Spectroscopic Characterization of Disulfiram and Nicotinic Acid after Biofield Treatment

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Abstract

Disulfiram is being used clinically as an aid in chronic alcoholism, while nicotinic acid is one of a B-complex vitamin that has cholesterol lowering activity. The aim of present study was to investigate the impact of biofield treatment on spectral properties of disulfiram and nicotinic acid. The study was performed in two groups i.e., control and treatment of each drug. The treatment groups were received Mr. Trivedi’s biofield treatment. Subsequently, spectral properties of control and treated groups of both drugs were studied using Fourier transform infrared (FT-IR) and Ultraviolet-Visible (UV-Vis) spectroscopic techniques. FT-IR spectrum of biofield treated disulfiram showed the shifting in wavenumber of C-H stretching from 1496 to 1506 cm⁻¹ and C-N stretching from 1062 to 1056 cm⁻¹. The intensity of S-S dihedral bending peaks (665 and 553 cm⁻¹) was also increased in biofield treated disulfiram sample, as compared to control. FT-IR spectra of biofield treated nicotinic acid showed the shifting in wavenumber of C-H stretching from 3071 to 3081 cm⁻¹ and 2808 to 2818 cm⁻¹. Likewise, C=C stretching peak was shifted to higher frequency region from 1696 cm⁻¹ to 1703 cm⁻¹ and C-O (COO⁻) stretching peak was shifted to lower frequency region from 1186 to 1180 cm⁻¹ in treated nicotinic acid.

UV spectrum of control and biofield treated disulfiram showed similar pattern of UV spectra. Whereas, the UV spectrum of biofield treated nicotinic acid exhibited the shifting of absorption maxima (λmax) with respect of control i.e., from 268.4 to 262.0 nm, 262.5 to 256.4, 257.5 to 245.6, and 212.0 to 222.4 nm.

Over all, the FT-IR and UV spectroscopy results suggest an impact of biofield treatment on the force constant, bond strength, and dipole moments of treated drugs such as disulfiram and nicotinic acid that could lead to change in their chemical stability as compared to control.

Keywords: Disulfiram; Nicotinic acid; Biofield treatment; Fourier transform infrared spectroscopy; Ultraviolet spectroscopy

Introduction

Disulfiram [bis(diethylthiocarbamoyl)disulphide] is an antabuse drug, being used clinically as an aid to the treatment of chronic alcoholism. It is the first drug approved by US Food and Drug Administration to treat the alcohol addiction [1]. Alcohol (ethanol) transforms into acetaldehyde by alcohol dehydrogenase enzyme, which further oxidized to acetic acid by acetaldehyde dehydrogenase (ADH) enzyme [2]. Disulfiram inhibits the ADH enzyme. As a result, the blood concentration of acetaldehyde increases and causes an unpleasant effect, thus increase the patient’s motivation to remain abstinent [3]. In addition to this, disulfiram is reported for protozoacidal effect in vitro study [4,5]. Recently, disulfiram has shown the reactivity to latent HIV-1 expression in a primary cell model of virus latency and presently it is assessed in a clinical trial for its potential to diminish the latent HIV-1 reservoir in patients combination with antiretroviral therapy [6].

Nicotinic acid or niacin is one of the B-complex vitamins (Vitamin B3) that has cholesterol lowering activity. Recent studies showed that therapeutic doses of nicotinic acid induce a profound alteration in plasma concentration of several lipids and lipoproteins, resulting in a greater ability to increase high-density lipoprotein (HDL) cholesterol [7]. Nicotinic acid favorably affects apolipoprotein (apo), very-low-density lipoprotein (VLDL), low-density lipoprotein (LDL) and HDL [7,8].

