

# Patient experiences with interdisciplinary treatment for substance dependence: an assessment of quality indicators based on two national surveys in Norway

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**Purpose:** The quality of health care is often measured using quality indicators, which can be utilized to compare the performance of health-care providers. Conducting comparisons in a meaningful and fair way requires the quality indicators to be adjusted for patient characteristics and other individual-level factors. The aims of the study were to develop and test a case-mix adjustment model for quality indicators based on patient-experience surveys among inpatients receiving interdisciplinary treatment for substance dependence, and to establish whether the quality indicators discriminate between health care providers.

**Patients and methods:** Data were collected through two national surveys involving inpatients receiving residential treatment in Norway in 2013 and 2014. The same questionnaire was used in both surveys, and comprised three patient-experience scales. The scales are reported as national quality indicators, and associations between the scales and patient characteristics were tested through multilevel modeling to establish a case-mix model. The intraclass correlation coefficient was computed to assess the amount of variation at the hospital-trust level.

**Results:** The intraclass correlation coefficient for the patient-reported experience scales varied from 2.3% for “treatment and personnel” to 8.1% for “milieu”. Multivariate multilevel regression analyses showed that alcohol reported as the most frequently used substance, gender and age were significantly associated with two of the three scales. The length of stay at the institution, pressure to be admitted for treatment, and self-perceived health were significantly related to all three scales. Explained variance at the individual level was approximately 7% for all three scales.

**Conclusion:** This study identified several important case-mix variables for the patient-based quality indicators and systematic variations at the hospital-trust level. Future research should assess the association between patient-based quality indicators and other quality indicators, and the predictive validity of patient-experience indicators based on on-site measurements.

**Keywords:** quality of health care, health care quality indicator, case-mix adjustment, patient satisfaction, survey

## Introduction

The systematic measurement and reporting of quality indicators in health care have become common in several countries during the last few decades.<sup>1,2</sup> The reasons for the increased focus on health care performance are often linked, and include both the increasing expense and the variations in the quality of health care.<sup>3</sup> Patient-reported experiences (also called user experiences) constitute parts of the frameworks for quality indicator projects for the Nordic Council, the OECD, and the WHO.<sup>2,4,5</sup> The quality

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indicators used in Norway are meant to reflect aspects of the quality of health care, as a means to inform policy-makers, health care leaders, health personnel, and patients about health care performance.<sup>6</sup> The Norwegian quality indicator system consists of several indicators of the quality of health care within different areas, using both patient administrative data and patient-reported experiences.<sup>7</sup>

Patient-experience surveys elicit a view of the quality of health care that cannot be provided by other clinical or administrative assessments. When measuring patient experiences on a national or other large scale, it is useful to be able to compare between health care providers or hospital trusts. While such comparisons may be informative when choosing a provider or allocating resources, patient characteristics not related to the quality of health care may influence how certain patient groups evaluate this quality. This makes it necessary to develop and apply appropriate case-mix adjustments to ensure that such comparisons are valid.<sup>8–10</sup> In addition, different patient groups have different probabilities of responding to the surveys, which results in some groups being underrepresented in the resulting sample. A common approach to compensate for nonresponse is to weight data to reduce bias related to certain groups being underrepresented.<sup>11</sup> Weighted results are meant to be more representative for the complete patient population; that is, not only to those who actually replied to the survey. The aim of making case-mix adjustments is to increase the accuracy of the results when comparing different health care providers.<sup>10</sup>

There is no apparent consensus on how to develop or use quality indicators, with the approach often being determined by the objective of performing particular measurements.<sup>6</sup> However, one step-by-step guide provided by the Agency for Healthcare Research and Quality suggests that the performance of such quality indicators should be judged according to: face validity, precision, minimum bias, construct validity, fostering real quality improvement, and application,<sup>12</sup> each of these factors being related to data quality, validity, reliability, applicability, and feasibility.<sup>6</sup>

In 2013, the Norwegian Directorate of Health commissioned a national patient-experience survey involving inpatients receiving interdisciplinary treatment for substance dependence.<sup>13</sup> In Norway, the regional hospital trusts are decreed by law to offer interdisciplinary treatment for substance dependence. This is a service for patients with substance dependence that include both medical, psychological and social work professionals, and the treatment focuses on both patients' health problems and social situations. The

service includes residential, day, polyclinic and ambulatory services.<sup>14</sup> In the agreement between that directorate and the Norwegian Institute of Public Health (NIPH), the directorate underlined that it wanted to test the possibility of developing quality indicators based on the national survey. The NIPH is responsible for conducting national patient-experience surveys, and has a long history of conducting national surveys and reporting corresponding quality indicators based on patient experiences. However, after testing the data material from this population as quality indicators, the NIPH concluded that the number of respondents at each institution was too small for reporting proper indicators. It was therefore decided to expand the sample with another national survey of the same population, which was conducted 1 year later.<sup>15</sup>

