





**Figure 1.** Schematic representation of the tack and peel measurement setup. This graphic depicts the schematic setup for tack tests (**a**) and peel tests (**b**).



**Figure 2.** Exemplary peel measurements performed with SSA patches attached to excised murine tympanic membranes. Overview about a typical measurement performed to analyze peel strength of patches on the tympanic membrane (**A**). The measurement consists of three different steps: (1) In the attachment phase, the SSA patch and control materials are applied to the tympanic membrane. In this phase, the contact between patch and tympanic membrane is not uniformly reached. (2) In the adhesion phase, the patches are uniformly adhered to the tympanic membrane by the application of slight pressure all over the entire surface. This phase is separately shown in **B**. (3) In the removal phase, the patch is peeled off the tympanic membrane while trying to maintain a constant speed. This phase is separately shown in **C**. Here, the removal phase is represented by the data points between 92 s and 94 s.



**Figure 3.** Cellular response to the protein coating of the elastomer films after seven days of storage in ddH<sub>2</sub>O before cellular seeding. L929 murine fibroblasts were cultured on SSA films after storage for seven days in water. Before storage, the films were incubated with albumin (**A1**, **A1.1**), fibronectin (**B1**, **B1.1**), and fibrinogen (**C1**, **C1.1**). Controls were non-treated (pristine) films (**D1**, **D1.1**). Phase contrast pictures were taken after a culture period of 48 h (**A1**, **B1**, **C1**, **D1**). Additionally, after fixation, immunocytochemistry was performed with an anti-phospho-FAK<sup>Tyr397</sup> antibody to visualize focal adhesion contacts (red). Additionally, the actin cytoskeleton (green) and cellular nuclei (blue) were visualized after staining with Alexa 488 conjugated phalloidin and Hoechst Dye 33342 (A1.1, B1.1, C1.1, and D1.1). Number of independent experiments: n = 3 for the fluorescence analysis; n=4 for the determination of the cellular area. Scale bar in A1, B1, C1, D1 = 100 µm; scale bar in A1.1, B1.1, C1.1, D1.1 = 25 µm.

## $\label{eq:stable} \begin{array}{l} \textbf{Table S1.} \mbox{ Corresponding table to Figure 4.} \\ test parameters: single SSA MG 7-9800 layers on glass \\ preload stress 13 \pm 5 kPa; approach velocity: 30 \mbox{ } \mu m/s; detachment velocity: \\ 10 \mbox{ } \mu m/s; hold time (in contact): 1s \end{array}$

		glass smooth	1		glass rough	
a]	water	fibronectin	untreated	water	fibronectin	untreated
s [kP	$22.9 \pm 3.2$	$23.8 \pm 3.7$	$24.9\pm6$	$29.9 \pm 4.4$	$29.6 \pm 3.1$	31.8 ± 5
ll-off stres		fibrinogen 23.6 ± 6.2			fibrinogen 29.76 ± 3.1	
nd		albumin			albumin	
	1 1 1	$27.1\pm6.1$			$32.4 \pm 4.5$	
-		alass smooth	1		olass rough	
/m <sup>2</sup> ]		glass smooth	1 www.two.at.ad	sustau	glass rough	www.washad
mJ/m²]	water	glass smooth fibronectin	untreated	water	glass rough fibronectin	untreated
n [m]/m²]	<i>water</i> 932 ± 256	<b>glass smooth</b> fibronectin 1106 ± 458	n untreated 926 ± 226	<i>water</i> 1159 ± 191	<b>glass rough</b> <i>fibronectin</i> 1655 ± 681	<i>untreated</i> 1510 ± 500
of separation [mJ/m²]	water 932 ± 256	glass smooth fibronectin 1106 ± 458 fibrinogen 948 ± 243	n untreated 926 ± 226	water 1159 ± 191	<b>glass rough</b> fibronectin 1655 ± 681 fibrinogen 1231 ± 250	untreated 1510 ± 500
ork of separation [m]/m²]	water 932 ± 256	glass smooth fibronectin 1106 ± 458 fibrinogen 948 ± 243 albumin	untreated 926 ± 226	water 1159 ± 191	glass rough fibronectin 1655 ± 681 fibrinogen 1231 ± 250 albumin	untreated 1510 ± 500

Table S2. Corresponding table to Figure 5.test parameters: composite films (SSA MG 7-9800 layers and Sylgard 184) on PET foilpreload stress 13 ± 5 kPa; approach velocity: 30  $\mu$ m/s; detachment velocity: 10  $\mu$ m/s;hold time (in contact): 1s

glass roughuntreatedglass roughwaterfibronectinuntreated $37.4 \pm 7.8$ $40 \pm 5.5$ $43 \pm 8$ $40 \pm 5.5$ $43 \pm 8$ $1320 \pm 512$ $1438 \pm 483$ $2870 \pm 512$	e <i>ted</i> 2340
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Table S3. Corresponding table to Figure 6. test parameters: composite films (SSA MG 7-9800 layers and Sylgard 184) initial peel angle: ≈ 90°; detachment velocity: 10 µm/s

		glass rough		[N/m
ted .01	untreate 358 ± 10	fibronectin 286 ± 151	water 373 ± 141	el strength
				peel st

## Table S4. Corresponding table to Figure 7.

test parameters: composite films (SSA MG 7-9800 layers and Sylgard 184), silicone strip, single layer Sylgard 184. The films were manually applied to the tympanic membrane and peeled off using tweezers.

