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Facile coating of low-molecular-weight stretchable adhesive films leveraging carbodiimide-to-urea conversion and gallic acid for enhanced adhesion†

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We present a simple method for bonding an elastomer substrate using a stretchable, low-molecular-weight adhesive. By mixing 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide (EDC) and gallic acid on an aminated surface, EDC is converted into a urea derivative. This derivative features both hydrogen bond donor and acceptor functionalities, combined with bio-inspired adhesive gallol groups, which collectively enhance adhesiveness through extensive hydrogen bonding. Importantly, the adhesive preserves the stretchability of the elastomer.

1. Introduction

Adhesives play a vital role in the robust attachment of biomedical devices to biological tissues or living bodies. Numerous stretchable devices have been reported for biosensing^{1–8} and therapeutic applications.^{9–11} In most cases, these devices require an adhesive for long-term use on the target tissue. Potential adhesive materials for stretchable substrates are either polymeric materials (e.g. polydopamine,^{11,12} graphene oxide-based bioadhesives,⁷ polyethylene glycol-lactide acid diacrylate hydrogels,⁸ and gallol-functionalized hydrogels^{13–15}) or low-molecular-weight materials (e.g. ionic liquids¹⁶ and supramolecular materials^{17,18}). Although polymeric adhesives have strong adhesive performance due to abundant covalent and cross-linked structures,^{7,8} these curable adhesives require careful surface treatment and have a risk of impairing the flexibility or stretchability of the elastomer substrates. In contrast, the adhesiveness of low-molecular-weight adhesives (LMWAs) can be tuned based on their various *in situ* non-covalent interactions, and they have larger absolute adhesion areas owing to a

higher degree of freedom for molecular diffusion.¹⁹ Although LMWAs are strongly adhesive, most are also highly viscous.^{16–18} This high viscosity of LMWAs may impair the stretchability of the substrate. Therefore, further optimization of non-covalent bonds of LMWAs is necessary to balance adhesiveness with the stretchability of elastomeric substrates.

To ensure the stretchability of LMWAs, we aimed to increase the formation of additional non-covalent bonds from the precursor solution by applying the conversion of carbodiimide to urea derivatives. Liang *et al.* created an adhesive that used urea derivatives as hydrogen bond donors for efficient adhesion.²⁰ Urea derivatives are often produced as by-products from carbodiimides such as 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide (EDC) during amide bond formation. EDC has also tertiary amino hydrochloride groups that can be used as hydrogen bond acceptors.²⁰ Therefore, in the present study, we used EDC as an adhesive precursor solution. In addition to exploiting this conversion from carbodiimide to urea derivatives, we introduced bio-inspired adhesive-functional gallol groups as auxiliary hydrogen bond donors.^{15,21} They were used to supplement the hydrogen bonds from urea and amino groups for stronger adhesiveness.²²

Herein, we report a facile method for coating a low-molecular-weight stretchable adhesive on an aminated substrate of polydimethylsiloxane (PDMS), one of the versatile elastomers (Fig. 1a). EDC and gallic acid (GA) are mixed on the surface of PDMS so that the EDC is converted to 1-ethyl-3-(3-dimethylaminopropyl)urea (EDU) for the formation of an adhesive layer with numerous hydrogen bond acceptors and donors from gallol, urea, and amino groups (Fig. 1b and c). The simultaneous modification of the surface of the PDMS substrate by gallol *via* amino groups produces an adhesive substrate (Fig. 1b).

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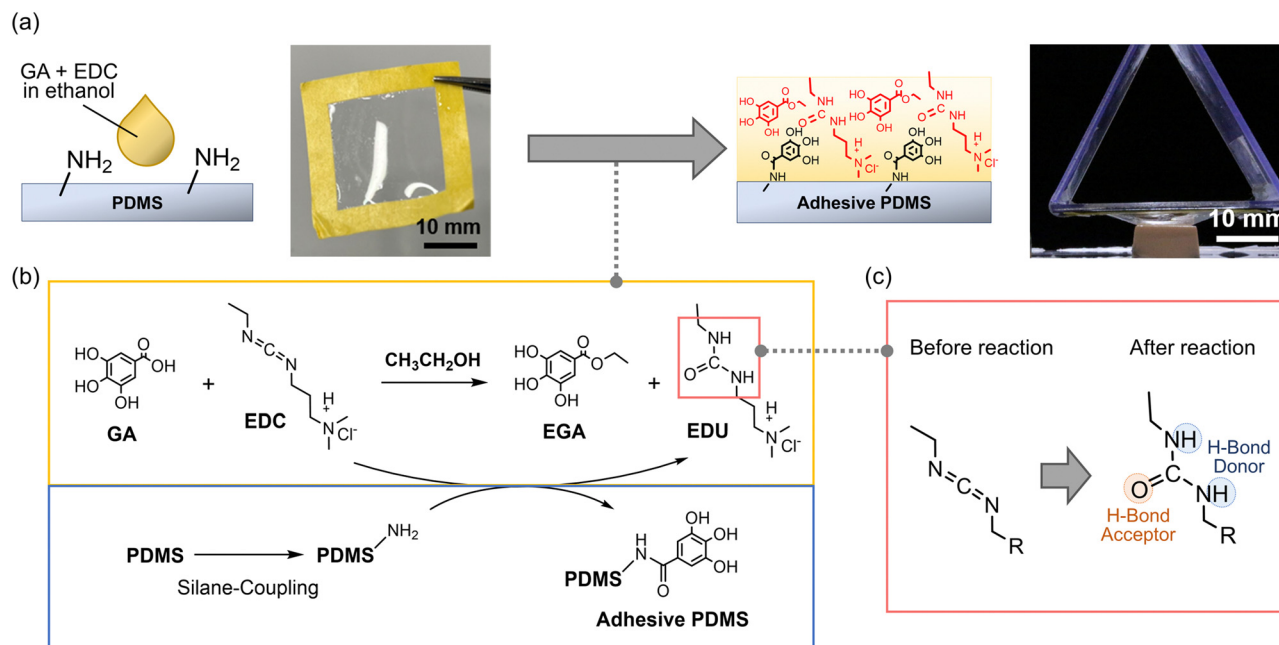


