Biocompatibility of orthodontic adhesives in rat subcutaneous tissue

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Received: February 2, 2009 - Modification: September 5, 2009 - Accepted: February 16, 2010

ABSTRACT

Objective: The objective of the present study was to verify the hypothesis that no difference in biocompatibility exists between different orthodontic adhesives. Material and Methods: Thirty male Wistar rats were used in this study and divided into five groups (n=6): Group 1 (control, distilled water), Group 2 (Concise), Group 3 (Xeno III), Group 4 (Transbond XT), and Group 5 (Transbond plus Self-Etching Primer). Two cavities were performed in the subcutaneous dorsum of each animal to place a polyvinyl sponge soaked with 2 drops of the respective adhesive in each surgical loci. Two animals of each group were sacrificed after 7, 15, and 30 days, and their tissues were analyzed by using an optical microscope. Results: At day 7, Groups 3 (Transbond XT) and 4 (Xeno III) showed intense mono- and polymorphonuclear inflammatory infiltrate with no differences between them, whereas Groups 1 (control) and 2 (Concise) showed moderate mononuclear inflammatory infiltrate. At day 15, severe inflammation was observed in Group 3 (Transbond XT) compared to other groups. At day 30, the same group showed a more expressive mononuclear inflammatory infiltrate compared to other groups. Conclusion: Among the orthodontic adhesive analyzed, it may be concluded that Transbond XT exhibited the worst biocompatibility. However, one cannot interpret the specificity of the data generated in vivo animal models as a human response.

Key words: Orthodontic adhesives. Biocompatibility testing. Inflammation.

INTRODUCTION

Orthodontic adhesives are used to provide effective union between composite and dental structure. However, the most common substances present in adhesive systems show a well defined cytotoxic effect. These adhesives are used in moist media and often on contaminated surfaces without compromising their adhesion, but different compounds can be released during the aqueous phase, such as non-polymerized free monomers from resin materials. Some authors have demonstrated that these free monomers caused apoptosis in cell culture. In vivo studies show that non-polymerized resin compounds released from dental adhesives cause a pulp inflammatory response, which is visible and chronic. Recent studies have shown presence of macrophages together with resin compounds following restorative procedure, in which a persistent and chronic inflammatory response was observed over a period of 300 days. In general, the adhesives used in orthodontics are chosen based on the research on mechanical assays and effectiveness in sealing the interface between tooth and orthodontic accessory. However, many research studies on the biocompatibility of dental materials are currently being performed. Taken together, all these findings are in accordance with the idea that the close proximity of orthodontic accessories to gingival and oral tissues makes this issue very important when choosing these adhesives. Therefore, the aim of the present work was to test the hypothesis that there is no difference in biocompatibility between the adhesives used for attaching orthodontic accessories.
MATERIAL AND METHODS

This study used 30 male adult Wistar rats weighing 250-350 g, which were divided into five groups of 6 animals each: Group 1 (control, distilled water), Group 2 (Concise, 3M Unitek Orthodontic Products, Monrovia, CA, USA), Group 3 (Xeno III, Dentsply/DeTrey, Konstanz, Baden-Württemberg, Germany), Group 4 (Transbond XT, 3M Unitek Orthodontic Products) and Group 5 (Transbond Self-Etching Primer, 3M Unitek Orthodontic Products) (Figure 1). The rats were anesthetized with intraperitoneal injection of sodium thiopental (50 mg/kg) (THIO, Cristália, Itapira, SP, Brazil). Two midline incisions of approximately 18-mm-deep surgical loci each. All animals received two PVA sponge implants (4.0 mm long x 2.0 mm diameter). The implants were approximately 18-mm-deep surgical loci each. All animals received two PVA sponge implants (4.0 mm long x 2.0 mm diameter). The implants were previously kept in 70% alcohol for 120 min, rinsed with sterile distilled water, autoclaved and then soaked with 2 drops of the respective adhesives. The implants in the sponges were photoactivated with a LED source unit (Radii, SDI, Baywater, Victoria, Australia) according to the application time recommended by the manufacturer. The light intensity of the curing unit (1000 mW/cm²) was checked immediately before each polymerization using a radiometer (Model 100, Demetron Research Corporation, Danbury, CT, USA). The surgical loci were sutured with 4.0 suture (Ethicon, Johnson & Johnson, São José dos Campos, São Paulo, Brazil) and then the animals received an injection of sodium dipyrone (0.3 mL/100 g Novalgina®; Sanofi-Aventis Farmacêutica LTDA, Suzano, SP, Brazil).

