Evaluation of the Clinical Performance of NuSmile Pedodontics Zirconia Crowns in Pulp-Treated Primary Teeth—2 Years Follow-Up Study.

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Abstract

Objectives There is a lack of information regarding the longevity of prefabricated crowns on primary teeth. Therefore, the objective of the present study was to evaluate the clinical success rate of NuSmile pediatric Zirconia crowns in both primary anterior and posterior teeth up to 2 years in terms of gingival health, levels of plaque, restoration failure, and marginal integrity.

Materials and Methods This interventional study evaluated the clinical performance of NuSmile primary Zirconia crowns on 232 (172 primary incisors and 60 primary molars) pulp-treated primary teeth. Statistical analysis included independent t-test and Kaplan–Meier survival method for which the level of significance was set up at p < 0.05.

Results Gingival and plaque index of the teeth restored with NuSmile Zirconia crowns (test tooth) compared with adjacent and antagonist teeth (control tooth) showed no statistically significant difference at all time intervals (p > 0.05). The Kaplan–Meier survival graph indicated that only 34% of NuSmile Zirconia incisor crowns and 86% of NuSmile Zirconia molar crowns have survived at the end of 24 months. Of the 172 NuSmile Zirconia incisor crowns placed, only 82 teeth survived at the end of 2 years and the majority of the crowns completely lost the Zirconia crown. The Kaplan–Meier graph also indicated that more than two-thirds of crowns, i.e., 90% incisor crowns and 76% of molar crowns, had good marginal integrity at the end of 24 months.

Conclusions Zirconia pediatric crowns preserve and maintain gingival health and have long-term survival rates with good retention and marginal integrity, indirectly preventing secondary caries. Hence, Zirconia pediatric crowns are a well-suited restoration of choice for primary teeth rehabilitation.

Keywords
► Zirconia pediatric crowns
► NuSmile
► clinical success
► primary teeth
► incisors
► molars

Introduction

In today’s modern society, both parents and children are cosmetically conscious and increasingly concerned about the esthetics of their teeth. The child’s parents often influence the dental professionals treating the children in selecting dental restoration, and parents are getting more involved in clinical decision-making than ever before.¹ A recent study has also shown that children at the age of 6 years and above can

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appreciate the esthetics of their anterior teeth. Thus, there is a great preference for restorations that bring the primary tooth back to a healthy state in both appearance and function.

Management of extensive carious lesions and traumatized primary teeth has gradually shifted from extraction to full-coverage restorations. Extra-coronal full-coverage crowns are indicated in primary teeth with developmental defects, multi-surface caries, patients with high caries risk, fractured teeth, teeth where direct restoration tends to fail, after pulpal therapy, as abutments for space maintainers, and teeth with excessive wear.

Traditionally, composite strip crowns and stainless steel crowns (SSC) have been used as full-coverage restorations for grossly decayed and pulp-treated primary anterior teeth and molars. Strip crowns are esthetically better but are technique sensitive. Moisture and hemorrhage control is essential with strip crowns as it could lead to resin placement failure. SSC crowns have been successful for many years in terms of durability, retention, and function. The most significant problem was their poor esthetics limiting their use to the posterior segment only. However, some parents refuse SSC due to their black color, which is unappealing.

Pre-veneered stainless steel crowns (PSSC) combine conventional SSC’s durability with the esthetic appeal of a composite resin veneer. However, the significant concerns with PSSC were removing additional coronal tooth structure, inability to crimp the margins of the crowns before cementation, and loss of the esthetic acrylic facing.

Prefabricated pediatric Zirconia crowns offer an excellent alternative to full-coverage crowns when restoring deciduous teeth with a sizeable carious defect. These crowns have high flexural strength, allowing them to resist crack propagation. The additional benefits of these crowns are the ability to replace metals due to extremely high strength and toughness, higher resistance to chemicals, and superior erosion resistance. Zirconia crowns are biocompatible, autoclavable, and equal to or more durable than natural enamel.

