ACCELERATED TOBACCO SMOKE STAINING ON WATERBORNE ACRYLIC PAINTINGS CAUSED BY EXUDING SURFACTANTS: A STUDY WITH PY-GC/MS AND THM-GC/MS

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Abstract

The modern and contemporary artistic movements profited from the introduction of several innovative materials (pigments, binders and additives). Frequently these materials have been used without a thorough knowledge of their long term durability and the compatibility with other materials present. One example is observed in Raoul De Keyser’s canvas painting entitled Camping II (1969): brown stains have emerged over time on the white areas, while the blue and green areas remained intact. At that time, De Keyser painted most of his works with waterborne acrylic paints.

Microsamples from the bulk paint film were analysed using pyrolysis coupled to gas chromatography – mass spectrometry (py-GC/MS) with or without thermally-assisted hydrolysis and methylation (THM-GC/MS). Hydrophilic surfactants, which have been observed exuding from other paint films, were detected in the bulk of the waterborne acrylic-based white paint. In a second sampling campaign, small fractions of the brown exudate were successfully removed by aqueous extraction. Curiously, these extracts not only contained the sought-after surfactants, but also relatively high amounts of nicotine and other cigarette smoke-related components. It is suggested that the exuding hydrophilic surfactants form a suitable substrate to bind hydrophilic and hygroscopic smoke components, explaining the brown colour of the exudate.

The future restoration of Camping II, using a cleaning technique based on micro-emulsions, greatly benefits from the knowledge of the composition of the exudate and the processes that contribute to its formation.

1 Introduction

The diversity and availability of materials used in artists’ paints have grown enormously during the past century. A massive number of synthetic pigments, both organic and inorganic, and a number of new binding media have been introduced. Both traditional and new binding media have been supplemented with a range of additives to improve their paint characteristics. These minor components of modern commercial paints include products to modify the surface tension, stability, rheology and thickness of the paint, emulsifiers, UV absorbers, biocides and many more. However, the long-term ageing stability of these newly introduced pigments, binders and additives, and their interaction with other paint components is not always known. Frequently, this results in premature failing of these relatively young paint systems.

The thorough understanding of the physicochemical processes behind the different forms of degradation is essential for an optimal preservation of the art object. This way, the best suited exhibition and storage conditions can be selected in order to prevent or at least slow down degradation phenomena. Also, it aids in selecting the most appropriate conservation or restoration treatments.
The exuding surfactants and other hydrophilic components that are dragged along are seen to form semi-crystalline rounded crystals at the surface, but sometimes form patchy or even complete surface layers. This can lead to visual changes in gloss or enhanced dirt pick-up5.

The brown stains on the white areas in De Keyser’s Camping II visually deviate from the described semi-crystalline or matt surface effects observed in PEO surfactant exudates, but a relationship cannot be excluded. In this paper, the exact nature of the brown stains and reason for their formation are determined. To attain these goals, samples from the affected and unaffected paints, and the brown exudate are collected and analysed with py-GC/MS. To improve the detection of compounds with labile carboxylic acids, also THM-GC/MS is done on the bulk paint samples. In THM-GC/MS, these carboxylic acids are simultaneously converted into methyl esters and pyrolysed. Finally, the initial stages of a restoration campaign in the near future is described.

2 Experimental

2.1 Samples

For the determination of the pigments, binding media and additives, two microsamples were collected from the affected white and the unaffected blue painted areas: small fragments were scraped off at the side edges of the painting by using an ophthalmic scalpel. In a second phase, two extracts from the brown stains were collected following a protocol described by Ormsby5: 50 µL milli-Q water was gently pipetted onto the paint surface and left in contact with the stain for two minutes (figure 2). Afterwards, the drops of water were carefully collected using a micropipette. Prior to py-GC/MS analysis, the water was evaporated using a vacuum centrifuge.

Many of the documented problems, however, are related to the surfactants that are used to suspend (“emulsify”) the acrylic copolymers into the aqueous medium. In most cases non-ionic polyethoxylate-based surfactants (PEO) have been used, which are known to potentially migrate to the film surface through complex mechanisms involving water evaporation, interfacial tension reduction, diffusion and capillary forces and are influenced by several internal and external factors3,5,6. An extensive review of existing scientific research on this topic is given by Ormsby and Learner6. Evidently, the type and quantity of the surfactant play a decisive role, such as the length of the PEO-chains (the longer these are, the more hydrophilic and thus the more likely they will exude), as well as the copolymer blend and the quantity of acid monomer present.

