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5 Dual internal functionalization of imogolite nanotubes as evidenced

6 by optical properties of Nile red

7 **Abstract.** Hybrid imogolites have been prepared by combining methyltrimethoxysilane with various
8 organosilanes (Si-R) to obtain nanotubes with a bi-functional internal cavity. With an initial doping
9 ratio Si-R/Si of 5%, nanotubes remain the major product in the samples and their diameter is not
10 modified compared to pristine methyl imogolite (Imo-CH₃). The nanotube wall's internal polarity
11 and reactivity is significantly changed as evidenced by using a solvatochromic dye (Nile red). This
12 research opens many possibilities to bring selectivity to the internal cavity or to manipulate the
13 properties of trapped organic molecules.

14 1. Introduction.

15 Imogolite (Imo) is a natural aluminosilicate nanotube (Yoshinaga and Aomine, 1962). Its chemical
16 composition is (OH)₃Al₂O₃Si(OH) (Cradwick et al. 1972). The nanotubes are single-walled with a
17 large curvature. The internal diameter of synthetic Imo is about 1 nm with a wall thickness of 0.6 nm.
18 The curvature of the nanotube is partly due to the mismatch in distance between the relaxed size of
19 the silicon tetrahedra (2.6 nm) and the O-O distances of the lacuna of a flat aluminum dioctahedric
20 sheet (2.9 nm) where Si establishes three Si-O-Al bonds. This size mismatch alone is however not
21 sufficient to explain the curvature. The intramolecular OH bonds between the internal silanol groups
22 are also responsible for the stabilization of a zig-zag configuration of Imo and contribute to their very
23 high curvature (Lee et al. 2011).

24 Since their discovery and their first synthesis more than forty years ago, many modifications of the
25 wall composition as well as external and internal surfaces have been achieved (Arancibia-Miranda
26 and Escudey 2016). In particular, the group of Bonelli unraveled a one-pot process to produce hybrid
27 Imo nanotubes where the internal silanol groups are completely replaced by Si-CH₃ groups (Bottero
28 et al. 2011). This synthesis produces nanotubes with a slightly larger internal diameter (*ca.* 1.8 nm),
29 probably due to the absence of intramolecular hydrogen bonds. The hybrid Imo (Imo-CH₃) form
30 stable dispersions but are always associated in small bundles (Boyer et al. 2014). Boyer et al. (2014)
31 also showed that the inside of the hybrid nanotubes has an electronic density lower than water. Thus,
32 the cavity of the Imo-CH₃ is hydrophobic and small enough to prevent water from wetting the internal
33 pore surface (Liao et al., 2018). However, Imo-CH₃ are able to trap small organic molecules in their
34 cavity (Amara et al. 2015; Picot et al. 2016).

35 In this paper, the possibility to incorporate an additional organic function within the internal cavity
36 of Imo-CH₃ nanotubes is established. This functionalization is obtained through the partial
37 replacement of the Si-CH₃ groups by other Si-R groups. We attempted to substitute 5% of the Si-CH₃
38 groups, which corresponds to one Si-R group per nanotube circumference. This ratio was however
39 difficult to detect and to quantify with standard analytical techniques. Nevertheless, the reactivity of
40 the internal cavity towards organic molecules is strongly modified by this additional
41 functionalization, as illustrated by using Nile red (NR), a solvatochromic hydrophobic dye.

42 2. Materials and Methods

43 Imo-CH₃ and functionalized Imo-CH₃ with a modified internal organic layer were prepared using a
44 protocol inspired by Bottero *et al.* (2011). Briefly, 80 mM aluminum-tri-sec butoxide (ASB) was
45 added to HCl_{aq} solution with a HCl/Al molar ratio of 0.5 and stirred during one hour. Then, silicon
46 precursors were added to obtain a Si/Al molar ratio of 0.575. Methyltrimethoxysilane is used as the
47 main silane. For the new surface function of the internal organic layer, trimethoxysilanes are used
48 bearing the following functional groups: 3-bromopropyl, chloromethyl, 3-aminopropyl, vinyl and 3-
49 mercaptopropyl. An initial ratio of 5%, defined as the molar ratio between Si-R silane and total
50 amount of Si, was used. For vinyl group, higher ratios (20 and 50%) were also tested. The dispersions
51 were kept in an oven for 5 days at 90°C and then dialyzed against MilliQ water using a 10 kDa
52 membrane until the conductivity of the external phase reaches 4 μS·cm⁻¹. In the following, the pristine
53 Imo-CH₃ is called methyl. The 5% doped Imo are called by the name of the end group of the
54 additional organosilane, i.e. bromo, chloro, amino, vinyl and mercapto.

