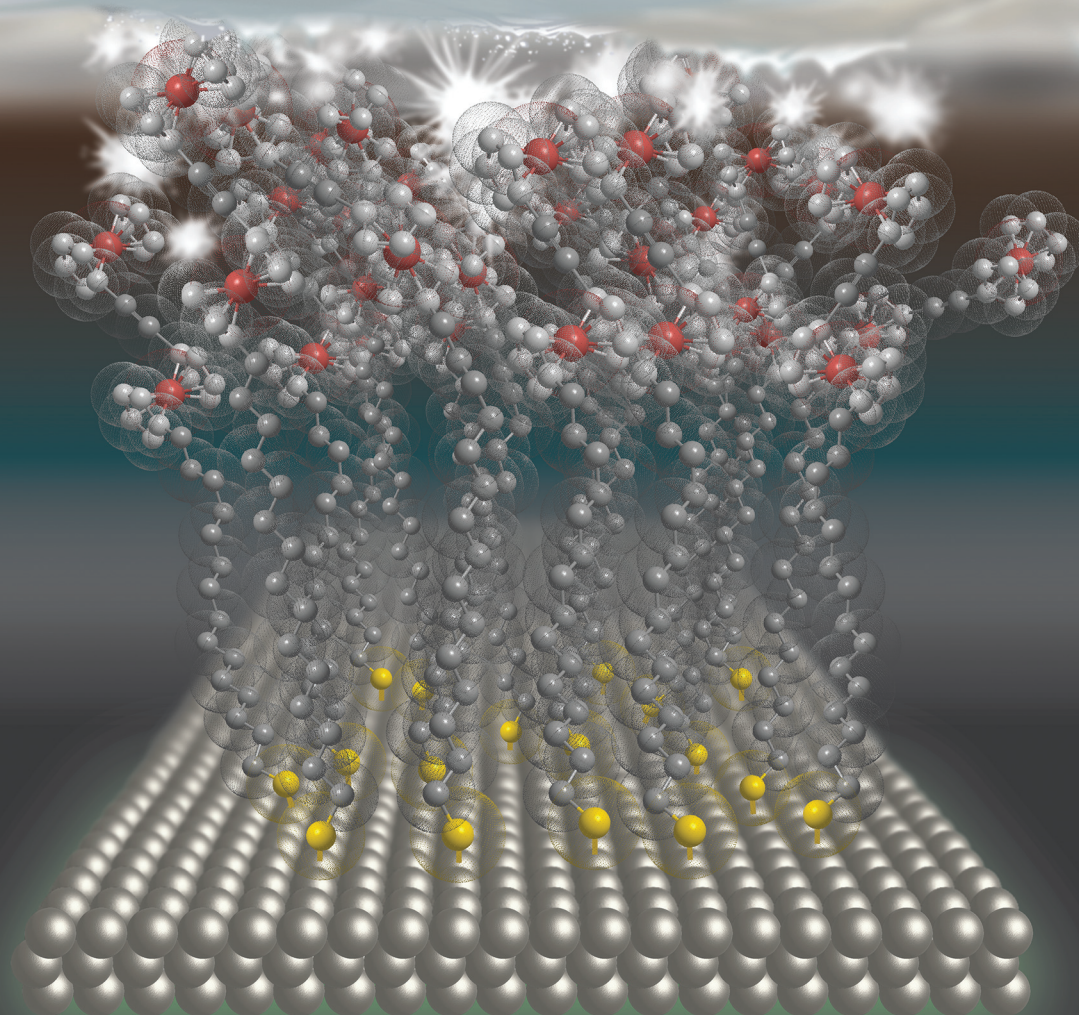


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Supramolecular tunnelling junctions with robust high
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Supramolecular tunnelling junctions with robust high rectification based on assembly effects†

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The performance of large-area molecular diodes can in rare cases approach the lower limit of commercial semiconductor devices but predictive structure–property design remains difficult as the rectification ratio (R) achieved by self-assembled monolayer (SAM) based diodes depends on several intertwined parameters. This paper describes a systematic approach to achieve high rectification in bisferrocenyl-based molecular diodes, $\text{HSC}_n\text{Fc}-\text{C}\equiv\text{C}-\text{Fc}$ ($n = 9-15$) immobilised on metal surfaces (Ag, Au and Pt). Experiments supported by molecular dynamics simulations show that the molecular length and bottom electrode influence the SAM packing, which affects the breakdown voltage (V_{BD}), the associated maximum R (R_{max}), and the bias at which the R_{max} is achieved ($V_{\text{sat},R}$). From the electrical characterisation of the most stable $\text{Pt}-\text{SC}_n\text{Fc}-\text{C}\equiv\text{C}-\text{Fc}/\text{GaO}_x/\text{EGaIn}$ junctions, we found that V_{BD} , $V_{\text{sat},R}$ and R_{max} all scale linearly with the spacer length of C_n , and that R_{max} for all the SAMs consistently exceeds the “Landauer limit” of 10^3 . Our data shows that the robust switching of $\text{M}-\text{SC}_n\text{Fc}-\text{C}\equiv\text{C}-\text{Fc}/\text{GaO}_x/\text{EGaIn}$ junctions is the result of the combined optimisation of parameters involving the molecular structure, the type of metal substrate, and the applied operating conditions (bias window), to create stable and high-performance junctions.

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Introduction

Molecular diodes are attractive building blocks for nanoscale electronic circuits,^{1–5} and their development has been a focal point since the landmark paper by Aviram and Ratner proposing the first molecular diode in 1974.⁶ The performance of a

diode is usually expressed in terms of its rectification ratio (R), which defines how much electrical current density (J , in A cm^{-2}) traverses the junction in the ON vs. OFF states. R is simply the ratio of $|J_{\text{on}}|/|J_{\text{off}}|$, i.e., $R = |J(-V)|/|J(V)|$, where V is the applied voltage in V, and R is the ratio of absolute J at forward bias to that at reverse bias. Ideal diodes would allow all current to pass through the diode in the ON state, but not in the OFF state. However, this ideal behaviour is not reached as a small amount of current (leakage current) always flows also in the OFF state. In molecular diodes, this leakage current is dictated by many factors, resulting in smaller values for R than ideal cases. Macroscopic semiconductor-based diodes operate at R of $\geq 10^5$, while molecular diodes have been generally limited to orders of magnitude smaller values of R because of the molecular resistance in the ON state and large leakage currents (due to defects or molecular disorder) in the OFF state.^{7–16} Recently, exceptionally well-performing molecular diodes have been reported with R of several thousand based on rigid biferrocenes with interlocked cyclopentadienyl rings in SAM based junctions or nonadiyne in single-molecule junctions,^{17,18} and even of several tens of thousands in switchable junctions, which operate with a different mechanism via directional counterion migration in SAMs of viologens.¹⁹