The NIPH usually samples 400 respondents from each participating unit (eg, hospital) when measuring the experiences that patients have with health care. However, the institutions offering interdisciplinary treatment for substance dependence in Norway are substantially smaller than this, which necessitated some adjustments to the usual methods employed when conducting patient-experience surveys and reporting the results from these surveys. There is a growing body of literature when it comes to assessing patient experiences and patient reported outcomes with different types of treatment for substance dependence. Some of these efforts focus on developing and validating relevant tools.<sup>16–23</sup> Within the literature, some evidence can be found regarding patient characteristics that should be considered for case-mix adjustments.<sup>24–26</sup> However, several of these studies are conducted in more narrow populations than the Norwegian one, which are including all patients treated for substance dependence, hence suggesting that a more-explorative approach was needed in the current setting.

The aim of this study was to establish whether the experiences of inpatients can be used as a basis for national quality indicators of interdisciplinary treatment for substance dependence. The objective was twofold: (i) to use the data material from the national surveys to develop a feasible case-mix adjustment model, and (ii) to establish whether the quality indicator scores vary across health care providers, hence testing the ability of the indicators to discriminate between levels of patient-reported quality.

## Material and methods

### Sample and data collection

All public and private residential institutions with a contract with regional health authorities were included in the national

surveys. The included institutions represented a range of treatments, dependencies, programs, number of patients and length of stay, while detoxification institutions were excluded. The surveys were conducted as national measurements, and it was mandatory for the institutions to participate. The NIPH established contact with each of the institutions through the four regional health authorities, which provided contact information to their underlying hospital trusts, which again provided information on the leaders at all the institutions working with interdisciplinary treatment for substance dependence. The NIPH could from there establish contact with health personnel at each institution, and together with them plan and conduct the surveys. All patients staying at the institutions, aged 16 years and older, were invited to participate. Individual patients could be excluded based on ethical considerations by personnel at each institution.

In 2013, 1,245 patients were staying at the residential institutions during the data collection. Twelve of the patients were excluded due to ethical considerations, while 163 patients were not available for participation (due to, eg, doctor's visit, leave from the institution). Hence, 1,070 inpatients from 98 institutions were eligible for inclusion in 2013. In 2014, 23 patients of 1,279 were excluded due to ethical considerations, while 143 patients were not available, leaving 1,113 patients eligible for participation.

The data were collected using a cross-sectional design on a single day decided by the institutions during a single designated week decided by the NIPH. The health personnel at each institution were responsible for handing out and collecting the answered questionnaires, making this an on-site survey. The questionnaires were distributed to the institutions in prepacked envelopes containing an information sheet, the questionnaire, and a return envelope before the day of data collection. The health personnel was instructed to give one envelope to each consenting patient. Patients were informed that participation was voluntary, and that no one would know their answers to the questions. Pre-established practice with treating a filled-out and returned questionnaire as consent to participate was employed. The surveys were conducted anonymously, and carried out as a quality assurance project. No demographic information other than the background items in the questionnaire was collected. The participating institutions did not provide information to the NIPH about their respondents, other than the number of eligible patients and the reasons for

ineligibility. One of the consequences of this procedure is a lack of information regarding nonrespondents.

## Measures

The questionnaire used in the survey was the Patient Experiences Questionnaire for Interdisciplinary Treatment for Substance Dependence (PEQ-ITSD), which consists of 51 closed-ended questions. The data quality, validity, and reliability of the PEQ-ITSD have been comprehensively described, tested, and reported on previously.<sup>17</sup> Most of the questionnaire items are scored on a 5-point scale, ranging from "not at all" to "to a very large extent." The questionnaire-scale was linearly converted to a scale from 0 to 100, where a higher score indicated a better outcome. To obtain a scale score, a respondent had to answer at least half of the questions constituting each scale.

