liberal consent process has always raised the important question of adequate patient protection and the ethical conformity of these projects [4, 10, 11]. Various research groups such as Brown et al. or Cassidy et al. have thoroughly investigated the different factors (age, gender, and ethnicity) influencing Informed Consent choice and patient recruitment into clinical trials [12, 13]. However, none have investigated the factors influencing a patients General Consent choice concerning routinely collected data in unspecified research projects. Therefore, the aim of this paper will be to investigate the demographic and medical factors influencing General Consent choice in USZ patient population.

Methods

This cross-sectional study was conducted at the University Hospital Zurich, Switzerland. We used completely anonymous data which does not require informed consent by participants or approval by the ethics committee. This conforms to both local law and research policies of the University Hospital Zurich [14]. Our study adhered to STROBE guidelines (STrengthening the Reporting of OBServational studies in Epidemiology) [15].

Study population

Our dataset was derived from the clinical record system, KISIM, in which all records from 01/2018 to 07/2019 were included (figure 1). We included all inpatients and outpatients, from any clinical unit.

Outcomes, exposures and data management

 Routinely collected health data (age, gender, diagnosis, number of visits, and General Consent choice) were anonymously extracted from the clinical data platform, exported to raw data files, which in turn were imported into a separate database management system. Number of visits is defined as in- and outpatient visits to the USZ after the first consultation. We used structured query language (SQL) statements for database management and processing so that only minor data restructuring steps were necessary during statistical analysis.

Depending on the patient’s choice, they were assigned to one of the three General Consent status groups: Yes, No, or Not issued. Not issued General Consents (GC status = not issued) included forms which were not handed out to the patients or were not returned to hospital administration. General Consent status was used to stratify the population and perform comparative statistical analysis to describe the sample characteristics according to their category (figure 1).

General Consent collection at the University Hospital Zürich

Clinical departments at the University Hospital Zürich have similar processes for collection of General Consent. Generally, data derived from the clinical record system (KISIM) showed that clinical departments with high patient turnover had lower General Consent collection rates (GC not issued) due to increased workload. Collection rates were also highly influenced by consultation time (time spent with patient at first contact) as well as the effort and organisation of the administrative teams. Furthermore, departments with vulnerable patient populations showed low General Consent collection rates. This included the geriatric department, maternity ward, and the psychiatric department.

Monthly internal General Consent quality assessment reports derived from the clinical record system (KISIM) only showed slight differences in General Consent behaviour between in- and outpatients. Since January 2021, the monthly General Consent “yes” proportion was stable at 83% for outpatients and 83–86% for inpatients, respectively. The process for General Consent collection is re-evaluated in all departments on a regular basis and corrective measures are implemented. Internal quality documents provided by the General Consent project management team of the USZ suggest that the General Consent collection rate is increasing in most departments of the USZ.

Statistical analysis

We investigated the potential associations between demographic and medical factors and General Consent status, using multinomial and logistic regression models (tables 1, 2 and 3). In the first model, the “GC =not issued” group was used as the reference group whereas in the second model the “GC = no” group was used as the reference. The models analysed potential associations between the outcome, General Consent status, and the exposures which included number of diagnoses (sum of ICD-10 codes and free text diagnoses using name mapping tables), visits (included both inpatient and outpatient stays), gender, and

![Figure 1: General Consent data processing workflow.](image-url)
age, in which the latter was calculated in decades (age/10). At the USZ, General Consent collection is known to be dependent on the “clinic of admission”. We controlled (co-founder) for the 43 clinics in our model. We conducted all tests as 2-sided and determined p-values of ≤0.001 to be indicative for statistical significance. Statistical analyses were performed using the software R, version 3.5.0 (R Foundation for Statistical Computing, Vienna, Austria) and the stargaze package [16] was used to build our regression model.

Additionally, observations were made about the distribution of General Consent status over the different age groups of the USZ patient population through descriptive statics (figures 2 and 3).

Ethics approval and consent to participate
This study used anonymous data and does not require approval of an ethics committee [14].

Results

Population overview
Overall, a total of 239,733 patients were included in our analysis (figure 1) of which the majority had not been issued a General Consent (GC = not issued, 51.3%). 116,691 (48.7%) patients signed a General Consent of which 79.7% had agreed (GC = yes) and the minority, 20.2%, had declined (GC = no) to the use of their data/samples. The total population consisted of 125,668 female (52.4%) and 114,065 male (47.6%) individuals in which the mean age, number of diagnoses and number of visits were 48.34 years (SD: 21), 2.33 diagnoses (SD: 2.43) and 1.69 readmission (SD: 2.39), respectively (table 4).

Multinomial logistic regression models
Respondents in our study population visited one of the 43 different USZ clinics, who were responsible for the collection of General Consent from their patients. Patients were treated predominantly in the ophthalmology department (23,987, 10%), the dermatology department (24,017, 10%) and the emergency department (22,724, 9.5%). The regression models (tables 1, 2 and 3) controlled for the 43 different inter-clinic collection rates.