Fig. 1 (a) Schematic showing the modification of a PDMS film by casting GA and EDC in ethanol to enhance the adhesiveness of the PDMS film, (b) the reaction scheme, and (c) accompanying substitution of carbodiimide to urea which can form hydrogen bonds (PDMS = polydimethylsiloxane; GA = gallic acid; EDC = 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide).

When applied to an aminated PDMS synthesized through hydrophilization by plasma treatment²³ followed by a silane coupling reaction²⁴ (Fig. S1, ESI[†]), the adhesive enhanced the adhesiveness of PDMS fabricated by bar-coating. The adhesive coating does not sacrifice the stretchability of the original PDMS. The simultaneous adhesiveness and stretchability are attributed to the energy dissipation and robust cohesion ensured by hydrogen bonds in the adhesive layer.

2. Materials and methods

2.1. Materials

Polyvinyl alcohol (PVA, M_w : 13 000–23 000) was purchased from Kanto Chemical (Tokyo, Japan). Polyethylene terephthalate (PET, 12 cm width, Lumirrors L-25T60) was purchased from Toray Industry Inc. (Tokyo, Japan). Ethyl acetate, hexane, super-dehydrated ethanol (99.5%), dimethyl sulfoxide-*d*₆ (DMSO-*d*₆, 99.9%, containing 0.05 vol% tetramethylsilane (TMS)), and 1-ethyl-3-(3-dimethylaminopropyl)urea (EDU) were purchased from FUJIFILM Wako Pure Chemical (Osaka, Japan). Polydimethylsiloxane (PDMS, SILPOT 184) was purchased from Dow-Toray Specialty Materials (Tokyo, Japan). 3-Aminopropyltrimethoxysilane (APTMS), gallic acid (GA), 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC), and ethyl gallate (EGA) were purchased from Tokyo Chemical Industry Co., Ltd (Tokyo, Japan). Scotch tape (243J Plus) was purchased from 3M Company (MN, USA). The glass substrate (no. 5 thickness, 30 × 30 mm²) was purchased from Matsunami Glass (Osaka, Japan). The nylon mesh (HC-15) was purchased from Sefar Holding AG (Thal, Switzerland). 70% Ethanol

(alcohol preparations ET-N) was purchased from UENO FOOD TECHNO INDUSTRY, Ltd (Tokyo, Japan). The bio-skin model was purchased from Beaulax Co., Ltd (Saitama, Japan).

2.2. Fabrication of the PDMS films

A PET film was coated with a sacrificial layer comprising PVA solution (7 wt% in water) using a roll-to-roll gravure printing system (Tabletop Mini-Labo Test Coater, Yasui Seiki Co., Ltd, Tokyo, Japan) under 30 rpm gravure roll rotation and 1.3 m min⁻¹ line speed with in-line thermal treatment at 80 °C. The PVA film was coated with a PDMS solution (the ratio of the base to the curing agent was 10:1 (w/w)) (30 and 60 wt% in a mixed solvent of hexane/ethyl acetate = 4:1 (w/w)) using a desktop coater (TC-1, Mitsui Electric Co., Ltd, Tokyo, Japan) and a wireless bar coater (OSP-100-L60, OSG System Products Co., Ltd, Aichi, Japan) at a moving speed of 10 mm s⁻¹ with a platform at 25 °C to fabricate the 9.4 and 40 μm PDMS films on PVA. The 133 μm thick film was fabricated in a similar way using a 100 wt% PDMS solution and double-layered scotch tape to raise the height of the processing bar. The PDMS with various thicknesses were crosslinked by heating at 100 °C for 1 h. Scotch tape was cut to a square frame with inner dimensions of 20 × 20 mm² and outer dimensions of 30 × 30 mm² using a laser processing machine (VLS2.30DT, Universal Laser System, Inc., Kanagawa, Japan). Subsequently, the PDMS/PVA film was peeled off from the PET film using the frame of scotch tape attached around the side of the PDMS film (20 × 20 mm²). Finally, the PVA sacrificial layer in the detached PDMS/PVA film was dissolved in water to obtain the PDMS film.