Identical versions of the questionnaire were used in the surveys performed in 2013 and 2014. The questionnaire comprises three scales: "treatment and personnel," "milieu," and "outcome", which were derived from exploratory factor analyses, and has shown good internal consistency reliability, test-retest reliability and construct validity. The variation in the three scale scores and the regression models' variance constitute the main outcomes in the following analyses. The "treatment and personnel" scale consists of 12 items concerning topics such as the relationships of patients to personnel, their access to personnel, and patient centeredness. The "milieu" scale consists of five items relating to the safety experienced by patients at the institution, how they were received when they arrived, and their opinion of the meals quality. The "outcome" scale also consists of five items, which asks the patients whether they consider that the time spent at the institution has helped them as it should and has prepared them adequately for their life after discharge.

The demographic variables collected through the respondents' replies to the questionnaire, ie, the variables tested as relevant case-mix adjusters, were: gender, marital status, education, self-perceived physical and mental health, most frequently used drug or substance prior to this admission, length of stay at this institution, number of previous admissions, age, and age when they developed the substance dependence. In addition to the mentioned variables, a new variable, mixed use, was computed. This variable corresponds to the number of substances that each respondent reported using most frequently before admission, with a possible value from 1 to 6. This variable acted as a proxy for the severity of dependence.

Given the smallness of the sample at each institution, the scales were inspected at several levels of health care. The participating institutions could be grouped into hospital trusts or private organizations as well as in regional hospital trusts. Grouping the institutions in this way allowed us to assess the standard error of the scale scores, and thereby determine the health care level at which the statistical uncertainty was too large. We used the criterion employed in other national patient-experience surveys performed in Norway, which is to compute scale scores only where the standard error is lower than 6. This criterion leads to that the hospital trust level was chosen to represent the provider level in this study, meaning public hospital trusts or private organizations with more than one underlying unit (institution).

Results of national surveys reported by the NIPH are usually weighted to address nonresponse. However, the present survey did not include this information. The results were weighted based on self-reported length of stay at the institution in order to compensate for the larger probability of being sampled when having stayed longer at the institution.

## Analysis

Due to the nested nature of the data (ie, the respondents were nested within the hospital trusts), and violation of the prerequisite that the data are statistically independent, multilevel modeling was used in all regression analyses.<sup>27</sup> For multilevel modeling to be necessary, it is often stated that the ICC should be larger than 0.01,<sup>28</sup> and that the design effect should be larger than 2.<sup>29</sup> In line with these recommendations, we calculated both the ICC and design effect.

Associations between independent variables and the scales were assessed with a merged data set containing data from 2013 to 2014. The analyses only included institutions that had respondents from both survey years. The tested variables and the resulting model were based on (i) the scientific literature, (ii) the effect of background variables on the scales, (iii) the distribution of the background variables across the institutions, and (iv) suggestions from external experts that we sought advice from.

The multilevel analyses assessed the amount of variation at the hospital-trust level. In the null model, the hospital trusts were entered as random intercepts. This model was used to calculate the proportion of the variance explained at the hospital-trust level based on the ICC. The design effect was calculated based on the mean number of responses across the hospital trusts and the ICC for each of

the scales. The full model included hospital trusts as random intercepts and all patient variables that were statistically significant on a 0.05-level with at least one of the scales in bivariate models as fixed effects. Mixed use and survey year were entered into the multivariate model regardless of results from the bivariate analyses. The variance explained by the full model was calculated as described by Snijders and Bosker.<sup>27</sup> Based on findings from the multilevel analysis, we calculated adjusted scores for all hospital trusts on the three scales. Variables that were statistically significant on a 0.05-level on at least two of the scales in the multivariate analysis were included in the case-mix model.

The statistical analyses were conducted using SPSS (version 24.0; SPSS Inc., Chicago, IL, USA) and R software.<sup>30</sup>

## Results

In total, 978 inpatients in 2013 and 1,017 inpatients in 2014 responded to the national survey, giving a total merged data set of 1,995 respondents. The response rate for both years was 91%. The analyses were applied to 1,452 respondents from 21 hospital trusts, whose sample sizes ranged from 25 to 154. Table 1 shows that the mean age in the merged sample at the hospital-trust level was 36 years, and 69% of the respondents were male. Most (82%) of the respondents reported being single and 89% had finished primary school. The mean age when the respondents had developed substance dependence was 20 years, and the most frequently used substances prior to the current admission were alcohol, cocaine/amphetamine and cannabis, closely followed by medication. Sixty-four percent of the respondents reported their physical health to be good, very good or excellent, with 56% reporting this for the question about mental health. About half of the respondents had been at the institution for <3 months, and 68% had one or more previous admissions to a residential institution.