Higher number of diagnoses and visits lead to a statistically significant increase in the collection of General Consent (GC = no and GC = yes) (tables 1 and 2). The female gender was found to associate with decreased odds in positive General Consent choice (more GC = no) (table 3) whereas age and number of diagnoses was found to associated with increased odds in positive General Consent choice (more GC = yes) (table 3).

Age distribution
Multinomial logistic regression predicts that higher age associates with increased odds in positive General Consent choice (tables 1 and 3, figure 2).

Further investigation shows, there are differences in General Consent choice and collection depending on the age group (figures 3 and 4). Overall, more General Consents were not issued in younger patients (<18 years) and geriatric patients (80+) (figures 3 and 4).

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR</th>
<th>CI</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (in 10 year intervals)</td>
<td>1.07</td>
<td>1.06; 1.08</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Sex (female)</td>
<td>1.02</td>
<td>1.00; 1.04</td>
<td>0.018</td>
</tr>
<tr>
<td>Number of diagnoses</td>
<td>1.55</td>
<td>1.54; 1.57</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Visits</td>
<td>1.11</td>
<td>1.10; 1.11</td>
<td>&lt;0.001</td>
</tr>
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</table>

Reference group: “not issued”

CI = confidence interval; OR = odds ratio of positive General Consent choice (consented)

<table>
<thead>
<tr>
<th>Variable</th>
<th>OR</th>
<th>CI</th>
<th>p value</th>
</tr>
</thead>
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<tr>
<td>Age (in 10 year intervals)</td>
<td>1.01</td>
<td>1.00; 1.02</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Sex (female)</td>
<td>1.23</td>
<td>1.25; 1.33</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Number of diagnoses</td>
<td>1.52</td>
<td>1.51; 1.53</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Visits</td>
<td>1.10</td>
<td>1.09; 1.12</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Reference group: “not issued”

CI = confidence interval; OR = odds ratio of negative General Consent choice (declined)

<table>
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<tr>
<th>Variable</th>
<th>OR</th>
<th>CI</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (in 10 year intervals)</td>
<td>1.05</td>
<td>1.0; 1.07</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Sex (female)</td>
<td>0.79</td>
<td>0.76; 0.81</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Number of diagnoses</td>
<td>1.02</td>
<td>1.02; 1.03</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Visits</td>
<td>1.00</td>
<td>1.00; 1.01</td>
<td>0.013</td>
</tr>
</tbody>
</table>

Reference group: “declined”

CI = confidence interval; OR = odds ratio of positive General Consent choice (consented)
Discussion

In this cross-sectional study, we used patient data stored in the clinical information system of the University Hospital Zurich from 01/2018 to 07/2019 to investigate the association of demographic and medical factors on General Consent choice. The female gender associated with increased negative General Consent choice whereas age was associated with positive General Consent choice. Additionally, number of diagnoses and visits were predictive for both positive as well as negative General Consent status.

The use of routinely collected data/samples in research is becoming increasingly more important and will continue to do so in the future. General Consent is a simplified consent process, in which the patients agree to the further use of their data/samples for multiple different research projects, which are deemed low-risk [7]. General Consent simplifies the conduct of research projects with data/sam-

![Figure 2: Age in General Consent status groups.](image2)

![Figure 3: General Consent status distribution in age categories.](image3)
amples and ultimately leads to the promotion of cooperative research, as patients no longer need to consent to each project specifically. The difficulty lies in improved access to health-related data/samples and simultaneously maintaining patient privacy, rights, and ethical conformity [11]. Investigation of the factors influencing a patient’s General Consent choice may help improve the information of these target groups in future.

Previous literature has found that women are historically under-represented in clinical research [17], which is corroborated by our results. In clinical trials the concerns of these women included lack of information, strenuous study procedures, interference with personal life and a discomfort with the clinical environment [12]. Since collection of General Consent does not require additional visits from patients, previous research may have underestimated the other effects connected to patient consent such as lack of information (on data privacy, regulations, and general science), incomprehension, and distrust in clinical institutions.

Additionally, our investigation showed that older patients were more willing to say yes to General Consent, contrary to previous findings [18]. Studies have shown that there are challenges recruiting elderly patients with the most frequent concerns found to be travelling to the investigative institution [13]. Since General Consent does not require additional travel, it may explain the increased willingness of these patients to participate in General Consent studies. Clinical trials in older patients are often difficult to conduct due to comorbidities, co-medications and differences in pharmacokinetics and pharmacodynamics [13]. The use of these data/samples in research may prove critical in overcoming these challenges in future. An additional factor to consider may be that younger patients are more aware of the data/sample privacy risks, which may lead to the decline in positive General Consent choice in comparison to the older population [19,20]. General Consent choice may also vary in different age groups. Our results show there are higher rates of “not issued” General Consents in very young (1-10 years) and old patients (+85) whose consent forms are often signed by a parent or surrogate. The signature of surrogates raises not only a practical issue but also an ethical concern which we wish to investigate in future.

