Entangled biphoton generation in myelin sheath

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Consciousness within the brain hinges on the synchronized activities of millions of neurons, but the mechanism responsible for orchestrating such synchronization remains elusive. In this study, we employ cavity quantum electrodynamics (cQED) to explore entangled biphoton generation through cascade emission in the vibration spectrum of C-H bonds within the lipid molecules' tails. The results indicate that the cylindrical cavity formed by a myelin sheath can facilitate spontaneous photon emission from the vibrational modes and generate a significant number of entangled photon pairs. The abundance of C-H bond vibration units in neurons can therefore serve as a source of quantum entanglement resources for the nervous system. The finding may offer insight into the brain's ability to leverage these resources for quantum information transfer, thereby elucidating a potential source for the synchronized activity of neurons.

I. INTRODUCTION

Understanding the intricacies of the human brain and its functions has perennially posed an intriguing and challenging puzzle. The synchronization of neurons within the cerebral cortex serves as the foundation for diverse neurobiological processes[1], closely linked to anomalies in brain function and brain diseases[2]. Notably, Parkinson's disease manifests a loss of neural activity synchronization in regions with damaged neurons[3]. Despite these observations, the mechanisms underpinning precise synchronization of neural activity remain unknown[4], necessitating interdisciplinary research, particularly in the realms of neuroscience and quantum physics.

In recent decades, quantum computing, harnessing the unique features of quantum entanglement, has witnessed remarkable success[5]. Experiments validating nonlocal correlations in quantum entanglement[6] have enabled quantum computation to outpace classical counterparts in tasks such as the Shor[7] and Grover[8] algorithms. Quantum computing's applicability to neuroscience was initially proposed by Penrose, who suggested a role for microtubules in quantum computation within the brain[4], and further explored for example, by Fisher who proposed nuclear spins as mediators[9]. Despite experimental deviations from predictions of these models[10, 11], the nonlocal correlations inherent in quantum entanglement remain captivating.

Recent studies highlight the role of photon as a quantum object not only in plants and bacteria but also in animal life activities[12]. Examples include mid-infrared (MIR) photons from ATP hydrolysis driving DNA replication[13] and polaritons formed by visible light coherently and resonantly coupled to the excitons of chlorophyll molecules in chloroplasts facilitating efficient energy transfer in photosynthesis[14]. Ultra-weak photon emission (UPE) in living organisms, traditionally consid-ered as metabolic by-products, is now implicated in neuronal function[15]. Moreover, MIR photons at 53.7 THz modulate K^+ ion channel activity, neuronal signaling, and sensorimotor behavior[16]. Photons released during the TCA cycle in neurons resonantly couple to C-H bond vibrons in lipid molecules, altering possibly the dielectric constant of the membrane to enhance action potential conduction[17]. These findings, though each may require further scrutiny, offer an alternative perspective on the significance of light in neural activity.

Myelin sheath, a lipid membrane encasing the outer side of a neuron's axon, provides energy to axon, enhances action potential conduction, and acts as an insulator in the nervous system[18]. Abnormal myelin function or damage to the myelin structure is strongly associated with neurodegenerative diseases such as multiple sclerosis and Alzheimer's disease[19]. Myelin sheath is generally regarded only as an insulator. However, emerging evidence suggests myelin's plasticity, indicating its role beyond insulation and its potential to promote neural phase synchronization[20].

This study demonstrates that the vibrational spectrum of C-H bonds in lipid molecule's tails, within cylindrical cavities formed by myelin sheath encasement, can generate quantum entangled photon pairs through cascade radiation from the second excited state to the ground state. In Sec.II, we establish the axon's well-defined cylindrical structure under the myelin sheath and discuss the quantization of the electromagnetic field within the cylindrical cavity. We show that, within the infrared region and under the dipole approximation, two-photon processes in the vibrational spectrum are predominantly governed by cascade radiation from dipole interactions in Sec.III. Utilizing Schmidt analysis, we assess the degree of quantum entanglement in biphoton systems and exemplify the potential for generating quantum entanglement in neural systems using real structural data of myelinated neuron from experiments in Sec.IV. Leveraging the nonlocal correlation properties of quantum entanglement, one may speculate that quantum entanglement will effectively synchronize neuronal activity throughout the brain, shedding light on the 'synchronization' puzzle

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in consciousness.

II. CYLINDRICAL CAVITY FORMED BY MYELIN SHEATH

As shown in Figure.1(a), oligodendrocytes in the central nervous system (CNS) form myelin sheaths by wrapping lipid-rich membranes around axons, and because of their high content of myelin, which is white in colour, axons surrounded by myelin sheaths in the CNS are called white matter[21]. More than half of the human brain is white matter, which supports the rapid and simultaneous transmission of information between numerous grey matter areas of the CNS[20]. The formation of microcavities is crucial in exciton-polaron mediated energy transfer in the recent progress of explaining the mechanism of high efficient energy transfer[14, 22]. The total reflection of light satisfying the evanescent wave condition on the inner wall of the cavity causes the confinement of photons by the cavity. The myelin sheath, which generally consists of 10^2 lipid bilayers (Figure.1(c)) wrapped around the axon, serves mainly as an insulator in neurons and can have a better confinement effect on photons due to the formation of polaritons in the myelin sheath[17]. For simplicity and without loss of generosity, it is reasonable to consider the outer wall of the myelin sheath as a perfect conductor wall, considering only that electromagnetic fields do not leak outside the myelin sheath. Concentrically wrapped around the periphery of the axon, myelin together with the axon forms a cylinder-like structure, and based on this fact, we consider myelinated neuronal axons as cylinders (Figure.1(b)). The thickness of the myelin sheath is generally in the range of 1∼3 μm [23]. The ratio of the length between two neighbouring nodes of Ranvier to the thickness of the myelin sheath is ∼100 , and the width of the nodes of Ranvier is generally in the range of 1∼2 μm[24], so that we can ignore the gaps caused by the nodes of Ranvier, and consider the multiple myelin sheaths merged together as a whole. In this way, we view the myelin-coated axonal portion of the entire myelinated neuron as a conductor-walled cylinder, as shown in Figure.1(b). The C-H bond dipole in the lipid tail is between the inner diameter a and outer diameter b.