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point wire. After drying, the wire is introduced in the pyrolysis unit, which is maintained at 250 °C. Pyrolysis is set at 625 °C (determined by the wire used) for 10 s. The split injector is at 250 °C with a split ratio of 8:1. The GC oven programme was 50 °C for 1 min, then ramped at 50 °C/min to 100 °C, at 5 °C to 180 °C and at 10 °C/min to 320 °C, which is maintained for 5 min. The mass spectrometer records the m/z 35-500 range (scan time 0.50 s) in positive electron impact modus.

For completeness, the paint samples are analysed with dispersive micro-Raman spectroscopy mainly for pigment identification, and with FTIR microscopy, which generally gives a general characterisation of the binding media. The micro-Raman spectrometer is a Renishaw Invia equipped with a 785 nm diode laser excitation source and a direct-coupled microscope with 50x objective lens. A Bruker Vertex 70 FTIR system is used, equipped with a Hyperion 3000 microscope in transmission modus and using a diamond cell.

3 Results

3.1 White paint sample

The pigments in the white paint were identified with micro-Raman spectroscopy (figure 3) as titanium white (TiO₂; rutile form) and barium sulphate (BaSO₄), which is often observed as a filler in contemporary paints. In addition, the FTIR spectrum (figure 4) showed the presence of acrylates in the binding medium. The highest correlation in our FTIR-library was of the methylation agent TMAH during pyrolysis (figure 5). These are characteristic breakdown products of poly(ethyl acrylate-co-methyl methacrylate) or p(EA-MMA), which is typical for early formulations of acrylic emulsion paints between WW II and the mid-1980s11. It was and is available under different commercial names, such as Plextol B-500 (Röhm), Primal AC-234, AC-634, AC-33 and AC-22 (Rohm & Haas). A number of minor components can be identified as characteristic for paint additives. Without methylation, these are dibutyl phthalate (RT 18.32 min), p-tert-octyl phenol (RT 11.28 min) and a di-tert-butyl phenol (RT 9.39 min). Di-n-butyl phthalate (DnBP) is a common plasticiser, while p-tert-octyl phenol is a pyrolysis product of octyl phenyl polyethoxy ethanol7, a common PEO surfactant to suspend the acrylics in the aqueous medium. It is commercially available under the name Triton (e.g. Triton X-405 and X-305). The di-tert-butyl phenols are a group of UV absorbers and antioxidants; the isomers share the same mass spectrum and cannot be distinguished. 3,5-di-tert-butyl phenol, for example, is known to be present in the Plextol product range (e.g. Plextol D-498)7.

With methylation using TMAH, the methyl esters of hexadecanoic (palmitic) acid, octadecanoic (stearic) acid and dehydroabietic acid (DHA) are detected (respectively at RT 17.81, 20.83 and 22.96 min). The former two are typical components of animal and vegetal oils, while the latter is a marker for colophony. However, other crucial markers for oils and colophony...
are absent or below detection limits, suggesting that hexadecanoic acid, octadecanoic acid and DHA are added separately; their salts are often used in contemporary paints to optimise certain characteristics of the paints (rheology, gloss, etc.). In the presence of TMAH, DnBP is converted into dimethyl phthalate (DMP; RT 8.17 min).

### 3.2 Blue paint sample

The pigment in the blue paint sample was identified as phthalocyanine blue using Raman microscopy (figure 3). Several forms of phthalocyanine blue exist, jointly called Pigment Blue 15:x (PB15:x), where x can be 0, 1, 2, 3, 4 or 6 corresponding to the specific crystalline form. Whether the exact form can be determined using Raman spectroscopy is a controversial topic14–16, and is therefore omitted here. FTIR analysis of the sample indicates the presence of poly(vinyl acetate) (PVAc) as binder (figure 4).

The binder was confirmed with py-GC/MS, where, although not completely resolved, its typical pyrolysis products13 acetic acid and benzene are observed. Figure 6 shows both components using selective ion plotting (m/z 43+45+60 for acetic acid with RT 0.88 min, m/z 78 for benzene with RT 1.03 min). Styrene and phenyl glyoxal (resp. RT 2.00 and 2.35 min) seem to be related compounds through incomplete pyrolysis (in comparison with Learner13). PVAc gives rise to hard and brittle paint layers unless high amounts (up to 20%) of plasticisers are added. DnBP is a typical external plasticiser for PVAc13, but is known to migrate out of the paint film over time and thus slowly increase brittleness, while creating a tacky surface. That is why often internal plasticisers that copolymerise within the PVAc were introduced in the 1960s13. In this case a vinyl versatate resin (VeoVa) seems to have been added, although only a single peak is observed (RT 5.13 min) contradicting the broad band of non-resolved peaks observed by Learner13.

