**Evaluation of the Effect of Consciousness Energy Healing Treatment on Physicochemical and Thermal Properties of Flutamide**

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**Abstract**

Flutamide is used primarily to treat the prostate cancer, which blocks the action of exogenous testosterone by binding to the androgen receptor. The aim of this study was to evaluate the impact of the Trivedi Effect®-Consciousness Energy Healing Treatment on the physicochemical and thermal properties of flutamide using the modern analytical technique. The flutamide sample was divided into control and treated part. The control part did not receive any Energy Treatment, while the treated part received the Biofield Energy Treatment remotely by a renowned Biofield Energy Healer, Dahryn Trivedi. The PXRD peak intensities and crystallite sizes were significantly altered ranging from -29.24% to 1.79% and 0.44% to 31.21% respectively; however, the average crystallite size was significantly increased by 19.13% in the treated sample compared with the control sample. The particle size values were altered by 0.12%(d₁₀) 2.9%(d₅₀), 7.52%(d₉₀), and 5.81%(D(4,3)), thus, the specific surface area was decreased by 0.55% in the treated sample compared to the control sample. The melting point and latent heat of fusion of the treated sample were increased by 0.63% and 5.31%, respectively compared to the control sample. The total weight loss was decreased by 1.19%; however, the residue amount was increased by 1.18% in the treated flutamide compared with the control sample. The Biofield Energy Treatment might have generated a new polymorphic form of flutamide which would be more efficacious in the pharmaceutical formulations for the treatment of prostate cancer.

**Keywords:** Flutamide; Consciousness Energy Healing Treatment; The Trivedi Effect®; Complementary and Alternative Medicine; PXRD; Particle size; DSC; TGA/DTG

Gravimetric Analysis.

Introduction

Flutamide is an acetyl-, nonsteroidal antiandrogen (NSAA) having the chemical name, 2-methyl-N-[4-nitro-3 (trifluoromethyl) phenyl] propanamide [1-3]. Capsules of eulexin contain flutamide, orally active anti-androgen, which is used primarily to treat men with prostate cancer [4]. This can be used as independent drug or is used with other medications as well as with radiation treatments [5]. Testosterone, a natural hormone, helps the prostate cancer to grow and spread. The role of flutamide is to block the effects of testosterone, thus slowing down the growth and spread of prostate cancer [6]. Irregular consumption of this medicine or stopping medications before it completely gets cured, could allow the cancer to spread more rapidly. Overdose may cause hypoactivity, piloerection, slow respiration, ataxia, and/or lacrimation, anorexia, tranquilization, emesis, and methemoglobinemia [7]. However, the flutamide overdose ordinarily associated with symptoms or considered to be life-threatening has not been established.

Study of the physicochemical properties of a pharmaceutical product regarding its dissolution and absorption is crucial for the formulation. It was observed that the Biofield Energy Healing Treatment (the Trivedi Effect®) has the considerable impact on various properties such as particle size, surface area, and other chemical and thermal behaviour of pharmaceutical/nutraceutical compounds [8-11]. The Trivedi Effect® is a natural and scientifically proven phenomenon in which a person can harness this inherently intelligent energy and transmit it anywhere on the planet through the possible mediation of neutrinos [8]. Every living organism possesses this kind of unique energy surrounding the body known as the “Biofield Energy”, which is infinite, para-dimensional electromagnetic field. Biofield (Putative Energy Fields) based Energy Healing Therapies have been reported with significant outcomes against various disease conditions [12]. The National Institutes of Health/National Center for Complementary and Alternative Medicine (NIH/NCCAM) recommend and included the Energy therapy under the Complementary and Alternative Medicine (CAM) category along with other therapies, which include homeopathy, Ayurvedic medicine, naturopathy, Tai Chi, Qi Gong, acupuncture, acupressure, Reiki, healing touch, hypnotherapy, Rolfing, etc. The CAM has been accepted by the most of the U.S. population with several advantages [13, 14]. The Trivedi Effect®-Consciousness Energy Healing Treatment has been widely reported with astounding capability to alter the characteristic properties of the several non-living materials and living object(s), i.e. metals and ceramic, organic compounds, nutraceutical/pharmaceuticals, and crops [9-11,15-23]. The Consciousness Energy Healing Treatment has also enhanced the bioavailability of pharmaceutical/nutraceutical compounds [24-26]. This study was designed to determine the impact of the Trivedi Effect®-Consciousness Energy Healing Treatment on the physicochemical and thermal properties of flutamide using powder X-ray diffraction (PXRD), particle size analysis (PSA), differential scanning calorimetry (DSC), and thermogravimetric analysis (TGA)/Differential thermo gravimetric analysis (DTG).