The rats were kept in cages and fed balanced food and water. After 7, 15, and 30 days, the animals were anesthetized and submitted to excisional biopsy at the implantation area so that enough surrounding normal tissue could be collected. Each group consisted of 6 rats with two implants, thus resulting in 12 samples per group (Table 1). Next, the animals were sacrificed by cervical dislocation.

After being fixed in 4% formaldehyde (Milyn solution) for 24 h, the samples were inserted into paraffin and then 6-µm-thick histological sections were cut and stained with hematoxylin and eosin. The inflammatory responses induced by the adhesives were examined with a light microscope and classified as mild, moderate, and severe17,19. The biocompatibility of the materials was determined according to the ISO 10993-313 standard.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Adhesive Primer</th>
<th>Composition</th>
<th>Manufacture</th>
</tr>
</thead>
<tbody>
<tr>
<td>Concise</td>
<td>Concise® Orthodontic Adhesive</td>
<td>Resin A: Bis-glycidyl-methacrylate (Bis-GMA), triethylene-glycol-dimethacrylate (TEGDMA);</td>
<td>3M Unitek, Monrovia, CA, USA</td>
</tr>
<tr>
<td>Xeno III</td>
<td>Xeno® III Single Step Self Etching Dental Adhesive</td>
<td>Fluid A: 2-hydroxyethyl methacrylate (HEMA), Purified water, Ethanol, Toluene hydroxybutylate (THB), Amorphous silica.</td>
<td>Dentsply DeTrey, Konstanz, Baden-Wurttemberg, Germany</td>
</tr>
<tr>
<td>Transbond</td>
<td>Transbond® XT Primer</td>
<td>Fluid B: Phosphoric acid modified methacrylate (Piro-EMA), Phosphazen mono-fluoride (PEM-F), Urethane dimethacrylate, Toluene hydroxybutylate (THB), camphoroquinone, Ethyl-4-dimethylaminobenzoate.</td>
<td>3M Unitek, Monrovia, CA, USA</td>
</tr>
<tr>
<td>TP Sep</td>
<td>Transbond® Adhesive Plus Self Etching Primer (SEP)</td>
<td>Mono and di-HEMA phosphates, camphoroquinone, distilled water, aminobenzoate, potassium hexafluoride titanate, Butylhydroxytoluene, methylparaben, and propylparaben.</td>
<td>3M Unitek, Monrovia, CA, USA</td>
</tr>
</tbody>
</table>

Figure 1- Composition of the tested adhesive primers
RESULTS

vessels and circulatory changes (dilatation and edema) around and within the cavity as a result of the material implantation in all four groups of adhesive systems (Figure 2AB). Groups 3 (Xeno III) and 4 (Transbond XT) showed the most observed due to the presence of sponge. In Group 2 (Concise), there was formation of granuloma and presence of multinuclear giant cells (Figure 2AB), which indicates the beginning of a repair process (Table 2).

At the end of the 15-day period, it was observed for all adhesive systems compared to that at day 7, except for Transbond XT (Figure 3A-B), which showed presence of neutrophils and abscess formation at the region where the material was implanted, thus indicating a very toxic effect on the tissue. Presence of granuloma and multinuclear giant cells were observed in Groups 3 (Xeno III) and 5 (Transbond SEP) as well as in the Control Group (Figure 3A-B) (Table 2).

After 30 days, the Control Group and all four processes characterized by discrete mononuclear cells.