55 Details on the characterization methods are given as supplementary information.

56

57 3. Results and Discussions

58 For all added silanes, nanotubes are obtained as the main product. The small angle X-ray scattering
59 (SAXS) signals obtained after dialysis for all the samples are reported in Figure 1a. The main feature
60 of the SAXS curves is the presence of clear oscillations due to a monodisperse diameter of the
61 nanotubes. The positions of the second and third intensity minimum of these oscillations, that are
62 very sensitive to the nanotube diameter, are similar for all nanotubes. However, the shape of the
63 curves changes. For vinyl- and mercapto-Imo, the slope of the low q range scattering signal is
64 modified and the position of the first intensity minimum is shifted to a larger q value. This shift may
65 be due to a modification of the internal electronic density or to the presence of a high proportion of
66 side products (allophane, proto-Imo or aluminum hydroxide platelets) (Picot *et al.* 2018). Methyl-,
67 chloro-, bromo- and amino-Imo display very similar SAXS curves. Cryo-TEM pictures of the bromo-,
68 amino- and mercapto-Imo (Figure 1b-d) clearly show the presence of side-products: Gibbsite platelets

69 (highlighted by black arrows in Figure 1c and 1d) and many proto-Imo and allophane. Cryo-TEM
70 also suggests that the bromo-Imo (Figure 1b) contains much more proto-Imo and allophane than the
71 amino-Imo, for example (Figure 1c).

72 The IR spectra of the synthesized hybrid Imos are reported in Figure 2. The dominant tubular nature
73 of the products is clearly visible with the presence of a doublet in the 960–900 cm^{-1} range (Boyer et
74 al. 2014; Amara et al. 2015; Picot et al. 2016). These bands are attributed to the Si–O stretching
75 vibrations. In the Imo local structure, the three Si–O bonds are not all equivalent with respect to the
76 nanotube axis. Therefore, two components at 910 and 955 cm^{-1} are observed depending on the angle
77 between the tube axis and the Si–O bonds. This is not the case for allophane or proto-Imo for which
78 a single band at 940 cm^{-1} is observed. The relative intensity of the two bands (910/955 cm^{-1}), linked
79 to the proportion of side products and notably to the presence of proto-Imo and/or allophane (Picot et
80 al. 2018), is changing from one hybrid to another depending on the introduced silane. On this basis,
81 the hybrids with the most side products should be the chloro- and mercapto-Imo whereas the methyl-,
82 amino- and vinyl-Imo should contain less of these. Moreover, for vinyl- and mercapto-Imo, the
83 presence of aluminum hydroxide is evidenced by additional peaks at 1020 cm^{-1} and in the 3700–3400
84 cm^{-1} range. The characteristic peaks of methyl-Imo are also noticed at 782, 1275, 2918 and 2976 cm^{-1}
85 for the Si–C stretching and methyl rocking, CH_3 symmetric deformation in Si– CH_3 and finally
86 symmetric and asymmetric stretching of C–H in CH_3 , respectively (Bottero et al. 2011; Boyer et al.
87 2014; Liao et al. 2017). For all introduced groups, slight modifications of these later bands are visible
88 (inset Figure 2).

89 In contrast, vibration bands signaling for the incorporation of added silanes are not clearly visible on
90 the IR spectra, as could be expected for their low relative content (5% maximum only). Only the vinyl
91 groups are visible with the typical =C–H stretching at 3080 and 3060 cm^{-1} (see inset of Figure 2).
92 Moreover, two other contributions at 1615 and 1420 cm^{-1} associated to C=C stretching and to the
93 bending in the plane of =C–H, respectively, are identifiable (inset (b) Figure S1) (Larkin 2011;
94 Silverstein et al. 2015). The relative intensity of these four peaks increases with the initial amount of
95 the vinyl-bearing silane, supporting their attribution.

