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Potential and Drawbacks of Raman (Micro)Spectrometry for the Understanding of Iron and Steel Corrosion

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1. Introduction

Raman scattering theory and first observation date back to the first decades of the 20th Century but the technique did not generalise until the 70s, with the development of lasers technology. Two major breakthroughs occurred in the 80s with the replacement of monochannel PM detectors by multichannel CCDs (Delhaye et al., 1996) and in the 90s with the rejection of Rayleigh elastic scattering by photonic crystals (Notch filter) or multilayer coatings (Edge filter) rather than monochromator(s). These latter technological changes improved the sensitivity by many orders of magnitude, which allowed either cutting the counting time (allowing mapping or real time monitoring) or reducing the illumination power (thus allowing for the analysis of black compounds). The latest developments concern the miniaturization of solid laser sources and the replacement of electronic boxes controlling the CCD detector by softwares uploaded on a common laptop, leading to portable Raman instruments. All these developments and the increase of the Raman instrument production lowered the price and made the technique more and more available, even for in-line/at-line/on-line control.

Among the many interactions of light with matter, Raman scattering is particularly well suited to the multiscale analysis of ill-organized heterogeneous solids as the corrosion films (Gouadec & Colomban, 2007a; ibidem, 2007b). The Raman probe being for interatomic bonds themselves, the technique offers a “bottom-up” approach to study nanomaterials and amorphous compounds which best works in the case of imperfect crystals with strong covalent bonds (Fig. 1) such as those typically produced by metal corrosion. Raman signal results from the interaction of a monochromatic coherent light (laser beam) with electronic and vibrational levels of atomic bonds (Long, 1977; Lewis & Edwards, 2001; Gouadec & Colomban, 2007 for a more complete theoretical description). The interaction with the electronic levels is often described as virtual; this is true for non-coloured samples or non-absorbent for the excitation laser line, but wrong for absorbent materials, leading to (pre)resonance Raman features. Consequently, peak intensity will depend on the exciting wavelength and 2nd order Raman features could be present (harmonics, combinations). Theory predicts both elastic (so called Rayleigh scattering) and inelastic (namely Raman) contributions in the scattered electric field. The latter occurs only if vibrations change bond polarizability, which is a second rank tensor containing the crystal symmetry. Raman scattering is
complementary to infrared absorption with the advantage of much narrower peaks: the Raman peak shape is thus very informative because the Raman probe is very local. On the other hand, the IR spectroscopy probes, the instantaneous dipole moments which are subjects of much longer distance interactions. Consequently, IR bands are often very broad.

Raman spectroscopists in general refer the vibrational modes, the phonons, by i) their wavenumber \( \nu = \frac{\nu}{c} \) (c the light speed, \( \nu \) in cm\(^{-1}\) unit, here after and usually noted \( \nu \)) and expressed it in cm\(^{-1}\), however energy (meV), wavelength (nm) are frequency (THz) units can also be used and ii) their symmetry (total symmetric modes are the strongest ones).

![Fig. 1. The basics of the information to be extracted from a Raman spectrum (after Gouadec et al., 2010).](image)

The polarization of a sample illuminated with light (electric field \( E_0 \); frequency \( \nu_l \)) has the following form:

\[
\mathbf{P} = \alpha \times E_0 \cos(2\nu_l t)
\]

In Eq. (1), \( \alpha \) represents the polarizability tensor, which depends on matter vibrations (the oscillations of atoms and molecules around their equilibrium positions). The polarization can be expressed as a function of the atomic displacement (normal coordinates) using a Taylor approximation, thus predicting elastic scattering (\( \nu = \nu_l \), the exciting laser wavenumber) and inelastic (\( \nu = \nu_l \pm \nu_{\text{vib}} \)) scattering by atomic vibrations. The former is called Rayleigh scattering and the latter, which occurs only if vibrations change polarizability (\( \partial \alpha / \partial Q \neq 0 \)), is Raman scattering. Other terms correspond to Hyper and higher orders Raman scattering (Long, 1977; Gouadec & Colomban, 2007a).

