Polyphenol-Enriched Extract of Lacquer Sap Used as a Dentine Primer with Benefits of Improving Collagen Cross-Linking and Antibacterial Functions

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ABSTRACT: Commercial dentin adhesive systems are applied to restorations due to their resistant bonding properties, but they suffer from the lack of bioactivity and are prone to hydrolysis. Therefore, to overcome these limitations, an eco-friendly natural monomer, urushiol, was adopted to be a primer in dentin bonding due to its interaction with collagen and antibacterial activity, preventing further hydrolysis development. First, urushiol was determined to be capable of improving the biological stability of dentin collagen through cross-linking. Using high-fidelity analytical chemistry techniques, such as Fourier transform infrared spectroscopy, we quantified the effects of urushiol on collagen molecules. It could also effectively decrease weight loss after collagenase ingestion by improving the stability of dentin. Moreover, urushiol inhibited Streptococcus mutans growth as well as its biofilm formation. Finally, we demonstrated that the urushiol primer could improve the bonding strength, particularly after aging. The cross-linking and antibacterial functions of urushiol have provided promising developmental prospects for biomaterials in dentin adhesion.

KEYWORDS: urushiol, antibacterial activity, cross-linking, primer, bonding strength

1. INTRODUCTION

A hybrid layer, composed of infiltrated resin monomers and demineralized dentin collagen is the key to dentin bonding. In the etch-and-rinse mode, however, this resin–dentin composite is vulnerable to enzymatic degradation with time as the collagen fibrils that are incompletely infiltrated by monomers and the current bonding resins are not able to totally remove the free and slackly bound water within the collagen matrix. Moreover, the cured adhesive layer may act as a semipermeable membrane that allows water diffusion from the bound dentine to the hybrid layer, leading to the reliable long-term dentin bonding being challenged. Endogenous proteases, including the matrix metalloproteinases (MMPs) and cysteine cathepsins, are exposed and reactivated after etching, and progressively degrades collagen and results in the loss of adhesive restoration retention.

From the relationship between the dentin structure and adhesive, the collagen structure of the demineralized dentin is unavoidably subjected to degradation, whatever the diverse situations in oral. Enhancement of collagen cross-links has been proven to increase the resin bonding. External collagen cross-linking agents may cause supplementary inter- and intramolecular cross-links, improving the bonding durability of the demineralized collagen network in coronal dentin along with its resistance to enzymatic degreda-
of years ago.\(^\text{30}\) Urushiol, a natural monomer obtained from the raw lacquer sap that has a catechol group and a linear unsaturated side chain,\(^\text{31}\) not only has antibacterial and antioxidant properties\(^\text{32}\) but also excellent corrosion and water resistance.\(^\text{33,34}\) Its antibacterial activity may be related to attacking the outer membrane and bleb formation on bacteria.\(^\text{35}\)

Moreover, the catechol group of urushiol may have the ability to promote collagen cross-linking with its intrinsic antibacterial activity, which can be used in dentistry. Although urushiol has been used for many years, almost no study has explored its effects as a primer on adhesive bonding and the structure of demineralized dentin. In addition, a primer should have comparable interests, and more profitably the benefit is available to etch-and-rinse adhesive, as collagen is an important part of the interface. The motivation of this research is the potentials of the promotion effect of the collagen cross-linking primer in dentin bonding. Herein, we established for the first time an urushiol primer that can simultaneously exert collagen cross-linking and antibacterial effects to achieve durable dentin bonding in a complex oral environment even after aging. The antibacterial activity of urushiol was demonstrated against \textit{S. mutans}. Furthermore, to obtain a more representative estimation of the urushiol primer, we performed the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay on the \textit{S. mutans} biofilm on the dentin surface treated by the urushiol primer. The outcomes indicate that pretreating the dentin surface with the novel multifunctional primers provides a potential reliable operative strategy for dentists to enhance the durability of dentin bond strength.

2. MATERIALS AND METHODS

2.1. Materials. Urushiol was purchased from Wuhan National Lacquer Co., Ltd. (Wuhan, China). Collagenase with a molecular weight of \(\sim 110\) kDa was obtained from Sigma-Aldrich (Beijing, China). Ethanol, dimethyl sulfoxide (DMSO), glutaraldehyde (GA), \(\text{CaCl}_2\), sodium azide, and phosphate buffer saline were purchased from Aladdin (Shanghai, China). The \textit{S. mutans} ACTT UA159 (Maryland) was incubated in the brain heart infusion (BHI) broth from QingDaoHopebio-Technology Co., Ltd. (Qingdao, China) supplemented with 5% sucrose at 37 °C with 5% CO\(_2\) for 48 h. The hydroxyproline assay kit was purchased from Solarbio (Shanghai, China). Resin composite Filtex Z350 XT and Adper Single Bond 2 were obtained from 3 M (St.Paul).