96 As an alternative method to evidence the successful doping of the internal organic layer, it can be
97 assumed that the presence of an additional chemical group could change the polarity of the cavity.
98 Therefore, NR, a solvatochromic hydrophobic molecule, is used as a probe of the modification of the
99 internal organic layer (Hess et al. 2013; Martin et al. 2016; Singappuli-Arachchige et al. 2016). The
100 optical properties of NR are very sensitive to its environment, displaying a bathochromic shift of its
101 absorption maximum of more than 100 nm when it is transferred from an apolar to a polar solvent
102 (Dutta et al. 1996; Dias et al. 1999; Hazra et al. 2004; Jose and Burgess 2006). NR was transferred
103 from a dodecane solution to an aqueous dispersion of the Imo- CH_3 through a liquid/liquid extraction

104 (Picot et al. 2016). It was noticed that no phase transfer of the dye occurred if pure water or an aqueous
105 dispersion of Imo (Imo-OH) were used (Figure S2). Imo-OH and Imo-CH₃ have almost the same
106 external surface covered with Al₂-(μOH) groups. The fact that no significant adsorption of NR is
107 observed on Imo-OH, even after prolonged contact time, disqualifies an external surface adsorption
108 on Imo-CH₃ and explains the very efficient liquid-liquid extraction of NR. Based on such experiment
109 and on the results obtained by Amara et al. (2015), the conclusion is that NR molecules are mainly
110 trapped in the internal cavity of Imo-CH₃ nanotubes.

111 The UV-vis spectra of the dispersions obtained after extraction of the dye by the different doped
112 hybrid Imo are significantly changed (Figure 3). In dodecane (an apolar solvent), NR exhibits a
113 yellow color with an absorption maximum at 491 nm. In polar ones like ethanol, it is red/pink with
114 the absorption maximum shifted to 549 nm. After NR extraction, the nanotube dispersions have a
115 strong coloration going from pink for amino-Imo to blue for mercapto-Imo. The measured
116 bathochromic shift of the dye absorption maximum within the nanotubes compared to the initial
117 dodecane solution is, in all cases, larger than the one observed with ethanol. For methyl-Imo, this
118 maximum is at 556 nm and it shifts up to 589.5 nm for the mercapto-Imo. Thus, upon interaction with
119 hybrid nanotubes, NR has clearly taken a color typical of a polar solvent. A change in the spectral
120 shape is also noticed for almost all samples compared to methyl-Imo, with up to three distinct
121 absorption bands being clearly identified. The main emission band of NR extracted from dodecane
122 by the different doped hybrid Imo also showed a significant shift of its maximum as a function of the
123 doping moiety (figure S3)

124 . Importantly, by measuring the dye concentration in the dodecane phase at the end of the extraction
125 process, it was possible to determine the concentration of the extracted NR inside the different
126 nanotubes. This concentration is 1.25 mg of dye per gram of Imo for all doped hybrid nanotubes.

127 Two effects can explain these observations. First, theoretical predictions of Elliott et al.(2017) using
128 DFT calculations suggested that the nanocavity of Imo has an intrinsic polarity that could be strongly
129 influenced by modifications of the internal surface. The present results would therefore constitute an
130 experimental confirmation of their calculation. Second, an effective 5% doping corresponds to one
131 organosilane group every 5.5 Å. The determined NR concentration within the nanotube (1.25 mg·g⁻¹)
132 corresponds to approximately one molecule every 360 Å along the Imo nanotube axis. Hence, the
133 doping organosilane dye ratio would be 65. Therefore, direct physical or chemical interactions
134 between NR and the doping organic moieties are highly plausible even at 5% doping. It should also
135 be noticed that the side products of the imogolite synthesis (allophane, proto-imogolite or gibbsite)
136 might also interact with the dye and contribute to its optical response, a point that will require further
137 studies.