The signal intensity is predicted with the following formula:

\[
I_{\text{Raman}} \propto I_{\text{laser}} \nu_{\text{laser}}^4 |e_0 \mathbf{\alpha} \mathbf{e}|^2 d\Omega
\]

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In Eq. (2), \( e_o \) and \( e_s \) are unit vectors indicating the laser polarization and direction of observation, respectively, whereas \( d\Omega \) represents the solid angle of light collection, at a maximum when high magnification, high aperture number microscope objectives are used. Classical electromagnetic theory predicts Raman peaks should have a Lorentzian shape (Long, 1977; Lewis & Edwards, 2001; Gouadec & Colomban, 2007a). Isotropic disorder leads to a distribution of Lorentzian usually described by a Gaussian shape. Actually more complex shapes (e.g. asymmetric ones in the case of anisotropic disorder, Fig. 1) occur and complex laws should be used (Havel et al., 2004; Gouadec & Colomban, 2007a; Havel et al., 2007) to describe the Raman signature of some nanophased materials or when defects break the phonon propagation (Havel et al., 2007; Chi et al., 2011).

The scattering intensity varies by orders of magnitude depending on the bond polarisability (the more covalent the bonds, the higher the number of electrons involved and the higher the Raman peak intensity), the crystal symmetry and the exciting wavelength. As polarizability (5 second rank tensor) changes drastically from one bond to another, Raman intensity may not be used to measure the relative amounts of different phases without preliminary calibration. Consequently, minor phases or even traces could have a stronger Raman signature than some major phases. The preferential orientation of certain phases, common for surface grown phases enhances some peaks and calibration cannot be efficient.

The absorption of the laser light by coloured phases can be very high and thus the penetration depth can be less than a few tenths of nm (Gouadec & Colomban, 2001; Havel & Colomban, 2006). Furthermore light absorption may involve strong local heating and thus phase transformation towards more stable ones, crystallization of amorphous ones or oxidation (de Faria et al., 1997; de Faria & Lopez, 2007; Cvejic et al., 2006). Raman analysis appears very sensitive to answer some questions on a given material whereas can be neither useless to study some others. Mapping (see further) and quantitative analysis should be then performed with caution.

Since Raman instruments were made available in the 70s, attempts to characterize the corrosion products of iron-based artefacts were performed. Most of studies concerned pure iron or low carbon content alloys but reference spectra are now available for large majority of the most common corrosion products: haematite, (Beattie & Gilson, 1970), magnetite (Morke et al., 1980), lepidocrocite (Thibeau et al. 1978). First series of corrosion studies flourish during the 80’s (Farrow, 1980; Farrow & Nagelberg, 1980; Keiser et al., 1982; Hugot-Le Goff & Pallotta, 1985; Naouer et al. 1985; Ohtsuka et al., 1986; Boucherit et al. 1989, Dünnwald & Otto, 1989). Then, high sensitivity instruments made it possible to study black and low crystallinity/amorphous films that were easily transformed into the stable ones by laser heating (Gouadec et al. 2001; Mazetti & Thistlethwaite, 2002; Cvejic et al., 2006: Gouadec & Colomban, 2007; Gouadec et al., 2010). More recently, mapping allowed getting a semi-quantitative global view (Neff et al., 2005; Neff et al., 2006 ; Monnier et al., 2010). A great effort was made to obtain reference phases and their signature, especially by controlled electrochemically synthesis (Savoy et al., 2001; Sinard et al., 2001; Joinet et al., 2002; Legrand et al. 2003, Refait et al. 2003, Poupard et al. 2006, Pineau et al. 2008; Dubois et al. 2008).

In this chapter we address the advantages of Raman spectroscopy and mapping, with particular attention to the intrinsic experimental and conceptual drawbacks of the methods as well as possible ways to overcome them.
2. Phases issued from metal corrosion and their Raman signature

2.1 Phase and structure relationship

The corrosion of metal results from the reaction with anions and the formation of new bonds at the surface. The simplest case is the oxidation (formation of Fe(OH)$_3$) and electronic conductivity as well as the presence of liquid electrolytes such as water are prominent parameters. We will discuss the Raman signature of iron-based compounds but all the phases (oxides, hydroxides, carbonates, sulphates, etc.) of any element used in the alloy must be considered because corrosion may promote their formation and trace phases may have a signature sufficiently strong to be detected.