The monomers, sesquimers, dimers and trimers of p(EA-MMA) are present in the blue paint sample as well, although at a much lower abundance compared with the white paint sample. Without selective ion plotting only the most intense peak, that of the EA-EA-EA trimer, can be observed in the top spectrum of figure 6. This suggests the blue paint has a small fraction of the acrylic white paints mixed in to attain a slightly lighter blue hue. Titanium white and barium sulphate are weak Raman scatterers compared to PB15:x, which exhibits a high Raman response; this explains why these were not observed with Raman spectroscopy in the blue mixture.

With methylation using TMAH, the same additives were found as in the white paint sample: the methyl esters of hexadecanoic acid, octadecanoic acid and DHA. The relatively high amounts of DMP are the result of methylation of the relatively high amounts of DnBP that were already observed in the non-methylated pyrogram.

### 3.3 Aqueous extracts of the brown stains

Even though the brown stains were locally removed as a consequence of the aqueous extraction procedure (figure 2), the extracts contained very little material detectable by py-GC/MS. This suggests that the brownish layer is extremely thin. After evaporation and redissolving in ethanol, the complete sample was introduced in the pyrolysis unit (figure 7).
Unfortunately, there was no sample left for THM-GC/MS.

Many of the unresolved peaks in the pyrogram could not be identified. The pyrolysis product of the potential exuding octyl phenyl polyethoxy ethanol surfactant, p-tert-octyl phenol, could only be observed when plotting m/z 135. The larger peaks in the pyrogram include nicotine at RT 6.82 min and nicotyrine at RT 9.32 min. Both components (figure 8) were identified by using the NIST 02 MS library. Nicotyrine has been reported as an oxidation product\(^\text{17}\) and as a pyrolysis product\(^\text{18}\) of nicotine. Several other components are most likely also related to tobacco smoke, but could not be identified. At RT 15.04 min, tri(2-chloroethyl)phosphate elutes; this is a common flame retardant, plasticiser and viscosity regulator that is often used in acrylics\(^\text{19}\). It is not clear if this product originates from within the paint layer.

4 Discussion

4.1 Hypothesis for the brown stain formation

Both titanium white and acrylic emulsions have long been widely used in artists’ paints because of their relatively high stability. Degradation of acrylics under influence of UV-B irradiation is a well-studied phenomenon, but is unlikely to take place in a museum environment. Nonetheless, a cause within the paint film is presumed, since particularly the waterborne acrylic white paint is affected, while the blue waterborne (mainly) PVAc paint and other colours are not. An exudate of one of the additives, or a related breakdown product must be at the core of the problem.

A PEO type surfactant was detected in the bulk of the acrylic emulsion paint using py-GC/MS which is known for its tendency to migrate to the paint surface. The process of this phenomenon is not yet fully understood, but the migration of surfactants, most likely together with other hydrophobic paint additives, involves desorption during the coalescence of the polymer particles in the drying process\(^\text{20}\). As a result, semi-crystalline structures or less visible surface layers are formed on the paint surface, changing the visual aspect of the paints, such as the gloss. This is a known and well-studied phenomenon. As far as the authors know, however, no brown stains have been described.

The surfactant-rich hydrophilic surface layer is suspected of filtering out hydrophilic and hygroscopic air pollutants. Although a smoking ban has existed for many years in the museum, it is possible that this painting was subjected to a smoky environment at some time in the past: in the painter’s workshop, an art gallery, the museum itself or during its loan in a public official’s office. We believe that the hygroscopic nicotine and related products in cigarette smoke found a suitable substrate in the surfactant-rich exudate at the paint surface, forming a viscous, brown surface layer. This would explain the flow-like patterns that are observed.

In the areas with little to no acrylates, particularly in the blue PVAc-based area, little to no PEO surfactant exudates are detected, creating a less suitable substrate for cigarette smoke deposits.