2.2. Collagen Cross-Linking. The collagen cross-linking test process is illustrated in Figure 1.

2.2.1. Preparation of the Urushiol Solution. Urushiol (0.5, 1, and 3 wt %) was dissolved in ethanol. The structure of urushiol is shown in Figure 2. Glutaraldehyde was dissolved at 5 wt % in distilled water as the positive control group. Deionized water was the blank control group. The solutions were buffered to pH 7.2.

2.2.2. Preparation of the Dentin Specimens. Non-carious human third molars were obtained from the patients of certain ages (18 < age < 30 years) and approved by the Ethics Committee of the Stomatology Hospital of Jilin University. The root and occlusal portions of teeth were cut away by a slow-speed water-cooled diamond saw (Kejing, China) to obtain a 1.2 mm thick slab in the middle, which was cut into beams (1 mm \(\times\) 2 mm \(\times\) 6 mm). Dentin slabs and beams were immersed in 10% phosphoric acid for 48 h to be completely demineralized and then characterized by digital radiography. The samples were randomly distributed into 7 groups and then immersed in 1 mL of each solution \((n = 15)\). After biomodification, the samples were rinsed with excessive deionized water and dried in a desiccator for 24 h.

2.2.3. Fourier Transform Infrared (FTIR) Spectra. The deminer- alized dentin collagen beams were respectively soaked in deionized water and in the experimental urushiol solutions for 1 min and 24 h, and then rinsed and dried. The biomodification effect of samples was detected by Fourier transform infrared (FTIR) spectra at a resolution of 4 cm\(^{-1}\). The slabs were placed on the diamond crystal top plate of an attenuated total reflectance (ATR) accessory with a gauge force of 100 N. The range of all spectra was between 650 and 3700 cm\(^{-1}\).

2.2.4. Thermogravimetric Analysis (TGA) Test. The beams were treated with distilled water (control), GA and urushiol solutions for 24 h. Thermogravimetric analysis was performed under a nitrogen
atmosphere using a thermogravimetric analyzer (TA instrument). Temperature increased at the rate of 10 °C/min from 35 to 700 °C.

2.2.5. Contact Angle (CA). The hydrophilicity of the slabs (n = 5) was determined with a goniometer (Dataphysics OCA-20, Germany) by dropping 6 μL of deionized water onto the slab surface, which showed both left and right contact angles from the shape of the water drop at a room temperature of 25 °C.

2.2.6. Hardness. The Vickers hardness numbers (VHNs) of the samples were gauged using a Vickers hardness tester with a load of 50 g for 15 s. Each sample had five indentations on the surface, which were spaced at least 100 μm apart.

2.2.7. Swelling Ratio. The specimens were swollen by immersion in water at room temperature and equilibrated overnight in phosphate-buffered saline (PBS) (pH 7.4). The samples were weighed immediately after removing excess water from the surface with filter paper. Dentin specimens were then immersed in a large amount of deionized water to remove the buffer salts and dried to a constant weight. The swelling ratio was the weight of the swollen sample compared to the weight of the dried sample.

2.2.8. Collagenase Treatment. TESCA buffer was prepared by adding 11.5 g of N-tris(hydroxymethyl)-methyl-2-aminoethanesulfonic acid, 53 mg of CaCl₂·2H₂O, and 50 mg of sodium azide in distilled water to 1000 mL of deionized water and adjusting the pH to 7.4. Then, 100 μg of collagenase with a molecular mass of ~110 kDa was added to the TESCA buffer at the final concentration of 0.1% (w/v).

