



**HAL**  
open science

## Raman Spectroscopy: an elegant and effective method, from single crystals to the planet Mars

Philippe Colomban

► **To cite this version:**

Philippe Colomban. Raman Spectroscopy: an elegant and effective method, from single crystals to the planet Mars. Asian Journal of Physics , 2023, 32 (1-2), pp.1-4. hal-04090559

**HAL Id: hal-04090559**

**<https://hal.science/hal-04090559>**

Submitted on 5 May 2023

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Available @ <https://asianjournalofphysics.com/volume-31-no-8-aug-2022-8-2-2-2-2/>

[https://asianjournalofphysics.com/Raman-Spectroscopy\\_an-elegant-and-effective-methodfrom-single-crystals-to-the-planet-Mars/](https://asianjournalofphysics.com/Raman-Spectroscopy_an-elegant-and-effective-methodfrom-single-crystals-to-the-planet-Mars/)

## **Raman Spectroscopy: an elegant and effective method, from single crystals to the planet Mars**

Philippe Colomban

Illuminated by a monochromatic light wave (e.g. a laser), matter interacts with this wave in several ways: most of the energy is re-emitted without change in wavelength in all directions (elastic Rayleigh scattering); a much weaker part is emitted after having received or given up energy to the material by coupling between the magnetic electric vectors of the wave with the variations of the electronic distributions (in other words the chemical bonds) and more rarely magnetic. This inelastic scattering is called the Raman Effect, named after the scientist who first published his evidence 95 years ago and for this won the Nobel Prize in 1930.

Ceramist by first training, I first practiced the synthesis of single crystals and therefore crystallography, a method which required patience to mount and orient the crystals, to make the recordings of the hundreds of Bragg spots in Weissenberg or precession chamber for different orientations in order to determine the class of symmetry then to collect on a diffraction instrument with 3 or 4 circles as many spots as possible in order to solve the structure after many trials: at the time (1975) this required the perforation of hundreds of cards and the printing of many views in Fourier cross-sections using the biggest computer of the time to see if the electronic residues did not reveal partially occupied sites. Days of exposure were required for each film and months of work to solve a structure.

It was also the time of the first Ar<sup>+</sup> or Kr<sup>+</sup> ion plasma lasers giving high power (1 W) making it possible to obtain good spectra on a spectrometer with triple monochromator stage, at high resolution (0.5 cm<sup>-1</sup>), over a large spectral window, from ~3 cm<sup>-1</sup> to more than 4000 cm<sup>-1</sup>. The optical adjustment was an art: for a crystal positioned in a cryostat, the adjustment could take several hours, a good spectrum also requiring several hours, the photomultiplier detectors being not very sensitive and should not be dazzled by the laser: we spent all the day to adjust and at night to record the 'good' spectrum. A measurement campaign depending on the temperature required several days. The new possibility offered by triple-monochromators to access to very low wavenumbers made it possible to study the soft modes of certain phase transitions of ferroelectric materials, the damped modes of ionic superconductors, the tinny signature of polaritons, etc. competing with neutron spectroscopy and, to a certain extent also with NMR, which alone could then provide this information, on condition of having access to the few centers in the world with high-performance instruments and crystals of the necessary composition and (very big) size, it is say ten times or more larger than those required by the Raman analysis.

The strength of Raman analysis was in collecting the entire polarizability tensor components (90° setting) by changing the orientation of the electric vector of the laser beam and scattered light

according to the procedure and notation of S.P.S. Porto [1] to access an understanding of the structure as good as in crystallography, and even better concerning the static and dynamic disorder. The analysis in group theory of the structure supposedly well explained in the work of Poulet & Mathieu [2] made it possible by comparing with the measurements to validate or not the hypothesis. Indeed a structure based on the collection of Bragg spots symmetrizes too much the real structure and simplifies reality, because the imperfect and local orders lead to diffuse diffusions not measured by diffraction and whose study requires very specific techniques such as the method of Laüe under polychromatic X-rays, very greedy in counting time, or TEM diffraction, requiring very specialized expertise. On the contrary, despite the long counting times of the 1980s, Raman scattering was the - in my view - most elegant method of understanding solids. The study of the forms of peaks, of the dependencies in temperature, in pressure, as well as the comparison between the symmetry seen by the lattice modes and that seen by the vibrational units (group of the site of Baghavantam) confronted with the conclusions of the crystallography gives a more reliable view of 'real' matter.