4.2 Conservation strategy

The main goal in the restoration of Camping II is to remove the brown stains present in the white painted areas without damaging the acrylic paint layer. After sample collection using the drop of water to extract material for the analysis of the brown stains, white spots became visible (figure 2). This showed that the cigarette smoke could be easily dissolved in an aqueous cleaning system. Acrylic painted surfaces, however, are very sensitive to water, but also to solvents, heat and vibration.

Aqueous and solvent-based cleaning systems involve a risk of partial removal of surfactant and other constituents from the bulk of the emulsion paints\(^\text{8}\). The water can also penetrate into the pores and induce temporarily swelling of the paint. Aqueous preparations can be used on these delicate surfaces if they are isotonic and buffered at about pH 6. The challenge is to determine accurately the isotonic point of these paints. Wolbers\(^\text{20}\) for instance uses small blocks of agarose gel to “sample” the surface pH and conductivity. Ideally, the conductivity should always be adapted to that of an individual colour or paint layer. Environmental factors can influence the cleaning process on these acrylics: both temperature and relative humidity play a role in the migration of surfactants and soiling materials to the paint surface. Creating a stable environment during the cleaning process also helps to improve the cleaning efficacy. By raising the relative humidity to 60% or greater, the surfactant and soils dissolved in them will migrate to the surface and will therefore be easier to clean.

So-called micro-emulsions have been developed to increase cleaning efficacy while limiting the contact of aqueous materials on these paints\(^\text{20}\). These micro-emulsion systems are composed with surfactants that have a low hydrophilic-lipophilic balance; this helps to prevent the loss of surfactants that are inherent to emulsion paints. Micro-emulsions are liquid dispersions of water and oil that are made homogenous, transparent (or translucent) and thermodynamically stable by the addition of relatively large amounts of a surfactant and a cosurfactant. The micro-emulsion droplets have a diameter in the range of 10-100 nm;
much smaller than in conventional emulsions. For acrylics, a surfactant-rich oil phase that coexists with the surfactant-poor aqueous phase should be used (water-in-oil micro-emulsion, figure 9). This can be referred to as "dry cleaning".

In addition, a dimethyl siloxane-based solvent will be used to protect portions of paints that might otherwise be affected by water-based preparations\(^\text{20}\). This is a very low-aliphatic structure, which has even a lower ionic strength than white spirit. This solvent can be placed on top of the paint surface, on which we subsequently apply the micro-emulsion. As long as the dimethyl siloxane-based solvent is not completely vapourised, it is safe to work on the object since it repels the water. Afterwards, removal of the micro-emulsion will be done by rinsing with dimethyl siloxane as well. Because of the surfactants that are present in the micro-emulsion this step is crucial.

Different cleaning problems with acrylic painted surfaces can be approached by adapting the water-in-oil micro-emulsion. Raising the relative humidity will also aid in capturing the soils present in the surfactant of the acrylic emulsion. In the near future, a series of tests using water-in-oil micro-emulsions will commence. Once the restoration treatment of *Camping II* is completed, the work will be restored to its former glory without any brown nicotine spots that disrupt the monochromic white paint.

5 Conclusion

The brown stains that are observed on the white painted areas of Raoul De Keyser’s *Camping II* are related to the use of a titanium white waterborne \((p\text{-EA-MMA})\) acrylic paint, while the other areas, based primarily on PVAc paints, remain unaffected by the staining process. The surfactant used to suspend and stabilise the \((p\text{-EA-MMA})\) in the aqueous medium was identified as octyl phenyl polyethoxy ethanol by py-GC/MS. Depending on a series of parameters, this hydrophilic PEO-type surfactant has a tendency to migrate from the bulk to the surface of the paint. It has been detected with py-GC/MS in aqueous extracts of the brown stains that were locally removed using an aqueous extraction procedure. It is hypothesised that the brown colour of the exudate is caused by cigarette smoke components absorbed onto the surfactant-rich surface layer. This is substantiated by the relatively high amounts of (hydrophilic) nicotine and related components found by py-GC/MS analysis of the extracts.

Most conservation treatments designed for traditional oil paintings are not suited for acrylic emulsion paintings. Characterisation of the subtle and unique nature of the surfaces of acrylic emulsion paint films is crucial to minimise the risks associated with conservation procedures, such as cleaning. A cleaning method to remove only the brown stains, presumably caused by cigarette smoke, can therefore be developed. In the near future, a series of tests with water-in-oil micro-emulsions will be carried out: these are the safest and most effective cleaning systems on this important acrylic painting from Raoul de Keyser.

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7 References

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