The multilevel bivariate regression models showed that all but five variables of 16 had a significant association with at least one of the scales (results not shown). The five variables were (i) heroin/morphine as the most frequently used drug prior to admission, (ii) other as the most frequently used drug prior to admission, (iii) mixed use, (iv) marital status, and (v) education. All of the other background variables were associated with at least one of the scales at the 0.05 level. Table 2 shows the multilevel multivariate regression model and that patients reporting alcohol as their most frequently used substance before admission had significantly better experiences on the

**Table I** Sample descriptives at the hospital trust level in 2013, 2014, and both years combined

	2013		2014		2013+2014	
	n	%	n	%	n	%
Gender						
Male	450	66.0	489	71.0	939	68.5
Female	232	34.0	200	29.0	432	31.5
Marital status						
Married/cohabiting	132	19.5	116	17.0	248	18.2
Single	545	80.5	566	83.0	1,111	81.8
Education						
Primary school	282	41.7	271	40.1	553	40.9
Secondary school	321	47.4	334	49.5	655	48.4
University or college	74	10.9	70	10.4	144	10.7
Self-perceived physical health						
Excellent	45	6.6	54	7.8	99	7.2
Very good	141	20.7	146	21.2	287	20.9
Good	241	35.4	248	36.0	489	35.7
Quite good	162	23.8	171	24.8	333	24.3
Poor	92	13.5	70	10.2	162	11.8
Self-perceived mental health						
Excellent	32	4.7	40	5.8	72	5.3
Very good	107	15.7	124	18.1	231	16.9
Good	228	33.5	229	33.4	457	33.5
Quite good	190	27.9	190	27.7	380	27.8
Poor	123	18.1	103	15.0	226	16.5
Most frequently used drug/substance prior to this admission						
Alcohol	321	44.8	337	45.9	658	45.3
Medication	330	46.0	304	41.4	634	43.7
Cannabis	323	45.0	332	45.2	655	45.1
Cocaine/amphetamine	348	48.5	367	49.9	715	49.2
Heroin/morphine	200	27.9	213	29.0	413	28.4
Other	95	13.2	83	11.3	178	12.3
Length of stay at this institution						
0–2 weeks	95	13.3	96	13.2	191	13.3
3–11 weeks	264	37.1	284	39.0	548	38.1
3–6 months	203	28.5	219	30.1	422	29.3
7–12 months	107	15.0	99	13.6	206	14.3
>12 months	43	6.0	30	4.1	73	5.1
Previous admissions						
No	211	30.9	229	33.4	440	32.2
Yes, once	180	26.4	174	25.4	354	25.9
Yes, twice	130	19.1	113	16.5	243	17.8
Yes, 3–5 times	97	14.2	117	17.1	214	15.6
Yes, >5 times	64	9.4	53	7.7	117	8.6
	<b>n</b>	<b>years</b>	<b>n</b>	<b>years</b>	<b>n</b>	<b>years</b>
Age	678	35.9 (mean)	667	36.1 (mean)	1,345	36.0 (mean)
Age when substance dependence developed	669	19.6 (mean)	673	19.8 (mean)	1,342	19.7 (mean)



**Table 2** Results from multilevel regression models of the associations between independent variables and scale scores

Variable	Treatment and personnel		Milieu		Outcome	
	Estimate	P	Estimate	P	Estimate	P
Most frequently used drug/substance prior to this admission						
Alcohol	3.764	0.006	2.014	0.086	3.638	0.020
Medication	1.052	0.512	-0.747	0.586	1.724	0.345
Cannabis	-1.904	0.211	-1.503	0.248	-3.201	0.065
Cocaine/amphetamine	2.581	0.081	2.842	0.024	2.025	0.230
Mixed use <sup>a</sup>	-0.463	0.626	0.638	0.432	0.203	0.851
Length of stay at this institution <sup>b</sup>	2.249	0.000	-1.490	0.001	2.434	0.000
Pressured/forced by others to be admitted for treatment	-0.919	0.025	-1.062	0.002	-1.651	0.000
Gender						
Male	-	-	-	-	-	-
Female	2.623	0.020	0.117	0.903	3.730	0.004
Age	0.177	0.003	0.251	0.000	0.055	0.420
Age when substance dependence developed	0.044	0.537	-0.024	0.687	0.139	0.083
Self-perceived physical health <sup>b</sup>	-1.830	0.000	-1.347	0.002	-1.592	0.007
Self-perceived mental health <sup>b</sup>	-1.869	0.000	-1.596	0.000	-2.837	0.000
Year						
2013	-	-	-	-	-	-
2014	-0.352	0.726	0.072	0.933	0.169	0.882

**Notes:** <sup>a</sup>Sum of the respondent replies to which substance was most frequently used before admission, and indicates how many substances each respondent reported to have used (range =1–6); <sup>b</sup>numerical variables.