Our previous studies showed that the performance of SAM-based diodes depends on the choice of the bottom electrode,²⁰ electrode–molecule anchoring group,²¹ molecular structure,^{12,22}

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[†]Electronic supplementary information (ESI) available: Synthetic procedures, analytical data of the molecules, details of the SAM formation and characterisation data (CV, XPS, UPS, NEXAFS), details of the MD calculations, electrical characterisation data and details of the fitting. See DOI: <https://doi.org/10.1039/d4nr01514b>
[‡]These authors contributed equally to this work.


and tilt angle of the active group,⁹ which all impact the supra-molecular packing of the SAM and its ability to effectively transport current in the ON state while blocking leakage currents in the OFF state.²³ Even seemingly subtle changes in the molecular structure can have profound macroscopic implications as nicely demonstrated by well-known odd–even effects.^{24,25} SAMs of molecules on Ag with odd numbers of methylene units (n_c = odd) pack better than the ones with even numbers (n_c = even), which results in odd–even effects for surface coverages, electrical behaviours and the SAM packing properties predicted from molecular dynamics (MD) simulations. For example, the odd–even effects in packing energies of SAMs derived from *n*-alkanethiolates (where an odd or even number of CH₂ moieties determines the orientation of the functional terminal group) lead to odd–even effects in exchange kinetics during SAM formation,²⁶ surface dipole and work-function,²⁷ dielectric constant²⁸ and optical properties.²⁹ Such odd–even effects also result in current oscillations in molecular junctions^{22,30–32} and impact molecular diode performance.³⁰

The mechanism of rectification of molecular diodes derived from S(CH₂)₁₁Fc (Fc is ferrocenyl),^{10,22,33–41} or derivatives such as S(CH₂)₁₁Fc₂,¹¹ is well understood. At positive bias, the highest occupied molecular orbital (HOMO) does not participate in charge transport resulting in low conductivity, while at negative bias, the HOMO falls in the conduction window leading to high conductivity. We showed recently that molecular junctions of Pt–SC₁₅Fc–C≡C–Fc//GaO_x/EGaIn (EGaIn stands for eutectic alloy of gallium and indium) can rectify with $R = 6.3 \times 10^5$ by an electrostatic mechanism that maximises the number of molecules contributing to charge transport in only the forward direction of applied bias when the diodes are ON,⁴² allowing them to bypass the Landauer limit for single-level dominated charge transport.^{42–44} In these diodes, the SAM structure changes with applied electric field (or applied bias) due to the electrostatic forces between the SAM and the top EGaIn electrode. This effect led to an increase in the number of conducting molecules with increasing applied bias for only one bias polarity, boosting the diode performance. This example highlights that molecular diode performance can be bias dependent.

To rationally design molecular diodes that are stable and operate with high R for potential future applications in nanoscale electronics, it is important to understand the strengths and weaknesses of such systems by systematically taking both electronic and (supra)molecular structure of the SAMs into consideration. Here, we go beyond that first SC₁₅Fc–C≡C–Fc diode demonstration. Being aware that very long molecules become highly resistive, and very short molecules give disordered SAMs (resulting in large leakage currents at reverse bias when the diodes should block the current), we identified an optimal range of molecular lengths to obtain well-performing molecular diodes by systematically exploring the alkyl chain length-dependence of the SC_{*n*}Fc–C≡C–Fc junctions with $n_c = 9–15$. Furthermore, the choice of the bottom electrode, either Ag, Au or Pt, results in changes in energy level align-

ments, differentiating electronic effects from molecular effects in the M–SC_{*n*}Fc–C≡C–Fc//GaO_x/EGaIn junctions, but also affects the stability of the junctions. We identified Pt as the bottom electrode of choice and we determined the breakdown voltage (V_{BD}) and subsequently measured those junctions at the maximum possible bias range. We show that R depends on the applied bias range and that the maximal possible bias range depends on the length of the alkyl chains and type of electrode material. The larger V_{BD} allows us to increase the operating bias window, leading to an improved value of R . This work shows that the design of optimally performing junctions is complicated by several intertwined factors that all need to be optimised, illustrating that it is still challenging to rationally design molecular junctions.

Results and discussion

The molecular tunnelling junctions

The junctions consist of thiolate–metal contacts with the bottom electrode and non-covalent interactions with the EGaIn top electrode (Fig. 1). The variations of the alkyl chain length and the type of metal substrate (Fig. 1b) allow us to probe odd–even effects of the SAM, the corresponding electrical characteristics, the evolution in the trend of charge transport rates, and the corresponding rectification ratios, as well as characterising the stability of the junctions as measured by the breakdown voltage.

The HSC_{*n*}Fc–C≡C–Fc precursors were synthesised and characterised (see ESI†). The melting points of the HSC_{*n*}Fc–C≡C–Fc molecules show an odd–even effect with a similar trend as observed for the corresponding monoferrocenyl thiols (HSC_{*n*}Fc),⁴⁵ *i.e.*, the odd numbered derivatives melt at lower temperature than the even numbered ones highlighting an odd–even effect in the packing structure. Such odd–even effects are well known for *n*-alkanes, in the solid and the liquid states.^{46,47} Since the non-templated assembly already shows an odd–even effect, naturally odd–even effects are also expected in the monolayer packing structure. Fig. 1 illustrates the different SC_{*n*}Fc–C≡C–Fc SAMs with $n_c = 9–15$ on metallic substrates M (M = Au, Ag, or Pt). The terminal Fc–C≡C–Fc moiety follows an odd–even effect in the tilt angle of the head group (α) with respect to the surface normal. This odd–even effect originates from the number of methylene units in the alkyl chain (n_c) and the fixed M–C–S angle, as illustrated in Fig. 1a.²⁴ However, in SAMs with large head groups, the SAM-packing structure may be driven by head group – head group interactions.^{18,42,48}

Surface characterisations of the SAMs

We prepared stable SAMs on Ag, Au, and Pt, and characterised them using cyclic voltammetry (CV), X-ray photoelectron spectroscopy (XPS), ultra-violet photoelectron spectroscopy (UPS), and near-edge X-ray absorption fine spectroscopy (NEXAFS). We have previously characterised M–SC₁₅Fc–C≡C–Fc by these techniques.⁴² Here, we provide XPS, UPS, and NEXAFS datasets



