An Overview of Children’s Oral Health-Related Quality of Life Assessment: From Scale Development to Measuring Outcomes

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Key Words
Clinically meaningful difference · Dental caries · Minimally important difference · Oral health-related quality of life · Quality of life

Abstract
The objectives of this paper are to present an overview of children’s oral health-related quality of life and include specific applications for using quality of life assessment in dental research. The process of developing pediatric oral health-related quality of life measures, in particular the Child Oral Health Impact Profile, is outlined. Examples of children’s oral health-related quality of life measurement in caries research are also provided. Quality of life outcomes are presented and discussed in the context of caries research. Lastly, the relevance of measuring clinically meaningful difference in the context of measuring outcomes research is highlighted with recommendations for future research.

Why Is OHRQoL Important and How Can It Be Used in Research?
OHRQoL plays an important role in understanding subjective patient evaluations of and experience with oral healthcare. The subjective evaluation of OHRQoL reflects people’s comfort when eating, sleeping and engaging in social interaction; their self-esteem; and their satisfaction with care, and sense of self [Sischo and Broder, 2011a].
fication with respect to their oral health’ [DHHS, 2000]. Consonant with the biopsychosocial model of health, OHRQoL is the result of an interaction between oral health conditions, social and contextual factors [Locker et al., 2005], and the rest of the body [Atchison et al., 2006].

Incorporating OHRQoL creates a shift from traditional medical/dental criteria to assessment and care that focus on a person’s social and emotional experience and physical functioning. Thus OHRQoL evaluations have the potential to enhance evaluation, clinical research and care in a number of ways including needs assessment of a population or a specific clinical group. For example, a recent study among children with orofacial clefts found that individuals without health insurance and those representing ethnic minorities had lower OHRQoL and a higher rate of unmet surgical needs than their privately insured counterparts [Broder et al., 2012a]. Here it is shown that the patient’s perspective captured via OHRQoL assessments can illuminate health disparities among vulnerable youth with orofacial clefts. This information can allow researchers and clinicians to better define appropriate treatment goals and outcomes resulting in important benefits for individual patients, community-based dental practices, clinical research, and potentially public health policy.

Another way OHRQoL can be used is to inform and refine care by incorporating such assessments into oral health service planning for discrete populations like school-aged children. In this application, the Child Oral Impacts on Daily Performances was used as an indicator to prioritize children’s sociodental needs among 11- to 12-year-old children in Thailand [Gherunpong et al., 2006].

OHRQoL assessment can also be used as an outcome or evaluative measure across specialty areas, including pediatric caries research. OHRQoL provides a unique perspective on dental care from the child participant or from an observer like a parent. Traditional ‘objective’ criteria such as dental decay, missing teeth, and filled teeth (DMFT) fail to include subjective assessments (i.e., patient satisfaction, symptom reduction, and increased functional and emotional well-being) from the patient. Taking OHRQoL impacts into account, however, can differentiate need and help prioritize care for vulnerable populations [Gherunpong et al., 2006]. This information is important as most studies indicate a modest yet significant correlation between unmet needs like dental decay and children’s OHRQoL.

### Developing Child OHRQoL Measures

Specific issues arise when developing OHRQoL measures for children. Because oral health is ‘strongly age-dependent’ [John et al., 2004], it is not surprising that differences in OHRQoL have been found between children and adults [Tapsoba et al., 2000]. While many instruments exist to measure adult OHRQoL designing instruments specifically for children and adolescents allows researchers to identify and examine OHRQoL factors that are unique to these populations (i.e., self-image, social acceptance, and school environment) [McGrath et al., 2004; Rumsey and Harcourt, 2004; Broder, 2007].

Several validated instruments currently exist to measure children’s OHRQoL, including the Child Perception Questionnaire, the Early Childhood Oral Health Impact Score, Pediatric Oral Health-Related Quality of Life, Child Oral Impacts on Daily Performances, and the Child Oral Health Impact Profile (COHIP; see table 1 for an overview of these instruments).