2.2.9. Mass Change. Demineralized dentin beams were respectively soaked in different solutions for 24 h and then weighed (M₀). Further weight measurements were performed after collagenase degradation (Mᵢ) for 48 h. After treatment, the beams should be dried in a vacuum desiccator with silica gel at room temperature for 72 h. The mass change (Wᵢ₀) was described by the percentage weight loss of individual samples, calculated by the following formula

\[ Wᵢ₀ = [(M₀ - Mᵢ) \times 100%] / M₀ \]  

(1)

2.2.10. Hydroxyproline Release. The collagenolytic activity was evaluated by measuring the hydroxyproline content in collagenase solution. Measurement of hydroxyproline in each sample was performed according to the instructions. Briefly, a 0.06 mL sample (collagenase solution) was mixed with 0.06 mL of reagent A. After a 20 min interval, 0.06 mL of reagent B and 0.12 mL of deionized water were added. Then, the samples were incubated at 60 °C for 20 min and then 0.2 mL of sample solution of each group was dropped into a 96-well plate. By the microplate reader, the absorbance was read at 560 nm.

2.2.11. Scanning Electron Microscopy (SEM). Specimens that were treated by different solutions and after collagenase degradation were air-dried and then sprayed with gold. The micro-morphologies images of samples were observed by scanning electron microscopy (SEM).

2.3. Antibacterial Activities. 2.3.1. Bacterial Strains and Cultivation Characteristics. S. mutans (UA159) was used for measuring the antibacterial activities of 0, 0.1, 0.5, 0.7, and 1 wt % urushiol. After 24 h of culture with the brain heart infusion (BHI), the bacterial strain was diluted to 1 × 10⁶ CFU/mL.

2.3.2. Colony Counting Assay. The growth conditions of bacteria were assessed by the colony counting method. S. mutans were inoculated in the BHI broth with or without different concentrations of urushiol at 37 °C. Approximately 4 × 10⁵ bacterial cells were inoculated into 1 mL of BHI medium with different concentrations of urushiol. At 0, 30, 60, and 90 min of bacterial incubation, the BHI agar plates were coated with the medium and colony-forming units (CFU) were counted. The test was repeated three times.

2.3.3. Antibacterial Evaluation by SEM. Scanning electron microscopy (SEM) graphs were analyzed using 24-well plates with glass slides. Before SEM imaging, S. mutans was treated with urushiol for 48 h at room temperature. The medium was extracted, and the wells were gently washed twice with sterilized deionized water. Samples were placed in a 2.5% glutaraldehyde solution overnight. Then, the specimens were dehydrated in different concentrations of ethanol washes (50, 70, 90, and 100% for 30 min) and air-dried. Before being visualized by SEM (S4800, Tokyo, Japan), the samples were sprayed with gold. The samples were observed in three views randomly for imaging.

2.4. Urushiol as the Primer. Figure 3 presents the process of applying the primer.

2.4.1. Preparation of the Dentin Specimens and Primer Agents. Fifty freshly non-caries molars stored in 0.5% chloramine-T at 4 °C were randomly divided into eleven groups (n = 10). The occlusal portion of the teeth was cut away using a diamond saw to expose mid-depth dentin. The standard smear layer was prepared by grinding the dentin surface with 600-grit silicon carbide sandpaper. Dentin surfaces were etched with 37% phosphoric gel for 15 s and then deeply rinsed with excessive deionized water.
Urushiol (0.1, 0.5, 0.7, and 1 wt %) dissolved in ethanol and DMSO were used as the primer agents.

2.4.2. Degree of Conversion. The effect of urushiol primers on the polymerization of commercial adhesive was explored by the degree of conversion (DC) test with a Fourier transform infrared spectrometer \((n = 5)\). After urushiol pretreatment for 1 min, adhesives were applied and cured according to instructions, which were then immediately tested by FTIR. The DC values were estimated in accordance with the variation of the absorbance peak intensity of carbon–carbon double bonds \((\text{C} = \text{C}, \text{peak at } 1636 \text{ cm}^{-1})\) before and after curing, using the phenyl \((\text{peak at } 1608 \text{ cm}^{-1})\) as the internal reference. The DC was estimated by the following formula

\[
\text{DC} \% = 100 \times \left(1 - \frac{(A_{1637}/A_{1608}) \text{peak area after curing}}{(A_{1637}/A_{1608}) \text{peak area before curing}} \right)
\]

2.4.3. Contact Angle. The dentin slabs were polished with 1000-grit silicon carbide sandpaper and ultrasonically rinsed twice for 5 min to eliminate the residue. The samples were etched with phosphate gel for 15 s and washed with deionized water for 15 s. Then, 6 μL of deionized water was dropped onto the slab surface using a contact angle meter (DataPhysics OCA-20).

