This is an Accepted Manuscript, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. This Accepted Manuscript will be replaced by the edited, formatted and paginated article as soon as this is available.

You can find more information about Accepted Manuscripts in the Information for Authors.

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal’s standard Terms & Conditions and the Ethical guidelines still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this Accepted Manuscript or any consequences arising from the use of any information it contains.
New Composite Thixotropic Hydrogel Composed of a Low-Molecular-Weight Hydrogelator and a Nanosheet

Yutaka Ohsedo,*a Masashi Oono, b Kowichiro Saruhashi, b Hisayuki Watanabe a,b and Nobuyoshi Miyamoto c

A two-component composite comprised of low-molecular-weight hydrogelator and the nanosheet Laponite®, a synthetic layered silicate, showed thixotropic behaviour, whereas each single component did not show thixotropic behaviour at the same concentration. This mixing enhancement effect may be attributed to the improved quality of the gel network.

Molecular hydrogels composed of crystalline, fibrous networks of low-molecular-weight gelators (LMWGs)1,2 and water have received considerable attention as new soft materials due to their resemblance to biological compounds that enable formation of molecular hydrogels. In particular, they are considered to have potential application as biocompatible drug delivery materials.3 Moreover, the multi-responsiveness4 and thixotropy (shear thinning)5,6 of molecular hydrogels and organogels have attracted much attention, both as new aspects of soft matter and as clues to enhancing the poor mechanical properties of known molecular gels and increase their use in medical, electroactive and photoactive applications.4 Although, LMWGs feature well-defined molecular structures and various molecular designs similar to those of polymer hydrogel materials, the available list of LMWGs for molecular hydrogels remains limited at present. Therefore, to broaden the scope of known molecular hydrogels, new methods for adding enhanced functionality to assembled networks of LMWGs must be investigated and new LMWG’s must be explored. We discovered that mixing-induced thixotropy of LMWGs with different alkyl chains (commercially available alkylhydrazides, alkylamides and alkylureas) can serve as a new method for enhancing the functionality of molecular organogels.7

Herein, we report a new method for functionalization of molecular hydrogels by the formation of a composite of an LMWG and Laponite®, a commercially available inorganic silicate rheology modifier, which is a type of inorganic nanosheet that exists in the form of nanodiscs (diameter: 30 mm, thickness: 1 nm).8 Although several systems that involve the mixing-induced enhancement of gel properties or mixed gel formulations,9,10 including organic/inorganic composite hydrogels, have been studied,11-13 our route involves mixing Laponite with a hydrogelator, i.e. palmitoyl-glycine-histidine hydrogelator (Scheme 1).14 Moreover, no thixotropic behaviour has yet been reported for LMWG/nanosheet composite gels.12
Initially, the hydrogelation ability of PalGH/Laponite composites was tested. For ease of handling, a non-viscous 1 wt% Laponite aqueous solution was used. The composites were prepared by heating and cooling a mixture of PalGH powder and Laponite aqueous solution. As shown in Figs 1(a) and (c), respectively, only PalGH in water formed a hydrogel when incorporated at a concentration above 0.5 wt%, while even a 2 wt% Laponite solution remained a liquid. For the PalGH/Laponite composites (Fig. 1(b)), a hydrogel with slightly increased clarity compared to that of the PalGH gel at 0.5 wt% was formed, possibly due to an increase in the homogeneity of the hydrogel. Moreover, rheometry measurements for the PalGH 1 wt%/Laponite 1 wt% composite confirmed the formation of a softer hydrogel than the corresponding PalGH hydrogel (Fig. S1, ESI†). Notably, the composite gels maintained their gel states for at least six months.

In the two-component composite hydrogel comprising 1 wt% of PalGH and Laponite aqueous solution, respectively, thixotropic behaviour was observed. The individual components did not exhibit such behaviour at the same concentration (Figs 1(d) and 1(f)). Moreover, a 2 wt% PalGH hydrogel exhibited thixotropic behaviour but required 2 h for recovery to the gel state after shaking (Fig. S2, ESI†). Furthermore, 1 and 2 wt% Laponite aqueous solutions remained in the liquid state. These results suggested that a new network structure that is different from the network structures of individual components may form upon mixing.

To evaluate the thixotropic behaviour of the composite hydrogels, the rheometry of the gels was investigated using step-shear measurements (Fig. 2). The PalGH 1 wt%/Laponite 1 wt% composite hydrogel exhibited a large recovery of \( G' \) and \( G'' \) and reverted to the gel-like state \((G' > G'') \) after repeated application of a large deformation shear. Although repeated recovery of the PalGH gel was also observed, unlike in the thixotropic test (Fig. 2(b)), this recovery did not directly affect the macro-scale reconstruction of the hydrogel. Moreover, the Laponite solution showed liquid-like behaviour \((G' < G'') \) at this concentration. Thus, these results clearly demonstrate qualitatively the mixing enhancement effect on the thixotropic properties of the composite.

