Synthesis of organized inorganic crystal assemblies

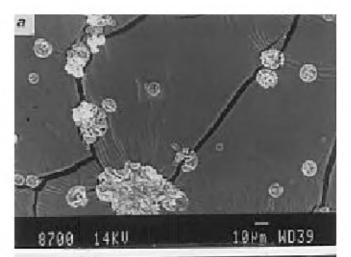
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Organized crystalline assemblies of cobalt salt and γ -iron oxide have been produced by *in situ* matrix-mediated biomimetic route. The process makes use of an organized supramolecular matrix and produces cobalt chloride crystals with characteristic morphology of *coccolith of alga* and nacreous structure of *Pinctada martensii*. Crystals of γ -iron oxide have been produced with typical morphology of aragonite spherulites in regenerated shell of *Pomaceae paludosa*.

NATURE has synthesized many inorganic crystals in living organisms with the help of biopolymers. The natural route consists of pre-organization of a supramolecular matrix, mineralization of it through interfacial molecular recognition leading to the nucleation, growth and organization of the inorganic phase. A signature of the natural synthesis is the production of the same inorganic phase with several distinctly different morphologies. For example, calcium carbonate crystals can grow in the form of plates, rods or complex tubules¹. Recent efforts in the laboratory, to duplicate this natural method of natural synthesis, have come to be known as biomimetic methods for synthesis². An understanding of the mechanism involved in such matrix-mediated synthesis has been recognized to be of great potential in the production of engineering materials. Thus, catalyst particles of controlled size and morphology, magnetic materials with appropriate anisotropy, highly porous materials, composites and well-organized crystallite assemblies can be produced through biomimetic synthesis. A variety of technically important inorganic materials such as CaCO₃, CdS, TiO₂, Fe₂O₃, Fe₃O₄, nano-sized copper and silver have been produced by the biomimetic route³⁻⁸. While these materials exhibited much of the sophistication expected in terms of shape, size and orientation, however, complex and unusual morphologies found in living organisms could not be produced. The present communication reports the synthesis of cobalt chloride and iron oxide particles with morphologies that are characteristic of the coccolith of alga, nacreous structure of P. martensii and aragonite spherulites in the regenerated shell of P. paludosa⁹. In this respect, these results open up the exciting possibility of imitating nature more closely through the control of experimental conditions and producing several morphologies of the inorganic material in the same system.

In the present investigation polyvinyl alcohol (PVA) has been used as the supramolecular matrix. Being a water-soluble polymer with a monoclinic crystal structure, PVA having rigid carbon backbone and inter-chain hydrogen





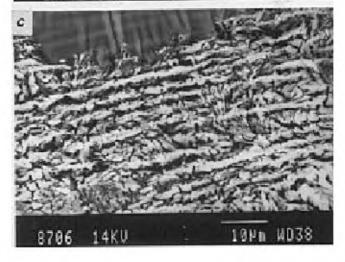


Figure 1. *a*, Random distribution of cylindrical-shaped cobalt salt particles in a supernatant PVA film; *b*, Growth of cobalt salt precipitate with morphology typical of *coccolith of algae*; *c*, Cobalt salt precipitate with morphology similar to nacreous structure of *P. martensii*.

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bonds, has the ability to solvate. It can also undergo cross-linking and a series of gelation reactions. The gel can be crystalline and the degree of crystallinity is a function of time. The presence of inorganic salts is also known to enhance gelation. Aqueous solutions containing different concentrations of PVA were produced by dissolving the polymer in deionized water at 90°C. The concentrations varied between 2.5 wt% and 10 wt% of PVA. Two different inorganic salts, viz. CoCl₂· 6H₂O and FeSO₄· 7H₂O were employed in the present study. The salts were dissolved in deionized water at room temperature to prepare 0.5 to 1.0 molar solutions and were filtered. Salt solutions were mixed with the aqueous solution of PVA in various proportions, stirred and allowed to react for a time period extending from 15 min to 15 days. Samples were drawn at regular intervals either by dipping a clean glass slide or by lifting the supernatant layer after the lapse of about a week or more. In the case of iron sulphate solutions, the polymer films with iron salt were reacted with a cold solution of sodium borohydride and dried under controlled oxygen atmosphere. The borohydride reduces the iron salt to the metallic state through a combined process of hydrolysis and reduction¹⁰. The nano-sized metal particles were then re-oxidized at room temperature to yield γ-Fe₂O₃ in a single step. The films so obtained were examined under scanning electron microscope (SEM). The