The Hamiltonian of the quantised electromagnetic fields and the electric field inside the cylindrical cavity can be written respectively as[25]

$$\boldsymbol{E}(\boldsymbol{R},t) = \sum_{s\sigma} f_{s\sigma}(a_{s\sigma}(t) + a_{s\sigma}^{\dagger}(t))\boldsymbol{u}_{s\sigma}(\boldsymbol{R}), \quad (1)$$

$$H_R = \sum_{s\sigma} \hbar\omega_{s\sigma}\left(a_{s\sigma}^{\dagger}a_{s\sigma} + \frac{1}{2}\right), \quad (2)$$

where $\boldsymbol{u}_{s\sigma}$ and $a_{s\sigma}$ are respectively the vector mode functions and annihilation operator and $f_{\omega} = \sqrt{\hbar\omega_{s\sigma}/(2\epsilon_0)}$. $s = (m, \mu, n)$ is the mode index and $\sigma = 0, 1$ represents polarization.

The vector mode functions $\boldsymbol{u}_{s\sigma}(\sigma = 0, 1)$ take the form

$$\boldsymbol{u}_{s1} = \nabla \times \nabla \times \boldsymbol{e_z}\psi_{s1}, \quad (3)$$

$$\boldsymbol{u}_{s2} = i\omega_{s2}\nabla \times \boldsymbol{e_z}\psi_{s2}, \quad (4)$$

where ψ_{s1} and ψ_{s2} have the form

$$\psi_{s1} = c_{s1}J_m(\frac{\chi_{s1}R}{b})e^{im\varphi}\cos\left(\frac{n\pi z}{L}\right), \quad (5)$$

$$\psi_{s2} = c_{s2}J_m(\frac{\chi_{s2}R}{b})e^{im\varphi}sin(\frac{n\pi z}{L}). \quad (6)$$

$\chi_{s1} \equiv \chi_{m\mu1}$ is the μth zero point of the first kind of Bessel function J_m, $\chi_{s2} \equiv \chi_{m\mu2}$ is the μth zero of J'_m(the derivative of J_m). $c_{s\sigma}$ is a normalization constant, $c_{s1} = \sqrt{(2c^2b^2)/(V\alpha_{s1}\chi_{s1}^2\omega_{s1}^2)}$, $c_{s2} = \sqrt{(2b^2)/(V\alpha_{s2}\chi_{s2}^2\omega_{s2}^2)}$, Here $\alpha_{s1} \equiv \alpha_{m\mu1} = J_{m+1}^2(\chi_{m\mu1})$, $\alpha_{s2} \equiv \alpha_{m\mu2} = J_m^2(\chi_{m\mu2}) - J_{m+1}^2(\chi_{m\mu2})$. d is the diameter of the cylindrical cavity, L is the length of the cylindrical cavity. The eigen modes of the cavity are determined by

$$\omega_{s\sigma} = c\sqrt{\left(\frac{\chi_{s\sigma}}{b}\right)^2 + \left(\frac{n\pi}{L}\right)^2}. \quad (7)$$

III. RADIATIVE CASCADE WITHIN THREE-LEVEL MORSE OSCILLATOR

In this section, we first introduce morse oscillator which is extensively used for depicting the vibration of chemical bonds. Then we discuss two-photon processes via radiative cascade by dipole interaction. For obtaining actual wave function of biphoton, we use the real size data of myelinated neuron by experiments.

III.1. Chemical Bonds Described by Morse Oscillators

In contrast to the rotation of a chemical bond and its electronic energy levels (Figure.2(b)), the transitions in the vibrational energy spectrum are typically on the order of 0.1 eV, placing them in the mid-infrared band. This band lies in the range that has a significant influence on neural activity. The vibrations of chemical bonds are generally anharmonic oscillators, and the energy levels are not equally spaced. In order to describe anharmonic vibrations, we adopt Morse oscillators[26] to describe actual chemical bond vibrations as shown in Figure.2(b). The advantage of using a Morse oscillator is that its determinant Schrödinger equation has an analytical solution, which makes it very convenient for us to calculate the dipole moment of the chemical bond. the potential function of Morse oscillator can be written as[27]