The Child Perception Questionnaire was the first instrument specifically designed to measure children’s OHRQoL [Locker et al., 2005]. Two age forms of the Child Perception Questionnaire currently exist – one for 8- to 10-year-olds and another for 11- to 14-year-olds. It is not clear, however, whether the two measures are continuous and can be used in longitudinal studies when children age out of the younger version. The Early Childhood Oral Health Impact Score [Pahel et al., 2007], which uses caregiver reports for proxies, is targeted for preschool children who can have a variety of dental, orthodontic, and craniofacial conditions. The Pediatric Oral Health-Related Quality of Life is a relatively new OHRQoL instrument for preschool and school-aged children. While it shows promise for inclusion in OHRQoL research, particularly in its ability to capture the impact of oral conditions like caries on both general and vulnerable populations [Huntington et al., 2011], it has yet to undergo testing for its evaluative properties in a longitudinal study. The Child Oral Impacts on Daily Performances was originally developed to measure OHRQoL in 12-year-old Thai children [Gherunpong et al., 2004], although it has since been validated in a number of languages and has undergone extensive psychometric assessments. It focuses on the negative impact of oral conditions on daily performances and has been used in several population-based studies [Yusuf et al., 2006]. The COHIP is the first children’s OHRQoL instrument to incorporate both positive and negative health impacts [Sischo and Broder, 2011a], therefore has the potential to
measure more than the absence of a condition but can measure positive attributes or enhanced well-being (e.g., confidence) as a result of care. Three versions of the COHIP (child, caregiver, and teacher) are currently being used in an ongoing multicenter, longitudinal study of children’s QoL [Broder, pers. commun.]. Additionally, a short form (19 items) of the COHIP has recently been validated. Short forms are quicker to administer and therefore facilitate utilizing QoL assessment in clinical studies [Broder et al., 2012b]. All of these instruments have undergone forward and reverse translation and are available in multiple languages.

The overall goal in OHRQoL instrument development can vary depending on the research goal. For example, a discriminative instrument should not contain questions to which all respondents with the disease would respond in a like manner (e.g., items to which virtually all or none of the respondents answer similarly should be deleted). Items that are strongly influenced by factors other than the disease of interest should be excluded. Including items that correlate with one another will ensure high internal consistency, which is important for sound statistical analysis. On the other hand, an evaluative tool for clinical trials should have test items that are sensitive to change as a result of treatment.

Another consideration for measurement development and selection involves change in health in the context of treatment, a phenomenon which is referred to as response shift. Response shift encompasses changes within people regarding internal standards, values, or conceptualization of health-related QoL, particularly when health state changes. It can also affect standard psychometric indices, such as reliability and validity [Schwartz, 2010]. Acknowledging and assessing the degree to which response shifts do occur in the context of oral health and oral health treatment [Gregory et al., 2005] can inform development of measures designed to reveal when it may be occurring. This is particularly an important consideration when evaluating OHRQoL for groups known to have a lengthy and often complicated treatment such as adults with edentulism [Kimura et al., 2012], people with disabilities, and youth with extensive oral health care needs [Reissman et al., 2012].

Regardless of the application, items must be clear and relevant to the target group and calibrated to the appropriate reading level for the targeted age group using an established methodology (e.g., Flesch-Kincaid). Length and formatting of a questionnaire (e.g., line shading) can also be critical, especially for children, in order to reduce error and fatigue [Schwartz, 2010].

Achieving these goals requires that questionnaire development be an iterative process that includes a literature review, item generation, face and content validity testing, item-impact analyses, qualitative interviewing as well as consideration of theoretical issues used to nominate items for inclusion and elimination [Broder et al., 2007].