2.4.4. Microtensile Bond Strength Test. As the control group, Adper Single Bond 2 was cured for 20 s followed the steps of the manufacturer’s instructions. Then, a resin composite build-up was constructed with two 2-mm increments and was light-cured for 40 s. The samples were stored in deionized water for 24 h at 37 °C. The specimens were cut parallel to the long axis into small rods with cross-sectional area of 1.0 mm² using a diamond saw under flowing water, and at least 6 small rods were expected to be cut out of each tooth. Sample rods were attached to the microtensile molds and the test was performed by a universal testing machine at a crosshead speed of 1 mm/min until failure. The cross-sectional area of the rods was measured with a digital caliper to the nearest 0.01 mm, and the microtensile strength (MPa) was the ratio of the maximum load to the cross-sectional area of the sample.

2.4.5. Aging. Half of the bonded samples were randomly selected to be aged (5 °C for 30 s and 55 °C for 30 s) for 5000 cycles in the thermocycling device (PTC2c; Proto-tech, Portland, ME). The μTBS test was performed as described above.

2.4.6. Nanoleakage. After thermocycling, the bonded samples were cut into 1 mm thick pieces parallel to the long axis of the tooth, and the middle two pieces were selected to be submerged in the 50% ammoniated silver nitrate solution for 24 h. The specimens were rinsed with deionized water, and then were submerged in the photo development solution with fluorescent irradiation for 8 h. After polished with SiC sandpapers (1000-, 1200-, 1500-, 2000- and 2500-grit), samples were then ultrasonically washed with deionized water for 30 min. The samples were air-dried after dehydration and then observed by SEM. In addition, the surfaces of the samples were determined by energy dispersive spectrometry (EDS).

![Fig. 4. FTIR spectra of collagen treated with 0.5 wt % (A), 1 wt % (B), and 3 wt % urushiol (C).](https://doi.org/10.1021/acsbiomaterials.1c01287)
2.4.7. Antibacterial Activity of Urushiol Primer. 2.4.7.1. Preparation of the Dentin Disk. In accordance with the μTRS test, ethanol-0.7 wt % urushiol and DMSO-0.7 wt % urushiol groups were chosen to test the antibacterial activity of the urushiol primers. The root and occlusal portions of the non-carious teeth were cut away by the slow-speed diamond saw to obtain 0.5 mm thick dentin disks, which were then polished with 600-, 1200-, 2000-, 3000-grit SiC sandpapers under water and diamond paste (0.5 μm) and washed ultrasonically.

2.4.7.2. Bacterial Culture and Biofilm Preparation. Dentin disks were etched by 37% phosphate gel for 15 s, and then disinfected with UV for 1 h each side before biofilm preparation. The samples were pretreated with two primers for 1 min in 24-well plates and 1 mL of S. mutans suspension was added dropwise which was diluted to 10^8 CFU/mL in advance. After culture in an incubator to form biofilm-coated specimens at 37 °C for 24 h, the specimens were gently rinsed with sterile phosphate buffer saline (PBS) twice.

2.4.7.3. Antibacterial Evaluation by MTT Assay. The biofilm-coated specimens (n = 9) were transferred to a new 24-well plate and incubated at 37 °C after adding 1 mL of 0.5 mg/mL MTT solution dropwise to each well to estimate the biofilm metabolic activity. The absorbances of plates were measured using a microplate reader at 570 nm with five readings for each sample. The experiment was repeated three times.

2.4.7.4. SEM. The biofilm-coated specimens (n = 3) were fixed overnight by 2.5% glutaraldehyde solution and dried with ethanol gradient (50, 70, 90, and 100% for 20 min), then sprayed with gold. AIII/Amide III band centered at 1235 cm⁻¹ was attributed chiefly to the stretching vibrations of C–N–O peptide groups in the glycine backbone and proline side chains. In accordance with the Space Structure of Collagen.

3. RESULTS

3.1. Collagen Cross-Linking. 3.1.1. Effect of Urushiol on the Space Structure of Collagen. Figures 4 and 5 show the FTIR spectra of untreated dentin collagen and urushiol-treated dentin collagens. The amide A and B bands at 3300 and 3071 cm⁻¹, generally corresponded to the stretching vibrations of N–H groups, respectively. The peak at 1633 cm⁻¹ was attributed chiefly to the stretching vibrations of C=O peptide groups in the amide I band. C–N stretching at 1544 cm⁻¹ was observed in amide II. Amide III band centered at 1235 cm⁻¹ was attributed to the C–N stretching and N–H bending vibrations from amide linkages, in addition to the wagging vibrations of CH₂ groups in the glycine backbone and proline side chains. It should be noted that the peak of the amide I band related to the triple helix structure of collagen retained its position. However, for amide A, I and II bands were broadened to some extent.

