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Thermally-induced agglomeration tailors the stability of Pt SAs on TiO₂ and use in photocatalytic H₂ generation†

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Thermally-induced agglomeration of Pt single atoms (SAs) on anatase TiO₂ thin films is exploited for tuning their activity in photocatalytic H₂ generation. After thermal treatment at temperatures ranging from 350 to 650 °C in an Ar atmosphere, a more than fivefold higher photocatalytic activity is achieved under optimized conditions. The Pt species obtained after annealing at different temperatures are monitored *via ex situ* and *in situ* transmission electron microscopy. For *in situ* measurements, a gas cell system operating at 700 mbar is utilized. Structural data obtained in the temperature range from 250 to 950 °C demonstrate the co-existence of SA species and agglomerates over the whole temperature range, and give fundamental insights into thermal and light-induced agglomeration processes, including nucleation, growth and ripening of Pt species on TiO₂. Under optimized annealing conditions Pt is present as a mixture of SAs, 2D rafts and nanoparticles. Data show that annealing can stabilize the Pt species against light-induced agglomeration and detailed analysis reveals such SAs (realized under optimized annealing conditions) provide the most active sites (in comparison with various agglomerates). We further emphasize the importance of conducting atomic-scale structural characterization both before and after photocatalysis to accurately establish the process–structure–property–performance relationships of SAs in photocatalysis.

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Engineering of atomically dispersed metal sites has recently drawn tremendous research attention, with clearly foreseeable applications in the fields of catalysis, energy conversion and storage, and biomedicine. Particularly, in heterogeneous catalysis and photocatalysis, where supported active noble-metal clusters play a crucial role in the performance, the feasibility of downscaling their dimensions to the size of single atoms (SAs) offers remarkable benefits for the cost-efficiency and potentially the reactivity of the resulting catalysts.^{1–5} Owing to maximized exposure of isolated atoms to the reaction medium/interface, SA active sites are far superior to traditional nanoparticles (NPs) in terms of atomic utilization efficiency.^{6–9} At the

same time, they offer some promising features, such as unique selectivity in catalyzing certain reactions,^{10–12} as well as tunable electronic properties and activity depending on the coordination environment^{13,14} and could thus prospectively further advance the solar-to-hydrogen efficiency in photocatalytic water splitting which recently already exceeded 9%.¹⁵

Despite the significant progress made in recent years, a major obstacle to the applicability of many SA-based catalytic systems remains their stability.¹⁶ Due to their high surface energy as well as weak metal–support bonding, most supported noble metal SAs show a tendency to agglomerate under reaction conditions.^{17–19} The consequences can be clearly detrimental not only due to the loss of active surface area, but also due to size-related effects on the adsorbate binding at the active sites – *i.e.*, the latter could potentially become inactive in the desired reactions (such as H₂ evolution). On the other hand, recent investigations on light-induced agglomeration of Pt on TiO₂ (ref. 17) reveal a stable activity although only ~6% of the Pt stays in its SA state.

The rationale behind this finding is not clear, but studies which aim at identifying the nature of top-performing Pt species reveal that not all Pt SAs contribute equally to the photocatalytic activity²⁰ and that under usual photocatalytic HER conditions, low Pt loading is good enough to achieve a maximized co-

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catalytic effect.²¹ This can have several reasons, *e.g.*, the varying intrinsic activity of different SAs due to the varying Pt–support interaction or a higher activity of agglomerated Pt species as compared to their SA state. In the literature various species such as ~ 1 nm Pt clusters,^{22–25} 0.2 nm PtO species,²⁶ rafts below a size of ~ 10 atoms²¹ and dual atom sites^{27–29} have been identified to contribute primarily to the total activity. Additionally, the co-existence of SAs with rafts and intrinsic defects can have beneficial effects.^{30–33} In contrast, larger Pt agglomerates, predominantly consisting of Pt⁰ species (metallic nanoparticles), seem to not substantially contribute to the overall photocatalytic activity.^{20,21,23} Thus, studying gentle thermally-induced SA agglomeration might be an expedient strategy to extract beneficial and detrimental Pt configurations for photocatalytic applications.

Surprisingly, studies on thermally-induced agglomeration of SAs on TiO₂ and the connection of agglomerates with activity are scarce.^{5,24} Regarding thermal agglomeration, it has been established that the agglomeration depends on both the annealing atmosphere and the metal–support interaction.^{16,23} The observed effects indicate phenomena including agglomeration into amorphous clusters or crystalline nanoparticles,^{23,34–37} NP redispersion into SAs,^{35,38–42} or encapsulation of Pt species into a TiO_x shell.^{23,37,43,44} For thermally-induced agglomeration of Pt to nanoparticles on SiO₂ as well as Al₂O₃ it was found that a particle size dependent activation energy for Pt detachment⁴⁵ and Pt attachment⁴⁶ is needed to describe the observed agglomeration behavior during Ostwald ripening.^{47,48} Notably, Ostwald ripening as an acting mechanism and some studies on ceria-based SA catalysts⁴⁹ underscore the importance of a competition between agglomeration and dispersion of Pt species. Accordingly, for Pt on TiO₂ for thermally-induced agglomeration of Pt, co-existence of agglomerated and dispersed Pt species may be expected – however, a systematic study with atomic resolution is missing.

Therefore, in this study, we systematically investigate the influence of inert Ar annealing conditions over a wide temperature range on the Pt atomic surface configurations as well as the resulting photocatalytic activity of Pt on TiO₂ thin films. The Pt on the TiO₂ thin film model system enables comprehensive structural and chemical characterization *via* high-resolution scanning transmission electron microscopy (HR-STEM) and correlative X-ray photoelectron spectroscopy (XPS), both revealing the co-existence of dispersed and agglomerated Pt species with different ratios and varying cluster sizes at different temperatures. For a better understanding of the acting agglomeration mechanism, *in situ* electron microscopy in a gas cell (DENSsolution climate chip) was conducted in an Ar atmosphere (700 mbar) up to 950 °C. The *in situ* data elucidates the first steps of agglomeration, including non-classical nucleation of Pt SAs, the detachment of Pt from already existing NPs and the changing temperature-dependent balance between dispersed SA and agglomerated Pt species with a high degree of redispersion at elevated temperatures. Finally, we investigate the Pt configuration of the pre-annealed and as-deposited samples after photocatalysis, shedding light on the impact of the thermally pre-agglomerated state on its stability under

illumination. The correlation of the structural characterization with the photocatalytic activity suggests that SAs, which are preserved after both thermal annealing and photocatalysis, represent super-performing catalytic sites that provide the main catalytic activity.