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**Table 1. Child OHRQoL measures**

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Date of creation</th>
<th>Age range</th>
<th>Items, n</th>
<th>Short form</th>
</tr>
</thead>
<tbody>
<tr>
<td>Child Perceptions Questionnaire (11–14) [Jokovic et al., 2002]</td>
<td>2002</td>
<td>11–14</td>
<td>37</td>
<td>yes (16 items)</td>
</tr>
<tr>
<td>Child Perceptions Questionnaire (8–10) [Jokovic et al., 2002]</td>
<td>2004</td>
<td>8–10</td>
<td>25</td>
<td>no</td>
</tr>
<tr>
<td>Child Oral Impacts on Daily Performances [Gherunpong et al., 2004; Yusuf et al., 2006]</td>
<td>2004</td>
<td>10–12</td>
<td>8</td>
<td>no</td>
</tr>
<tr>
<td>Child Oral Health Impact Profile (COHIP) [Broder et al., 2012b]</td>
<td>2007</td>
<td>7–18</td>
<td>34</td>
<td>yes (19 items)</td>
</tr>
<tr>
<td>Pediatric Oral Health-Related Quality of Life [Huntington et al., 2011]</td>
<td>2011</td>
<td>2–12</td>
<td>20</td>
<td>no</td>
</tr>
</tbody>
</table>
et al. [1989, 1996] and Christie et al. [1993] have written extensively on this process, and more about how this multistage process was used in the development of the COHIP can be found elsewhere [Broder and Wilson-Genderson, 2007; Broder et al., 2007, 2012b]. Validity and reliability testing must be reported to demonstrate psychometric worthiness of the instrument. Scale reliability, which was measured using Cronbach’s alpha coefficient, was excellent (0.91 for the overall score) as was the test-retest reliability of the Overall COHIP (ICC = 0.84) computed for a subset of participants who did not report a health change. Discriminant and convergent validity were also supported by the comparisons between and associations within the four groups of children. Effective QoL measures should reveal covariation with the severity of dental disease – thus demonstrating sensitivity within a disease group (e.g., dental caries) [Yusuf et al., 2006].

OHRQoL in Pediatric Caries Research

To demonstrate the value OHRQoL can bring to caries research, we recently analyzed OHRQoL COHIP data from a sample of underserved pediatric patients who were being evaluated/treated for caries as part of a larger study [Markowitz et al., 2012].

The sample included 102 schoolchildren from Newark, N.J. The participants assented to participate and the caregivers gave informed consent in accordance with IRB regulations. The clinical exam was performed in a mobile van donated by the Colgate-Palmolive Company using methods and procedures that have been vigorously calibrated and tested [Fine et al., 2007].

Demographics collected included age, gender, ethnicity and grade in school. A dental examination was performed and the presence of decayed, missing, and filled surfaces (DMFS) was entered on the patient’s chart and these data were entered into an electronic database. In these examinations, visual examination and the dental explorer were used to detect caries. In performing tactile examination, the explorer was used to remove plaque and to detect defects on the tooth surfaces as an adjunct to visual inspection. Teeth found to have occlusal pit and fissure sealants were classified as filled. Only teeth that were extracted due to caries were designated as missing. Permanent second molars were examined if their entire occlusal surface was erupted and could be examined; otherwise they were charted as unerupted. Clinical information included number of DMFS. We coded number of decayed surfaces as either less than or equal to 5 or greater than 5 (the latter representing the upper third of the distribution). The number of missing teeth was negligible, so this information was not used in the analysis.

The Child Oral Health Impact Profile-Short Form (COHIP-SF) 19 [Broder et al., 2012b] has 19 items in three subscales: Oral Health, Functional Well-Being, and Socio-Emotional Well-Being. (1) Oral Health is comprised of specific oral symptoms that are not necessarily related to one another (e.g., pain, spots on teeth). (2) Functional Well-Being included items related to the child’s ability to carry out specific everyday tasks or activities (e.g., speaking clearly, chewing). (3) Socio-Emotional Well-Being (Social-Emotional-School-Self) included items pertaining to peer interactions, mood states, school environment, and positive feelings about the self. All participants were instructed as follows: read carefully each statement and choose the answer that best describes you in the past 3 months regarding your teeth, mouth, or face. Responses were recorded as ‘never’ = 0, ‘almost never’ = 1, ‘sometimes’ = 2, ‘fairly often’ = 3, and ‘almost all of the time’ = 4. Scoring of the negatively worded items was reversed. Higher COHIP-SF 19 scores reflect more positive OHRQoL, while lower scores reflect lower OHRQoL (see table 2 for COHIP-SF item details).