2.3. Measurement of the thickness of the PDMS films

The PDMS film was dried on a glass substrate and the film was scratched to make the glass surface exposed. The thickness of the film was calculated based on the difference of height between the film surface and glass surface measured using a profilometer (Dektak XT-S, Bruker BioSpin, Kanagawa, Japan, 0.1 nm resolution) with a stylus (12.5 μm radius).

2.4. Adhesive modification of the PDMS films with GA and EDC

Each PDMS film was transferred on a nylon mesh and surface-treated using a plasma cleaner (PDC-001, HARRICK PLASMA, Ithaca, NY, USA) at 7 W for 10 min to introduce hydroxy groups onto the PDMS surface.²³ Each film was then immersed in an APTMS solution (10 wt% in methanol) for 30 min. Subsequently, each PDMS film was washed with methanol and dried. GA (0.65 mol L⁻¹) and EDC (1 eq. vs. GA) in super-dehydrated ethanol were cast on the PDMS film and left for 1 h in a vacuum to obtain the adhesive PDMS (adhesive layer: 40 μm). We also prepared adhesive PDMS with a thinner adhesive layer (4.5 or 10 μm) by using 0.065 or 0.325 mol L⁻¹ GA instead of 0.65 mol L⁻¹ GA.

2.5. Fourier-transform infrared (FT-IR) spectroscopy analysis

To investigate the surface structure of the modified PDMS film, the adhesive layer of the adhesive PDMS film was washed with ethanol to obtain PDMS-GA to expose the modified surface. Subsequently, the FT-IR spectra of the freestanding films of pristine PDMS, PDMS-NH₂ (after reaction with APTMS), and PDMS-GA were obtained using a Compact FT-IR Spectrometer (ALPHA II-P, C295-W/D, Bruker BioSpin, Kanagawa, Japan).

2.6. Electrospray ionization-mass spectrometry (ESI-MS)

Samples were prepared by diluting pure GA, EDC, and the adhesive layer, which comprised the products from GA and EDC, by 10⁶ times with methanol. The ESI-MS spectra of the samples were obtained using an electrospray ionisation time-of-flight mass spectrometry system (microTOF II, Bruker BioSpin, Kanagawa, Japan). Positive mode ESI was used for the measurement of pure GA, EDC and the adhesive layer, and negative mode ESI was used for the measurement of the adhesive layer.

2.7. Proton nuclear magnetic resonance (¹H NMR) spectroscopy

The adhesive layer, which comprised the products from GA and EDC, was concentrated using a rotary evaporator (N-1100, Tokyo Rikakikai Co., Ltd, Japan) to remove the ethanol. Ethyl acetate was added to the products from GA and EDC followed by water. After stirring thoroughly, the aqueous layer was separated from the organic layer. The products in the aqueous and organic layers were concentrated under vacuum, and DMSO-*d*₆ was added to each product to dissolve it. The ¹H NMR spectrum of each product was obtained using a nuclear magnetic resonance apparatus (Agilent 400-MR spectrometer at 400 MHz, Agilent Technologies International Japan, Ltd, Japan).

2.8. Measurement of the thickness of the adhesive layer

Adhesive PDMS (PDMS: 40 μm) was prepared on a glass substrate (560 μm). The thickness of the adhesive PDMS and glass substrate was measured using a digimatic micrometer (227 Series Adjustable Measurement Force Digimatic Micrometers, MISUMI Group Inc., Tokyo, Japan). The thickness of the adhesive layer was calculated by subtracting the sum of the thicknesses of the PDMS and the glass substrate (600 μm) from the value measured using the digimatic micrometer.

2.9. Ultraviolet-visible (UV-Vis) analysis

The UV-Vis spectra of the films of pristine PDMS (40 μm) and adhesive PDMS (PDMS: 40 μm) were obtained using a GENESYS 50 Spectrophotometer (Thermo Fisher Scientific K. K., Tokyo, Japan).

2.10. Tensile tests

Tensile tests were carried out on the films of pristine PDMS (40 μm) and adhesive PDMS (PDMS: 40 μm) using a tensile tester (EZ Test EZ-SX, SHIMADZU Co., Ltd) equipped with a 100 N load cell. Each film was stretched at a strain rate of 10 mm min⁻¹. The elastic modulus was estimated from the slope of the stress-strain curve from 2% to 5% of the strain.

2.11. Tack separation tests with a custom jig

The adhesion performances of the films of the pristine PDMS (40 μm) and adhesive PDMS (PDMS: 40 μm , adhesive layer: 40 μm) were determined by tack separation tests using a tensile tester (EZ Test EZ-SX, SHIMADZU Co., Ltd) equipped with a 100 N load cell. Our method is based on a tack separation test^{11,25-27} and we modified this method to measure the adhesiveness of the adhesive-modified composite film rather than just the adhesive. Each film supported by a square frame (internal area: 20 × 20 mm²) comprising a polystyrene plate was attached to the surface of the skin model (10 × 10 × 5 mm³) wiped with 70% ethanol in advance. After applying a downward force of 0.1 N to the film against the skin model for 5 s, the force was decreased to 0 N and the film was kept to be attached to the skin model for 10 min at room temperature. After 10 min, the frame was lifted (5 mm min⁻¹) until the film was detached from the skin model. From the obtained profile of the separation displacement *versus* the separation force, the adhesion energy of the film was calculated using the following equation:

$$W = \int_0^x f ds$$

where W , x , f , and s represent the adhesion energy, the displacement when the film was detached, the force required to lift the film against the skin model, and the displacement, respectively. The same tests were conducted by using pristine PDMS (9.4 or 133 μm) and adhesive PDMS (PDMS: 9.4 or 133 μm , adhesive layer: 40 μm) and adhesive PDMS (PDMS: 40 μm , adhesive layer: 4.5 or 10 μm).