Since the introduction of primary Zirconia crowns, several studies have reported their clinical success. Severely mutilated primary anterior teeth showed that NuSmile Zirconia crowns (NuSmile Ltd., Houston, TX, United States) offered superior esthetics and durability with remarkable gingival responses up to 24 months. A clinical study on the wear behavior of primary enamel against Zirconia crowns demonstrated the lowest wear rate of primary enamel. A retrospective study that evaluated the clinical success and parental satisfaction showed a considerable percentage (89%) of parents were highly satisfied by the crowns’ size, color, and form. Additionally, a study on three randomized controlled anterior aesthetic full-coverage crowns showed Zirconia crowns to be highly retentive compared with resin composite strip crowns and PSSC. To date, most clinical studies of prefabricated crowns have been conducted on anterior teeth. Therefore, the objective of the present study was to evaluate the clinical success rate of NuSmile pediatric Zirconia crowns in both primary anterior and posterior teeth up to 2 years in terms of gingival health, plaque levels, restoration failure, and marginal integrity.

Materials and Methods

Study Protocol

This interventional study was done between 2014 and 2017 in accordance with the Declaration of Helsinki (as revised in Edinburgh 2000). It evaluated the clinical performance of NuSmile primary Zirconia crowns on 75 children (40 males and 35 females) aged 2 to 8 years. The study included healthy and cooperative children managed by non-pharmacological and pharmacological behavioral management techniques. Children with no history of systemic illness and parents who signed the consent on behalf of their children and were willing to follow-up during the entire course of the study formed the study cohort. Inclusion criteria for the study were primary teeth with sufficient tooth structure and expected to survive for at least 2 years. Grossly carious non-restorable clinical crown, primary tooth with root resorption, tooth exfoliated within 1 year of crown placement, special need children, and parents not willing to come for follow-up were excluded from the study.

Pulpal status was assessed clinically and confirmed by pre-operative periapical radiographs. Around 232 (172 primary incisors and 60 primary molars) pulp-treated primary teeth received a prefabricated Zirconia crown. The study cohort consisted of 52 pulpotomy and 180 pulpectomy cases. Two clinicians (T.W. and D.D.) with experience of more than 20 years of clinical practice performed pulp treatment and crown placement in the same appointment. Teeth with non-vital pulp that required two visit pulpectomy were done, and the crowns were cemented on the day of root canal obturation. Endodontic procedures for pulpally involved teeth were performed as per the best policy clinical practice guidelines from the American Academy of Pediatric Dentistry Reference Manual 2020.

The same operator who placed the crown recorded clinical parameters such as gingival index, plaque index, restoration

<table>
<thead>
<tr>
<th>Primary teeth</th>
<th>Maxillary teeth</th>
<th>Mandibular teeth</th>
<th>Pulpotomy</th>
<th>Pulpectomy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central incisors</td>
<td>104</td>
<td>–</td>
<td>–</td>
<td>104</td>
</tr>
<tr>
<td>Lateral incisors</td>
<td>68</td>
<td>–</td>
<td>–</td>
<td>68</td>
</tr>
<tr>
<td>First primary molars</td>
<td>8</td>
<td>17</td>
<td>22</td>
<td>3</td>
</tr>
<tr>
<td>Second primary molars</td>
<td>5</td>
<td>30</td>
<td>30</td>
<td>5</td>
</tr>
</tbody>
</table>

Table 1 Distribution of NuSmile Zirconia crowns as per the type of teeth
failure, and marginal integrity at 6, 12, 18, and 24 months. Both operators were standardized to evaluate index scores. The gingival index was recorded using a blunt periodontal probe, and a plaque disclosing swab was used to assess the plaque index. The gingival health and plaque scores were evaluated and compared with adjacent and opposing teeth. The evaluation of restoration failure and marginal integrity was done clinically with visual assessment of restoration, according to the United States Public Health Service criteria (Table 2).