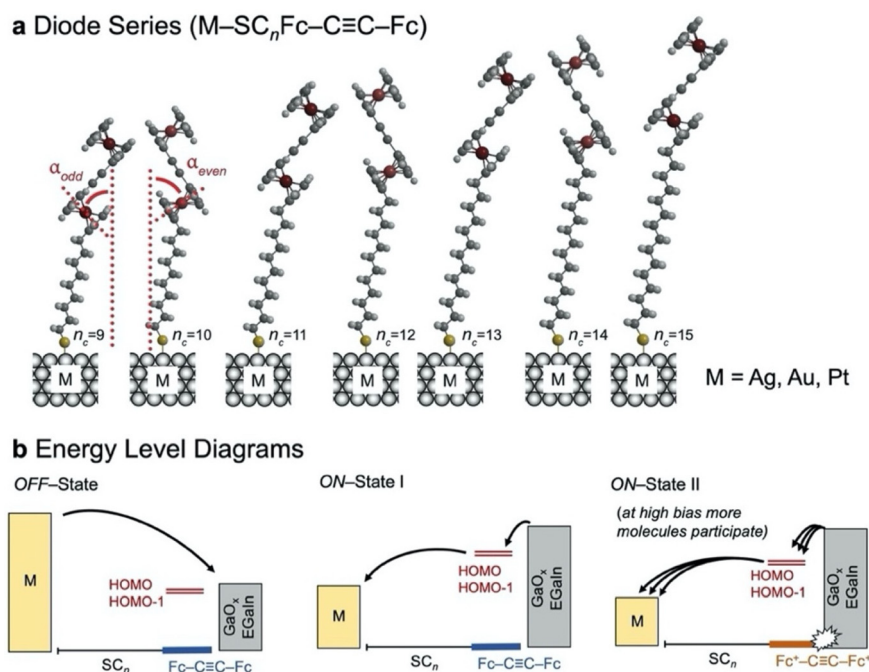


Fig. 1 (a) Representation of the series of $\text{HSC}_n\text{Fc-C}\equiv\text{C-Fc}$ ($n_c = 9-15$) immobilised on a noble metal surface (M). The odd and even numbered diodes differ in orientation and tilt angle (α_{odd} and α_{even}) of the $\text{Fc-C}\equiv\text{C-Fc}$ head group. (b) Energy level diagrams for the diode in the OFF-state, the ON-state I and the ON-state II at which the current flow is enhanced due to Coulomb interactions between the oxidised positively charged $\text{Fc}^+-\text{C}\equiv\text{C-Fc}^+$ and the negatively charged EGaIn-electrode increasing the number of molecules involved in charge transport as indicated by the multiple arrows. The $\text{Fc-C}\equiv\text{C-Fc}$ head group provides two near-degenerate levels HOMO and HOMO-1 for charge transport.

for two further derivatives, $\text{M-SC}_{14}\text{Fc-C}\equiv\text{C-Fc}$ and $\text{M-SC}_{13}\text{Fc-C}\equiv\text{C-Fc}$ (see ESI† for details), and CV data for all SAMs on Au and Pt. In addition, all SAMs were characterised with MD simulations.

Fig. 2a and b shows the measured surface coverages (Γ_{CV} , in nmol cm^{-2}) of $\text{SC}_n\text{Fc-C}\equiv\text{C-Fc}$ SAMs on Au and Pt derived from CV (Fig. S2 and S3 show the CV curves and Tables S1 and S2† the corresponding data). We have previously characterised $\text{M-SC}_{15}\text{Fc-C}\equiv\text{C-Fc}$ (M = Au, Pt) by CV, and the data of this derivative was taken from our previous work.⁴² The Γ_{CV} values show distinct odd-even effects on both Au and Pt surfaces, with odd numbers of n_c giving higher Γ_{CV} , indicating a more ordered and tightly packed structured SAM than the $n_c = \text{even}$ derivatives.

On the other hand, previous work showed that for SAMs of HSC_nFc derivatives on Au this effect is largely reversed with respect to Pt (driven by differences in the M-S-C bond angles of 104° for Au and close to 180° for Pt)³² as most odd-numbered members of the series exhibited lower Γ_{CV} .^{22,49} For the current $\text{Fc-C}\equiv\text{C-Fc}$ systems, the odd-even effect on Au and Pt electrodes was alike, *i.e.*, odd numbered SAMs on both metals gave higher values of Γ_{CV} than the even numbered SAMs. Furthermore, we did not detect back-bending in the CV curves up to $n_c = 15$ (see ESI, Fig. S2 and S3†), which is in contrast to the series of HSC_nFc showing substantial back-bending for the longer derivatives ($n_c \geq 14$), resulting in additional peaks in the CV curves.^{22,49} Instead, for the current series, we observed

a gradual increase in the Γ_{CV} as a function of n_c . The observed differences are likely due to packing effects related to the large head group. The data suggests that the packing of the $\text{SC}_n\text{Fc-C}\equiv\text{C-Fc}$ SAMs is driven by the large $\text{Fc-C}\equiv\text{C-Fc}$ group, overruling the effects of the M-S-C bond angles that directed the packing and properties of the SC_nFc series.^{22,49} Our observations are supported by MD calculations of the SAM heights ($d_{\text{SAM,MD}}$, in nm) and surface coverages (Γ_{MD} , in nmol cm^{-2}) on Pt (Fig. 2c). As expected for well-packed SAMs, the $d_{\text{SAM,MD}}$ increases with n_c as the molecule becomes longer. The SAM structural order, as quantified by the damping of root-mean-square fluctuation (RMSF) values of non-hydrogen atoms, improves with length and shows a clear odd-even effect (Fig. 2d). The improved structuring in odd-numbered SAMs on Pt is reflected also in the computed odd-even packing energy differences (ΔE , in kcal mol^{-1}), with large components coming from the head groups (Fig. 2e). The MD calculations show that the odd-even effect in order and ΔE is reflected in the tilt angles α of the head groups (Fig. S10 for SAMs on Pt and Fig. S11† for the SAMs on Au).

