Synthesis and properties of latent pigment nano- and micro-encapsulated in high-durability inorganic capsules

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Abstract
Preparation and properties of latent pigment nano- and micro-encapsulated in inorganic capsule have been investigated. Sol-gel derived C$_6$H$_5$-SiO$_2$ sphere capsules containing organic pigment were prepared using solvent-soluble latent pigment. By controlling the sol-gel reaction of C$_6$H$_5$-Si(OC$_2$H$_5$)$_3$ as starting material by acidic substituent into an insoluble organic pigment [16]. A latent pigment pigment is made soluble in various solvents by introducing an organic containing organic pigment were synthesized, and the properties of synthesized by the sol-gel method. In that manner, inorganic MCs microencapsulated in inorganic capsule have been investigated. Sol-gel derived C$_6$H$_5$-SiO$_2$ sphere capsules containing organic pigment were prepared using solvent-soluble latent pigment. By controlling the sol-gel reaction of C$_6$H$_5$-Si(OC$_2$H$_5$)$_3$ as starting material by acidic catalyst, followed by basic catalast, it was possible to prepare spherical inorganic capsules containing organic pigment with micro- and nano-size. Multi-colored inorganic sphere capsules were prepared by using the three primary colors of latent pigments (Qn-BOC, Indigo-BOC, and PY-BOC). Colored microcapsules (MCs) with excellent heat resistance, light resistance, and coloring property were obtained. Inorganic micro- and nano-capsules of Qn-BOC showed strong fluorescence-emission characteristics.

Introduction
Microcapsules (MCs) are rapidly being developed as materials for applications such as information recording, cosmetics, coatings, and food stuffs. In particular, as for microencapsulation of functional organic dyes, it is expected to be applied to, for example, display materials, information-recording materials, microsphere lasers, biomaterials such as drug delivery system and in-vivo fluorescent dye materials, and it is a field in which great development is expected in the future [1-13]. Conventionally, MCs of functional organic dyes have a structure in which an organic dye forms a core that is encased by an organic-polymer shell. However, due to the low durability of the dye, such MCs have been insufficient in terms of performance. Although organic pigments with good durability have been achieved, they are not soluble in a solvent, so it is difficult to make them into microparticles and microencapsulate them, and good performance has not been achieved [8]. In general, mechanical pulverization by a bead mill or the like is mainly used for micronizing an organic pigment [14,15]. A point of issue with that processing, however, is that the resulting particle shape is non-uniform, and spherical MCs cannot be obtained. Furthermore, when organic pigments are used, organic polymers are used for the shell structure; consequently, heat resistance, light resistance and mechanical durability (such as hardness) are low.

The present study aimed to microencapsulate high-durability and high-performance functional organic pigment in inorganic polymer. In particular, organic-solvent-soluble latent pigment was microencapsulated in inorganic MCs formed from inorganic polymer synthesized by the sol-gel method. In that manner, inorganic MCs containing organic pigment were synthesized, and the properties of the MCs were investigated. Starting as an organic pigment, a latent pigment is made soluble in various solvents by introducing an organic substituent into an insoluble organic pigment [16]. A latent pigment has the property that it can be returned to the original pigment by heat treatment. For that reason, it can be easily introduced into a capsule because it has solubility like a dye being introduced into a film, and after the pigment is transformed by heat treatment, it manifests high durability. Organic-pigment-containing inorganic MCs (MCs) were synthesized by combining the latent pigment with an inorganic polymer synthesized by the sol-gel method. As for the inorganic polymer (synthesized by the sol-gel method), phenyltriethoxysilane was used as the starting material, and MCs with spherical shape were made by coating an organic pigment with an inorganic polymer (C$_6$H$_5$-SiO$_2$; Ph-SiO$_2$), in which a phenyl group is chemically bonded to a siloxane skeleton. In this manner, organic-pigment-containing inorganic MCs with higher durability than MCs formed with organic resin could be developed. In addition, it was possible to make MCs of various colors by using latent pigments of the three primary colors as coloring material.

Experiment
Synthesis of latent pigment
The latent pigment was synthesized with three primary-color pigments as coloring materials, namely, magenta-colored quinacridone (Pigment Yellow 19: Dainichiseikakougyo Co. Ltd.), indigo (for cyan) (Tokyo Chemical Inc.), and a diazo yellow (for yellow) (Pigment Yellow 93: Dainichiseikakougyo Co., Ltd.), by a previously reported method [17].