The densest phase is wustite because of its high Fe/O ratio (Fe$_{10}$O, space group Fm$ar{3}$m, density: ~6, black). Two other dense phases are built with two available ways of oxygen atom packing: magnetite (Fe$_{12}$O$_{16}$ spinel structure with space group Fd-3m, cubic ABCABC packing, density 5.18, black) and haematite (Fe$_{2}$O$_{3}$ corundum structure with space group R-3c, hexagonal ABAB packing, density 5.23, dark red, Froment et al., 2008). When oxygen vacancies are formed - that decreases the density - the oxygen layer packing is preserved but some protonation may occurs: i) maghemite ($\gamma$ FeO$_x$, space group P4$_3$2 preserves the cubic spinel structure but its density lowers to 4.87, brown), ii) goethite ($\alpha$ FeOOH, space group 2/m2/m2/m, density : 4.3 to 3.3, ochre to black ) and lepidocrocite ($\gamma$ FeOOH, Cmcm, density :~4, dark red) ; these phases retain the initial framework made of oxygen atom layers but the structure becomes more open due to oxygen defects. Furthermore, some oxygen atoms may be replaced by Cl ones (akaganeite $\beta$ FeOOH, density : 3.8 to 3.6, orange), especially at the phase surface. Other disordered phases are observed : ferrihydrites (Fe$^{iii}$/OOH, density : 4.2, yellow), ferricydrites (Fe$^{iii}$/OH)$_6$ 9H$_2$O, ABACA or mixed ABA ACA packing, density : 3.8, brown), hydroxychlorides, $\beta$ Fe$^{ii}$/OH)$_2$Cl or the so-called "green rust" (Fe$^{ii}$/$_{1-x}$Fe$^{iii}$/$_x$(OH)$_2$ Cl$_y$ (SO$_4$)$_z$, nH$_2$O, AABCC packing, a variant of the ABC cubic packing, green). White rust consists in the iron lamellar hydroxides (main phase: $\beta$ Fe$^{ii}$/OH)$_2$, P-3m1, ABAB compact packing, white). Carbonates ( siderite, Fe$^{ii}$CO$_3$, density ~3.9; Fe$^{ii}$(CO$_3$)(OH)$_2$, and some other mixed frameworks) may also form. Sulfates (Fe$^{ii}$SO$_4$) result from sulphuric acid treatment in the finishing of steel before coating or plating.

Galvanized steel sheets received ZnFe or more complex (ZnNiMn, etc…) coatings that enlarge the variety of phases to be formed (Bernard et al., 1993; Marchebois et al., 2002a; ibidem, 2002b; Tomandl et al., 2004; Yadav et al., 2004; Hernandez et al., 2006; Refait et al., 2007; Colomban et al., 2008; Dubois et al., 2008) : smithsonite ZnCO$_3$, Zn(OH)$_2$ and complex phases like hydrozincite Zn$_3$(OH)$_4$(CO$_3$)$_4$ ("white corrosion"), ZnCl$_2$(OH)$_4$ SO$_4$, 5 H$_2$O, phosphates like Zn$_3$(PO$_4$)$_2$, H$_2$O, ZnS ("black corrosion").

In the same way, water and/or high temperature resistant steels such as Ni and Cr-rich and the corresponding oxides/hydroxides are formed : $\alpha$ CrOOH, Cr$_2$O$_3$, NiO, spines, … (Beatie & Gilson, 1970; Bernard et al., 1993; Zuo et al., 1996; Delichère et al., 1997; Colomban et al., 1999; Maslar et al., 2001). Reference spectra of common phases can be also found in review books (see e.g. Karr Jr, 1975; Nakamoto, 1997) and the data on parent compounds are very useful for the identification of solid solutions and ill-crystallized compounds (Delichère et al., 1988; Desilvestro et al., 1988; Colomban et al., 1999). Table 1 lists the most frequent corrosion products and their characteristic Raman fingerprints.
## Table 1. Raman fingerprint of main corrosion products.