Data Analysis

Descriptive statistics for all measures were computed, including means and standard deviations (SD) for continuous and frequencies for categorical data. General Linear Regression Modeling was used to compare the COHIP-SF scores for the decayed groups and separately for the filled groups on Oral Health, Functional Well-Being, Emotional Well-Being and the Overall COHIP-SF. All data manipulation and analyses were conducted with SAS 9.2, Cary, N.C., USA.

Results

Data relevant to these analyses were collected from 102 participants. Seven participants had an insufficient number of responses on the COHIP and were not included in these analyses. Average age of the participants was 12.8 (SD = 1.2); 52.8% were female. Grade in school ranged from 6 to 10 (25.3% 6th grade, 33.3% 7th, 29.9% 8th, 6.9% 9th, and 4.6% 10th grade). Self-reported ethnicity was as follows: 53.9% Latino, 34.8% Black, 1% White, and 10% other.

Clinical Data. Decayed surfaces averaged 4.4 (SD = 3.9) with a wide range (0–17) and filled surfaces were similarly distributed with an average of 3.3 (SD = 4.4), range 0–24.
The average subscale scores were Oral Health 12.0 (SD = 4.7), Functional Well-Being 13.5 (SD = 3.2), Socio-Emotional Well-Being 28.5 (6.7) and Overall COHIP-SF 57.5 (13.0).

**COHIP-SF Scores by Clinical Severity.** There were significant differences between the groups having fewer caries (as measured by the number of decayed surfaces) when compared to those having more caries on Oral Health \(F(1, 94) = 6.1, p < 0.02\), Functional Well-Being \(F(1, 94) = 11.9, p < 0.001\) and the Overall COHIP-SF \(F(1, 94) = 7.3, p < 0.009\). The scores on the Socio-Emotional Well-Being subscale were directionally consistent with those having more caries having lower OHRQoL. However, this result failed to reach conventional level of statistical significance \(F(1, 94) = 3.5, p < 0.06\). Post hoc power analysis suggests that our power to detect significant differences on the Socio-Emotional Well-Being subscale with the present effect size and sample size was fairly low (power = 0.68) suggesting a larger sample may have yielded a more significant \(p\) value less than 0.05. Comparisons of COHIP-SF scores for those having greater numbers of filled surfaces compared to lower number of filled surfaces revealed no significant differences on any subscale or the Overall COHIP-SF (data not shown). These findings are consistent with other caries data and OHRQoL research [Varni et al., 2001; Locker and Allen, 2002; Broder and Wilson-Genderson, 2007].

### Clinically Meaningful Change versus Statistical Significance

Another application of OHRQoL research involves changes in oral health as a result of specific treatment [Johnson, 2008]. In their recent study on the effect of fluoride varnish on caries among schoolchildren in rural Brazil, Arruda et al. [2011] conducted a double-blind, randomized, placebo-controlled trial with 379 children between 7 and 14 years who attended three schools in Brazil. Each school was visited four times at 6-month intervals when recruitment, dental exams, and fluoride varnish applications were completed. An interviewer-administered questionnaire was used to collect information from children regarding sociodemographic characteristics and oral health behavior (e.g., toothbrushing). Four calibrated dentists performed caries assessment exams using probes and mirrors (no radiographs) on permanent teeth only. Crude caries increments of decayed and filled surfaces (DFS) were compared between the fluoride varnish (5% sodium fluoride) and the placebo group. After 12 months of follow-up, 210 participants completed the study. Those in the varnish group had significantly lower DFS increments than did those in the control group (10.8 vs. 13.3; \(p < 0.01\)), with PF of 40% (95% CI: 34.3–45.7%; \(p < 1.01\)). Thus, applications of 5% NaF varnish are recommended as a public health measure in high-caries-risk populations. However, it is unknown if this statistically significant difference translates into clinically meaningful change for the children who participated in the study.