2.12. Rheology measurements

The viscosity, storage modulus, and loss modulus were measured at 25 °C using a rheometer (Modular Compact Rheometer MCR102e, Anton Paar, Graz, Austria) with a laminator with a diameter of 20 mm and a cone angle of 2°. The flow sweep measurements (viscosity *vs.* shear rate) of the adhesive layer, which was synthesized from GA and EDC in ethanol, pure EDU, and a mixture of pure EGA and EDU (EGA + EDU), were conducted at shear rates of 10^{-2} to 10^2 s⁻¹ at 25 °C. The temperature sweep measurements (viscosity *vs.* temperature) for each sample were conducted in the temperature range 15 to 50 °C at a shear rate of 10 s⁻¹. The frequency sweep measurement (frequency *vs.* storage and loss moduli) of the adhesive layer was conducted in the frequency range from 10^{-1} to 10^2 rad s⁻¹ at a shear strain of 1%.

3. Results and discussion

3.1. Structure analysis of modified PDMS films

We investigated the chemical structure of the film surface using FT-IR spectroscopy. We obtained FT-IR spectra of pristine PDMS, PDMS-NH₂ (after reaction with APTMS), and PDMS-GA (after reaction with GA and EDC, and washing the adhesive layer with ethanol) to determine the chemical structure of the film surface. The FT-IR spectra confirmed the introduction of amino groups by the silane coupling reaction, and subsequently the introduction of gallol groups by reaction with GA and EDC on the PDMS film surface were achieved (Fig. S2, ESI[†]). We attributed the bands at 3200–3500 and 1570 cm⁻¹ to N–H stretching vibrations and N–H bending vibrations,²⁸ respectively, and confirmed the successful introduction of amino groups onto the PDMS films. The bands at 3100–3300 and 1610 cm⁻¹ were attributed to the O–H stretching vibrations of gallol groups and the C=O stretching vibrations of amide bonds,¹⁴ respectively, and confirmed the successful introduction of GA on the surfaces of the PDMS films.

3.2. Structure analysis of the adhesive layer

We investigated the chemical structures of the compounds in the adhesive layer using ESI-MS and ¹H NMR spectroscopy. The thickness and transparency of the adhesive layer were, respectively, evaluated using a digimatic micrometer and UV-Vis analysis. The adhesive layers on the films were analyzed by ESI-MS and ¹H NMR spectroscopy to identify the specific synthesized compounds. The ESI-MS spectra of pure GA, pure EDC, and the compounds in the adhesive layer dissolved in methanol were measured (Fig. S3, ESI[†]). The peak (*m/z* = 193, 363) in GA solution is, respectively, attributed to GA monomer and GA dimer, which gained a sodium ion (Fig. S3a, ESI[†]), and the peak (*m/z* = 156) in an EDC solution is attributed to protonated EDC (Fig. S3b, ESI[†]). The peak derived from the cation (*m/z* = 174) which was observed in positive mode ESI-MS measurement of the adhesive layer was attributed to the protonated EDU (Fig. S3c, ESI[†]), and the peak derived from the anion (*m/z* = 197) which was observed in negative mode ESI-MS

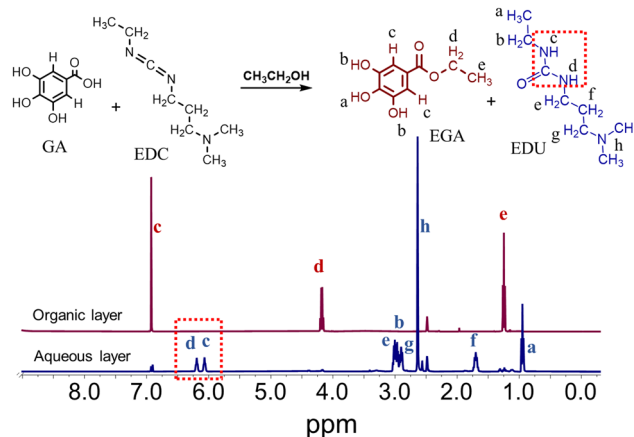


Fig. 2 Reaction scheme in the adhesive layer and the results of ¹H NMR spectroscopy of the compounds in the adhesive layer after separation by extraction (¹H NMR = proton nuclear magnetic resonance; EGA = ethyl gallate; EDU = 1-ethyl-3-(3-dimethylaminopropyl)urea).