<table>
<thead>
<tr>
<th>Compound</th>
<th>Formula/structure</th>
<th>Characteristic Wavenumber</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wustite</td>
<td>FeO</td>
<td>655</td>
<td></td>
</tr>
<tr>
<td>Haematite</td>
<td>Fe₂O₃ / Corundum</td>
<td>~1320, 290, 220</td>
<td></td>
</tr>
<tr>
<td>Magnetite</td>
<td>Fe₃O₄ / spinel</td>
<td>670</td>
<td></td>
</tr>
<tr>
<td>Maghemite</td>
<td>γFe₂O₃·H₂O / spinel</td>
<td>670-720, ~1400</td>
<td>Easily transforms into haematite under laser beam idem</td>
</tr>
<tr>
<td>Ferrihydrite</td>
<td>Fe₅HO₆·9H₂O</td>
<td>710, ~1380</td>
<td></td>
</tr>
<tr>
<td>Goethite</td>
<td>αFeOOH</td>
<td>390</td>
<td></td>
</tr>
<tr>
<td>Lepidocrocite</td>
<td>γFeOOH</td>
<td>250, 1300</td>
<td></td>
</tr>
<tr>
<td>Akaganeite</td>
<td>βFeOOH (Cl)</td>
<td>310, 390, 720</td>
<td></td>
</tr>
<tr>
<td>Ferroxyhyte</td>
<td>δFeOH</td>
<td>680, ~1350</td>
<td></td>
</tr>
<tr>
<td>Hydroxychloride</td>
<td>βFe₂(OH)₃Cl</td>
<td>160, 423</td>
<td></td>
</tr>
<tr>
<td>Green rust</td>
<td></td>
<td>430-510</td>
<td></td>
</tr>
<tr>
<td>Iron chloride</td>
<td>FeCl₂</td>
<td>610</td>
<td></td>
</tr>
<tr>
<td>Zinc chloride</td>
<td>ZnCl₂</td>
<td>80, 248</td>
<td></td>
</tr>
<tr>
<td>Zinc oxide</td>
<td>ZnO</td>
<td>100, 540-580</td>
<td></td>
</tr>
<tr>
<td>Zinc hydroxide</td>
<td>Zn(OH)₂</td>
<td>470</td>
<td></td>
</tr>
<tr>
<td>Zinc carbonates</td>
<td>ZnCO₃</td>
<td>1095, 370</td>
<td></td>
</tr>
<tr>
<td>White rust</td>
<td>3Zn(OH)₂ 2ZnCO₃</td>
<td>1050, 385</td>
<td></td>
</tr>
<tr>
<td></td>
<td>4Zn(OH)₂ ZnCl₂ (OH)₄</td>
<td>910, 3455-3486</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Zn(OH)₄ Cl₂ SO₄ 5H₂O</td>
<td>955, 208, 292</td>
<td></td>
</tr>
<tr>
<td></td>
<td>ZnSO₄ 3 Zn(OH)₂ 3H₂O</td>
<td>961, 1007, 463</td>
<td></td>
</tr>
<tr>
<td>Zinc phosphate</td>
<td>ZnPO₄</td>
<td>996</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Zn₃(PO₄)₂·H₂O</td>
<td>1055, 1150</td>
<td></td>
</tr>
<tr>
<td>Manganese oxide</td>
<td>MnO₂</td>
<td>~600</td>
<td></td>
</tr>
<tr>
<td>Nickel oxide</td>
<td>NiO</td>
<td>~510</td>
<td></td>
</tr>
<tr>
<td>Chromium oxide</td>
<td>Cr₂O₃</td>
<td>351, 551, 609</td>
<td></td>
</tr>
<tr>
<td>Chromium hydroxide</td>
<td>CrOOH (Cr₂O₇ · nH₂O)</td>
<td>485</td>
<td></td>
</tr>
<tr>
<td>Mackinawite</td>
<td>FeS</td>
<td>~280-300</td>
<td></td>
</tr>
<tr>
<td>Greigite</td>
<td>Fe₃S₄</td>
<td>~350-360</td>
<td></td>
</tr>
</tbody>
</table>

### 2.2 Understanding Raman signature

If the analytic approach is useful (identification and quantification of the phases formed by corrosion), a more comprehensive understanding of the phase structures and relationship is necessary for a comprehension of the reaction scheme and the prospect of regulating them. The recording of the Raman signatures will depend on the phase colour – i.e. their electronic band structure – that determines the light penetration and the intensity of the scattered...
signal as well as on the phase structure: the higher the symmetry, the lower number of Raman peaks (Lewis & Edwards, 2001; Gouade & Colomban, 2007a).

Fig. 2 shows the Raman signature of the main phases encountered in the corrosion layers. Typical intensity ratios in comparable recording conditions are:

Lepidocrocite (1) > wustite, haematite, goethite, hydroxychloride, ferrihydrites, ferroxyhite (~1/3) > akaganeite (~1/4) >> magnetite, maghemite (~1/10).

Fig. 2. Raman signature of the main (crystalline) iron oxihydroxides observed in corrosion films (after Colomban et al., 2008).

It is thus clear that the analysis of a mixture will strongly lower the contribution of the latter phases.

