Competing quantum spin liquids, gauge fluctuations, and anisotropic interactions in a breathing pyrochlore lattice

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We use the projective symmetry group analysis to classify the quantum spin liquids on the $S = 1/2$ pyrochlore magnet with a breathing anisotropy. We find 40 $Z_2$ spin liquids and 16 $U(1)$ spin liquids that respect the $F43m$ space group and the time reversal symmetry. As an application, we consider the antiferromagnetic Heisenberg model, which is proposed to be the dominant interaction in the candidate material $Ba_3Yb_2Zn_3O_{11}$. Focusing on the $U(1)$ spin liquid ansatze, we find that only two of them are physical when restricted to this model. We present an analytical solution to the partition mean field theory for each of these two $U(1)$ spin liquids. It is revealed that one of them has gapless, while the other one has gapped, spinon excitations. The two $U(1)$ spin liquids are equal in energy regardless of the degree of breathing anisotropy, and they can be differentiated by the low-temperature heat capacity contribution from the quadratically-dispersing gapless spinons. We further show that the latter is unaffected by fluctuations of the $U(1)$ gauge field within the random phase approximation. Finally, we demonstrate that a small Dzyaloshinskii-Moriya interaction lifts the degeneracy between the two $U(1)$ spin liquids, and it eventually causes the lattice to decouple into independent tetrahedra at strong coupling. While current model parameters for $Ba_3Yb_2Zn_3O_{11}$ place it indeed in the decoupled regime, other candidate materials may be synthesised in the near future that realize the spin liquid states discussed in our work.

I. INTRODUCTION

Pyrochlore magnets with antiferromagnetically coupled local moments are exemplary frustrated systems that provide fertile grounds for the exploration of spin liquid physics in three dimensions [1–4]. For instance, those with a strong Ising anisotropy realize classical [5–12] or quantum [13–17] spin ice, which displays emergent electromagnetism and magnetic monopoles. Further enriching the research into pyrochlore magnets is the abundance of synthetized materials, many of which possess spin interactions that deviate significantly from the spin ice Hamiltonian but nevertheless exhibit unusual behaviors [18–22].

In this work, we study a less symmetric variant of the pyrochlore lattice known as the breathing pyrochlore lattice [23–29], which breaks the inversion symmetry but retains all other symmetries of its regular counterpart. Pictorially, the up and down tetrahedra that share vertices with one another have different sizes. Breathing pyrochlore materials, such as spinel oxides [30–36] and $Ba_3Yb_2Zn_3O_{11}$ [37–42], with $S = 3/2$ and $S = 1/2$ local moments respectively, were first synthetized and investigated in the context of frustrated magnetism as early as 2012. Recently, there has been a revival of interest in breathing pyrochlore magnets due to proposals that they may stabilize spin liquid phases that are characterized by rank-2 $U(1)$ gauge fields (i.e., the emergent electromagnetic fields are tensors instead of vectors) and fractonic excitations [43–45], and that host an emergent axion field and a $\theta$-term coupled to the emergent QED [46].

Roughly coinciding with these works is the successful classification of symmetric $Z_2$ and $U(1)$ quantum spin liquids on the regular pyrochlore lattice [47–50], within the frameworks of bosonic and fermionic parton mean field theories. Such a symmetry-based classification is yet to be extended to the breathing pyrochlore lattice, a task which is taken up by this study. A better understanding of the lower-rank spin liquids is important and interesting in its own right, and, together with the aforementioned rank-2 spin liquids, they will provide a basis for future investigations into exotic phases of matter in breathing pyrochlore magnets.

Here, we classify the possible $Z_2$ and $U(1)$ quantum spin liquids in the $S = 1/2$ breathing pyrochlore magnet, via the projective symmetry group (PSG) analysis [51–57] based on the complex fermion mean field theory [58–61]. Both the spatial symmetries of the breathing pyrochlore lattice, i.e., the space group $F43m$, and the time reversal symmetry are enforced. This results in 40 $Z_2$ spin liquids and 16 $U(1)$ spin liquids. We then explain how these quantum spin liquids are related to those in the regular pyrochlore lattice previously classified by Ref. 48. In particular, we explicitly demonstrate that all of the 16 $U(1)$ spin liquids in the regular pyrochlore lattice are special cases of the 8 $U(1)$ spin liquids in the breathing pyrochlore lattice.

As an application of the PSG classification results, we consider the antiferromagnetic (AFM) Heisenberg model and look for physical $U(1)$ spin liquid ansatze that have nonzero bond parameters throughout the lattice. There are only two such ansatze out of 16, which we label $U(1)_0$ and $U(1)_\pi$, with trivial and projective realizations of translational symmetries respectively. Interestingly, their corresponding mean field theories admit analytical solutions, owing largely to the existence of flat bands [26, 27]. We find that, independently of the ratio between the interactions on the small and large tetrahedra that characterize the breathing anisotropy, the two spin liquids are exactly degenerate. However, the $U(1)_0$ state has gapless spinon excitations with a quadratic dispersion at low energies, while the $U(1)_\pi$ state has gapped spinon excitations, so they can in principle be distinguished by thermodynamic measurements [62].

For instance, the exponential (power law) dependence on temperature of the heat capacity may be used to infer the presence (absence) of an excitation gap. We further investigate how the coupling of the $U(1)$ gauge field to the gapless
spinons [63–75] may modify the low temperature heat capacity by means of an effective field theory. Within the random phase approximation and the small momentum limit, we find that the heat capacity of the gauge photon has the same scaling \( C(T) \sim T^{3/2} \) as that of bare spinons, suggesting that the effect of gauge fluctuations are not important.

The candidate spin model of the \( S = 1/2 \) breathing pyrochlore material \( \text{Ba}_3\text{Yb}_5\text{Zn}_3\text{O}_{11} \) (abbreviated as BYZO) consists of an AFM Heisenberg interaction and a Dzyaloshinskii-Moriya (DM) interaction roughly five times smaller in magnitude [39, 40]. For a more realistic model, we thus add a subleading DM interaction to the AFM Heisenberg interaction and study its effects on the two \( U(1) \) spin liquids. We find that a finite DM interaction lifts the degeneracy between them and favors the \( U(1)_0 \) state. When the DM interaction becomes sufficiently strong, it drives the system into isolated tetrahedra and neither spin liquids survive. In particular, neither the \( U(1)_0 \) state nor the \( U(1)_{\pi} \) state can be stabilized in the parameter space relevant for BYZO, where the interactions on the large tetrahedra are very weak (i.e., near the decoupled limit). This result corroborates the fact that single tetrahedron modelling is well suited to understand the physics of BYZO, as previously demonstrated by highly accurate fits to inelastic neutron scattering spectra [39–42]. We expect that the \( U(1)_0 \) and \( U(1)_{\pi} \) spin liquids will be relevant to spin models closer to the Heisenberg limit or with a less severe breathing anisotropy. Materials that satisfy these criteria may be synthesized in the near future, given the growing interests in breathing pyrochlore magnets.

The rest of the paper is organized as follows. In Sec. II, we discuss the symmetry of the breathing pyrochlore lattice and set up conventions for the coordinate system. In Sec. III, we briefly introduce the complex fermion mean field theory, before presenting the \( Z_2 \) and \( U(1) \) quantum spin liquids that result from the PSG analysis. In Sec. IV, we consider applications to the AFM Heisenberg model. Focusing on the \( U(1) \) spin liquids, we first argue that there are only two physical ansatze, \( U(1)_0 \) and \( U(1)_{\pi} \), due to constraints from the PSG (Sec. IV A). Then we present analytical solutions to the corresponding mean field theories (Sec. IV B). Taking into account the coupling of gapless spinons to the gauge field, we construct an effective field theory for the \( U(1)_0 \) state, from which we derive the low temperature heat capacity (Sec. IV C). After that, we study the effects of adding a Dzyaloshinskii-Moriya interaction (Sec. IV D). In Sec. V, we summarize our work, discuss its relations to existing theoretical studies, and outline potential future directions.

II. SYMMETRY

The pyrochlore lattice is a three dimensional network of corner sharing tetrahedra. By convention, each tetrahedron can be categorized as “up” or “down” according to its spatial orientation. These two species are related by an inversion symmetry about a site (i.e., a vertex of some tetrahedron). The inversion symmetry can be broken by introducing a breathing anisotropy, so that the up and down tetrahedra now have different sizes, see Fig. 1a. The resulting structure is called the breathing pyrochlore lattice, which belongs to the space group \( F43m \) (No. 216) [37, 39]. The underlying Bravais lattice is the face centered cubic (FCC) lattice, with the primitive translation vectors

\[
\mathbf{a}_1 = \frac{a}{2}(\hat{y} + \hat{z}), \quad \mathbf{a}_2 = \frac{a}{2}(\hat{z} + \hat{x}), \quad \mathbf{a}_3 = \frac{a}{2}(\hat{x} + \hat{y}),
\]

where \( a \) is the lattice constant that defines the FCC unit cell. We choose, for concreteness, the up (down) tetrahedra to be the smaller (larger) ones. The coordinates of a site on the breathing pyrochlore lattice can be expressed as

\[
\mathbf{r} = r_1 \mathbf{a}_1 + r_2 \mathbf{a}_2 + r_3 \mathbf{a}_3 + \mathbf{d}_s \equiv (r_1, r_2, r_3; s),
\]

where \( r_i \in \mathbb{Z} \), \( s \in \{0, 1, 2, 3\} \) indexes the four sites of a unit cell, and the sublattice coordinates \( \mathbf{d}_s \) are

\[
\begin{align*}
\mathbf{d}_0 &= \rho a(\hat{x} + \hat{y} + \hat{z})/8, \\
\mathbf{d}_1 &= \rho a(\hat{x} - \hat{y} - \hat{z})/8, \\
\mathbf{d}_2 &= \rho a(-\hat{x} + \hat{y} - \hat{z})/8, \\
\mathbf{d}_3 &= \rho a(-\hat{x} - \hat{y} + \hat{z})/8.
\end{align*}
\]

Here \( 0 < \rho < 1 \) parametrizes the breathing anisotropy, with smaller \( \rho \) giving a greater difference between the sizes of small and large tetrahedra. Note that with \( \rho = 1 \) in (3a)-(3d), we restore the inversion symmetry and thus recover the regular pyrochlore lattice.

The point group of \( F43m \) is the tetrahedral group \( T_d \) of 24 elements, which are best visualized by embedding a tetrahedron in a cube [76, 77], see Fig. 1b. These 24 elements can be generated by just two of them, namely \( C_3 \), a rotation by \( 2\pi/3 \) about the \([111]\) axis, and \( \sigma \), a reflection across the plane perpendicular to \([011]\) and containing the origin. For example, the fourfold rotoreflection consisting of a rotation by \( \pi/2 \) about the \([100]\) axis and then a reflection across the \( yz \) plane can be expressed as \( S_4 = C_2^3\sigma C_3^2 \). Below we list the

![Diagram of Breathing Pyrochlore Lattice](image-url)
action of the space group generators on a generic site with the coordinates \((r_1, r_2, r_3; s)\).

\[
\begin{align*}
T_1 : (r_1, r_2, r_3, s) &\rightarrow (r_1 + 1, r_2, r_3, s), \\
T_2 : (r_1, r_2, r_3, s) &\rightarrow (r_1, r_2 + 1, r_3, s), \\
T_3 : (r_1, r_2, r_3, s) &\rightarrow (r_1, r_2, r_3 + 1, s), \\
C_3 : (r_1, r_2, r_3, s) &\rightarrow (r_3, r_1, r_2, C_3(s)), \\
\sigma : (r_1, r_2, r_3, s) &\rightarrow (-r_1 - r_2 - r_3, r_2, r_3, \sigma(s)).
\end{align*}
\]

When \(s = 0, 1, 2, 3\), \(C_3(s) = 0, 2, 3, 1\) and \(\sigma(s) = 1, 0, 2, 3\).

Apart from the space group, we also consider time reversal symmetry, which is present in a model with only bilinear spin interactions (or, more generally, involving only products of an even number of dipolar operators). We denote the time reversal operator by \(T\).

The commutation relations between these symmetry generators constitute what are called the algebraic identities, which are listed in App. A. The algebraic identities will be crucial to the projective symmetry group (PSG) analysis discussed in the next section.

### III. COMPLEX FERMION MEAN FIELD THEORY

Complex fermion mean field theory [58–61] and its gauge structure have been discussed extensively in numerous references [51–57]. We only outline here some key steps that are instrumental to the methodology of this work.

