Raman Spectroscopy: a Test Novel Technique for Quality Control In the Acetylsalicylic Acid

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Abstract Raman spectroscopy (RS) is an inelastic scattering technique that is being recently applied in pharmaceutical applications. In this work this technique was applied to analyze the differences in concentrations of a commercial pharmaceutical form of an acetylsalicylic acid derivative present in tablets manufactured by five different laboratories (A, B, C, D and E). Meyer brand acetylsalicylic acid reagent was used to obtain spectral data as active principle. Five samples of different laboratories were also analyzed. We used a Raman microscopy device Horiba-JY HR800 with an excitation source of 830 nm and 17 mW. Raw data were analyzed by an arithmetic difference between the active substance from each different laboratory sample. Principal component analysis (PCA) was also applied to the spectral set to verify the differences in each pharmaceutical form of acetylsalicylic acid. We found a correlation of 0.7 between laboratory A and the active substance, and 0.98 between all spectra of the same laboratory A, we conclude that this drug was elaborated with better quality control. Results indicate that Raman spectroscopy can be applied in the pharmaceutical industry to determine adulteration or alteration in pharmaceutical formulations. We have demonstrate that RS together with PCA can be used as a fast and nondestructive method to determine the characteristics of the molecules present in commercially available drugs with very short acquisition times of few seconds.

Keywords Pharmaceutical, Raman spectroscopy, Principal Components, Drug

1. Introduction

RS is an analytical technique based on the effect of the inelastic scattering of light arising from changes in the polarizability of electron cloud in the molecule[1-4]. This technique has become important in recent years due to its variety of applications in areas such as mineralogy, forensics, biology and medicine[5-9]. It is currently used for the characterization of drugs, bone structures, organic pesticides, among others[10-12]. RS has an important role in the analysis stage of the design of pharmaceutical products and production process. Spectroscopy applications in this field are the supervision control of industrial manufacturing processes to determine the distribution of active pharmaceutical ingredients and excipients in different stages of a development cycle. This technique offers unparalleled discrimination of materials because it is capable of analyzing liquid and solid samples. Furthermore, this tool is particularly suitable when used along with other analytical techniques because it provides a non-destructive analysis which requires little or no sample preparation.

Most drugs are solid formulations, tablets and capsules. It is known that the state of a drug can have a significant impact on bioavailability and stability of it[13], therefore the identification and characterization of its properties salt formation, solvate formation, polymorphism, and degree of crystallinity become important in quality control. Pharmaceutical substances are subject to change in chemical composition and concentration during processing or storage of the formulation[14]. For the analysis of drugs apply a variety of methods such as colour tests, microcrystal tests, High Performance Liquid Chromatography (HPLC), UV spectroscopy, coupled gas chromatography mass spectrometry, infrared spectroscopy (IR) spectroscopy nuclear magnetic resonance (NMR), X-Ray diffraction, among others[15-21].

One of the best drug is acetylsalicylic acid, it is widely used as analgesic and antipyretic. This compound is considered unstable, have crystalline appearance and hydrolysable in water[22].

This compound has a crystalline appearance at room temperature. During the manufacture of this drug there may be several abnormalities that could not be detected during the process, changing the concentration or the substances that compose it, resulting in an adulterated product.

Most of the counterfeit drugs takes place in Asian countries. The variety of these products has been extended
due to its dissemination through the Internet, covering a bewildering variety of drugs, both branded and generic.

According to the World Health Organization, more than 50% of the purchase of drugs is done through websites established[23].

The application of the RS has become important in various fields of study, particularly in the pharmaceutical industry as it offers great advantages such as no sample preparation, allows the study of liquid and solid substances, the spectra can be obtained without the destruction of the sample[24]. This tool can be helpful in the identification of raw materials and for the quantitative determination of active substances in different formulations[25].

The use of RS greatly strengthens the process analytical technologies, which unlike laboratory studies are faster and require no manual handling of samples. The implementation of the RS in pharmaceutical activities can bring great benefits as the evaluation of product quality, making this a fast, accurate result and decreasing costs[26].