Effect of temperature on photocatalytic H₂ activity of Pt SA decorated TiO₂

To study the impact of thermal Ar annealing on the photocatalytic activity of Pt loaded TiO₂ samples, first 20 nm thick TiO₂ films were sputtered on Si/SiO₂ (*cf.* Fig. S1†) and annealed at 450 °C to crystallize the thin films as described above. The crystallized anatase films were then immersed in a 2 mM H₂PtCl₆ solution for 24 h leading to highly dispersed Pt SAs on the surface.^{41,50} The films were then annealed in an Ar atmosphere in a temperature range from 350 to 650 °C. Subsequently the photocatalytic activity was determined in a quartz glass reactor under LED illumination at 275 nm at a power of 30 mW. Here, charge carriers (e⁻/h⁺ pairs), which further migrate to the surface and react with the surrounding aqueous environment, are generated in the anatase thin films. The resulting hydrogen evolution data, which is displayed in Fig. 1 as well as in Fig. S2,† reveals a clear impact of the annealing conditions on the activity. Surprisingly, the activity is higher than that of the untreated sample independent of Ar temperature. Note that the Raman spectra (*cf.* Fig. S3†) of the thin films after annealing at different temperatures reveal no changes in the TiO₂ anatase fingerprint nor the co-existence of other TiO₂ phases such as rutile, brookite, or TiO₂(B). In this context, other phases would influence the thermal and light-induced agglomeration behavior due to different bonding strengths due to their inherently different surface energetics.⁵¹ Notably, the activity increases with increasing annealing temperature and peaks at 550 °C with an activity which is ~ 5.7 fold higher than that of the non-treated sample. In the following, we will elucidate the

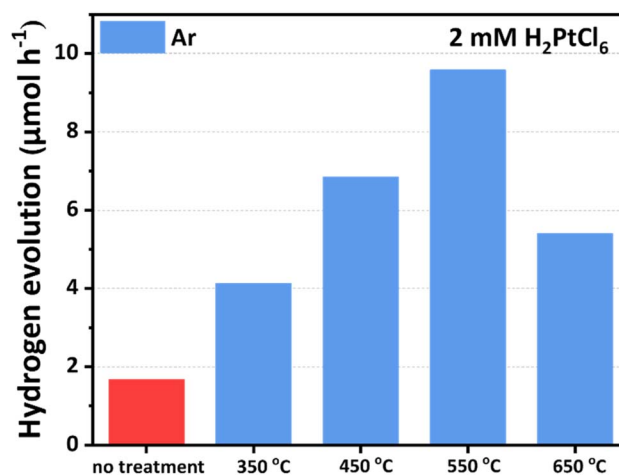


Fig. 1 Impact of thermal annealing in an Ar atmosphere on the photocatalytic activity of Pt dispersed on flat TiO₂ films.



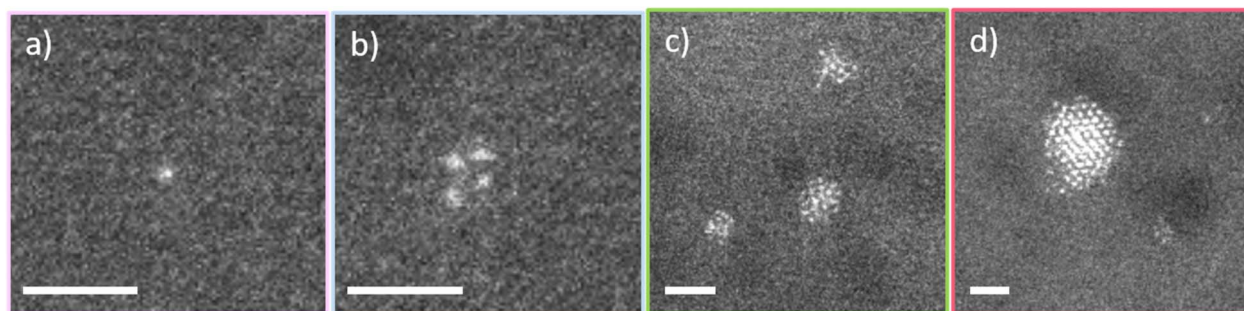


Fig. 2 Different types of Pt species typically co-existing on the TiO_2 support: (a) SA, (b) a few-atom raft, (c) rafts exhibiting a disordered structure, and (d) a NP showing lattice fringes (the scale bar corresponds to 1 nm).

corresponding Pt deposit morphology changes by the annealing and photocatalytic process by *ex situ* and *in situ* STEM investigations.

Impact of temperature on the thermal agglomeration behavior of Pt SAs on TiO_2

For high-angle annular dark-field (HAADF) imaging, 20 nm thick TiO_2 films were deposited and treated in the same way as the thin films for the activity measurements reported above. In the following we will discriminate between SAs, few-atom as well as

larger amorphous rafts and crystalline Pt nanoparticles. Typical candidates of those species are exemplarily shown in Fig. 2, here the Z dependent contrast⁵² in HAADF scanning transmission electron microscopy (STEM) imaging yields a good contrast even for Pt SAs supported on 20 nm TiO_2 , which consequently appear bright in the micrographs. To prove the correlation of Pt with the bright spots in HAADF imaging, energy dispersive X-ray spectroscopy (EDXS) mapping of a typical TiO_2 surface was conducted (*c.f.* Fig. S4†) revealing a clear correlation between the Pt signal and HAADF NPs. The evolution in the Pt configuration after annealing is captured in Fig. 3. Similarly to Farnesi *et al.*⁴⁹ for Pt on ceria-based catalysts, a mixture of SAs and assemblies can be found independent of the annealing temperature. Additionally, a larger