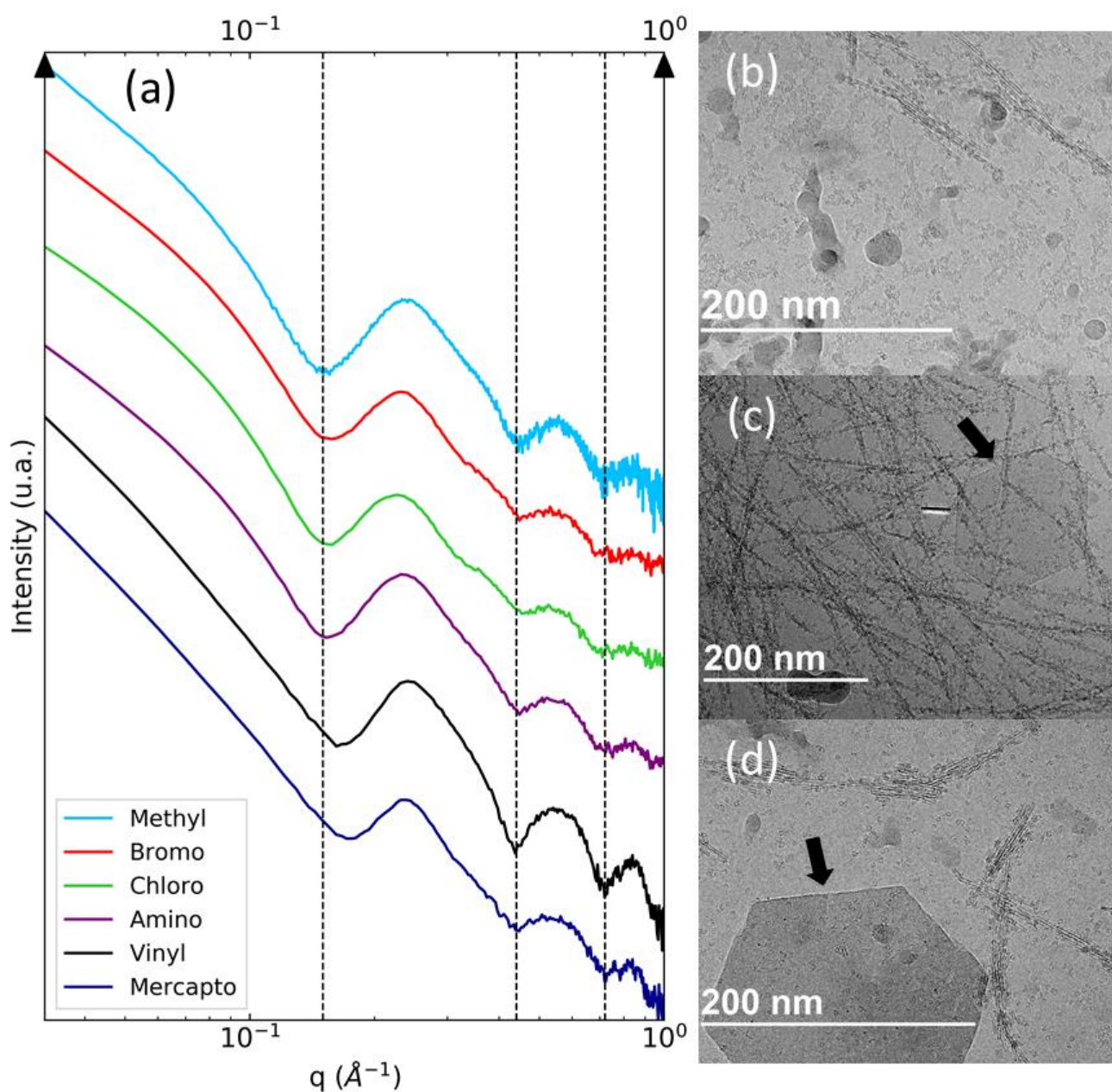
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139 [4. Conclusion](#)

140 Hybrid nanotubes with an internal Si-CH₃ surface doped with various Si-R chemical groups have
141 been obtained. The doping agent does not significantly modify the affinity of Nile red for the internal
142 cavity. However, the polarity of the nanotube wall is strongly modified by the presence of the doping
143 organosilanes, as evidenced by the strong variations of NR optical responses after entrapment. This
144 demonstrates that the proposed synthesis allows for the preparation of Imo nanotubes with a bi-
145 functionalized internal surface and opens many possibilities to improve the specificity of the cavity
146 towards targeted organic molecules and/or to tune the properties of encapsulated species, with
147 foreseeable applications in energy harvesting, remediation and sensing devices.