$$V(r) = D_e\left(e^{-2w(r-r_e)} - 2e^{-w(r-r_e)}\right). \quad (8)$$

where r_e is bond length at equilibrium. D_e and w represent the depth and the width of the potential well respectively. Its stationary Schrödinger equation $H_P|\Psi_\nu\rangle =$
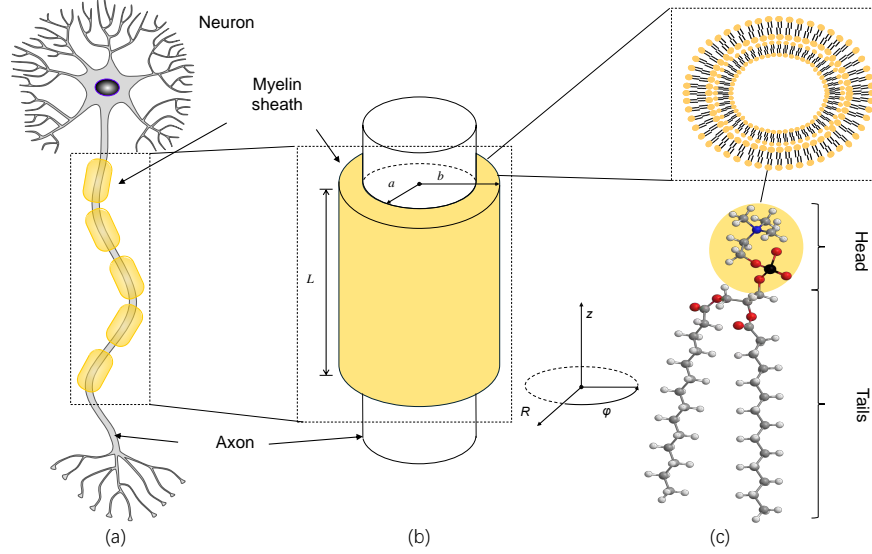
FIG. 1: Cylindrical cavity formed by myelin sheath. (a) Axons of myelinated neuron have several segments of myelin sheaths wrapped in lipid membranes of different oligodendrocytes. The gap between the two segments of the myelin sheath is called node of Ranvier and is usually 1~2 μm in length, whereas each segment of the myelin sheath is typically around 100 μm in length. The gap in the node of Ranvier is negligible for the whole axon. (b) Consider the entire myelin-coated axon as a cylinder, with the radius of the axon as a, the radius of the entire cylinder as b, and the thickness of the myelin sheath as $d = b - a$. The length of the cylinder is denoted by L. Let the central axis of the axon be the z-axis to establish an axial coordinate system. (c) Phospholipid molecules, as the major component of myelin, have tails consisting of a large number of C-H bonds.

$E_\nu |\Psi_\nu\rangle$ has an analytical solution, which is

$$|\Psi_\nu\rangle = N_\nu z^{\lambda-\nu-\frac{1}{2}} e^{-\frac{1}{2}z} L_\nu^{(2\lambda-2\nu-1)}(z), \qquad (9)$$

$$E'_\nu = -\left(\lambda - \nu - \frac{1}{2}\right)^2 \qquad (10)$$

$$\nu = 0, 1, \cdots, [N] = [\lambda - \frac{1}{2}],$$

with variable substitutions $z = 2\lambda e^{-w(r-r_e)}$ and $E'_\nu = [(2m)/(w^2\hbar^2)]E_\nu$. In the above, $L_\nu^{(2\lambda-2\nu-1)}(z)$ is a generalized Laguerre polynomial, $[\lambda - \frac{1}{2}]$ represents the largest integer not exceeding $[\lambda - \frac{1}{2}]$, and the normalization constant $N_\nu = \sqrt{[n!(2\lambda-2\nu-1)]/[\Gamma(2\lambda-\nu)]}$ with $\Gamma(x)$ being the gamma function.

The matrix elements of positional operators also have the following analytical forms[28]

$$\langle \Psi_\nu|r|\Psi_{\nu'}\rangle = \frac{2(-1)^{\nu'-\nu+1}}{(\nu'-\nu)(2N-\nu-\nu')} \times \sqrt{\frac{(N-\nu)(N-\nu')\Gamma(2N-\nu'+1)\nu'!}{\Gamma(2N-\nu+1)\nu!}}. \qquad (11)$$

III.2. Biphoton Wave Function via Cascade Radiation

The Hamiltonian of the interaction of matter with an electromagnetic field is usually written as[29]

$$H_I = -\frac{e}{m}\boldsymbol{p}\cdot\boldsymbol{A} + \frac{e^2}{2m}\boldsymbol{A}^2 + \hbar g^s \boldsymbol{s}\cdot\boldsymbol{B}, \qquad (12)$$