We note that the experimental odd-even effect in Γ_{CV} cannot be observed in the modelling of Γ_{MD} , which is due to the protocol of the modelling procedure. The SAM model was created by allowing the molecules to move freely on the surface until they form packed monolayers and this equilibrated physisorbed configuration is then fixed to the surface by switching on the metal-thiolate bonds. Although this method works well



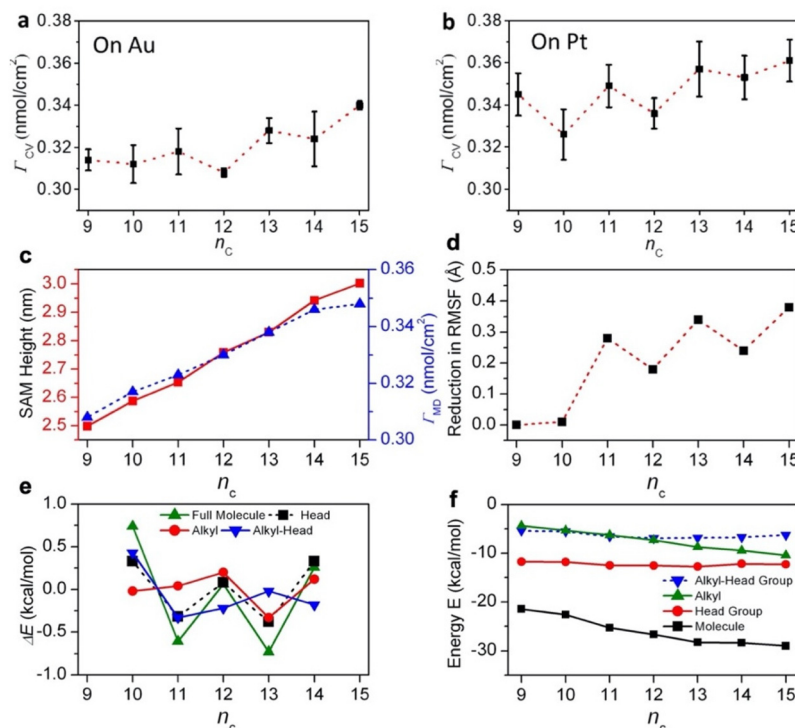


Fig. 2 Surface coverage (Γ_{CV}) of $SC_nFc-C\equiv C-Fc$ SAMs on Au (a) and Pt (b) derived from CV. $Au-SC_nFc-C\equiv C-Fc$ and $Pt-SC_nFc-C\equiv C-Fc$ were used as working electrodes, saturated $Ag/AgCl$ as reference electrode, and Pt as counter electrode. A 1.0 M $HClO_4$ aqueous solution was used as supporting electrolyte and the scan rate was 1.00 V s^{-1} . The error bars represent the standard deviation from measurements of three samples for each kind of SAM. The red dotted lines serve as guides for the eye only. Results from the molecular dynamics calculations for the $Pt-SC_nFc-C\equiv C-Fc$ SAMs. (c) SAM heights ($d_{SAM,MD}$) and surface coverages (Γ_{MD}) on Pt. (d) Computed damping of the head group motion with increasing n_c . (e) Computed odd-even effects in packing energy differences (ΔE) for the full molecule and projected on to the different parts of the molecules, i.e., the alkyl chain, the head group, and alkyl-head cross interactions. (f) Computed packing energy components (E) for the different components of the molecules of the $Pt-SC_nFc-C\equiv C-Fc$ series.

to capture different types of odd-even effects in the chemisorbed SAMs (such as oscillations in tilt angles and molecular packing energies), it does not capture the more subtle odd-even effects in surface coverages, which would require a more gradual “strengthening” of the metal–thiolate bonds in the model as the SAM forms. The computed packing energy contributions (E) of different components of the molecules (Fig. 2f), show that the combination of head group and alkyl group stabilisation promotes creation of tightly-packed, upright SAMs for all $n_c = 9$ –15 with gradual improvement in stability for longer chains. The simulations predict that at higher $n_c = 12$ –15, the alkyl chain is long enough and its packing strong enough to replace alkyl-head group cross-interactions as the primary driving force assisting head group packing in stabilising the SAMs.

Electrical characterisation of the SAMs

We measured $J(V)$ curves and determined R across the $M-SC_nFc-C\equiv C-Fc/GaO_x/EGaIn$ junctions following the procedure reported previously.^{22,42} From statistically large numbers of $J(V)$ curves, we determined the Gaussian log-average values of J , $\langle \log_{10}|J| \rangle_G$, at each measured V to construct the $\langle \log_{10}|J| \rangle_G$ vs. V curves. Details of data recording and ana-

lysis are given in the ESI,[†] and the $J(V)$ curves and corresponding histograms are shown in Fig. S14–S16.[†] We have previously collected $J(V)$ curves of $M-SC_{15}Fc-C\equiv C-Fc$ ($M = Ag, Au, Pt$), and the data of this derivative was taken from our previous work.⁴²

For the three SAM series on Ag, Au and Pt (Fig. 3a–c), the $\langle \log_{10}|J| \rangle_G$ decreased with increasing n_c , which is as expected as the tunnelling barrier width increases with increasing n_c . We observe a clear odd-even oscillation on Pt (Fig. 3c) as the odd-numbered derivatives produce smaller values of J than the even numbered ones. For junctions on Ag (Fig. 3a), this odd-even effect is weakly present but absent for junctions on Au (Fig. 3b).

All the junctions rectify electrical current at $\pm 1.0\text{ V}$ and we observe a gradual increase of R with increasing n_c , i.e., from $n_c = 9$ to 15 on all three metal surfaces (Fig. 3d–f). For instance, the R for SAMs on Ag increased by approximately an order of magnitude from $\langle \log_{10}R \rangle_G$ of 2.0 ± 0.5 to 3.0 ± 0.3 . This increase in R is likely due to the increasing surface coverage at increasing n_c along with the improvement in molecular packing, and so reduced leakage currents.^{22,23} As established above by the Γ_{CV} and the MD simulations, the longer chain derivatives pack better as the alkyl–alkyl inter-



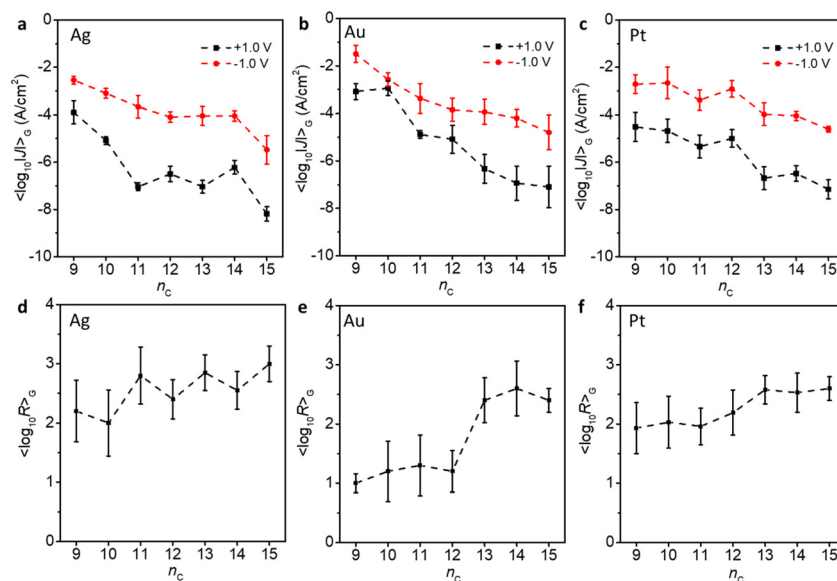


Fig. 3 Measured electrical properties $\langle \log_{10}|J| \rangle_G$ (a–c) and $\langle \log_{10}R \rangle_G$ (d–f) vs. n_c for M–SC_nFC–C≡C–Fc//GaO_x/EGaIn junctions at low bias of ± 1.0 V. The respective metals are indicated in each panel. The error bars represent the $\sigma_{\log, G}$ from the Gaussian fit of $\log_{10}|J|$ and $\log_{10}R$.