We first represent spins in terms of complex fermions,

\[
S_i = \sum_{\alpha\beta} f_{i\alpha}^\dagger \frac{[\tilde{\gamma}]_{\alpha\beta}}{2} f_{i\beta}, \tag{5}
\]

so that the Hamiltonian, which is assumed to be bilinear in the spins, is quartic in the fermions. We then perform a mean field decoupling to obtain a Hamiltonian quadratic in the fermions. For concreteness, let us consider the antiferromagnetic Heisenberg model,

\[
H = \sum_{ij} J_{ij} S_i \cdot S_j, \quad J_{ij} > 0 \tag{6}
\]

from which we obtain

\[
H_{\text{MF}} = \sum_{ij} \frac{J_{ij}}{4} \left[ \chi_{ij} (f_{i1}^\dagger f_{j1} + f_{i1}^\dagger f_{j1}) + \text{h.c.} - |\chi_{ij}|^2 \right]
\]

\[
+ \Delta_{ij} (f_{i1}^\dagger f_{j1} - f_{i1} f_{j1}) + \text{h.c.} - |\Delta_{ij}|^2 \right]
\]

\[
+ \sum_l \left[ \lambda_l^{(3)} (n_l - 1) + (\lambda_l^{(1)} + i\lambda_l^{(2)}) f_{i1}^\dagger f_{i1} + \text{h.c.} \right], \tag{7}
\]

where \(n_l = f_{i1}^\dagger f_{i1} + f_{i1}^\dagger f_{i1}\) is the number operator, and \(\chi_{ij}\) and \(\Delta_{ij}\) are variational parameters of the singlet hopping and pairing channels respectively. On-site Lagrange multipliers \(\lambda_l^{(1,2,3)} \in \mathbb{R}\) are introduced to enforce the single occupancy constraint (i.e., one fermion per site), as the representation (5) has enlarged the original Hilbert space \(H = \otimes_i \{|\uparrow_i\rangle, |\downarrow_i\rangle\}\) by allowing zero or double occupancies, which are unphysical.

Anisotropic spin interactions, such as the Dzyaloshinskii-Moriya interaction that will be discussed in Sec. IV D, usually require triplet hopping and pairing channels [78–81] in the parton representation. Importantly, the projective symmetry group classification of quantum spin liquids (discussed in Sec. III A) does not depend on the particular spin model, but only on the symmetries of the system.

The parton representation (5) introduces an \(SU(2)\) gauge redundancy, in which the mean field Hamiltonian (7) is invariant under a symmetry \(X\) of the system only up to a gauge transformation \(G_X \in SU(2)\). In other words, the symmetries of the system are realized projectively at the mean field level. The \(2 \times 2\) matrix of variational parameters,

\[
u_{ij} = \frac{J_{ij}}{4} \left( \chi_{ij} - \Delta_{ij} \right), \tag{8}
\]

by symmetry and \(SU(2)\) gauge redundancy, obeys

\[
\nu_{X(i)X(j)} = G_X (X(i)) \nu_{ij} G_X^\dagger (X(j)), \tag{9}
\]

for any space group element \(X\).

The time reversal symmetry \(T\), being an antiunitary operator, requires some care. We can choose a gauge such that [51]

\[
u_{ij} = -G_T (i) \nu_{ij} G_T^\dagger (j). \tag{10}
\]

The \(2 \times 2\) matrix \(\nu_{ij}\) for the on-site terms is similarly defined, see (B5), and it also obeys (9) and (10). Further details of the complex fermion mean field theory can be found in App. B.

#### A. Projective symmetry group analysis

It was first proposed in Ref. 51 that, given a particular set of symmetries \(\{X\}\), the different possible sets of gauge transformations \(\{G_X\}\) provide a means to classify quantum spin liquids. Compound operators of the form \(G_X \in SU(2)\) constitute the so called projective symmetry group (PSG). This forms the basis of the PSG analysis [52–57].

The form of \(G_X \in SU(2)\) is not arbitrary but restricted by the symmetries of the system. To understand this, we introduce a special subgroup of PSG known as the invariant gauge group (IGG), which consists of pure gauge transformations that leave the mean field ansatz invariant. The algebraic identities (A1a)-(A1n), such as \(T_2^{-1} T_1^{-1} T_2 T_1 = e\), constrain \(G_X\) via, for example

\[
(G_T G_2)^{-1} (G_T G_1)^{-1} (G_T G_2) (G_T G_1) \in \text{IGG} \tag{11}
\]

This is because for expressions like the left hand side of (11), the net effect of the symmetry operators is an identity, so what remains must amount to a pure gauge transformation that leaves the mean field ansatz invariant (take \(X = e\) in (9), for instance). When both hopping and pairing terms are present in the Hamiltonian, the IGG is \(\{+1, -1\}\), and the resulting spin liquids are called \(\mathbb{Z}_2\) spin liquids. When only hopping terms are present in the Hamiltonian, the IGG is \(\{e^{i\theta}\} \mid 0 \leq \theta < 2\pi\},\) and the resulting spin liquids are called \(U(1)\) spin liquids.

We summarize below the results of the PSG classification of the \(\mathbb{Z}_2\) and \(U(1)\) spin liquids. Details of the calculations can be found in Apps. C and D.
TABLE I. In conjunction with (12a)-(12f) and (14a)-(14f), this table lists all the possible $\mathbb{Z}_2$ and $U(1)$ spin liquids. For all $\mathbb{Z}_2$ spin liquids, $g_C(1,2,3)=1$ and $g_T(s)=i\tau_2$. For all $U(1)$ spin liquids, $n_T=1$, $\varphi_C(s)=0$, $\varphi(0,1,2)=0$, and $\varphi(s)=0$. Below $q_0 \in \{0,1,2\}$ and $p_T, q_\mathbf{T}, p_\mathbf{T}, p_\mathbf{C}, q_\mathbf{C}, \in \{0,1\}$.

\[
\begin{array}{ccc|ccc|cc}
& g_C(0) & g_C(1,2) & g_T(3) & \varphi_C & \varphi & \varphi_T & \varphi_T \\
\hline
(\pm 1, \pm 1, -1, +1, 0) & 1 & 1 & & & & & \\
(\pm 1, \pm 1, -1, -1, +1) & 1 & i\tau_2 & & & & & \\
(\pm 1, \pm 1, -1, -1, -1, +1) & & & & & & & \\
\end{array}
\]

1. $\mathbb{Z}_2$ spin liquids

We find 40 fully symmetric $\mathbb{Z}_2$ spin liquids on the breathing pyrochlore lattice, which are distinguished by the gauge transformations

\[
\begin{align*}
G_T(r_1, r_2, r_3; s) &= 1, \\
G_C(r_1, r_2, r_3; s) &= \eta_{r_1 c}^{r_1 c}, \\
G_T(r_1, r_2, r_3; s) &= \eta_{r_1 r_2}^{r_1 r_2}, \\
G_C(r_1, r_2, r_3; s) &= \eta_{r_1 r_2}^{r_1 r_2} g_C(s), \\
G_T(r_1, r_2, r_3; s) &= \eta_{r_1 r_2}^{r_1 r_2} g_T(s), \\
G_T(r_1, r_2, r_3; s) &= g_T(s),
\end{align*}
\]

with $\eta_{\ldots}$ and $g_X(s)$ given in Table I.

2. $U(1)$ spin liquids

For $U(1)$ spin liquids, the gauge transformations have the specific form [48, 51, 54, 56]

\[
G_X(r_1, r_2, r_3; s) = (i\tau_3)^{n_X} e^{i\phi_X(r_1, r_2, r_3; s)\tau_3},
\]

\[
n_X \in \{0,1\}, \phi_X \in \{0,2\pi\}.
\]

We find 16 fully symmetric $U(1)$ spin liquids on the breathing pyrochlore lattice, which are distinguished by the gauge transformations

\[
\begin{align*}
\phi_T(r_1, r_2, r_3; s) &= 0, \quad n_T = 0, \\
\phi_T(r_1, r_2, r_3; s) &= r_1 T_T, \quad n_T = 0, \\
\phi_T(r_1, r_2, r_3; s) &= (r_2 - r_1) T_T, \quad n_T = 0, \\
\phi_C(r_1, r_2, r_3; s) &= r_1 T_T + \varphi_C(s), \quad n_C = 0, \\
\phi_C(r_1, r_2, r_3; s) &= r_1 T_T + \varphi_C(s), \quad n_C = 0, \\
\phi_C(r_1, r_2, r_3; s) &= \frac{r_2 (r_2 + 1)}{2} + (-1)^{\mathbf{T}_T} r_3 (r_3 + 1) \quad \text{and} \quad \phi_T(r_1, r_2, r_3; s) = \varphi_T(s).
\end{align*}
\]

We have found 16 fully symmetric $U(1)$ spin liquids in the breathing pyrochlore lattice. Intriguingly, the regular pyrochlore lattice has the same number of $U(1)$ spin liquids [48]. In this subsection, we would like to further investigate the relation between the $U(1)$ spin liquids of the regular and breathing pyrochlore lattices. Let A and B be two systems such that the symmetry group of B is a subgroup of the symmetry group of A. Generally, we expect the spin liquids in A to be included, as special cases, among the spin liquids in B [55]. In our case, the breathing pyrochlore lattice is obtained from the regular pyrochlore by breaking the inversion symmetry, so we can think of A (B) as the regular (breathing) pyrochlore lattice.

The PSG analysis for the regular pyrochlore lattice is done with the set of space group generators $\{T_1, T_2, T_3, C_\mathbf{6}, S\}$, where $C_\mathbf{6}$ is a sixfold rotoinversion and $S$ is a twofold nonsymmetric screw [48]. They can be used to construct the point group generators of the breathing pyrochlore lattice, via $C_3 = C_\mathbf{6}^{-1}$ and $\sigma = (C_\mathbf{6})^4 S C_\mathbf{6}^{-1}$. Therefore, to see the relation between the spin liquids in the breathing and regular pyrochlore lattices, we can compare the gauge transformation parts of $G_T, G_T, G_C, G_C, G_\sigma, \ldots$ for $G_\tau$ found in Sec. IIIA to those of $G_T, G_T, G_T, G_C, G_\sigma, \ldots$ for $G_\tau$ found in Sec. IIIA to those of $G_T, G_T, G_T, G_C, G_\sigma, \ldots$ for $G_\tau$ found in Ref. 48, respectively [55].

Leaving the detailed calculations to App. D 1, the final result is stated as follows. All 16 $U(1)$ spin liquids of the regular pyrochlore lattice are continuously connected to the 8 $U(1)$ spin liquids of the breathing pyrochlore lattice with $\theta_T = 0$, in the fashion of a two-to-one mapping. The other 8 $U(1)$ spin liquids in the breathing pyrochlore lattice with $\theta_T = \pi$ have no correspondence. A similar exercise can be carried out to clarify the relation between the $\mathbb{Z}_2$ spin liquids in the regular and breathing pyrochlore lattices, but we shall not report it here for simplicity.

IV. ANTIFERROMAGNETIC HEISENBERG MODEL

$\text{Ba}_3\text{Yb}_2\text{Zn}_6\text{O}_{11}$ (abbreviated as BYZO) is so far the only existing $S=1/2$ breathing pyrochlore magnet [37–42]. There are other breathing pyrochlore materials but with higher spins [30–36], and they do not fall into the classification scheme presented in this work. The proposed spin model for BYZO consists of a dominant antiferromagnetic (AFM) Heisenberg inter-
action and a subleading Dzyaloshinskii-Moriya interaction on isolated (small) tetrahedra [39, 40, 82]. The inter-tetrahedron coupling, i.e., the interactions on the large tetrahedra, are not well understood at the moment but expected to be insignificant, due to the rather large ratio $\sim 2$ between the bond lengths of the large and small tetrahedra.

As an application of the PSG classification results, we first study the pure AFM Heisenberg model on the breathing pyrochlore lattice, and then consider the effects of adding a Dzyaloshinskii-Moriya interaction. For simplicity, we consider only the fully symmetric $U(1)$ spin liquid candidates.

We denote the nearest neighbor Heisenberg interactions on the up (down) tetrahedron by $J_1$ ($J_2$). The mean field Hamiltonian of the AFM Heisenberg model is given by (7), which describes a $\mathbb{Z}_2$ spin liquid as both hopping and pairing terms are present (one can think of it as the spinon analog of a BCS superconductor). We set the pairing terms to be zero for a $U(1)$ spin liquid Hamiltonian (essentially a chargeless free fermion model). Notice that both $U(1)$ spin liquids with gapless and gapped spinons are allowed to exist in 3+1 dimensions, while in 2+1 dimensions $U(1)$ spin liquids with gapped spinons are unstable to monopole proliferation [83].

A. Physical spin liquid ansatze

As discussed in Sec. IIIA, the mean field ansatze $u_{ij}$ on different bonds are related by the (projective) symmetries of the system via (9), which reduces the number of independent $\chi_{ij}$ and/or $\Delta_{ij}$. On the other hand, a symmetry that maps a bond onto itself, e.g., the time reversal symmetry via (10), may impose additional constraints on the bond. If the symmetry, or the conflicting requirements arising from different symmetries, forces $u_{ij}$ = 0 everywhere, then we get a zero Hamiltonian, which is unphysical and thus discarded [52, 54]. We also exclude the case of $u_{ij}$ ≠ 0 on one species of the tetrahedra (say up) and $u_{ij}$ = 0 on the other (say down). This describes a system of decoupled tetrahedra, which reduces to a 4-spin problem. While the decoupled tetrahedron model is interesting, it does not require complex fermion mean field theory, and we do not consider it explicitly in this work where we focus on spin liquid states with full spatial connectivity.