The Raman signature of any compound that structure is built with one or many strong covalent-bonded vibrational units, can be separated in four groups (Gouade & Colomban, 2007a):

i. (symmetric) stretching modes, for instance Fe$^{II}$-O and/or Fe$^{III}$-O modes peaking in the 400-700 cm$^{-1}$ region for oxides, 300-600 cm$^{-1}$ for chlorides, 200-400 cm$^{-1}$ for sulphides. In symmetric modes, the strongest ones, only oxygen atoms move and thus the peak wavenumber mainly depends on the Fe-O distance (Vunicic-Vasic et al., 2006). These modes are very sensitive to oxygen vacancies that broaden the peaks.

ii. bending modes peak at lower energy, namely 400-500 cm$^{-1}$ range for oxides or less for chlorides, sulphides,... They are very sensitive to the short range disorder in the first neighbouring shell (1-5 nm around the chemical bond) and their broadness can be very informative on the short-range (dis)order. Because the mean symmetry often breaks the symmetry of the vibrational units, many components are frequently observed.
iii. librational (orientational oscillations, 150-400 cm\(^{-1}\) range) and lattice modes (<200 cm\(^{-1}\)). Librational modes are very sensitive to the short range disorder of the vibrational entities (e.g. FeO\(_4\) tetraedron); lattice modes reflect the long range crystalline order, like X-ray diffraction, and strongly depend on atom mass (\(\sqrt{M}\)).

iv. additional features (combination, harmonics,...) consisting bands between 1000 and 1500 cm\(^{-1}\): their origin lies in the interaction between electronic (and magnetic, see e.g. (Morke et al., 1980)) levels and the light because the laser beam wavelength interacts with the electronic levels. The high intensity of these bands facilitates the phase identification. On the other hand the high light-matter coupling leads to a strong heating under the laser spot: temperatures close to 500°C can be easily achieved. The temperature does not depend on the laser power and objective magnification only but also on the heat dissipation (Gouadec et al., 2001): the dissipation is often better for small aggregates but matrix effects are important. Measurements at low temperature (liquid N\(_2\)) or in water should be preferred.

Taking into account the above rules it is obvious that the narrow low wavenumber peaks of lepidocrocite, hematite, goethite and some hydrochlorides are characteristic of crystalline phases. The strong intensity of their lattice modes is typical for layered materials. Replacing atoms by heavier ones (O/Cl for instance in hydrochlorides) shifts the lattice modes toward lower wavenumbers and broadens the different components. The relationships between haematite, lepidocrocite, goethite and akaganeite on one side (both phases having AB derived packing, Fig. 2), and on the other hand magnetite, maghemite, ferrihydrites (ABC packing) are obvious (Colomban et al., 2008). Partial substitution modifies the bandwidth (Cvejic et al., 2006).

Detection of “pure” hydroxides (white rust), carbonates and sulphates (black rust) is very easy because of the narrowness of their stretching mode (1020-1100 cm\(^{-1}\) range) and because reference spectra are well documented (Karr Jr, 1975; Nakamoto, 1997). Furthermore, high power of illumination can be used for non-coloured phases. Actually, in most of the practical cases, corrosion films are made of complex intergrowths at the nanoscale, giving rise to mixed oxyhydroxycarbonates/sulfates/etc. with complex Raman signatures because of the solid solutions formation, intimate mixtures of ill-crystallised phase with a very small grain size. Synthesis and characterisation of reference compounds are complex (handling in CO\(_2\)- and/or water-free media, redox control…). Fig. 4 shows spectra recorded on galvanized steel sheets corroded in real conditions. In most of the cases fluorescence, a phenomenon in competition with Raman scattering but more efficient, gives rise to a strong background for real samples (galvanised steel coupons have been cut from vehicles that have been in circulation for years in severe weather conditions). The fluorescence often due to microbiological activity can be eliminated by H\(_2\)O\(_2\) or ~500°C thermal treatment (furnace heating or laser cleaning). On the other hand, spectra free of fluorescence background are easier recorded on coupons accelerated corroded at the laboratory. Indeed, reagents are simpler and cleaner. However, the most stringent point is that formed phases are often different. It is clear that many improvements on the procedure to conduct accelerated tests, more representative for real corrosion have to be made.

For specific applications special coatings like Cr(Al)N, NiCrAlY (Colomban et al., 1999) or polyaniline films (Delichère et al., 2007; Bernard et al., 2002; Nguyen et al., 2004) are used. This requires special database.

Note that Raman analysis is possible through a layer, inorganic (enamel, ceramic matrix (Wu et al, (1997); Gouadec et al.(1998)) or organic one (paint, e.g. Colomban et al., 2008). Typically, good Raman signatures are obtained up to 20 to 40 \(\mu\)m below the surface.

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Fig. 3. Example of the decomposition of the Raman spectrum of a corroded film using the 3 signatures of pure reference phases. The appropriate program was then ran to adjust the data in all the mapped area with these combination spectra and to image the protective ability index in all the studied area (adapted from Gouadec et al., 2010). Note the very narrow 250 cm\textsuperscript{-1} peak characteristic of Lepidocrocite layer structure and the strong 2\textsuperscript{nd} order features at ~1350-1400 cm\textsuperscript{-1} of spinel derived ill crystallized phases.

