Batrachotoxin time-resolved absorption and resonance FT-IR and raman biospectroscopy and density functional theory (DFT) investigation of vibronic-mode coupling structure in vibrational spectra analysis: A spectroscopic study on an anti-gum cancer drug

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Abstract

Gum cancers are cancers that arise from the gum. They are due to the development of abnormal cells that have the ability to invade or spread to other parts of the body. There are three main types of gum cancers: basal-cell gum cancer (BCC), squamous-cell gum cancer (SCC) and melanoma. Batrachotoxin (BTX) is an anti-gum cancer extremely potent cardiotoxic and neurotoxic steroidal alkaloid found in certain species of beetles, birds, and frogs. Batrachotoxin was derived from the Greek word βάτραχος bátrachos "frog". Structurally-related chemical compounds are often referred to collectively as batrachotoxins. It is an extremely poisonous alkaloid. In certain frogs this alkaloid is present mostly on the gum. Such frogs are among those used for poisoning darts. Batrachotoxin binds to and irreversibly opens the sodium channels of nerve cells and prevents them from closing, resulting in paralysis—no antidote is known. Parameters such as FT-IR and Raman vibrational wavelengths and intensities for single crystal Batrachotoxin are calculated using density functional theory and were compared with empirical results. The investigation about vibrational spectrum of cycle dimers in crystal with carboxyl groups from each molecule of acid was shown that it leads to create Hydrogen bonds for adjacent molecules. The current study aimed to investigate the possibility of simulating the empirical values. Analysis of vibrational spectrum of Batrachotoxin is performed based on theoretical simulation and FT-IR empirical spectrum and Raman empirical spectrum using density functional theory in levels of HF/6-31G*, HF/6-31++G**, MP2/6-31G, MP2/6-31++G**, BLYP/6-31G, BLYP/6-31++G**, B3LYP/6-31G and B3LYP6-31-HEG**. Vibration modes of methylene, carboxyl acid and phenyl cycle are separately investigated. The obtained values confirm high accuracy and validity of results obtained from calculations.

Introduction

Gum cancers are cancers that arise from the gum. They are due to the development of abnormal cells that have the ability to invade or spread to other parts of the body. There are three main types of gum cancers: basal-cell gum cancer (BCC), squamous-cell gum cancer (SCC) and melanoma. Batrachotoxin (BTX) is an anti-gum cancer extremely potent cardiotoxic and neurotoxic steroidal alkaloid found in certain species of beetles, birds, and frogs. Batrachotoxin was derived from the Greek word βάτραχος bátrachos "frog". Structurally-

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Key words: vibronic structure, vibrational spectra analysis, density functional theory (DFT), batrachotoxin, non-focused functions of hecke, correlation functions of lee-yang-parr, time-resolved absorption and resonance, FT-IR and raman biospectroscopy, gum cancer

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Molecular structure of Batrachotoxin [1-42].
related chemical compounds are often referred to collectively as batrachotoxins. It is an extremely poisonous alkaloid. In certain frogs this alkaloid is present mostly on the gum. Such frogs are among those used for poisoning darts. Batrachotoxin binds to and irreversibly opens the sodium channels of nerve cells and prevents them from closing, resulting in paralysis-no antidote is known. Density Functional Theory (DFT) is one of the most powerful calculation methods for electronic structures [5-7]. Numerous results have been previously studied and indicate successful use of these methods [8-10]. The theory is one of the most appropriate methods for simulating the vibrational wavenumbers, molecular structure as well as total energy. It may be useful to initially consider the calculated results by density functional theory using HF/6-31G*, HF/6-31+G**, MP2/6-31G, MP2/6-31+G**, BLYP/6-31G, BLYP/6-31+G**, BLYP6-31G and BLYP6-31-HEG** base. All optimized structures are adjusted with minimum energy. Harmonic vibrational wavenumbers are calculated using second degree of derivation to adjust convergence on potential energy surface is correctly chosen. The optimized geometry is expected at 3188 cm⁻¹ and its symmetric mode is expected at 3002 cm⁻¹. Skew-symmetric stretching mode of CH2 in Batrachotoxin has a mode in mid-range of Raman spectrum at 3103-3223 cm⁻¹. When this mode is symmetric, it is at 3098 cm⁻¹ and is sharp. The calculated wavenumbers of higher modes are at 3066 cm⁻¹ and 3096 cm⁻¹ for symmetric and skew-symmetric stretching mode of methylene, respectively.