Therefore, having specified the spin model and thus the mean field parameters, we may find that some of the PSGs in Sec. IIIA are unphysical or irrelevant. As an application of the PSG classification results, we first study the pure AFM Heisenberg model on the breathing pyrochlore lattice, and then consider the effects of adding a Dzyaloshinskii-Moriya interaction. For simplicity, we consider only the fully symmetric $U(1)$ spin liquid candidates.

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A. Physical spin liquid ansatze

As discussed in Sec. IIIA, the mean field ansatze $u_{ij}$ on different bonds are related by the (projective) symmetries of the system via (9), which reduces the number of independent $\chi_{ij}$ and/or $\Delta_{ij}$. On the other hand, a symmetry that maps a bond onto itself, e.g., the time reversal symmetry via (10), may impose additional constraints on the bond. If the symmetry, or the conflicting requirements arising from different symmetries, forces $u_{ij}$ = 0 everywhere, then we get a zero Hamiltonian, which is unphysical and thus discarded [52, 54]. We also exclude the case of $u_{ij}$ ≠ 0 on one species of the tetrahedra (say up) and $u_{ij}$ = 0 on the other (say down). This describes a system of decoupled tetrahedra, which reduces to a 4-spin problem. While the decoupled tetrahedron model is interesting, it does not require complex fermion mean field theory, and we do not consider it explicitly in this work where we focus on spin liquid states with full spatial connectivity.

Therefore, having specified the spin model and thus the mean field parameters, we may find that some of the PSGs in Sec. IIIA are unphysical or irrelevant. As an application of the PSG classification results, we first study the pure AFM Heisenberg model on the breathing pyrochlore lattice, and then consider the effects of adding a Dzyaloshinskii-Moriya interaction. For simplicity, we consider only the fully symmetric $U(1)$ spin liquid candidates.

We denote the nearest neighbor Heisenberg interactions on the up (down) tetrahedron by $J_1$ ($J_2$). The mean field Hamiltonian of the AFM Heisenberg model is given by (7), which describes a $\mathbb{Z}_2$ spin liquid as both hopping and pairing terms are present (one can think of it as the spinon analog of a BCS superconductor). We set the pairing terms to be zero for a $U(1)$ spin liquid Hamiltonian (essentially a chargeless free fermion model). Notice that both $U(1)$ spin liquids with gapless and gapped spinons are allowed to exist in 3+1 dimensions, while in 2+1 dimensions $U(1)$ spin liquids with gapped spinons are unstable to monopole proliferation [83].
FIG. 2. The mean field ansatze of the two possible \( U(1) \) spin liquids, the (a) \( U(1)_0 \) and (b) \( U(1)_\pm \) states, in the antiferromagnetic Heisenberg model. They contain \( N = 4 \) and \( N = 16 \) sites per unit cell respectively, which are labelled by \( s = 0, \ldots, N - 1 \). Green (red) bonds on the down tetrahedra indicate \( \chi_{ij} = +\chi_2 (\chi_{ij} = -\chi_2) \). All bonds on the up tetrahedra have \( \chi_{ij} = +\chi_1 \) and they are colored in green. In (b), the unit cell is enlarged in the \( a_3 \) and \( a_1 \) directions, due to the projective realization of the translational symmetry.

Since the Hamiltonian contains only singlet hopping channels, fermions with up and down spins do not mix, and the \( 2N \times 2N \) matrix \( D_k \) is block diagonal, consisting of two identical copies of the \( N \times N \) matrix \( d_k \). The energy eigenvalues of \( D_k \) can thus be obtained by diagonalizing \( d_k \) and imposing a twofold degeneracy. The single occupancy constraint in real space, in which each site is occupied by a fermion, is translated to a half filling constraint in momentum space, in which the lower (upper) half of the energy eigenstates are occupied (empty) in the ground state. Note that if we impose the half filling constraint by hand, then we no longer have to explicitly introduce the Lagrange multiplier \( \lambda_1^{(5)} \).

Furthermore, we note that the global gauge transformation \( \bar{i} \tau_1 \) flips the signs of both \( \chi_1 \) and \( \chi_2 \) (which can be thought of as a particle hole transformation), while leaving the total energy invariant (as it should). We thus assume \( \chi_1 < 0 \) without loss of generality. While the overall sign is not important, we will see later that the relative sign between \( \chi_1 \) and \( \chi_2 \) does make a difference. Finally, we define \( \tilde{\chi}_1 = -J_1\chi_1/4 > 0 \) and \( \tilde{\chi}_2 = -J_2\chi_2/4 \) for convenience. We also require both \( \chi_1 \) and \( \chi_2 \) to be nonzero for the physical solutions that we are interested in (see the discussion in Sec. IV A).

For the \( U(1)_0 \) state, diagonalizing \( d_k \) yields the four eigenvalues \( \epsilon_0, \epsilon_0, \epsilon_+, \epsilon_- \), where

\[
\epsilon_0(k) = -\tilde{\chi}_1 - \tilde{\chi}_2, \\
\epsilon_\pm(k) = \tilde{\chi}_1 + \tilde{\chi}_2 \pm \sqrt{4\tilde{\chi}_1^2 + 4\tilde{\chi}_2^2 - 2\tilde{\chi}_1\tilde{\chi}_2[2 - f(k)]},
\]

with

\[
f(k) = \cos(k_1 - k_2) + \cos(k_2 - k_3) + \cos(k_3 - k_1) \\
+ \cos k_1 + \cos k_2 + \cos k_3, \quad k_i \in [0, 2\pi).
\]

The maximum and minimum of \( f(k) \) are 6 and \(-2\) respectively, an information that will be useful later. The same dispersions (23a) and (23b) have been obtained in different contexts, namely as eigenvalues of (i) the interaction matrix in the classical analysis of the breathing pyrochlore Heisenberg model [23] and (ii) the tight binding model on the breathing pyrochlore lattice with real hopping integrals [27].

We first assume that \( \chi_2 > 0 \), i.e., \( \chi_1 \) and \( \chi_2 \) have the same sign. We see that \( \epsilon_+(k) > \epsilon_0(k) \) for all \( k \), so the entire + band is unoccupied. On the other hand,

\[
\min_k \epsilon_-(k) = \tilde{\chi}_1 + \tilde{\chi}_2 - \sqrt{4\tilde{\chi}_1^2 + 4\tilde{\chi}_2^2 + 8\tilde{\chi}_1\tilde{\chi}_2}, \\
= -\tilde{\chi}_1 - \tilde{\chi}_2,
\]

which implies \( \epsilon_0(k) \leq \epsilon_-(k) \) for all \( k \) (equality holds only when \( k = 0 \)). The \( \pm \) bands are thus well separated from, and higher in energy than, the two 0 bands throughout the Brillouin zone except at \( k = 0 \). Then, filling the lower half of the energy eigenstates, i.e., all the flat bands, the total energy of a system with \( N \times N \) sites is given by

\[
E_S = \sum_k \left[ 4\epsilon_0(k) + 6\left( \frac{J_1}{4} \tilde{\chi}_1^2 + \frac{J_2}{4} \tilde{\chi}_2^2 \right) \right] \\
= \sum_k \left[ 4 \left( \frac{J_1}{4} \chi_1 + \frac{J_2}{4} \chi_2 \right) + 6 \left( \frac{J_1}{4} \chi_1^2 + \frac{J_2}{4} \chi_2^2 \right) \right],
\]

(25)

where the factor of 4 instead of 2 in front of \( \epsilon_0(k) \) takes into account the two spin flavors (recall that we have two copies of \( d_k \)).

Next, we show that if \( \tilde{\chi}_2 < 0 \), i.e., \( \chi_1 \) and \( \chi_2 \) have opposite signs, then the total energy is always higher than the previously analyzed case of \( \tilde{\chi}_2 > 0 \). Without loss of generality, we may assume \( |\tilde{\chi}_1| \geq |\tilde{\chi}_2| \), for if it is otherwise, we can perform the global gauge transformation \( \bar{i} \tau_1 \) and interchange \( \tilde{\chi}_1 \) and \( \tilde{\chi}_2 \). For clarity, we rewrite the eigenvalues (23a) and (23b) as

\[
\epsilon_0(k) = -|\tilde{\chi}_1| + |\tilde{\chi}_2|, \\
\epsilon_\pm(k) = |\tilde{\chi}_1| - |\tilde{\chi}_2| \pm \sqrt{4|\tilde{\chi}_1|^2 + 4|\tilde{\chi}_2|^2 + 2|\tilde{\chi}_1||\tilde{\chi}_2|[2 - f(k)]},
\]

One can again see that \( \epsilon_+(k) > \epsilon_0(k) \) for all \( k \), so the entire + band is unoccupied. On the other hand,

\[
\max_k \epsilon_-(k) = |\tilde{\chi}_1| - |\tilde{\chi}_2| - \sqrt{4|\tilde{\chi}_1|^2 + 4|\tilde{\chi}_2|^2 + 8|\tilde{\chi}_1||\tilde{\chi}_2|} \\
= -|\tilde{\chi}_1| + |\tilde{\chi}_2|, \\
\min_k \epsilon_-(k) = |\tilde{\chi}_1| - |\tilde{\chi}_2| - \sqrt{4|\tilde{\chi}_1|^2 + 4|\tilde{\chi}_2|^2 + 8|\tilde{\chi}_1||\tilde{\chi}_2|} \\
= -|\tilde{\chi}_1| - 3|\tilde{\chi}_2|,
\]

which implies \( \epsilon_0 \geq \epsilon_-(k) \) for all \( k \) (equality holds only when \( k = 0 \)). The \( \pm \) bands are thus well separated from the two 0 zero bands, with the + (−) band lying above (below) the flat bands. Therefore, filling the lower half of the energy eigenstates corresponds to filling the − band and one of the 0 bands. The total energy for a system of \( N \times N \) sites is given...
dispersing bands are (un)occupied.

bands (4 sublattices \( \times 2 \) spins). At half filling, all flat (dispersing) bands are (un)occupied. (d-f) The spinon dispersion of the \( U(1)_0 \) spin liquid, at (d) \( J_2/J_1 = 0.8 \), (e) \( J_2/J_1 = 0.5 \), and (f) \( J_2/J_1 = 0.2 \). The flat band (indicated by darker color) is 16-fold degenerate, while each of the four dispersing bands (indicated by lighter color) is 4-fold degenerate, totalling 32 bands (16 sublattices \( \times 2 \) spins). At half filling, all flat (dispersing) bands are (un)occupied.

by

\[
E_A = \sum_k \left[ 2\varepsilon_-(k) + 2\varepsilon_0(k) + 6 \left( \frac{J_1}{4} \vert \chi_1 \vert^2 + \frac{J_2}{4} \vert \chi_2 \vert^2 \right) \right] \\
> \sum_k \left[ 2 \min_k \varepsilon_-(k) + 2 \left( -\frac{J_1}{4} \vert \chi_1 \vert + \frac{J_2}{4} \vert \chi_2 \vert \right) \right] \\
+ 6 \left( \frac{J_1}{4} \vert \chi_1 \vert^2 + \frac{J_2}{4} \vert \chi_2 \vert^2 \right) \\
= \sum_k \left[ -4 \left( \frac{J_1}{4} \vert \chi_1 \vert + \frac{J_2}{4} \vert \chi_2 \vert \right) + 6 \left( \frac{J_1}{4} \vert \chi_1 \vert^2 + \frac{J_2}{4} \vert \chi_2 \vert^2 \right) \right] \\
= E_S.
\]

The inequality in the second line is strict because there is at least one point, \( k = 0 \), that does not saturate the lower bound of \( \varepsilon_-(k) \). We have thus shown that \( E_S < E_A \), i.e., the total energy for \( \tilde{\chi}_2 < 0 \) is always higher than that for \( \tilde{\chi}_2 > 0 \), given that \( \tilde{\chi}_1 > 0 \). We can therefore exclude the case where \( \chi_1 \) and \( \chi_2 \) have opposite signs.

Finally, minimizing (25) with respect to \( \chi_1 \) and \( \chi_2 \), \( \partial E_S/\partial \chi_{1,2} = 0 \), yields \( \chi_{1,2} = -1/3 \), regardless of the values of \( J_1 \) and \( J_2 \) as long as they are positive. The ground state energy per site is thus \(- (J_1 + J_2)/24 \), as in the \( U(1)_0 \) state. We have thus shown the robust “1/3 quantization” of the mean field parameters in the ground states of the two \( U(1) \) spin liquids, and established their degeneracy, independent of the ratio \( J_2/J_1 \) as long as both \( J_1 \) and \( J_2 \) are positive.

