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The effect of polarity on the molecular exchange dynamics in imine-based covalent adaptable networks†

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Covalent adaptable networks (CANs) are a rising type of polymeric materials that consist of covalently cross-linked polymer chains, but with the inclusion of dynamic covalent bonds, and that can perform bond exchange reactions under equilibrium control. The dynamic behaviour of these exchange reactions within a polymer matrix has been established to be a key parameter in the control of the material properties. Therefore, in order to fully control the macroscopic material properties of CANs, understanding the underlying molecular exchange processes of these dynamic covalent bonds is essential. In this work, we studied the effect of polarity in polyimine-based CANs, and considered not only the network response itself, but also the – so far often overlooked – effect on the exchange dynamics. By combining results from kinetic studies and material analysis we were firstly able to show a distinct correlation between the presence of polar domains in the molecular structure and the thermal and dynamic mechanical properties of the materials. More importantly, the presence of polar domains also greatly affected the exchange kinetics of the dynamic imine bonds. On the molecular level, we showed that the imine exchange could be greatly enhanced (up to 20 times) when polar groups were present near the reactive imine species. As a result, in our polymer materials we established a tuneable range of phase transition temperatures from glass-to-rubber and rubber-to-liquid over roughly 100 °C as a result of either presence or absence of polar groups in the polymer matrix. Furthermore, detailed analysis in the stress relaxation behaviour of the polyimine materials revealed three relaxation processes, which we could attribute to the relaxation in network topology, to the imine exchange on a local level, and to the imine exchange as a result of diffusion through the polymer network. From this analysis we were also able to illustrate the effect of polarity on the polymer network to each of the three relaxation mechanisms.

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Introduction

Vitrimer-like materials are a relatively new type of polymer that exhibit the strength of thermosets *via* a crosslinked network structure, but are still able to be recycled, reshaped, and even self-healed, as a result of dynamic covalent bonds within their molecular structure.^{1–5} These dynamic covalent bonds can perform bond exchange reactions under equilibrium control^{6,7} to construct covalent adaptable networks (CANs) when built into a polymer matrix.^{8–10} By stimulating the exchange reactions of the dynamic covalent bonds, for example by introduction of heat or light, a molecular flow is introduced in the material.¹¹ Several types of bond exchange reactions have been studied over the last decade, such as

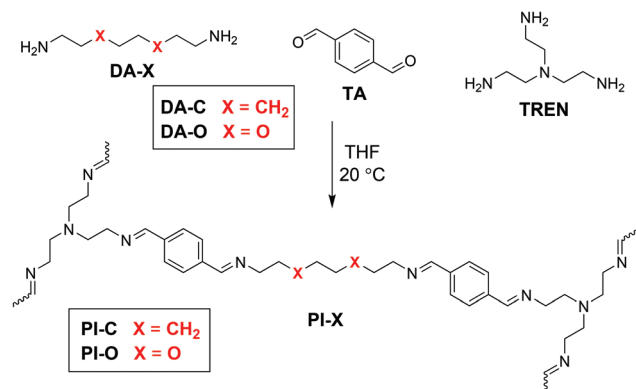
transesterifications,^{1,12} boronic ester exchange,^{13–16} 1,2,3-triazolium exchange,^{17,18} diketoenamine exchange,^{19,20} (alkyl)urea exchange,^{21–23} urethane exchange,^{24–28} thiol-ene/yne exchange,^{29,30} and imine exchange.^{31–34} Individually, each type of bond exchange reaction possesses different dynamics according to the mechanism of the exchange reaction.³⁵ Therefore, CANs can be synthesised with selective properties based on the type of bond exchange reaction. However, not only the type of bond exchange reaction is determinative for the dynamic (mechanical) response of the material, but also the composition of the polymer matrix plays a definitive role,³⁶ which includes for example the crosslinking density³⁷ or steric effects.²² For many polymeric materials, the effect of such network characteristics has been evaluated for their influence on the material properties at the macroscopic level.^{21,24,38} However, as the dynamic mechanical properties of these materials are directly influenced by the underlying molecular exchange reaction,^{22,39,40} further understanding of the specific effects of polymer com-