Fig. 4. Example of Raman signatures recorded in different spots of a corroded galvanized steel sheet with the sequences a) steel/Zn-based coating/phosphate coating/epoxy-TiO\textsubscript{2} paint and b) steel/ZnFe coating/ZnNiMn coating/epoxy-TiO\textsubscript{2} paint; accelerated (Acc) or real (R) corrosion conditions (after Colomban et al., 2008).
2.3 Semiquantitative analysis and phase mapping
Raman mapping is most valuable to point out subtle composition and structure modifications taking place from one place of a sample to another. The images are obtained by extracting pertinent parameters from the hyperspectral map, which is a collection of individual Raman spectra, each being associated with a given point on the sample surface (Turrell, 1996; Colomban, 2003).

The great advantage of the automatic mapping procedure is the achievement of a more representative view of the present phases. However, the visual selection of the spot to be analysed is very subjective and leads to neglect places where the optical focus view through the microscope objective looks poor (low contrast, roughness, ...).

Using the parameter(s) extracted from each spectrum (peak intensity, wavenumber shift but also grain size, defect concentration, etc.), it is possible to build a map of the investigated area the so-called smart image (Colomban, 2003). The following requirements are mandatory to record and exploit Raman mapping (Gouadec et al., 2010):

i. the analysed area must be horizontal and the roughness smaller than the vertical height of the laser spot (typically, ~10 µm for x100, ~50 µm for x50 magnification objective,...)
ii. the horizontal resolution, i.e. the laser spot diameter (less than ~1 µm for x200, ~5 µm for x100, ~10 µm for x50 magnification objectives) combined to the stage step (~0.1µm or more) must be compatible with the material/phase/grain size.
iii. the analyzed area must be chosen in order to be representative of the different topological features but limiting the number of spectrum to be collected and processed to built smart images (Colomban, 2003; Havel et al., 2004; Havel & Colomban, 2006; Gouadec et al., 2010).

The image resolution will thus depend on the optical parts (objective and spectrometer), the laser beam quality (alignment, fundamental mode), the sample (parasite refractions at interfaces, roughness), Raman signature contrast, light absorption and penetration in the sample as well as the stage motion step, usually up to 0.1 µm...

The mapping has been used to understand the long term atmospheric and in soil corrosion taking advantage of the big thickness and large grain size of corrosion films (tenths of microns) present in heritage buildings (cathedrals, churches,...) and archaeological artefacts (Neff et al., 2005; ibidem, 2006; Monnier et al., 2010). Archaeological and Cultural Heritage artefacts are however considered as good analogues for the understanding and prediction of iron alloys corrosion behaviour in soil and in atmosphere, and hence to determine the lifetime of over-containers used to protect the vitrified nuclear waste.

The Raman technique has also been used to compare the thinner (~1-3 µm thick) corrosion films obtained on automotive galvanized steels in accelerated corrosion tests, at the laboratory, and in real use (intense corrosion for used cars in severe mountain weathering conditions (temperature, water, salt (Colomban et al., 2008)). Raman analysis in the bottom of the pits formed by the corrosion is possible using long focus, large aperture high quality objectives.

3. New challenges
The current challenges are i) the development of data bases of complex mixed and nanosized phases, ii) the improvement of procedures/data treatment for two dimensional (2D) and three dimensional (3D) Raman mapping and iii) a better understanding of the 2nd order Raman signature of coloured phases in order to obtained smart imaging of pertinent...
parameters, for instance to image the phase amount ratio that has been proposed to establish if the corrosion is passive or active (Kashima et al., 2000).

3.1 Mixed oxides and oxyhydroxycarbonates
Although the reference signatures of pure iron oxides are well established, those of the mixed compounds like spinels associating Fe, Mn, Cr, Ni,… atoms or of oxyhydroxycarbonates, hydrated or not, are very poorly documented (Colomban et al., 1999; Colomban et al. 2008; Cvejic et al., 2006). These phases may be unstable under the laser spot and may transform into more stable ones having different structure. Furthermore, for many films it is not established if the matter consists in a mixture, a phase or nanoparticles with another phase covering the grain surface. In many cases, because of the sample colour and its absorption of the laser light by electronic level, the interaction with electronic levels is anymore virtual and Raman (pre)resonance phenomenon is observed: the Raman intensity concentrates in some modes, a (small) wavenumber shift is observed and harmonic/combination second order peaks become visible. Note that in situ measurements under controlled electrochemical condition allow to record nice vibrational signature. However the representativity of these signatures is not straightforward because the real conditions are more complex what can promote the formation of other minor phases with stronger Raman cross section.