The spinon spectra of the \( U(1)_0, \pi \) spin liquids at several values of \( J_2/J_1 \) along some high symmetry cuts in the Brillouin zone [84] are plotted in Figs. 3a to 3f. In the \( U(1)_0 \) spin liquid, the spinon gap closes at the \( \Gamma \) point, while in the \( U(1)_\pi \) spin liquid the spinons are always gapped; these hold throughout the range \( J_2/J_1 \in (0, 1) \)

We remark that, in the isotropic limit \( J_2/J_1 = 1 \), our \( U(1)_0 \) and \( U(1)_\pi \) states respectively reduce to the uniform and \( (\pi, \pi) \) states of Ref. 85, where they are also found numerically to be degenerate. Note that our energy is smaller by an overall factor of 4 than that in Ref. 85 due to different schemes of the mean field decoupling.

\[ C. \text{ Low Temperature Heat Capacity} \]

Although the \( U(1)_0 \) and \( U(1)_\pi \) spin liquids are degenerate, there is a qualitative distinction between their spinon spectra, namely the presence or absence of an excitation gap. This allows them to be differentiated by specific heat measurements, for example. Considering the contribution from the spinons
alone, i.e., ignoring photons and visons, the heat capacity \( C(T) \) is expected to show a power law (exponential) dependence on temperature \( T \) for the gapless (gapped) case at low temperatures.

The spinon excitation spectrum of the \( U(1) \) spin liquid becomes gapless at the \( \Gamma \) point \( (\mathbf{k} = 0) \), where the flat bands \( \epsilon_0(x) \) touch the dispersing bands \( \epsilon_- (\mathbf{k}) \), as shown in Sec. IV B. Assuming \( \chi_{1,2} \equiv -J_{1,2} \chi_{1,2} / 4 > 0 \), so that the flat (dispersing) bands are filled (empty), we perform a small \( \mathbf{k} \) expansion in the vicinity of the \( \Gamma \) point to the lowest nontrivial order,

\[
\epsilon_- (\mathbf{k} \approx 0) \approx \tilde{x}_1 + \tilde{x}_2 - \sqrt{4(\tilde{x}_1 \tilde{x}_2)^2 - \tilde{x}_1 \tilde{x}_2 |\mathbf{k}|^2} \\
\approx -\tilde{x}_1 + \tilde{x}_2 + \frac{\tilde{x}_1 \tilde{x}_2}{4(\tilde{x}_1 \tilde{x}_2)} |\mathbf{k}|^2, \tag{28}
\]

where \( |\mathbf{k}|^2 = k_x^2 + k_y^2 + k_z^2 \) and we have used

\[
\begin{pmatrix}
k_1 \\
k_2 \\
k_3
\end{pmatrix} = \frac{1}{2} \begin{pmatrix}
0 & 1 & 0 \\
1 & 0 & 1 \\
0 & 1 & 0
\end{pmatrix} \begin{pmatrix}
k_x \\
k_y \\
k_z
\end{pmatrix}. \tag{29}
\]

Therefore, at low energies the spinon excitations follow a quadratic dispersion, which gives rise to a heat capacity \( C(T) \sim T^{3/2} \) at low temperatures if we neglect the effects of the \( U(1) \) gauge field.

However, gauge fluctuations may be important in a spin liquid with gapless spinons and lead to a singular correction to the heat capacity coefficient \( C(T)/T \) for small \( T \). For instance, it has been established that, if spinons in a \( U(1) \) spin liquid form a sufficiently large Fermi surface, when their coupling to the gauge field is taken into account, \( C(T) \) scales as \( T^{2/3} \) and \( T \ln(1/T) \) in 2D \([71, 72, 75]\) and 3D \([70, 86, 87]\) respectively, in contrast to \( T \) for bare spinons. Here, we do not have a Fermi surface, but a flat-quadratic band touching. To understand how gauge fluctuations may modify \( C(T) \) in the \( U(1) \) spin liquid, we construct an effective field theory that includes the interaction between the low energy spinons and the gauge field via minimal coupling. Going to the continuum limit, the Lagrangian and the partition function are \([74, 75]\)

\[
\mathcal{L} = \overline{\psi}_\sigma (i \gamma^\mu \partial_\mu - i a_0) \psi_\sigma + \frac{1}{2m} \overline{\psi}_\sigma (i \gamma^\mu \partial_\mu - a^2) \psi_\sigma, \tag{30a}
\]

\[
Z = \int \mathcal{D}a \mathcal{D}\psi_\sigma \mathcal{D}\overline{\psi}_\sigma e^{-\int \mathcal{D}r \mathcal{D}x \mathcal{L}}, \tag{30b}
\]

where \( \tau \) is the imaginary time, \( \psi_\sigma \) and \( \overline{\psi}_\sigma \) are the spinon fields with \( \sigma \) being the spin flavor, \( a_0 \) and \( a \) are the temporal and spatial components of the \( U(1) \) gauge field respectively, and \( a \equiv (a_0, \mathbf{a}) \). The effective mass \( m \) of the spinons is determined by the coupling constants \( J_{1,2} \) and by the mean field parameters \( \chi_{1,2} \) through (28). Note the absence of the Maxwell term

\[
\mathcal{L}_g = \frac{1}{8} f_{\mu\nu} f^{\mu\nu}, \quad f_{\mu\nu} = \partial_\mu a_\nu - \partial_\nu a_\mu, \tag{31}
\]

in (30a) as \( g \) is proportional to the charge gap and we are working in the insulating phase \([74, 75]\). Furthermore, since the flat bands are momentum independent, they contribute a constant (taken to be zero here) to the total energy and act as a reservoir of spinons that can be thermally excited to the quadratic spectrum. Note that the fields are functions of spacetime, for example \( \psi_\sigma \) in (30a) should be understood as \( \psi_\sigma (x) \).

Integrating out the spinons in (30b) and using the random phase approximation \([73–75, 88]\), we obtain an effective Lagrangian for the \( U(1) \) gauge field,

\[
\mathcal{L}_{\text{eff}} = - \sum_{i,j \in \{x,y\}} a_i (q) \Pi_{ij} (q) a_j (q), \tag{32a}
\]

\[
\Pi_{ij} (q; l = 0) = - c_1 \sqrt{T} \delta_{ij}, \tag{32b}
\]

\[
\Pi_{ij} (q; l \neq 0) = \delta_{ij} \left( -c_2 T^{3/2} + c_3 T^{5/2} \frac{|q|^2}{v_f^4} \right), \tag{32c}
\]

in the small \( |q| \) limit. Some remarks are in order. The first one is about conventions and notations. We have chosen the Coulomb gauge \( \nabla \cdot a = 0 \) and labeled the two transverse components of \( a \) by \( \chi \) and \( \gamma \). The inverse photon propagator is identified as \( \Pi_{ij} (q) \), where \( q = (q_x, q_y) \) is the four-momentum, \( v_f = 2\pi l T \) is the Matsubara frequency, and \( \mathbf{q} \) is the momentum of the photon. The coefficients \( c_i \) are positive and depend only on \( m \). The second one is that, unlike the case of a fermion Fermi surface \([70, 74, 88]\), we do not have a Fermi wavevector \( k_F \) relative to which the smallness of \( |q| \) can be defined. The natural dimensionless parameter that enables a small \( |q| \) expansion in our case is \( |q|/\sqrt{m T} \ll 1 \). Finally, \( \Pi_{ij} (q) \) assumes one of the two forms, either (32b) or (32c), according to whether \( l \) is zero or finite. Details of the calculations as well as extended discussions can be found in App. F.

In a \( U(1) \) spin liquid with a spinon Fermi surface, one finds \( \Pi_{ij} (q) = \delta_{ij} (\Gamma q_l / |q| - \chi |q|^2) \), with positive constants \( \Gamma \) and \( \chi \) that are determined by the details of the spinon dispersion \([70]\). This is in contrast with our \( \Pi_{ij} \) (32b) and (32c), where the coefficients \( c_i \) are multiplied by powers of \( T \) so that they vanish at \( T = 0 \), as expected because there is no thermally excited spinon in the quadratic spectrum at absolute zero.

From (32a) we can calculate the partition function and the free energy as \([70, 74]\)

\[
Z = \int \mathcal{D}a e^{-\frac{\sum \mathcal{L}_{\text{eff}}}} , \tag{33a}
\]

\[
F = - T \ln Z, \tag{33b}
\]

which yield, for the “static” \( (l = 0) \) and the “dynamic” \( (l \neq 0) \) contributions respectively,

\[
F_{\text{sta}} = T \sum_q \ln \left( c_1 \sqrt{T} |q|^2 \right), \tag{34a}
\]

\[
F_{\text{dyn}} = T \sum_{l \neq 0} \ln \left( c_2 T^{3/2} - c_3 T^{5/2} \frac{|q|^2}{v_f^4} \right). \tag{34b}
\]

To evaluate (34a) and (34b), we change the summation over \( \mathbf{q} \) to an integral, which has an upper limit of \( \sqrt{m T} \) in line
with the small \( q \) assumption, and regularize the summation over nonzero \( l \) with the Riemann zeta function. Details of the calculation are presented in App. F.

The heat capacity is then given by \( C(T) = -T(\partial^2 F/\partial T^2) \). In the low temperature limit, we find \( C(T) \sim T^{3/2} \) to leading order. We thus conclude that \( U(1) \) gauge fluctuations do not modify the scaling of the bare spinon heat capacity with temperature, unlike the case of a Fermi surface.

### D. Effects of Dzyaloshinskii-Moriya interaction

For the pure AFM Heisenberg model, we have seen in Sec. IVB that regardless of the strength of breathing anisotropy, we are always able to obtain spatially connected solutions from the mean field theories with the \( U(1)_0 \) and \( U(1)_\pi \) ansatze (even if we take \( J_2/J_1 \rightarrow 0^+ \), as long as \( J_2 \) is not strictly zero). Moreover, these two spin liquids are degenerate throughout the range \( J_2/J_1 \in (0, 1) \). In this subsection, we investigate how the stability and degeneracy of the \( U(1)_0 \) and \( U(1)_\pi \) spin liquids are affected by a finite Dzyaloshinskii-Moriya interaction.

The spin Hamiltonian is now given by

\[
H = \sum_{n=1}^{2} \sum_{\gamma} \left( J_n S_i \cdot S_j + D_n \mathbf{d}_i \cdot \mathbf{d}_j \right), \tag{35}
\]

where, as before, the subscripts 1 or 2 on the interactions indicate whether they belong to the up or down tetrahedra, and \( \mathbf{d}_i = \mathbf{d}_{\gamma i} \) are unit vectors that depend only on the sublattice indices \( s \) and \( t \) of sites \( i \) and \( j \) [39, 43],

\[
\mathbf{d}_{01} = \frac{-\hat{y} + \hat{z}}{\sqrt{2}}, \quad \mathbf{d}_{02} = \frac{-\hat{x} + \hat{z}}{\sqrt{2}}, \quad \mathbf{d}_{03} = \frac{-\hat{x} + \hat{y}}{\sqrt{2}},
\]

\[
\mathbf{d}_{12} = \frac{-\hat{x} - \hat{y}}{\sqrt{2}}, \quad \mathbf{d}_{23} = \frac{-\hat{x} - \hat{z}}{\sqrt{2}}, \quad \mathbf{d}_{31} = \frac{-\hat{z} - \hat{y}}{\sqrt{2}}. \tag{36}
\]

In line with the proposed spin models of BYZO in Refs. 39 and 40, we choose \( D_1 < 0 \). We further assume \( D_2 = (J_2/J_1) D_1 \). Using the parton representation of spins (5), the Dzyaloshinskii-Moriya interaction involves both the singlet and triplet channels (see App. G). If we require the singlet channels associated with the dominant AFM Heisenberg interaction to be nonvanishing, we still have \( U(1)_0 \) and \( U(1)_\pi \) as the only two \( U(1) \) spin liquid ansatze to be considered upon adding the Dzyaloshinskii-Moriya interaction.

For either ansatz, the mean field theory of the model (35) contains only two additional variational parameters, \( E_1^\gamma \) and \( E_2^\gamma \), on top of \( \chi_1 \) and \( \chi_2 \) that are already present in the Heisenberg limit (see App. G). The theory is no longer analytically tractable, so we solve it computationally by iterating the self-consistency equations, e.g., \( \chi_{ij} = \sum_{\gamma} (f_{\alpha i} f_{\gamma j}) \) (which is done in momentum space, see Ref. 80 for example), until the mean field parameters converge. We study the parameter region defined by \( J_2/J_1 \in (0, 1) \) and \( D_1/\sqrt{2} J_1 \in [-0.5, 0] \).

Our result, which is plotted in Fig. 4a, has two main features. First, in the presence of a sufficiently small Dzyaloshinskii-Moriya interaction, the degeneracy between the \( U(1)_0 \) and \( U(1)_\pi \) spin liquids is lifted, such that the \( U(1)_0 \) spin liquid is lower in energy. The Dzyaloshinskii-Moriya interaction also gaps out the spinon spectrum of the \( U(1)_0 \) spin liquid. In particular, the highest occupied band and the lowest unoccupied band no longer touch at the \( \Gamma \) point, see Fig. 4b. Second, as the Dzyaloshinskii-Moriya interaction increases in magnitude, there exists a critical value \( |D_1|_c \) above which the mean field theory favors a solution of isolated tetrahedra, instead of a spatially connected \( U(1) \) spin liquid. The red star indicates the parameters of the candidate spin model for BYZO [39, 40], where \( |D_1|/J_1 \approx 0.2 \) \& \( J_2/J_1 \approx 0.28 \). Filled (empty) bands are indicated by darker (lighter) blue lines.