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To investigate the role of polarity-induced effects by EO moieties, we made use of a CAN consisting of dynamic imine bonds, as we recently reported that the imine exchange was very sensitive to electronic effects and charge distribution at the imine bond.³⁴ Furthermore, imines are favourable materials as they can be readily synthesised from a condensation reaction between an aldehyde and amine at room temperature, without a catalyst, and with water as the only side product.^{52,53} While earlier work on polyimine CANs already described the effect of solvent polarity,^{54,55} and moisture sensitivity,⁵⁶ the quantification of polar effects in the polymer matrix itself on the imine exchange has not been addressed in great detail yet. Therefore, in this study we synthesised different polyimine materials with varying polar, EO-based moieties (also in different components) of the polymer network (Scheme 1). With a combination of kinetic studies using small-molecule analogues by NMR analysis, and temperature-dependent rheology experiments on the network polymers, we were able to dissect the effect of the polarity in the material on both the macroscopic level, and on the molecular level of the bond exchange. Furthermore, we obtained a better understanding in the relaxation behaviour of CANs, as we found the stress relaxation process to proceed in three individual phases, which we could attribute to (1) chain distribution within the polymer network, (2) imine exchange at a local level, and (3) imine exchange as a result of diffusion through the polymer matrix.

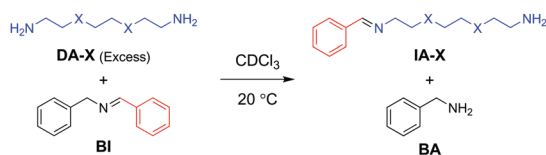
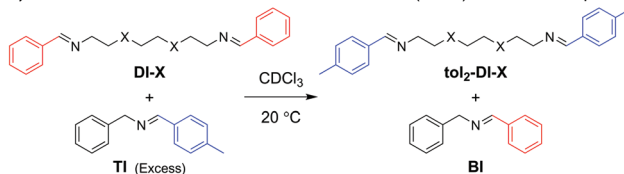


Results & discussion

Small-molecule bond exchange studies

For the initial kinetic evaluation of the bond exchange reaction, two same-sized diamine monomers (**DA-X**) were selected (Scheme 1). The first consisted of a fully carbon octyl chain (**DA-C**), and the second included two oxygen atoms on the 3' and 6' position of the chain (**DA-O**) to represent the EO moiety. Once the amines have reacted with an appropriate aldehyde, imines are formed. These imines can then perform bond exchange reactions *via* either a transimination reaction with a primary amine, or *via* imine metathesis with another imine.⁵³ In this section, the study of the kinetics of both these exchange reactions will be divided in four parts: (1) transimination for which the tuneable monomer acts as the nucleophile, (2) transimination for which the tuneable monomer acts as the electrophile, (3) metathesis for which the tuneable monomer acts as the nucleophile, and (4) metathesis for which the tuneable monomer acts as the electrophile (Scheme 2). The material that is added in excess acts as the attacking moiety. The excess is required to obtain a *pseudo* first-order reaction towards formation of the new imine, as a 1:1 ratio would result in an equilibrium between starting compound and product. In the transimination reactions (Scheme 2A and B) the primary amine acts as the nucleophile, which will attack on the C end of the imine bond.⁵⁷ For the imine metathesis reactions (Scheme 2C and D) the material that is added in excess is considered as the attacking party (nucleophile), as the exchange reaction of the starting material with the excess material will be the most prominent until new equilibrium is reached.

For the first transimination reaction, in which **DA-X** is the nucleophile (Scheme 2A), a benzyl imine (**BI**) was synthesised and dissolved in CDCl₃. An excess of **DA-X** was then added to push the reaction towards formation of the imine on the tuneable diamine chain (**IA-X**) and freeing the benzylamine (**BA**) from the starting benzyl imine (**BI**) in a *pseudo* first-order fashion. The conversion over time was followed using ¹H NMR

A) Transimination for which the tuneable monomer (**DA-X**) acts as nucleophileB) Transimination for which the tuneable monomer (**DA-X**) acts as electrophileC) Imine metathesis for which the tuneable monomer (**DA-X**) acts as nucleophileD) Imine metathesis for which the tuneable monomer (**DA-X**) acts as electrophile

Scheme 2 Overview of imine exchange reactions. (A) Transimination with **DA-X** as nucleophile, (B) transimination with **DI-X** as electrophile, (C) imine metathesis with **DI-X** as nucleophile, and (D) imine metathesis with **DI-X** as electrophile.

by integration of the imine signals of **BI** and **IA-X**. The data (Fig. 1A) were fitted using the equation for first order reaction kinetics to obtain the reaction rate constant (k) for the exchange reaction (see ESI†). The determined k values for all exchange reactions have been combined in Table 1.