Synthesis of inorganic micro- and nano-capsules
2.40 g (9.98 mmol) of phenyltriethoxysilane (PTES; Shin-Etsu Chemical Co., Ltd.) was mixed with 3.00 g of HCl (adjusted to pH 3), and the mixture was stirred at room temperature for 24 hours. As the
polymerization reaction progressed, the solution became an opaque emulsion. When the solution was allowed to stand for 72 hours, it became separated into two layers: one layer of C₆H₅-SiO₂ polymer and one layer of hydrochloric-acid aqueous solution. The upper layer (hydrochloric-acid aqueous solution) was removed, and a SiO₂ polymer sol having a phenyl (Ph) group was obtained by a sol-gel reaction. The Ph-SiO₂ polymer sol was dropped into NH₄OH solution (adjusted to a constant amount of 13 mol/L) by using a micropipette and stirred for 30 minutes, after which MCs settled out. The produced MCs were separated and dried at 100°C for 30 minutes. Then, 5- to 80-wt% solutions in which the Ph-SiO₂ polymer sol was dissolved in 1,4-dioxane were prepared, and those solutions were used to prepare MCs in the same manner as described above.

**Synthesis of latent-pigment-containing inorganic micro- and nano-capsules**

Latent pigment was added to and mixed with a solution prepared by dissolving the Ph-SiO₂ polymer sol (prepared as described in Section 2.2) in 1,4-dioxane to give a homogeneous solution. When that solution was added dropwise to an ammonia solution (13 mol/L), MCs containing the latent pigment were generated. The MCs were taken out and dried at 100°C for 30 minutes. They were then heat-treated at 160°C for 30 minutes to convert the latent pigment into an organic pigment encapsulated in the inorganic polymer MCs.

**Evaluation**

To determine the molecular structure of the prepared Ph-SiO₂ MCs, an infrared absorption spectrum was measured (with a Shimadzu FTIR-8400) by the KBr method. Differential thermogravimetric analysis was also performed with a Shimadzu TG-50. The surface morphology of the MCs was observed by optical microscope and scanning electron microscope (SEM; KEYENCE VE-8800). A light-resistance test (an accelerated ultraviolet exposure test) was then performed. In detail, MCs were spin-coated on a quartz substrate, irradiated with ultraviolet light (wavelength: 365 nm; intensity: 614 μW/cm²) for a certain time, and the change in the light-resistance property was tracked by using a visible-ultraviolet transmission spectrum (Shimadzu UV-2450).

**Result and discussion**

**Preparation and properties of inorganic MCs**

The synthetic route for the MCs is shown in Figure 1. To prepare inorganic MCs with high durability, PTES (C₆H₅Si(OCH₃)₃) is used as a starting material, and an inorganic-polymer precursor (in which a phenyl group is bonded to a siloxane skeleton) is synthesized by the sol-gel method with an acidic catalyst using HCl. Then, MCs are formed by developing this precursor in NH₄OH (which is a basic catalyst). After the inorganic polymer is generated as straight chains by using acidic catalyst, three-dimensional condensation is promoted by an accelerating reaction with an alkali catalyst, and spherical MCs are subsequently obtained. By introducing a phenyl group into the inorganic polymer network, compatibility with a latent pigment having an aromatic ring is improved, incorporating the pigment into the inorganic polymer becomes easy, and the coloring property of the pigment is improved. By accelerating the two-step sol-gel reaction with the acidic and basic catalysts, a spherical inorganic capsule was successfully synthesized.

The procedure for preparing the MCs is shown schematically in Figure 2. When PTES was added in a hydrochloric-acid aqueous solution of pH 3 and stirred, they did not mix and formed an emulsion state. When this emulsion-forming reaction was continued for a certain period of time in this state, a polymerization reaction (by the sol-gel reaction of PTES) proceeded, and a PTES sol was formed. After standing for 72 hours, the PTES sol and the HCl aqueous solution phase separated; accordingly, after this sol solution was extracted and added in aqueous NH₄OH, spherical MCs were formed. In the case that a latent pigment was included (as described later), a solution of latent pigment and PTES sol dissolved in 1,4-dioxane was added in the NH₄OH. In this way, MCs containing a latent pigment were formed, and by heat-treating the MCs to convert the latent pigment into an organic pigment, and organic-pigment-containing MCs were obtained.

Optical-microscope and SEM images of the prepared inorganic MCs (heat-treated at 160°C) that do not contain the latent pigment are shown in Figure 3. As shown in Figures 3(a) and 3(b), spherical MCs with smooth surface and sizes in the range of 30 to 100 μm were obtained. An observation image of the internal structure of an inorganic microcapsule with a diameter of 300 μm taken by SEM (i.e., the fracture surface of an inorganic microcapsule) is shown in Figure 3(c). It is clear from the image that the capsule is composed of a core-shell structure in which the inside is hollow. And the latent pigment is mainly contained in this hollow interior. The thickness of the inorganic coating on the hollow exterior shell depended on the size of the inorganic capsule, and when the capsule diameter was large, the thickness of the coating also tended to be large.