### V. DISCUSSION

In summary, we have investigated the possible quantum spin liquids in the \( S = 1/2 \) breathing pyrochlore magnet using the complex fermion mean field theory and the projective symmetry group (PSG) analysis. We identify 40 \( Z_2 \) spin liquids and 16 \( U(1) \) spin liquids that are subjected to the \( F_43m \) space group of the breathing pyrochlore lattice as well as the time reversal symmetry. As an application, we consider the antiferromagnetic (AFM) Heisenberg model, and identify the physical \( U(1) \) spin liquid ansatze. Most of the 16 \( U(1) \) states are constrained by PSG to have vanishing bond parameters, leaving only two cases, which we label \( U(1)_0 \) and \( U(1)_\pi \). Their corresponding mean field theories admit analytical solutions, as shown in the text. We find that the \( U(1)_0 \) state has gapless spinon excitations, while the \( U(1)_\pi \) state exhibits a spinon gap.
these two states are however degenerate regardless of the ratio $J_2/J_1$ between the interactions on the large and small tetrahedra. While the spinon contribution to the heat capacity of the $U(1)_{_{S^+}}$ state is exponentially suppressed at low temperatures in the presence of an excitation gap, the quadratically dispersing spinons in the $U(1)_{_{S^T}}$ state give rise to a heat capacity contribution $C(T) \sim T^{3/2}$ at low temperatures. Using an effective field theory, we demonstrate that such power law dependence is unchanged by small momentum gauge fluctuations within the random phase approximation.

The degeneracy between the $U(1)_{_{S^T}}$ and $U(1)_{_{S^+}}$ states is lifted by a finite Dzyaloshinskii-Moriya interaction, which favors the $U(1)_{_{S^T}}$ state and gaps out its spectrum. It is also found that when the DM interaction becomes sufficiently large, neither $U(1)_{_{S^T}}$ spin liquids survive as the bond parameters on the large tetrahedra tend to zero upon solving the self consistent mean field equations iteratively. The parameter region that favors such a decoupled tetrahedron solution contains the candidate spin model [39, 40] for the material Ba$_3$Yb$_2$Zn$_5$O$_{11}$ (BYZO), whose interactions on the large tetrahedra appear to be orders of magnitude smaller than those on the small tetrahedra. The $U(1)_{_{S^T}}$ and $U(1)_{_{S^+}}$ spin liquids are thus more relevant for spin models closer to the Heisenberg limit or with a weaker breathing anisotropy. While BYZO is the only $S = 1/2$ breathing pyrochlore material discovered so far, similar materials with parameters that span the phase diagram in our work may be synthesized in near future.

The AFM Heisenberg model on the breathing pyrochlore lattice had been investigated in several prior works, first as a route to understand the same model but on the regular pyrochlore lattice, and later for its own sake when breathing pyrochlore materials made their appearance in experimental laboratories. It is therefore worth briefly discussing our work in relation to the existing literature.

Starting from decoupled tetrahedra with one $S = 1/2$ moment per vertex, Refs. 89 and 90 treat the intertetrahedron interaction perturbatively up to third order, and arrive at an effective Hamiltonian with one pseudospin-1/2 degree of freedom per tetrahedron and interactions that involve three nearby pseudospins. A mean field approximation, which essentially treats the pseudospins as classical vectors, yields a partially dimerized ground state, leaving one disordered pseudospin for every four pseudospins. This is different from our approach, which is nonperturbative, and our spin liquid ansatz preserves the lattice symmetry, while the dimerized state breaks it. Such a perturbative reconnection method first appeared in Ref. 91 (see also Ref. 92) for $S = 1/2$, and it was also extended to $S = 1$ and $S = 3/2$ in Ref. 25.

Ref. 24 uses gauge mean field theory [15, 16] to study the $S = 1/2$ XXZ model on the breathing pyrochlore lattice. While gauge mean field theory provides an excellent description of the quantum spin ice state near the Ising limit, it is unclear how accurately the theory captures other types of quantum spin liquid in a more generic parameter region. For example, it predicts an antiferromagnetically ordered state in the AFM Heisenberg limit, where the strong frustration warrants the consideration of possible quantum spin liquid candidates, which can be systematically classified by PSG.

Ref. 23 studies the Heisenberg model on the breathing pyrochlore lattice in the classical limit, where the signs of $J_1$ and $J_2$ are allowed to be different. Interestingly, the classical ground state depends only on the signs of the interactions but not on their relative magnitude. In particular, for $J_1 > 0$ and $J_2 > 0$, the ground state is a Coulombic spin liquid [93] subject to the divergenceless condition that the vector summation of the four spins on every tetrahedron is zero, which is the continuous version of the 2-in-2-out ice rule. This gives us reason to look for quantum spin liquids in the corresponding $S = 1/2$ model.

More interestingly, a recent $SU(2)$ density matrix renormalization group study [94] of the $S = 1/2$ AFM Heisenberg model on the regular pyrochlore lattice suggests a disordered ground state with spontaneously broken inversion symmetry, as evidenced by a difference in energy between the two species of tetrahedra. Assuming that none of the remaining symmetries is broken, the quantum spin liquids classified in this work may even be considered as candidate ground states of the regular pyrochlore lattice.

We close by mentioning several potentially interesting future directions. While we have focused on the $U(1)$ spin liquids in this work, it will also be useful to conduct a thorough examination of the $Z_2$ spin liquids, though they are greater in number. Besides, we have only considered here time reversal symmetric spin liquid ansatzes. It may be worthwhile to study ansatze that break time reversal symmetry, like the monopole flux state in Ref. 85, as they may give rise to stable chiral spin liquids. Moreover, it is known that mean field theory does not give highly accurate energetics, and advanced numerical techniques, specifically variational Monte Carlo with Gutzwiller projection, may be employed to obtain more reliable estimates of the energies of the various spin liquids, magnetic orders, and other candidate ground states. Our PSG analysis lay the foundation on which the variational wave functions of the quantum spin liquids can be constructed for such comparisons.

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**Appendix A: Algebraic identities**

In this appendix, we list the algebraic identities of the breathing pyrochlore lattice:

\[ T_2^{-1}T_1^{-1}T_3T_1 = e, \]  
\[ T_3^{-1}T_2^{-1}T_3T_2 = e, \]  

(A1a)  

(A1b)
\[ T^{-1}_1 T^{-1}_2 T_3 = e, \quad (A1c) \]
\[ C^{-1}_1 T^{-1}_2 C_2 T_1 = e, \quad (A1d) \]
\[ C^{-1}_1 T^{-1}_3 C_3 T_2 = e, \quad (A1e) \]
\[ C^{-1}_1 T_1 C_3 T_3 = e, \quad (A1f) \]
\[ \sigma^{-1}_i T_i \sigma T_i = e, \quad (A1g) \]
\[ \sigma^{-1}_2 T_1 \sigma T_2 = e, \quad (A1h) \]
\[ \sigma^{-1}_3 T_3 \sigma T_3 = e, \quad (A1i) \]
\[ C_3 = e, \quad (A1j) \]
\[ \sigma^2 = e, \quad (A1k) \]
\[ (\sigma C_3)^4 = e, \quad (A1l) \]
\[ O^{-1} T^{-1}_1 OT = e, \quad (A1m) \]
\[ T^2 = e, \quad (A1n) \]

where \( O \) in (A1m) denotes any space group generator, and \( e \) is the identity element.

**Appendix B: Complex fermion mean field theory**

Using the parton representation (5), the antiferromagnetic Heisenberg model (6) can be written as, up to some constant,

\[ H = -\sum_{ij} J_{ij} \left( \hat{\psi}^\dagger_{i} \psi_{j} + \hat{\Delta}_{ij} \right), \quad (B1) \]

where

\[ \hat{\psi}_{i} = f_{i}^\dagger \psi_{i}, \quad (B2a) \]
\[ \hat{\Delta}_{ij} = f_{i}^\dagger f_{j}^\dagger - f_{i} f_{j}, \quad (B2b) \]

are the singlet hopping and pairing operators respectively. A mean field decoupling then leads to (7).

Ref. 60 first noted an alternative form of (5),

\[ S_{i} = \frac{1}{4} \text{Tr} \left[ \Psi_{i}^\dagger \sigma \Psi_{i} \right], \quad \Psi_{i} = \left( f_{i}^\dagger \right) \begin{pmatrix} f_{i1}^\dagger & f_{i2}^\dagger & f_{i3}^\dagger \end{pmatrix}, \quad (B3) \]

from which it is easy to see that an SU(2) gauge transformation \( G_{i} \), such that \( \Psi_{i} \rightarrow \Psi_{i} G_{i} \), leaves the spin operator invariant. Note that \( G_{i} \) preserves the anticommutation relation of fermions. On the other hand, the mean field Hamiltonian (7) can be written as

\[ H_{MF} = \sum_{ij} \text{Tr} \left[ \Psi_{i} u_{ij} \Psi_{j}^\dagger \right], \quad (B4) \]

where \( u_{ij} \) is defined in (8) and the on-site \( u_{ii} \) is given by

\[ u_{ii} = -\frac{1}{2} \begin{pmatrix} \lambda_{i}^{(1)} & \lambda_{i}^{(1)} + i \lambda_{i}^{(2)} & \lambda_{i}^{(3)} \\ \lambda_{i}^{(1)} - i \lambda_{i}^{(2)} & -\lambda_{i}^{(3)} \end{pmatrix}. \quad (B5) \]

As stated in the main text, due to the SU(2) gauge redundancy, the mean field Hamiltonian is invariant under a symmetry \( X \) of the system only up to a gauge transformation \( G_X \in SU(2) \). From Ref. 54,

\[ H_{MF} \xrightarrow{X} \sum_{ij} \Psi_{X(i)} u_{ij} \Psi_{X(j)}, \quad (B6) \]

we see that the mean field ansatz should obey (9).

**Appendix C: Classification of \( \mathbb{Z}_2 \) spin liquids**

We solve

\[ G_{T_2}^{(1)}(i)G_{T_2}^{(i)}(i)G_{T_2}^{1}(i)G_{T_2}^{(i)}(i) = \eta_{T_2}, \quad (C1a) \]
\[ G_{T_2}^{(1)}(i)G_{T_2}^{1}(i)G_{T_2}^{(i)}(i)G_{T_2}^{(i)}(i) = \eta_{T_2}, \quad (C1b) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1c) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1d) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1e) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1f) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1g) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1h) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1i) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1j) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1k) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1l) \]
\[ G_{C_1}^{(1)}(C_1^{(1)})(i)G_{C_1}^{(i)}(i)G_{C_1}^{(i)}(i)G_{C_1}^{(1)}(i) = \eta_{C_1}, \quad (C1m) \]
\[ [G_{T_2}^{(i)}] = \eta_{T_2}, \quad (C1n) \]

for \( G_X(r_1, r_2, r_3; s) \), where each \( \eta_{\ldots} \) on the right hand side is either +1 or −1.