To obtain information about the network that provides the enhanced mechanical properties of the composite hydrogels, differential scanning calorimetry (DSC) measurements were performed (Table S1 and Fig. S3, ESI†). The DSC curves of the PalGH 1 wt%/Laponite 1 wt% composite hydrogel underwent ordinal sol-to-gel and gel-to-sol changes with a similar \( \Delta H \). Compared to the peaks for the pure PalGH solution, the peaks for the composite hydrogel were shifted to lower temperature. Aqueous 1 wt%–3 wt% Laponite solutions showed no peaks, probably due to the dominant nature of the liquid. This result suggests that the composite hydrogel forms a new, possibly finer network structure than that of the pure PalGH.

To investigate the microstructure of the composite hydrogels, scanning electron microscopy (SEM) images of the corresponding xerogels and a dried Laponite solution were obtained (Fig. 3). The PalGH (1 wt%)/Laponite (1 wt%) composite xerogel exhibited a sub-micrometer-scale network fibre structure with a micrometer-scale network sheet structure. These structures appeared to originate from the PalGH fibre network and aggregated Laponite silicate discs, respectively, as

---

**Fig. 2** Periodic step-shear test results for (a) the PalGH 1 wt%/Laponite 1 wt% composite hydrogel and (b) the PalGH gel and Laponite solution.

**Fig. 3** SEM images of PalGH, composite and Laponite xerogels: (a, d) PalGH (1 wt%), (b, e) PalGH (1 wt%)/Laponite (1 wt%) composite, (c, f) Laponite (3 wt%). The values in parentheses denote the concentration of the hydrogels before generation of the xerogels.

**Fig. 4** Kratky-type plots of SAXS data of hydrogels and Laponite aqueous solution (see also Figs. S3 and S4, ESI†).
determined by comparing the SEM images of the single components with that of the composite. Notably, the composite xerogel contained finer fibres than the Laponite xerogel (Figs 3(b) and 3(c)), possibly because of the presence of PalGH, which might prevent aggregation of the silicate nanodiscs.

To small structure information on the crystalline fibres in the hydrogels, small-angle X-ray scattering (SAXS) results were obtained, and the corresponding Kratky-type plots$^{14,15}$ are shown in Fig. 4. The slope of $–2$ for the Laponite confirms the fibre-like crystals (such as those in the single-component PalGH) fibres. The interdigitated packing structure of PalGH derivatives has been extensively covered in the literature.$^{16}$ These results suggest that the composite hydrogel is composed of a network of PalGH fibre-like crystals (such as those in the single-component PalGH hydrogel) and well-dispersed Laponite.

The mixing enhancement effect on the thixotropic behaviour may be explained by assuming the formation of an improved network of the PalGH in the presence of Laponite. Note that reduced aggregation of Laponite in the presence of PalGH, the same peaks corresponding to rigid PalGH moieties and an interdigitated network were detected via FT-IR analysis (no significant peaks were observed in the spectrum). Given that the PalGH and Laponite do not interact with each other, it is possible that the bundle formation of PalGH fibre bundles may be reduced in the homogeneous presence of Laponite resulting in the formation of a high density network of finer PalGH fibres (In Fig. 3, it is possible that the finer fibres readily aggregated during the freeze-dry process). Subsequently, this new improved network may exhibit thixotropic behaviour, unlike the individual 1 wt% components.

In conclusion, we demonstrated that a new composite system exhibits thixotropic behaviour, unlike the individual 1 wt% of PalGH and Laponite. This effect may be attributed to the formation of an extended network with crosslinking between the fibres of each component. We are currently investigating the application of this mixed strategy to other systems.

Notes and references

$^1$ Advanced Materials Research Laboratory, Collaborative Research Division, Art, Science and Technology Center for Cooperative Research, Kyushu University 4-1 Kyushinmachi Nishi-ku Fukuoka 819-0388 (Japan). Fax: (+81) 92-400-4382; Tel: (+81) 92-400-4381; E-mail: phsedo@astec.kyushu-u.ac.jp.

$^2$ Nissan Chemical Industries, Ltd., 2-10-1 Tsuboinishi Funahashi Chiba 274-8507 (Japan).

$^3$ Department of Life, Environment and Materials Science, Fukuoka Institute of Technology, 3-30-1, Wajiro-Higashi, Higashi-ku, Fukuoka 811-0295 (Japan).

$^4$ Electronic Supplementary Information (ESI) available: See DOI: 10.1039/c000000x/