where $\boldsymbol{p}$ and $\boldsymbol{A}$ are the momentum of the matter and vector potential of electromagnetic field, respectively, and g^s is the gyromagnetic ratio of the particle. The last two terms describe two-photon processes under strong electromagnetic fields and strong interactions[29], respectively, which should be ignored considering that strong field environments are not usually existed in neurons. Under the long-wave approximation (or dipole approximation), the first term can be written as $-\boldsymbol{D}\cdot\boldsymbol{E}$, where $\boldsymbol{D}$ and $\boldsymbol{E}$ are the dipole moment of the particle and electric field respectively. The process of emitting two photons by jumping from the first excited state to the ground state under dipole interaction is forbidden, so we will consider the process of emitting two photons from the second excited state to the ground state by cascade radiation[30], as shown in Figure.2(c).
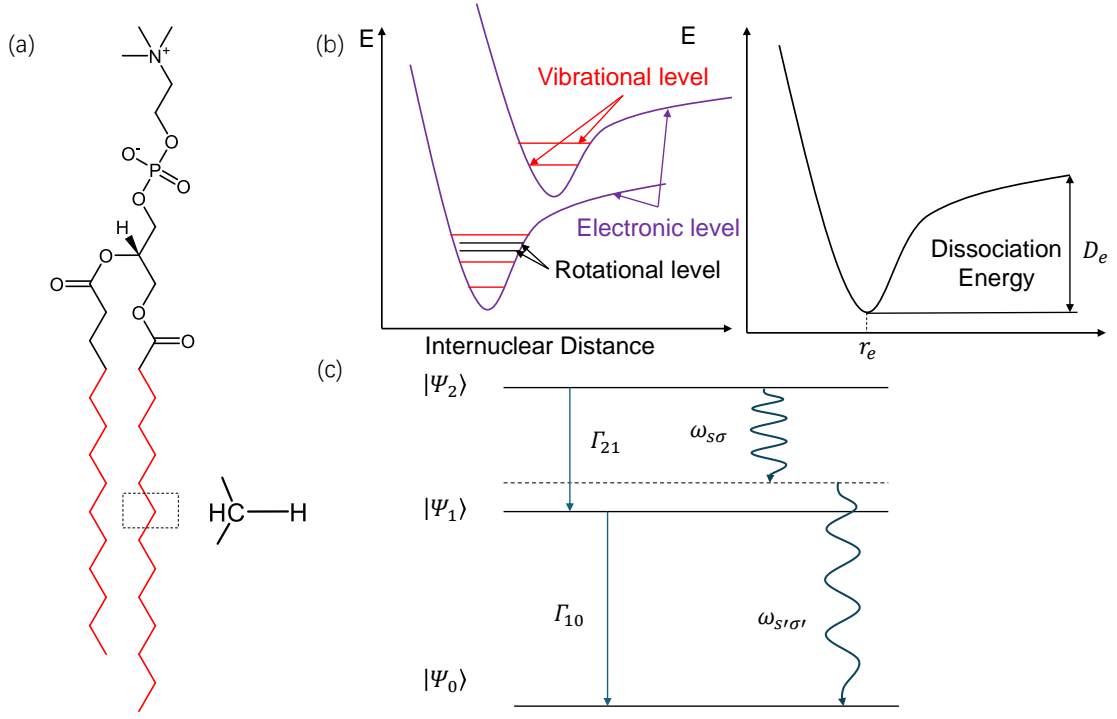
FIG. 2: Cascade emission from vibrational spectrum. (a) A skeletal formula of a phospholipid molecule, with the red part representing the carbon chain in the tail, consisting of several methylene groups and two methyl groups. (b) Left: Schematic diagram of molecular energy levels, with electronic energy levels indicated in purple, rotational energy levels in black, and vibrational energy levels in red. Right: Vibrational energy levels are generally represented by Morse oscillators. The lowest point of the potential energy corresponds to the equilibrium bond length of the chemical bond r_e, and the depth of the potential energy characterizes the dissociation energy D_e. (c) Cascade emission in three-level system. This figure represents a three-level system returning to the ground state from the second excited state by emitting photons with frequencies $\omega_{s\sigma}$ and $\omega_{s'\sigma'}$. Γ_{21} and Γ_{10} are the transition rates(or the reciprocal of lifetime) between states.

Considering the rotating wave approximation, the interacting Hamiltonian under the Dirac picture can be written as

$$H_I(t) = \hbar \sum_{s\sigma} \left[g_\omega^* \sigma_+^{(1)} a_{s\sigma} e^{i(\omega_{21}-\omega_{s\sigma})t} + h.c. \right]$$
$$+ \hbar \sum_{s'\sigma'} \left[g_{\omega'}^* \sigma_+^{(2)} a_{s'\sigma'} e^{i(\omega_{10}-\omega_{s\sigma})t} + h.c. \right], \quad (13)$$

where $\sigma_+^{(1)} = |\Psi_2\rangle \langle\Psi_1|$, $\sigma_+^{(2)} = |\Psi_1\rangle \langle\Psi_0|$, and g_ω and $g_{\omega'}$ are states $|\Psi_2\rangle$ to state $|\Psi_1\rangle$ and state $|\Psi_1\rangle$ to $|\Psi_0\rangle$ state transition coupling constant which take the form of

$$g_\omega = \frac{f_\omega \langle\Psi_1|\boldsymbol{D}|\Psi_2\rangle \cdot \boldsymbol{u}_{s\sigma}}{\hbar}, \quad (14)$$

$$g_{\omega'} = \frac{f_{\omega'} \langle\Psi_0|\boldsymbol{D}|\Psi_1\rangle \cdot \boldsymbol{u}_{s'\sigma'}}{\hbar}. \quad (15)$$

Similarly to the Jaynes-Cummings model[31], we let the total number of excitations between the atom and the cavity mode be conserved, and let the vibrational energy levels and the state vectors of the photons be of the form

$$|\Psi(t)\rangle = c_0(t) |\Psi_2, 0\rangle + c_1(t) \sum_{s\sigma} |\Psi_1, \omega_{s\sigma}\rangle$$
$$+ c_2(t) \sum_{s'\sigma'} |\Psi_0, \omega_{s\sigma}, \omega_{s'\sigma'}\rangle. \quad (16)$$

Substituting Eq.(16) into the time-dependent Schrödinger equation, we have three equations of motion for the probability amplitudes

$$\dot{c}_0 = -i \sum_{s\sigma} g_\omega^* c_1 e^{i(\omega_{21}-\omega_{s\sigma})t}, \quad (17)$$

$$\dot{c}_1 = -i g_\omega c_0 e^{i(\omega_{21}-\omega_{s\sigma})t} - i \sum_{s'\sigma'} g_{\omega'}^* c_2 e^{i(\omega_{10}-\omega_{s'\sigma'})t}, \quad (18)$$

$$\dot{c}_2 = -i g_{\omega'} c_1 e^{i(\omega_{10}-\omega_{s'\sigma'})t}. \quad (19)$$

We begin by looking at the first two equations, and in order to simplify the system of equations, we first integrate Eq.(19) and substitute into Eq.(18) without considering c_0 to get

$$\dot{c}_1 = -\sum_{s'\sigma'} |g_{\omega'}|^2 \int_0^t dt' e^{i(\omega_{10}-\omega_{s'\sigma'})(t-t')} c_1(t'). \quad (20)$$