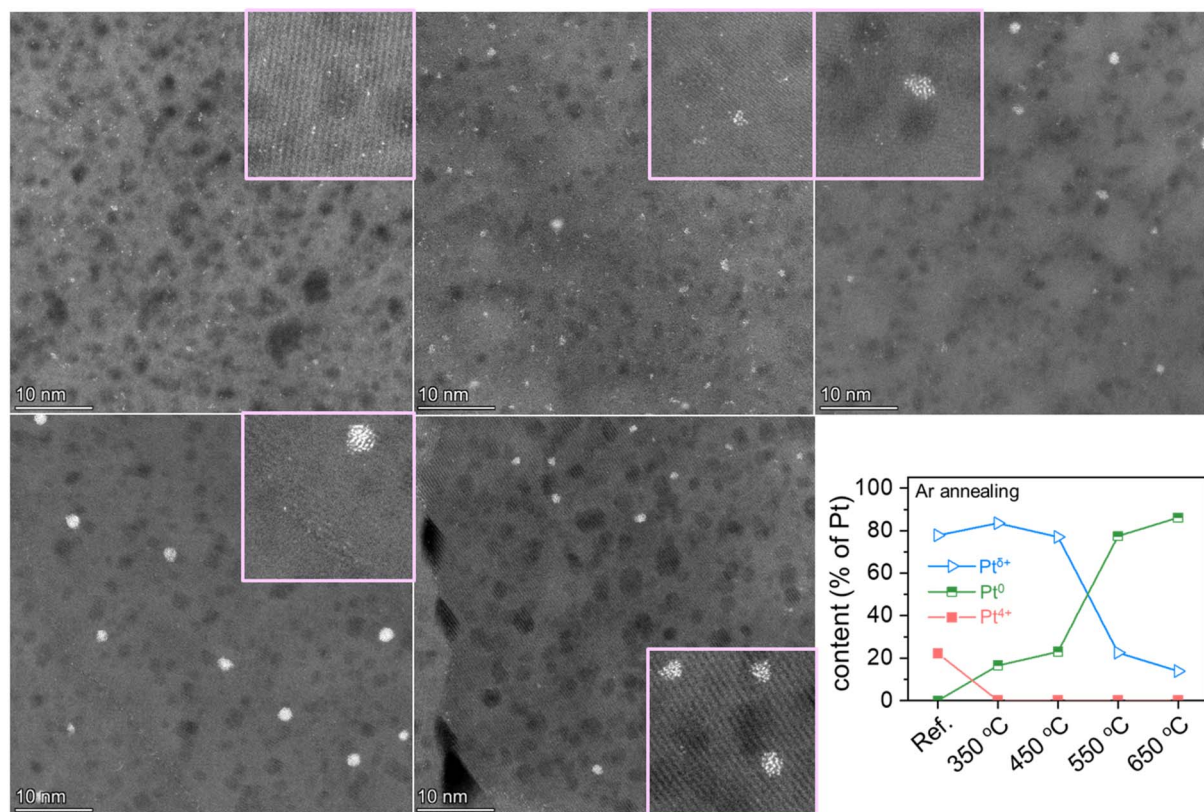


Fig. 3 Impact of thermal annealing in an argon atmosphere on the Pt dispersion and agglomeration on flat TiO_2 films (top left to bottom right: as-deposited; 350 °C, 1 h; 450 °C, 1 h; 550 °C, 1 h; 650 °C, 1 h; Pt^{5+} to Pt^0 ratio as determined by XPS).



Table 1 Percentage of Pt^{δ+}, density of SAs, rafts and NPs as well as the average NP size after different pre-annealing procedures

	As-deposited	350 °C, 1 h	450 °C, 1 h	550 °C, 1 h	650 °C, 1h
Pt ^{δ+} % (XPS)	77.8	83.5	77	22.6	13.9
#SA/μm ²	9 × 10 ⁴	1 × 10 ⁵	8 × 10 ⁴	6 × 10 ⁴	3 × 10 ⁴
#Rafts/μm ²	1 × 10 ⁴	5 × 10 ³	1 × 10 ⁴	2 × 10 ⁴	3 × 10 ³
#NPs/μm ²	0	1 × 10 ⁴	1 × 10 ³	1 × 10 ⁴	3 × 10 ³
d(NPs)/nm	n/A	1.6	1.7	2.3	1.8

degree of Pt agglomeration is visible with increasing temperature. Remarkably and similar to light-induced¹⁷ or current-induced¹⁸ agglomeration, Pt SAs always co-exist with their agglomerates even after annealing at 650 °C for 1 h. The co-existence of at least two species is also reflected *via* XPS (*cf.* Fig. 3, S5† and Table 1). Here, the XPS Pt 4f_{7/2} peak shifts gradually from ~72.6 eV to lower binding energies around ~71 eV. In accordance with the literature,⁵³ we assign one species to individual atom species Pt^{δ+} at 72.6 eV and another species to metallic Pt⁰. The position of the Pt⁰ peak deviates from that of a metal foil (~70.6 eV), and this deviation depends on the size of the Pt nanoparticles. This is because a larger fraction of Pt atoms at the surface experience a different electronic environment.¹⁷ Additionally, the strong metal-support interaction (SMSI) between small Pt NPs and the TiO₂ support under thermal treatment in an O₂-free atmosphere may contribute to the shift of Pt 4f peaks from their standard positions, as reported in the literature.^{54,55} Consequently, the position of the Pt⁰ peak was used as a fitting parameter centered around ~71 eV. The resulting analysis of the XPS spectra yields a co-existence of Pt^{δ+} and Pt⁰ species with a decreasing Pt^{δ+} fraction upon annealing. In order to quantify the degree of agglomeration with respect to the Pt species present, several STEM images for each sample were segmented by the Trainable Weka Segmentation plugin of ImageJ (*cf.* Methods section). The resulting density and size of Pt species are summarized in Table 1 revealing a decrease of the SA density in line with XPS analysis.

In general, the SA density decreases with increasing annealing temperature, still leaving 3 × 10⁴ SAs per μm² on the TiO₂ support even after annealing at 650 °C. In addition, rafts as well as NPs with an average size of around 2 nm can be found on the support after thermal annealing. Note that a correlation of the as-deposited and annealed Pt configurations would at this stage suggest, for instance, that the SAs are not the active species and hint more at the NPs. This conclusion is, however, invalid since the Pt configuration upon illumination will change within minutes,¹⁷ and thus the correlation of Pt surface-configuration and activity can only be made by investigating the Pt species after photocatalysis (*vide infra*).

Monitoring SAs and their thermal agglomeration behavior *in situ*

To shed further light on the thermal agglomeration process of Pt on TiO₂ on the atomic scale, *in situ* HAADF-STEM studies were performed aided by a gas cell setup schematically shown in Fig. S6.† Here, 20 nm thick amorphous TiO₂ layers were sputtered on the supporting TEM membranes (40 nm thick SiN_x)

embedded within temperature-controlled silicon microchips (DENSsolutions). Next, an Ar atmosphere (700 mbar) was established within the sealed space above the TiO₂ surface, while the well-controlled heating⁵⁶ of the TiO₂-covered areas was directly provided by the heating elements within the microchips. Heating to 400 °C in this setup leads to crystallization of TiO₂ layers in the anatase phase, as confirmed by *in situ* selected area electron diffraction analysis (SAED, Fig. S7†).