Furthermore, our results suggest that multi-morbid patients seem more likely to agree to General Consent which may be explained through the increased medical severity in this patient population. This is a positive step forward as pa-

Table 4:
Baseline characteristics for General Consent status groups consented, declined, and not issued (total n = 239,733).

<table>
<thead>
<tr>
<th>Variable</th>
<th>Consented</th>
<th>Declined</th>
<th>Not issued</th>
</tr>
</thead>
<tbody>
<tr>
<td>n (%)</td>
<td>93,089 (38.8)</td>
<td>23,602 (9.8)</td>
<td>123,042 (51.3)</td>
</tr>
<tr>
<td>Age (median, IQR)</td>
<td>52.00 [36.00, 68.00]</td>
<td>47.00 [34.00, 62.00]</td>
<td>45.00 [30.00, 62.00]</td>
</tr>
<tr>
<td>Sex (n, %)</td>
<td>Male 44,732 (18.7)</td>
<td>9,643 (4.0)</td>
<td>59,690 (24.9)</td>
</tr>
<tr>
<td></td>
<td>Female 48,357 (20.1)</td>
<td>13,959 (5.8)</td>
<td>63,352 (26.4)</td>
</tr>
<tr>
<td>n diagnoses</td>
<td>2.00 [1.00, 4.00]</td>
<td>2.00 [1.00, 3.00]</td>
<td>1.00 [1.00, 2.00]</td>
</tr>
<tr>
<td>Visits</td>
<td>1.00 [1.00, 2.00]</td>
<td>1.00 [1.00, 2.00]</td>
<td>1.00 [1.00, 2.00]</td>
</tr>
</tbody>
</table>

n = number, *IQR = Interquartile range

Figure 4: General Consent status in all ages.
tients with multiple diagnoses are preferably not recruited into clinical trials [21]. Number of visits was found to associate with both negative as well as positive General Consent choice. This is in line with USZ General Consents processes. General Consent is usually signed by the patients at their first visitation and if this does not occur the form will be handed out at the next visitation. In the context of our investigation, we would also like to discuss the contemporary ethical and legal concerns regarding General Consent. Current discussions about the conformity of the General Consent have raised the question if patients are being adequately informed about the use of their data and samples in research [10, 11]. However, increasing the length, complexity, legal and medical jargon of consent forms is known to decrease understanding amongst patient populations [22]. We therefore suggest a targeted and individual nationwide communication concept, such as online information platforms and “expert chats” for specific and/or general questions regarding the patient’s interest. A national effort will increase the trust in the scientific community, identify patient concerns as well as find shortcomings in the General Consent collection processes. All communication tools and platforms should explicitly and coherently inform patients about the general concepts of research and science; the benefits of this research; give an overview on current projects working with General Consent data/samples; have detailed explanations on the risks of data privacy and security; and include up to date information on ethical, legal, and regulatory requirements. Furthermore, the collection of General Consent requires a large administrative effort which has been established through hospital processes. This study may serve as a basis to approach these target groups and increase interest and participation of the general population in research projects. The study we conducted has several strengths: To our knowledge, this is one of the first large and systematic investigations looking into the factors influencing General Consent choice in a population. We considered in- and out-patients from all clinics of the USZ over a period longer than a year, thus limiting selection bias. The exposures were investigated in a very large sample size, which decreased the margin of error. There are however limitations: due to the large sample size of our data, we may have found small effects, which are not medically or practically relevant. We did not investigate if these patient populations are also underrepresented in actual human research projects. Furthermore, our data did not include other important confounders such as nationality, religion, and profession. In a subsequent study we wish to investigate additional exposures such as nationality and socio-economic factors and the impact of the COVID-19 pandemic on General Consent behaviour. If feasible, we will also include a second university hospital into the analysis to increase the generalisability of our findings. Further areas of interest include if this effect is seen in other countries, investigation of the patients’ comprehension of General Consent, and important efforts to increase comprehension through adaptive consent designs such as dynamic and eConsent [23].

Conclusion

In conclusion, General Consent is a necessary development towards a future in which large repositories of data and samples will be accessible for research. It will be a fine line navigating patient protection and privacy regulations in addition to promoting research with a simplified consent process. Investigation into the factors influencing General Consent may improve the targeted information these patient populations may receive as well as guarantee ethical conformity.

Availability of data and materials section

The datasets generated and/or analysed during the current study are not publicly available due to the protection of patient data but are available from the corresponding author on reasonable request.

Acknowledgements

Authors’ contributions: AG structured and acquired the data, completed the statistical analysis and designed the graphics/figures. AB supported AG and collected additional information. RG supervised the planning, conduct and completion of the project. FJ verified legal and regulatory statements made in manuscript. AG and AB wrote the first draft of manuscript and FJ and RG thoroughly reviewed and edited it. AB made all changes for the review process. We thank Cyril Simmen for the initial review of the manuscript and we thank Prof. Gabriela Senti for the final review.

Financial disclosure

The project did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Potential competing interests

All authors have completed and submitted the International Committee of Medical Journal Editors form for disclosure of potential conflicts of interest. No potential conflict of interest was disclosed.

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