Meanwhile, besides the above changes, the FTIR absorption ratios of CH₂ scissoring to 1450 cm⁻¹, attributed to AIII/4050 theretofore, are also regarded as a measure of protection of the integrity of collagen triple helices. As is shown in Figure 4, the ratios for 3% urushiol-treated collagen slightly decreased from 1.0 for pure collagen.

3.1.2. Thermal Stability of Collagen with Urushiol Treatment. The thermal stability of collagen with urushiol treatment was evaluated by TGA analysis. As shown in Figure 4, the mass loss of collagen without urushiol treatment at 350 °C was about 44.6%, but the weight losses were 39.2 and 40.3% of 3 wt %-urushiol and GA groups. The results for urushiol-modified collagen films of the GA group and 3 wt % urushiol group had the least weight loss at 350 °C.

3.1.3. Contact Angles. As is shown in Figure 5, the addition of urushiol caused the initial contact angle to increase by approximately 20°. The TA-treated collagen had the highest contact angle (P < 0.05). All of the treated collagen had a higher contact angle than pure collagen (P < 0.05).

3.1.4. Hardness. Figure 8 illustrates the surface hardness measurements of the urushiol-treated collagen at different concentrations. The hardness value of the GA group was the highest of all groups (P < 0.05). The specimens treated with the urushiol solutions exhibited significantly higher HVs than...
No significant differences were revealed in HV between different urushiol-treated groups.

**3.1.5. Swelling Ratio.** Figure 9 presents the swelling ratio of the urushiol-treated and untreated collagen beams. The use of urushiol solution at different concentrations significantly decreased the swelling ratio \( (P < 0.05) \) except for the 0.5 wt % urushiol group. The GA group mostly reduced the swelling ratio \( (P < 0.05) \).

**3.1.6. Mass Change.** All of the urushiol-treated groups and GA-treated groups exhibited significantly decreased weight loss \( (P < 0.05) \) (Figure 10). As is shown in Figure 10, with increasing urushiol content, the samples exhibited a lower weight loss than the control group \( (P < 0.05) \). Nevertheless, when the concentration increased to 1 wt %, no significant difference among the higher urushiol content groups was observed. The weight loss of the control group was the highest \( (P < 0.05) \).

**3.1.7. Hydroxyproline.** An in vitro biodegradation assay indicated that the proteins of the control group exposed to bacterial collagenase were almost entirely lost after 48 h. The urushiol- and GA-treated collagen were strongly resistant to collagenase degradation (Figure 11). The HYP release values of the urushiol-treated collagen were lower than that of the untreated collagen regardless of the concentration \( (P < 0.05) \). Among all groups, the HYP release of the 3 wt % urushiol group and the GA group was the lowest.

**3.1.8. SEM Evaluations.** The representative SEM images of dentine surfaces from each group after 1 min or 24 h urushiol ingestion with the lowest and highest concentrations are presented in Figure 12. In pictures D1 and D2 of the control group, the dentin tubules were clear and smooth. After 0.5 wt % urushiol treatment for 1 min, the walls of dentin tubules were wrinkled, as shown in pictures A1 and A2. However, after 24 h, all of the treated collagen showed distinct collagen cross-linking.
After collagenase digestion, the collagen matrix of the control group appeared to collapse, and the dentin structure of the control group could not be seen. However, in the pictures of the 0.5 wt % urushiol group, dentin tubules could be identified, but the dentin collagen was destroyed (Figure 13). Specimens treated with a 3 wt % urushiol preconditioner showed a relatively complete dentin tubule structure.

3.2. Antibacterial Test. 3.2.1. Antibacterial Activity Test. As is shown in Figure 14, no viable S. mutans microorganisms were discovered after a 30 min culture with four concentrations of urushiol.

3.2.2. SEM. Figure 15 shows typical morphological micrographs of S. mutans of the control and urushiol groups. SEM microscopy provided the images of S. mutans. Urushiol (0.1 wt %) affected the membranes of the bacteria. With the increasing concentrations of urushiol, the internal substance of bacteria leaked out. Moreover, the content of normal bacteria decreased. In the control group, the bacteria had a complete shape. When bacteria were treated with 0.7 wt % urushiol, almost none of the bacteria were alive. Samples treated with urushiol exhibited marked differences in morphology compared with control samples.

