Hydrocarbon-Based Statistical Copolymers Outperform Block Copolymers for Stabilization of Ethanol−Water Foams

James Jennings,* Rebekah R. Webster-Aikman, Niall Ward-O’Brien, Andi Xie, Deborah L. Beattie, Oliver J. Deane, Steven P. Armes,* and Anthony J. Ryan*

Cite This: ACS Appl. Mater. Interfaces 2022, 14, 39548−39559

ACCESS

ABSTRACT: Well-defined block copolymers have been widely used as emulsifiers, stabilizers, and dispersants in the chemical industry for at least 50 years. In contrast, nature employs amphiphilic proteins as polymeric surfactants whereby the spatial distribution of hydrophilic and hydrophobic amino acids within the polypeptide chains is optimized for surface activity. Herein, we report that polydisperse statistical copolymers prepared by conventional free-radical copolymerization can provide superior foaming performance compared to the analogous diblock copolymers. A series of predominantly (meth)acrylic comonomers are screened to identify optimal surface activity for foam stabilization of aqueous ethanol solutions. In particular, all-acrylic statistical copolymers comprising trimethylhexyl acrylate and poly(ethylene glycol) acrylate, P(TMHA-stat-PEG), confer strong foamability and also lower the surface tension of a range of ethanol−water mixtures to a greater extent than the analogous block copolymers. For ethanol-rich hand sanitizer formulations, foam stabilization is normally achieved using environmentally persistent silicone-based copolymers or fluorinated surfactants. Herein, the best-performing fully hydrocarbon-based copolymer surfactants effectively stabilize ethanol-rich foams by a mechanism that resembles that of naturally-occurring proteins. This ability to reduce the surface tension of low-surface-energy liquids suggests a wide range of potential commercial applications.

KEYWORDS: statistical copolymers, polymeric surfactants, surface activity, foam stabilization, micellization, X-ray scattering

■ INTRODUCTION

Polymeric surfactants are important for many industrial sectors, including detergents for home and personal care, printing inks, oil recovery, latex syntheses via emulsion polymerization, cosmetics, and as excipients for drug formulations.1−5 Polymeric surfactants can offer significant advantages over their small-molecule counterparts, including very low critical micellar concentrations (CMC), fine control over the hydrophilic−lipophilic balance (HLB), and enhanced emulsion stability.3 Block copolymer architectures in which hydrophilic and hydrophobic comonomers are spatially segregated into two or more blocks are particularly common. For example, Pluronics comprising distinct blocks of poly(ethylene oxide) (PEO) and poly(propylene oxide) (PPO) have found widespread use in many of the above industrial applications.4,5 Traditionally, the synthesis of block copolymer surfactants requires multiple steps and stringent reaction conditions using techniques such as anionic polymerization.6 However, the development of pseudo-living radical polymerization techniques in the 1990s has allowed the preparation of many functional block copolymers using less synthetically demanding conditions.7,8 Nevertheless, the specialty reagents that are required for such syntheses are typically expensive and toxic. In principle, this problem can be mitigated by targeting longer copolymer chains. Unfortunately, high molecular weight block copolymer surfactants (>15 kg mol−1) usually lead to lower surface activity.9−12 Thus, linear (multi)block copolymer architectures may not be optimal for polymeric surfactants.13 Amphiphilic polymeric surfactants with branched, star-like, or dendritic architectures can also exhibit high surface activity.1 However, such architectures typically involve much more complex synthetic protocols and require reagents that are not readily applicable to industrial-scale synthesis.

Nature often employs proteins as polymeric surfactants, with amphiphilic character conferred by the hydrophilic and hydrophobic amino acid side-chains that decorate the polypeptide backbone.14 Stochastic evolutionary design has produced highly surface-active proteins that can reduce the
surface tension of water to 25 mN m\(^{-1}\),\(^{15}\) which is rarely achieved when using synthetic hydrocarbon-based surfactants. The surface activity of certain proteins is exploited to prepare ultrastable foams. For example, the African tree frog creates robust protein-stabilized foams that can survive harsh environmental conditions and hence protect its spawn.\(^{14,16,17}\) Protein-
based surfactants can be used for fire-retardant foams and also in the food and beverage industry for the stabilization of edible emulsions and foams.

Notwithstanding the precise secondary and tertiary structures often formed by amphiphilic proteins, the distribution of hydrophilic and hydrophobic amino acids along polypeptide backbones resembles that of a statistical (or random) synthetic copolymer. Amphiphilic statistical copolymers have already shown considerable promise as emulsifiers. However, numerous reports suggest that amphiphilic statistical copolymers are inferior to their block copolymer analogues in terms of surface activity. Recently, the self-assembly of amphiphilic statistical copolymers and alternating copolymers has been explored but the precise relationship between chemical structure, the formation of colloidal aggregates in aqueous solution, and surface activity remains unclear. However, there is little doubt that the synthesis of such copolymers should be amenable to industrial scale-up.

Herein, we evaluate a library of amphiphilic hydrocarbon-based copolymers in terms of their ability to stabilize foams generated from various ethanol–water mixtures. By screening a wide range of hydrophobic/hydrophilic copolymer pairs and targeting various copolymer compositions, we identify new copolymer surfactants that can stabilize foams for ethanol-rich solutions required for hand sanitizer formulations. Foam stabilization experiments and surface tensiometry measurements demonstrate that such statistical copolymers are more surface-active than the analogous diblock copolymers. Furthermore, small-angle X-ray scattering (SAXS) studies indicate that statistical copolymers undergo unimolecular self-assembly in ethanol–water mixtures, implying an unfolding mechanism at the air–water interface that resembles that of natural protein surfactants. New environmentally-friendly foam stabilizers for ethanol–water mixtures are particularly timely given the unprecedented growth in the use of alcoholic hand sanitizer formulations. Foam stabilization experiments and surface tensiometry measurements show that statistical copolymers are more active than the analogous diblock copolymers. Additionally, small-angle X-ray scattering (SAXS) studies indicate that statistical copolymers undergo unimolecular self-assembly in ethanol–water mixtures, implying a unfolding mechanism at the air–water interface that resembles that of natural protein surfactants. New environmentally-friendly foam stabilizers for ethanol–water mixtures are particularly timely given the unprecedented growth in the use of alcoholic hand sanitizer formulations.

We screened hydrophobic comonomers with differing alkyl side-chains and degrees of branching because such structural features are known to influence the surface activity of small-molecule surfactants. Meanwhile, hydrophilic components were selected from readily available non-ionic or anionic comonomers. With the exception of a single acrylate–acrylamide copolymer, synthones were performed using all-acrylic (or all-methacrylic) formulations to ensure comparable copolymer reactivities and hence the formation of statistical (rather than blocky or gradient) copolymers.

Comonomer feed mass fractions ranging from 0.40 to 0.60 were targeted to generate a range of HLB values, with $^{1}H$ NMR analysis confirming that the final copolymer compositions were close to those used in the feed (Figure S1 and Table S1). The resulting copolymers are denoted using the convention $M_xA-N$, where $M$ and $N$ refer to the hydrophobic and hydrophilic monomers, respectively, $x$ is the mass fraction of the hydrophobic copolymer used in the feed, and $A$ is the copolymer architecture (either statistical [stat] or block).