The exact mechanism of nicotinic acid activity is unknown. However, new findings indicate that nicotinic acid inhibits directly and non-competitively to the triglycerides synthesis enzyme i.e., hepatocyte diacylglycerol acyltransferase-2, which causes acceleration of intracellular hepatic apo B degradation and thus decrease secretion of VLDL and LDL [9]. Several evidence suggest that nicotinic acid administered either alone or in combination with other cholesterol-lowering medicines can reduce the risk of cardiovascular and atherosclerosis diseases. The clinical uses of nicotinic acid are somewhat limited due to some harmless but unpleasant side effects like cutaneous flushing phenomenon, nausea, vomiting and headache [10]. The chemical and physical stability of pharmaceutical drugs or products are most desired attributes of quality that potentially affect the efficacy, safety and shelf life of drugs [11]. Hence, it is essential to find out an alternate approach, which could enhance the stability of drugs by altering the structural and bonding properties of these compounds.

Contemporarily, biofield treatment is reported to alter the spectral properties of various pharmaceutical drugs like paracetamol, piroxicam, metronidazole, and tinidazole; likewise physical, and structural properties of various metals i.e., tin, lead etc. [12-14]. The conversion of mass into energy is well known in literature for hundreds of years that mass into energy is well known in literature for hundreds of years that was further explained by Hasenohrl and Einstein [15,16]. According to Maxwell JC, every dynamic process in the human body had an electrical significance, which generates magnetic field in the human body [17].

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This electromagnetic field of the human body is known as biofield and energy associated with this field is known as biofield energy [18,19]. Mr. Trivedi has the ability to harness the energy from environment or universe and can transmit into any living or nonliving object around this Globe. The object(s) always receive the energy and responding into useful way, this process is known as biofield treatment.

Mr. Mahendra Kumar Trivedi’s biofield treatment (The Trivedi Effect) has considerably changed the physicochemical, thermal and structural properties of metals and ceramics [14,20,21]. Growth and anatomical characteristics of some plants were also increased after biofield treatment [22,23]. Further, biofield treatment has shown the significant effect in the field of agriculture science [24,25] and microbiology [26,27].

Considering the impact of biofield treatment on physical and structural property of metals and ceramics, the present study was aimed to evaluate the impact of biofield treatment on spectral properties of disulfiram and nicotinic acid. The effects were analyzed using Fourier transform infrared (FT-IR) and Ultraviolet-Visible (UV-Vis) spectroscopic techniques.

Materials and Methods

Study design

The disulfiram and nicotinic acid (Figure 1) samples were procured from Sigma-Aldrich, MA, USA; and each drug was divided into two parts i.e., control and treatment. The control samples were remained as untreated, and treatment samples were handed over in sealed pack to Mr. Trivedi for biofield treatment under laboratory condition. Mr. Trivedi provided this treatment through his energy transmission process to the treated groups without touching the sample [12,13]. The control and treated samples of disulfiram and nicotinic acid were evaluated using FT-IR and UV-Vis spectroscopy.

FT-IR spectroscopic characterization

FT-IR spectra were recorded on Shimadzu’s Fourier transform infrared spectrometer (Japan) with frequency range of 4000-500 cm\(^{-1}\). The FT-IR spectroscopic analysis of both control and treated samples of disulfiram and nicotinic acid were carried out to evaluate the impact of biofield treatment at atomic level like force constant and bond strength [28].

UV-Vis spectroscopic analysis

UV spectra of disulfiram and nicotinic acid were recorded on Shimadzu UV-2400 PC series spectrophotometer with 1 cm quartz cell and a slit width of 2.0 nm. The analysis was carried out using wavelength in the range of 200-400 nm. The analysis was performed to determine the effect of biofield treatment on structural properties of treated drugs [28].

Results and Discussion

FT-IR spectroscopic analysis

Vibrational spectral assignment was performed on the recorded FT-IR spectra (Figure 2) based on theoretically predicted wavenumber and presented in Table 1. The FT-IR spectrum of control disulfiram sample (Figure 2a) showed the characteristic vibrational peak at 2975 cm\(^{-1}\) that was assigned to C-H (CH\(_3\)) stretching. Another characteristic peak observed at 1496 cm\(^{-1}\) was attributed to C-H symmetrical deformation vibrations. The absorption peaks appeared at 1351-1457 cm\(^{-1}\) was assigned to CH\(_2\)-CH\(_3\) deformations. The vibrational peaks at 1273 cm\(^{-1}\) and 1151-1195 cm\(^{-1}\) were assigned to C=S stretching and C-C skeletal vibration, respectively. Further, IR peaks observed at 967-1062 cm\(^{-1}\) and 817-914 cm\(^{-1}\) were attributed to C-N stretching and C-S stretching, respectively. The vibrational peaks appeared at 554-666 cm\(^{-1}\) was assigned to S-S dihedral bending. The FT-IR data of control disulfiram was well supported by the literature data [29].