3.2 Controlling the analysed area
Most of the phases formed on the metal surface are coloured and hence absorb the light. Consequently the penetration depth of the light depends on the absorption coefficient. For dark phase the depth can be very limited and the information obtained from a mapping is perturbed: the analysed thickness will vary from spot to spot and correction is not possible. From a practical point of view, microscope objectives are usually characterized by the numerical aperture NA:

\[ NA = n \sin(\theta_{\text{back}}) \] (3)

In Eq (3), \( \theta_{\text{back}} \) is the maximum collection angle for the backscattered light and \( n \) is the refractive index in the medium between the sample and the microscope lens (Fig. 5). The numerical aperture is a key parameter because it sets the resolution \( R \) of the microscope, defined as the shortest spacing for two points on a sample surface to be resolved with \( \lambda \) wavelength observation (see references in Gouadec & Colomban, 2007a; Gouadec et al., 2010):

\[ R = 0.6 \frac{\lambda}{NA} \] (4)

Since the optics of ultraviolet light devices are expensive and not very efficient, the lasers offering violet (~450 to ~400nm) or deep violet (~365 nm) excitations are preferred to obtain the best resolution. Actual laser beams are not perfectly parallel (this has been exaggerated in Fig. 5) and their focusing through a microscope lens gives an elongated volume called the focal domain or focal cylinder. The diameter \( \phi(z) \) of the focal domain at \( z \) axial coordinate must be defined arbitrarily as the electric field obviously does not drop to zero for a definite \( r \) distance away from the optical axis. The radial decrease of the electric field in a laser beam actually obeys a Gaussian law. Similarly to the lateral resolution, the in depth \( \Delta z \) or axial \( (\sim \phi(z)) \) resolution of Raman spectroscopy can not be defined unambiguously. Indeed, the laser intensity does not drop to zero for a given \( z \) value and one has to choose an arbitrary
Potential and Drawbacks of Raman (Micro)Spectrometry for the Understanding of Iron and Steel Corrosion

The Raman efficiency shows order of magnitude variation that makes the choice very difficult. Δz also depends on the generally unknown refractive index of the studied compound.

In confocal Raman microscopy set up, pinholes are placed in the microscope at intermediate image planes, resulting in a better in-depth discrimination power but a great increase of the counting times, 10 times and even more! In that case, the integrated intensity coming from a given plane perpendicular to the optical axis is no longer a constant but, rather, decreases by 50% between the focal plane (z=0) and Δz_{confocal} (the half width of the so-called point spread function). The use of confocal pinholes goes with a lateral resolution improvement of about one third but confocal microscopy interest mainly consists in the possibility to select sample layers axially. However, even with the best dry (n=n_{air}=1, NA=0.95) or oil immersion (n=1.51, NA=1.4) objectives, and in the most favourable case of a violet excitation (e.g. λ~407 nm), Δz_{spot} will remain above 275 nm (Eq. (4)). The use of high quality (expansive) objectives is often more efficient than the confocal setting. Note bias polishing enlarge the topological resolution and micronic films can be easily analysed by the Raman technique (Gouadec et al., 2001).

Consequently the spatial repartition and relative proportion deduced from a Raman mapping may be interpreted with caution. The information can be however very useful to characterise the evolution of the corrosion film as a function of time, temperature, external parameters, etc.

Fig. 5. a) Intensity distribution of the illuminating laser spot as a function of the de-focus (after Colomban, 2003); b) schematic comparison of the focus in a non-absorbing medium with and without the use of a confocal pinhole in the optical system. Note the very limited penetration for coloured material (after Gouadec & Colomban, 2007a).

3.3 Understanding at the nanostructure scale

The decrease of the grain size (D, see Fig. 6) makes that atoms at the particle surface or atoms having their chemical bonding perturbed by the vicinity of the surface (t: distance where the surface perturbs the structure and chemical bonds) become more important than the bulk atoms when the particle size drops below 30 to 5nm as a function of elements and chemical bonding. Vacancy concentration is maximal close to the surface and adsorbed species can be present: hydroxylation, water, etc.
The disorder modifies the Raman signature: symmetry exclusion can disappear leading to the activity of new components. The most prominent effect is the modification of the Raman peaks shapes. A reliable analysis of the Raman spectra requires the use of appropriate modelling to describe the band shape of the different spectral components (Gouadec & Colomban, 2007a, Chi et al. 2010).