To assess the surface activity of each surfactant in various ethanol–water mixtures, we designed a high-throughput assay that required minimal copolymer. This protocol involved dissolving each copolymer in pure ethanol at 0.5% w/v prior to the successive addition of known aliquots of ultrapure water, with foamability being measured by vortex agitation at each ethanol–water composition. Foam stabilization was initially examined in pure ethanol ($\gamma_{\text{solvent/air}} = 22.3 \text{ mN m}^{-1}$) and then for a series of ethanol–water mixtures containing 11, 20, 27, 34, 39, or 47% w/w water (at the highest water content $\gamma_{\text{solvent/air}} = 29.0 \text{ mN m}^{-1}$). Although the copolymer concentration necessarily decreases across this series, stronger foaming was typically observed at higher water contents (which correspond to higher $\gamma_{\text{solvent/air}}$ values). The foam height ratio was used as a proxy for foamability, while foam lifetime was used as a metric for foam stability (see the Experimental section). Representative digital photographs of foams generated using the best-performing statistical copolymer surfactant are shown in Figure S2, alongside other copolymer surfactant foams for comparison.

The foam stabilization assay enables highly surface-active formulations to be rapidly identified, thus circumventing the need for laborious surface tension measurements on more than 30 copolymers. This assay also enables the comparison of these new statistical copolymer surfactants with commercially available hydrocarbon-based surfactants such as Triton X-100, sodium dodecyl sulfate, and Pluronic F127. The three commercial surfactants did not produce stable foams for any of the ethanol–water mixtures investigated in this study. Thus, any statistical copolymer surfactant producing a stable foam under such conditions clearly exhibits strong surface activity well beyond that of typical hydrocarbon-based surfactants. In this context, we note that silicone-based block copolymers are not expected to offer environmental advantages in various commercial applications.

### RESULTS AND DISCUSSION

**Screening Binary Combinations of Hydrophilic and Hydrophobic Comonomers.** During the synthesis of amphiphilic statistical copolymers via free-radical copolymerization in ethanol–water mixtures, we observed that the resulting reaction mixtures were prone to foaming when subjected to manual agitation (hand shaking), with similar surface activity being observed in the presence of organic solvents such as THF during work-up and purification. This serendipitous discovery led us to investigate the surfactant properties of amphiphilic statistical copolymers in binary ethanol–water mixtures relevant to hand sanitizer formulations. Conventional free-radical polymerization (FRP) is particularly well-suited for the synthesis of statistical copolymers using a wide range of (meth)acrylic and (meth)acrylamide monomers. Their broad commercial availability means that such monomers are applicable to high-throughput syntheses to generate libraries of amphiphilic copolymer surfactants. Accordingly, we performed a series of FRP syntheses employing numerous combinations of hydrophilic and hydrophobic comonomers (Scheme 1 and Table 1).

We screened hydrophobic comonomers with differing alkyl side-chains and degrees of branching because such structural features are known to influence the surface activity of small-molecule surfactants. Meanwhile, hydrophilic components were selected from readily available non-ionic or anionic comonomers. With the exception of a single acrylate–acrylamide copolymer, synthones were performed using all-acrylic (or all-methacrylic) formulations to ensure comparable copolymer reactivities and hence the formation of statistical (rather than blocky or gradient) copolymers.

Comonomer feed mass fractions ranging from 0.40 to 0.60 were targeted to generate a range of HLB values, with $^{1}H$ NMR analysis confirming that the final copolymer compositions were close to those used in the feed (Figure S1 and Table S1). The resulting copolymers are denoted using the convention $M_xA-N$, where $M$ and $N$ refer to the hydrophobic and hydrophilic monomers, respectively, $x$ is the mass fraction of the hydrophobic copolymer used in the feed, and $A$ is the copolymer architecture (either statistical [stat] or block).

To assess the surface activity of each surfactant in various ethanol–water mixtures, we designed a high-throughput assay that required minimal copolymer. This protocol involved dissolving each copolymer in pure ethanol at 0.5% w/v prior to the successive addition of known aliquots of ultrapure water, with foamability being measured by vortex agitation at each ethanol–water composition. Foam stabilization was initially examined in pure ethanol ($\gamma_{\text{solvent/air}} = 22.3 \text{ mN m}^{-1}$) and then for a series of ethanol–water mixtures containing 11, 20, 27, 34, 39, or 47% w/w water (at the highest water content $\gamma_{\text{solvent/air}} = 29.0 \text{ mN m}^{-1}$). Although the copolymer concentration necessarily decreases across this series, stronger foaming was typically observed at higher water contents (which correspond to higher $\gamma_{\text{solvent/air}}$ values). The foam height ratio was used as a proxy for foamability, while foam lifetime was used as a metric for foam stability (see the Experimental section). Representative digital photographs of foams generated using the best-performing statistical copolymer surfactant are shown in Figure S2, alongside other copolymer surfactant foams for comparison.

The foam stabilization assay enables highly surface-active formulations to be rapidly identified, thus circumventing the need for laborious surface tension measurements on more than 30 copolymers. This assay also enables the comparison of these new statistical copolymer surfactants with commercially available hydrocarbon-based surfactants such as Triton X-100, sodium dodecyl sulfate, and Pluronic F127. The three commercial surfactants did not produce stable foams for any of the ethanol–water mixtures investigated in this study. Thus, any statistical copolymer surfactant producing a stable foam under such conditions clearly exhibits strong surface activity well beyond that of typical hydrocarbon-based surfactants. In this context, we note that silicone-based block copolymers are
typically used to generate foams from the ethanol-rich ethanol—water mixtures required for effective hand sanitizer formulations.\(^5\)

A summary of the initial foaming assay data obtained for statistical copolymer surfactants is presented in Table 1. Entries are shaded using a color scale to indicate enhanced foamability (or foam height ratio, \(F_x\) and foam lifetime (\(t_x\) seconds). Clearly, the nature of the hydrophilic and hydrophobic acrylic comonomers has a significant influence on the foam stabilization performance of these copolymer surfactants in various ethanol—water compositions. For example, the P(BuA-stat-PEGA) copolymer produced no discernible foam for any binary solvent mixture, while P(LA-stat-PEGA), P(OA-stat-AA), and P(OAA-stat-AA) copolymers only produced foams at the highest water content (47% w/w). The remaining copolymer surfactants can be ranked according to their relative foamability and foam stability performance across the range of ethanol—water mixtures as follows: P(TMHA-stat-PEGA) > P(OAA-stat-PEGA) > P(TMHA-stat-AA) > P(OA-stat-PEGA) > P(EHA-stat-PEGA) > P(BoA-stat-PEGA).

P(TMHA-stat-PEGA) copolymers significantly outperformed all other surfactants in terms of their foamability and foam stability. Thus, we hypothesized that using TMHMA, which is the methacrylic analogue of TMHA, should lead to similar performance for the analogous all-methacrylic statistical copolymer surfactants. Accordingly, P(TMHMA-stat-PEGMA), P(TMHMA-stat-GMA), and P(TMHMA-stat-MAA) were prepared targeting the same copolymer architectures and evaluated using the same foaming assay (see blue entries in Table 1). Surprisingly, each of these copolymers exhibited significantly lower foamability than the equivalent all-acrylic copolymer. For example, the P(TMHMA-stat-GMA) and P(TMHMA-stat-PEGMA) copolymers only displayed foamability at higher water contents (43—47% w/w). We will revisit these unexpected observations later.