**Infrared-absorption spectrum and thermogravimetric analysis of inorganic MCs**

Infrared spectra of the fabricated MCs (PTES-MCs) and PTES starting material are compared in Figure 4. In the case of the PTES-MCs, absorption due to the –CH stretching vibration of –OEt (C₂H₅) from 2900 cm⁻¹ to near 2800 cm⁻¹ seen in PTES disappeared, and wide absorption due to Si-O-Si vibration was observed at 1000 to 1200 cm⁻¹. This finding suggests that the condensation reaction proceeded by the sol-gel reaction, and an inorganic polymer was formed. Moreover, absorption due to CH vibration of the phenyl group was observed near 3100 cm⁻¹ and absorptions of C=O and Si-Ph bonds of the phenyl group were observed near 1600 cm⁻¹ and around 1430 cm⁻¹, respectively, indicating the MCs have a molecular structure in which a phenyl group is bonded to an inorganic skeleton of SiO₂. In addition, absorption due to Si-OH was observed at 3660 cm⁻¹, suggesting that an -OH group was formed in part of the inorganic skeleton of SiO₂.

The results of thermogravimetric analysis (TG) of PTES-MCs and polystyrene (PS) (which is an organic polymer) are compared in Figure 5. In the case of PTES-MCs, although a slight weight loss was observed from around 400°C, a significant weight loss was confirmed from 500°C to 650°C (namely, about 56%). That weight-loss percentage roughly agrees with the value (i.e., 52.6%) given when the phenyl group in C₆H₅-SiO₂ burns off, resulting in weight loss. This result suggests that the PTES-MC has a structure in which a phenyl group directly binds to a siloxane skeleton. Above 650°C, the phenyl group was completely combusted, and only the SiO₂, skeleton remained. On the contrary, in the case of PS, weight loss began from 200°C, and a large weight loss was observed at 300 to 400°C. At 400°C, combustion of PS was complete; that is, weight loss was 100%. This result demonstrates that the prepared MCs have very high heat resistance compared with the organic polymer used for conventional microencapsulation, i.e., PS.

**Particle-size control of MCs and factors affecting particle size**

The size of the microcapsule particles depended on the concentration of the inorganic-polymer precursor used during preparation of the
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Figure 1. Formation of microcapsule containing latent pigment by using acid and basic sol-gel method

Figure 2. Preparation procedure of microcapsule (MC)
PTES-MCs and the amount of PTES sol dripped into the ammonia-catalyst solution. Accordingly, by controlling those conditions, it was possible to control MC particle size. Change in diameter of MCs with raw-material concentration (PTES sol) is plotted in Figure 6. The dropping amount at each PTES-sol concentration was fixed at 1.2×10⁻² g. Although diameter of 300 µm was obtained at PTES concentration of 100 wt%, when PTES-sol concentration was reduced in the order 80%-60%-40%-20%-10%, MC particle diameter was reduced in the order 150 µm→120 µm→90 µm→50 µm→20 µm→5 µm.

Change in mean particle diameter with PTES-sol concentration with PTES dropping weight as a parameter is plotted in Figure 7. When dropping weight was reduced from 1.2×10⁻² g to 3.0×10⁻³ g, mean particle diameter was significantly decreased. In other words, nanometer-sized (i.e., sub-micron level) particles were obtained. At PTES concentration of 60%, particle diameter dramatically was reduced, namely, from 120 to 5 µm. At PTES concentrations of 40% and 20%, particle diameters were 2 and 1 µm, respectively. At the PTES concentration of 10%, nanometer-sized MCs (i.e., diameter of about several hundred nanometers) were obtained. When the PTES sol was dropped into the base catalyst (NH₄OH), particulate MCs were formed immediately. It is presumed that since the PTES sol is insoluble in an aqueous solution, the sol-gel reaction proceeds on the surface in a three-dimensional direction as soon as the droplet of the sol contacts the base-catalyst surface. And since the size of the droplet is determined by the volume of sol dropped, the sol-droplet volume has a significant effect on the size of the MCs formed.

Observation images (taken by SEM) of micrometer-sized and nanometer-size MCs are shown in Figure 8. It is clear from these SEM images that both the micrometer-sized and nanometer-sized MCs are spherical. MCs and NCs (nanocapsules) show significantly different heat resistance. Change in the surface morphology of MCs and NCs heat-treated at 180, 200, and 220°C are shown in Figure 9. In this temperature range, the MCs show no change in morphology; in contrast, the NCs start to melt and fuse together when heated above 200°C. It is considered that the difference in the two types of capsules in terms of thermal energies is derived from the difference in specific surface areas of the particles. Such reduction of the melting point by this so-called “nano-sizing” is a phenomenon also known to occur in the case of other particles (e.g., fine metal particles) [18].