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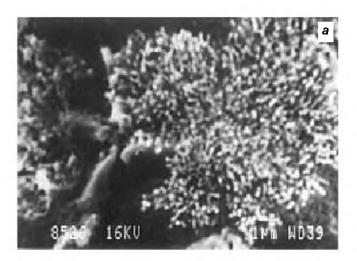
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Figure 2. *a*, EDAX analysis of the cobalt salt precipitate; *b*, XRD of the sample with ring-shaped aggregates of cobalt salt in polymer.

The composition and structure of the film constituent were confirmed by X-ray diffraction (XRD).

The pink-coloured supernatant films as well as the films coated on glass slides containing cobalt salt developed localized regions of blue colour under the polymer surface, when preserved in dry atmosphere at room temperature for several days. The SEM study of the films containing cobalt salt showed randomly distributed particles with a hollow cylindrical morphology (Figure 1 a). Examination of the film at a higher magnification showed that these particles are the aggregates of crystals arranged in a circle around a rim of PVA (Figure 1 b). This morphology is identical to the coccolith of the alga, coccolithus huxleyi, made of radial growth of calcite in a biopolymer⁹. On the other hand, the films obtained on glass slide dipped into the PVA-cobalt chloride solution immediately after mixing, showed a distinctly different microstructure (Figure 1 c). In this case, a nacreous structure of cobalt salt aligned in rows perpendicular to polymer crystals was obtained. This also resembled the nacreous growth of P. martensii lamellae composed of aragonite crystals⁹. Both the above morphologies have been synthesized through the biomimetic route for the first time substituting CaCO₃ by cobalt salt.



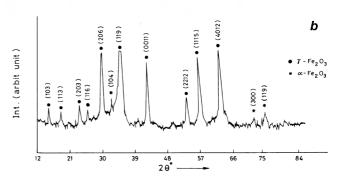


Figure 3. a, γ-iron oxide particles having growth arrangement similar to aragonite spherulites in regenerated shell of P. paludosa; b, XRD of the produced γ-iron oxide particles.

In a recent publication, crystallization of a hexamer duplex [d (TGCGCG) and d (CGCGCA)] in similar ringshaped crystals of hexadeoxyribonucleotide has been reported in the presence of cobalt hexamine¹¹. However, the EDAX analysis has indicated that the ring-shaped samples as well as rod-shaped samples produced by the present investigators are rich in cobalt and chlorine (Figure 2 a). XRD of the ring-shaped aggregates encapsulated in polymer confirmed the presence of CoCl₂ and CoCl₂·2H₂O phases embedded in PVA matrix (Figure 2 b). However, the XRD of the film coated on the glass slide showed a diffused broad peak characteristic of the glassy substrate. In the case of the films of the PVA-FeSO₄ system, γ-Fe₂O₃ formed with a range of morphologies extending from diffusion-limited aggregates to spherulite structures. Of these, the spherulite-like morphology of γ-Fe₂O₃ in PVA resembled aragonite spherulites in the regenerated shell of *P. paludosa*⁹ (Figure 3 a). Formation of the phase was confirmed by XRD which showed the presence of γ-Fe₂O₃ as a major phase along with α-Fe₂O₃ (Figure 3 b).