Another advantage is that the RS can be combined with sensitive statistical tools such as PCA, which is a technique for deriving a reduced set of orthogonal linear projections of a single collection of correlated variables[27-29].

2. Procedure

A Raman spectrometer LabRAM HR 800 (Horiba Jobin Yvon) with a laser excitation source of 830 nm was used, an integrated Olympus microscope with a 100X objective and a power of 17 mW. The Raman spectrometer calibration was performed using a silicon sample until adjusting the Raman intensity characteristic to 520 cm$^{-1}$.

Five samples of acetylsalicylic acid from different laboratories were purchased in different pharmaceutical shops. They were analyzed along with a sample of Meyer acetylsalicylic acid technical grade that was used as active substance. For each sample, five spectra were obtained by pointing the laser at different regions of the drug, collecting a total of 35 Raman spectra. The acquisition was carried out in the range from 200 to 2000 cm$^{-1}$ with a 0.6 cm$^{-1}$ resolution with an exposure time of 10 sec and an integration time of 5 s.

A baseline correction was carried out to the entire spectral data by the least squares method to eliminate background noise, following the Peng method[30], as shown in Figure 1.

All Raman spectra were normalized to the greatest intensity peak.

![Figure 1. Baseline correction and comparing average Raman spectra](image)

All Raman spectra processed were analyzed in the range of 200 to 2000 cm$^{-1}$. All the processed Raman spectra were evaluated by principal components analysis and also a correlation matrix was obtained to verify the similarities between different laboratories and the active substance.

On the other hand, a first procedure performed was that for every five spectra obtained for each sample they were analyzed, averaged and compared between each laboratory including the active substance, as shown in Figure 1. Data processing was performed using the commercial software MatLab 2011, Minitab 16 and Origin version 8.5.

3. Results and Discussions

Figure 2 shows comparatively each spectral average of each laboratory sample evaluated with respect to the active substance.

It is spectrally observed that the drugs that correspond to the laboratories A and C are very similar to the active substance. Moreover, the drug of laboratory E shows very significant differences with respect to the active substance.
The data set with baseline correction was assessed by principal component analysis (PCA) allowing plot a new coordinate space to observe spectral differences in all tested drugs. In Figure 3 (PC1 vs. PC2) and Figure 4 (PC1 vs. PC3) shows that laboratories A and C are the best features conserved with respect to the active substance. Moreover, drugs which correspond to laboratories B and D differ significantly from the active substance but there is homogeneity among them. Laboratory E differs significantly with respect to the active substance, but also resulted in inhomogeneous regions analyzed in this drug, this means that, in the drug compaction the concentrations were not homogeneous.

The Raman bands associated with each intensity that showed differences are shown in Table 1. When comparing the Raman spectra of samples taken from different laboratories, similar features were found between each drug sample. This denotes significant variations and indicates that
these differences are due to changes in the chemical composition of the sample.

Table 1. Raman peak positions and vibrational mode assignment (Abbreviations: ν, stretching vibration; δ, bending vibration)

| Raman shift (cm⁻¹) | Assignments                  |
|-------------------|------------------------------|
| 281 to 290        | δ(CC), Aliphatic chains      |
| 420 to 430        | ν (S=S)                      |
| 550               | ν (C-Cl)                     |
| 747               | ν (C-Br)                     |
| 785               | ν (C(aliphatic)-S)           |
| 1040 to 1050      | ν ((C)SO(-C))                |
| 1080 to 1090      | ν (C(aromatic)-S)            |
| 1290-1295         | ν (CC), Aliphatic chains, alicyclic |
| 1605              | ν (C=N)                      |

It was identified a difference in the range of 281-290 cm⁻¹, where there are C-C bonds of aliphatic chains. In the range of 420-430 cm⁻¹ the differences associated with S-S bonds were also found. In this case the molecule of active substance to be studied (acetylsalicylic acid) does not have S-S bonds, we think it is the excipient or vehicle that contains the sample. So also in the Raman shift 550 cm⁻¹, there is a shift and this is a C-Cl bond, which indicates a different composition of the acetylsalicylic acid.