Including QoL assessment when evaluating clinical change begins to address our understanding of the qualitative impact the treatment has on the patient. In one study of preschool children with early childhood caries (ECC), changes in OHRQoL after treatment were reported. Using both patient and parental proxy assessments, Filstrup et al. [2003] found that children with ECC have significantly lower OHRQoL than their non-ECC counterparts. Further, they also reported that following dental treatment, those children with ECC showed significant improvement in their OHRQoL compared to their baseline measurements. Likewise, Huntington et al. [2011] found that children with ECC had significantly higher OHRQoL following surgical treatment at both their 3-month and 6-month follow-up appointments. In fact,

### Table 2. Items in COHIP-SF

<table>
<thead>
<tr>
<th>Oral Health</th>
<th>Functional Well-Being</th>
<th>Socio-Emotional Well-Being</th>
</tr>
</thead>
<tbody>
<tr>
<td>Had pain in your teeth/toothache</td>
<td>Had difficulty eating foods you would like to eat</td>
<td>Been unhappy or sad</td>
</tr>
<tr>
<td>Had discolored teeth or spots on your teeth</td>
<td>Had trouble sleeping</td>
<td>Felt worried or anxious</td>
</tr>
<tr>
<td>Had crooked teeth or spaces between your teeth</td>
<td>Had difficulty saying certain words</td>
<td>Avoided smiling or laughing with other children</td>
</tr>
<tr>
<td>Had bad breath</td>
<td>Had difficulty keeping your teeth clean</td>
<td>Felt that you look different</td>
</tr>
<tr>
<td>Had bleeding gums</td>
<td></td>
<td>Been worried about what other people think about your ...</td>
</tr>
</tbody>
</table>

1 Questions finish with ‘because of your teeth, mouth, or face’.

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by 6 months, the OHRQoL of children who had surgical treatment for the ECC was ‘indistinguishable’ from the study control group.

However, even statistical significance in QoL measures does not identify whether the change achieved has a qualitative impact on the patient [Tsakos et al., 2011]. Since a statistically significant change might not translate into real effects on patients’ lives, there has been a shift in QoL outcomes research to measure clinically meaningful change or the Minimally Important Difference (MID) [Masood et al., 2012]. MID is defined as ‘the smallest difference in score in the domain of interest which patients perceive as beneficial and which would mandate, in the absence of troublesome side effects and excessive cost, a change in the patient’s management’ [Jaeschke et al., 1989]. Instead of defining change on the basis of a statistical test of mean scores, MID uses the subjective perspective of the patient to determine what kind or how much change is meaningful to her/him [Juniper et al., 1994; Cella et al., 2002].

Two methods currently exist to measure MID: anchor methods and the standard error of measurement (SEM). Anchor methods use an independent standard (or ‘anchor’) that is interpretable and correlates at the 0.50 level (or higher) with the actual change score within the QoL instrument [Guyatt and Jaeschke, 1997; Cella et al., 2002]. They are intended to ‘measure a patient’s change score against clinically relevant or outside changes, such as expected changes caused by time, therapy, known disease diagnoses, or life events’ [Wyrwich et al., 1999a]. Anchors divide subjects into groups that reflect no change, small positive changes, large positive changes, small negative changes, or large negative changes [Revicki et al., 2008]. While many different anchors could be chosen for the analysis, the best anchors are those that identify subjects who have changed ‘to a small but meaningful degree’ [Revicki et al., 2008]. Problems arise, however, when (1) retrospective self-reports, which are subject to recall bias, are used as anchors (as is most often the case), and (2) trying to establish the amount of change that is a ‘reasonable indicator of minimal change’ [Revicki et al., 2008]. These problems make anchor-based methods of determining the MID less than ideal.