**Table 3** Summary statistics on model variance

Scale	Var( $U_{0j}$ ) <sup>a</sup>	Var( $R_{ij}$ ) <sup>b</sup>	Total variance explained (%) <sup>c</sup>	ICC <sup>d</sup>	Design effect
Treatment and personnel	7.55	298.01	7.62	0.023	2.34
Milieu	23.29	218.99	6.97	0.081	5.69
Outcome	20.69	382.26	7.72	0.048	3.90

**Notes:** <sup>a</sup>Hospital-trust-level variance. <sup>b</sup>Patient-level residual variance. <sup>c</sup>Total variance explained calculated only for the individual level. <sup>d</sup>ICC calculated from the multilevel regression null models with hospital trusts as random intercepts.

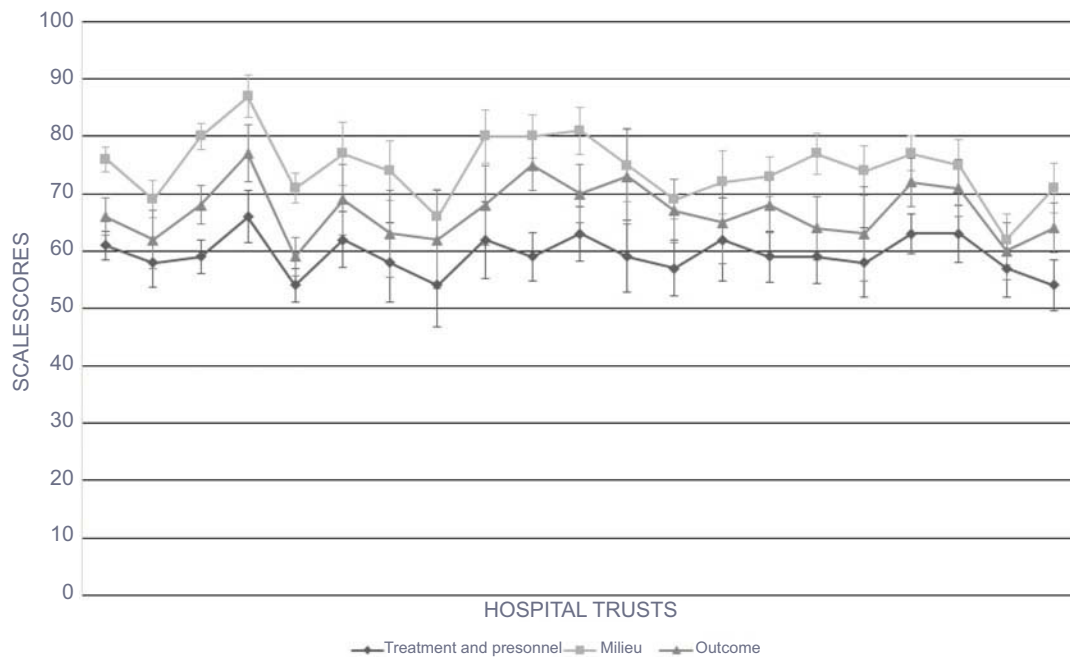
“treatment and personnel” and “outcome” scales when other factors were controlled for. Respondents with longer stays at the institutions reported better experiences with “treatment and personnel” and “outcome,” but worse experiences on the “milieu” scale. Being pressured or forced by others to be admitted for treatment had a negative impact on all three scales. Females had significantly better experiences than males regarding “treatment and personnel” and “outcome,” while older patients reported better experiences on the “treatment and personnel” and “milieu” scales. Patients reporting worse health (both physical and mental) reported worse experiences on all three scales.

Table 3 shows that the ICC varied from 2.3% for “treatment and personnel” to 8.1% for “milieu”, which indicate meaningful variance in the scale scores at the hospital-trust level, and the need for multilevel modeling.