measurement of the adhesive layer was attributed to EGA, which lost a proton (Fig. S3d, ESI[†]). Considering the peaks of GA and EDC disappeared in the adhesive layer, we demonstrated GA and EDC are mostly reacted and converted to EDU and EGA. Next, the compounds in the adhesive layer, *i.e.* the products from GA and EDC, were separated into aqueous and organic layers by extraction, and the ¹H NMR spectrum of each compound was obtained. The obtained spectra shown in Fig. 2 reveal that EGA and EDU were present in the organic and aqueous layers, respectively. Therefore, the adhesive layer comprised EGA and EDU, which were synthesized from GA and EDC in ethanol. In summary, the adhesive PDMS consisted of a PDMS layer, the surface of which had been modified with gallol, and an adhesive layer containing EGA and EDU. It was also shown that the adhesive layer had 40 ± 4.3 μm thickness and the same level of transparency as the pristine PDMS measured using a digimatic micrometer and UV-Vis measurement (Fig. S4, ESI[†]). Therefore, the modification method would be applicable to biomedical devices with light-emitting systems.^{4,11}

3.3. Mechanical properties of adhesive PDMS

To confirm the stretchability of the adhesive based on abundant non-covalent bonds, the mechanical properties of the adhesive PDMS were evaluated by a tensile test. We used a pristine PDMS film (40 μm) and an adhesive PDMS film (80 μm (PDMS: 40 μm, adhesive layer: 40 μm)) for the measurement on a tensile tester. The films were stretched at a strain rate of 10 mm min⁻¹, and their elastic moduli were estimated from their stress-strain curves. The results are shown in Fig. 3a and the calculated elastic moduli are shown in Fig. 3b. The elastic modulus of the adhesive PDMS film (1.01 MPa) was lower than that of the pristine PDMS film (1.79 MPa). The observed reduction in stress for the adhesive PDMS compared to the pristine PDMS can be attributed to accounting for the adhesive layer's thickness in the stress calculation, despite both films being subjected to the same tensile force (Fig. S5, ESI[†]). This result means the elastic



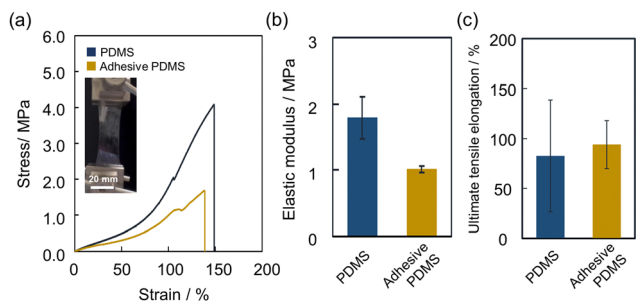


Fig. 3 (a) Results of tensile tests on a PDMS film and an adhesive PDMS film, and the image of stretching adhesive PDMS. (b) Calculated elastic moduli and (c) ultimate tensile elongation values of each PDMS film (PDMS = polydimethylsiloxane).

modulus of the PDMS substrate is dominant over the adhesive layer. The ultimate tensile elongation of the adhesive PDMS film (94.0%) was as large as that of the pristine film (82.6%) (Fig. 3c). Therefore, we conclude that the adhesive modification does not impair the original mechanical properties of the PDMS film.

3.4. Tack separation test of adhesive PDMS with a custom jig

Next, we conducted a tack separation test with a custom jig on a skin model to evaluate the adhesiveness of the adhesive-modified PDMS. First, the skin model (surface area: $10 \times 10 \text{ mm}^2$) was prepared on a plastic plate and pristine PDMS film ($40 \mu\text{m}$) or an adhesive PDMS film (PDMS: $40 \mu\text{m}$, adhesive layer: $40 \mu\text{m}$) was prepared and attached to the skin model for 10 min (Fig. 4a). Next, the PDMS film supported by a square frame was lifted until it detached from the skin model, and a profile of the separation displacement *versus* the separation force was obtained (Fig. 4b). The adhesion energy of each sample was obtained by calculating the integral value of the profile (Fig. 4c). The average adhesion energy value of the adhesive PDMS film was $27.4 \pm 3.40 \mu\text{J cm}^{-2}$,

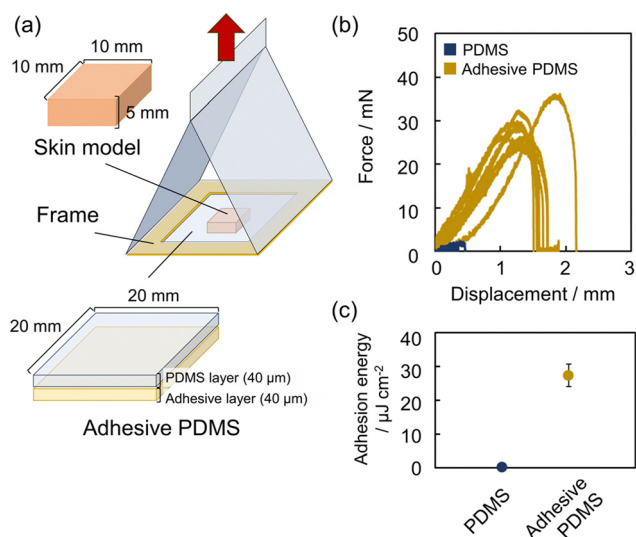


Fig. 4 (a) The schematic image of the tack separation test of adhesive PDMS with a custom jig. (b) Results of the tack separation test and (c) calculated adhesion energy values of pristine PDMS and adhesive PDMS (PDMS = polydimethylsiloxane).