Unlike anchor-based methods that rely on the use of an independent standard, SEM is a distribution-based method that is built upon the statistical properties of a study’s results [Wyrwich et al., 1999b]. SEM is calculated based on the sample standard deviation and the sample reliability coefficient and is a function of the precision of the given instrument [Wyrwich et al., 1999b]. In other words, it is based on the estimate of error in an individual’s score, which is inversely related to the reliability of a scale (i.e., the higher the SEM, the lower the scale reliability and vice versa). An important advantage of SEM is that it is relatively stable across populations with cutoffs based on confidence intervals. It also has the most potential for establishing benchmark scores that can be used to determine clinically meaningful change [Crosby et al., 2003].

Distribution-based indices such as SEM provide no direct information on the MID; instead, they establish a standardized metric for expressing an observed change [Revicki et al., 2008]. It is therefore important to verify the MID identified in a given instrument. One way to do this is by using a Global Assessment of Change scale. Our current ongoing longitudinal observational study of children undergoing secondary cleft surgery provides an example of using a global assessment scale to rate changes in OHRQoL. Participants were asked to rate changes in OHRQoL since their last clinic visit using a scale by domain [Juniper et al., 1994] and overall scale. Absolute global ratings were coded as follows: 1–3 = minimal, 4–5 = moderate, and 6–7 = large clinically important change [Wyrwich et al., 1999a]. This global assessment, along with data from the COHIP, was used to determine the MID and clinically meaningful change. Specifically, we explored associations between OHRQoL and the Global Assessment of Change at the participants’ follow-up visit. Means were compared using General Linear Modeling; analyses were conducted using SAS 9.2.

Methods
The sample is comprised of 384 school-aged English- or Spanish-speaking children ages 7–19 with cleft who were followed for care at one of six geographically diverse cleft/craniofacial centers. These centers include: Children’s Healthcare of Atlanta, Children’s Hospital of Philadelphia, Lancaster Cleft Palate Clinic, New York University, University of Illinois at Chicago, and University of North Carolina-Chapel Hill. The sites also represent both rural and urban locations. As per IRB-approved protocols, data were collected at the children’s regularly scheduled clinic visits with children and caregivers completing the COHIP packets and demographic information, respectively. Response rates across centers averaged 90% (range: 78–95%). (See earlier description of the COHIP.)

At the follow-up visit (approximately 1 year later), in addition to the other measures in the participants’ packets, the Global Assessment of Change was administered. Participants were asked to rate changes in OHRQoL since their last clinic visit using a scale by domain [Juniper et al., 1994] and overall scale. Absolute global ratings were coded as follows: 1–3 = minimal, 4–5 = moderate, and 6–7 = large clinically important change [Wyrwich et al., 1999a]. This global assessment, along with data from the COHIP, was used to determine the MID and clinically meaningful change. Specifically, we explored associations between OHRQoL and the Global Assessment of Change at the participants’ follow-up visit. Means were compared using General Linear Modeling; analyses were conducted using SAS 9.2.
al., 1994] and overall scale (table 3). A response of zero on either scale represents no notable clinical change. Response ratings that capture important clinical change were coded as follows: minimal was comprised of ratings of 1 to 3 (−3 to −1 and 1 to 3), moderate was comprised of ratings of 4 to 5 (−4 to −5 and 4 to 5), and large was comprised of ratings of 6 to 7 (−6 to −7 and 6 to 7) [Wyrwich et al., 1999a].

Data Analysis. Demographic data were computed (means and frequencies). Mean COHIP scores for the subscales and the Overall Total COHIP were compared for patients reporting no change, minimal, moderate, and large clinical change on Oral Health, Functional Well-Being, Self-Esteem and Total Change using General Linear Modeling with SAS 9.2.1.