In the manner explained above, by controlling PTES-sol concentration and volume of PTES-sol dropped into the base catalyst, it is possible to obtain MCs with size ranging from the sub-micron level to around 300 µm and to control particle diameter by controlling the synthesis conditions during fabrication of MCs. It will be possible to more precisely control particle size by centrifugation of the prepared particles. That matter will be reported in a future paper.

**Synthesis and properties of microencapsulated latent pigment**

As for synthesis of PTES-MCs, latent pigment dissolved in solvent (dioxane) was added to sol solution (formed by polymerization of PTES by acid catalyst), and when this mixed solution was dropped into a basic catalyst solvent, a microencapsulated latent pigment was formed. The chemical structures of the latent pigments used are shown in Figure 10. Three primary-color pigments, namely, quinacridone-BOC (for magenta), indigo-BOC (for cyan), and Pigment Yellow 93-BOC (PY-BOC) (for yellow), were selected for the latent pigmentation. In the three kinds of latent pigments, the hydrogen of the N-H group in each organic pigment molecule was substituted with a t-BOC (COOC(CH₃)₃) group, and all three kinds were soluble in an organic
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colored with the three primary-color materials, namely, red-purple (quinacridone, Qn), blue (indigo), and yellow (PY93), were obtained. It is clear that colors of the MCs differ considerably before and after the heat treatment and that the pigment is formed inside the MCs due to the heat treatment. Moreover, the MC surface after heat treatment was smooth, and porosity due to gas generated when the latent pigment transforms to the pigment was not observed on the surface of the MCs. By varying the mixing ratio of the three primary-color latent pigments of this color material, it is possible to develop MCs with various colors. Multi-colored MCs prepared by mixing the three primary-color latent pigments are shown in Figure 12. By systematically varying the mixing ratios of PY93/Qn, indigo/Qn, and PY93/indigo from 9: 1 to 1: 9, it was possible to produce inorganic MCs with various colors, including red, blue, and green (namely, the three primary colors of light). In addition, it was possible to prepare nanoencapsulated organic pigment by controlling the particle diameter during preparation of the inorganic MCs.

Light-resistance test of inorganic-material-microencapsulated organic pigment

Light resistance is one factor that becomes important when coloring materials are applied in optical materials. It is a particularly important property in cases such as when continuity of coloration (e.g., storage stability of a pigment) must be necessary. In light of that requirement, a light-resistance test on the prepared inorganic-microencapsulated organic pigment was performed. The result of that test on microcapsulated PTES-Qn pigment is shown in Figure 13 in comparison with PS encapsulated Acid Red (organic polymer encapsulated dye). The relationship between UV exposure time and change in maximum absorption wavelength for each sample is shown in Figure 13.

As a sample in which inorganic MCs containing Qn (PTES-MCs) were coated as a film on a glass substrate was prepared and tested. As a comparison target, polystyrene MCs (PS-MCs) containing a rhodamine dye called Acid Red 52 (sulphophorhodamine B sulphonylchloride) were also prepared and tested. Acid Red 52 was chosen because it is a typical magenta-coloring material and has an absorption peak at 560 to 550 nm in almost the same manner as that of Qn.

As for the test method, the samples were irradiated with ultraviolet rays (with wavelength of 365 nm and intensity of 614 μW/cm²), and the change with time of the transmittance of the films at 550 nm was recorded. One sample was irradiated for 24 hours, and the temporal change of the UV/VIS transmittance spectrum for the maximum absorption wavelength (Qn: 550 nm; Acid Red: 560 nm) of the pigment was calculated under the assumption that the transmittance at time zero was 100%.

Since the film was made by using MCs, transmittance varied according to location; despite that variation, the transmittance of the inorganic-material-microencapsulated Qn pigment varied only slightly.