The aforementioned setup was then utilized in the thermal studies of Pt SAs dispersed on the TiO₂ support (Fig. 4). Notably, upon introduction of an Ar atmosphere and heating, HAADF-STEM imaging of the Pt SAs becomes significantly more challenging. Most evidently, this is due to electron beam scattering effects within the cell, which includes ~70 nm thick SiN_x membranes and the ~1 μm thick Ar gas layer. Additionally, mobile SA species under elevated temperatures could be difficult to detect due to relatively slow image acquisition time (10 μs per pixel). Nevertheless, the structural evolution of immobilized Pt species (*e.g.* small nanoparticles, clusters or SAs immobilized on the support) under elevated temperatures can be reliably monitored *in situ*, owing to the significant difference in atomic number between Pt (78) and Ti (22).

Fig. 4 shows identical-location HAADF-STEM overview images of a Pt SA loaded (0.05 mM solution, reactive deposition⁴¹) TiO₂ layer heated in Ar in the temperature range from 250 °C to 950 °C (corresponding temperature profiles are provided in Fig. S8†). This location was specifically selected to show the influence of TiO₂ edges and grain boundaries (GB) in the polycrystalline TiO₂ structure on the thermally-induced processes. The images clearly reveal a strong Pt agglomeration tendency along the 2D defects, which can be subdivided into three stages: (1) nucleation of Pt agglomerates already at 250–350 °C; (2) further growth and sintering of the nuclei at 350–550 °C; (3) reduction in the NP density at 650–950 °C. This trend was further confirmed in additional series of HAADF-STEM studies obtained under the same conditions, but other magnifications or locations on the TEM grid (Fig. S9 and S10†).

Observed phenomena of thermally-induced SA agglomeration

The Pt agglomeration observed *in situ* is in line with the *ex situ* characterization of the 2 mM samples described above. The *in situ* data adds valuable information on the history of the existing NPs and reveals that initially formed agglomerates can dissolve, whereas others act as starting points for further NP growth. In this context, Pt agglomeration is well-documented in



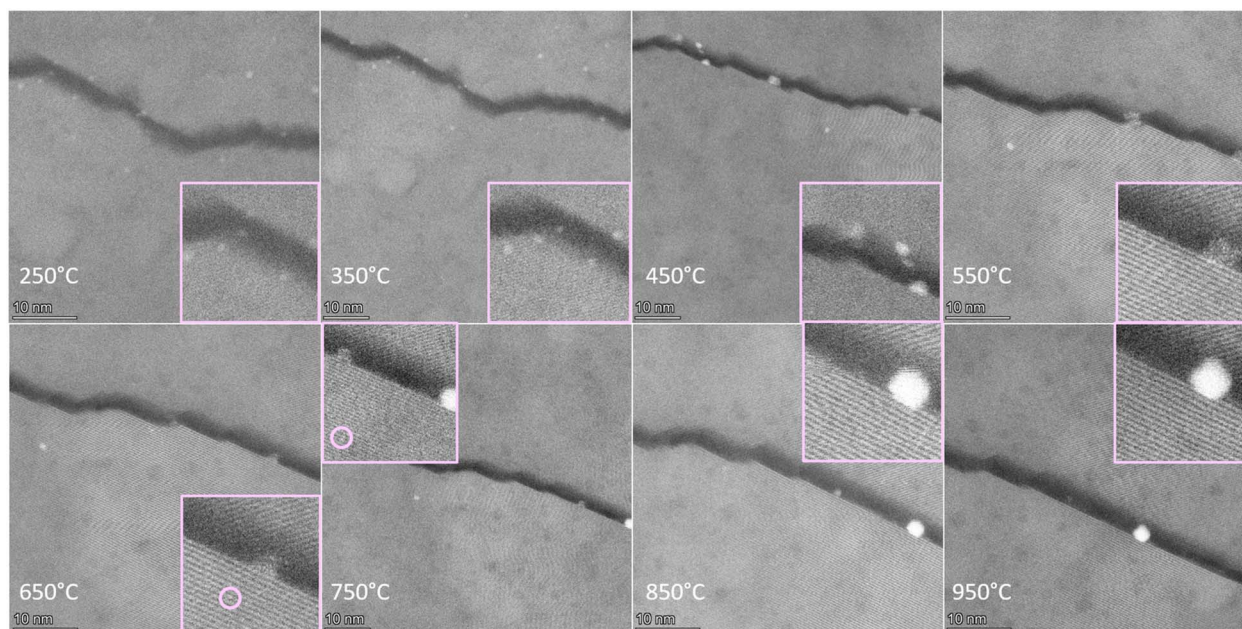


Fig. 4 Identical-location *in situ* HAADF-STEM images of the Pt SA decorated TiO₂ layer (using 0.05 mM H₂PtCl₆) heated in an Ar atmosphere (700 mbar) to different temperatures (250–950 °C). Pt species can be identified as bright spots on the TiO₂ surface (based on the significant difference in atomic number between Pt (78) and Ti (22), and the related scattering strength in HAADF-STEM imaging).

reductive atmospheres^{23,49,57} where gas adsorption weakens the metal–support interaction facilitating Pt SA destabilization and mobilization.^{23,57,58} In our case, utilizing an inert Ar atmosphere was crucial to prevent any external influence on the Pt–TiO₂ binding strength – thus the observed thermal destabilization of Pt SAs already at 250 °C confirms the intrinsically low diffusion barriers for Pt SAs on TiO₂.⁵⁹ It is also clear that the structural quality of the support plays a role in thermally-induced processes, based on preferential formation of NPs along the 2D defect. The latter can be ascribed to the defect-induced changes in the metal–support interaction, which either facilitate heterogeneous nucleation (*e.g.* *via* lowering the critical radius for nucleation⁶⁰) or provide higher mobility of SAs along extended defects⁶¹ (thus higher flux of Pt SAs towards the

nuclei). The observed dynamics of NP coarsening at higher temperatures is in line with the Ostwald ripening mechanism of metastable Pt clusters and NPs, which reduces their total surface energy.^{45–49,59} Notably, this process implies the co-existence of Pt SAs and NPs on the surface, and assumes a size-dependent activation energy for atom detachment.⁴⁶ Accordingly, in our experiments the existence of non-aggregated Pt SAs at different temperatures (650–950 °C) could be resolved in individual images presented in Fig. 5 in line with the *ex situ* data of the 2 mM samples presented above.