Clinical Procedure for NuSmile Zirconia Crown Preparation
Both operators prepared pulp-treated teeth for NuSmile Zirconia crowns as per the manufacturer’s guidelines on crown placement protocol. There was no formal training or standardization between the two operators for crown placements except following technical guide instructions mentioned and recommended for use and general information on NuSmile Zirconia crowns. Before the tooth preparation, the operator chose a well-fitting prefabricated crown based on the pulp-treated tooth’s original size. Local anesthesia was given using lidocaine 2% with epinephrine (1:80,000), especially when placing crowns without general anesthesia. Tooth structure of 1–1.5 mm was removed from the natural occlusal contours followed by 2 mm sub-gingival reduction with a special crown cutting kit provided by NuSmile. This opened up the interproximal contact areas and reduced the entire clinical crown by 20% (or 0.5–1.25 mm). All line angles of the prepared tooth were rounded. The crown’s fit was checked, and crowns were cemented with Ketac Cem luting cement. Excess cement was removed, and crowns were held in place until the cement had been set.

Results
Data were analyzed for descriptive data by SPSS version 20.0. Inferential statistics to compare gingival health scores/plaque level scores over the period between test tooth (NuSmile Zirconia crowns) and control tooth (adjacent/antagonist tooth) was done using independent samples t-test. The Kaplan–Meier survival method was used to analyze restoration failure and marginal integrity of Zirconia crown for 2 years. The 95% confidence interval (CI) was included for stating the relevance of the finding, and the level of significance was set up at \( p < 0.05 \). The exfoliated teeth restored with NuSmile Zirconia crowns were evaluated for restoration failure and marginal integrity.

Gingival Health and Plaque Scores
Tables 3 and 4 show the mean gingival health and plaque scores of normal and prefabricated Zirconia crowns at intervals up to 24 months. The oral health domain of clinical performance determined by the gingival and plaque index of teeth restored with NuSmile Zirconia crowns (test tooth) compared with adjacent and antagonist teeth (control tooth) had no statistically significant difference at all time intervals \( (p > 0.05) \). Fig. 1 shows plaque deposits on maxillary second molar compared with prefabricated Zirconia mandibular second crown, while Fig. 2 shows plaque deposits on right deciduous incisor compared with prefabricated Zirconia crown on the left incisor.

Restoration Failure
The prefabricated Zirconia crowns were assessed for restoration failure every 6 months for a 2-year follow-up period (Table 5). The incisors prefabricated Zirconia crowns exhibited an increased incidence of restoration failure than the

<table>
<thead>
<tr>
<th>Criteria</th>
<th>Grades</th>
<th>Description</th>
<th>Author</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gingival health</td>
<td>0</td>
<td>No obvious signs of inflammation</td>
<td>Löe and Silness</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>Mild marginal gingivitis tissue slightly reddened and edematous</td>
<td>gingival index</td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Moderate marginal gingivitis tissue obviously reddened and edematous</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>Severe gingivitis tissue is very swollen: spontaneous bleeding</td>
<td></td>
</tr>
<tr>
<td>Plaque index</td>
<td>0</td>
<td>No plaque</td>
<td>The United States Public Health Service</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>Film at the gingival margin</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Moderate accumulation</td>
<td></td>
</tr>
<tr>
<td></td>
<td>3</td>
<td>Abundance of plaque</td>
<td></td>
</tr>
<tr>
<td>Restoration failure</td>
<td>0</td>
<td>Crown appears normal: no cracks, chips, or fractures</td>
<td>The United States Public Health Service</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>Small but noticeable area of loss of material</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Complete loss of the crown</td>
<td></td>
</tr>
<tr>
<td></td>
<td>E</td>
<td>Exfoliated</td>
<td></td>
</tr>
<tr>
<td>Marginal integrity</td>
<td>0</td>
<td>No detectable margin</td>
<td>The United States Public Health Service</td>
</tr>
<tr>
<td></td>
<td>1</td>
<td>Detectable margin</td>
<td></td>
</tr>
<tr>
<td></td>
<td>2</td>
<td>Infra-occlusion</td>
<td></td>
</tr>
<tr>
<td></td>
<td>E</td>
<td>Exfoliated</td>
<td></td>
</tr>
</tbody>
</table>
molar Zirconia crowns. The Kaplan–Meier 2-year survival estimate of the NuSmile Zirconia crowns incisor group was 62.1% (95% CI: 58.5–65.5), and the molar crowns group was 88% (95% CI: 78–85) survival probability (►Fig. 3). Results show that incisor and molar significantly influence the duration of survival time of Zirconia crown, individually.