This is still an exact equation. Next we make some approximations. The dipole moment of the C-H bond in the tail of the lipid molecule is overwhelmingly perpendicular to the radial direction, and for the sake of computational convenience, we assume that the dipole moment is only in the z-direction. It is obvious that $\boldsymbol{u}_{s2}$ has no z component from Eq.(4), and we only need to find out the z component of $\boldsymbol{u}_{s1}$. By expanding Eq.(3) we get $\boldsymbol{u}_{s1} = \nabla(\nabla \cdot \boldsymbol{e_z}\psi_{s1}) - \nabla^2 \boldsymbol{e_z}\psi_{s1} = k_{s1}^2\boldsymbol{e_z}\psi_{s1} + \nabla(\partial_z\psi_{s1})$. Then we have

$$\boldsymbol{u}_{s1,z} = \boldsymbol{e_z}\sqrt{\frac{2c^2\chi_{s1}^2}{Vb^2\alpha_{s1}\omega_{s1}^2}}J_m(\frac{\chi_{s1}R}{b})\cos(\frac{n\pi z}{L})e^{im\varphi}. \quad (21)$$

Now we have explicit form of the coupling constant

$$g_{\omega'} = d_{01}\sqrt{\frac{c^2\chi_{s'1}^2}{\hbar\epsilon_0 Vb^2\alpha_{s1}\omega_{s'1}}}J_{m'}(\frac{\chi_{s'1}R}{b})\cos(\frac{n'\pi z}{L})e^{im\varphi}, \quad (22)$$

where d_{01} is the matrix element of the dipole moment $\boldsymbol{D}$ which is $\langle\Psi_0|\boldsymbol{D}|\Psi_1\rangle$.

In general the length of the myelin sheath is much larger than the radius of the axon, at which point the energy level gap of the cavity mode is very small, and we can convert the summation into integral in Eq.(20), in other words

$$\lim_{L\to\infty}\frac{1}{L}\sum_{n'} = \frac{1}{c\pi}\int_{c\chi_{s'1}/b}^{\infty}\frac{d\omega_{s'1}\omega_{s'1}}{\sqrt{\omega_{s'1}^2 - c^2\chi_{s'1}^2/b^2}}. \quad (23)$$

Now Eq.(20) becomes

$$\dot{c}_1 = -\sum_{m'\mu'}\frac{d_{01}^2 c\chi_{s'1}^2}{\hbar\epsilon_0\pi b^4\alpha_{s1}}J_{m'}^2(\frac{\chi_{s'1}R}{b})\cos^2(\frac{n'\pi z}{L})\int_{c\chi_{s'1}/b}^{\infty}d\omega_{s'1}\int_0^t dt'\frac{e^{i(\omega_{10}-\omega_{s'1})(t-t')}}{\sqrt{\omega_{s'1}^2 - c^2\chi_{s'1}^2/b^2}}c_1(t'). \quad (24)$$

In the emission spectrum, the frequency of light associated with the emitted radiation is going to be centered about the eigen vibration frequency ω_{10}. The quantity $\omega_{s'1}$ varies little around $\omega_{s'1} = \omega_{10}$ for which the time integral in Eq.(24) is not negligible. We can then replace $\omega_{s'1}$ by ω_{10} and the lower limit in the $\omega_{s'1}$ integration by $-\infty$. Then we have

$$\int_{-\infty}^{\infty}d\omega_{s'1}e^{i(\omega_{10}-\omega_{s'1})(t-t')} = 2\pi\delta(t-t'). \quad (25)$$

Since the term $\sqrt{\omega_{s'1}^2 - c^2\chi_{s'1}^2/b^2}$ in the denominator has no physical meaning at less than 0, so that a step function will be added to Eq.(24). The Eq.(24) becomes

$$\dot{c}_1 = \frac{3\Gamma_{10}^{(0)}}{\xi_{10}^3}\sum_{m'\mu'}\frac{\chi_{s'1}^2}{\alpha_{s'1}\sqrt{\xi_{10}^2 - \chi_{s'1}^2}}J_{m'}^2(\chi_{s'1}R/b) \\ \times \cos^2(\sqrt{\xi_{10}^2 - \chi_{s'1}^2}z/L)\theta(\xi_{10} - \chi_{s'1})c_1(t), \quad (26)$$

where $\xi_{10} = \omega_{10}a/c$, $\theta(x) = 1(x>0), 0(x<0)$ and

$$\Gamma_{10}^{(0)} = \frac{e^2|\langle\Psi_0|\boldsymbol{D}|\Psi_1\rangle|^2\omega_{10}^3}{3\pi\hbar\varepsilon_0 c^3} \quad (27)$$

is the transition rate in free space. By using Eq.(4-12) in [30], we have

$$\dot{c}_1 = -\frac{\Gamma_{10}}{2}c_1, \quad (28)$$

where Γ_{10} is the transition rate between $|\Psi_1\rangle$ and $|\Psi_0\rangle$ in cavity. In this way the physical meaning of Eq.(26) is clear: at $\xi_{10} < \chi_{011}$, transition between $|\Psi_1\rangle$ and $|\Psi_0\rangle$ is forbidden. Using the same approach similar to Weisskopf-Wigner approxiamation mentioned in [30], we can transform the summation on the right-hand side of Eq.(17) and Eq.(18) to be expressed in terms of transition rates and get

$$\dot{c}_0 = -\frac{\Gamma_{21}}{2}c_0, \quad (29)$$

$$\dot{c}_1 = -ig_\omega c_0 e^{i(\omega_{21}-\omega_{s\sigma})t} - -\frac{\Gamma_{10}}{2}c_2, \quad (30)$$