The apparent decline in NP density at 650–950 °C with simultaneous observations of Pt SAs strongly suggests the possibility of thermally-induced NP redistribution into SAs. Accordingly, upon cooling down the supports from 950 °C to

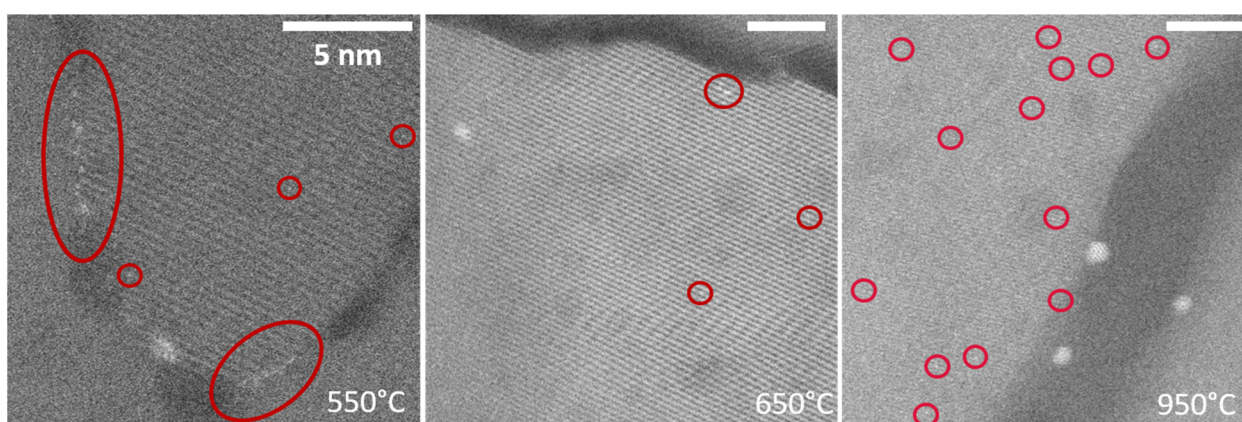


Fig. 5 High-magnification *in situ* HAADF-STEM observations of Pt SAs (highlighted) at different temperatures (following the same procedure as in Fig. 4). The scale bar in all images corresponds to 5 nm.



room temperature, we found abundant Pt SAs on the surface (Fig. S11†) and the preserved configuration of Pt agglomerates (Fig. S12†). This finding agrees with a previous report of the redispersion of Pt NPs under inert conditions and elevated temperatures.⁶² To check if the observed redispersion into SAs also results in a high photocatalytic activity, a sample was annealed at 950 °C in an Ar atmosphere. The hydrogen evolution rate (Fig. S13†) is, however, lower as compared to all other samples. This effect might stem from a strong interaction of Pt with TiO₂ potentially leading to a TiO₂ overgrowth of Pt or Pt

diffusion into the thin film and could be further investigated in the future.

Interplay of SAs and a NP: SA detachment and attachment monitored *in situ*

To gain further insights into the thermally-induced transformations and redispersion of Pt NPs supported on TiO₂, we

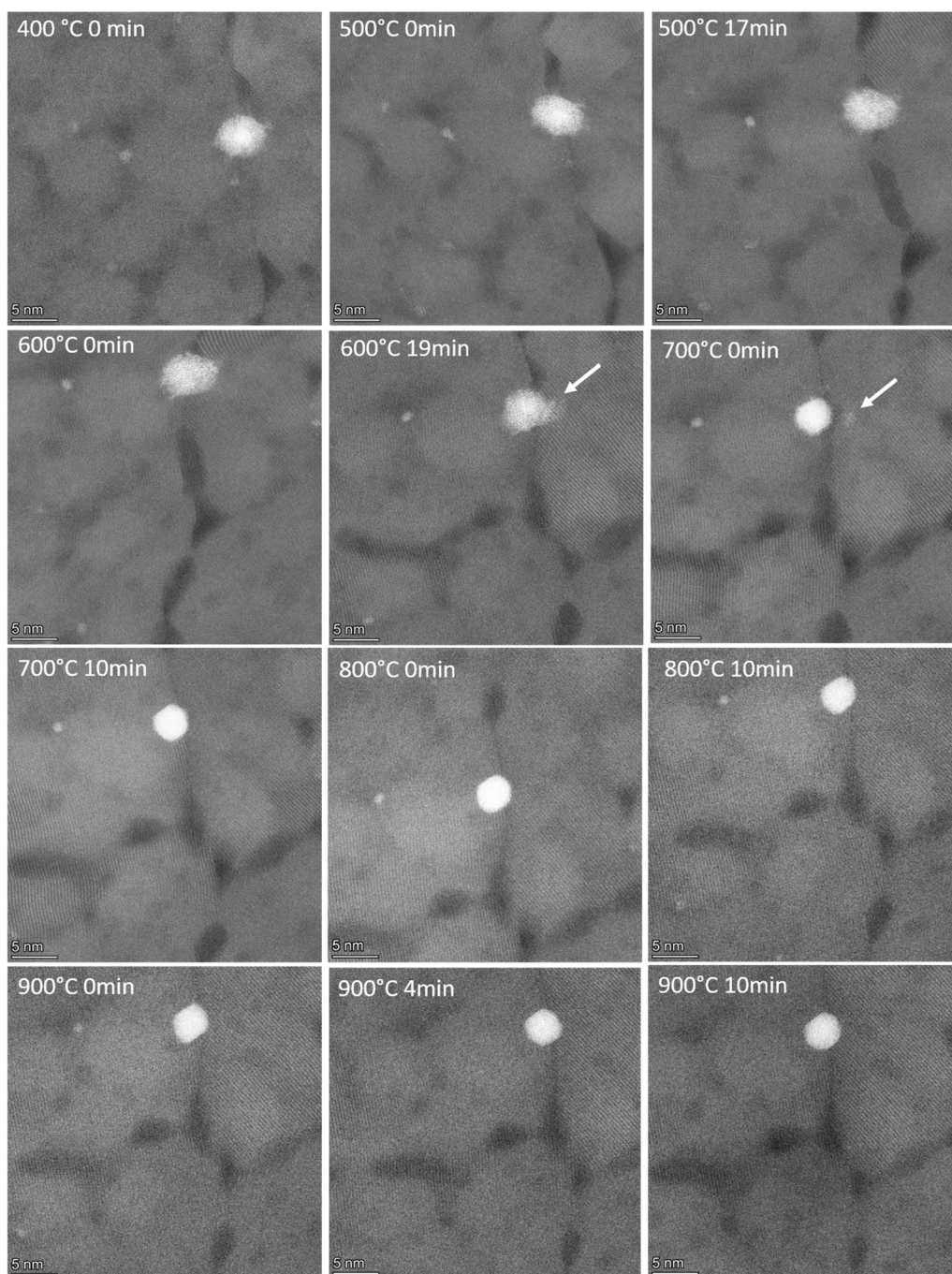


Fig. 6 High-magnification identical-location *in situ* HAADF-STEM imaging series showing the evolution of a Pt nucleus and its surroundings at 400–900 °C. The white arrow indicates the detachment of a Pt cluster from a larger particle.