First, notice that for a given \( X \) we are free to multiply \( G_X \) by an element of IGG. We exploit this IGG freedom of \( G_{T_2} \) to fix \( \eta_{T_2} = 1 \), of \( G_{T_2} \) to fix \( \eta_{T_2} = 1 \), of \( G_{T_2} \) to fix \( \eta_{C_1} = 1 \), and of \( G_{C_1} \) to fix \( \eta_{C_1} = 1 \). Then, using

\[ G_{X}(i) \rightarrow W(i) G_{X}(i) W^{\dagger}(X^{-1}(i)), \quad W(i) \in SU(2), \quad (C2) \]

we fix

\[ G_{T_2}(r_1, r_2, r_3; s) = 1, \]
\[ G_{T_2}(0, r_2, r_3; s) = 1, \]
\[ G_{T_2}(0, 0, r_3; s) = 1. \]
Then, (C1a) yields
\[ G_{T_2}^{ij} (r_1 - 1, r_2, r_3; s) G_{T_2} (r_1, r_2, r_3; s) = \eta_{1T_1}^* \] (C4)

Eqs. (C1b) and (C1c) yield
\[ G_{T_1} (r_1, r_2, r_3; s) = \eta_{1T_1}^* G_{T_1}(r_1, r_2, r_3; s) \]
and
\[ G_{T_1} (0, r_2, r_3; s) = \eta_{1T_1}^* G_{T_1}(0, r_2, r_3; s) \]
respectively, so
\[ G_{T_1} (r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \] (C5)

Eqs. (C1d), (C1e), and (C1f) yield
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* G_C(r_1, r_2, r_3; s), \] (C6a)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* G_C(r_1, r_2, r_3; s), \] (C6b)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* G_C(r_1, r_2, r_3; s), \] (C6c)

which further lead to
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g_C(s), \] (C7a)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \eta_{3T_3}^* g_C(s), \] (C7b)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \eta_{3T_3}^* g_C(s), \] (C7c)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \eta_{3T_3}^* g_C(s), \] (C7d)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \eta_{3T_3}^* g_C(s), \] (C7e)
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \eta_{3T_3}^* g_C(s), \] (C7f)

where \( g_X(s) = G_X(0, 0; s) \). The right hand sides of these six equations must be equal to each other, which forces \( \eta_{1T_1} = \eta_{2T_2} = \eta_{3T_3} \). It follows that
\[ G_{C_1}(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g_C(s), \] (C8)

Eq. (C1j), with \( i = (r_1, r_2, r_3; 0) \), yields
\[ \eta_{1T_1}^* \eta_{2T_2}^* [g_C(0)]^3 = 1. \] (C9)

Since the right hand side has no coordinate dependence, we must have \( \eta_{C_1T_1} = 1 \), which is anticipated since (A1d) is implied by (A1e) and (A1f). On the other hand, with \( i = (r_1, r_2, r_3; s = 1, 2, 3) \), we get
\[ g_C(3) g_C(2) g_C(1) = 1. \] (C10)

Eq. (C1g) yields
\[ G_\sigma (r_1, r_2, r_3; s) = \eta_{1T_1}^* G_\sigma (0, r_2, r_3; s). \] (C11)

Subsequently, (C1h) and (C1i) yield
\[ G_\sigma (0, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g_\sigma(0, r_2, r_3; s), \] (C12a)
\[ G_\sigma (0, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g_\sigma(0, r_2, r_3; s), \] (C12b)

which together imply
\[ G_\sigma(r_1, r_2, r_3; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g_\sigma(0, r_2, r_3; s). \] (C12c)

Eq. (C1k), with \( i = (r_1, r_2, r_3; 0) \) yields
\[ \eta_{1T_1}^* g_\sigma(1) g_\sigma(0) = \eta_\sigma, \] (C13)

which implies \( \eta_\sigma T_1 = 0 \). We also have
\[ [g_\sigma(2)]^2 = \eta_\sigma, [g_\sigma(3)]^2 = \eta_\sigma, \] (C14)

with \( i = (r_1, r_2, r_3; s = 2, 3) \). Finally, (C11) yields
\[ g_\sigma(0) g_C(1) g_\sigma(3) g_C(3) g_\sigma(2) g_C(2) g_\sigma(1) g_C(0) = \eta_\sigma C_1. \] (C11)

We now proceed to the parts that involve the time reversal symmetry. Eq. (C1m) with \( O = T_1, T_2, T_3 \) yields
\[ G_T(r_1, r_2; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g(T), \] (C15a)
\[ G_T(r_1, r_2; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g(T), \] (C15b)
\[ G_T(r_1, r_2; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g(T), \] (C15c)

which together imply
\[ G_T(r_1, r_2; s) = \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* g(T). \] (C16)

Eq. (C1m) with \( O = C_3 \) and \( i = (r_1, r_2, r_3; 0) \) yields
\[ \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \delta_{C_3}(0) \eta(T) \delta_{C_3}(0) = \eta_{C_3}, \] (C17)

which implies \( \eta_{1T_1} = \eta_{2T_2} = \eta_{3T_3} \). We also have
\[ g_C^{\dagger}(2) g_C^{\dagger}(2) g_C(2) g_C(2) \eta_\sigma(1) = \eta_{C_3}, \] (C18a)
\[ g_C^{\dagger}(3) g_C^{\dagger}(3) g_C(3) g_C(3) \eta_\sigma(2) = \eta_{C_3}, \] (C18b)
\[ g_C^{\dagger}(1) g_C^{\dagger}(1) g_C(1) g_C(1) \eta_\sigma(3) = \eta_{C_3}, \] (C18c)

with \( i = (r_1, r_2, r_3; s = 1, 2, 3) \). Eq. (C1m) with \( O = \sigma \) and \( i = (r_1, r_2, r_3; 0) \) yields
\[ \eta_{1T_1}^* \eta_{2T_2}^* \eta_{3T_3}^* \delta_{\sigma}(0) \eta(T) \delta_{\sigma}(0) = \eta_{\sigma}, \] (C19)

which implies \( \eta_{1T_1} = 1 \). We also have
\[ g_\sigma^{\dagger}(1) g_\sigma^{\dagger}(1) g_\sigma(1) g_\sigma(1) \eta_\sigma(1) = \eta_{\sigma}, \] (C20a)
\[ g_\sigma^{\dagger}(2) g_\sigma^{\dagger}(2) g_\sigma(2) g_\sigma(2) \eta_\sigma(2) = \eta_{\sigma}, \] (C20b)
\[ g_\sigma^{\dagger}(3) g_\sigma^{\dagger}(3) g_\sigma(3) g_\sigma(3) \eta_\sigma(3) = \eta_{\sigma}, \] (C20c)

with \( i = (r_1, r_2, r_3; s = 1, 2, 3) \). Eq. (C1n) yields
\[ [g_\sigma(T)]^2 = \eta_\sigma. \] (C21)

a. Grand summary

Before we proceed to gauge fixing, it is worthwhile to recollect the results obtained thus far. They are (12a)-(12f) shown in the main text, together with the "sublattice constraints":
\[ g_C(0) = 1, \]
\[ g_C(3) g_C(2) g_C(1) = 1, \] (C22a)
\[ g_C(0) = 1, \] (C22b)
The relation between $SU(2)$ transformations and $SO(3)$ rotations, without changing any previously fixed gauge which is proportional to identity. On the other hand, the most generic solution to (C22a) is $g_{c_1}(0) = \exp[i(2\pi q_0/3){\mathbf{n}}_0 \cdot \hat{\tau}]$, where $q_0 \in \{0, 1, 2\}$ and $\mathbf{n}_0$ is a unit vector. Eq. (C22f) leads to $g_{c_3}(3) = -\eta_{c_1}g_{c_1}^\dagger(0)g_{c_3}(2)$. (C22e) then gives $g_{c_3}(3)g_{c_3}(2)g_{c_3}(1)=\eta_{c_1}g_{c_1}$. This leads to a global gauge transformation of the form $e^{i\theta\hat{\tau}_2}$, i.e., a uniform gauge rotation about the $\tau_2$ axis, which does not affect any of the previously fixed gauges. Thus $g_{c_1}(0) = \exp[i(2\pi q_0/3){\tau}_3]$ and $g_{c_3}(3) = \eta_{c_1}g_{c_1}(i\tau_2)\exp[i(2\pi q_0/3){\tau}_3]$.

**Case 2.** $\eta_{c_1} = -1$. Eq. (C22o) implies $g_{c_1}(0) = \hat{\mathbf{n}}_1 \cdot \hat{\tau}$ and $g_{c_1}(1, 2, 3) = \hat{\mathbf{n}}_1 \cdot \hat{\tau}$, where $\hat{\mathbf{n}}_1$ are unit vectors and $\hat{\tau} = (\tau_1, \tau_2, \tau_3)$. We rotate $\hat{\mathbf{n}}_1$ to $(0, 1, 0)$, such that $g_{c_1}(1, 2, 3) = i\tau_2$ by a global gauge transformation, without changing any previously fixed gauge which is proportional to identity. (Note: $\hat{\mathbf{n}}_1$ here is unrelated to that defined in 1.2. We merely recycle the notation. As this is quite clear from the context, we will not give such warnings should similar situations arise later.)

**Case 2.1.** $\eta_{c_1} = +1$. Eqs. (C22d) and (C22e) imply $g_{c_3}(2) = \xi_2$ and $g_{c_3}(3) = \xi_3$ respectively, with $\xi_{2, 3} \in \{+1, -1\}$. We perform a sublattice gauge transformation $W_0 = \xi_2g_{c_1}^\dagger(0)$, $W_{1, 2, 3} = 1$ to fix $g_{c_3}(0) = \xi_2$, without affecting the previously fixed gauges. It follows from (C22c) that $g_{c_1}(2) = \xi_2$. We use the IGG freedom of $G_{c_1}$ to fix $g_{c_1}(0, 1, 2) = 1$ and $g_{c_3}(2) = \xi_2\xi_3$. Since the product $\xi_2\xi_3 = \pm 1$, let us just call it $\xi_3$ for simplicity. Eq. (C22f) yields $g_{c_1}(0) = \xi_3\eta_{c_1}g_{c_1}$, which together with (C22a) implies $g_{c_1}(0) = \xi_3$ and $\xi_3 = \eta_{c_1}$. (Note that the procedures outlined above are exactly same as those in Case 1.1.) Finally, (C22m) or (C22n) imply $\eta_{c_1} = 1$, which together with (C22k) or (C22l) imply $g_{c_1}(0) = i\tau_2$.

**Case 2.2.** $\eta_{c_1} = -1$. Eqs. (C22d) and (C22e) imply $g_{c_3}(2) = \hat{\mathbf{n}}_2 \cdot \hat{\tau}$ and $g_{c_3}(3) = \hat{\mathbf{n}}_3 \cdot \hat{\tau}$ respectively, where $\hat{\mathbf{n}}_{2, 3}$ are unit vectors.

**Case 2.2.1.** $\eta_{c_1} = +1$. From (C22m), we see that $g_{c_3}(2)$ commutes with $\tau_2$, so $g_{c_3}(2) = \xi_2\hat{\tau}_2$, $\xi_2 = \pm 1$. Similarly, from (C22n) we have $g_{c_3}(3) = \xi_3(i\tau_2)$, $\xi_3 = \pm 1$. We then perform a sublattice dependent gauge transformation $W_0 = \xi_2g_{c_3}^\dagger(0)$, $W_{1, 2, 3} = 1$ to fix $g_{c_3}(0) = \xi_2$. It follows from (C22c) that $g_{c_3}(1) = -\xi_2$. Using the IGG freedom of $G_{c_3}$, we can eliminate $\xi_2$ and redefine $\xi_2\hat{\tau}_3 \rightarrow \xi_3$ as in Case 2.1. Eq. (C22f) yields $g_{c_1}(0) = \xi_3\eta_{c_1}g_{c_1}$, which together with (C22a) imply $g_{c_1}(0)$ is $1$ and $\xi_3 = \eta_{c_1}$. Finally, (C22k) implies $g_{c_1}(0) = i\tau_2$. To render the solutions in a neater form, we further perform a sublattice dependent gauge transformation $W_0 = i\tau_2$, $W_{1, 2, 3} = 1$ such that $g_{c_1}(0, 1) \rightarrow i\tau_2$, while others are unaffected.

**Case 2.2.2.** $\eta_{c_1} = -1$. We first perform a sublattice dependent gauge transformation, $W_0 = g_{c_3}^\dagger(0)$, $W_{1, 2, 3} = 1$ to fix $g_{c_3}(0) = 1$. It follows from (C22c) that $g_{c_3}(1) = -1$. Eq. (C22k) or (C22l) then implies $g_{c_1}(0) = -\xi_2$. From (C22m), we see that $g_{c_3}(2)$ anticommutes with $\tau_2$, which implies that $\hat{\mathbf{n}}_2$ cannot have a finite $\tau_2$ component, i.e., it has the form $(\sin \theta, 0, \cos \theta)$. We further rotate $\hat{\mathbf{n}}_2$ to $(0, 0, 1)$ via a global gauge transformation $\exp(-i\theta\tau_2/2)$, which does not affect any of the previ-
ously fixed gauges. Thus $g_{\sigma}(2) = i\tau_3$. The most generic solution to (C22a) is $g_{C_3}(0) = \exp[i(2\pi q_0/3)\hat{n}_0 \cdot \mathbf{T}]$, where $q_0 = 0, 1, 2$ and $\hat{n}_0$ is a unit vector. But (C22g) requires $g_{C_3}(0)$ to commute with $i\tau_2$, so $\hat{n}_0 = (0, \pm 1, 0)$, and we can further specialize to the plus sign without loss of generality. Finally, (C22f) yields $g_{\sigma}(3) = \eta_{\sigma C_3}(i\tau_2)\exp[i(2\pi q_0/3)\tau_2]$, which anticommutes with $\tau_2$ as required by (C22n). To render the solutions in a neater form, we further specify a sublattice dependent gauge transformation $W_0 = i\tau_3$, $W_{1,2,3} = 1$, such that $g_{C_3}(0) \rightarrow \exp[-i(2\pi q_0/3)\tau_2]$, $g_{\sigma}(0,1) \rightarrow i\tau_3$, and $g_{\sigma}(0) \rightarrow i\tau_2$, while others are unaffected.

We will exclude the solutions with $\eta_{\sigma} = +1$ (Case 1) because they have $g_{\sigma}(i) = 1$ for all sites $i$, which forces $u_{ij} = 0$ for any pair of sites $i$ and $j$ by (10). These solutions, which lead to vanishing mean field ansatze and thus a zero Hamiltonian, are unphysical. Let us count the remaining solutions (Case 2). Each of 2.1, 2.2.1, 2.2.2 has three $\mathbb{Z}_2$ variables $\eta_{T_1}, \eta_{T_2},$ and $\eta_{T_3}$. For 2.2.2, there is an additional $\mathbb{Z}_2$ variable $q_0$. There are in total $2^3 \times (1 + 1 + 3) = 40$ gauge inequivalent solutions, i.e., 40 possible $\mathbb{Z}_2$ spin liquids. They are listed in Table I.