The FT-IR spectrum of biofield treated disulfiram (Figure 2b) showed the vibrational peak at 2975 cm\(^{-1}\), which was assigned to CH\(_3\) stretching. Vibrational peak appeared at 1506 cm\(^{-1}\) was assigned to C-H symmetrical deformation vibrations. Likely, the IR peaks at 1350-1457 cm\(^{-1}\) were attributed to CH\(_2\)-CH\(_3\) deformations. The vibrational peaks appeared at 1273 cm\(^{-1}\) and 1151-1195 cm\(^{-1}\) were assigned to C=S stretching and C-C skeletal vibration, respectively. The IR peaks observed at 967-1056 cm\(^{-1}\) and 817-914 cm\(^{-1}\) were attributed to C-N stretching and C-S stretching, respectively. The vibrational peaks at 553-665 cm\(^{-1}\) were assigned to S-S dihedral bending.

Altogether, the FT-IR data of biofield treated disulfiram (Figure 2b) showed the shifting in frequency of some bonds with respect to control spectra like C-H symmetrical deformation vibrations frequency was shifted from 1496 (control) to 1506 (treated) cm\(^{-1}\). The frequency (\(v\)) of vibrational peak depends on two factors i.e., force constant (k) and

![Figure 1: Chemical structure of (a) Disulfiram and (b) Nicotinic acid.](Image 1)

![Figure 2: FT-IR spectra of Disulfiram (a) control and (b) treated.](Image 2)
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**Wave number (cm⁻¹) | Frequency Assignment**

| Control | Treated |
|---------|---------|
| 2975    | 2975    | CH₃ stretching |
| 1506    | 1506    | C-H symmetrical deformation vibrations |
| 1351-1457 | 1350-1457 | CH₃ and CH₂ deformation |
| 1273    | 1273    | C=S stretching |
| 1151-1195 | 1151-1195 | C-C skeletal vibrations |
| 967-1062 | 967-1056 | C-N stretching |
| 818-914  | 817-914  | C-S stretching |
| 554-666  | 553-665  | S-S stretching |

Table 1: FT-IR vibrational peaks observed in Disulfiram.

The vibrational spectral assignment of nicotinic acid was performed on the recorded FT-IR spectra (Figure 3) based on theoretically predicted wavenumber and presented in Table 2. The vibrational peaks appeared at 3071-2808 cm⁻¹ were assigned to C-H stretchings. The IR peaks observed at 1696-1710 cm⁻¹ and 1596 cm⁻¹ were assigned to C=O (COO⁻) asymmetric stretching and C=C stretching, respectively. Absorption peaks appeared at 1417, 1323, and 1301 cm⁻¹ were attributed to C=O symmetric stretching, C=O asymmetric stretching, and C-N stretching, respectively. The C-O (COO⁻) stretching peak was assigned to IR bend observed at 1186 cm⁻¹. Further, C-H in plane and out plane bending vibrations was assigned to peaks observed in the range of 1033-1114 cm⁻¹ and 642-812 cm⁻¹, respectively. The FT-IR data of control nicotinic acid was well supported by the literature data [32,33].

The FT-IR spectra of biofield treated nicotinic acid (Figure 3) showed the absorption bands at 2818-3081 cm⁻¹ that were assigned to C-H stretching. Vibrational peaks appeared at 1703-1714 cm⁻¹ and 1594 cm⁻¹ were assigned to C=O (COO⁻) asymmetric stretching and C=C stretching, respectively. Likewise, the IR peaks observed at 1417, 1324, and 1301 cm⁻¹ were assigned to C=N stretching, C=O (COO⁻) symmetric stretching, and C-N stretching, respectively. The IR absorption peak appeared at 1180 cm⁻¹ was attributed to C-O (COO⁻) stretching. Further, the C-H in plane and out plane bending vibrations was assigned to IR peaks observed at 1037-1116 cm⁻¹ and 642-812 cm⁻¹, respectively.