The use of simple Gaussian and Lorentzian band profiles is often preferred than the Voigt profile; many fitting modules misleadingly name Voigt profile the simple sum of one Lorentzian and one Gaussian, as both depend on three parameters rather than four. The exact position, intensity, width and lineshape of each band depend on many different parameters such as the actual chemistry (neighbouring inclusions, substitutions or vacancies), crystallinity, domain size, phase orientation and corresponding polarization effects or thermomechanical stress (anharmonic effects). In grains much larger than the wavelength, phonons propagate almost in the same way as in perfect "infinite" crystals. When the grain size falls below a few tens of nanometers, the Phonon Confinement Model (PCM) accounts for the phonons coherence length limitation by a weighed exploration of longitudinal optical dispersion curves (see Gouadec & Colomban, 2007a nd 2007b and references herein). Below a certain size, the very notion of collective vibrations disappears and the Elastic Sphere Model (ESM) takes over, using first principle description of low wavenumber vibrations in a "free-standing", homogeneous (constant density) and elastic sphere (Fig. 7). In this scheme, the wavenumbers of the two most intense modes are inversely proportional to the grain size (Fig. 6).

Fig. 6. Schematic of the relative contribution of skin/bulk matter for a particle as a function of its size: for a nanosized grain less than 1-10 nm in diameter the contribution of the surface atoms is very dominant (after Colomban, 2003).

Fig. 7. Modelling spherical nanocrystals vibrations as a function of the grain size (after Gouadec & Colomban, 2007)
3.4. Understanding the corrosion mechanisms and anticipating the evolution

Although the phenomenological description of corrosion has received attention since long time from the electrochemical and thermodynamical points of view, the understanding of the mechanisms from the solid state chemistry point of view remains very limited. For instance, many phases contain protons in different forms and data on proton conductivity are very scarce (Colomban, 1992). Vibrationnal spectroscopy can be very useful to document the structural changes associated to ion diffusion.

The action of micro-organisms was poorly documented from the spectroscopic point of view. Microbiologically influenced corrosion is used since millennia to improve the plasticity of kaolin and clays by chemical modifications of silicates. This results from both, the direct action of microorganisms and the indirect one via the species generated by their metabolic activity. For iron-based materials, the phenomenon is induced in anoxic environments by sulphate-reducing bacteria (SRB), microorganisms that produced sulphide species. Thus, in media where sulphides are not naturally present, the observation of Fe$^{II}$ sulphides (all sulphides have their stronger peak at ~250-350 cm$^{-1}$) inside the rust layer may be a clear indication that SRB were active and play a role in the corrosion process. The heterogeneity of the biofilm that covers the metal surface can lead to a galvanic interaction between regions of microbial activity and the surface. This leads to locally accelerated corrosion pits (Videla & Characklis, 1992; Marchal, 1999; Beech & Sunner, 2004; Little & Lee, 2007).

When metal is associated with organic materials (some paints, films used for "conservation" treatments of archaeological artefacts, …), consequence of microbiologically induced corrosion can be severe when contact with air: cracks occur due to the volume increase associated to iron oxidation and simultaneously sulphuric acid is produced. Raman study confirms that FeS (mackinawite) is very reactive towards oxygen (Remazeilles et al., 2010).

4. Conclusion

During decades Raman spectroscopy remains a useful technique for solid state physics and chemistry, giving valuable information about the structure and the reactivity of colourless single crystals. Then, clean polycrystalline, ill-crystallized samples can be analyzed. The high sensitivity of the modern instruments makes it possible to study any kind of samples if a palette of exciting laser wavelength is available to avoid detrimental pollution of the Raman information by strong fluorescence phenomenon. Very recently the potential of algorithmic methods to extract pertinent information from the spectra dominated by the fluorescence has been demonstrated (Widjaja et al., 2010) and the application of the Band-Target Entropy Minimization (BTEM) or similar techniques to the “poor” Raman signature of certain corrosion films appears very interesting.

5. References


Ohtsuka, T.; Kubo, K. & Sato N. (1986). Raman spectroscopy in thin corrosion films on iron at 100 to 150°C in air, Corrosion-NACE (National Association of Corrosion Engineers), 42, [8], 476-481.


In the last few years the automobile design process is required to become more responsible and responsibly related to environmental needs. Basing the automotive design not only on the appearance, the visual appearance of the vehicle needs to be thought together and deeply integrated with the “powerâ€ developed by the engine. The purpose of this book is to try to present the new technologies development scenario, and not to give any indication about the direction that should be given to the research in this complex and multi-disciplinary challenging field.

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