which was 130 times larger than that of pristine PDMS films ($(2.11 \pm 1.30) \times 10^{-1} \mu\text{J cm}^{-2}$). The order of the adhesion energy value was comparable to that of the value obtained using a previous method using polydopamine to enhance the adhesiveness of PDMS to chicken muscle¹¹. This phenomenon can be attributed to the energy dissipation of the compounds in the adhesive layer, which facilitated an increase in the film's absolute adhesion area by effectively infiltrating the surface of the adherents. We prepared adhesive PDMS with different thicknesses of the adhesive layer (4.5 ± 3.0 and $10 \pm 3.1 \mu\text{m}$) by changing the concentration of the precursor solution of GA and EDC and a tack separation test of them was conducted (Fig. S6, ESI†). We found that the adhesion energy of adhesive PDMS increased as the thickness of the adhesive layer became thicker, and it hit the limit at the point around $10 \mu\text{m}$ thickness. This indicates that a thicker adhesive layer is required for efficient energy dissipation for strong adhesiveness but a too thick adhesive layer cannot contribute to strong adhesiveness because of cohesive failure.²⁹ We also prepared adhesive PDMS with different thicknesses of the PDMS layer (9.4 ± 5.0 , 40 ± 3.8 , and $133 \pm 5.9 \mu\text{m}$) and confirmed that our adhesive modification can improve the adhesiveness of the PDMS films regardless of the thickness and mechanical properties (Fig. S7, ESI†). The adhesion energy as well as the data variation of adhesive PDMS ($133 \mu\text{m}$) was larger because the peeling behavior was considered inconsistent throughout the experiment. When using a thicker film, a larger force is required to pull up and deform the margin of the PDMS film in the phase before the peak of maximum force³⁰ and the adhesion energy was calculated as a larger value. The increased data variation observed in the thicker film is attributed to the slight tilt of the adhesive PDMS film when attached to the skin model. This tilt causes a heterogeneous distribution of vertical pulling force. Although the thinner film is more stretchable and can remain attached on the skin model even if the pulling force is different on both sides of the tilt, the thicker film cannot remain attached and easily peels off, resulting in larger data variations due to the tilt of the film.

3.5. Viscoelasticity of the adhesive layer and the mechanism of adhesiveness

Next, we measured the viscoelasticity of the adhesive layer (Fig. 5a) using a rheometer to elucidate the mechanism underlying the adhesiveness of the film. For example, we sought to identify the chemical bonds formed by the functional groups that were required for proper cohesion performance and the energy dissipation required for a large adhesion area between the PDMS and the adherents. Adhesive layers comprising EGA and EDU, pure EDU, and a mixture of pure EGA and EDU (EGA + EDU) were prepared on a glass plate by each and investigated using a rheometer. The flow sweep measurements (viscosity *versus* shear rate) and temperature sweep measurements (viscosity *versus* temperature) are shown in Fig. 5b and Fig. S8 (ESI†), respectively. It is supposed that the larger decrease in the viscosity of EDU under an increasing shear rate and temperature was attributable to the dissociation of hydrogen bonds.^{18,31} However, it is suggested that the smaller decrease in the viscosity of the adhesive layer and the EGA + EDU layer



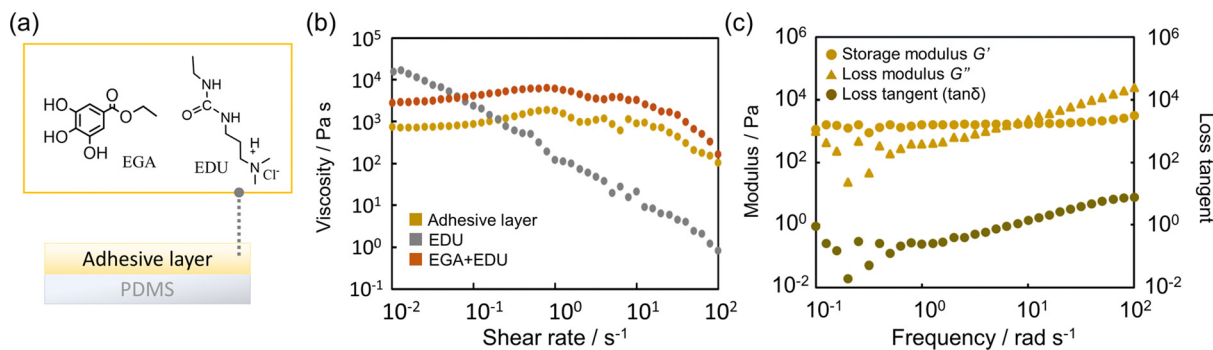


Fig. 5 (a) The compounds in the adhesive layer. (b) Flow sweep measurements (viscosity vs. shear rate) of the adhesive layer, EDU, and EGA + EDU and (c) frequency sweep measurements (frequency vs. storage (G') and loss (G'') moduli and loss tangent) of the adhesive layer (EDU = 1-ethyl-3-(3-dimethylaminopropyl)urea; EGA = ethyl gallate).

