Endodontic treatment of primary teeth in the general dental practice

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The most common endodontic treatment of primary teeth is pulpotomy, which is used in the case of carious, iatrogenic or traumatic damage of the pulp. As prerequisite for such a procedure, the patient has to be free of clinical symptoms, according to the DGZMK.1 As per UK Guideline 2 and Weisshaar,3 this condition is still met in cases of temporary discomfort or short, spontaneous pain.

Various materials are used for pulpotomy and pulpectomy. The standard preparation for pulpotomy is still Formocresol—19% formaldehyde, 35% cresol, 15% glycerine, 31% distilled water—used in a 1:5 dilution. Weisshaar3 reported that 92.4% of paediatric dentists in Canada and 76.8% of paediatric dentists worldwide used Formocresol for pulpotomy in 1989. While there have always been objections to the use of formaldehyde, Tagger and Tagger4 state that Formocresol pulpotomy assists the retention of primary teeth and outperforms any other current method.

In their 2002 statement, the DGZMK advised using calcium hydroxide—Ca(OH)₂—for pulpotomy.1 The 2006 UK Guideline however considers this treatment inappropriate.2 Furthermore, the UK Guideline emphasised that pulpotomy with 15.5% ferric sulphate (Astringedent, Ultradent) is as effective as a five-minute treatment with Formocresol. However, ferric sulphate only acts haemostatically when applied to the bleeding pulp stump for 15 seconds. Subsequently, ferric sulphate is removed from the pulp cavity and zinc oxide-eugenol is directly applied to the root canals. The UK Guideline cites a study that achieved 55% internal resorption—as with Ca(OH)₂—and 71% obliteration of the root canal radiologically with this pulpotomy method.2 Peng et al.5 demonstrated equivalent results in a meta-analysis of 11 studies that compared Formocresol and ferric sulphate pulpotomies. The ferric sulphate pulpotomies demonstrated a clinical success rate of 78 to 100% and a radiological success of 42 to 97%. For inclusion

<table>
<thead>
<tr>
<th>Table I</th>
<th>N₂-therapy of primary teeth 1992–1998.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total</td>
<td>n</td>
</tr>
<tr>
<td>1 VitA</td>
<td>559</td>
</tr>
<tr>
<td>2 Dev/MoA</td>
<td>216</td>
</tr>
<tr>
<td>3 MoA</td>
<td>343</td>
</tr>
<tr>
<td>4 r.c.f./Fist</td>
<td>66</td>
</tr>
<tr>
<td>5 r.c.f. kons</td>
<td>493</td>
</tr>
<tr>
<td>no-shows after treatment</td>
<td>71</td>
</tr>
<tr>
<td>without no-shows</td>
<td>423</td>
</tr>
<tr>
<td>0–23 months</td>
<td>17</td>
</tr>
<tr>
<td>≥ 24 months</td>
<td>54</td>
</tr>
<tr>
<td>Failure</td>
<td>28</td>
</tr>
<tr>
<td>0–23 months</td>
<td>22</td>
</tr>
<tr>
<td>≥ 24 months</td>
<td>6</td>
</tr>
</tbody>
</table>

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in the analysis, the teeth were required to have remained in situ for at least 12 months in order to avoid the inclusion of early failures in the baseline studies.

According to Einwag, the 90 to 100% clinical success rate of pulpotomies using N2 Endodontic cement (Ghimas) is the same as that achieved using the five-minute Formocresol technique, which is not surprising owing to their similar composition. Prior to the EU certification of 14 June 1998, N2 contained 7% formaldehyde, which was subsequently reduced to 5%.

Bürkle reported a low technical sensitivity and good clinical results for all aldehyde-containing materials, such as Formocresol, glutaraldehyde and N2.

Materials and method

A prerequisite for treatment was the restorability of the respective teeth. An exclusion criterion was increased mobility without advanced root resorption. In accordance with these, the following endodontic treatments between 1992 and 1998 were recorded. X-rays were only routinely available for mortal amputations and root-canal treatments of gangrenous primary teeth. Where possible, follow-up X-rays of the latter were taken.

Vital amputations (pulpotomies), mortal amputations and root-canal fillings were performed using the root-canal filling material N2 under relatively dry conditions. For the root-canal filling, N2 powder and N2 liquid were mixed to a creamy consistency and applied near the apex following preparation using a manual reamer or mechanical HERO 642 endodontic files (MICRO-MEGA).

For vital or mortal amputations, N2 was mixed to a relatively solid consistency. First, deep carious defects were removed with an excavator. The remaining caries was excavated using a thick spherical bur without water-cooling, followed by cavity preparation with elimination of the coronal pulp using a turbine.