Materials and Methods

Chemicals and Reagents

The flutamide was purchased from Tokyo Chemical Industry Co. Ltd. All other chemicals used during the experiments were of analytical grade available in India.

Consciousness Energy Healing Treatment Strategies

The flutamide was the test sample divided into two parts. One part of flutamide was considered as control sample (no Biofield Energy Treatment was provided). Consequently, the second part of flutamide was treated with the Trivedi Effect®-Energy of Consciousness Healing Treatment remotely under standard laboratory conditions for 3 minutes and was called the Biofield Energy Treated flutamide sample. This Biofield Energy Treatment was provided through the healer’s unique energy transmission process by the renowned Biofield Energy Healer, Dahryn Trivedi, USA, to the second part of the test sample. Further, the control sample was treated with a “sham” healer. The “sham” healer did not have any knowledge about the Biofield Energy Treatment. After the treatment, the Biofield Energy Treated and untreated sample were both kept in sealed conditions and characterized using PXRD, PSA, DSC, and TGA techniques.

Characterization

Powder X-ray Diffraction (PXRD) Analysis: The PXRD analysis of flutamide was performed with the help of Rigaku MiniFlex-II Desktop X-ray diffractometer (Japan) [27,28]. The average size of individual crystallites was calculated from XRD data using the Scherrer’s formula (1)
G = k\lambda/\beta\cos\theta \quad (1)

Where k is the equipment constant (0.94), G is the crystallite size in nm, \(\lambda\) is the radiation wavelength (0.154056 nm for Kα 1 emission), \(\beta\) is the full-width at half maximum (FWHM), and \(\theta\) is the Bragg angle [29]. The % change in crystallite size (G) of flutamide was calculated using the following equation 2:

\[
\% \text{ change in crystallite size } = \left( \frac{G_{\text{Treated}} - G_{\text{Control}}}{G_{\text{Control}}} \right) \times 100 \quad (2)
\]

Where \(G_{\text{Control}}\) and \(G_{\text{Treated}}\) are the crystallite size of the control and the Biofield Energy Treated samples, respectively.

**Particle Size Analysis (PSA):** The particle size analysis of flutamide was conducted on Malvern Mastersizer 2000, from the UK with a detection range between 0.01 \(\mu\)m to 3000 \(\mu\)m using wet method [30,31]. The sample unit (Hydro MV) was filled with a dispersant medium (sunflower oil) and operated the stirrer at 2500 rpm. The PSA analysis of flutamide was performed to obtain the average particle size distribution. Where, \(d\) (0.1) \(\mu\)m, \(d(0.5)\) \(\mu\)m, \(d(0.9)\) \(\mu\)m represent particle diameter corresponding to 10%, 50%, and 90% of the cumulative distribution. \(D(4,3)\) represents the average mass-volume diameter, and SSA is the specific surface area (m\(^2\)/g). The calculations were done by using software Mastersizer Ver. 5.54.

The percent change in particle size (d) for at below 10% level \((d_{10})\), 50% level \((d_{50})\), 90% level \((d_{90})\), and \(D(4,3)\) was calculated using following equation 3:

\[
\% \text{ change in particle size } = \left( \frac{d_{\text{Treated}} - d_{\text{Control}}}{d_{\text{Control}}} \right) \times 100 \quad (3)
\]

Where, \(d_{\text{Control}}\) and \(d_{\text{Treated}}\) are the particle size (\(\mu\)m) at below 10% level \((d_{10})\), 50% level \((d_{50})\), and 90% level \((d_{90})\) of the control and the Biofield Energy Treated samples, respectively. Percent change in surface area (S) was calculated using following equation 4:

\[
\% \text{ change in surface area } = \left( \frac{S_{\text{Treated}} - S_{\text{Control}}}{S_{\text{Control}}} \right) \times 100 \quad (4)
\]

Where \(S_{\text{Control}}\) and \(S_{\text{Treated}}\) are the surface area of the control and the Biofield Energy Treated flutamide, respectively.