148 [Acknowledgements.](#)

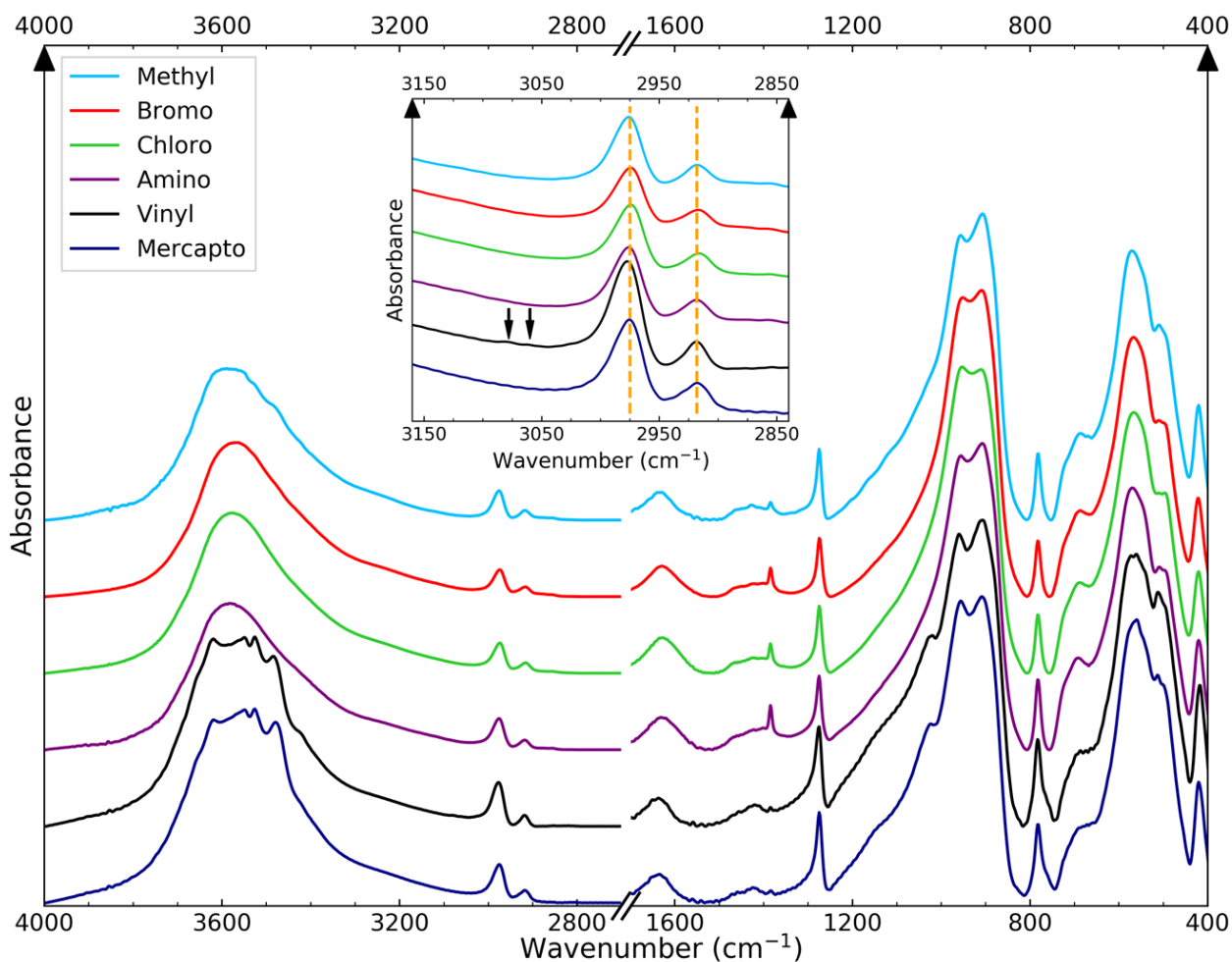
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157

Figure 1 - SAXS curves (a) of methyl-Imo (cyan) and doped nanotubes: bromo (red), chloro (green), amino (purple), vinyl (black) and dark blue (mercapto). Dashed lines highlight the position of the first, second and third intensity minimum. Cryo-TEM images of bromo- (b), amino- (c) and mercapto-Imo (d). Black arrows highlight the presence of aluminum hydroxide platelets.