Out of 172 anterior Zirconia crowns that were followed up at 6 months, 19 crowns were lost completely (►Fig. 4). At 24 months review, only 148 incisor crowns were available, out of which 32 teeth had a complete loss of Zirconia crown. The corresponding figures for molar Zirconia crowns were only four crowns out of 54 lost entirely after 2 years.

**Marginal Integrity**

The 2-year marginal integrity calculated with the Kaplan–Meier method revealed the NuSmile Zirconia crowns incisor group to be 94.5% (95% CI: 92.5–96.5). Kaplan–Meier 2-year marginal integrity estimate of the NuSmile Zirconia crowns incisor group was 92% (95% CI: 96.5–98) and the molar crowns group was 84.5% (95% CI: 86.5–96) survival probability. The data show that incisor and molar significantly influence the time taken to lose marginal integrity, individually (►Fig. 5). Prefabricated Zirconia crowns were checked for their marginal adaptation every 6 months for 2 years (►Table 6). The NuSmile Zirconia incisor group had 16

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Table 3 Comparison of gingival health scores of normal and prefabricated Zirconia crowns at 6-, 12-, 18-, and 24 months intervals

<table>
<thead>
<tr>
<th>Duration</th>
<th>Incisor</th>
<th>Molar</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Normal crown</td>
<td>Zirconia crown</td>
</tr>
<tr>
<td>6 months</td>
<td>Mean 1.45, SD 0.55</td>
<td>Mean 1.35, SD 0.45</td>
</tr>
<tr>
<td>12 months</td>
<td>Mean 1.50, SD 0.55</td>
<td>Mean 1.35, SD 0.50</td>
</tr>
<tr>
<td>18 months</td>
<td>Mean 1.50, SD 0.45</td>
<td>Mean 1.40, SD 0.50</td>
</tr>
<tr>
<td>24 months</td>
<td>Mean 1.55, SD 0.55</td>
<td>Mean 1.35, SD 0.60</td>
</tr>
</tbody>
</table>

Abbreviations: ns, not significant; SD, standard deviation.

Table 4 Comparison of plaque scores of normal and prefabricated Zirconia crowns at 6-, 12-, 18-, and 24 months intervals

<table>
<thead>
<tr>
<th>Duration</th>
<th>Incisor</th>
<th>Molar</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Normal crown</td>
<td>Zirconia crown</td>
</tr>
<tr>
<td>6 months</td>
<td>Mean 1.05, SD 0.60</td>
<td>Mean 0.89, SD 0.45</td>
</tr>
<tr>
<td>12 months</td>
<td>Mean 1.20, SD 0.60</td>
<td>Mean 0.95, SD 0.50</td>
</tr>
<tr>
<td>18 months</td>
<td>Mean 1.20, SD 0.60</td>
<td>Mean 1.00, SD 0.40</td>
</tr>
<tr>
<td>24 months</td>
<td>Mean 1.20, SD 0.60</td>
<td>Mean 0.95, SD 0.45</td>
</tr>
</tbody>
</table>

Abbreviations: ns, not significant; SD, standard deviation.

Fig. 1 Deposits of plaque on maxillary second molar compared with prefabricated Zirconia mandibular second crown.