**Differential Scanning Calorimetry (DSC):** The DSC analysis of flutamide was performed with the help of DSC Q200, TA instruments. A sample of \(~1-5\) mg was loaded to the aluminium sample pan at a heating rate of 10\(^\circ\)C/min from 30\(^\circ\)C to 350\(^\circ\)C [30, 31]. The % change in melting point (T) was calculated using the following equation 5:

\[
\% \text{ change in melting point } = \left( \frac{T_{\text{Treated}} - T_{\text{Control}}}{T_{\text{Control}}} \right) \times 100 \quad (5)
\]

Where \(T_{\text{Control}}\) and \(T_{\text{Treated}}\) is the melting point of the control and treated samples, respectively.

The Percent change in the latent heat of fusion (\(\Delta H\)) was calculated using the following equation 6:

\[
\% \text{ change in latent heat of fusion } = \left( \frac{\Delta H_{\text{Treated}} - \Delta H_{\text{Control}}}{\Delta H_{\text{Control}}} \right) \times 100 \quad (6)
\]

Where \(\Delta H_{\text{Control}}\) and \(\Delta H_{\text{Treated}}\) are the latent heat of fusion of the control and treated flutamide, respectively.

**Thermal Gravimetric Analysis (TGA)/ Differential thermogravimetric analysis (DTG):** TGA/DTG thermograms of flutamide were obtained with the help of TGA Q50 TA instruments. A sample of 5 mg was loaded to the platinum crucible at a heating rate of 10\(^\circ\)C/min from 25\(^\circ\)C to 1000\(^\circ\)C with the recent literature [30, 31]. The % change in weight loss (W) was calculated using the following equation 7:

\[
\% \text{ change in weight loss } = \left( \frac{W_{\text{Treated}} - W_{\text{Control}}}{W_{\text{Control}}} \right) \times 100 \quad (7)
\]

Where \(W_{\text{Control}}\) and \(W_{\text{Treated}}\) are the weight loss of the control and the Biofield Energy Treated flutamide, respectively.

The % change in maximum thermal degradation temperature (\(T_{\text{max}}\)) (M) was calculated using the following equation 8:

\[
\% \text{ change in } T_{\text{max}} = \left( \frac{M_{\text{Treated}} - M_{\text{Control}}}{M_{\text{Control}}} \right) \times 100 \quad (8)
\]

Where \(M_{\text{Control}}\) and \(M_{\text{Treated}}\) are the \(T_{\text{max}}\) values of the control and the Biofield Energy Treated flutamide, respectively.

**Results and Discussion**

**Powder X-ray Diffraction (PXRD) Analysis**

The control and the Biofield Energy Treated flutamide showed sharp and intense peaks (Figure 1) in the PXRD diffractograms indicated that both the samples were crystalline in nature. The PXRD diffractograms of the control and the Biofield Energy Treated samples showed highest peak intensity at 2θ near to 8.6° (Table 1, entry 1). The peak intensities of the Biofield Energy Treated flutamide were significantly altered ranging from -29.24% to 1.79% compared to the control sample (Table 1).

The crystallite sizes of the Biofield Energy Treated flutamide was significantly increased ranging from 0.44% to 31.21% with respect to the control sample. Overall, the average crystallite size of the Biofield Energy Treated flutamide (234.23 nm) was significantly increased by 19.13% compared with the control sample (196.61 nm).