$$\dot{c}_2 = g_{\omega'}c_2 e^{i(\omega_{10}-\omega_{s'\sigma'})t}, \quad (31)$$

where Γ_{21} and Γ_{10} are respectively

$$\Gamma_{21} = \frac{6\Gamma_{21}^{(0)}}{\xi_{21}^3}\sum_{m\mu}\frac{\chi_{s1}^2}{\alpha_{s1}\sqrt{\xi_{21}^2 - \chi_{s1}^2}}J_m^2(\chi_{s1}R/b) \\ \times \cos^2(\sqrt{\xi_{21}^2 - \chi_{s1}^2}z/L)\theta(\xi_{21} - \chi_{s1}), \quad (32)$$

$$\Gamma_{10} = \frac{6\Gamma_{10}^{(0)}}{\xi_{10}^3}\sum_{m\mu}\frac{\chi_{s1}^2}{\alpha_{s1}\sqrt{\xi_{10}^2 - \chi_{s1}^2}}J_m^2(\chi_{s1}R/b) \\ \times \cos^2(\sqrt{\xi_{10}^2 - \chi_{s1}^2}z/L)\theta(\xi_{10} - \chi_{s1}). \quad (33)$$

We are only interested in the system of $t \gg \Gamma_{21}^{-1}, \Gamma_{10}^{-1}$, in this case $c_0(\infty)$ and $c_1(\infty)$ both tend to 0, and $c_2(\infty)$ is

$$c_3(\infty) = \frac{g_\omega g_{\omega'}}{\left[i\left(\omega_{s'\sigma'} + \omega_{s\sigma} - \omega_{20}\right) - \frac{\Gamma_{21}}{2}\right]\left[i\left(\omega_{s'\sigma'} - \omega_{10}\right) - \frac{\Gamma_{10}}{2}\right]}. \tag{34}$$

Subsituting Eq.(34) into Eq.(16), we obtain the biphoton state

$$|\Omega\rangle = \sum_{s\sigma,s'\sigma'} \frac{-g_\omega g_{\omega'}\,|\omega_{s\sigma}, \omega_{s'\sigma'}\rangle}{\left[(\omega_{s'\sigma'} + \omega_{s\sigma} - \omega_{20}) + \frac{i\Gamma_{21}}{2}\right]\left[(\omega_{s'\sigma'} - \omega_{10}) + \frac{i\Gamma_{10}}{2}\right]}. \tag{35}$$

In order to obtain the specific biphoton wave function, we need to first obtain the transition rates between the second excited state and the first excited state and between the first excited state and the ground state. Then we use data from [32]. And subsequently, by using the relationship between N and the energy levels

$$\omega_{10} = \frac{E_1 - E_0}{\hbar} = \frac{w^2\hbar}{2m}\left(2N + 3\right) = 0.33\text{eV},$$

$$\omega_{20} = \frac{E_2 - E_0}{\hbar} = \frac{w^2\hbar}{m}\left(2N + 2\right) = 0.64\text{eV}, \tag{36}$$

$$\omega_{21} = \frac{E_2 - E_1}{\hbar} = \frac{w^2\hbar}{2m}\left(2N + 1\right) = 0.31\text{eV},$$

we can find $N = 19.6$, which in turn allows us to find the coordinate operator matrix elements by using Eq.(11). Then the dipole moments are $d_{10} = 0.16$ eÅ and $d_{21} = 0.23$ eÅ.

Experimental results show that the radius of axons in the brain or CNS is generally 2∼6 μm, and the thickness of myelin sheaths is generally 1∼3 μm[23, 24]. then let the radius of exposed axons (inner diameter of myelinated neurons) be 2 μm, and the thickness ranges from 0 to 3 μm. Assuming that the C-H bonding is widely distributed with the inner diameter a to the outer diameter b of axons, we calculate the average of Bessel functions and cos functions in Eq.(33) and Eq.(32)

$$\int_a^b J_m^2(\chi_{s1}R/b)\mathrm{d}R, \quad \int_0^L \cos^2(\frac{n\pi z}{L})\mathrm{d}z. \tag{37}$$

Then we can observe the variation of the transition rates with myelin thickness as shown in Figure.3(a). Each peak in the figure corresponds to a zero point of the Bessel function $\chi_{m\mu}$. When the cavity radius is less than 2.45 μm, Γ_{21} and Γ_{10} are quite small, which is due to the fact that the dipoles are all clustered at the axon boundaries $R/b \approx 1$, and so the coupling constant tends to 0. The coupling between the electromagnetic field and the vibrational modes is then negligible, and the vibrational modes change without any relationship between the excitation or non-excitation of the electromagnetic field.

The biphoton wave function can be calculated using the transition rates and the coupling constants. Considering that the linewidth is much smaller than the vibrational frequency, the allowed modes are almost all at resonance. So, for practical calculations, we will disregard frequencies greater than 0.39 eV and less than 0.26 eV. From the coefficient plots (Figure.3(b)) of the normalised biphoton wave functions it can be seen that the allowed modes are all very close to resonance or at resonance, and only a few modes are significantly allowed.