The design effect was larger than 2 for all three scales. The proportion of the explained variance in the full model was 7.62% for “treatment and personnel,” 6.97% for “milieu,” and 7.72% for “outcome.”

Figure 1 displays the variation in results from the 21 participating hospital trusts, comprising hospital trusts or private organization with underlying departments/institutions. Each line represents one scale, while the results for each hospital trust appearing as points on the lines in the same order for all three scale scores. The scale scores for “treatment and personnel” varied from 54 to 66, while those for “milieu” varied from 62 to 87 and, those for “outcome” varied from 59 to 77. The mean national scale scores for the merged material were 61 for “treatment and personnel,” 75 for “milieu,” and 68 for “outcome.” One hospital trust obtained a score that was significantly lower than the national mean for “treatment and personnel”

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**Figure 1** Variations in mean scale scores with 95% confidence intervals across hospital trusts.<sup>a</sup>

**Note:** <sup>a</sup>Somewhat fewer respondents compared to other results due to the R analyses eliminating cases with missing data on one or more variables in the case-mix adjustment model.

( $P < 0.01$ ). For “milieu,” four hospital trusts scored lower than the national mean ( $P < 0.01$  to  $P < 0.001$ ) and two had higher scores ( $P < 0.05$  to  $P < 0.001$ ). Four hospital trusts varied significantly from the national mean for “outcome,” with two in each direction ( $P < 0.05$  to  $P < 0.001$ ).

## Discussion

This study explored the effects of patient characteristics on different aspects of the experiences of patients receiving residential treatment for substance dependence, with the aim of developing a feasible case-mix model to enable fairer comparisons between health care providers. The results showed that there is a need to adjust for several of the available variables, such as length of stay at the institution, pressure from others to be admitted for treatment, and self-perceived physical and mental health. Furthermore, alcohol reported as the most frequently used substance, being female, and increasing age were positively associated with two of the three scales. The results also showed that the indicators can be used to discriminate between health care providers.

The multilevel models showed that the scale scores should be adjusted for self-perceived physical and mental health. For both the “treatment and personnel” and “milieu” scales, comprising items related to aspects related to the residential institution only, self-perceived health

should function well as a case-mix variable. However, for the “outcome” scale, which measures the experiences of patients with aspects related to the treatment results, self-perceived health was omitted from the case-mix adjustment, even though the two variables were significantly associated with the scale. The reason for omitting self-perceived health is that both of these variables can be viewed as outcome variables, since the health of the patients might be expected to change due to them receiving treatment at the institutions.

In addition to the previously mentioned variables, the full case-mix model also included the computed variable of mixed use. Professionals in the field of substance-dependence treatment were approached for advice on the variables they thought should be included in the testing—their suggestions were age, marital status, gender, and mixed use. Mixed use was not significantly associated with any of the three scales, but the variable was kept in the model to compensate for the “outcome” scale not being adjusted for self-perceived health, and also functioning as a proxy for severity.

Age and self-perceived health have been found to be significantly associated with patient experiences and/or satisfaction in several populations, with patients who rate their health more positively and/or are older reporting satisfaction or better experiences.<sup>31–33</sup> This is consistent with the findings of the present study. Other variables that

have been demonstrated (although more inconsistently) to be associated with the experiences of patients include education, marital status, income, and gender,<sup>31–33</sup> but only gender was found to be significantly associated with the scale scores in the current study.

In line with the above-mentioned findings in other populations as well as those in the current study, a study conducted in the USA found that both self-perceived physical health and mental health were the strongest predictors of survey scores for patients receiving behavioral health care, such as treatment for mental illness or treatment for alcohol or drug use.<sup>9</sup> Other variables reported as significant were education, age, and race/ethnicity, along with whether or not the patients were treated for alcohol or drug use. Education was not significantly associated with the scale scores in the current study; however, in the Norwegian sample, few respondents reported having a university- or college-level education, and the potential effect of differences in education level could be weaker. The PEQ-ITSD currently does not include questions on race or ethnicity, so these variables were not tested in the current study.