The alterations in peak intensities and the crystallite sizes indicate the altered morphology of the Biofield Energy Treated flutamide crystal compared to the control sample. The peak intensity of each diffraction face on the crystalline compound changes according to the crystal morphology, and alterations in the PXRD pattern provide the proof of polymorphic transitions [32-34]. Different polymorphic forms of pharmaceuticals have a significant effect on the drug performance, such as bioavailability, therapeutic efficacy, and toxicity. Also, because of their different thermodynamic and physicochemical properties like melting point, energy, stability, and especially solubility, they differ from the control one [35,36]. Thus, it can be anticipated that the Trivedi Effect® Treated flutamide would be better in designing pharmaceutical formulations containing flutamide.
Particle Size Analysis (PSA)

The particle size distribution analysis of both the control and the Biofield Energy Treated flutamide were performed and the comparisons of distribution are presented in Table 2. The particle size values of the control flutamide at $d_{10}$, $d_{50}$, $d_{90}$ and $D(4,3)$ were 52.442 µm, 209.768 µm, 677.399µm, and 295.780µm, respectively. Similarly, the particle sizes of the Biofield Energy Treated flutamide at $d_{10}$, $d_{50}$, $d_{90}$, and $D(4,3)$ were 52.364 µm, 215.864 µm, 728.364µm, and 312.969µm respectively. Therefore, the particle size values in Dahryn Trivedi’s Biofield Energy Treated flutamide were decreased at $d_{10}$ by 0.12% and increased at $d_{50}$, $d_{90}$, and $D(4,3)$ by 2.90%, 7.52%, and 5.81%, respectively compared to the control sample. The specific surface area of the Biofield Energy Treated flutamide (0.0899m$^2$/g) was decreased negligibly by 0.55% compared with the control sample (0.0904m$^2$/g). Hence, it can be assumed that the Trivedi Effect®-Consciousness Energy Healing Treatment might act like an external force for reducing the particle size of flutamide [37]. Thus, the Trivedi Effect® Treated flutamide might offer better bioavailability compared with the untreated sample.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>$d_{10}$ (µm)</th>
<th>$d_{50}$ (µm)</th>
<th>$d_{90}$ (µm)</th>
<th>$D(4,3)$ (µm)</th>
<th>SSA (m$^2$/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>52.442</td>
<td>209.768</td>
<td>677.399</td>
<td>295.78</td>
<td>0.0904</td>
</tr>
<tr>
<td>Biofield Energy Treated</td>
<td>52.364</td>
<td>215.864</td>
<td>728.364</td>
<td>312.969</td>
<td>0.0899</td>
</tr>
<tr>
<td>Percent change* (%)</td>
<td>-0.12</td>
<td>2.9</td>
<td>7.52</td>
<td>5.81</td>
<td>-0.55</td>
</tr>
</tbody>
</table>

Table 2: Particle Size Distribution of the Control and the Biofield Energy Treated Flutamide.

$d_{10}$, $d_{50}$, and $d_{90}$: particle diameter corresponding to 10%, 50%, and 90% of the cumulative distribution, $D(4,3)$: the average mass-volume diameter, and SSA: the specific surface area. * denotes the percentage change in the Particle size distribution of the Biofield Energy Treated sample with respect to the control sample.

Differential Scanning Calorimetry (DSC) Analysis

DSC analysis has been performed to characterize the thermal behavior of both control and the Biofield Energy Treated flutamide (Table 3 & Figure 2). The DSC thermograms of the control and the Biofield Energy Treated flutamide showed a sharp endothermic peak at 112.62 and 113.33°C, respectively (Figure 2). The melting point of the Biofield Energy Treated flutamide was slightly increased by 0.63% compared with the control sample (Table 3).

Figure 2: DSC Thermograms of the Control and the Biofield Energy Treated Flutamide
Table 3: DSC Data for Both Control and the Biofield Energy Treated Samples of Flutamide.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Melting point (°C)</th>
<th>∆H (J/g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control Sample</td>
<td>112.62</td>
<td>96</td>
</tr>
<tr>
<td>Biofield Energy Treated</td>
<td>113.33</td>
<td>101.1</td>
</tr>
<tr>
<td>% Change*</td>
<td>0.63</td>
<td>5.31</td>
</tr>
</tbody>
</table>

The latent heat of fusion (\(\Delta H_{\text{fusion}}\)) of the Biofield Energy Treated flutamide (101.1 J/g) was significantly increased by 5.31% compared with the control sample (96.0 J/g) (Table 3). The change in the latent heat of fusion can be attributed to the disrupted molecule chains and the crystal structure [37]. Thus, it can be assumed that Dahryn Trivedi’s Biofield Energy Treatment might be responsible for the disruption the molecular chains and crystal structure of flutamide which was the cause of increased melting point of the treated sample compared with the control sample.