3.3. Urushiol as a Primer. 3.3.1. Degree of Conversion. As shown in Figure 16, the DC of the primer showed just over 50% polymerization throughout the analysis. The degree of conversion of the ethanol samples increased slowly to approximately 57%. However, that of the DMSO groups decreased gradually. The DC of the ethanol-1% urushiol group was 59.73 ± 0.73% and the DC of the DMSO-1 wt % urushiol group was 47.86 ± 0.41%. There was a significant difference between the control group and other experimental groups ($P < 0.05$).

3.3.2. Contact Angle Test. As Figure 17 shows, the average contact angles (mean ± standard deviation, SD) for the control, 1 wt % urushiol in ethanol and DMSO solvents were 26.4 ± 2.41°, 39.38 ± 1.02°, and 31.12 ± 0.65°, respectively. Compared with the blank control group, the contact angle of the pure ethanol-treated dentin increased significantly ($P < 0.05$). However, the contact angle of the pure DMSO group was lower than pure collagen ($P < 0.05$). The 0.1 wt % urushiol groups had a lower contact angle than the other three urushiol-treated groups that showed gradually increasing contact angles.

3.3.3. μTBS. Table 1 shows the μTBS data of all groups. In the immediate groups, the DMSO-1 wt % urushiol group has the highest bond strength ($P < 0.05$). Nevertheless, among the thermocycling groups, there were no significant differences between the 1 wt % urushiol in the ethanol and DMSO groups. After aging the bond strengths of the ethanol-0.7 wt % and ethanol-1 wt % treated groups were not affected. After 5000 thermocycling cycles of the aging test, the bond strength of all groups decreased to different degrees. The control group had the lowest bond strength among all groups.

Different lowercase letters in the same column imply statistically significant differences ($P < 0.05$; vertical comparisons); different capital letters in the same line imply statistically significant differences ($P < 0.05$; horizontal comparisons).

3.3.4. Nanoleakage. After thermocycling, the Ag had infiltrated into the hybrid layer and the resin tag of the control group, which was confirmed by EDS mapping. As shown in Figure 18, the ethanol-0.7 wt % and DMSO-0.7 wt % groups had less nanoleakage, especially the DMSO primer group.

3.3.5. Antibacterial Activity by the MTT Assay. The results of the MTT assay are shown in Figure 19. The S. mutans biofilm of the 0.7 wt % urushiol-containing ethanol/DMSO pretreated groups had lower metabolic activity ($P < 0.05$).

3.3.6. SEM. Figure 20 shows typical SEM photos of S. mutans biofilms in the control and 0.7 wt % urushiol groups. The bacterial biofilm was observed to cover the entire dentin surface in the control group. After the treatment of dentin samples with 0.7 wt % urushiol-containing ethanol/DMSO primers, few intact bacteria grew, and they were dispersed along the dentin surface. More destroyed bacteria could be seen on the surface.

3.3.7. Cytotoxicity. The toxicity of the urushiol primer toward L929 cells was assessed. As is illustrated in Figure 21, the relative cell viabilities of different groups were all higher than 75%. The cell culture time of 48 h had higher cell viability than the 24 h culture time.
4. DISCUSSION

In the present study, urushiol dissolved in ethanol and DMSO as primers improves the resin–dentin bond strength, especially after aging. The desirable functions of urushiol are achieved.

The FTIR spectra of proteins can correspond to their structures. The triple helix of collagen was maintained due to the unchanged peak at 1633 cm$^{-1}$, which suggested that the treatment did not destroy the structure of collagen. As is shown in the FTIR spectra, the amide A, I and II were broadened, which might involve hydrogen bonding interactions between urushiol and collagen. The mechanism of urushiol interaction with the collagen molecule by hydrogen bonds is shown in Figure 22. Hydrogen bonds have an important effect on thermal stability and triple helix

Figure 12. Representative SEM images of the dentine surfaces of different groups after 1 min and 24 h treatment. GA represents glutaraldehyde.

Figure 13. SEM graphs of the dentin surfaces of different groups after collagenase ingestion. GA represents glutaraldehyde.