Overall, the FT-IR data of biofield treated nicotinic acid (Figure 3) showed the shifting in wavenumber of some bonds with respect to control sample. For instance, the C-H stretching towards higher frequency region i.e., from 3071 to 3081 cm⁻¹ and 2808 to 2818 cm⁻¹. This could be due to increase in force constant of C-H bond. Likewise, a slight upstream shifting in C=O stretching peak from 1710 to 1714 cm⁻¹ and 1696 to 1703 cm⁻¹ in treated nicotinic acid also suggests an increase in force constant of C=O bond in treated sample as compared to control. Contrarily, a slight downstream shifting in wavenumber of treated nicotinic acid from 1186 to 1180 cm⁻¹ C-O (COO⁻) stretching; and from 1033 to 1037 cm⁻¹ (=C-H in plane bending) suggests the decrease in force constant of C=O bond and decrease in rigidity of -=C-H bond in treated sample as compared to control.

**UV-Vis spectroscopy**

![UV-Vis spectra](image)

Figure 3: FT-IR spectra of Nicotinic acid (a) control and (b) treated.

**Table 2: FT-IR vibrational peaks observed in Nicotinic acid.**
UV spectra of control and treated disulfiram showed a similar pattern of UV spectra with absorption maxima ($\lambda_{\text{max}}$) of 219.8, 250.2, and 281.6 nm in control and 220.8, 249.4, and 281.2 nm in treated sample. This indicates no significant change in the UV spectral property of treated disulfiram with respect to control sample. The UV spectra of control and treated nicotinic acid are showed in Figure 4. The UV spectrum of treated nicotinic acid (Figure 4) exhibited the shifting of absorption maxima ($\lambda_{\text{max}}$) from 268.4 to 262.0 nm, 262.5 to 256.4 nm, 257.5 to 245.6 nm, and 212.0 to 222.4 nm. The existing literature on principle of UV spectroscopy suggests that a compound can absorbs UV light due to presence of either or both conjugated $\pi$ and $\pi^*$-bonding systems ($\pi-\pi^*$ transition) and nonbonding electron system ($n-\pi^*$ transition) in the compound. The UV absorption phenomenon occurred when electrons travelled from low energy orbital (i.e., $\sigma$, $\pi$, and $n$) to high energy orbital (i.e., $\sigma^*$ and $\pi^*$). There is certain energy gap between $\sigma-\sigma^*$, $\sigma-\pi^*$, $\pi-\pi^*$ and $n-\pi^*$ orbitals. When this energy gap altered, the wavelength ($\lambda_{\text{max}}$) was also altered respectively [28]. Based on this, it is speculated that, due to influence of biofield treatment, the energy gap between $\pi-\pi^*$ and $n-\pi^*$ transition in nicotinic acid might be altered, which causes shifting of wavelength ($\lambda_{\text{max}}$) in treated nicotinic acid as compared to control. To the best of our knowledge, this is the first report showing an impact of biofield treatment on structural properties like force constant, bond strength, dipole moment of disulfiram and nicotinic acid.

**Conclusion**

The FT-IR data of biofield treated disulfiram showed an alteration in the wavenumber of C-H, C=O, and C-O stretching, and C-H bending were altered in biofield treated nicotinic acid, with respect of control. Also, the peak intensity at 553-665 cm$^{-1}$ (S-S dihedral bending) was increased in biofield treated disulfiram, as compared to control. This alteration in wavenumber referred to alteration in the force constant and bond strength of respective group. The UV spectral data of biofield treated nicotinic acid also support the possible change in the structural property with respect of control.

In conclusion, the results suggest a significant impact of biofield treatment on structural property like force constant, bond strength, dipole moment, and energy gap between bonding and nonbonding orbital of treated drug with respect to control.

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