Fig. 2 Deposits of plaque on right deciduous incisor compared with prefabricated Zirconia crown on the left incisor.
crowns that were infra-occluded and re-erupted within 2 years, and the NuSmile Zirconia molar group had 14 crowns with detectable loss of marginal integrity (∼Figs. 6 and 7). The results showed that the NuSmile Zirconia incisor crown had greater marginal integrity in the molar group. ∼Figs. 8,9,10 show various aspects of prefabricated Zirconia crown placement.

Discussion

Pediatric dentistry has undergone a paradigm shift from clinical decision-making to a patient-sensitive treatment
Childhood caries is a public health problem requiring multisectoral coordination for effective management.\(^4,13,14\) The preformed crown is a promising method of providing substantial shelter for endodontically treated primary teeth.\(^4,15\) Although an array of prefabricated economic crowns is available in the market, esthetics and the most significant pitfall can be efficiently satisfied by pediatric Zirconia crowns.\(^16\) In this study, NuSmile prefabricated Zirconia crowns were placed more on the anterior teeth than molars. This indicated high esthetic demand currently emerging as an expected standard of care among parents and children, qualifying Zirconia crown as an excellent alternative for other pediatric restorative options. Only a handful of studies have assessed the clinical performance of Zirconia crowns.\(^1,3,10,17–20\) Moreover, reports of longitudinal evaluations are sparse.

Any metric to be justifiably validated in dentistry needs to be comprehensively drafted to evaluate its survivability, marginal integrity, and effect on surrounding oral structures. Hence, the purpose of this work was to efficiently investigate the longitudinal detailing of NuSmile Zirconia crowns’ clinical performance for 2 years. Zirconia pediatric crown requires extensive subgingival preparation; hence, assessing its biocompatibility with gingiva is of paramount importance. In this work, better gingival scores consistently over 2 years were evidenced. This finding was similar to other studies.

**Table 6 Marginal integrity of Zirconia crowns at 6-, 12-, 18-, and 24 months intervals**

<table>
<thead>
<tr>
<th>Follow up Period</th>
<th>6th months</th>
<th>12th months</th>
<th>18th months</th>
<th>24th months</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type of Crowns</td>
<td>Incisors</td>
<td>Molars</td>
<td>Incisors</td>
<td>Molars</td>
</tr>
<tr>
<td></td>
<td>(n=172)</td>
<td>(n=60)</td>
<td>(n=162)</td>
<td>(n=60)</td>
</tr>
<tr>
<td>Score 0</td>
<td>172</td>
<td>60</td>
<td>155</td>
<td>55</td>
</tr>
<tr>
<td>Score 1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Score 2</td>
<td>0</td>
<td>0</td>
<td>7</td>
<td>0</td>
</tr>
<tr>
<td>Score E</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

Abbreviations: ns, not significant; SD, standard deviation.

Note: Score 0, no detectable margin; Score 1, detectable margin; Score 2, infra-occlusion; Score E, exfoliated.

**Fig. 7** Loss of marginal integrity on lower deciduous first molar for prefabricated Zirconia crown.

**Fig. 8** (A) Pre-treatment 51, 52, 61, and 62. (B) Prefabricated Zirconia crown on 52, 51, 61, and 62. (C) Prefabricated Zirconia crown on 51 infra-occluded at 12 months.
reported earlier.\textsuperscript{21–23} The average plaque index was higher for the control group than the test group comprising NuSmile prefabricated Zirconia crowns. Decreased plaque score reported in this work was consistent with some previous studies.\textsuperscript{11,21–23}

The biological outcome measured in terms of mean gingival health scores corresponded to the results of plaque level scores for 2 years. This was in line with previous findings that polished and smooth surfaces result in less plaque accumulation and minor gingival irritation.\textsuperscript{24,25} Furthermore, the manufacturing method of NuSmile prefabricated Zirconia crown utilizes a proprietary injection molding technique and hand-polishing method, which lowers its lower surface roughness (Ra 2.8) and higher mean gloss (Gs 42.7).\textsuperscript{26}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{image1.png}
\caption{(A) Pre-treatment 51, 52, 61, and 62. (B) Prefabricated Zirconia crown on 52, 51, 61, and 62. (C) Infra-occluded tooth 52 at 12 months.}
\end{figure}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{image2.png}
\caption{(A) Pre-operative 61 trauma, (B) Post-operative prefabricated Zirconia crown on 61, (C) Infra-occluded at 12 months, and (D, E) Tooth re-eruption.}
\end{figure}
The above parameters might have resulted in lesser plaque accumulation and decreased gingival inflammation.