The length of stay at the institution was significantly associated with all of the scale scores: a longer stay was associated with more-positive evaluations on both the “treatment and personnel” and “outcome” scales, but with more negative evaluations on the “milieu” scale. This might be due to patients who have stayed in an institution for a longer time becoming more familiar with the personnel and the institution as a whole, forming relationships and trust, and being able to experience some benefits from their treatment that patients with shorter stays have not yet experienced. However, patients with longer stays may also become more tired of the institution and other patients, hence affecting the “milieu” score negatively. Another possible explanation is that patients who generally are more positive and have better experiences and outcomes stay longer in treatment.

Even though length of stay has been demonstrated to be a predictor of the results in these surveys, its suitability as a case-mix adjustor has been discussed. Ideally, a case-mix adjustor should be specific to individual patients and also outside the control of health care providers.<sup>31</sup> The length of stay can clearly be influenced by health care providers, and a shorter length of stay in hospitals has been shown to negatively affect the quality of health care.<sup>34</sup> However, the point in our study is not to adjust for the actual length of stay for discharged patients, but to correct for the fact that institutions and hospital trusts at the time of measurement had different patient-mix concerning length of stay. The time of

measurement (a single day in a specific week) was decided by the external research organization, and so was outside the control of the different health care providers. Thus, differences in the length of stay at the time of measurement might be considered a valid and important case-mix adjustor in our study.

To be pressured or forced by others to be admitted for treatment had a negative impact on the scores on all the scales. The question did not distinguish who applied the pressure/force and may, therefore, be given a high score by all patients regardless of the legal status on their admission (voluntary vs involuntary) or the reason for seeking treatment. Previous research has shown that patients receiving psychiatric residential treatment who report higher levels of coercion (both perceived and documented) either at admission or during treatment report lower satisfaction or worse experiences with treatment.<sup>35,36</sup>

The present results also indicated that respondents reporting alcohol as their previously most-used substance reported more positive experiences which therefore influenced the scores. Our external experts described this as a possible effect of age, in that people with alcohol dependence are generally older than people with other types of dependence. However, the current study identified each of alcohol dependence and age as predictors for scale scores when controlling for the other.

The data materials in national surveys are usually weighted to minimize the potential bias of nonresponses. As stated above, no information were collected concerning the background of patients or administrative information other than what the respondents replied in the questionnaire. This prevented the standard weighting procedure, where information on nonrespondents is compared to that on respondents in order to weight respondents to make them more representative of the total population. However, the high response rate of 91% strengthens the generalizability of the findings and reduces the need to implement weighting procedures based on responses.

The surveys were conducted as cross-sectional studies, and the results are therefore a description of the current status of the residential institutions. This means that some patients would have already been at the institutions for several weeks or even months, while others had arrived more recently. This way of recruiting patients and collecting data leads to a skewness in the possibility of being sampled to the surveys, since patients who have stayed for a longer time will have a higher probability of being sampled, and so those responding to the survey may have a longer stay on average. The data materials from these surveys are therefore



weighted based on the length of stay, with patients with longer stays having smaller weights.

## Implications

When measuring quality and reporting quality indicators based on patient experiences in similar populations as the one surveyed in this study, the following variables are suggested case-mix adjusters: alcohol as the most frequently used substance, length of stay, pressure from others to be admitted for treatment, gender, age, and self-perceived physical and mental health. We also recommend adjusting for mixed use. Given several measuring points, one should also consider adjusting for this. Case-mix adjustments based on these suggestions should be considered when comparing between health care providers and to enhance the clinical legitimacy. Information on nonrespondents should be considered collected as a means to weight the results for response, especially in studies where the response rate is relatively low.

One of the main objectives of case-mix adjustments is to ensure fairer comparisons between health care providers based on survey scores. This is achieved by predicting how health care providers would score or rate if the populations were more standardized.<sup>31</sup> Case-mix adjustments usually have a relatively small impact,<sup>9</sup> and the need for such adjustments has been discussed.<sup>38</sup> However, some still argue that even small adjustments show that the appropriate measures have been taken to ensure valid comparisons, and to reduce the probability of health care providers attracting “easier” populations to improve their scores, such as patients who are older or in better health.<sup>9,38</sup>

Quality indicators are often reported based on patient administrative data and/or registries. However, patient experiences and satisfaction have become an acknowledged and important way to measure aspects of the quality of health care, including in treatment for substance dependence.<sup>37</sup> The three scales tested in this study are the first quality indicators based on patient experiences reported nationally in Norway for this population, and they have been shown to effectively discriminate between health care providers, as indicated by the relatively large variations among the hospital trusts. In addition, it appears that the scale scores are somewhat correlated for each hospital (Figure 1). Furthermore, the scales can be used as quality indicators given their relevance to patients, and they have been demonstrated to be valid and reliable.<sup>17</sup> The scales provide an important tool for use when measuring and discussing the quality of interdisciplinary treatment for substance dependence in Norway, and the current study has

shown how the survey scores should be adjusted based on patient characteristics in order to ensure valid and fair comparisons between hospital trusts.