actions become more pronounced with increasing number of n_c , which translates into better diode performance (Fig. 2). A strong odd–even effect was evident for $\langle \log_{10}R \rangle_G$ on Ag (Fig. 3d), with the odd-numbered derivatives operating at consistently higher $\langle \log_{10}R \rangle_G$. These effects were less pronounced on Pt (Fig. 3f) and partially reversed on Au (Fig. 3e). The reason could be that the surface morphologies of template-stripped Ag, Au, and Pt differ from each other.^{42,50,51} This observation agrees with our earlier work where we also showed that the odd–even effect on Au is smaller than on Ag, likely due to the fact that the tilt angle of the molecular backbone on Ag (and Pt) is smaller than on Au.³² In light of the surface characterisation described above, the trend of R on Au indicates a transition around $n_c = 12$ which can be explained by a transition from an alkyl chain packing driven SAM to a head group packing dominated SAM (Fig. 2 and S11†). Probably for this series, the tilt angle of the molecular backbone (the alkyl chain) of the SAM of 30° (as opposed to reported tilts of 0–10° for SAMs on Ag and Pt)⁴⁸ explains why such a clear transition was not visible for the other two types of SAMs.

Electrical stability of the junctions

Above we have shown that the monolayers on Pt are well-organised and densely-packed and that SAMs with $n_c = \text{odd}$ performing better than those with $n_c = \text{even}$. We have established previously that the SAMs on Pt substrates are more stable and can withstand larger applied bias than those with Ag and Au electrodes.⁴² For these reasons, we focus on junctions on Pt with $n_c = \text{odd}$ to study how the value of R depends on applied voltage. To do so, we need to establish the maximum bias range we can apply for each value of $n_c = \text{odd}$ before a junction electrically shorts.

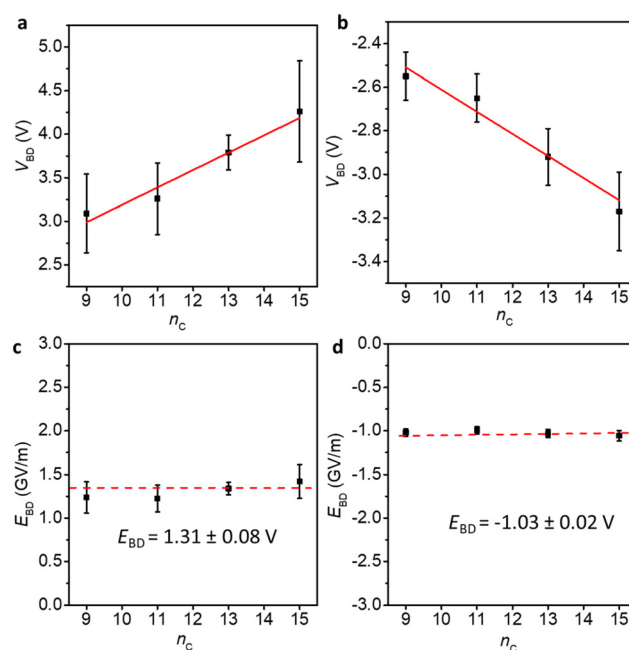


Fig. 4 Breakdown voltages V_{BD} (a and b) and breakdown fields E_{BD} (c and d) for junctions of Pt–SC_nFC–C≡C–Fc//GaO_x/EGaIn ($n_c = 9, 11, 13, 15$) at positive and negative bias, respectively. V_{BD} represents the breakdown voltage, E_{BD} represents the breakdown field. The error bars in V_{BD} represent the standard deviation (σ) from the Gaussian fits in Fig. S17.† The V_{BD} of $n_c = 15$ is taken from our previous report.⁴² The error bars in E_{BD} are calculated from the error bars of V_{BD} ($E_{BD} = V_{BD}/d_{SAM, MD}$).

Fig. 4 shows the V_{BD} (the applied maximum bias at which the junctions start to short) and the associated electric fields (E_{BD} , $E_{BD} = V_{BD}/d_{SAM, MD}$). We have previously determined the break-down voltage of Pt–SC₁₅FC–C≡C–Fc, and the data of this



derivative was taken from our previous work.⁴² The general finding is that the V_{BD} increases linearly with increasing n_c at both positive and negative bias directions, while the electric fields across the junctions remain constant, with E_{BD} of 1.31 ± 0.08 and -1.03 ± 0.02 GV m⁻¹ at positive and negative voltages, respectively. This further confirms that increasing the n_c improves the stability of the Pt-SC_nFc-C≡C-Fc//GaO_x/EGaIn junctions in terms of V_{BD} . This finding is in agreement with the breakdown behaviour of *n*-alkylthiolate based EGaIn junctions,⁵² and silicon based molecular wires of varying lengths in single-molecule junctions.⁵³

High bias operation

Through determination of the breakdown voltages, we have established the maximum safe bias windows in which the junctions are stable and do not short. Now we apply this knowledge to measure junctions of those derivatives at the maximum bias range. Fig. 5 shows the $\langle \log_{10}|J| \rangle_G$ vs. applied bias V for Pt-SC_nFc-C≡C-Fc//GaO_x/EGaIn junctions ($n_c = 9, 11, 13$, and 15) at high bias and the corresponding values of R vs. V which are also summarised in Table 1.

As shown in Fig. 5i, $\langle \log_{10}R \rangle_G$ for junctions at the highest applied bias ($\langle \log_{10}R \rangle_{G,max}$) increases linearly with increasing n_c , which indicates that the longer alkyl chain ensures a high V_{BD} , meaning that a high forward voltage can be applied without suffering from an increase in leakage current at large reverse bias. Interestingly, the voltage at which the R saturates ($V_{sat,R}$) also shows a linearly positive dependence on the number of n_c (Fig. 5j).

The onset voltage at which R begins to evolve ($V_{onset,R}$), may be used as a semi-quantitative guide to determine when HOMO and HOMO-1 begin entering the conduction window, which is approximately 0.3 V for all junctions (with the exception for junctions with $n_c = 15$ which are too resistive and the current is dominated by capacitive current near 0 V, and therefore we cannot determine the onset voltage reliably).⁴² In principle, the $V_{sat,R}$ represents the bias at which the HOMO and HOMO-1 have fully entered the conduction window, and therefore R no longer increases with increasing bias.⁴² Calculations by Zhang and co-workers indicated that other frontier orbitals, up to HOMO-5, can potentially enter the bias window and contribute to the conduction in (theoretical) single-molecule junctions of the SC_nFc-C≡C-Fc.⁵⁴ It would be interesting to investigate in more detail the potential role of deeper orbitals in our large-area junctions.