For the second transimination reaction, in which **DI-X** is the electrophile (Scheme 2B), imines **DI-C** and **DI-O** were first synthesised. The imines were then dissolved in CDCl_3 and an excess of benzylamine (**BA**) was added to push the reaction towards the formation of the benzyl imine (**BI**) and transformation of the imines of the tuneable chains back to amine monomer (**DA-X**). The conversion over time (Fig. 1B) was followed using ^1H NMR by integration of the imine signals of **DI-X** and **BI**. Reaction rate constants (k) were obtained using the same procedure as before and are also presented in Table 1.

For the first metathesis reaction, in which **DI-X** acts as nucleophile (Scheme 2C), a toluene imine (**TI**) was first synthesised and dissolved in CDCl_3 . An excess of the tuneable imine chain (**DI-X**) derived from benzaldehyde was then added to push the reaction towards the formation of the benzyl imine (**BI**). The conversion was followed over time (Fig. 1C), and k (Table 1) was determined following the same procedure as before.

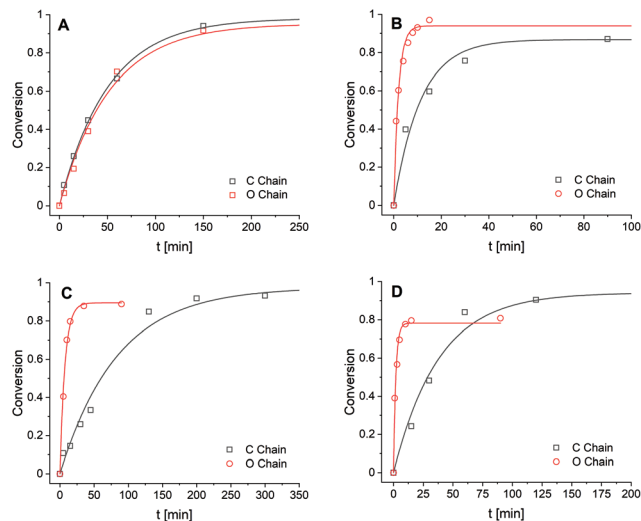


Fig. 1 Conversion as function of time for (A) transimination with **DA-X** as nucleophile, (B) transimination with **DI-X** as electrophile, (C) imine metathesis with **DI-X** as nucleophile, and (D) imine metathesis with **DI-X** as electrophile. Data points were fitted according to first-order kinetic model. Note the different timescale on the x-axis for each plot.

For the second metathesis reaction, in which **DI-X** acts as electrophile (Scheme 2D), **DI-X** was dissolved in CDCl_3 and reacted with an excess of toluene imine **TI** to push the reaction towards the formation of the bis-toluene imine chain (**tol2-DI-X**). The conversion was followed over time (Fig. 1D), and k (Table 1) was determined following the same procedure as before.

From the combined data (in Table 1) we observed that for the transimination reaction in which the **DA-X** acts as a nucleophile, that the k values were similar for the apolar and polar compound, which thus implies that the presence of the more polar EO moiety in the material has no observable effect on the reaction rate. However, for the transimination in which **DI-X** acts as electrophile, we observed that the ether-containing material yielded a five-fold higher reaction rate. These results together suggest that the reactivity of the amine is not affected by the presence or absence of the polar EO moiety in the material, but the reactivity of the imine is affected: the presence of the ether moiety enhances the reaction rate five-fold compared to the alkyl chain. The observation that imines are more affected by the polar effect than the amines might be explained by their prominent charge separation *via* the (conjugated) $\text{C}=\text{N}$ double bond. The transimination reaction can also be catalysed by trace amount of acid (which could even be water) *via* protonation of the imine nitrogen before formation of the aminal intermediate, and can therefore result in faster exchange.⁵⁷ As this charged species would be stabilised better in polar media, in addition to the availability of oxygen lone pairs to transfer the H^+ , this could explain the enhanced reaction rate for the EO-containing material. This acid-catalysed reaction would also only affect the reactivity of the imine, which could explain why no difference in reaction rate was