### Appendix D: Classification of $U(1)$ spin liquids

We solve

\begin{align*}
G_{T_3}^\dagger(T_1^{-1}(i))G_{T_3}^\dagger(i)G_{T_3}(i)G_{T_3}(T_1^{-1}(i)) &= e^{i\theta_{T_3T_3}}, \\
G_{T_1}^\dagger(T_1^{-1}(i))G_{T_1}^\dagger(i)G_{T_1}(i)G_{T_1}(T_1^{-1}(i)) &= e^{i\theta_{T_1T_1}}, \\
G_{T_1}^\dagger(T_3^{-1}(i))G_{T_1}^\dagger(i)G_{T_1}(i)G_{T_1}(T_3^{-1}(i)) &= e^{i\theta_{T_1T_3}}, \\
G_{C_3}^\dagger(T_2^{-1}(i))G_{C_3}^\dagger(i)G_{C_3}(i)G_{C_3}(T_2^{-1}(i)) &= e^{i\theta_{C_3T_2}}, \\
G_{C_3}^\dagger(T_3^{-1}(i))G_{C_3}^\dagger(i)G_{C_3}(i)G_{C_3}(T_3^{-1}(i)) &= e^{i\theta_{C_3T_3}}, \\
\sigma_{\sigma}(T_1^{-1}(i))G_{\sigma}(i)G_{\sigma}(T_1^{-1}(i)) &= e^{i\theta_{\sigma\sigma}}, \\
\sigma_{\sigma}(T_2^{-1}(i))G_{\sigma}(i)G_{\sigma}(T_2^{-1}(i)) &= e^{i\theta_{\sigma\sigma}}, \\
\sigma_{\sigma}(T_3^{-1}(i))G_{\sigma}(i)G_{\sigma}(T_3^{-1}(i)) &= e^{i\theta_{\sigma\sigma}}, \\
\sigma_{\sigma}((\sigma C_3)^{2}(i))G_{\sigma}(i)G_{\sigma}(i) &= e^{i\theta_{\sigma\sigma}}, \\
\sigma_{\sigma}((\sigma C_3)^{2}(i))G_{\sigma}(i)G_{\sigma}(C_3^{-1}(i)) &= e^{i\theta_{\sigma\sigma}}, \\
G_{C_3}^\dagger(C_3(i))G_{C_3}^\dagger(i)G_{C_3}(i) &= e^{i\theta_{C_3C_3}}, \\
G_{C_3}^\dagger(i)G_{C_3}^\dagger(i)G_{C_3}(i)G_{C_3}(i) &= e^{i\theta_{C_3C_3}}, \\
G_{\sigma}(i)G_{\sigma}(i)G_{\sigma}(O^{-1}(i)) &= e^{i\theta_{\sigma\sigma}}, \\
\left[G_{\sigma}(i)\right]^2 &= e^{i\theta_{\sigma\sigma}},
\end{align*}

for $G_X(r_1, r_2, r_3; s)$, where each $\theta_{\sigma\sigma}$ on the right hand side is a continuous variable in the interval $[0, 2\pi)$. For $U(1)$ spin liquids, the gauge transformations have the specific form (13).

For a symmetry operator $X$ that appear an odd number of times in an algebraic identity, if any other symmetry operators appear an even number of times in the same algebraic identity, we must have $n_X = 0$, otherwise the equality of the corresponding equation in (D1a)-(D1n) will not hold. Therefore, (D1b) or (D1l) force $n_{T_1} = 0$, and (D1j) forces $n_{C_3} = 0$. Since $G_{T_1}$ and $G_{C_3}$ do not carry $i\tau_1$ with them, (D1d) and (D1f) then force $n_{T_2} = 0$ and $n_{T_3} = 0$ respectively. Next, we use the IGG freedoms (see App. C) of $G_{T_1}, G_{T_2},$ and $G_{T_3}$ to fix $\theta_{T_1T_1} = 0$, $\theta_{C_3T_3} = 0$, and $\theta_{\sigmaT_1} = 0$. Notice that equalities involving $U(1)$ variables, which are abundant in this appendix, are defined modulo $2\pi$.

Through (C2) we fix

\begin{align*}
\phi_{T_1}(r_1, r_2, r_3; s) &= 0, \\
\phi_{T_2}(0, r_2, r_3; s) &= 0, \\
\phi_{T_3}(0, 0, r_3; s) &= 0.
\end{align*}

Eq. (D1a) yields $-\phi_{T_1}(r_1, r_2, r_3; s) + \phi_{T_1}(r_1, r_2, r_3; s) = \theta_{T_1T_1}$, or

\begin{equation}
\phi_{T_1}(r_1, r_2, r_3; s) = r_1\theta_{T_1T_1}.
\end{equation}

Eqs. (D1b) and (D1c) respectively yield

\begin{align*}
\phi_{T_1}(r_1, r_2, r_3; s) &= -r_1\theta_{T_1T_1} + \phi_{T_1}(0, r_2, r_3; s), \\
\phi_{T_2}(r_1, r_2, r_3; s) &= r_2\theta_{T_1T_2} + \phi_{T_2}(r_1, 0, r_3; s),
\end{align*}

so

\begin{equation}
\phi_{T_1}(r_1, r_2, r_3; s) = r_2\theta_{T_1T_2} - r_1\theta_{T_1T_3}.
\end{equation}

Eqs. (D1d), (D1e), and (D1f) yield

\begin{align*}
\phi_{C_3}(r_1, r_2, r_3; s) &= r_2\theta_{C_3T_1} + r_1r_2\theta_{T_1T_2} + \phi_{C_3}(r_1, 0, r_3; s), \\
\phi_{C_3}(r_1, r_2, r_3; s) &= -r_3r_1\theta_{T_1T_1} + r_2r_3\theta_{T_1T_2} - \theta_{T_1T_1} + \phi_{C_3}(r_1, r_2, 0; s) + \phi_{C_3}(r_1, r_2, r_3; s),
\end{align*}

which further lead to

\begin{align*}
\phi_{C_3}(r_1, r_2, r_3; s) &= r_2\theta_{C_3T_1} + r_1r_2\theta_{T_1T_2} - r_3r_1\theta_{T_1T_1} + \phi_{C_3}(r_1, 0, r_3; s), \\
\phi_{C_3}(r_1, r_2, r_3; s) &= r_2\theta_{C_3T_1} + r_1r_2\theta_{T_1T_2} - r_3r_1\theta_{T_1T_1} + \phi_{C_3}(r_1, 0, r_3; s) + \phi_{C_3}(r_1, r_2, 0; s) + \phi_{C_3}(r_1, r_2, r_3; s),
\end{align*}

where $\phi_{X}(s) = \phi_{X}(0, 0, 0; s)$. The right hand sides of these six equations must be equal to each other, which forces $\theta_{T_1T_1} = \theta_{C_3T_3} = \theta_{\sigmaT_1}$. Eq. (D1j) with $i = (r_1, r_2, r_3; 0)$ yields

\begin{equation}
(r_1 + r_2 + r_3)\theta_{C_3T_1} + 3\phi_{C_3}(0) = \theta_{C_3}.
\end{equation}
Since the right hand side has no coordinate dependence, we must have $\theta_{C,T_1} = 0$, which is anticipated since (A1d) is implied by (A1e) and by (A1f). We also have
\[
\varphi_{C_1}(3) + \varphi_{C_1}(2) + \varphi_{C_1}(1) = \theta_{C_1},
\] (D9)
with $i = (r_1, r_2, r_3; s = 1, 2, 3).
We now proceed to the parts that involve $\sigma$.

**Case 1.** $n_{i\sigma} = 0$. Eq. (D1g) yields
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = -r_1\theta_{\sigma T_1} + \phi_{\sigma}(0, r_2, r_3; s).
\] (D10)
Subsequently, (D1h) and (D1i) yield
\[
\phi_{\sigma}(0, r_2, r_3; s) = \frac{r_2(r_2 + 1)}{2} + r_2(2r_1 + r_3 + 1) \theta_{T_2 T_1}
+ r_2(\theta_{\sigma T_2} - \theta_{\sigma T_1}) + \phi_{\sigma}(0, 0, r_2, r_3; s),
\]
\[
\phi_{\sigma}(0, r_2, r_3; s) = -\frac{r_3(r_3 + 1)}{2} + r_3(2r_1 + r_2 + 1) \theta_{T_2 T_1}
- r_3\theta_{\sigma T_1} + \phi_{\sigma}(0, r_2, 0; s),
\]
which further lead to
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = \left[\frac{r_2(r_2 + 1)}{2} + r_2(2r_1 + r_3 + 1)\right] \theta_{T_2 T_1}
+ (2r_1 + 1)(r_2 - r_3) + r_2 r_3 \theta_{T_2 T_1}
- (r_1 + r_2 + r_3)\theta_{\sigma T_1} + r_2\theta_{\sigma T_2} + \varphi_{\sigma}(s),
\]
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = \left[\frac{r_3(r_3 + 1)}{2} - r_3(2r_1 + r_2 + 1)\right] \theta_{T_2 T_1}
+ (2r_1 + 1)(r_2 - r_3) - r_2 r_3 \theta_{T_2 T_1}
- (r_1 + r_2 + r_3)\theta_{\sigma T_1} + r_2\theta_{\sigma T_2} + \varphi_{\sigma}(s),
\]
which further lead to
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = \left[\frac{r_2(r_2 + 1)}{2} - \frac{r_3(r_3 + 1)}{2}\right] \theta_{T_2 T_1}
+ (2r_1 + 1)(r_2 - r_3) + r_2 r_3 \theta_{T_2 T_1}
- (r_1 + r_2 + r_3)\theta_{\sigma T_1} + r_2\theta_{\sigma T_2} + \varphi_{\sigma}(s),
\]
\[\phi_{\sigma}(0, r_2, r_3; s) = \frac{r_3(r_3 + 1)}{2} + r_3(3r_2 - 1) \theta_{T_2 T_1}
- r_3\theta_{\sigma T_1} + \phi_{\sigma}(0, r_2, 0; s),\]
which further lead to
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = \left[\frac{r_2(r_2 + 1)}{2} - \frac{r_3(r_3 + 1)}{2}\right] \theta_{T_2 T_1}
- r_2 + r_3 + r_2 r_3 \theta_{T_2 T_1}
- (r_1 + r_2 + r_3)\theta_{\sigma T_1} + r_2\theta_{\sigma T_2} + \varphi_{\sigma}(s),
\]
\[
\phi_{\sigma}(r_1, r_2, r_3; s) = \left[\frac{r_2(r_2 + 1)}{2} - \frac{r_3(r_3 + 1)}{2}\right] \theta_{T_2 T_1}
- r_2 + r_3 - 3r_2 r_3 \theta_{T_2 T_1}
- (r_1 + r_2 + r_3)\theta_{\sigma T_1} + r_2\theta_{\sigma T_2} + \varphi_{\sigma}(s).
\]
The right hand sides of these two equations must be equal to each other, which forces $\theta_{T_2 T_1} = 2p_{T_2 T_1}/4$, $p_{T_2 T_1} \in \{0, 1, 2, 3\}$.
Eq. (D1k) with $i = (r_1, r_2, r_3; 0)$ yields
\[- (2r_1 + 2r_3)\theta_{\sigma T_1} - \varphi_{\sigma}(1) + \varphi_{\sigma}(0) = \theta_{\sigma} + \pi, \]
which implies $\theta_{\sigma T_1} = 0$. The additive factor of $\pi$ on the right hand side comes from $(i r_1)^2 = -1$. We also have
\[- \varphi_{\sigma}(2) + \varphi_{\sigma}(2) = \theta_{\sigma} + \pi \]
with $i = (r_1, r_2, r_3; 2)$, which implies $\theta_{\sigma} = \pi$.
Eq. (D1l) with $\sigma^{-1}(i) = (r_1, r_2, r_3; 0)$ yields
\[2(r_1 + r_2)\theta_{\sigma T_2} - \varphi_{\sigma}(0) - \varphi_{\sigma}(1) + \varphi_{\sigma}(3) + \varphi_{\sigma}(3)
- \varphi_{\sigma}(2) - \varphi_{\sigma}(2) + \varphi_{\sigma}(1) + \varphi_{\sigma}(0) = \theta_{\sigma C_1}, \]
which implies $\theta_{\sigma T_2} = p_{\sigma T_2} \pi$, $p_{\sigma T_2} \in \{0, 1\}$. We also have
\[- \varphi_{\sigma}(3) - \varphi_{\sigma}(3) + \varphi_{\sigma}(2) + \varphi_{\sigma}(2)
- \varphi_{\sigma}(1) - \varphi_{\sigma}(0) + \varphi_{\sigma}(0) + \varphi_{\sigma}(1) = \theta_{\sigma C_1} \]
with $\sigma^{-1}(i) = (r_1, r_2, r_3; 1)$, which implies $\theta_{\sigma C_1} = p_{\sigma C_1} \pi$, $p_{\sigma C_1} \in \{0, 1\}$.
We now proceed to consider the parts that involve $\mathcal{T}$.