carried out *in situ* identical-location observations of a Pt NP (~5 nm in diameter) at elevated temperatures, as presented in Fig. 6. Clearly, in the temperature range of 400–600 °C the NP appears as a metastable aggregate with ill-defined and overall unstable shape. Consequently, at the intermediate stage of heating (600 °C, 19 min–700 °C, 0 min) the metastable Pt agglomerate cleaves off a small cluster (marked by a white arrow). The detached Pt cloud appears to be spatially separated from the original NP by the grain boundary, and further disappears upon exposure to higher temperatures (700 °C, 10 min). This again suggests the competition between SA aggregation and redispersion in the Pt SA-TiO₂ system, although the latter process is activated at significantly higher temperatures. While such competition is implied in the Ostwald ripening mechanism, here it is experimentally verified by identical-location *in situ* HAADF-STEM with atomic resolution. The fact that smaller clusters are more prone to fragmentation/redispersion as compared to the larger agglomerates⁶³ agrees with a size-dependent activation energy for detachment (proportional to R^{-1} (ref. 45 and 46)) as a result of size-dependent surface energy.

With an increase in temperature from 600 °C to 700 °C (Fig. 6), a partially amorphous round-shaped and more flat Pt aggregate³⁰ (as revealed by STEM imaging) transforms into a compact, three-dimensional crystalline NP with distinct facets. Such reshaping and faceting of the investigated NP driven by the reduction of the surface energy is a common phenomenon for supported metal particles on oxide substrates,⁶⁴ and the size-dependent transition from a metastable liquid-like structure (at 400–600 °C) to the crystalline-ordered state (at 700–900 °C) is also in line with observations in colloidal systems at room temperature.⁶⁵

Overall, the combined *ex situ* and *in situ* data reveal that the Pt configuration can be systematically tuned by thermal annealing. Pt tends to agglomerate *via* non-classical⁶⁰ heterogeneous nucleation and further agglomerates towards NPs. In addition, Ostwald ripening, the detachment of Pt species from existing agglomerates and the co-existence of SAs with NPs is observed over the whole temperature range. Thus, we can identify three major mechanisms: (i) attachment of Pt towards agglomerates, (ii) detachment of Pt from agglomerates, and (iii) SA diffusion on the TiO₂ support. While the attachment process is rather fast, the co-existence of SAs and agglomerates suggests that the detachment rate is on a similar timescale as compared to the residence time of SAs on the support. Here, the detachment rate, which is a temperature- as well as size- and morphology-dependent quantity, and the residence time are decisively influenced by the Pt binding energy to the agglomerate and the support, respectively. Thus, tuning the metal-support interaction⁶⁶ by changing the charge state of Pt^{57,67,68} or the coordination environment by, *e.g.*, generating defects at the support Pt interface^{30,53,69} can shift the SA-agglomerate balance towards the SA side. Alternatively, increasing the detachment rate by changing the surface or interface energies of Pt agglomerates by, *e.g.*, the annealing atmosphere^{49,57} will lead to a similar effect. Another way to increase the mean free path for diffusion and thus the residence time on the support is tuning

the Pt concentration: Here, very dilute SA configurations are found to be stable against sintering as well described in the literature.^{16,70} In line with these findings, we observed almost no Pt thermal aggregation at lower SA loading (produced using 0.005 mM precursor solution) on the TiO₂ surface (Fig. S14†).

Pre-annealing as a remedy against light-induced agglomeration and identification of active sites

While thermal agglomeration is activated by temperature, light-induced and current-induced agglomeration are activated by a weakening of the metal-substrate interaction due to H adsorption.^{17,18} Thus, it is interesting to see how light-induced agglomeration interacts with thermally pre-agglomerated species. For that, the as-deposited as well as the annealed samples were investigated after photocatalysis under illumination at 275 nm (30 mW) for 3 h. The resulting Pt configurations are exemplarily shown in Fig. 7 and summarized after segmentation (*vide supra*) in Table 2. Interestingly the SA density after photocatalysis is the lowest for the least active sample; thus, light-induced agglomeration acted the most on the non-annealed/as-deposited sample. One likely reason can be the generation of oxygen vacancies (O_vs) during thermal annealing in an Ar atmosphere, which has been found for powder samples by electron paramagnetic resonance (EPR) in the past.⁷¹ Here, EPR reveals a maximum density of O_vs after annealing at 500 °C, those vacancies are then potential sites stabilizing SAs during photocatalysis. As can be seen by, *e.g.*, the average NP size or the final SA density, all samples underwent further agglomeration, while the raft density is reduced (except for 650 °C). In contrast to the solely thermally agglomerated state, the most active samples (450 and 550 °C) exhibit the lowest NP density, thus NPs can be excluded as main active sites. Similarly, the raft density is the highest for 650 °C and quite low for 450 °C, also indicating that rafts alone are not the most efficient activity contributors. All annealed samples, however, exhibit a higher SA density as compared to the as-deposited sample indicating that SAs are responsible for the activity. We also note that the 650 °C sample, although being in a similar range to the other most active samples, exhibits the highest SA density, which is a strong indication that not all SAs exhibit the same activity, in line with previous findings.^{20,21} Interestingly, the correlation with XPS data (*cf.* Fig. 7 and Table 2) reveals that the most active samples exhibit the highest abundance of Pt^{δ+} species. However, the amount of Pt^{δ+} detected by XPS follows the activity trend, but decreases significantly compared to the density of SAs observed by STEM (*cf.* Table 1 and Fig. S15†). While a fairly linear relationship between SA density and Pt^{δ+} species exists after annealing, this relationship breaks down following photocatalysis. This suggests that not all structural SAs (STEM) are in the same electronic state (XPS), highlighting the importance of stabilizing SAs at the appropriate substrate sites to maximize the efficiency of the co-catalyst.⁵ In this context, recent studies revealed a massive impact of the exposed TiO₂ surface on the Pt SA coordination and