Having observed inferior foamability for AA- or MAA-based surfactants relative to neutral PEGA-based copolymers, we sought to understand how surface activity varied with solution pH and ionic strength. Thus, an additional series of foaming assays was performed for P(TMHA\(_{0.5}\)-stat-PEGA), P(TMHA\(_{0.5}\)-stat-AA), and P(TMHA\(_{0.5}\)-stat-MAA) in which the aqueous component of the ethanol—water mixture comprised either 1.0 M NaCl or was adjusted to either pH 3 or pH 10 (using HCl or NaOH, respectively). Compared to the assay involving ultrapure water (Table 1), the foam stability of P(TMHA\(_{0.5}\)-stat-PEGA) was marginally reduced when adjusting the aqueous component pH to pH 10 but remained essentially unchanged in the presence of 1.0 M NaCl or after adjusting the aqueous component pH to pH 3 (Table S2). On the other hand, the foamability conferred by P(TMHA\(_{0.5}\)-stat-AA) was marginally reduced in the presence of 1.0 M NaCl or at pH 3 but remained essentially unchanged at pH 10. Finally, P(TMHMA\(_{0.5}\)-stat-MAA) could only stabilize foam in the presence of 1.0 M NaCl. Overall, the poor foamability of copolymers containing ionizable AA or MAA repeat units was not significantly enhanced whether these moieties are fully protonated (pH 3), fully deprotonated (pH 10), or charge-screened (1.0 M NaCl). This suggests that the surface activity is not significantly affected by the degree of ionization of the carboxylic acid repeat units under these conditions. Moreover, P(TMHA\(_{0.5}\)-stat-PEGA) remained highly surface-active and was able to stabilize foams in the presence of each type of modified aqueous phase (Table S2).

Furthermore, similar foaming performance was obtained even after the deliberate addition of up to 8% comonomer by mass, which suggests that foaming is not caused by small-molecule impurities (Table S3). The non-ionic nature of this copolymer surfactant ensures good performance over a range of conditions, which is likely to be an advantage for many industrial applications. Finally, P(TMHA-stat-PEGA) copolymers were also prepared containing TMHA mass fractions of either 0.25 or 0.75 but in each case, the copolymer foamability was significantly poorer than that observed for the formulations shown in Table 1.

Each of the statistical copolymers presented in Table 1 was prepared by FRP, as indicated by their relatively high dispersions (\(D = 1.9—2.4\), see Table S1). Further copolymers [hereinafter denoted as R-P(TMHA\(_{0.4,0.5,0.6}\)-stat-PEGA)] were prepared using reversible addition–fragmentation chain transfer (RAFT) polymerization to understand the influence of chain length distribution on foam stabilization performance. GPC analysis of RAFT-synthesized copolymers confirmed that they possessed significantly narrower molecular weight distributions relative to FRP-synthesized copolymers (Figure S3A). Foam stabilization assays using these lower dispersity copolymers indicated similar foamability and foam stability as that obtained using P(TMHA\(_{0.4,0.5,0.6}\)-stat-PEGA) (see Table 2). This suggests that foamability is insensitive to the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions. RAFT polymerization also provides good control over the copolymer molecular weight distribution under these conditions.
marginaly lower foamability for all ethanol–water compositions investigated, suggesting slightly lower surface activity. Clearly, using RAFT polymerization to gain better control over the copolymer $M_n$ and produce narrower molecular weight distributions does not lead to superior performance for P(TMHA-stat-PEGMA) copolymers in foam stabilization experiments.

There are relatively few studies of the effect of copolymer architecture on surface activity that explicitly compare statistical and block copolymer architectures.\textsuperscript{25–28} Herein, we prepared diblock copolymer analogues (i.e., PTMHA\textsubscript{0.4–0.6}–block–PPEGA) of the most surface-active statistical copolymers P(TMHA\textsubscript{0.4–0.6}–stat–PEGMA) using RAFT polymerization.\textsuperscript{7,8} Sequential monomer addition enabled the production of well-defined diblock copolymers with $M_n$ values of 11.8–12.9 kg mol\textsuperscript{−1} and relatively narrow molecular weight distributions ($D = 1.2–1.3$), as judged by GPC analysis (see Figure S4 and Table S1). A relatively low copolymer $M_n$ was targeted in view of observations made by Matsuoka et al., who found that diblock copolymers with $M_n > 15$ kg mol\textsuperscript{−1} became surface-inactive.\textsuperscript{11,12} Furthermore, PTMHA\textsubscript{0.4}–block–PPEGA, PTMHA\textsubscript{0.5}–block–PPEGA, and PTMHA\textsubscript{0.6}–block–PPEGA exhibited comparable GPC $M_n$ values to those obtained for the polydisperse statistical copolymers prepared by conventional FRP ($M_n = 11.5–13.2$ kg mol\textsuperscript{−1}). Nevertheless, foam stabilization assays conducted in ethanol–water mixtures using PTMHA\textsubscript{0.4–0.6}–block–PPEGA indicated very poor foamability for all copolymer compositions compared to P/(TMHA\textsubscript{0.4–0.6}–stat–PEGMA). Foamability was only observed for PTMHA\textsubscript{0.6}–block–PPEGA for the most water-rich compositions (see Table 2), while foam heights and lifetimes were significantly lower than those observed for the equivalent statistical copolymer.

**Surface Tension Analysis.** According to the foamsability assays, several statistical copolymer surfactants can stabilize long-lasting foams in low surface energy solvents (<29 mN m\textsuperscript{−1}). Notably, the analogous diblock copolymers, and also a range of commercial surfactants, do not perform as effectively under such conditions. This suggests that these statistical copolymer surfactants are more surface-active than many other hydrocarbon-based surfactants. In several prior studies, tensiometry was used to show that block copolymers reduced the interfacial tension ($\gamma_{\text{solvent/air}}$) to a greater extent than statistical/random copolymers, thus indicating superior surfactant performance.\textsuperscript{7–9}

To correlate performance in the foamability assay with surface activity, we conducted a series of ring tensiometry measurements using P(TMHA\textsubscript{0.5}–stat–PEGMA) and PTMHA\textsubscript{0.5}–block–PPEGA in various ethanol–water mixtures (Figure 1A). Copolymer stock solutions were prepared at 5.0 g dm\textsuperscript{−3} in ethanol and $\gamma_{\text{solvent/air}}$ was determined at 25 °C in various ethanol–water mixtures prepared by serial dilution with water to replicate the foaming assay conditions.

At the highest ethanol mass fractions (79–100% w/w), $\gamma_{\text{solvent/air}}$ was only marginally lower relative to the pure binary solvent mixtures for each copolymer (Figure 1A). However, at 57% w/w ethanol (for which we measured $\gamma_{\text{solvent/air}} = 26.21 \pm 0.03$ mN m\textsuperscript{−1}), the statistical copolymer lowered the surface tension to 24.56 $\pm$ 0.01 mN m\textsuperscript{−1}, whereas the diblock copolymer only led to a reduction to 25.26 $\pm$ 0.03 mN m\textsuperscript{−1}. This observation is consistent with the foaming assay: only the statistical copolymer was able to produce a stable foam at this ethanol–water composition. The difference in $\gamma_{\text{solvent/air}}$ between the statistical and diblock copolymers gradually increased with addition of further water. At 44% w/w ethanol (not studied in the foamsability assay), PTMHA\textsubscript{0.5}–block–PPEGA produced the same $\gamma_{\text{solvent/air}}$ as that for the pure solvent mixture (ca. 28.2 mN m\textsuperscript{−1}). This implies that these copolymer chains no longer occupied the interface under such conditions.