compared with the EDU layer was due to stronger non-covalent bonds than hydrogen bonds. These bonds include electrostatic interactions between the cationic amino groups in the EDU and the anionic phenol groups in EGA or hydrophobic interactions, which contributed to the adhesiveness. However, further research on the relationships between functional groups and adhesiveness is required. Next, frequency sweep measurements (frequency vs. storage and loss moduli) revealed that G' was larger than G'' at low frequencies, but a gel-sol transition occurred at 7.09 rad s^{-1} , after which G'' exceeded G' ($\tan \delta > 1$) at higher frequencies (Fig. 5c). Together with the adhesion mechanism, these phenomena suggest that the adhesive layer on the PDMS had high cohesive strength at smaller frequencies and high peel strength at higher frequencies.³² It is possible that a loss tangent close to 1 facilitated the proper diffusion of the compounds that contributed to a larger adhesion area between the skin model and the PDMS, as well as strong adhesiveness.^{33,34} This phenomenon could be attributed to energy dissipation by weak non-covalent bonds in the adhesive layer.³⁵ It has been reported that a larger cohesion and a larger G' are more important factors for stronger adhesiveness than a large G'' .³⁶ Therefore, we conclude that the adhesive PDMS had stronger adhesiveness at lower frequencies and under slow retraction. Furthermore, the larger G'' at higher frequencies meant that the adhesive layer could be easily detached by fast retraction. Therefore, it could be described as a pressure-sensitive adhesive. In short, we found that our adhesive layer applies to elastomer films in terms of balancing cohesion and molecular diffusion due to energy dissipation for effective adhesiveness and detachment.

4. Conclusions

In conclusion, we demonstrated a facile method for fabricating an adhesive layer comprising the low-molecular-weight compounds; EGA and EDU for maintaining the original mechanical properties of PDMS. Our results suggested that abundant hydrogen bonds as well as many other kinds of non-covalent bonds among the gallol, urea, and tertiary amino groups contributed to both energy dissipation and strong cohesion for effective adhesiveness. To further strengthen the adhesive performance, the conjugation of low-molecular-weight compounds to

the polymer backbone to improve cohesion for water resistance is also required. This versatile method could be applied not only to amine-bearing substrates but to a variety of elastomers to which amines can be introduced.³⁷ Our method for modifying adhesives onto elastomer films does not sacrifice stretchability and could be applied to thicker films, despite thinner films being preferred to obtain larger adhesiveness.^{11,38} This method expands the possibility of the application of many kinds of thin film devices for electroencephalographic analysis,² electromyographic analysis,³ measuring electrical signals in plants,⁶ and so on. The present research provides guidelines for the molecular design and modification of adhesives for biomedical devices and will contribute to their long-term use.