Interestingly, a clear transition can be seen around 1.8 to 2.0 V, $n_c = 9, 11$ and 13 , but the curves still continue to increase at higher voltages. The behaviour is different for junctions with $n_c = 15$ which show a continuous increase in the current all the way to the maximum applied voltage of 3.0 V. This behaviour can be explained as follows. We have reported before⁴² that at forward bias, when the HOMO and HOMO-1 fall in the conduction window (Fig. 1b, ON-State I), the bisferrocene is oxidised resulting in electrostatic attraction between the ferrocenium cations and the negatively charged top-electrode (Fig. 1b, ON-State II). This electrostatic attraction is

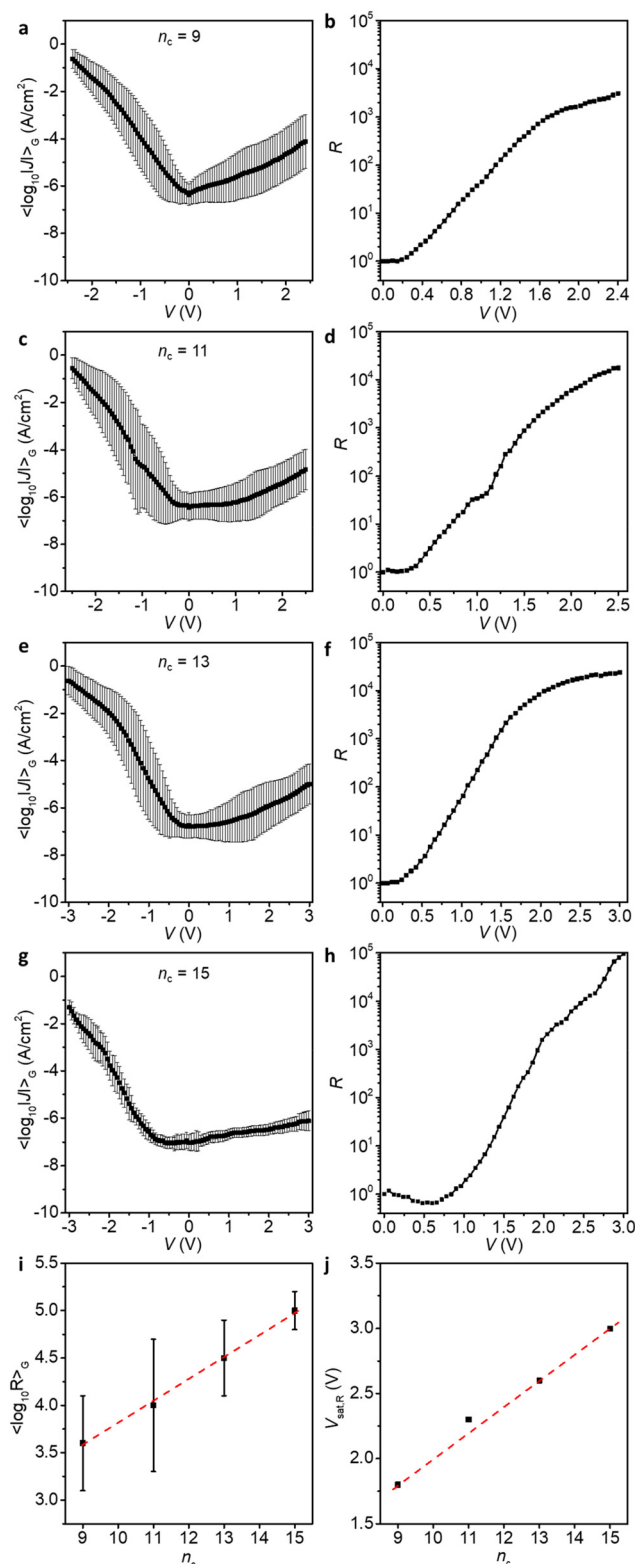


Fig. 5 Curves of $\langle \log_{10}|J| \rangle_G$ vs. applied bias V (a, c, e and g) and R against applied bias (b, d, f and h) of Pt-SC_nFc-C≡C-Fc//GaO_x/EGaIn ($n_c = 9, 11, 13$ and 15) at bias of $\pm 2.4, \pm 2.5, \pm 3.0$, and ± 3.0 V, respectively. Error bar represents the $\sigma_{\log,G}$ at each bias. (i) $\langle \log_{10}R \rangle_G$ for junctions of Pt-SC_nFc-C≡C-Fc//GaO_x/EGaIn ($n_c = 9, 11, 13$ and 15) at the highest applied bias (as in panels a, c, e and d). (j) The plot of $V_{sat,R}$ vs. n_c . The error bars in panel i are $\sigma_{\log,G}$ and the dashed red lines in i and j are guides to the eye.



Table 1 R for Pt-SC_{*n*}Fc-C≡C-Fc//GaO_x/EGaIn ($n_c = 9, 11, 13$, and 15) at high bias

n_c	9	11	13	15
Bias [V]	±2.4	±2.5	±3.0	±3.0
R	4.0×10^3	1.0×10^4	3.1×10^4	1.0×10^5

voltage driven, and for $n_c = 15$ clearly the largest voltages of up to 3 V are accessible explaining the superior performance in terms of the value of R of the diode with $n_c = 15$. We did not investigate molecular diodes with $n_c > 15$ as already at $n_c = 15$ the currents measured in the low voltage ranges were below the detection limit of our system.