<p>| Table 1- Distribution of the groups according to type of adhesive system and sacrifice day |
|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|-----------------------------------------------|</p>
<table>
<thead>
<tr>
<th>Days</th>
<th>Control</th>
<th>Concise</th>
<th>Xeno III</th>
<th>Transbond XT</th>
<th>Transbond Plus SEP</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Days</td>
<td>Rats</td>
<td>Samples</td>
<td>Rats</td>
<td>Samples</td>
</tr>
<tr>
<td>7 days</td>
<td>2</td>
<td>4</td>
<td>2</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>15 days</td>
<td>2</td>
<td>4</td>
<td>2</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>30 days</td>
<td>2</td>
<td>4</td>
<td>2</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>Total</td>
<td>6</td>
<td>12</td>
<td>6</td>
<td>12</td>
<td>6</td>
</tr>
</tbody>
</table>

SEP: Self-Etching Primer

Figure 2- Photomicrographs of histological samples after 7 days of implantation. A: evidence of acute inflammatory infiltrate with predominance of polymorphonuclear cells (1,000x magnification; scale: 100 µm). B: presence of granulation tissue (1,000x magnification; scale: 100 µm).

Figure 3- Photomicrographs of histological samples after 15 days of implantation. A: formation of granulomas with multinuclear giant cells (1,000x magnification; scale: 100 µm). B: areas of intense cell formation and deposition of collagen fibers (1,000x magnification; scale: 100 µm).
The quality and specificity of the data generated by in vivo models is questioned and criticized in literature, depends of use of a biological system that reproduces as close as possible the metabolic behavior of the target organ for the toxic effect of xenobiotics and of choice of appropriate parameters to evaluate toxic effects. Evaluating the biocompatibility of orthodontic adhesives by means of subcutaneous implants in rats is of great value as the tissue response in rats is similar to that expected when the same material is applied to the gingival tissue surrounding the area to receive orthodontic accessories. Several studies have assessed the biocompatibility of dental materials. However, methodological divergence exists. In the present study, polyvinyl sponges saturated with the respective adhesives were inserted into rats subcutaneous tissue and then light cured in an attempt to simulate actual clinical procedures. Costa, et al. (1999) have used polyvinyl sponges saturated with adhesives that had not been photoactivated after surgical implantation, allowing the adhesive and their monomers to be in close contact with the subcutaneous connective tissue. Therefore, not only a cytotoxic effect of the dental adhesive was observed but also a persistent inflammatory reaction resulting from the resin infiltration, discrete vascular alterations, and formation of granuloma with multinuclear giant cells and fibrous tissue formation around the samples (Figure 4). Presence of granulation tissue around the sponges and cell proliferation with deposition of collagen fibers correspond to the repair process and fibrosis, respectively. Therefore, all adhesives were shown to be biocompatible on long-term basis (Table 2).
compounds. According to the authors, these materials do not seem to be suitable for direct application to connective tissue.

Studies\textsuperscript{15,21,22} have demonstrated the cytotoxicity of the compounds of adhesive systems, which can be explained by the different compositions, mechanisms, and application procedures as well as by methodological variations\textsuperscript{1}. However, it is clear that the choice for a given adhesive system should be based on its biocompatibility.

The most often studied method for \textit{in vivo} analysis of biocompatibility relies on inflammation\textsuperscript{17}. Analyzing and rating the inflammation phenomena in different experimental groups makes it possible to establish the best biocompatibility by placing the material in contact with vascularized tissues and observing the different reactions. It is also important to use an innocuous substance in the control group in order to facilitate data interpretation\textsuperscript{17}.

The inflammatory response begins with a more intense reaction to both surgical procedures and implanted foreign body, and because such a reaction is not specific, the first post-surgical hours are not taken into account. After 7 days, a more organized inflammatory reaction was expected due to the adhesive rather than the surgical procedure.

The decrease in the inflammatory intensity relies on the control of the host defense system, which organizes itself to limit the aggressive action from the compounds existing in the adhesives and to localize the inflammatory reaction. It was possible to differentiate the inflammation intensity between the experimental groups, mainly regarding the higher level of cytotoxic from Xeno III and Transbond XT after 7 days.