The Zirconia crowns were cemented with conventional glass ionomer cement due to the limited control of hemostasis as the preparation for Zirconia crowns is 1–2 mm subgingival. However, as per manufacturer recommendations, the luting cement of choice for prefabricated zirconium crowns is bioactive resin-based glass ionomer cement which requires good gingival bleeding control prior to cementation. Numerous published studies have highlighted the success of resin-reinforced glass ionomers, but a recent study conducted concluded that packable conventional glass ionomer cement is more retentive than bioactive cement for the cementation of primary Zirconia crowns.

The survival rate measured in terms of restoration failure for NuSmile prefabricated Zirconia restoration in this work reported no chipping or fracturing. Only loss of crowns was evaluated at each follow-up visit. The endodontic failures were not evaluated and, hence, not included. The absence of porcelain-veneered restoration and monolithic Zirconia preparation, which was said to exhibit higher resistance to fracture loads than layered Zirconia crowns and metal-ceramic crowns. The restoration failure in terms of complete loss of crown was higher for incisor crowns than for the molar prefabricated Zirconia group. This may be due to increased surface area and anatomy of posterior teeth compared with its anterior counterpart that offer better retention. More overstudies reported that it was comparatively convenient to fit Zirconia crown in molars. In all cases, restoration failure happened at the crown-to-teeth interface and not on the cement-to-crown interface as no luting cement remained on the tooth structure after the debonding of Zirconia crowns.

Marginal integrity is an integral component of clinical performance. It has many subsets of factors—the first being marginal fit. The excellent finish line of the study group could have contributed to a high marginal fit in this study. The second factor is the vertical gap between the restoration and the prepared tooth abutment. Creating a favorable area for bacterial accumulate in the marginal gap jeopardizes the longevity of the restoration, leading to the development of secondary caries. No evidence of secondary caries in this study substantiates good marginal fit and a minimal vertical gap in this pediatric crown.

In the present study, traumatized teeth that were infraoccluded and restored with the Zirconia crowns re-erupted along with the natural tooth, spontaneously suggesting a favorable prognosis. However, periodic clinical and radiographic follow-up should be performed to prevent pulp infection of the intruded teeth and possible disturbances to the budding permanent tooth.

The strength of this research work is a sufficient sample size and comprehensive clinical performance evaluation longitudinally. Limitations of the present study were that both patients under general and local anesthesia formed the study cohort that could have resulted in mixed clinical outcomes challenging the external validity. Since it is a retrospective study, potential confounding factors such as patient selection, operator bias, and lack of standardization between operators can have an effect on the results. Moreover, further long-term studies among different commercially available Zirconia crowns can help us understand the differences in survival and biological outcomes in each group and foster amendments as needed. To summarize, pediatric Zirconia crowns have demonstrated better gingival health, decreased plaque accumulation, and reasonable survival rates longitudinally, especially for molars. Further research on various brands of Zirconia crowns and methods for longer retention of crowns in anterior teeth should be planned.

**Conclusion**

Based on the findings of this study, we conclude that Zirconia pediatric crowns preserve and maintain gingival health. They also prevent microbial plaque accumulation similar to the natural tooth. These pediatric crowns have long-term survival rates with good retention and marginal integrity, indirectly preventing secondary caries. Furthermore, these crowns do not intervene with the path of eruption of infra-occluded traumatized teeth. Hence, Zirconia pediatric crowns are a well-suited restoration of choice for primary teeth rehabilitation.

**Conflict of Interest**

None declared.

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