Characterisation and Mapping of active pharmaceutical ingredients and excipients in a tablet BY RAMAN and infrared spectroscopy on the same microspectrometer

The objective of the measurements is to identify the unknown components and their respective dispersion within pharmaceutical tabs by Raman and FT-IR microspectrometry.

Conditions of measurements

The LabRAM IR provides a unique functionality to collect Raman and infrared spectra on the same microscope. This strongly facilitates the operation and handling of the samples. The Raman spectra have been acquired by using an excitation at 633 nm. The IR spectra have been acquired by using an ATR objective and the spectral resolution is 4 cm\(^{-1}\). The spectra have been adjusted taking account of the effect of depth penetration.

Identification of the different ingredients

Several point measurements have been carried out in different areas of the sample and two typical spectra have been obtained. The main Raman bands have been pointed out and a spectral search using an electronic library enables to identify the components characterized by these two spectra (Fig.1) as being Lactose and Starch.

In order to improve the chemical description of the sample, a Raman mapping has been acquired on the area presented below (Fig.2). The area is 35x35 µm\(^2\) large and one spectrum has been recorded every 1µm.

The Raman mapping (Figure 3) shows the spatial distribution of the compounds and allows us to identify another ingredient characterized by the spectrum plotted Fig.4. For this third spectrum, the spectral library did not identify the component. However, some useful information can be extracted from this spectrum as the main Raman band pointed out at 1005 cm\(^{-1}\) characterizes a substituted aromatic group.
Some complementary measurements have been made by FTIR:

**Complementary FT-IR measurements**

Spectra have been measured in different areas of the sample and the two spectra displayed below could be obtained.

![Spectra](image)

**Figure 5: Two characteristic FTIR spectra obtained on different points of analysis on the tablet.**

FTIR spectrum (b) can be correlated to our reference spectrum of lactose. This FTIR measurement confirms the results obtained by the Raman mapping technique showing that the lactose is present.

On spectrum (a), some additional bands can be noticed. In particular, the infrared bands around 3030 cm$^{-1}$ are due to the stretching vibration of C-H bonds in alkenes or aromatic compounds.

A substraction has been made between the two spectra to extract the IR spectrum of the unidentified third entity (Fig. 6). That allows us to make a spectral search and identify the spectrum as characteristic of diphenhydramine.

**Conclusion**

The different components of this pharmaceutical sample can be characterized easily by Raman spectroscopy and the comparison with spectra from our libraries allowed us to identify some of these molecules. A good spectral discrimination could be obtained as the characteristic spectral fingerprints have exclusive features. The Raman imaging technique is a very powerful tool as shown here, as it also enables to characterize the spatial distribution of the components with a high spatial resolution.

The complementary results obtained by FTIR spectroscopy on the sample confirm the presence of lactose and allows the identification of the third component also observed by Raman spectroscopy.

The combination of FTIR and Raman analysis on the same spectrometer represents a powerful analytical tool to readily record both IR and Raman vibrational information from a same sample position.