## Limitations

One of the limitations of this study was that the only available case-mix variables were patient-reported responses to the questionnaire. It is probable that other patient characteristics can impact the experiences of inpatients, and therefore should have been included in the analyses. Although this limitation is quite common when attempting to measure quality,<sup>39</sup> it is a topic that needs further exploration and which presupposes the availability of other background variables. Further developments of these measures in the given patient population should consider the availability of possibly important independent variables, such as patient administrative data and institution characteristics, eg, treatment programs or planned treatment duration.<sup>40</sup>

Another limitation was the lack of information on the health of respondents at admission. This made it impossible to know whether the self-reported health answered in the questionnaire had changed from admission and, if it had, in what direction. A possible solution is to include transition questions that either ask the respondents about their health status prior to admission, or ask them to compare their current health status with that prior to admission. Changes in health might be interpreted as an outcome in itself, but could also be used to validate the patient-reported experience scales, especially the outcome scale.

The cut-off for reporting quality indicator scores was set at the level of hospital trusts with more than one underlying unit, and with merged data from two identical surveys. The rationale for this approach was to ensure a sufficient number of respondents per unit. Since not all residential institutions in Norway are organized together under hospital trusts or are part of a larger private organization, some institutions do not receive results on the indicators and so were excluded from the present analyses. This means that even though they have a relatively large patient population, some institutions will not receive case-mix-adjusted indicator scores. A solution could be to merge even more data sets (covering more survey years), but the results from several years might be difficult to use and interpret, and was deemed unnecessary in the present study.

The present surveys were conducted as two cross-sectional surveys performed in consecutive years. The included patients might, therefore, have been at the institutions for very different durations, which is why the data were both weighted and adjusted for the length of stay.

Another approach would have been to collect data at discharge. Such an approach has been discussed previously, and deemed a more-time-consuming method for data collection, since many institutions are small and their patient flows are therefore relatively low; in addition, it could mean losing patients that drop out of treatment.<sup>17</sup> We are currently working on how to collect the experiences of patients at discharge in a fashion that is feasible for all involved. Furthermore, assessing the predictive validity of on-site patient-based indicators will be important for future research, preferably using a longitudinal design.

## Conclusion

This study found that the three scales comprising the PEQ-ITSD function well as patient-experience-based quality indicators at the hospital-trust level, given that data from 2 years were merged and implementation of the proposed case-mix model. The variations in scale scores across health care providers show that the indicators can be used to discriminate between different providers, suggesting the possibility of improving scale scores through improvement work. Future research should assess the associations between patient-based quality indicators and other quality indicators, and the predictive validity of patient-experience indicators based on on-site measurements.

## Ethics approval and informed consent

Data were collected anonymously, with no registration of the patients being surveyed. The project was run as part of the national program and was an anonymous quality assurance project. According to the Norwegian Regional Committees for Medical and Health Research Ethics, research approval is not required for quality assurance projects. The Norwegian Social Science Data Services states if the information used are anonymous, the project is not subject to notification ([http://www.nsd.uib.no/per\\_sonvern/en/notification\\_duty/meldeskjema?eng](http://www.nsd.uib.no/per_sonvern/en/notification_duty/meldeskjema?eng)). Hence, no ethics approval was needed in this project. Patients were informed that participation was voluntary and that they would remain anonymous. In accordance with all the patient surveys in the national program, health professionals at the institutions could exclude individual patients for special ethical considerations. Since no notification or ethics approval was needed, the NIPH obtained signed agreements with all the participating

institutions, describing the project and both the institutions' and NIPH's responsibility in data collecting, handling, analyzing and reporting. Previously established guidelines concerning consent through a returned questionnaire were applied.

## Data availability

The datasets generated and analyzed during the current study are not publicly available due to this being one part of an ongoing PhD-project at the NIPH and the University of Oslo. Request for permission to use the data must be directed to the NIPH.

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## Author contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; gave final approval of the version to be published; and agree to be accountable for all aspects of the work.

## Disclosure

The authors report no conflicts of interest with this work.

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