X	Transimination (nucleophilic)	Transimination (electrophilic)	Metathesis (nucleophilic)	Metathesis (electrophilic)
CH ₂	0.33 ± 0.01	1.68 ± 0.33	0.20 ± 0.02	0.44 ± 0.07
O	0.32 ± 0.03	8.88 ± 0.92	2.30 ± 0.17	8.69 ± 1.20

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Fig. 3 Molecular structure of the tuneable dialdehyde monomer, for which X is either CH_2 (AL-C, apolar chain) or O (AL-O, polar chain).

With two different aldehyde (AL-X) and two diamine (DA-X) monomers we could synthesise four different polyimines (crosslinked with TREN via the same procedure as before): both diamine and dialdehyde are polar (*i.e.* EO-containing linker, PI-OO), both diamine and dialdehyde are apolar (*i.e.* alkyl linker, PI-CC), the diamine is polar and the dialdehyde is apolar (PI-OC), the diamine is apolar and the dialdehyde is polar (PI-CO).

A temperature sweep experiment was performed on the four polyimine samples using the same rheology setup as before to determine the phase transition temperatures (Fig. 4). From the results we can draw a clear conclusion that the increasing polarity lowers both phase transition temperatures. In the extreme case where both the aldehyde and amine chains are polar (PI-OO) the phase transition of glass to rubber was observed at a temperature as low as 1 °C and the phase transition from rubber to liquid at 37 °C. If we compare this to the other extreme where both monomers were apolar (PI-CC), we observed a phase transition from glass to rubber at 119 °C, and a phase transition from rubber to liquid above 150 °C. This means that the phase transition temperatures were increased by more than 100 °C by simply replacing the polar linkers for apolar ones (that vary only in two of the eight atoms that make up the linker). When only one of the monomers – either the dialdehyde or diamine – was polar, we observed that the phase transition temperatures were found in between the two extremes. Comparing the PI-OC and PI-CO individually shows that the phase transition from glass to rubber is similar (57 °C for PI-OC and 59 °C for PI-CO), but the phase transition from rubber to liquid occurred at slightly higher temperatures for PI-OC (83 °C) compared to PI-CO (68 °C). This means that the presence of polar domains in either the diamine or dialde-



Fig. 4 Visual representation of the glass (blue), rubber (yellow) and liquid (red) phases of the PI-XX materials as a function of the temperature.

hyde monomer decreases the phase transition temperatures of the polyimine materials, with the addition that the presence of EO moieties in the dialdehyde monomer is slightly more effective in lowering the rubber-to-glass phase transition, and therefore shortening the rubber phase of the material. This slight difference between the effect of polar groups in either the aldehyde or amine monomer could possibly be explained by other, secondary interactions within the network, *e.g.* in relation to the aromatic moieties being present next to the EO moiety.

Stress relaxation of polyimine networks

Next to the thermal properties, the stress relaxation behaviour of PI-C and PI-O was also investigated by applying a 1% deformation to the materials and following the relaxation modulus ($G(t)$) over time until it reached zero again. In this relaxation process, we observed multiple relaxation phases, as can be seen in the normalised stress relaxation plots (Fig. 5). Please note that for PI-C and PI-O different temperatures needed to be selected in order to study the relaxation behaviour. The relaxation curves could not be accurately fitted to a one-component Maxwell model that is commonly used for the determination of relaxation times (τ) in CANs.⁶⁰ However, the data could be fitted with a three-component function (see ESI†), revealing that three individual modes of relaxation (with corresponding relaxation times) occur within the material: a fast, intermediate and slow process. The relaxation curves for PI-O and PI-C were measured at several temperatures within their rubber state (Fig. S14†) and at every given temperature the three relaxation times were determined (Table 2 for PI-O and Table 3 for PI-C).