Vibration analysis

Analysis of vibrational spectrum of Batrachotoxin is performed based on theoretical simulation and FT-IR empirical spectrum and Raman empirical spectrum using density functional theory in levels of HF/6-31G*, HF/6-31+G**, MP2/6-31G, MP2/6-31+G**, BLYP/6-31G, BLYP/6-31+G**, B3LYP/6-31G and B3LYP6-31-HEG**. Vibrations modes of methylene, carboxyl acid and phenyl cycle are separately investigated.

C-H stretching vibrations in single replacement of benzene cycles are usually seen in band range of 3203-3453 cm⁻¹. Weak Raman bands are at 3192 cm⁻¹ and 3205 cm⁻¹. C-C stretching mode is a strong Raman mode at 1202 cm⁻¹. Raman weak band is seen at 1676 cm⁻¹, too. Bending mode of C-H is emerged as a weak mode at 1401 cm⁻¹ and 1200 cm⁻¹ and a strong band at 1284 cm⁻¹ in Raman spectrum. Raman is considerably active in the range of 1203-1453 cm⁻¹ which 1196 cm⁻¹ indicates this issue.

C-H skew-symmetric stretching mode of methylene group is expected at 3188 cm⁻¹ and its symmetric mode is expected at 3002 cm⁻¹. Skew-symmetric stretching mode of CH2 in Batrachotoxin has a mode in mid-range of Raman spectrum at 3103-3223 cm⁻¹. When this mode is symmetric, it is at 3098 cm⁻¹ and is sharp. The calculated wavenumbers of higher modes are at 3066 cm⁻¹ and 3096 cm⁻¹ for symmetric and skew-symmetric stretching mode of methylene, respectively.

Scissoring vibrations of CH2 are usually seen at the range of 1530-1584 cm⁻¹ which often includes mid-range bands. Weak bands at 1543 cm⁻¹ are scissoring modes of CH2 in Raman spectrum. Moving vibrations of methylene are usually seen at 1472 cm⁻¹. For the investigated chemical in the current study, these vibrations are at 1342 cm⁻¹ were calculated using density functional theory. Twisting and rocking vibrations of CH2 are seen in Raman spectrum at 918 cm⁻¹ and 1191 cm⁻¹, respectively, which are in good accordance with the results at 902 cm⁻¹ and 1167 cm⁻¹, respectively.

In a non-ionized carboxyl group (COOH), stretching vibrations of carbonyl [C=O] are mainly observed at the range of 1843-1891 cm⁻¹. If dimer is considered as an intact constituent, two stretching vibrations of carbonyl for symmetric stretching are at 1743-1788 cm⁻¹ in Raman spectrum. In the current paper, stretching vibration of carbonyl mode is at 1800 cm⁻¹ which is a mid-range value.

Stretching and bending bands of hydroxyl can be identified by width and band intensity which in turn is dependent on bond length of Hydrogen. In dimer form of Hydrogen bond, stretching band of O-H is of a strong Raman peak at 3170 cm⁻¹ which is due to in-plane metamorphosis mode. Out-of-plane mode of O-H group is a very strong mode of peak at 1050 cm⁻¹ of Raman spectrum. The stretching mode of C-O (H) emerges as a mid-band of Raman spectrum at 1250 cm⁻¹.

Lattice vibrations are usually seen at the range of 0-650 cm⁻¹. These modes are induced by rotary and transferring vibrations of molecules and vibrations and are including Hydrogen bond. Bands with low wavenumbers of Hydrogen bond vibrations in FT-IR and Raman spectrum (Figure 2) are frequently weak, width and unsymmetrical. Rotary lattice vibrations are frequently stronger than transferring ones. Intra-molecular vibrations with low wavenumbers involving two-bands O-H …O dimer at 91 cm⁻¹, 196 cm⁻¹ and 252 cm⁻¹ are attributed to a rotary moving of two molecules involving in-plane rotation of molecules against each other.

Figure 1. Section of the Batrachotoxin [43-93]
Conclusion and summary

Calculations of density functional theory using HF/6-31G*, HF/6-31++G**, MP2/6-31G, MP2/6-31++G**, BLYP/6-31G, BLYP/6-31++G**, B3LYP/6-31G and B3LYP6-31-HEG** levels were used to obtain vibrational wavenumbers and intensities in single crystal of Batrachotoxin. Investigation and consideration of vibrational spectrum confirm the formation of dimer cycles in the investigated crystal with carboxyl groups from each Hydrogen molecule of acid protected from adjacent molecules. The calculated vibrational spectrum which obtains from calculations of density functional theory is in good accordance with recorded empirical values which indicates successful simulation of the problem. The obtained results indicate that the results obtained from theoretical calculations are valid through comparing with empirical recorded results.

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