Frequently, the remaining pulp bleeds from the canals. Light bleeding can be disregarded, but N2 must be firmly applied to the cavity for several minutes in the case of heavier bleeding. Owing to the formaldehyde level, the bleeding can be stopped relatively quickly. The blood-soaked N2 is then removed from the cavity and replaced with freshly mixed N2 of a solid consistency.
The mortal amputations were attributed partly to Toxavit-devitalisation and partly to treatment of gangrenous teeth. Devitalised teeth were treated in the same manner as vital teeth. Gangrenous teeth were treated with different methods. Simple mortal amputations were performed in a manner similar to a vital amputation. Root-canal fillings (partial or complete) were simply filled with paste. In some cases, the roots were finished through artificial fistulation, which was done interradicularly—mostly with a Cavit lifter through the mucosa—and in a few cases after the mucosa had been shifted with a turbine bur.

All endodontic measures, including the fillings, were performed in one appointment. Additional visits were only necessary when massive bleeding occurred, when a composite filling was required and in devitalisation cases. In the first two cases, the cavity was filled with excess N2 or with zinc oxide-eugenol following N2 application. N2 application and immediate amalgam sealing of the cavity without lining materials is still the most time-saving method. In exceptional cases, stainless-steel crowns served as definitive restoration.

_Results_

A total of 559 primary teeth were treated endodontically between 1992 and 1998. I treated about 39% of these myself. Pulpotomies were conducted on 460 primary molars, 29 to Toxavit devitalisation with subsequent mortal amputation was performed on 29 primary molars. Of the 70 gangrenous primary molars treated, 37 were treated by simple mortal amputation. In 22 cases, root filling or mortal amputation was followed by artificial fistulation. Root-canal treatment of 11 gangrenous primary molars was performed without fistulation. The following gives the average age of the patients who underwent a pulpotomy:

- maxillary first molar: 7 years, 6 months
- maxillary second molar: 7 years, 5 months
- mandibular first molar: 6 years, 6 months
- mandibular second molar: 6 years, 6 months

After treatment, 11.8% (n = 66) of the young patients did not return to the practice, which left only 88.2% of the original patient group (n = 493) for further treatment.
observation in the practice. During the observation period, clinical failure (pain, swelling and fistulas) was registered in 5.7% \((n = 28)\) of the patients. The symptoms of clinical failure occurred within 23 months after treatment in 78.6% of patients \((n = 22)\). In one patient, these symptoms occurred only several hours after devitalisation followed by mortal amputation. Of the extractions done, 76.1 per cent \((n = 54)\) occurred 24 months or more post-treatment. Problems with tooth eruption of or enamel damage to the permanent teeth were not observed.

In summary, we observed that the number of extractions and failures in the observation period was twice as high in necrotic teeth when comparing vital amputation with treatment of gangrenous teeth (column 1 compared to column 3 to 5 in Table I).

**Discussion**

The number of young patients that did not return to the dental practice after treatment may be attributed to the discomfort of the treatment. However, the DMS III (Third German Oral Health Study) found a social attachment to the dentist ("always the same dentist") as high as 92.5% in 12-year-old patients in West Germany in 1997.\(^8\) It is possible that a significant number of those who did not return were clinical failures, prompting the patients’ parents to consult another dentist, but this is rather unlikely.

Huber\(^6\) analysed 179 N2 vital amputations in 105 patients and documented a failure rate of 9.5%. The age of tooth loss corresponded to the average age of premolar eruption. The N2 pulpotomies had a low failure rate of 4.4% in this study. This was confirmed by Einwag, who gives the clinical success rate of N2 pulpotomies as 90 to 100%.\(^6\)

Although not documented in detail in this study, pain anamnesis appears to be insignificant in determining the success of a N2 pulpotomy. Acute exacerbations, as often observed following pulpotomy with Ca\((OH)\)_\(_2\), almost never occurred after N2 pulpotomy. Against this background, it is incomprehensible that German universities continue to favour Ca\((OH)\)_\(_2\) pulpotomies. The UK Guideline is a step ahead in this respect.\(^2\)

It is important for the dental practice that the outcome of the treatment be guaranteed, that both the young patient and the dentist be comfortable with one another, that the treated tooth remains *in situ* until development of the permanent teeth and that the erupted permanent teeth are not damaged.

Dental practitioner Garry\(^10\) aptly expresses the use of N2 in endodontic treatment of primary teeth, which reflects my personal opinion: "When endodontic treatment is recommended for primary teeth, the concept 'careful selection of cases' is often used to warn the practitioner against treating gangrenous teeth. The technique herein does not require 'careful selection' for treatment [...]. The goal of endodontic intervention on primary teeth is not to obtain radiographic images which meet standards used in treating adult teeth, but to retain deciduous teeth as long as possible as entities and space maintainers.”

**Editorial note:** A list of references is available from the publisher.