Figure 14. Numbers of CFUs in terms of the culture time of S. mutans
ND: S. mutans not discovered, CFU: colony-forming unit.
conformation of collagen. The FTIR spectra in Figures 4 and 5 demonstrate that the different concentrations of urushiol treatment have significant effects on cross-linking. In FTIR spectra, the effect of time on collagen cross-linking was not obvious.

The range of 200–500 °C is involved in the degradation of the collagen chains, which is related to the preservation of the triple helix conformation of collagen, which is essential for the mechanical properties and biocompatibility of collagen-based materials. With urushiol treatment, the stability of treated collagen is enhanced, which indicates that improving the cross-linking of urushiol on collagen has distinct effects.

To investigate the relevance between the microstructure of urushiol-modified collagen and its surface properties, the contact angle of the dentin surface was tested. Urushiol contains phenol hydroxyl groups, which combine with the carboxyl and amino groups in collagen to form hydrogen bonds. Following this, the hydrophilic groups of collagen decrease and the hydrophobicity of collagen increases. The cross-linking of collagen leads to dehydration, which causes a higher contact angle of the dentin surface, and dehydration is the real reason why the cross-linking effect could enhance the stability of collagen.

The results of the surface hardness measurements manifest that urushiol could improve the mechanical properties of collagen by the cross-linking effect. All treated collagen samples had lower water sorption than the untreated collagen except for the 0.5 wt % urushiol group (Figure 9). The collagen fibers stimulated by exogenous cross-linking are dense leading to the reduction in the swelling ratio. The treatment of urushiol could affect the water distribution of the dentin matrix and significantly decrease water sorption.

Owing to the presence of the triple helix structure, collagen fiber has a certain ability to resist endogenous proteases. The binding site of collagenase is situated at a deep gap with a width of approximately 0.5 nm. Some proteolytic enzymes cannot enter such a narrow region. Figure 10 indicates that although the concentration of urushiol is low, treated collagens are durable and could withstand collagenase degradation. Type I collagen consists of approximately 10% hydroxyproline in quality, while most other proteins hardly contain this amino acid. As a result, the hydroxyproline release test is widely used. In this study, upon the urushiol treatment of the collagen, the release of HYP decreased significantly regardless of the concentration. The results are in accordance with the weight loss test. The SEM images indicate after collagenase degradation, urushiol-treated collagen presents a preservation effect as in the images in Figure 13.

Table 1. μTBS of the Experimental and Control Groups at 24 h and 6 m

|     | group          | immediate       | aging          |
|-----|----------------|-----------------|----------------|
|     | ethanol        | 44.73 ± 2.91    | 37.13 ± 2.25   |
|     | ethanol-0.1%   | 54.73 ± 2.89    | 50.37 ± 1.43   |
|     | ethanol-0.5%   | 56.58 ± 1.47    | 52.97 ± 0.87   |
|     | ethanol-0.7%   | 58.66 ± 1.20    | 57.77 ± 1.58   |
|     | ethanol-1%     | 59.95 ± 0.95    | 58.69 ± 0.61   |
|     | DMSO           | 53.05 ± 2.96    | 38.05 ± 1.38   |
|     | DMSO-0.1%      | 56.46 ± 1.82    | 49.75 ± 1.56   |
|     | DMSO-0.5%      | 59.84 ± 3.15    | 55.10 ± 0.59   |
|     | DMSO-0.7%      | 63.53 ± 1.46    | 60.16 ± 1.26   |
|     | DMSO-1%        | 65.10 ± 1.14    | 61.64 ± 2.51   |
|     | control        | 27.53 ± 3.28    | 22.87 ± 3.32   |

**MPa, mean ± SD.**
Urushiol is a kind of herbal medicine for the treatment of bacteria and cancer. S. mutans is chosen for study because it is widely known as the major bacteria in caries. No S. mutans was discovered after 30 min of culture with urushiol (Figure 14). In the primer part, on the basis of MTT results, the metabolism of the S. mutans biofilm in 0.7 wt % urushiol dissolved in two solvents groups was lower in comparison with the control group. Based on the result and SEM images of Figures 19 and 20 show that urushiol has antibacterial activity even at low concentrations.

In fact, the mechanism of several antimicrobial drugs in clinical treatment involves preventing the synthesis of the bacterial cell wall. Urushiol is a polyphenol with a catechol group. The cell wall of bacteria seems to be the major molecular target for the antibacterial effect of most polyphenols. Polyphenols can destroy the morphology of bacteria and structural integrity of intracellular matrix. One underlying mechanism related to the degradation of the bacterial cell wall could be the accumulation of certain essential surface proteins that leads to its inactivation. The targets of

Figure 18. Nanoleakage images of the 0.7 wt % urushiol primer groups and the control group.