There are several reports available in the literature on the synthesis of agglomeration-free fine-sized transition metal powders using PVA and other hydrophilic polymers ^{12,13}. However, the present study highlights the possibility of producing inorganic materials with morphologies similar to those observed in living organism through biomimetic synthesis. This can be exploited to develop materials for magnetic and humidity sensors with high sensitivity.

- 1. Calvert, P. and Mann, S., J. Mater. Sci., 1988, 23, 3801-3815.
- Addadi, L. and Weiner, S., Angew. Chem., Int. Ed. Engl., 1992, 31, 153–169.
- 3. Berman, A. and Charychi, D., Adv. Mater., 1999, 11, 296-300.
- Mann, S., Heywood, Brig, R., Rajam, S., Derck, J. and Birchall, Nature, 1988, 334, 692-695.
- Sobon, C. A., Bowen, H. K., Broad, A. and Calvert P., J. Mater. Sci. Lett., 1987, 6, 901–904.
- Baskaran, S., Song, L., Liu, J., Chen, Yuan L. and Graff, Gordon L., J. Am. Ceram. Soc., 1998, 81, 401–408.
- Mel, L., Schnablegger, H. and Mann, S., Nature, 1999, 402, 393–395.
- Joshi, S. S., Patil, S. F., Iyer, V. and Mahamuni, S., Nanostruct. Mater., 1998, 10, 1135–1144.
- 9. Watabe, N., J. Crystal Growth, 1974, 24/25, 116-122.
- Freitag, W. O., Sharp, T. A., Baltz, A. and Suchodolski, V., J. Appl. Phys., 1979, 50, 7801–7803.
- Satheesh Kumar, P. and Gautham, N., Curr. Sci., 1999, 77, 1076– 1078.
- Savinova, E. R., Chuvilin, A. L. and Parmon, V. N., J. Mol. Catal., 1988, 48, 217–229.
- Hirai, H., Wakabayashi, H. and Komiyama, M., Bull. Chem. Soc. Jpn., 1986, 59, 367–372.

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Detection of areal snow cover changes over Antarctica using SSM/I passive microwave data

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SSM/I (Special Sensor Microwave Imager) passive microwave data of two different months over the Antarctic subcontinent have been used to (i) detect the areal extent of snow cover, and (ii) to monitor the changes in the snow cover in this region. Minimum accumulation of snow over Antarctica occurs in February, and hence the detection of snow cover during January and February should indicate a depletion phase. An overall 14% decrease of snow has been observed during this period. Corresponding data for accumulation/depletion of snow indicates a matching of general trend. SSM/I data, thus, can be used as a very promising tool for monitoring of snow over the Antarctic region since it is not affected by cloud cover/darkness.

ANTARCTICA is the fifth largest continent on the earth (area approximately 14 million sq km), which surrounds the South Pole. It is almost totally covered by ice, devoid of any permanent human population. Ice-covered region over Antarctica constitutes almost 4.5% of the earth's surface area. It is also the coldest continent with ambient air temperature varying much below the freezing point of water (-80° C to -30° C) throughout the year. It is enriched with valuable minerals, coal and petroleum. Antarctica contains about 90 per cent of the world's ice, and if melted, it would raise the ocean by around 60 m. Antarctica is also the highest continent on the earth, because of the massive thickness of ice which overlays the land. The thickness of ice at places is beyond 4500 m (average thickness is approximately 2100 m). Snow and sea ice are dynamic geophysical features on the earth's surface. Together with land ice, they influence the local and global climate and oceanic regimes. Moreover, polar ice masses, and the ice sheets in particular, are sensitive indicators of climate change, which it is feared may result from an increase in atmospheric 'greenhouse gases' such as methane and CO₂, the latter caused by burning of fossil fuels. Study on ozone hole over Antarctica has attracted global attention due to its multifaceted impact over global warming and related hazards due to increased penetration of UV rays on the earth through the ozone hole. The seasonal sea ice covers of both polar regions may react to an increase in atmospheric temperature by reducing their areal extent¹. Based on the study with Nimbus 7 (USA)

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