From the relaxation times of PI-O (Table 2) we observed a clear temperature dependency for each of the three relaxation processes: τ_1 changed by a factor ~ 3.5 over a temperature range of 35 °C, τ_2 by a factor ~ 8 , and τ_3 by a factor ~ 40 . When we compared these observations to the relaxation times of PI-C (Table 3), we found distinct differences. Firstly, we observed that both τ_1 and τ_3 seem to be independent of the temperature



Fig. 5 Normalised three-step relaxation curves for PI-C at 140 °C (red) and PI-O at 20 °C (blue).



Table 2 Relaxation times (τ) for the three relaxation processes of PI-O at several temperatures within the rubber domain of the material. Full table including errors is given in Table S1†

T (°C)	τ_1 (ms)	τ_2 (ms)	τ_3 (ms)
20	13.7	4.4×10^2	18.4×10^3
25	13.2	3.3×10^2	9.6×10^3
30	12.1	2.3×10^2	5.1×10^3
35	9.0	1.4×10^2	2.8×10^3
40	7.3	1.0×10^2	1.6×10^3
45	5.5	0.74×10^2	1.0×10^3
50	5.0	0.63×10^2	0.67×10^3
55	4.1	0.56×10^2	0.46×10^3

Table 3 Relaxation times (τ) for the three relaxation processes of PI-C at several temperatures within the rubber domain of the material. Full table including errors is given in Table S2†

T (°C)	τ_1 (ms)	τ_2 (ms)	τ_3 (ms)
140	1.0	1.12×10^2	49×10^3
145	1.0	0.94×10^2	37×10^3
150	1.0	0.83×10^2	32×10^3
155	1.0	0.77×10^2	36×10^3
160	1.0	0.70×10^2	38×10^3
165	1.0	0.65×10^2	48×10^3
170	1.1	0.59×10^2	47×10^3
175	1.1	0.56×10^2	48×10^3

for **PI-C**. However, τ_2 was dependent on the temperature, as the relaxation time changed by a factor ~ 2 over a temperature range of 35 °C.

Based on the observed three-step relaxation process and determined τ values as a function of the temperature, we propose three modes of relaxation to each of the relaxation phases, with corresponding τ . The initial response (τ_1) of the material when it is exposed to a stress was attributed to chain rearrangements in the material.^{61,62} This process is generally very fast, but quickly reaches its maximum potential due to the crosslinked structure of the polymer network.^{8,63} Therefore, in order to fully relax the materials, additional relaxation through bond exchange is required.⁶⁴ We do however see that this additional relaxation through bond exchange proceeds on two different timescales (τ_2 and τ_3). This separation in two timescales could be attributed to imine exchange on a local level (τ_2), meaning exchange between imines groups that are in close proximity, and secondly to imine exchange through diffusion (τ_3). The latter occurs on a longer timeframe, as this mechanism requires the imine groups to diffuse through the network as well as find partnering imine groups to perform the bond exchange. The concept of diffusion and flow in the materials could be provided by reptation theory,^{65,66} which applies to random Brownian movement of polymer strains in entangled macromolecular structures,⁶⁷ and could be extrapolated to dynamic polymer networks as well.⁶⁸ More flexible chains would show an enhanced reptation process (better “slithering” of chains),⁶⁸ which in turn promotes the availability of imines as they move better through the medium.

Regarding the mentioned three relaxation mechanisms, we can then further discuss the observations on the stress relaxation behaviour between the polar **PI-O** and apolar **PI-C** material. As can be seen in Table 2, each of the relaxation phases for **PI-O** was significantly influenced by the temperature, whereas for **PI-C** (Table 3) only τ_2 showed a clear temperature response. The temperature response of the imine exchange is expected, although the absence of a clear temperature response for the imine exchange through diffusion (τ_3) is peculiar. An explanation for this absence might be found in earlier work on cluster formation of exchanging polar groups in apolar polymer networks.⁵⁰ The diffusion of imine groups in the apolar **PI-C** material might be limited as the polar imine groups tend to cluster together. This would mean that the local imine exchange becomes the main exchange pathway as the imine groups are already in close reach to each other, and the diffusion of imine groups through the polymer matrix is limited. The observation that τ_2 showed stronger temperature response for the polar **PI-O** material (factor ~ 8 over 35 °C range) over the apolar **PI-C** material (factor ~ 2 over 35 °C range) suggests that the increasing polarity in the polymer network also enhances the thermal response of the imine exchange. As τ_1 was found to be independent of the temperature for the apolar **PI-C**, this suggests that the glass transition of the material is mainly caused by the imine exchange, whereas for the polar **PI-O** material the flexibility within the material might also contribute to the softening of the material. Lastly, this could also add to the observation that the τ_3 in **PI-O** showed a significant temperature response (factor ~ 40 over 35 °C) as both the imine exchange and network flexibility are enhanced by increasing the temperature.