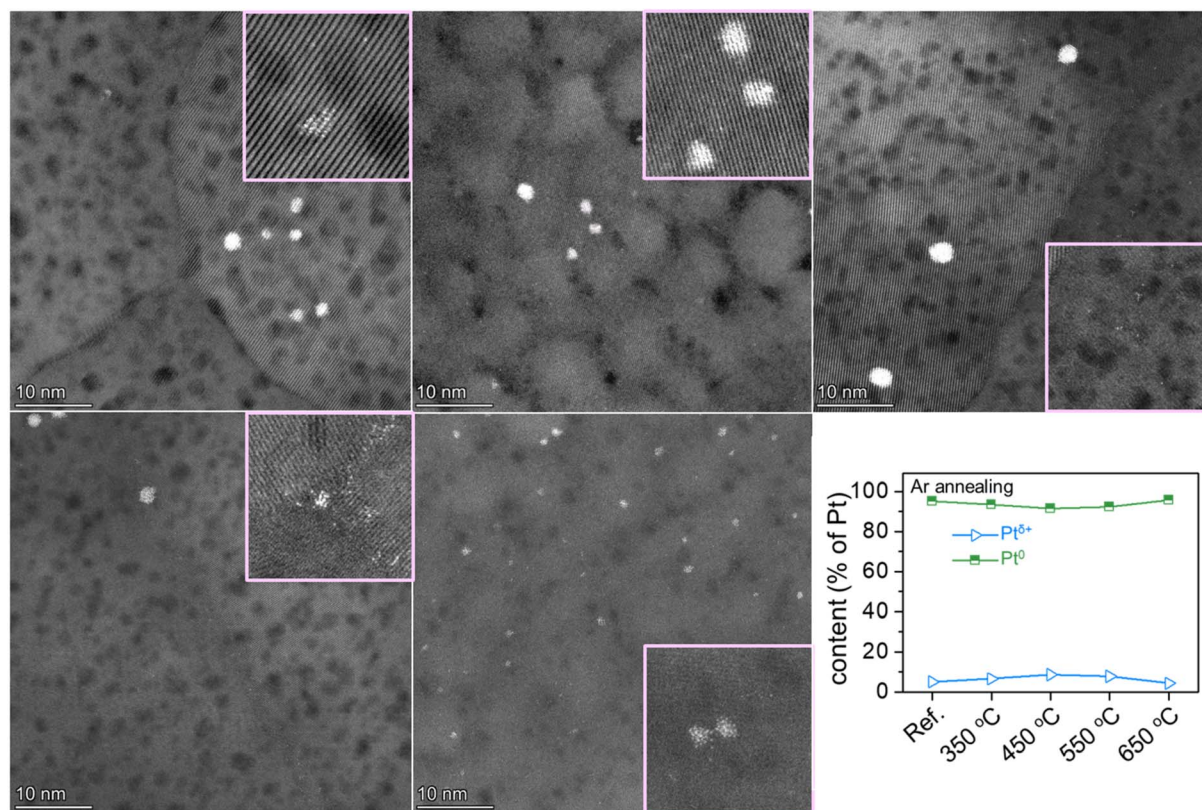


Fig. 7 Impact of thermal annealing in argon on the light-induced agglomeration behavior of Pt species on flat TiO₂ films (top left to bottom right: as-deposited; 350 °C, 1 h; 450 °C, 1 h; 550 °C, 1 h; 650 °C, 1 h; Pt⁰⁺ to Pt⁰ ratio as determined by XPS).

Table 2 Percentage of Pt⁰⁺, density of SAs, rafts and NPs as well as the average NP size after different pre-annealing procedures and photocatalysis for 3 h (275 nm, 30 mW)

	As-deposited	350 °C, 1 h	450 °C, 1 h	550 °C, 1 h	650 °C, 1 h
Pt ⁰⁺ % (XPS)	5	6.6	8.6	7.7	4.3
#SA/μm ²	1 × 10 ⁴	5 × 10 ⁴	4 × 10 ⁴	6 × 10 ⁴	6 × 10 ⁴
#Rafts/μm ²	2 × 10 ³	5 × 10 ³	3 × 10 ³	8 × 10 ³	2 × 10 ⁴
#NPs/μm ²	1 × 10 ³	4 × 10 ³	1 × 10 ³	2 × 10 ³	8 × 10 ³
d(NPs)/nm	1.8	2.4	2.9	3.1	1.8

activity.^{58,72} As described above O_vs may play a key role in (i) stabilizing SAs in potentially the right coordination environment and (ii) acting as co-catalytic sites as recently reported for Pd³⁺ and Pt³⁺ on TiO₂ supports. Thus, whereas agglomeration in electro- and photocatalysis takes place close to room temperature due to H adsorption weakening the binding energy to the support and enhancing the self-diffusion of Pt by orders of magnitude,⁷³ thermal agglomeration requires elevated temperatures to overcome the Pt binding energy towards the TiO₂ substrate sites. In this context, elevated temperatures will lead to the generation of additional point defects such as O_vs on the support, which serve as additional anchoring sites for SAs partially stabilizing the system. Thermal annealing also helps to guide the SAs to the most stable anchoring sites on the substrates, since here the binding energy is the highest and consequently the Pt surface diffusion kinetics are the lowest.

Conclusion and outlook

In conclusion, our study demonstrates the feasibility of thermal treatments for fine-tuning the state of Pt as SAs, rafts, or 3D nanoparticles, and thus presents a tool for optimizing the Pt-speciation towards a maximum photocatalytic activity. We find that the photocatalytic activity of a Pt SA decorated sample annealed at 550 °C is more than five times higher as compared to that of the as-deposited one. *Ex situ* HRSTEM investigations reveal that the Pt dispersion and agglomeration state can be tuned by the treatment temperatures. While in any case a combination of SAs, rafts and NPs exists, overall, higher annealing temperatures lead to larger NPs (but still co-existing with SAs on the support). *In situ* investigations in the gas cell reveal details of the microscopic mechanism of agglomeration. At 250 °C, a high density of small 2D SA clusters (rafts) is formed