Fitting employing a single-level tunnelling model

We successfully modelled the behaviour of the junctions employing a theoretical model for the Landauer formulism developed previously.⁴² Eqn (1) describes a modified version of the Landauer single-level model that accounts for the multiple junctions present in a SAM junction.

$$I = \frac{n(V)q}{\hbar} \int_{-\infty}^{\infty} dE dE' D_{E'}(E) G_e \Gamma(E') [f_L(E) - f_R(E)] \quad (1)$$

Further, a function was used to describe the number of molecules contributing to conduction as a function of the bias voltage. The individual components of eqn (1) have been explained in detail in our previous work,⁴² and are further outlined in the ESI (pages S62–S63†). Fitting was conducted by implementing several constraints and the low and high bias regimes were treated slightly differently to account for the differences in the onset voltage (see pages S64–S68† for details). Overlays of the fits of the $J(V)$ curves and the experimental results, for the whole series of diodes, are shown in Fig. S18.† Overall, a strong agreement between the theoretical calculations and the experimental data was evident. The data shows as well that the asymmetry across the junctions increased with the molecular length, and that the molecule-electrode coupling varied as a function of molecular length. This further confirms that the molecular potentiometer rule applies in the current series, by demonstrating that an asymmetric position of the molecular orbitals involved in the charge transport in the junction results in a high rectification ratio. In addition, this agrees with theoretical models developed for other rectifiers such as an alkyl cobaltocene,⁵⁵ and aryl systems with two alkylthiol anchoring groups of varying lengths.⁵⁶

Conclusions

In summary, we show that the evolution of R for SAM based molecular diodes depends on the molecular length, the bias range, and the type of metal used for the bottom electrode. We optimised these parameters to achieve consistently high R values of 10^3 – 10^5 , above the Landauer limit and approaching

the lower limit of CMOS performance. The different molecular lengths and metals affect the supramolecular structure of the SAM, and therefore impact the junction stability and leakage current. Increasing the lengths of the molecule from $n_c = 9$ to 15 resulted in a ~ 1.3 fold increase of the V_{BD} , therefore allowing to operate these stable molecular diodes at a high operating bias of $V = 3$ V. Remarkably, the tight surface packing ensured low leakage currents at reverse bias despite the high driving voltages. Subtle changes in the packing structure driven by odd–even effects as well as the important role of the bottom electrode material affect diode performance in accordance with previous findings,^{9–12,18,22,57} with Pt electrodes supporting densely packed, stable SAMs. Our findings give new insights into why diodes that “look good on paper” often showed disappointing performance, with our dataset showing that systematic improvement requires simultaneous tuning of both applied bias and (supra)molecular effects induced by supramolecular and molecule–substrate interactions. We therefore believe that our findings will inform future studies aiming to improve molecular electronic devices.

Author contributions

M.R. synthesised the compounds. X.C. and L.Y. characterised the compounds electrically. L.W. and M.R. characterised the SAMs by CV. X.Y. characterised the SAMs by XPS, UPS and NEXAFS. P.A.C. and D.T. performed the MD calculations. C.N., R.A. and E.D.B. performed the fitting. C.A.N. conceptualised and led the study. M.R., X.C. and C.A.N. wrote the manuscript, and all authors commented on the manuscript and discussed the data.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

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References

- 1 J. C. Cuevas and E. Scheer, *Molecular Electronics, An Introduction to Theory and Experiment*, World Scientific Publishing, 2010.
- 2 Y. Han and C. A. Nijhuis, *Chem. – Asian J.*, 2020, **15**, 3752–3770.
- 3 N. Xin, J. Guan, C. Zhou, X. Chen, C. Gu, Y. Li, M. A. Ratner, A. Nitzan, J. F. Stoddart and X. Guo, *Nat. Rev. Phys.*, 2019, **1**, 211–230.
- 4 D. Xiang, X. Wang, C. Jia, T. Lee and X. Guo, *Chem. Rev.*, 2016, **116**, 4318–4440.
- 5 M. Naher, M. Roemer, G. A. Koutsantonis and P. J. Low, in *Comprehensive Coordination Chemistry III*, ed. E. C. Constable, G. Parkin and L. Que Jr, Elsevier, Oxford, 2021, pp. 38–80.
- 6 A. Aviram and M. A. Ratner, *Chem. Phys. Lett.*, 1974, **29**, 277–283.
- 7 B. Capozzi, J. Xia, O. Adak, E. J. Dell, Z. F. Liu, J. C. Taylor, J. B. Neaton, L. M. Campos and L. Venkataraman, *Nat. Nanotechnol.*, 2015, **10**, 522–527.
- 8 M. Elbing, R. Ochs, M. Koentopp, M. Fischer, C. von Hanisch, F. Weigend, F. Evers, H. B. Weber and M. Mayor, *Proc. Natl. Acad. Sci. U. S. A.*, 2005, **102**, 8815–8820.
- 9 P. Song, S. Guerin, S. J. R. Tan, H. V. Annadata, X. Yu, M. Scully, Y. M. Han, M. Roemer, K. P. Loh, D. Thompson and C. A. Nijhuis, *Adv. Mater.*, 2018, **30**, 1706322.
- 10 C. A. Nijhuis, W. F. Reus and G. M. Whitesides, *J. Am. Chem. Soc.*, 2009, **131**, 17814–17827.
- 11 C. A. Nijhuis, W. F. Reus and G. M. Whitesides, *J. Am. Chem. Soc.*, 2010, **132**, 18386–18401.
- 12 L. Yuan, N. Nerngchamnong, L. Cao, H. Hamoudi, E. del Barco, M. Roemer, R. K. Sriramula, D. Thompson and C. A. Nijhuis, *Nat. Commun.*, 2015, **6**, 6324.
- 13 G. M. Morales, P. Jiang, S. Yuan, Y. Lee, A. Sanchez, W. You and L. Yu, *J. Am. Chem. Soc.*, 2005, **127**, 10456–10457.
- 14 I. Díez-Pérez, J. Hihath, Y. Lee, L. Yu, L. Adamska, M. A. Kozhushner, I. I. Oleynik and N. Tao, *Nat. Chem.*, 2009, **1**, 635–641.
- 15 A. R. Garrigues, L. Yuan, L. Wang, E. R. Mucciolo, D. Thompson, E. del Barco and C. A. Nijhuis, *Sci. Rep.*, 2016, **6**, 26517.
- 16 M. L. Perrin, E. Galan, R. Eelkema, F. Grozema, J. M. Thijssen and H. S. J. van der Zant, *J. Phys. Chem. C*, 2015, **119**, 5697–5702.
- 17 A. C. Aragonès, N. Darwish, S. Ciampi, F. Sanz, J. J. Gooding and I. Díez-Pérez, *Nat. Commun.*, 2017, **8**, 15056.
- 18 L. Yuan, R. Breuer, L. Jiang, M. Schmittel and C. A. Nijhuis, *Nano Lett.*, 2015, **15**, 5506–5512.
- 19 Y. Han, C. Nickle, Z. Zhang, H. P. A. G. Astier, T. J. Duffin, D. Qi, Z. Wang, E. del Barco, D. Thompson and C. A. Nijhuis, *Nat. Mater.*, 2020, **19**, 843–848.
- 20 L. Yuan, L. Jiang, B. Zhang and C. A. Nijhuis, *Angew. Chem., Int. Ed.*, 2014, **53**, 3377–3381.
- 21 L. Wang, L. Yuan, L. Jiang, X. Yu, L. Cao and C. A. Nijhuis, *J. Phys. Chem. C*, 2019, **123**, 19759–19767.
- 22 N. Nerngchamnong, L. Yuan, D.-C. Qi, J. Li, D. Thompson and C. A. Nijhuis, *Nat. Nanotechnol.*, 2013, **8**, 113–118.
- 23 D. Thompson and C. A. Nijhuis, *Acc. Chem. Res.*, 2016, **49**, 2061–2069.
- 24 F. Tao and S. L. Bernasek, *Chem. Rev.*, 2007, **107**, 1408–1453.
- 25 P. Cyganik, A. Terfort and M. Zharnikov, *Nano Res.*, 2023, **17**, 4231–4243.
- 26 T. Felgenhauer, H. T. Rong and M. Buck, *J. Electroanal. Chem.*, 2003, **550–551**, 309–319.
- 27 D. M. Alloway, M. Hofmann, D. L. Smith, N. E. Gruhn, A. L. Graham, R. Colorado, V. H. Wysocki, T. R. Lee, P. A. Lee and N. R. Armstrong, *J. Phys. Chem. B*, 2003, **107**, 11690–11699.
- 28 F. Ben Amara, E. R. Dionne, S. Kassir, C. Pellerin and A. Badia, *J. Am. Chem. Soc.*, 2020, **142**, 13051–13061.
- 29 Y. Feng, E. R. Dionne, V. Toader, G. Beaudoin and A. Badia, *J. Phys. Chem. C*, 2017, **121**, 24626–24640.
- 30 M. Baghbanzadeh, F. C. Simeone, C. M. Bowers, K.-C. Liao, M. Thuo, M. Baghbanzadeh, M. S. Miller, T. B. Carmichael and G. M. Whitesides, *J. Am. Chem. Soc.*, 2014, **136**, 16919–16925.
- 31 M. M. Thuo, W. F. Reus, C. A. Nijhuis, J. R. Barber, C. Kim, M. D. Schulz and G. M. Whitesides, *J. Am. Chem. Soc.*, 2011, **133**, 2962–2975.
- 32 L. Yuan, D. Thompson, L. Cao, N. Nerngchangnong and C. A. Nijhuis, *J. Phys. Chem. C*, 2015, **119**, 17910–17919.
- 33 R. Arielly, M. Vadai, D. Kardash, G. Noy and Y. Selzer, *J. Am. Chem. Soc.*, 2014, **136**, 2674–2680.
- 34 B. Cui, Y. Xu, G. Ji, H. Wang, W. Zhao, Y. Zhai, D. Li and D. Liu, *Org. Electron.*, 2014, **15**, 484–490.
- 35 D. Duche, U. Planchoke, F.-X. Dang, J. L. Rouzo, M. Bescond, J.-J. Simon, T. S. Balaban and L. Escoubas, *J. Appl. Phys.*, 2017, **121**, 115503.
- 36 H. Jeong, Y. Jang, D. Kim, W.-T. Hwang, J.-W. Kim and T. Lee, *J. Phys. Chem. C*, 2016, **120**, 3564–3572.
- 37 E. D. Mentovich, N. Rosenberg-Shraga, I. Kalifa, M. Gozin, V. Mujica, T. Hansen and S. Richter, *J. Phys. Chem. C*, 2013, **117**, 8468–8474.
- 38 L. Müller-Meskamp, S. Karthäuser, H. J. W. Zandvliet, M. Homberger, U. Simon and R. Waser, *Small*, 2009, **5**, 496–502.
- 39 S. Seo, E. Hwang, Y. Cho, J. Lee and H. Lee, *Angew. Chem., Int. Ed.*, 2017, **56**, 12122–12126.
- 40 J. Trasobares, D. Vuillaume, D. Théron and N. Clément, *Nat. Commun.*, 2016, **7**, 12850.
- 41 S. Wang, M.-Z. Wei, G.-C. Hu, C.-K. Wang and G.-P. Zhang, *Org. Electron.*, 2017, **49**, 76–84.