Data availability

The data that support the findings of this study are available from the corresponding author, Toshinori Fujie, upon reasonable request.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 A. Imai, S. Takahashi, S. Furubayashi, Y. Mizuno, M. Sonoda, T. Miyazaki, E. Miyashita and T. Fujie, *Adv. Mater. Technol.*, 2023, **8**(21), 2300300.
- 2 L. M. Ferrari, U. Ismailov, J.-M. Badier, F. Greco and E. Ismailova, *Npj Flex. Electron.*, 2020, **4**, 1–9.
- 3 K. Yamagishi, T. Nakanishi, S. Mihara, M. Azuma, S. Takeoka, K. Kanosue, T. Nagami and T. Fujie, *NPG Asia Mater.*, 2019, **11**, 1–13.
- 4 Z. Chen, R. T. Yin, S. N. Obaid, J. Tian, S. W. Chen, A. N. Miniovich, N. Boyajian, I. R. Efimov and L. Lu, *Adv. Mater. Technol.*, 2020, **5**(8), 2000322.
- 5 L. Lan, X. Le, H. Dong, J. Xie, Y. Ying and J. Ping, *Biosens. Bioelectron.*, 2020, **165**, 112360.
- 6 F. Meder, S. Saar, S. Taccola, C. Filippeschi, V. Mattoli and B. Mazzolai, *Adv. Mater. Technol.*, 2021, **6**, 2001182.
- 7 J. Deng, H. Yuk, J. Wu, C. E. Varela, X. Chen, E. T. Roche, C. F. Guo and X. Zhao, *Nat. Mater.*, 2021, **20**, 229–236.
- 8 Q. Yang, T. Wei, R. T. Yin, M. Wu, Y. Xu, J. Koo, Y. S. Choi, Z. Xie, S. W. Chen, I. Kandela, S. Yao, Y. Deng, R. Avila, T.-L. Liu, W. Bai, Y. Yang, M. Han, Q. Zhang, C. R. Haney, K. Benjamin Lee, K. Aras, T. Wang, M.-H. Seo, H. Luan, S. M. Lee, A. Brikha, N. Ghoreishi-Haack, L. Tran, I. Stepien, F. Aird, E. A. Waters, X. Yu, A. Banks, G. D. Trachiotis, J. M. Torkelson, Y. Huang, Y. Kozorovitskiy, I. R. Efimov and J. A. Rogers, *Nat. Mater.*, 2021, **20**, 1559–1570.
- 9 M. Takuma, H. Fujita, N. Zushi, H. Nagano, R. Azuma, T. Kiyosawa and T. Fujie, *Biomater. Sci.*, 2024, **12**, 3401–3410.
- 10 M. Saito, E. Kanai, H. Fujita, T. Aso, N. Matsutani and T. Fujie, *Adv. Funct. Mater.*, 2021, **31**, 2102444.
- 11 K. Yamagishi, I. Kirino, I. Takahashi, H. Amano, S. Takeoka, Y. Morimoto and T. Fujie, *Nat. Biomed. Eng.*, 2019, **3**, 27–36.
- 12 H. Lee, S. M. Dellatore, W. M. Miller and P. B. Messersmith, *Science*, 2007, **318**, 426–430.
- 13 M. A. Gwak, S. J. Lee, D. Lee, S. A. Park and W. H. Park, *Int. J. Biol. Macromol.*, 2023, **227**, 493–504.
- 14 S. N. Ramirez-Barron, S. Sanchez-Valdes, R. Betancourt, C. A. Gallardo, B. Puente-Urbina, O. S. Rodriguez-Fernández, M. G. Carneiro-da Cunha, M. T. dos Santos-Correia and Z. V. Sanchez-Martinez, *Int. J. Adhes. Adhes.*, 2021, **104**, 102749.
- 15 Q. Yang, L. Tang, C. Guo, F. Deng, H. Wu, L. Chen, L. Huang, P. Lu, C. Ding, Y. Ni and M. Zhang, *Chem. Eng. J.*, 2021, **417**, 127962.
- 16 J. Zhang, W. Wang, Y. Zhang, Q. Wei, F. Han, S. Dong, D. Liu and S. Zhang, *Nat. Commun.*, 2022, **13**, 5214.
- 17 X. Li, Y. Deng, J. Lai, G. Zhao and S. Dong, *J. Am. Chem. Soc.*, 2020, **142**, 5371–5379.
- 18 Q. Zhang, C.-Y. Shi, D.-H. Qu, Y.-T. Long, B. L. Feringa and H. Tian, *Sci. Adv.*, 2018, **4**, eaat8192.
- 19 C.-Y. Shi, Q. Zhang, H. Tian and D.-H. Qu, *SmartMat*, 2020, **1**(1), e1012.
- 20 Y. Liang, K. Wang, J. Li, Y. Zhang, J. Liu, K. Zhang, Y. Cui, M. Wang and C.-S. Liu, *Mater. Horiz.*, 2022, **9**, 1700–1707.
- 21 K. Zhan, C. Kim, K. Sung, H. Ejima and N. Yoshie, *Biomacromolecules*, 2017, **18**, 2959–2966.
- 22 G. P. Maier, M. V. Rapp, J. H. Waite, J. N. Israelachvili and A. Butler, *Science*, 2015, **349**, 628–632.
- 23 D. Bodas and C. Khan-Malek, *Sens. Actuators, B*, 2007, **123**, 368–373.
- 24 C.-C. Wu, C.-Y. Yuan and S.-J. Ding, *Surf. Coat. Technol.*, 2011, **205**, 3182–3189.
- 25 D. Astm, ASTM D2979.
- 26 B. S. Terry, A. C. Passernig, M. L. Hill, J. A. Schoen and M. E. Rentschler, *J. Mech. Behav. Biomed. Mater.*, 2012, **15**, 24–32.
- 27 T. Horii, K. Yamashita, M. Ito, K. Okada and T. Fujie, *NPG Asia Mater.*, 2024, **16**, 33.
- 28 H. Kim, H.-G. Kim, S. Kim and S. S. Kim, *J. Memb. Sci.*, 2009, **344**, 211–218.
- 29 L. F. M. da Silva, T. N. S. S. Rodrigues, M. A. V. Figueiredo, M. F. S. F. de Moura and J. A. G. Chousal, *J. Adhesion*, 2006, **82**, 1091–1115.
- 30 J. P. Phillips, X. Deng, R. R. Stephen, E. L. Fortenberry, M. L. Todd, D. M. McClusky, S. Stevenson, R. Misra, S. Morgan and T. E. Long, *Polymer*, 2007, **48**, 6773–6781.
- 31 J. Kajtna, B. Alič, M. Krajnc and U. Šebenik, *Int. J. Adhes. Adhes.*, 2014, **49**, 103–108.
- 32 F. Mazzeo, TA Instruments report RH082.
- 33 E. P. Chang, *J. Adhes.*, 1991, **34**, 189–200.
- 34 Y.-J. Wang, Y. He, S. Y. Zheng, Z. Xu, J. Li, Y. Zhao, L. Chen and W. Liu, *Adv. Funct. Mater.*, 2021, **31**, 2104296.
- 35 M. G. Mazzotta, A. A. Putnam, M. A. North and J. J. Wilker, *J. Am. Chem. Soc.*, 2020, **142**, 4762–4768.
- 36 M. Seong, S. Kondaveeti, G. Choi, S. Kim, J. Kim, M. Kang and H. E. Jeong, *ACS Appl. Mater. Interfaces*, 2023, **15**, 11042–11052.
- 37 J. Song, B. Winkeljann and O. Lieleg, *Adv. Mater. Interfaces*, 2020, **7**, 2000850.
- 38 T. Fujie, N. Matsutani, M. Kinoshita, Y. Okamura, A. Saito and S. Takeoka, *Adv. Funct. Mater.*, 2009, **19**, 2560–2568.