Methacrylate monomers such as TEGDMA, Bis-GMA, UDMA, and HEMA, which are largely used in the composition of dentinal adhesives, can cause cell lesion\textsuperscript{10,12}. TEGDMA, Bis-GMA, UDMA are hydrophobic monomers that are often associated with HEMA. Diffusion of these monomers can be facilitated because HEMA increases the hydrophilic characteristic of the material. Under such conditions, the hydrophobic monomers can reach the cells and damage them\textsuperscript{10,12,23}.

With respect to Xeno III, the presence of HEMA in association with ethanol seems to cause more cell damage. The ethanol in relation with the oral mucosa showed increased mucosal permeability\textsuperscript{13,24} and penetration of potential carcinogens across the mucosal permeability barrier\textsuperscript{13}. It has been reported that topical application of ethanol on the oral mucosa affects epithelial cell homeostasis\textsuperscript{2} and alters mucosal structure\textsuperscript{16}.

In the present study, a mild reaction was observed in the control group whose sponges were saturated with distilled water, whereas a moderate to severe reaction was found in all experimental groups. After 15 and 30 days, the inflammatory reaction was visible, thus allowing the biocompatibility of the materials to be rated in ascending order. In general, small necrotic areas with edema surrounded by cell proliferation, consequent deposition of collagen fibers, chronic inflammation, and decreased number of blood vessels were observed. The presence of multinuclear giant cells suggests the formation of granulomas due to the presence of sponge and/ or adhesive material. Therefore, these events are described as a favorable tissue response regarding the biocompatibility of the material.

At the end of the 30-day period, it was possible to observe that all adhesive systems showed good biocompatibility, although Transbond XT was found to be more aggressive compared to other groups as formation of abscess occurred at the implant region. Based on this finding, one can conclude that Transbond XT is the least biocompatible adhesive.

According to the literature, HEMA is an important toxic component released by most adhesive systems since several \textit{in vitro} studies have demonstrated a defined cytotoxicity of HEMA to the culture of cells\textsuperscript{1,25}. Methacrylate monomers, such as HEMA, are incorporated in the lipid bilayers of cell membranes which are solubilized by the unreacted monomers\textsuperscript{10}. This mechanism of action of uncured leached monomers on the cell membrane may be regarded as responsible for the high cytotoxicity of Transbond (Transbond XT, 3M Unitek) observed in the present investigation.

Traditionally, persulfate molecules have been used as initiators in redox water-based polymerization systems to decrease the amount of residual monomers after setting\textsuperscript{14}. The high cytotoxicity of adhesive systems is probably caused by leachable resin components, such as TEGDMA, Bis-GMA and HEMA, which has frequently been added to their chemical composition\textsuperscript{9}.

However, it may be speculated that some minor adhesive components released into the connective tissue, such as HEMA, which presents low molecular weight, might be removed by local lymphatic drainage. This hypothesis should explain why the inflammatory reaction decreased with time and the connective tissue healing occurred for all experimental materials at 30 days following the implantation.

Sohoeil, et al.\textsuperscript{22} (1994), after testing adhesives in pigs, have suggested that orthodontic adhesives can be potentially allergenic for human being, particularly the “no-mix” ones, and lead to adverse reactions in both patients and practitioners. Such cytotoxicity can last two years after polymerisation\textsuperscript{27}.

Thompson, et al.\textsuperscript{28} (1982) have concluded that even adequately mixed and set, the orthodontic adhesives showed great amounts of material that not had been cured (up to 14% of the material), thus resulting in potential toxicity. Therefore, in addition to adequately preparing and applying these products, the clinician should be careful not to expose skin, mucosa, and gingival to these
materials for long periods of time, particularly the subgingival and interproximal areas.

CONCLUSIONS

It is possible (and safer) to evaluate inflammatory and healing phenomena to characterize, and rate the experimental groups by comparing them to a control group. This allows us to state that Xeno III, Transbond SEP, and Concise adhesives had the best biocompatibility, since formation of chronic inflammation with peripheral healing phenomena and multinuclear giant cells around the samples were observed. However, one cannot interpret the specificity of the data generated in vivo animal models as a human response. The hypothesis was rejected and one can state that, among the adhesives studied, Transbond XT was found to have the worst biocompatibility.

REFERENCES