Investigation of Prevention, Protection and Treatment of Lopinavir Effectiveness on Coronavirus Disease–2019 (COVID–19) Infection Using Fourier Transform Raman (FT–Raman) Biospectroscopy

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ABSTRACT

Lopinavir is an antiretroviral of the protease inhibitor class. It is used against HIV infections as a fixed–dose combination with another protease inhibitor, ritonavir (lopinavir/ritonavir). In the current research, the stimulated FT–Raman biospectroscopy of liquid sample of Lopinavir was investigated. The stimulated FT–Raman diffractions emitted through focusing the second harmonic laser beam Nd:YAG into the sample were recorded by Echelle spectrometer and ICCD detector. Increasing the energy of laser beam from 2.6 (mJ) to 16 (mJ) was led to increase in stimulated FT–Raman signal but after breakdown threshold of liquid sample, more increasing of energy was led to decrease in stimulated FT–Raman signals and for energies higher than 20 (mJ), they were disappeared.

Keywords: FT–Raman Biospectroscopy, Stimulated FT–Raman Biospectroscopy, Lopinavir, Breakdown, Coronavirus Disease–2019, COVID–19, Infection, Protective and Therapeutic Effect, Potent Drug

1. INTRODUCTION:

FT–Raman biospectroscopy is a vibration biospectroscopy based on the influence of FT–Raman [1–47]. The influence of FT–Raman is elastically diffracting the electromagnetic ray due to rotational and vibrational transitions in molecules and its characteristic is changing the energy of diffracted beam photons compared to incident beam [48–95]. The difference between wavelength of incident beam light and diffracted light is related to molecular vibrations and is considered as exclusive “chemical finger print” of sample and can be used in identification of molecular compounds on a surface, into a liquid or into the air [96–142].

The stimulated FT–Raman diffraction is a non–linear effect [143–189]. If the pumping intensity exceeds the threshold of this effect, it observes [190–237]. The pumping threshold limit for stimulated FT–Raman depends on FT–Raman active material [238–285]. Regarding the spectral characteristics, stimulated FT–Raman can be distinguished from normal FT–Raman [286–333]. While the intensity of FT–Raman bands are several times smaller than pumping laser intensity in normal FT–Raman, the intensity of FT–Raman bands in stimulated FT–Raman can be similar to laser intensity and for most materials, only strongest FT–Raman bands of material are intensified and are dominant in the recorded spectrum of material [334–379].

In the current research, the stimulated FT–Raman spectrum are obtained through pumping the second harmonic beam laser Nd:YAG and it is performed by a spectrometer and detector. The resulted spectra and their characteristics are investigated here.

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3.5 (cm). The resulted emissions from this interaction filters by an optical system consisting some lens and optical fiber conducts to Eschelle spectrometer. The necessary time range for collecting spectra and its start time in ICCD detector controls by delay device. Optical emissions of sample collects and intensifies from the striking moment of laser to sample until 5 (ms) after that moment. Test was repeated five times for each energy level for laser energy from 2.4 (mJ) to 29 (mJ).

By increasing the energy of laser beam, the intensity of main bands of 1380 (cm−1) and 1461 (cm−1) also are increased and for energy levels higher than 8 (mJ), anti–Stokes FT–Raman band corresponding to 1380 (cm−1) intensifies in the spectrum and can be observed at left hand side of laser line in FT–Raman shift of ~1380 (cm−1). Recording the anti–Stokes band necessitates the occupation of corresponding vibration level through diffraction of Stokes FT–Raman (Table 1).

By more increasing the energy level higher than 16 (mJ), all four graphs of Figure (3) shows reduction in intensity. The reason for this reduction is creation of spark in the Lopinavir liquid due to increase in energy of laser more than the breakdown threshold of liquid. As a result of this spark, which creates in the center of liquid, laser beam absorbs by liquid and some part of it diffracts and only this part plays a role in creation of stimulated FT–Raman. By increasing the energy, beam has higher contribution in making the spark and the diffracted emission which reaches to detector decreases.

### Table (1): FT–Raman modes for Lopinavir.

| FT–Raman Shift (cm⁻¹) | FT–Raman Mode             |
|-----------------------|---------------------------|
| 1 940 (cm⁻¹)          | C–H Stretch               |
| 2 1127 (cm⁻¹)         | CH₂ Rocking               |
| 4 1339 (cm⁻¹)         | CH₂ Wagging               |
| 5 1380 (cm⁻¹)         | CH₂ Symmetric Stretch     |
| 7 1461 (cm⁻¹)         | C–H Asymmetric Stretch    |

Figure (1): Schematic of stimulated FT–Raman biospectroscopy test arrangement.

3. RESULTS AND DISCUSSION

Figure (2) shows the normal and stimulated FT–Raman spectra. Normal FT–Raman spectrum can be obtained when laser beam is not focused on the sample. When laser beam focuses on sample using a lens, non–linear effects stimulate and stronger bands of FT–Raman spectrum intensify up to some levels of laser intensity.

Figure (2): (a) Normal and (b) stimulated FT–Raman spectra for Lopinavir.
Figure (3): Peak intensity (a) band 940 (cm$^{-1}$), (b) band 1127 (cm$^{-1}$), (c) band 1339 (cm$^{-1}$), (d) band 1380 (cm$^{-1}$), (e) band 1461 (cm$^{-1}$) and (f) band –1380 (cm$^{-1}$) based on increase in energy level of beam focused on the liquid.

4. CONCLUSIONS AND SUMMARY

The stimulated FT–Raman biospectroscopy test was performed for liquid sample of Lopinavir. The main band at 1380 (cm$^{-1}$) shows an intensity level comparable to pumping laser intensity. The intensity of stimulated FT–Raman spectrum at 16 (mJ) energy level is the highest intensity in this test and more increasing the energy level reduces the intensity of spectrum. The reason for this reduction is creation of spark in the Lopinavir liquid due to increase in energy of laser more than the breakdown threshold of Lopinavir.

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