Surface tensiometry measurements were also conducted for copolymer concentrations of 0.1–5.0 g dm\textsuperscript{−3} to identify the critical micellar concentration (CMC) for P(TMHA\textsubscript{0.5}–stat–PEGMA) and PTMHA\textsubscript{0.5}–block–PPEGA in an ethanol–water composition containing 61.2% w/w ethanol (Figure 1B). The diblock copolymer reduced $\gamma_{\text{solvent/air}}$ between 2.7 and 5.0 g dm\textsuperscript{−3}, but for lower copolymer concentrations, $\gamma_{\text{solvent/air}}$ was equivalent to that of the pure solvent mixture (25.00 $\pm$ 0.02 mN m\textsuperscript{−1}). These two distinct regimes suggest that the diblock copolymer exhibits a CMC between 2.0 and 2.7 g dm\textsuperscript{−3}, while above this concentration, the solvent–air interface is saturated with copolymer chains and micelles are present in the bulk solution. Notably, the statistical copolymer reduced $\gamma_{\text{solvent/air}}$ by ca. 0.1–0.2 mN m\textsuperscript{−1} more than the diblock copolymer at every concentration investigated. Moreover, the former
copolymer exhibits no CMC over this concentration range and remained surface-active even at 0.1 g dm\(^{-3}\).

Bubble pressure tensiometry (BPT) measurements performed at 4.0 g dm\(^{-3}\) in the presence of 61.2% w/w ethanol (Figure 1C) enabled comparison of the dynamic interfacial behavior of these surfactants. Statistical copolymer surfactants produced a reduction in \(\gamma_{\text{solvent/air}}\) even for short surface aging times (<30 ms) and maintained relatively low \(\gamma_{\text{solvent/air}}\) values at longer surface aging times (>500 ms). These data suggest that statistical copolymer surfactants adsorb much faster at the solvent/air interface than block copolymer surfactants. To understand how the copolymer morphology in solution influences the chain dynamics, we studied various copolymer solutions using SAXS.

**Bulk Solution and Interfacial Self-assembly.** The consistently larger reduction in \(\gamma_{\text{solvent/air}}\) by P(TMHA\(_{0.5}\)-stat-PEGA) indicates that these copolymer chains adsorb more strongly at the interface than the analogous diblock copolymer chains. Establishing a sufficient surface tension gradient at the interface is a prerequisite for foam stabilization\(^{30-33}\) and accounts for the enhanced performance of the statistical copolymer in ethanol–water mixtures. The self-assembly of amphiphilic statistical copolymers in selective solvents is less well understood.\(^{30-33,47}\) The presence of hydrophilic and hydrophobic repeat units within the same copolymer chain results in intramolecular chain-folding such that hydrophilic pendant groups are presented at the copolymer/solvent interface, while hydrophobic groups remain buried within the cores. Importantly, this can lead to the formation of micelles with relatively low aggregation numbers; indeed, unimolecular aggregates can be obtained under certain conditions.\(^{31,33}\)

Clearly, these two distinct self-assembly mechanisms must influence the propensity of copolymer chains to populate an interface within a given timescale.

We studied the self-assembly behavior of the P(TMHA\(_{0.5}\)-stat-PEGA) and PTMHA\(_{0.5}\)-block-PEGA copolymers above the CMC of the latter (3.0% w/w) and at an intermediate ethanol fraction (61.2% w/w). Under such conditions, only the statistical copolymer formed a stable foam. SAXS was used to characterize the micelles that are formed in solution (Figure 2). The presence of intensity minima (\(q_{\text{min}}\)) in both scattering patterns indicated the presence of nanoparticles, while the low \(q\) gradient of approximately zero indicated a spherical morphology in both cases. The significantly higher \(q_{\text{min}}\) observed for P(TMHA\(_{0.5}\)-stat-PEGA) compared to PTMHA\(_{0.5}\)-block-PEGA suggests significantly different self-assembly behavior.
A satisfactory fit to the SAXS pattern recorded for the former copolymer was obtained using a well-known spherical micelle model \(^{48}\) by fitting the core radius based on an individual hydrophobic repeat unit and the shell \(R_s\) value based on the expected \(R_g\) of the PEG side-chain \(^{31}\) (see the Supporting Information for model parameters). It is assumed that the micelle shell mainly comprises solvated PEG side chains with low X-ray contrast, thus scattering is dominated by the hydrophobic TMHA-based micelle core. This SAXS model indicated the presence of spherical micelles with a mean core radius, \(R_c\), of 1.5 ± 0.7 nm, which is consistent with the size of unimolecular micelles formed by other amphiphilic statistical copolymers reported in the literature. \(^{31,35}\) The aggregation number (\(N_{agg}\)) could be estimated by comparing the micellar volume measured by SAXS with the expected volume occupied by a single copolymer chain. Using the \(M_w\) value determined by GPC, an \(N_{agg}\) of 1.18 was calculated (see the Supporting Information). The use of poly(ethyl methacrylate) calibration standards for the GPC measurements incurs a systematic error in \(M_w\) which most likely accounts for the modest deviation of \(N_{agg}\) from unity. Thus, these data suggest that P(TMHA\(_{0.5}\)-stat-PEGA) forms unimolecular micelles in ethanol–water mixtures. Using the same spherical micelle model, but with the core radius based on a PTMHA block, PTMHA\(_{0.5}\)-block-PEGA micelles were estimated to have a core radius of 6.3 ± 1.0 nm and an \(N_{agg}\) of 92, which are physically reasonable values for such an amphiphilic diblock copolymer. \(^{50}\)

To produce an air-in-liquid foam, sufficient surfactant must adsorb to create a surface tension gradient and stabilize the new interfacial area that is generated. For conventional surfactants, micelles must first dissociate to form individual solvent−air interface. Such micelle dissociation events introduce a relatively high energetic penalty. \(^{31}\) Alternatively, the relatively large diblock copolymer micelles could diffuse slowly to the interface and undergo adsorption-induced dissociation. Once adsorbed, the hydrophobic tails are oriented toward the gas phase, while hydrophilic head-groups remain solvated within the near-surface of the interface. However, the foam stabilization mechanism for the statistical copolymer surfactants reported herein involves a different mechanism. In this case, individual copolymer chains form micelles by chain folding such that the hydrophobic groups are located within the cores, while hydrophilic PEG groups form the solvated micelle shell (see the schematic cartoon shown in Figure 2). Such unimolecular micelles must undergo a conformational transition at the interface to enable the buried hydrophobic TMHA groups to interact with the gas phase, while the pendant PEG groups remain solvated in the aqueous milieu. Compared to multimolecular micelles formed by the diblock copolymer, unimolecular micelles can rapidly diffuse to the interface. Moreover, while the former must first dissociate into unimers, statistical copolymer chains can simply undergo a conformational transition at the interface to present their hydrophobic groups (Figure 2). This rearrangement has an entropic penalty associated with the transition from collapsed and compact chains to stretched conformations. However, this is more than offset by the gain in favorable enthalpic interactions, which enables efficient foam stabilization even with relatively low energy input (e.g., manual hand shaking).