Results

Youth participants averaged 12.6 years (SD = 2.9); 42% were female. The sample's ethnic composition included: 68% White, 13.3% Latino or mixed, 8.2% Black or African American and 10.5% Asian. Fifty-four percent had private insurance.

COHIP subscale (mean, SD) scores were: Oral Health (21.3, 5.5), Functional Well-Being (14.0, 3.5), Emotional Well-Being (22.7, 4.0), School (9.9, 2.6), Self-Esteem (21.6, 3.7) and Overall COHIP (89.5, 13.4). Note that although the SEM approach for assessing clinically meaningful change does permit participants to endorse a decrement in OHRQoL, nearly none (3 total in the present sample) of these participants did. Thus, in all cases included in this analysis, ratings of change indicate either no change or improvement in OHRQoL. Distribution (in percent) of participants in the Global Assessment of Change groups representing no, minimal, moderate, and large amounts of change were: Oral Health (32.4, 32.1, 18.4, 17.0), Functional Well-Being (47.5, 30.9, 11.6, 10.0), School (50.1, 27.9, 10.3, 11.6) and Overall change (54.0, 23.1, 11.2, 11.7) (table 4). There were significant differences on many COHIP subscales and overall for Oral Health, Functional Well-Being, Emotional Well-Being, School, Self-Esteem and Overall COHIP. Total global improvement was significantly associated with higher COHIP scores on Self-Esteem and Overall COHIP.

Discussion

COHIP scores reflect differences in OHRQoL among treatment-seeking patients based on the patients’ reports of clinically meaningful change. Although these results are preliminary, they represent an important methodology rarely utilized in oral health research. Our future work will seek to explore the role of other patient and clinical characteristics in the context of these associations.

To date, one published oral health report has utilized this MID methodology in a caries clinical study [Malden et al., 2007]. Caregivers in New Zealand complet-
ed the Parental-Caregiver Perceptions Questionnaire, an OHRQoL assessment, before their children had general anesthesia for dental caries and then completed a follow-up questionnaire again after 1–4 weeks (a global change rating scale was also included in the follow-up questionnaire). Comparisons between the baseline and follow-up data for the mean scores were examined and the MID was calculated for the overall scale and the subscales. The results revealed significant reductions in mean Parental-Caregiver Perceptions Questionnaire (improved OHRQoL) after treatment, with effect sizes ranging from moderate to large. Additionally, about two thirds of the children showed or exceeded the MID in their OHRQoL scores following their treatment.

Although studies in caries research that incorporate QoL outcomes are expanding [Sischo and Broder, 2011b], most studies report statistical difference using only change scores before and after treatment. Despite the widespread use of global change ratings in health-related QoL outcomes research across pediatric populations, its use in pediatric caries research is extremely limited.

Conclusions

Several published OHRQoL measures have been shown to have excellent psychometric properties and are suggested tools for oral health research. Currently, the gold standard in QoL research is to use school-aged children’s subjective reports as youth can reliably complete standard self-administered, condition-specific questionnaires [Feeny et al., 1998; Broder and Wilson-Genderson, 2007]. Reports have consistently shown modest yet significant correlations between clinical indices like dental decay with established OHRQoL instruments [Gherunpong et al., 2004; Locker et al., 2005; Broder et al., 2007; Huntington et al., 2011]. Although such measures are increasingly utilized in epidemiological as well as oral health clinical studies with pediatric populations, many clinical trials are not including these subjective evaluations in their research designs to measure outcomes [Sischo and Broder, 2011b]. Across patient groups in medicine, it is well documented that such QoL data often complement objective clinical data and may be useful in treatment decisions and measuring efficaciousness of care [Gherunpong et al., 2006]. Thus, during this era of evidence-based care coupled with reduced access to care among disadvantaged populations, utilizing OHRQoL assessments in epidemiological, observational, and clinical studies is recommended as an adjunct to dental caries research.

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Disclosure Statement

The authors declare that they have no competing interests.

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