Figure 19. Average OD570 values after S. mutans incubation on dentin surfaces for 24 h. Different letter cases represent significant differences at P < 0.05.

Figure 20. SEM images of the dentin surface with cultured S. mutans of the 0.7 wt % urushiol groups and the control group.
Polyphenols are not only proteins but also their ability to form various noncovalent interactions such as hydrogen bonds, van der Waals forces, and hydrophobic interactions with proteins and other molecules. Moreover, possible targets for polyphenols are membrane-bound enzymes, cell surface adhesion proteins, and cell wall polypeptides. The antibacterial activities of urushiol derivatives with alkyl chains of different lengths were tested in other studies. The results showed that alkyl side chains in urushiol derivatives inhibited food spoilage and microbial growth. In conclusion, the mechanism of the antibacterial activity of urushiol is related to the combined effects of the catechol group and the alkyl chain.

Urushiol can fulfill the role as a priming agent on dentin due to its bioactive functions such as collagen cross-linking enhancement and its antibacterial activity. Commonly with polymer restorative materials, a high degree of conversion has an important effect on the mechanical properties. The higher permeability caused by residual monomers accelerates hybrid layer degradation and reduces its sealing ability. According to the DC results, although urushiol has an inhibitory effect on polymerization, low concentrations have no distinct effect on the adhesive. After pure ethanol was applied to collagen, water on the dentin surface evaporated with the ethanol leading to a higher contact angle. DMSO is a water-soluble solvent; as a result, the contact angle is lower than pure collagen. However, with increasing concentrations, the contact angle increased, which might indicate that the collagen cross-linking effect was effective.

In this study, compared with controls the \( \mu \)TBS results demonstrate that the values of urushiol-pretreated groups are significantly higher, especially the DMSO-1 wt % and ethanol-0.7 wt % urushiol groups after aging. DMSO has the ability to modify the collagen structure and improve dentin wettability to enhance bonding effectiveness even under dry conditions. As DMSO is capable of breaking the water self-associative inclination, it could improve the dried collagen re-expansion. To some extent, the use of DMSO as a pretreatment could reduce the sensitivity of the technique. Moreover, DMSO-pretreatments significantly reduced the activity of endogenous collagenase in dentin. The ethanol wet-bonding (EBW) theory suggests that the water in the dentin matrix could be replaced by ethanol and enhance the penetration of hydrophobic resin monomers into the collagen network. Therefore, the high-grade hydrophobic hybrid layer can be formed to develop bonding strength and durability. Ethanol was chosen in this study as a solvent. The compound urushiol and ethanol may withstand the interfacial biodegradation resulting from intraoral temperature changes to improve bonding durability. Nanoleakage was first described as an occurrence in which porosity can be seen all around the base of the hybrid layer. The nanoleakage of the dentin adhesive interface after aging suggests hydrolysis, product exudation, and destruction of collagen fibers after polymerization. With the urushiol primer treatment, there were obviously lower sliver particles deposited on the hybrid layer, which indicated that the primer application had better interface sealing.

According to the CCK-8 results (Figure 21), the biocompatibility of primers was acceptable, and with increasing culture time, the cytotoxicity of primers decreased. Owing to the pressure of dentinal tubules, it may be hard for urushiol to penetrate the long dentin tubules, which prevents it from affecting the pulp cells. Because of the multiple effects of urushiol, it may have great potential in dentistry. In our study, the effect of the urushiol primer on adhesive–dentin bonding was assessed. The results show that the pretreatment of urushiol solutions could enhance the aging resistance of the interface of dentin bonding, which may result from improving collagen cross-links and inhibiting the growth of \( S. \) mutans. It is of great significance to promote the long-term stability of resin restoration.
5. CONCLUSIONS

Urushiol (0.5, 0.7, and 1 wt %) showed obvious antibacterial activity and collagen cross-linking. Moreover, primers of ethanol-0.7 wt % and DMSO-1 wt % solutions applied for 1 min were effective at improving the integrity of the bonding interface and bond strength after aging. Therefore, urushiol may have promising applications in dentin adhesion.

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Notes
The authors declare no competing financial interest.

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