Further evidence for the unfolding mechanism of statistical copolymer surfactants is provided by considering their chain mobility, for which the copolymer \(T_g\) is a proxy. Differential scanning calorimetry (DSC) studies indicate a \(T_g\) of −64 °C for P(TMHA\(_{0.5}\)-stat-PEGA) (see Figure S5A). Unfortunately, no \(T_g\) could be determined for the corresponding P(TMHA\(_{0.5}\)-stat-PTMHA) copolymer. However, DSC analysis of PTMHA and P(TMHA) homopolymers confirmed a significantly higher \(T_g\) for the methacrylic analogues (Figure S5B). SAXS analysis confirms that P(TMHA\(_{0.5}\)-stat-PEGA) also forms unimolecular micelles in ethanol−water mixtures (Figure S6) but surface tensiometry measurements indicate that this copolymer does not reduce the surface tension over the full range of ethanol−water mixtures studied (Figure 1D). Nevertheless, P(TMHA-stat-PEGMA) copolymer were still able to stabilize foams at higher water contents, although foamability and foam stability were much lower than those observed for P(TMHA-stat-PEGA) copolymers (Table 1). We hypothesize that in the case of P(TMHA\(_{0.5}\)-stat-PEGMA), chains can only occupy the interface when a significant amount of energy is applied (i.e., vortex mixing) to drive the transition from compact unimolecular micelles to an extended conformation at the interface (as shown in Figure 2). Given the lower flexibility of the methacrylic backbone, \(^{32}\) this transition cannot occur under the static conditions used in tensiometry. Clearly, chain mobility is a critical parameter that significantly affects the performance of statistical copolymer surfactants. This important insight also explains why P(OA-stat-PEGA), another copolymer containing a high \(T_g\) hydrophobic comonomer, \(^{52}\) is a much less effective foam stabilizer than P(TMHA\(_{0.5}\)-stat-PEGA).

According to the surface tensiometry data, the trimethyl-hexyl (TMH) groups within statistical copolymers can adsorb more efficiently and more rapidly at the interface than those within the analogous diblock copolymer chains. Hence, statistical copolymers produced lower surface tensions at all concentrations and for most ethanol−water mixtures. For maximum surface coverage by P(TMHA-block-PEGMA), the PTMHA chain should adsorb parallel to the interface with each of its TMH groups directed toward the air phase. However, such a conformation would incur a significant enthalpic penalty. It is much more likely that the PTMHA chain collapses to form a desolvated coil (Figure 2), hence a suboptimal proportion of TMHA groups is located at the interface. In contrast, the statistical copolymer chain can readily adopt an extended conformation such that the majority of its pendant hydrophobic TMH groups are adsorbed at the air phase, while its hydrophilic PEG groups remain solvated within the near-surface of the solution. This produces a significantly higher interfacial area per molecule for P(TMHA-stat-PEGA) compared to P(TMHA-block-PEGMA) and therefore enables stabilization of a greater foam volume.

Notably, similar micellar dimensions were observed for several other statistical copolymer surfactants examined in this study (e.g., P(TMHA\(_{0.5}\)-stat-PEGMA), P(LA\(_{0.5}\)-stat-PEGA), P(OA\(_{0.5}\)-stat-PEGMA), and R-P(TMHA\(_{0.5}\)-stat-PEGA), Figure S6). This suggests that a wide range of non-ionic statistical copolymer surfactants can form unimolecular micelles. However, given the much lower foamability observed for P(TMHA\(_{0.5}\)-stat-PEGMA), P(LA\(_{0.5}\)-stat-PEGA), and P-(OA\(_{0.5}\)-stat-PEGMA), such self-assembly behavior is not the sole prerequisite for interfacial activity. Empirically, we find that the most efficient surfactants contain hydrophobic comonomers comprising more than two methyl groups per repeat unit (i.e., TMHA, OA, and OAA). Prior studies of both
hydrocarbon and silicone-based small-molecule surfactants indicate that more branched hydrophobic moieties lower the surface tension to a greater extent.\textsuperscript{41,53} This is believed to be related to weaker tail–tail interactions and the large volumes occupied by bulky hydrophobic tails at an interface.\textsuperscript{54} The length of the spacer group between the backbone and methyl groups also appears to be an important determinant of surface activity: copolymers bearing methyl-rich \textit{BuA} repeat units cannot stabilize foams in ethanol–water mixtures (Table 1). Notably, our hydrocarbon-based surfactants compare favorably to perfluorinated surfactants as foam stabilizers for ethanol–water mixtures. More specifically, for a comparable surfactant concentration in 50\% v/v ethanol, our optimized P(TMHA-stat-PEGA) copolymer surfactant reduced the surface tension to 25.4 mN m\textsuperscript{-1} (vs 28 mN m\textsuperscript{-1} for a perfluorinated surfactant).

Moreover, this hydrocarbon surfactant also produced more stable foams. Interestingly, this unfolding mechanism resembles how some protein-stabilized foams are generated in nature:\textsuperscript{55} when subjected to mechanical stimulation (such as the “whipping” mechanism used by tree frogs\textsuperscript{11}), globular proteins can unfold in solution to expose buried hydrophobic amino acids that stabilize the air–water interface to produce long-lasting foams. We also note that the molecular weight of these statistical copolymers is within the range determined for hydrophobins (\textless 20 kg mol\textsuperscript{-1}), which are highly surface-active proteins found in filamentous fungi.\textsuperscript{59} As far as we are aware, this mechanism of foam stabilization has not been previously reported for synthetic copolymers. In principle, scanning methods such as small-angle neutron scattering\textsuperscript{60} could provide further insight regarding the interfacial copolymer structure and foam stabilization mechanism.

\section*{Conclusions}

According to various studies, block copolymers are preferred to statistical copolymers for surfactant performance.\textsuperscript{25–28} Nevertheless, herein we show that judicious selection of pairs of acrylic comonomers leads to superior surface activity for statistical copolymers. The influence of various structural parameters (e.g., the chemical nature of the hydrophobic and hydrophilic comonomers, copolymer composition, \textit{M}_\text{c}, copolymer dispersity, and copolymer chain mobility) on the foam stabilization performance of a series of copolymer surfactants was examined. A foamability assay was employed to rapidly identify the most appropriate comonomer pairs, with the chemical nature of the hydrophobic comonomer found to strongly influence surface activity. The strong foamability exhibited by P(TMHA-stat-PEGA) copolymers suggests that the combination of a highly branched hydrophobic comonomer with a non-ionic hydrophilic comonomer is optimal.

The high surface activity demonstrated herein for certain statistical copolymers is related to their highly mobile all-acrylic backbones and their formation of unimolecular micelles in solution. This facilitates fast diffusion to the air–water interface and rapid conformational rearrangement of each copolymer chain to enable its hydrophobic comonomer units to interact with the air phase. In contrast, multimolecular block copolymer micelles must either dissociate into unimers that then diffuse to the surface or whole micelles must slowly diffuse to the air–water interface prior to micellar adsorption.\textsuperscript{61} Thus, amphiphilic statistical copolymers exhibit superior foam stabilization for a range of ethanol-rich ethanol–water mixtures that are relevant for hand sanitizer formulations. Indeed, statistical copolymer surfactants can stabilize high-quality foams when extruding hand sanitizer formulations through a model soap dispenser (see Video S1).

In nature, evolutionary design has produced various highly surface-active proteins comprising hydrophilic and hydrophobic amino acids distributed along polypeptide backbones in non-blocky sequences. Herein, we show that the informed design of new hydrocarbon-based amphiphilic statistical copolymers can also lead to remarkable surface activity. Further optimization of the synthetic protocol (e.g., monomer feeding protocols, temperatures, initiator systems) is expected to produce further performance enhancements that may even exceed that of natural proteins. Importantly, these synthetic copolymer surfactants offer both cost and environmental advantages over the fluorocarbon- and siloxane-based surfactants that are currently widely used in the chemical industry.
eluent. Following vacuum treatment, the residual monomer was less than 2 mol % in all cases.