Overall, we can conclude that τ_1 can be linked to reptation and is affected by the flexibility of the polymer chains.⁶⁸ Additionally, the ease of reptation alters the diffusion and availability of exchanging imine groups, affecting τ_3 . Cluster formation of exchanging groups as a result of polarity in the polymer network might cause additional alterations to τ_3 , as well as to τ_2 .⁵⁰ The temperature dependency of the imine exchange further applies to both τ_2 and τ_3 . Lastly, our supporting kinetic data from the NMR experiments showed a clear correlation between the rate of imine exchange and presence of polar EO groups, which applies again to both τ_2 and τ_3 .

Determination of the relaxation behaviour of the **PI-XX** materials was harder to achieve as the materials were generally very brittle, with the exception of **PI-OO**, and would break easily at the required applied stress. Measurements in the liquid phase of the materials were also unsuccessful as relaxation times were too short (<1 ms) to properly analyse. Therefore, no in-depth stress relaxation analysis could be performed on these materials. Only for the **PI-OO** material (Table 4) we were able to analyse the relaxation behaviour, for which we observed a relatively similar trend in the three relaxation phases as the **PI-O** material (Table 2). This resemblance may result from the fact that both materials contain the polar EO linkers. Note, however, that the two materials have different crosslinking densities and relative polarity effects, as

Table 4 Relaxation times (τ) for the three relaxation processes of PI-OO at several temperatures within the rubber domain of the material. Full table including errors is given in Table S3†

T (°C)	τ_1 (ms)	τ_2 (ms)	τ_3 (ms)
15	11.0	3.7×10^2	12.2×10^3
20	10.0	2.7×10^2	7.3×10^3
25	8.0	1.7×10^2	4.0×10^3
30	5.7	1.1×10^2	2.2×10^3
35	4.1	0.76×10^2	1.3×10^3
40	2.7	0.58×10^2	8.6×10^2
45	2.3	0.52×10^2	6.4×10^2
50	1.4	0.50×10^2	4.8×10^2

the PI-O material contains the small TA dialdehyde monomer, and PI-OO contains the longer AL-O dialdehyde monomer that also contains an additional polar EO linker.

Conclusions

Ethylene oxide and similar ether groups are commonly applied in (dynamic) polymer materials to increase flexibility of the material. What should, however, not be overlooked in the case of CANs, is the polarity-induced effect of these ethylene oxide groups on the kinetics of the dynamic covalent bond exchange, which in turn results in alteration of the physical and thermal properties of the material. In this study, we showed that the dynamic exchange behaviour in polyimine CANs could be directly related to polarity effects in the polymer matrix. Small-molecule kinetic studies revealed that the reaction rate of imine exchange could be enhanced by introduction of more polar ethylene oxide groups when compared to apolar purely aliphatic carbon chains in the monomer. Additionally, from rheology studies on the crosslinked polymer materials we concluded that the addition of the more polar ethylene oxide groups resulted in a higher dynamic response in the materials, which was reflected in a decrease in the phase transitions from glass to rubber, and from rubber to liquid. Furthermore, extensive investigations into the stress relaxation behaviour of the materials revealed a gradual three-phase relaxation process. These were attributed to relaxation by chain redistribution within the polymer network, relaxation *via* imine exchange on local level, and imine exchange after diffusion through the network. This division in three relaxation processes within one material offers a better physical-chemical understanding of the dynamic behaviour of CANs and how molecular interactions within the polymer matrix relate to macroscopic properties of the material. These results firstly reveal the subtle, sometimes unexpected, effect that small changes in molecular structure can have on material properties, as reflected here in the particular changes in the multi-step relaxation processes. Moreover, we envision that this multi-step relaxation could also be observed in other CANs to achieve a more precise insight in, and ultimately control over, the molecular dynamics of the material.

Conflicts of interest

There are no conflicts to declare.

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