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The authors would like to correct errors in the legends of the published Fig. 2 and Fig. 3. The correct versions of Fig. 2 and Fig. 3 are shown below.

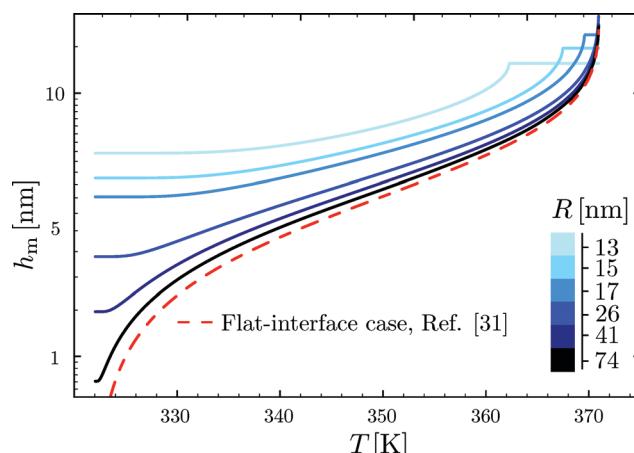


Fig. 2 Predicted surface mobile-layer thicknesses h_m of spherical polystyrene nanoparticles as a function of temperature T , according to eqn (16) and (17), for different sphere radii as indicated. We used the bulk glass-transition temperature $T_g^{\text{bulk}} = 371$ K,⁶⁹ and the onset temperature $T_c = 463$ K.^{57,70} We fixed the molecular size $\lambda_V = 3.7$ nm, and the Vogel temperature $T_V = 322$ K, to the values previously obtained for the thin-film geometry.³¹ Note that we replaced the $+\infty$ bound by 25 in eqn (16), and checked that it provides sufficiently precise numerical estimates. For comparison, the dashed line indicates the flat-interface result used for the thin-film geometry.³¹

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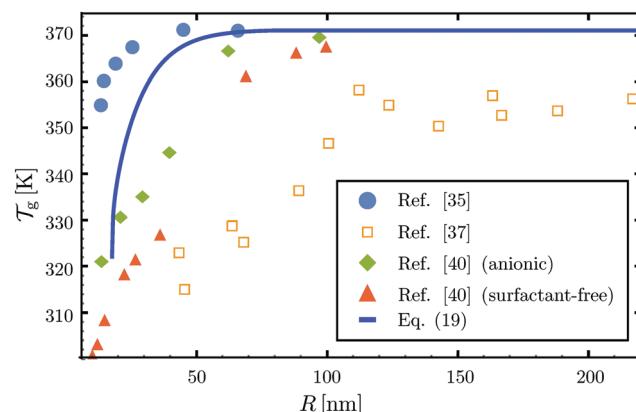


Fig. 3 Comparison between experimental data (symbols) for the reduced glass-transition temperature $T_g(R)$ of spherical polystyrene nanoparticles^{35,37,40} of radius R , and the theory (line) given by eqn (19) – that invokes eqn (16) through $\mathcal{F}(\nu) = f(2^{-1/3}, \nu)$. The fixed parameters are the bulk glass-transition temperature $T_g^{\text{bulk}} = 371$ K,⁶⁹ and the onset temperature $T_c = 463$ K.^{57,70} The two adjustable parameters are the molecular size $\lambda_V = 3.7$ nm, and the Vogel temperature $T_V = 322$ K, that were fixed to the values previously obtained for the thin-film geometry.³¹ Note that we replaced the $+\infty$ bound by 25 in eqn (16), and checked that it provides sufficiently precise numerical estimates.

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