Synthesis of a PTMHA-block-PEGA Diblock Copolymer by RAFT Solution Polymerization. AIBN (2.5 mg) and DDMAT (11 mg) were weighed into a 7 mL glass vial, followed by TMHA (0.25 g) and THF (0.50 mL). A magnetic flea was added to this vial, which was sealed using a rubber septum. After vortex mixing for 10 s to achieve a homogeneous solution, the reaction mixture was degassed on ice by sparging with nitrogen gas for 20 min. The vial was then immersed in an oil bath at 70 °C and the reaction mixture was stirred for 3 h. Then, the vial was removed from the oil bath, and an aliquot was extracted from the reaction solution for 1H NMR studies in CDCl₃ and GPC analysis (THF eluent). The residual monomer was less than 5%. PEGA (0.25 g) and THF (0.80 mL) were then added to the vial and the mixture was degassed on ice by sparging with nitrogen gas for 20 min. The vial was then submerged in an oil bath at 70 °C and the reaction mixture was stirred for a further 10 h. After removing the vial from the oil bath, the solvent and any unreacted monomer were removed overnight at 70 °C using a vacuum oven. The mean composition of the diblock copolymer was determined by 1H NMR spectroscopy analysis in CDCl₃ and its molecular weight and dispersity were assessed by GPC using THF eluent. Following vacuum treatment, the residual monomer was less than 2% in all cases.

Analytical Techniques. Following copolymer dissolution in CDCl₃, 1H NMR spectra were recorded using a Bruker AV1-400 MHz spectrometer with 64 scans being averaged per spectrum. Copolymer compositions were calculated in terms of the mass fraction of the hydrophobic comonomer (e.g., m_TMHA) by comparing the integrals for the oxymethylene ester groups in the (meth)acrylate repeat units (eq 1), which were located at sufficiently different chemical shifts for the hydrophilic (e.g., I_PEGA) and hydrophobic (e.g., I_TMHA) monomers.

$$m_{TMHA} = \frac{I_{TMHA} \times M_{TMHA}}{(I_{TMHA} \times M_{TMHA}) + (I_{PEGA} \times M_{PEGA})}$$ (1)

GPC analysis was conducted in THF at 30 °C (containing 2.0% v/v triethylamine and 0.05% w/v 3,5-di-tert-4-butylhydroxytoluene) using a flow rate of 1.0 mL min⁻¹. The GPC system comprised two Polymer Laboratories PL gel S µm Mixed C columns, an LC20AD ramped isocratic pump, and a Wellchrom K-2301 refractive index detector operating at 950 ± 30 nm. Calibration was achieved using a series of near-monodisperse poly(methyl methacrylate) standards ranging from M₉ ≈ 1.280–330,000 g mol⁻¹.

Foamability Assay. After purification under vacuum, the dry copolymer (5.0 mg) was weighed into a 7 mL glass vial and dissolved in ethanol (1.00 mL). The only exceptions were P(OAA-stat-PEG) and P(OAA-stat-AA), for which an aliquot of reaction solution (30 µL) following copolymerization was diluted with ethanol (970 µL). The ethanolic solution was agitated at 2850 rpm for 10 s using a vortex mixer (Cole-Palmer Vortex Mixer) and inspected for the appearance of bubbles. Ultrapure water, water adjusted to either pH 3 or 10, or 1 M NaCl, was then added in 0.10 mL aliquots, and the extent of foaming was evaluated by vigorous vortex mixing for 10 s. If the foam lifetime exceeded 5 s following agitation, then the foam height (h) from the top of the liquid to the top of the foam was measured five times and the maximum height was recorded (excluding any particularly large bubbles). To compare data for different experiments, the foamability (F) was calculated using eq 2, where V is the initial volume of surfactant solution and r is the vial radius. Thus, the foamability, or foam ratio, is simply given by the foam volume normalized with respect to the original solution volume.

$$F = \frac{h \pi r^2}{V}$$ (2)

Foam stability was arbitrarily defined as the time taken for the foam to break up to produce fewer than five bubbles at the surface of the liquid.

To test whether residual comonomers (<2% in all cases) had any effect on foamability, P(TMHAn₃-stat-PEGn) (which contained no detectable comonomer by 1H NMR spectroscopy) was deliberately contaminated with approximately 8% w/w PEGA and TMHA and then subjected to the same foaming assay. Essentially the same foamability and foam stability data were obtained within experimental error, suggesting that the surface activity of this copolymer was not affected by the residual comonomer.

Surface Tensiometry. Surface tension measurements were performed using the Du Noüy ring method on an automated Lauda TD3 ring/plate tensiometer equipped with a 90:10 platinum/iridium ring (ring radius = 0.955 cm). Solvent mixtures or surfactant solutions (40 mL) were added to a glass beaker and at least 10 measurements were recorded for each sample at 25 °C until a standard deviation of less than 0.07 mN m⁻¹ was obtained. For each copolymer, 200 mg was dissolved in ethanol (40 mL) and the surface tension of this solution was determined in the glass vessel. Following measurement, the copolymer solution was decanted into a jar, diluted with 8 mL water, and thoroughly mixed. The new solution (40 mL) was returned to the glass vessel for a second surface tension measurement (while maintaining the remaining solution in the jar). This protocol was repeated until the solution in the jar comprised 40 mL of water added to the initial 40 mL of ethanol (i.e., ethanol at +41.1% w/w). Concentration-dependent measurements were conducted by preparing an initial 5.0% w/w copolymer solution in an ethanol–water mixture containing 61.2% w/w ethanol, which was sequentially diluted using the same ethanol–water mixture, as described above.

Dynamic Surface Tensiometry. Measurements were conducted using a Krüss BP100 instrument using disposable polypropylene capillaries. The capillary radius was initially calibrated against ultrapure water. Dynamic surface tensions were determined on time scales ranging from 10 to 1000 ms and 10 measurements were averaged for time points. A thermostatted water bath was used to maintain the temperature at 20 °C. Dilute solutions were prepared by dissolving 4.0 g dm⁻³ copolymer in 50 mL of an ethanol/water mixture comprising 61.2% w/w ethanol. The copolymer was dissolved by vigorous stirring and each copolymer solution was allowed to equilibrate for at least 1 h prior to measurements.

Small-Angle X-ray Scattering Analysis. SAXS analyses of copolymer dispersions were conducted using a Xeuss 2.0 (Xenocs) SAXS instrument equipped with a FOX 3D multilayered X-ray mirror. X-rays were generated from a liquid gallium MetalJet X-ray source (Excillum, λ = 1.34 Å) and collimated using two sets of scatterless slits. The scattering intensity was measured on a hybrid pixel area detector (Pilatus 1M, Dectris) at a sample-to-detector distance of approximately 1.20 m (calibrated using a silver behenate standard). Two-dimensional (2D) SAXS patterns were reduced to 1D plots by azimuthal integration using a Foxrot software package.

SAXS Data Modeling. Particle radii were calculated using a simple scattering model for spheres reported in the literature, employing the parameters summarized in Table S4. The mean aggregation number (Nagg) was calculated from eq 3, i.e., from the ratio of the micelle core volume calculated from the particle radius measured by SAXS, R, to the theoretical micelle core volume, Vₙₜₐₘₜ, assuming that no solvent was located within the micellar core (which is reasonable given that PTMHA is insoluble in ethanol).

$$N_{agg} = \frac{3πR^3}{4V_{n_{_{TMHA}}}}$$ (3)

Vₙₜₐₘₜ could be estimated from $V_{n_{_{TMHA}}} = M_{TMHA}/(N_{X} \times \rho_{TMHA})$, where $M_{TMHA}$ is the calculated molecular weight of the TMHA fraction of the copolymer, $M_{TMHA} = M_{n} \times \rho_{TMHA}$, where $M_{n}$ is the number-average molecular weight measured by THF GPC and $\rho_{TMHA}$ is the TMHA mass fraction as measured by 1H NMR analysis. It was assumed that the density of the TMHA repeat units was equal to that of the monomer ($\rho_{TMHA} = 0.875$ g cm⁻³) for both the block and statistical copolymers. Although this assumption is a likely source of error in such calculations, in practice the polydispersity of the statistical copolymer chains and error arising from GPC analysis against PMMA standards are more significant contributions to the deviation in Nagg from unity.
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acsami.2c09910.