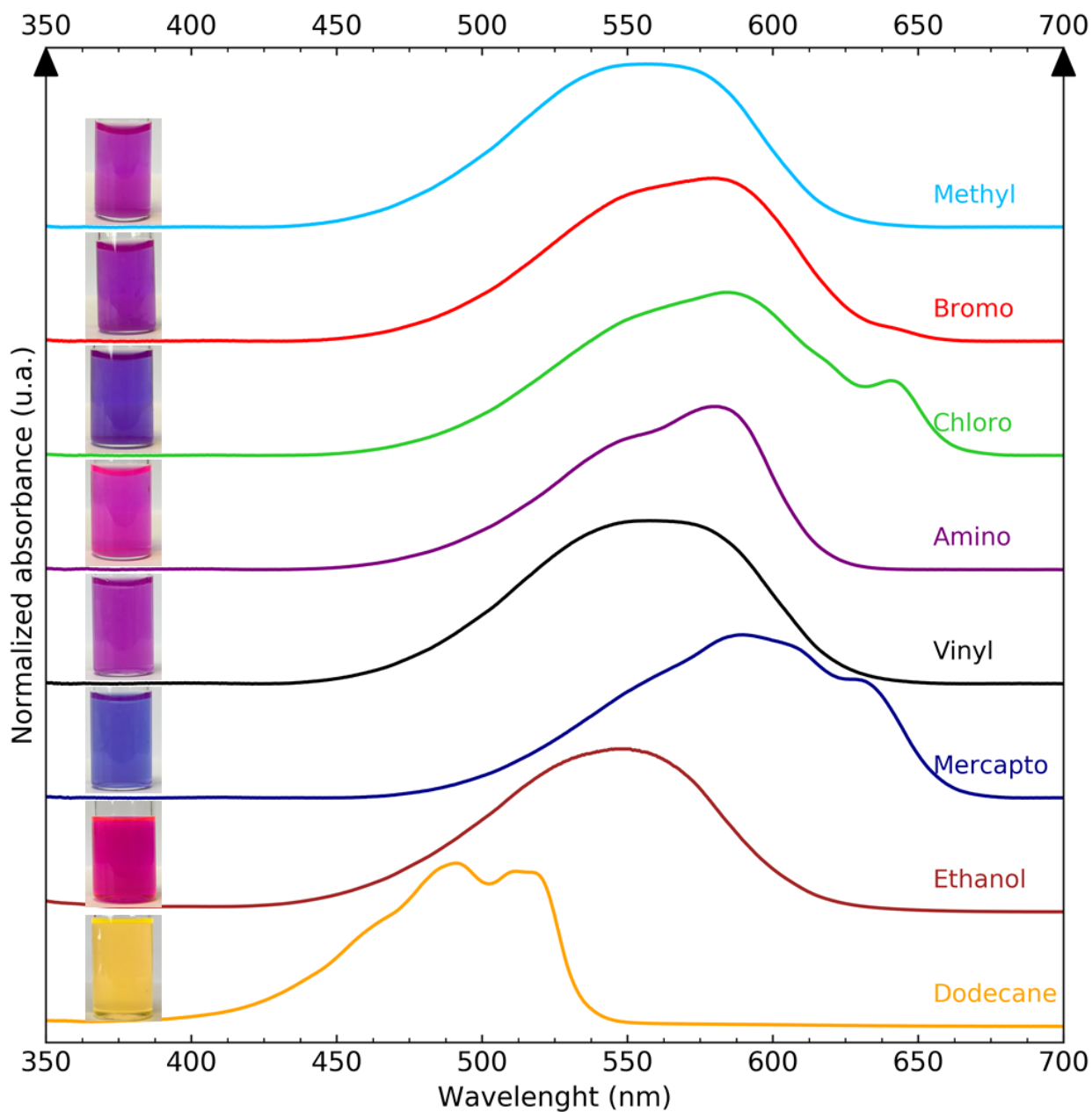
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Figure 2 - IR spectra of methyl-Imo (cyan) and doped nanotubes: bromo (red), chloro (green), amino (purple), vinyl (black) and dark blue (mercapto). Inset is a zoom of the 3150 – 2850 cm^{-1} area. Orange dashed lines denote the position of antisymmetric and symmetric stretching of C–H in CH_3 . The two arrows highlight the peaks of vinyl groups associated to the =C–H stretching.

160



161

Figure 3 - Normalized absorbance of NR extracted from dodecane by methyl-Imo (cyan) and doped nanotubes: bromo (red), chloro (green), amino (purple), vinyl (black) and mercapto (dark blue). Normalized absorbance of NR in ethanol (brown) and in dodecane (orange). Optical images of the dispersions are added above their corresponding curve.

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