In the band 747 and 785 cm⁻¹ there are differences and which do not correspond to the original molecule (acetylsalicylic acid), presenting links C-Br, and C-S in an aliphatic bond, respectively. In the region 1040-1050 cm⁻¹ are rings which correspond to the drug molecule.

Also we can find rings joined to oxygen, methyl carboxylic acids and even to the same carbon, as found in the range of 1080-1090 cm⁻¹, from 1290 to 1295 cm⁻¹, and 1605 cm⁻¹ respectively. To corroborate the comparisons between different laboratories and homogeneity in the group of the five Raman spectra in each of them, an analysis of best subsets in linear regression was applied, in which the response variable was one of the spectra of the active substance and it was selected the best five subsets of a variable. The results obtained are shown in Table 2. From Table 2, we corroborate that laboratories A and E are ones showing an important correlation with the active substance, the difference is that for each spectrum of the active substance the three spectrums of lab A appear consistently on the same amount of correlation, while the spectrums of lab E appear on the highest and lower five more significant correlations, which show again the lack of homogeneity quality on lab E.

Finally the spectral differences were observed by plotting the first three principal components, as shown in Figure 5. The dashed circle encloses the active substance and the two drugs that showed better similarities with respect to the active substance.

Table 2. Comparisons between laboratories A and E with respect to the active substance and the homogeneity in each group (Abbreviations: Lab., Laboratory; S, Spectrum)

| Active Substance | S1     | S2     | S3     | S4     | S5     |
|------------------|--------|--------|--------|--------|--------|
| R²               | Variable | R²     | Variable | R²     | Variable | R²     | Variable | R²     | Variable |
| 70.9             | Lab.E S5 | 63.8   | Lab.E S5 | 74.6   | Lab.E S5 | 74.0   | Lab.E S5 | 85.1   | Lab.E S5 |
| 54.8             | Lab.A S2 | 53.2   | Lab.A S2 | 65.1   | Lab.A S2 | 63.8   | Lab.A S2 | 72.7   | Lab.A S2 |
| 54.8             | Lab.A S4 | 53.2   | Lab.A S4 | 64.3   | Lab.A S4 | 63.1   | Lab.A S4 | 72.7   | Lab.A S4 |
| 53.5             | Lab.A S5 | 51.9   | Lab.A S5 | 62.9   | Lab.A S5 | 61.7   | Lab.A S5 | 71.6   | Lab.A S5 |
| 52.4             | Lab.E S4 | 51.0   | Lab.E S4 | 62.0   | Lab.A S3 | 60.7   | Lab.A S3 | 70.3   | Lab.A S3 |

Figure 5. Spectral distribution by plotting the principal tree first principal components, PC1 versus PC2 versus PC3. (Abbreviations A.S.: Active Substance, Lab: Laboratory)
We also observed that the correlation between Raman spectra of the laboratory A has a significant correlation about 0.7 with respect to the spectra of the active substance and show a correlation of at least 0.98 between themselves, as shown in Table 3, which means a good quality control in the development of this drug. For the laboratory E, the spectrum five shows a correlation between 0.82 and 0.93 with respect to the spectra of the active substance.

An statistical analysis was carry out, in which we found that the correlations between laboratory E are very variable, ranging from 0.51 to 0.9, which means that there is little homogeneity among them, as shown in Table 4. In this table is described the Raman spectra relations between them.

These market differences and these spectra indicate that the compaction process was not good enough.

4. Conclusions

We have demonstrate that Raman spectroscopy together with principal component analysis can be used as a fast and nondestructive method to determine the concentration and characteristics of the molecules present in commercially available drugs with very short acquisition times of few seconds. Here we demonstrated too that the power of this technique by discriminating between well manufactured (laboratory A sample) and poorly manufactured (laboratory E) drugs. The validity of this result is backed up by the use of a technical grade reagent employed as a reference. This non-destructive analysis could also be used as a quality control tool in the manufacturing processes of commercial drugs in short time analysis. It could also be possible to use this technique to help in the control and eventual reduction of the trade of counterfeit drugs.

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