that then act either as nuclei and grow further or dissolve with increasing temperature. Since the structure of the mainly 2D object is far from that of the final NP, we also reveal a non-classical nucleation pathway meaning that the initial nuclei exhibit a different structure as compared to Pt NPs in equilibrium. The dissolution of initially formed nuclei documented *in situ* by high-resolution imaging demonstrates the detachment of Pt atoms and assemblies from ill-defined nuclei as well as from the NP on an atomic scale. Here, an increasing number of initially formed clusters dissolved with increasing temperature, while others grow further into stable and crystalline Pt NPs. In addition, the *in situ* as well as *ex situ* structural data documents the importance of predominant heterogeneous nucleation taking place at grain boundaries during thermally-induced agglomeration. Moreover, we demonstrate that thermal pre-annealing decisively influences later light-induced agglomeration during photocatalytic applications. Here, the non-annealed sample is most prone to agglomeration within minutes,¹⁷ while annealed samples agglomerate, but to a much lower relative extent. Correlating the structural and electronic data after photocatalysis with the activity data indicates that SAs (in the right coordination environment and electronic state) are likely the most active sites during photocatalysis. The above consideration shows that – for a most effective use of Pt and a maximized activity and stability – on the one hand, strategies that shift the SA to agglomerate ratio towards the SA side are needed; on the other hand, the residual SAs can be trapped in a “right” structural environment (*e.g.* in the vicinity of $O_{v,s}$).

Methods

Formation of TiO₂ layers on Si/SiO₂ and SiN_x membranes

20 nm thick amorphous TiO₂ films were produced on the viewing windows of DENSsolutions climate chips as well as on Si/SiO₂ substrates by magnetron sputtering. The films were then crystallized *via in situ* heating in the climate system (700 mbar Ar atmosphere) for 30 min at 400 °C and within a conventional heating furnace for 1 h in an Ar atmosphere, respectively. Alternatively, the crystallization was achieved during the thermal studies with pre-loaded Pt SAs.

Deposition of Pt SAs on TiO₂ layers

The Pt SAs were produced on the crystalline TiO₂ layers *via* a facile reactive deposition method,^{17,21,53} whereas the *ex situ* samples and chips were immersed in aqueous 2 mM and 0.05 mM H₂PtCl₆:6H₂O solutions, respectively. All samples were kept in the dark for 1 h, while the backside was covered in order to only expose the TiO₂ functionalized side to the precursor solution. The membranes were subsequently washed with deionized water and dried in an N₂ atmosphere. In a separate experiment a lower Pt SA loading (0.005 mM H₂PtCl₆) on the amorphous TiO₂ layers was realized. The estimated Pt loading produced on the surface is equal to 0.14 at% (for 0.005 mM H₂PtCl₆), 0.47 at% (for 0.05 mM H₂PtCl₆), and 0.78 at% (for 2 mM H₂PtCl₆), respectively.²¹

Photocatalytic H₂ evolution

Photocatalytic H₂ evolution performance of the Pt SA-loaded TiO₂ anatase layers was evaluated under UV illumination. The samples were placed in a quartz reactor containing 50 vol% methanol (hole scavenger) aqueous solution (10 mL), which was subsequently purged with Ar for 15 min. After that, the reactor was sealed, and illumination with an LED ($\lambda = 275$ nm, power density of 20 mW cm⁻², exposure area = 1.2 cm × 1.2 cm) was carried out for a specified duration. A gas chromatograph (GCMS-QO2010SE, SHIMADZU) equipped with a thermal conductivity detector (TCD) was used to determine the amount of H₂ generated at specific time intervals.

SEM and XPS characterization

The surface and cross-section morphologies of the samples were investigated using a field-emission scanning electron microscope (FE-SEM, S-4800, Hitachi). The chemical composition of samples was analyzed by X-ray photoelectron spectroscopy (XPS, PHI 5600). All XPS spectra were shifted to a standard Ti 2p binding energy in anatase of 458.5 eV and the peak deconvolution was carried out by MultiPak software.

Ex situ STEM imaging

HAADF STEM imaging was conducted utilizing an aberration-corrected Thermo Fisher Scientific Spectra 200 operating at 200 keV. Images were taken at 960 kx as well as 1.8 Mx and various different positions each. For that a beam convergence angle of 15 mrad, a collection angle of 56–200 mrad, a beam current below 30 pA, and dwell times of 20 μ s and 30 μ s have been used.

In situ STEM imaging

High-resolution HAADF STEM imaging was performed at a double corrected FEI Titan Themis³ 300 operating at 300 kV. The beam convergence angle was 15.7 mrad. The collection angle was 38–200 mrad at a dwell time of 10 μ s and 30–184 mrad at a dwell time of 2 μ s for the 0.05 mM and 0.005 mM samples, respectively. To avoid drift due to charging of the membrane and to minimize the influence of the electron beam, the beam current was below the sensitivity limit of the screen of 30 pA. The DENSsolutions climate system was utilized for *in situ* heating and control of the atmosphere. In all *in situ* thermal studies, an Ar atmosphere with a pressure of 700 mbar and 0.3 mL min⁻¹ flow rate was maintained.

Segmentation of STEM images

For statistical analysis of the particle size and density, STEM images were segmented using the software (Fiji is Just) ImageJ utilizing the Trainable Weka Segmentation plugin within a human-in-the-loop approach. The segmentation was trained with respect to three classes, namely, SAs, agglomerates and background/substrate. Typically, 20 to 30 areas per class were chosen as reference, while the segmentation result was stepwise checked. Further, the resulting segmentation was analyzed with the particle analyzer of Fiji. Here, edges were excluded whereas



all other segmented SA and NP areas were analyzed. Subsequently, the results were classified within three classes: (i) SAs, (ii) rafts, and (iii) NPs. For that a maximum size/diameter of 4 Å corresponding to the (100) lattice spacing of anatase and 1 nm have been chosen to classify SAs and rafts as displayed in Tables 1 and 2, respectively. Everything beyond 1 nm is classified as NP.

Data availability

Related files are available online upon publication at <https://doi.org/10.5281/zenodo.14507308>.

Author contributions

J. Will: conceptualization, data curation, formal analysis, investigation, methodology, validation, visualization, writing – original draft, writing – review & editing. N. Denisov: formal analysis, investigation, visualization, writing – original draft, writing – review & editing. S. Qin: data curation, formal analysis, investigation, visualization, writing – original draft, writing – review & editing. M. Wu: investigation. Y. Wang: investigation. H. Kim: investigation. N. Karpstein: investigation, writing – review & editing. M. Dierner: formal analysis, investigation, visualization. P. Schmuki: conceptualization, funding acquisition, methodology, resources, supervision, writing – original draft, writing – review & editing. E. Spiecker: conceptualization, funding acquisition, resources, supervision, validation, writing – review & editing.

Conflicts of interest

The authors declare no competing financial interest.

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