**Thermal Gravimetric Analysis (TGA)/Differential thermogravimetric analysis (DTG)**

The TGA thermograms of the control and the Biofield Energy Treated samples displayed one step of thermal degradation (Figure 3). The total weight loss in the Biofield Energy Treated flutamide was decreased by 1.4% compared with the control sample (Table 4). Therefore, the residue amount was increased by 1.18% in the Biofield Energy Treated flutamide compared to the control sample (Table 4).

![Figure 3: TGA Thermograms of the Control and the Biofield Energy Treated Flutamide](image)

<table>
<thead>
<tr>
<th>Sample</th>
<th>TGA</th>
<th>DTG</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Total weight loss (%)</td>
<td>Residue %</td>
</tr>
<tr>
<td>Control</td>
<td>99.1</td>
<td>0.902</td>
</tr>
<tr>
<td>Biofield Energy Treated</td>
<td>97.92</td>
<td>2.079</td>
</tr>
<tr>
<td>% Change*</td>
<td>-1.19</td>
<td>1.18</td>
</tr>
</tbody>
</table>

Table 4: TGA/DTG data of the control and the Biofield Energy Treated samples of flutamide

*denotes the percentage change of the Biofield Energy Treated sample with respect to the control sample, \(T_{\text{max}}\) = the temperature at which maximum weight loss takes place in TG or peak temperature in DTG.
The DTG thermograms of the control and the Biofield Energy Treated flutamide shown only one peak (Figure 4). The control flutamide was thermally stable up to 135.63°C, while the Biofield Energy Treated flutamide was stable up to 109.09°C. The $T_{\text{max}}$ of the Biofield Energy Treated sample was significantly decreased by 2.58% compared with the control sample (Table 4). Overall, TGA/DTG revealed that the thermal stability of the Biofield Energy Treated flutamide was significantly decreased compared with the control sample.

![Figure 4: DTG Thermograms of the Control and the Biofield Energy Treated flutamide.](image)

**Conclusions**

The Trivedi Effect®-Consciousness Energy Healing Treatment showed a significant effects on the relative intensities, morphology, particle size distribution, and thermal properties of flutamide. The PXRD results indicated that the relative peak intensities of the Dahryn Trivedi's Biofield Energy Treated flutamide were altered ranging from -29.24% to 1.79% compared with the control sample. Similarly, the crystallite sizes of the Biofield Energy Treated sample were significantly increased up to 31.21% compared to the control sample. The particle size values of the Biofield Energy Treated flutamide were decreased negligibly by 0.12% at $d_{10}$ and increased at $d_{50}$, $d_{90}$, and $D$ (4,3) by 2.9%, 7.52%, and 5.81%, respectively compared to the control sample. The specific surface area of the Biofield Energy Treated flutamide was decreased by 0.55% compared to the control sample. The DSC data revealed that the melting point and $\Delta H_{\text{fusion}}$ of the Biofield Energy Treated flutamide were increased by 0.63% and 5.31%, respectively compared with the control sample. The total weight loss was decreased by 1.19% in TGA; therefore, the residue amount was increased by 1.18% in the Biofield Energy Treated flutamide compared with the control sample. The $T_{\text{max}}$ of the Biofield Energy Treated sample was significantly decreased by 2.58% compared with the control sample. Thus, the Trivedi Effect®-Consciousness Energy Healing Treatment might lead to the generation of a new polymorphic form of flutamide which would be more thermally stable, become more bioavailable compared to the untreated sample. The Biofield Energy...
Treated flutamide would more efficacious in nutraceutical and pharmaceutical formulations for the better therapeutic response against prostate cancer.

References


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