- 42 X. Chen, M. Roemer, L. Yuan, W. Du, D. Thompson, E. del Barco and C. A. Nijhuis, *Nat. Nanotechnol.*, 2017, **12**, 797–803.
- 43 D. Thompson, E. d. Barco and C. A. Nijhuis, *Appl. Phys. Lett.*, 2020, **117**, 030502.
- 44 N. Clement and A. Fujiwara, *Nat. Nanotechnol.*, 2017, **12**, 725–726.
- 45 N. Nerngchamnong, L. Yuan, D. C. Qi, J. Li, D. Thompson and C. A. Nijhuis, *Nat. Nanotechnol.*, 2013, **8**, 113–118.
- 46 K. Yang, Z. Cai, A. Jaiswal, M. Tyagi, J. S. Moore and Y. Zhang, *Angew. Chem., Int. Ed.*, 2016, **55**, 14090–14095.
- 47 R. Boese, H.-C. Weiss and D. Bläser, *Angew. Chem., Int. Ed.*, 1999, **38**, 988–992.
- 48 J. C. Love, L. A. Estroff, J. K. Kriebel, R. G. Nuzzo and G. M. Whitesides, *Chem. Rev.*, 2005, **105**, 1103.
- 49 N. Nerngchamnong, D. Thompson, L. Cao, L. Yuan, L. Jiang, M. Roemer and C. A. Nijhuis, *J. Phys. Chem. C*, 2015, **119**, 21978–21991.
- 50 L. B. Newcomb, I. D. Tevis, M. B. Atkinson, S. M. Gathiaka, R. E. Luna and M. Thuo, *Langmuir*, 2014, **30**, 11985–11992.
- 51 E. A. Weiss, G. K. Kaufman, J. K. Kriebel, Z. Li, R. Schalek and G. M. Whitesides, *Langmuir*, 2007, **23**, 9686–9694.
- 52 L. Yuan, L. Jiang and C. A. Nijhuis, *Adv. Funct. Mater.*, 2018, **28**, 1801710.
- 53 H. Li, T. A. Su, V. Zhang, M. L. Steigerwald, C. Nuckolls and L. Venkataraman, *J. Am. Chem. Soc.*, 2015, **137**, 5028–5033.
- 54 J.-M. Zhao, L.-Y. Chen, Y.-J. Li, N.-P. Shi, Y.-Z. Sun, H. Huang and G.-P. Zhang, *Phys. E*, 2021, **130**, 114691.
- 55 R. Liu, S. H. Ke, W. Yang and H. U. Baranger, *J. Chem. Phys.*, 2006, **124**, 024718.
- 56 P. E. Kornilovitch, A. M. Bratkovsky and R. Stanley Williams, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2002, **66**, 165436.
- 57 C. A. Nijhuis, W. F. Reus, J. R. Barber, M. D. Dickey and G. M. Whitesides, *Nano Lett.*, 2010, **10**, 3611–3619.