My first ICORS Raman Congress in 1978 allowed me to see and hear the pioneers and discover the potential of the newly discovered SERS. France occupied an important place in the production of Raman spectrometers with Coderg (T800), with Dilor (RTI, OMARS, LABRAM), with Jobyn-Yvon (RAMANOR, MOLE)[3]. Indeed, a revolution started with the replacement of the photomultiplier first with a linear diode array (OMARS-89 Dilor Instrument; Optical Multichannel Analyzer Raman Spectrophotometer), rapidly substituted by Charge Coupled Devices (CCD) matrices.[3 ] The gain obtained with these advanced detectors offers the possibility to decrease the power of illumination and thus to study coloured matter and the technique then started to be used in Material Science analyses. Michel Delhaye (The head of my Laboratory, the LASIR located in two town Thiais a suburban close to Paris and Lille, north of France) and Paul Dhamelincourt (LASIR) were the first to build a Raman microprobe, the MOLE (Molecular Optic Laser Examiner) in 1974.[4] The MOLE microprobe started to be commercialized in 1976 by ISA Jobin-Yvon (now HORIBA Scientific). The combination of optical microscopy in a back scattering setting with different laser sources covering from violet to red excitation with a double holographic gratings spectrometer equipped with an intensified advanced photomultiplier offered the possibility to analyse a large variety of organic and inorganic materials, even coloured, including thin films; the counting time required to achieve a "convenient" spectrum for a mineral, ceramic, glass, polymer, etc. decreased to about 1 hour. Very rapidly the MOLE found application in many fields. The first application to Cultural Heritage was presented by M.L. Dele, P. Dhamelincourt and H.J. Schubnel in 1978 at a meeting for the identification of gems.[5] At the same time C. Truchet, M. Delhaye and P. Dhamelincourt started coupling the Raman spectrometer with an electron microscope. More than 40 years will be necessary to have commercially available instruments.

A second revolution starts in 2000 with the availability of air-cooled lasers and Peltier-cooled CCD detectors, the only contingency was to have an electric plug for the electricity supply for the Raman spectrometer. We conducted the first studies with mobile Raman set-up in 2002 [5,6].

All these instrumental developments find their origin in those of the pioneers studying the light-matter interaction, in first C.V Raman and K.S. Krishnan, but also Léon Brillouin, A. Smekal, G. Lansberg and L. Mandelstam, P.A. Fleury and many others. The instruments are now compact enough to equip rovers exploring Mars [8] or to detect traces of drugs or pesticides.[9]

1. S.P.S. Porto, Laser Scattering with Laser Source, [http://spexspeaker.com/Documents/68-06\\_SPEXSpeaker.pdf](http://spexspeaker.com/Documents/68-06_SPEXSpeaker.pdf)
2. H. Poulet, J.P. Mathieu, *Vibration Spectra and Symmetry of Crystal*, Gordon & Breach, 1976.
3. F. Adar, M. Delhaye, E. Da Silva (2003). Evolution of Instrumentation for Detection of the Raman Effect as Driven by Available Technologies and by Developing Applications, Pittcon Conference, Water Symposium.  
[http://www.horiba.com/fileadmin/uploads/Scientific/Documents/Raman/waters\\_symposium\\_pittcon.pdf](http://www.horiba.com/fileadmin/uploads/Scientific/Documents/Raman/waters_symposium_pittcon.pdf)
4. M. Delhaye, P. Dhamelincourt (1975). Raman microprobe and microscope with laser excitation, *J. Raman Spectrosc.* **3**, 33-43.
5. H.G.M. Edwards, P. Vandenaeele, P. Coloman, *Raman Spectroscopy in Cultural Heritage Preservation*, Springer, 2023.
6. P. Colomban, V. Milande, L. Le Bihan (2004). On-site Raman Analysis of Iznik pottery glazes and pigments, *J. Raman Spectrosc.* **35**(7), 527-535.
7. P. Colomban, V. Milande, H. Lucas (2004). On-site Raman analysis of Medici porcelain, *J. Raman Spectrosc.* **35**(1), 68-72.
8. G. Lopez-Reyes, M. Veneranda, J.A. Manrique, A. González Martín, A. Moral, C. Perez-Canora, J.A. Rodríguez Prieto, A. Sanz Arranz, J. Saiz Cano, E. Lalla, M. Konstantinidis, O. Prieto-Ballesteros, J. Medina, M.A. González, E. Charro, J.M. Lopez, F. Rull, *J Raman Spectrosc* 2022, 53( 3), 382. <https://doi.org/10.1002/jrs.6281>
9. Y. Jiang, D.-W. Sun, H. Pu, Q. Wei, Surface enhanced Raman spectroscopy (SERS): A novel reliable technique for rapid detection of common harmful chemical residues, *Trends in Food Science & Technology*, 75 (2018) 10-22, <https://doi.org/10.1016/j.tifs.2018.02.020>