\(^1\)H NMR spectra for selected copolymers, GPC curves for P(TMHA-stat-PEGA) copolymers, representative digital photographs for foams, additional foamability assay data, raw SAXS data, DSC curves, and a video showing foam generation within a model foam dispenser when using a "hand sanitizer"-type copolymer solution (PDF)

Video demonstrating that statistical copolymer surfactants can stabilize high-quality foams when extruding hand sanitizer formulations through a model soap dispenser (AVI)

**REFERENCES**

1. Raffa, P.; Wever, D. A. Z.; Picchioni, F.; Broekhuis, A. A. Polymeric Surfaceactants: Synthesis, Properties, and Linksto Applications. Chem. Rev. 2015, 115, 8504–8563.

2. Torchilin, V. P. Structure and Design of Polymeric Surfaceactant-Based Drug Delivery Systems. J. Controlled Release 2001, 73, 137–172.

3. Laschewsky, A. Molecular Concepts, Self-Organisation and Properties of Polysoaps. In Polyoaps/Stabilizers/Nitrogen-15 NMR; Springer Berlin Heidelberg: Berlin, Heidelberg, 1995; pp 1–86.

4. Alexandridis, P.; Alan Hatton, T. Poly(Ethylene Oxide)—poly(Propylene Oxide)—poly(Ethylene Oxide) Block Copolymer Surfactants in Aqueous Solutions and at Interfaces: Thermodynamics, Structure, Dynamics, and Modeling. Colloids Surf., A 1995, 96, 1–46.

5. Schmolka, I. R. A Review of Block Polymer Surfactants. J. Am. Oil Chem. Soc. 1977, 54, 110–116.

6. Ryan, A. J.; Mai, S.-M.; Fairclough, J. P. A.; Hamley, I. W. Structures of Amphiphilic Block Copolymers in Their Liquid and Solid States Alexandridis, P.; Lindman, B. B.-T.- B. C., Eds.; Elsevier Science B.V.: Amsterdam, 2000; pp 151–167.

7. Moad, G.; Rizzardo, E.; Thang, S. H. RAFT Polymerization and Some of Its Applications. Chem. - Asian J. 2013, 8, 1634–1644.

8. Perrier, S. 50th Anniversary Perspective: RAFT Polymerization—A User Guide. Macromolecules 2017, 50, 7433–7447.

9. Ghosh, A.; Yusa, S. I.; Matsuoka, H.; Saruwatari, Y. Chain Length Dependence of Non-Surface Activity and Micellization Behavior of Cationic Amphiphilic Diblock Copolymers. Langmuir 2014, 30, 3319–3328.

10. Murugabooopathy, S.; Matsuoka, H. Surface Active to Non-Surface Active Transition and Micellization Behaviour of Zwitterionic Amphiphilic Diblock Copolymers: Hydrophobicity and Salt Dependenc. Polymers 2017, 9, No. 412.

11. Matsuoka, H.; Chen, H.; Matsumoto, K. Molecular Weight Dependence of Non-Surface Activity for Ionic Amphiphilic Diblock Copolymers. Soft Matter 2012, 8, 9140–9146.

12. Matsuoka, H.; Hachisuka, M.; Uda, K.; Onishi, T.; Ozoe, S. Why Ionic Amphiphilic Block Copolymer Can Be Non-Surface Active Comparison of Homopolymer, Block and Random Copolymers of Poly(styrene/2-sulfonate). Chem. Lett. 2012, 41, 1063–1065.

13. Bates, C. M.; Bates, F. S. 50th Anniversary Perspective: Block Polymers-Pure Potential. Macromolecules 2017, 50, 3–22.

14. Cooper, A.; Kennedy, M. W. Biofoams and Natural Protein Surfactants. Biophys. Chem. 2010, 151, 96–104.

15. Schor, M.; Reid, J. L.; MacPhee, C. E.; Stanley-Wall, N. R. The Diverse Structures and Functions of Surfactant Proteins. Trends Biochem. Sci. 2016, 41, 610–620.

16. Cooper, A.; Kennedy, M. W.; Fleming, R. I.; Wilson, E. H.; Videler, H.; Wokosin, D. L.; Su, T. J.; Green, R. J.; Lu, J. R. Adsorption of Frog Foam Nest Proteins at the Air-Water Interface. Biophys. J. 2005, 88, 2114–2125.

17. Cooper, A.; Vance, S. J.; Smith, B. O.; Kennedy, M. W. Frog Foams and Natural Protein Surfactants. Colloids Surf., A 2017, 534, 120–129.

18. Król, B.; Prochaska, K.; Chrzanowski, L. Biodegradability of Firefighting Foams. Fire Technol. 2012, 48, 173–181.

19. Rodriguez Patino, J. M.; Carrera Sánchez, C.; Rodriguez Niño, M. R. Implications of Interfacial Characteristics of Food Foaming Agents in Foam Formulations. Adv. Colloid Interface Sci. 2008, 140, 95–113.

20. Green, A. J.; Littlejohn, K. A.; Hooley, P.; Cox, P. W. Formation and Stability of Food Foams and Aerated Emulsions: Hydrophobs as Novel Functional Ingredients. Curr. Opin. Colloid Interface Sci. 2013, 18, 292–301.

21. Pegg, J. C.; Czajka, A.; Hill, C.; James, C.; Peach, J.; Rogers, S. E.; Eastoe, J. Alternative Route to Nanoscale Aggregates with a PH-Responsive Random Copolymer. Langmuir 2017, 33, 2628–2638.

22. Peng, F.; Ke, Y.; Lu, S.; Zhao, Y.; Hu, X.; Deng, Q. Anion Amphiphilic Random Copolymers and Their Performance as
Surfactants May Not Be Safe.

of Ingredients, Mechanisms of Action, Modes of Delivery, and Hand Hygiene Compliance.

Efficacy against Coronaviruses.

Impact of Hand Sanitizer Format (Gel/Foam/Liquid) and Dose

Copolymers into Size-Controlled and Thermoresponsive Micelles in Increasing Demand.

Biehl, R.; Allgaier, J. Controlled LCST Behavior and Structure Water.

Uniform Micelles.

Basheti, I. A.; Bonacucina, G.; Cespi, M.; Palmieri, G. F. Hand 2022

Consequences of the Monomer Sequence and Chain Flexibility on 2018,

Mykhaylyk, O. O. Control of Particle Size in the Self-Assembly of Macromolecules 2018, 51, 395–409.

Kostyurina, E.; De Mel, J. U.; Vasiyleva, A.; Krutova, M.; Frielinghaus, H.; Dulle, M.; Barnsley, L.; Förster, S.; Schneider, G. J.; Biehl, R.; Allgaier, J. Controlled LCST Behavior and Structure Formation of Alternating Amphiphilic Copolymers in Water. Macromolecules 2022, 55, 1552–1565.

Berardi, A.; Perinelli, D. R.; Merchant, H. A.; Bisharat, L.; Bashiti, I. A.; Bonacucina, G.; Cespi, M.; Palmieri, G. F. Hand Sanitizers amid CoVID-19: A Critical Review of Alcohol-Based Products on the Market and Formulation Approaches to Respond to Increasing Demand. Int. J. Pharm. 2020, 584, No. 119431.