IV. EVALUATING ENTANGLEMENT THROUGH SCHMIDT DECOMPOSTION

In the previous section we obtained the wave function of biphoton system. We now evaluate the entanglement degree of this system. In quantum mechanics, the pure state of a bipartite system is described by a wave function $|\Omega(x_1, x_2)\rangle$, which x_1 and x_2 represent any degrees of freedom of the two particles: coordinates, momentum, polarization, or frequency, etc[33]. In the discrete case, the wave function can be written as Eq.(35). According to Schmidt analysis[34], it can always be expanded with two adjoint basis

$$|\Omega\rangle = \sum_n \sqrt{\lambda_n}\,|k_1\rangle_n\,|k_2\rangle_n, \tag{38}$$

where $|k_1\rangle_n$ and $|k_2\rangle_n$ are two adjoint Schmidt modes. Representation of Eq.(38) is known as Schmidt decomposition. The expansion parameters $\sqrt{\lambda_n}$ in Eq.(38) are real and positive, and they obey the normalization condition $\sum_n \lambda_n = 1$. This decomposition is closely related to the singular value decomposition (SVD)[35]. We can consider the coefficients in Eq.(35) as a matrix $\boldsymbol{C}$, and then this matrix can always do SVD to obtain two unitary matrices $\boldsymbol{U}$ and $\boldsymbol{\Sigma}$ and a diagonal matrix Λ. The relation between them is

$$\boldsymbol{C} = \boldsymbol{U}\boldsymbol{\Lambda}\boldsymbol{\Sigma}. \tag{39}$$

The squares of the diagonal elements of matrix Λ then correspond to the expansion parameters λ_n. The distinguishing feature of the Schmidt decomposition is that when a particle is measured to be in one of the Schmidt modes $|k_1\rangle_n$, then the other particle must be in the adjoint mode $|k_2\rangle_n$ rather than the other mode, and the
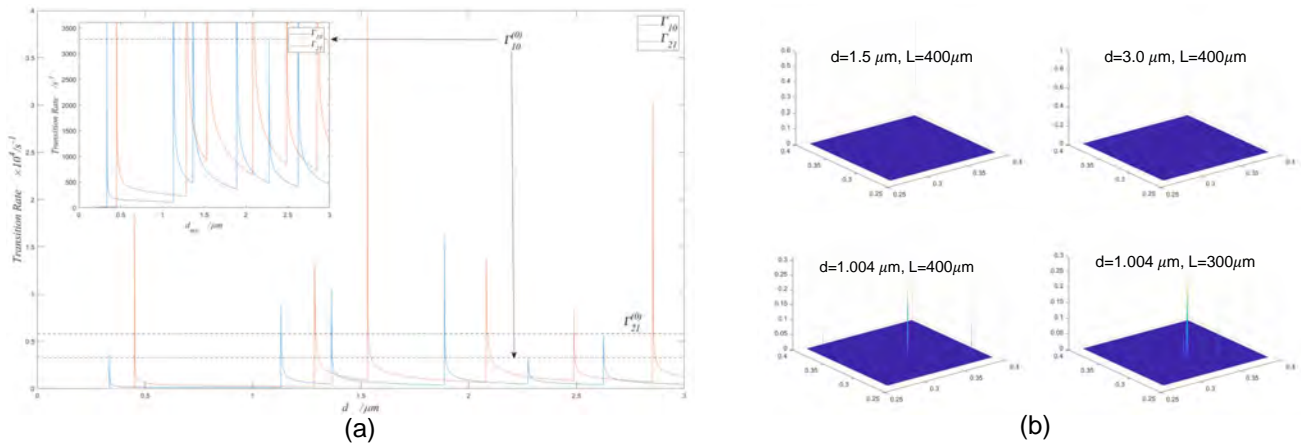
FIG. 3: (a) Transition rates Γ_{21} and Γ_{10}. Each peak correspond to ξ_{10} or ξ_{21} equals to $\chi_{m\mu}$. (b) Coefficients of biphoton wave functions at different sizes of cavity. X-axis and Y-axis are in unit of electron volts (eV). The highest peak always corresponds to the modes that close to ω_{10} and ω_{21}.

probability of finding such a pair of particles is given by λ_n. When two particles are independent of each other, they are not in entanglement. At this point, their wave functions can simply be written as

$$|\Omega\rangle = |k_1\rangle_n |k_2\rangle_n. \tag{40}$$

We get only one pair of Schmidt modes corresponding to the Schmidt decomposition of Eq.(38). When two particles are entangled, the Schmidt decomposition has more than one term. We can use the expansion parameters in Eq.(38) to define the von Neumann entropy to evaluate the manitude of entanglement, and the von Neumann entropy is defined as[34]

$$S = -\sum_n \lambda_n \log_2(\lambda_n). \tag{41}$$

When the Schmidt number has only one term, the entropy is 0, indicating that the system is not entangled. As defined, the maximum value of entropy is related to the number of states of the particle, and the maximum value of von Neumann entropy is $\log_2 n$ when the particle has n states.For normalisation purposes, the entropy involved in what follows is calculated in such a way that the base of the logarithmic function is taken to be the number of states n of the corresponding wave function.

The axon is usually wrapped by myelin sheath separated by several nodes of Raniver, each of which is usually $10\sim100$ μm[18, 23]. In the actual calculations, we neglect the total length of nodes of Raniver to be in the range of $200\sim500$ μm, which affects the denseness of electromagnetic modes in the microcavity. When the thickness of the myelin sheath is less than 0.45 μm, the transition rates Γ_{21} and Γ_{10} are extremely small due to the very small coupling constant. We consider that no biphoton state will be generated when the thickness of the myelin sheath is in the range of $0\sim0.45$ μm in the calculation. Electromagnetic modes at $(2,2,0)$ have energies exceed-

ing the eigen vibrational frequency ω_{10}, so modes exceeding χ_{221} are not considered here, and the results obtained are shown in Figure.4(a).