In contrast, as for the PS MCs containing Acid Red, transmittance decreased significantly with passage of time, and after 24 hours, it decreased by about 30%. This result can be explained by the fact that Acid Red (which is an organic dye) and PS (which is an organic resin) are decomposed by ultraviolet rays. It is clear from the above results that the Qn-containing inorganic MCs developed in this study show significantly improved light resistance compared to the PS MCs (organic resin) using conventional organic dye. Moreover, as for the MCs containing yellow (PY93) and cyan (Indigo) pigments, according to accelerated UV-irradiation tests, decrease in transmittance was not

Figure 6. Relationship between PTES concentration and mean diameter of MC

Figure 7. Relationship between PTES dropping weight and mean diameter of MC

Figure 8. SEM photographs of micro-size and nano-size capsules

Figure 9. SEM photographs of micro-size and nano-size capsules

Figure 10. SEM photographs of micro-size and nano-size capsules

Figure 11. SEM photographs of micro-size and nano-size capsules
of the organic-pigment-containing inorganic MCs made in this study is spherical, it is easy to pack them uniformly in a matrix and prepare dispersed solutions. The organic pigment, with high thermal and light resistance, is embedded in an inorganic network; consequently, the durability of the fabricated multicolored inorganic MCs is high. As a result, it is considered possible to develop it in a wide range of applications such as pigments for inkjets and toners and pigments for decoration.

**Fluorescence-emission characteristics of micro- and nano-encapsulated Qn-BOC latent pigment**

The latent pigment Qn-BOC showed strong fluorescence. Fluorescence emission spectrum of Qn-BOC in ethanol solution and observed, so it is concluded that the light resistance of the developed microencapsulated pigment is extremely high.

General organic pigments are insoluble in solvents, so fabricating a homogenous system with macromolecules used for microencapsulation is difficult; moreover, it is not easy to introduce them into a sol-gel film because organic pigment has an aromatic ring in a molecule and shows hydrophobic properties, compatibility to sol-gel solution synthesized in a polar solvent is not good.

A latent pigment is easy to introduce into the sol-gel derived inorganic MCs because it is soluble in a solvent, and Ph-SiO₂ (used in this study) has a phenyl group, so its compatibility with an organic pigment is high. Moreover, converting the latent pigment inside MCs to organic pigment is easy by heating at about 160°C. Since the shape

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**Figure 9.** Surface morphology of micro- and nano- capsules after heat treatment

**Figure 10.** Chemical structure of three primary colors of latent pigments

_Indigo-BOC_ _Cyan_ _Magenta_ _Quinacridone-BOC_
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Figure 11. Color change of micro capsules containing three primary colors of latent pigments by heat treatment

Figure 12. Multi-colored micro-capsules prepared by mixing three primary colors of latent pigments

emission state of Qn-BOC nanocapsules are shown in Figure 14. Although Qn-BOC shows absorption peak at 450 nm, when UV light with excitation wavelength of 284 nm (Qn-BOC has a strong absorption with molar absorption coefficient of log ε=5 at 284 nm) was irradiated onto the solution, an emission spectrum with a peak at 510 nm was obtained [Figure 14(a)]. Both the Qn-BOC-containing inorganic MCs and nanocapsules showed a stronger fluorescent property. The state (yellow-green) of fluorescence emission when UV light was irradiated onto the Qn-BOC nanocapsules is shown in Figure 14(b). By using the dispersed solution of nanocapsules with an inkjet, it was possible to print a character pattern. When the character pattern was irradiated with UV light, it emerged as fluorescent emission, and it was possible to define various character styles [Figure 14(c)]. Since complex-shaped patterns can be printed by inkjet, it is conceivable that the nanoparticles can be applied to fluorescent labeling, for example.

Conclusion

Three basic primary-color latent pigments (Qn-BOC, Indigo-BOC, and PY-BOC) as coloring materials and Ph-SiO₂ (synthesized by the sol-gel method) were used to synthesize organic-pigment-containing inorganic micro- and nanocapsules. By controlling the sol-gel reaction between acidic catalyst and basic catalyst, it was possible to prepare
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Figure 13. Durability against ultraviolet of micro-encapsulated Qn pigment

Figure 14. Fluorescence emission spectrum of Qn-BOC and emission state of nano-encapsulated Qn-BOC and inkjet patterned film on glass

Figure 13. Durability against ultraviolet of micro-encapsulated Qn pigment

Figure 14. Fluorescence emission spectrum of Qn-BOC and emission state of nano-encapsulated Qn-BOC and inkjet patterned film on glass

spherical inorganic capsules. Since the latent pigment was soluble in an organic solvent, it was easily taken into inorganic capsules and converted into an organic pigment by heat treatment. The organic pigment was enclosed in the inorganic capsules, and colored MCs with excellent heat resistance, light resistance, and coloring property were obtained. By changing the composition ratio of the three basic primary-color latent pigments, it was possible to create inorganic capsules with various colors. Inorganic micro- and nanocapsules of Qn-BOC showed strong fluorescence-emission characteristics, and various complicated fluorescent-pattern films were fabricated by the inkjet method using a dispersed solution containing the capsules.