Greenaway, R. E.; Ormandy, K.; Fellows, C.; Hollowood, T. Impact of Hand Sanitizer Format (Gel/Foam/Liquid) and Dose Amount on Its Sensory Properties and Acceptability for Improving Hand Hygiene Compliance. J. Hosp. Infect. 2018, 100, 195–201.

Golin, A. P.; Choi, D.; Ghahary, A. Hand Sanitizers: A Review of Ingredients, Mechanisms of Action, Modes of Delivery, and Efficacy against Coronaviruses. Am. J. Infect. Control 2020, 48, 1062–1067.

Mullin, C. A.; Fine, J. D.; Reynolds, R. D.; Fraizer, M. T. Toxicological Risks of Agrochemical Spray Adjuvants: Organosilicone Surfactants May Not Be Safe. Front. Public Health 2016, 4, No. 92.

Laubie, B.; Bonnafous, E.; Desjardin, V.; Germain, P.; Fleury, E. Silicone-Based Surfactant Degradation in Aqueous Media. Sci. Total Environ. 2013, 454–455, 199–205.

Cuzzucoli Crucitti, V.; Contreas, L.; Taresco, V.; Howard, S. C.; Dundas, A. A.; Limo, M. J.; Nisissak, T.; Williams, P. M.; Williams, P.; Alexander, M. R.; Wildman, R. D.; Muir, B. W.; Irvine, D. J. Generation and Characterization of a Library of Novel Biologically Active Functional Surfactants (Surfers) Using Combined High-Throughput Methods. ACS Appl. Mater. Interfaces 2021, 13, 43290–43300.

Alexander, S.; Smith, G. N.; James, C.; Rogers, S. E.; Guittard, F.; Sagisaka, M.; Eastoe, J. Low-Surface Energy Surfactants with Branched Hydrocarbon Architectures. Langmuir 2014, 30, 3413–3421.

Vazquez, G.; Alvarez, E.; Navaza, J. M. Surface Tension of Alcohol Water + Water from 20 to 50°C. J. Chem. Eng. Data 1995, 40, 611–614.

de Castro, M. T. F.; Koivisto, B. M.; Munoz, F. High Alcohol Content Foaming Compositions with Silicone-Based Surfactants. U.S. Patent 8309111B2, 2012.

Petkova, B.; Tcholakova, S.; Chenkova, M.; Golemanov, K.; Denkov, N.; Thorley, D.; Stoyanov, S. Foamability of Aqueous Solutions: Role of Surfactant Type and Concentration. Adv. Colloid Interface Sci. 2020, 276, No. 102084.

Mai, Y.; Eisenberg, A. Self-Assembly of Block Copolymers. Chem. Soc. Rev. 2012, 41, 5969–5985.

Blanazs, A.; Armes, S. P.; Ryan, A. J. Self-Assembled Block Copolymer Aggregates: From Micelles to Vesicles and Their Biological Applications. Macromol. Rapid Commun. 2009, 30, 267–277.

Shibata, M.; Matsumoto, M.; Hirai, Y.; Takenaka, M.; Sawamoto, M.; Terashima, T. Intramolecular Folding or Intermolecular Self-Assembly of Amphiphilic Random Copolymers: On-Demand Control by Pendant Design. Macromolecules 2018, 51, 3739–3745.

Pedersen, J. S. Form Factors of Block Copolymer Micelles with Spherical, Ellipsoidal and Cylindrical Cores. J. Appl. Crystallogr. 2000, 33, 637–640.

Oelmeier, S. A.; Dismer, F.; Hubbuch, J. Molecular Dynamics Simulations on Aqueous Two-Phase Systems - Single PEG-Molecules in Solution. BMC Biophys. 2012, 5, No. 14.

Matsuoka, H.; Maeda, S.; Kaesawa, P.; Matsumoto, K. Micellization of Non-Surface-Active Diblock Copolymers in Water. Special Characteristics of Poly(styrene)-Block-Poly(styrene sulfonate). Langmuir 2004, 20, 7412–7421.

Lund, R.; Willner, L.; Pipich, V.; Grillo, L.; Lindner, P.; Colmenero, J.; Richter, D. Equilibrium Chain Exchange Kinetics of Diblock Copolymer Micelles: Effect of Morphology. Macromolecules 2011, 44, 6145–6154.

Gibson, R. R.; Femyhough, A.; Musa, O. M.; Armes, S. P. RAFT Dispersion Polymerization of N,N-Dimethylacrylamide in a Series of n-Alkanes Using a Thermoresponsive Poly(Tert-Octyl Acrylamide) Steric Stabilizer. Polym. Chem. 2021, 12, 2165–2174.

Fabeyni, A.; Vittello, R.; Russo Krauss, I.; Illiano, M.; De Tommaso, G.; Amoresano, A.; Pinto, G.; Paduano, L.; Jones, C.; Di Serio, M.; D’Errico, G. Synthesis, Surface Properties, and Self-Agregation Behavior of a Branched N,N-Dimethylacrylamide Oxide Surfactant. J. Surfactants Deterg. 2019, 22, 115–124.

Czajka, A.; Hazell, G.; Eastoe, J. Surfactants at the Design Limit. Langmuir 2015, 31, 8205–8217.

Ning, B.; Wang, Y.; Zhang, M.; Bai, Y.; Tai, X.; Wang, W.; Wang, G. Surface and Foam Property of Perfluorooctyl Polyoxyethylene Ether Phosphate Salt in Aqueous-Ethanol System. J. Ind. Eng. Chem. 2021, 94, 217–224.

Ning, B.; Wang, Y.; Zhang, M.; Bai, Y.; Wang, W.; Wang, G. Surface Adsorption and Foam Performance of Sodium Perfluorooctyl Polyoxyethylene Ether Sulfate in Ethanol-Water Mixed System. J. Mol. Liq. 2021, 337, No. 116393.
(57) Brandani, G. B.; Vance, S. J.; Schor, M.; Cooper, A.; Kennedy, M. W.; Smith, B. O.; MacPhee, C. E.; Cheung, D. L. Adsorption of the Natural Protein Surfactant Rsn-2 onto Liquid Interfaces. Phys. Chem. Chem. Phys. 2017, 19, 8584−8594.

(58) Bayry, J.; Amanianda, V.; Guijarro, J. I.; Sunde, M.; Latgé, J. P. Hydrophobins-Unique Fungal Proteins. PLoS Pathog. 2012, 8, No. e1002700.

(59) Linder, M. B.; Szilvay, G. R.; Nakari-Setälä, T.; Penttilä, M. E. Hydrophobins: The Protein-Amphiphiles of Filamentous Fungi. FEMS Microbiol. Rev. 2005, 29, 877−896.

(60) Schmid, A. J.; Wiehemeier, L.; Jaksch, S.; Schneider, H.; Hiess, A.; Bogershausen, T.; Widmann, T.; Reitenbach, J.; Kreuzer, L. P.; Kühnhammer, M.; Löhmann, O.; Brandl, G.; Frielinghaus, H.; Müller-Buschbaum, P.; von Klitzing, R.; Hellweg, T. Flexible Sample Environments for the Investigation of Soft Matter at the European Spallation Source: Part i—the in Situ sans/Dls Setup. Appl. Sci. 2021, 11, No. 4089.

(61) An, S. W.; Su, T. J.; Thomas, R. K.; Baines, F. L.; Billingham, N. C.; Armes, S. P.; Penfold, J. Neutron Reflectivity of an Adsorbed Water-Soluble Block Copolymer: A Surface Transition to Micelle-like Aggregates at the Air/Water Interface. J. Phys. Chem. B 1998, 102, 387−393.