Owing to the discrete energy levels within the myelin cavity, instances where electromagnetic modes reside on both sides of the vibrational frequencies ω_{10} and ω_{21}—and are in proximity—lead to significant entanglement. This arises from photons being indistinguishable particles, implying that measuring the frequency of one photon imparts partial information about the other. Conversely, when a discrete electromagnetic mode precisely corresponds to ω_{10} and ω_{21}, it dominates, resulting in minimal entropy. The molecular coupling constant in Eq.(22) are about an order of magnitude of 10^7, and the transition rates (natural line width) is about an order of magnitude of 10^4, which is small and negligible compared to the frequencies of the photon and vibrational energy levels in the denominator, which are about 10^{14}. The discrete energy levels are solely determined by myelin geometry, making the creation and availability of entanglement entirely contingent on myelin size. In contrast to cascade radiation of the vibrational spectrum in free space, characterized by extremely narrow linewidths and continuous electromagnetic modes, photon cascade radiation within the myelin sheath has only one mode—photons corresponding to frequencies ω_{10} and ω_{21}.

V. DISCUSSIONS

As shown in Figure.4(b), the degree of entanglement is relatively high when the thickness of the myelin sheath is between $0.8\sim1.1$ μm. The radius of the axon is 2 μm, which corresponds to an inner/outer diameter ratio of $0.65\sim0.72$, which is close to the literature value of $0.6\sim0.8$[24, 36], and the entanglement decreases rapidly as the myelin thickness decreases beyond this ratio. The
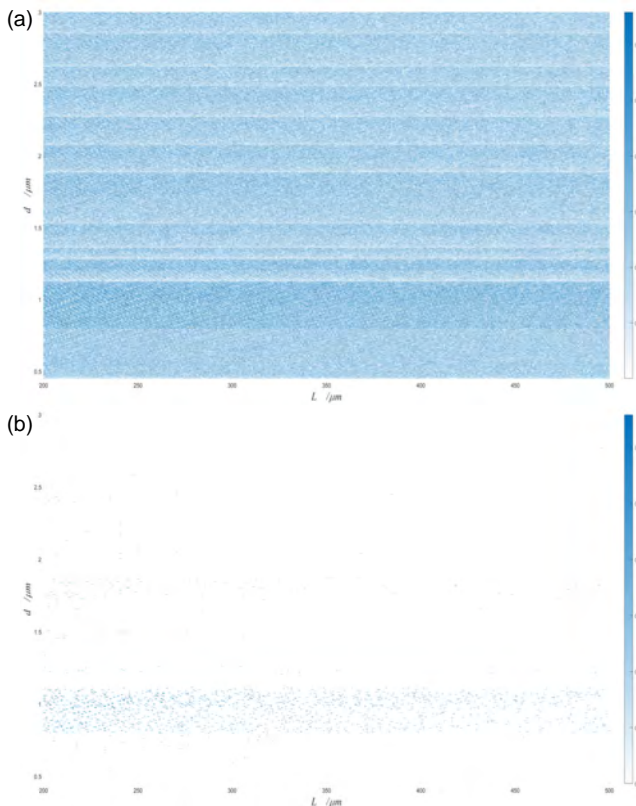
FIG. 4: von Neumann entropy at different sizes of the cavity. (a) Colors from white to dark blue represent entropy values from 0 to 0.65. Myelin thicknesses ranged from 0.45 to 3 μm, and lengths ranged from 200 to 500 μm. (b) The relationship between myelin thickness and the degree of entanglement can be better seen by erasing the points in (a) with entropy values below 0.5. Entanglement is substantial at thicknesses between 0.8 and 1.1 μm, and it decays rapidly as the thickness becomes smaller.

experimental results show that with age, myelin thickness becomes smaller and the probability of developing neurodegenerative diseases increases[37, 38], which is consistent with our calculations.

To summarize, the results of the cascade photons emission process by cQED and quantum optics indicate that biphotons in quantum entanglement can be released through cascade radiation on the vibrational spectrum of C-H bonds in the tails of lipid molecules inside cylindrical cavities encased by neural myelin. The presence of discrete electromagnetic modes due to the cavity structure formed by the myelin sheath, distinguishing it from the free-space continuous electromagnetic modes, results in the frequent production of highly entangled photon pairs permitted within the myelin cavity. Notably, due to the presence of microcavities, the coupling can be significantly enhanced compared to that in free space, indicating a higher probability of emitting photons. It should be noted that our model is very crude. The actual electromagnetic field should take into account the coupling of photons to the vibron ensembles, i.e. polaritons, which should be considered in future studies.

Finally, we would like to put the results into some speculations. It has been experimentally shown that the mid-infrared light energy at 53.53 THz has a modulating effect on the activity of K^+ ion channels, and the reason that its activity is not proportional to the number of photons[16] may be the influence of the superradiation of the vibrational system formed by eight C=O bonds. The photon energy released by one C-H bond vibration roughly corresponds to the vibrational energy of three C=O bonds, so that the vibrational collective spontaneous radiation of the C=O bonds is in a critical state about to reach its maximum value, and is very easy to be excited. Then, the entangled photon pairs emitted through the C-H bond vibrational spectrum cascade radiation might be able to link the K^+ ion channels at different locations together, affecting neural activity.

Since the myelin polariton (MP) contains a large number of vibronic states[17], the effect of thermal fluctuation on the states of the polariton is negligible. In recent years systems with similar systems such as cold-atom ensembles[39] were used as quantum memories, which can protect quantum entanglement between photons. The local entanglement generated in each neuron could be obtained in a larger region through entanglement swapping[6] and intercellular matter transfer, and the neurons in the brain may be corelated together through entanglement. It could be important for the synchronization of numerous in the brain. In this way, the resulting entangled photon pairs can propagate out in the neuromedullary sheaths thereby serving as a quantum communication resource in the nervous system.

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