**Case 1.** $n_{t\mathcal{T}} = 0$. Eq. (D1m) with $O = T_1, T_2, T_3$ yields
\[
\phi_{\mathcal{T}}(r_1, r_2, r_3; s) = -r_1\theta_{\mathcal{T}_1 \mathcal{T}} + \phi_{\mathcal{T}}(0, r_2, r_3; s), \]
\[
\phi_{\mathcal{T}}(r_1, r_2, r_3; s) = -r_2\theta_{\mathcal{T}_2 \mathcal{T}} + \phi_{\mathcal{T}}(r_1, 0, r_3; s), \]
\[
\phi_{\mathcal{T}}(r_1, r_2, r_3; s) = -r_3\theta_{\mathcal{T}_3 \mathcal{T}} + \phi_{\mathcal{T}}(r_1, r_2, 0; s), \]
which together imply
\[
\phi_{\mathcal{T}}(r_1, r_2, r_3; s) = -r_1\theta_{\mathcal{T}_1 \mathcal{T}} - r_2\theta_{\mathcal{T}_2 \mathcal{T}} - r_3\theta_{\mathcal{T}_3 \mathcal{T}} + \phi_{\mathcal{T}}(s). \]
Eq. (D1m) with $O = C_3$ and $C_3^{-1}(i) = (r_1, r_2, r_3; 0)$ yields
\[r_3 - r_1)\theta_{\mathcal{T}_1 \mathcal{T}} + (r_1 - r_2)\theta_{\mathcal{T}_2 \mathcal{T}} + (r_2 - r_3)\theta_{\mathcal{T}_3 \mathcal{T}} = \theta_{C_3 \mathcal{T}}, \]
(D25)
which implies $\theta_{T_1T} = \theta_{T_2T} = \theta_{T_3T}$, and subsequently $\theta_{C_3T} = 0$. We also have

\[
- \varphi_T(2) + \varphi_T(1) = 0, \quad \text{(D2a)}
\]
\[
- \varphi_T(3) + \varphi_T(2) = 0, \quad \text{(D2b)}
\]
\[
- \varphi_T(1) + \varphi_T(3) = 0, \quad \text{(D2c)}
\]

with $C_3^{-1}(i) = (r_1, r_2, r_3; s = 1, 2, 3)$.

**Case 1.1.** $n_{\sigma} = 0$. Eq. (D1m) with $O = \sigma$ and $\sigma^{-1}(i) = (r_1, r_2, r_3; 0)$ yields

\[
-(2r_1 + r_2 + r_3)\theta_{T_1T} - \varphi_T(1) + \varphi_T(0) = \theta_{\sigma T}, \quad \text{(D27)}
\]

which implies $\theta_{T_1T} = 0$. We also have

\[
- \varphi_T(2) + \varphi_T(2) = \theta_{\sigma T}, \quad \text{(D28)}
\]

with $\sigma^{-1}(i) = (r_1, r_2, r_3; 2)$, which implies $\theta_{\sigma T} = 0$.

Eq. (D1n) then yields

\[
2\varphi_T(s) = \theta_T. \quad \text{(D29)}
\]

**Case 2.1.** $n_{\sigma} = 1$. Eq. (D1m) with $O = \sigma$ and $\sigma^{-1}(i) = (r_1, r_2, r_3; 0)$ yields

\[
-(r_2 + r_3)\theta_{T_1T} + \varphi_T(1) + \varphi_T(0) = \theta_{\sigma T}, \quad \text{(D30)}
\]

which forces $\theta_{T_1T} = 0$. We also have

\[
2\varphi_T(2) = \theta_{\sigma T}, \quad \text{(D31a)}
\]
\[
2\varphi_T(3) = \theta_{\sigma T}, \quad \text{(D31b)}
\]

with $\sigma^{-1}(i) = (r_1, r_2, r_3; s = 2, 3)$.

Eq. (D1n) then yields

\[
2\varphi_T(s) = \theta_T. \quad \text{(D32)}
\]

**Case x.2.** $n_T = 1$.

**Case 1.2.** $n_{\sigma} = 0$. Eq. (D1m) with $O = T_1, T_2, T_3$ yields

\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} + \varphi_T(0, r_2, r_3; s), \quad \text{(D33a)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_2\theta_{T_2T} + \varphi_T(r_1, 0, r_3; s), \quad \text{(D33b)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_3\theta_{T_3T} + \varphi_T(r_1, r_2, 0; s), \quad \text{(D33c)}
\]

which together imply

\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} - r_2\theta_{T_2T} - r_3\theta_{T_3T} + \varphi_T(s). \quad \text{(D34)}
\]

Eq. (D1m) with $O = C_3$ and $i = (r_1, r_2, r_3; 0)$ yields

\[
(r_1 - r_2)\theta_{T_1T} + (r_2 - r_3)\theta_{T_2T} + (r_3 - r_1)\theta_{T_3T}
- 2\varphi_C(0) = \theta_{C_3T}. \quad \text{(D35)}
\]

which implies $\theta_{T_1T} = \theta_{T_2T} = \theta_{T_3T}$. We also have

\[
-2\varphi_C(1) - \varphi_T(1) + \varphi_T(3) = \theta_{C_3T}, \quad \text{(D36a)}
\]
\[
-2\varphi_C(2) - \varphi_T(2) + \varphi_T(1) = \theta_{C_3T}, \quad \text{(D36b)}
\]
\[
-2\varphi_C(3) - \varphi_T(3) + \varphi_T(2) = \theta_{C_3T}, \quad \text{(D36c)}
\]

with $i = (r_1, r_2, r_3; s = 1, 2, 3)$.

Eq. (D1m) with $O = \sigma$ and $i = (r_1, r_2, r_3; 0)$ yields

\[
(2r_1 + r_2 + r_3)\theta_{T_1T} - 2\varphi_C(0) + \varphi_T(0) + \varphi_T(1) = \theta_{\sigma T}. \quad \text{(D37)}
\]

which implies $\theta_{T_1T} = 0$. We also have

\[
-2\varphi_C(1) - \varphi_T(1) + \varphi_T(0) = \theta_{\sigma T}, \quad \text{(D38a)}
\]
\[
-2\varphi_C(2) - \varphi_T(2) + \varphi_T(2) = \theta_{\sigma T}, \quad \text{(D38b)}
\]
\[
-2\varphi_C(3) - \varphi_T(3) + \varphi_T(3) = \theta_{\sigma T}, \quad \text{(D38c)}
\]

with $i = (r_1, r_2, r_3; s = 1, 2, 3)$.

Eq. (D1n) then yields

\[
\pi = \theta_T, \quad \text{(D39)}
\]

where $\pi$ comes from $(i_{\tau_1})^2 = -1$.

**Case 2.2.** $n_{\sigma} = 1$. Eq. (D1m) with $O = T_1, T_2, T_3$ yields

\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} + \varphi_T(0, r_2, r_3; s), \quad \text{(D42a)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_2\theta_{T_2T} + \varphi_T(r_1, 0, r_3; s), \quad \text{(D42b)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_3\theta_{T_3T} + \varphi_T(r_1, r_2, 0; s), \quad \text{(D42c)}
\]

which further lead to

\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} - r_2\theta_{T_2T} - r_3\theta_{T_3T} + \varphi_T(s), \quad \text{(D43a)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} - r_2\theta_{T_2T} - r_3\theta_{T_3T} + \varphi_T(s) - 2r_2\theta_{T_2T}, \quad \text{(D43b)}
\]
\[
\phi_T(r_1, r_2, r_3; s) = -r_1\theta_{T_1T} - r_2\theta_{T_2T} - r_3\theta_{T_3T} + \varphi_T(s) - 2r_1\theta_{T_1T}, \quad \text{(D43c)}
\]

The right hand sides of these six equations must be equal to each other, which only allows $p_{T_1T} = 0.2$ in $\theta_{T_2T} = 2p_{T_2T}\pi/4$. We redefine $p_{T_2T}$ such that $\theta_{T_2T} = p_{T_2T}\pi$, $p_{T_2T} \in \{0, 1\}$.

Eq. (D1m) with $O = C_3$ and $i = (r_1, r_2, r_3; 0)$ yields

\[
(r_1 - r_2)\theta_{T_1T} + (r_2 - r_3)\theta_{T_2T} + (r_3 - r_1)\theta_{T_3T}
- 2\varphi_C(0) = \theta_{C_3T}. \quad \text{(D42)}
\]

which implies $\theta_{T_1T} = \theta_{T_2T} = \theta_{T_3T}$. We also have

\[
-2\varphi_C(1) - \varphi_T(1) + \varphi_T(3) = \theta_{C_3T}, \quad \text{(D43a)}
\]
\[
-2\varphi_C(2) - \varphi_T(2) + \varphi_T(1) = \theta_{C_3T}, \quad \text{(D43b)}
\]
\[
-2\varphi_C(3) - \varphi_T(3) + \varphi_T(2) = \theta_{C_3T}, \quad \text{(D43c)}
\]

with $i = (r_1, r_2, r_3; s = 1, 2, 3)$.

Eq. (D1m) with $O = \sigma$ and $i = (r_1, r_2, r_3; 0)$ yields

\[
(r_2 + r_3)\theta_{T_2T} - 2\varphi_C(0) + \varphi_T(0) + \varphi_T(1) = \theta_{\sigma T}. \quad \text{(D44)}
\]
which forces \( \theta_{T_1 T} = 0 \). We also have

\[
-2 \varphi_\sigma (1) + \varphi_T (1) + \varphi_T (0) = \theta_{\sigma T}, \quad (D45a)
-2 \varphi_\sigma (2) + 2 \varphi_T (2) = \theta_{\sigma T}, \quad (D45b)
-2 \varphi_\sigma (3) + 2 \varphi_T (3) = \theta_{\sigma T}, \quad (D45c)
\]

with \( i = (r_1, r_2, r_3; s = 1, 2, 3) \).

Eq. (D1n) then yields

\[
\pi = \theta_T, \quad (D46)
\]

where \( \pi \) comes from \( (i \tau_1)^2 = -1 \).

### a. Grand Summary

It is worthwhile to recollect the results obtained thus far before we proceed to gauge fixings. They are (14a)-(14f) shown in the main text and the “sublattice constraints” listed below in a case by case manner.

\[
3 \varphi_{C_1} (0) = \theta_{C_1}, \quad (D47a)
\sum_{s=1}^{3} \varphi_{C_1} (s) = \theta_{C_1}. \quad (D47b)
\]

#### Case 1. \( n_\sigma = 0 \).

\[
\theta_{T_1 T} = p_{T_1 T} \pi, p_{T_1 T} \in \{0, 1\}, \quad (D48a)
\theta_{\sigma T_2} = p_{\sigma T_2} \pi, p_{\sigma T_2} \in \{0, 1\}, \quad (D48b)
\varphi_\sigma (1) + \varphi_\sigma (0) = \theta_\sigma, \quad (D48c)
2 \varphi_\sigma (2) = \theta_\sigma, \quad (D48d)
2 \varphi_\sigma (3) = \theta_\sigma, \quad (D48e)
\sum_{s=0}^{3} [\varphi_\sigma (s) + \varphi_{C_1} (s)] = \theta_{\sigma C_1}. \quad (D48f)
\]

#### Case 1.1. \( n_T = 0 \).

\[
\varphi_T (0) = \varphi_T (1) = \varphi_T (2) = \varphi_T (3), \quad (D49a)
2 \varphi_T (s) = \theta_T. \quad (D49b)
\]

#### Case 1.2. \( n_T = 1 \).

\[
-2 \varphi_{C_1} (0) = \theta_{C_1} T, \quad (D50a)
-2 \varphi_{C_1} (1) - \varphi_T (1) + \varphi_T (3) = \theta_{C_1} T, \quad (D50b)
-2 \varphi_{C_1} (2) - \varphi_T (2) + \varphi_T (1) = \theta_{C_1} T, \quad (D50c)
-2 \varphi_{C_1} (3) - \varphi_T (3) + \varphi_T (2) = \theta_{C_1} T, \quad (D50d)
-2 \varphi_\sigma (0) - \varphi_T (0) + \varphi_T (1) = \theta_\sigma T, \quad (D50e)
-2 \varphi_\sigma (1) - \varphi_T (1) + \varphi_T (0) = \theta_\sigma T, \quad (D50f)
-2 \varphi_\sigma (2) = \theta_\sigma T, \quad (D50g)
-2 \varphi_\sigma (3) = \theta_\sigma T. \quad (D50h)
\]

#### Case 2. \( n_\sigma = 1 \).

\[
\theta_{\sigma T_2} = p_{\sigma T_2} \pi, p_{\sigma T_2} \in \{0, 1\}, \quad (D51a)
\varphi_\sigma (0) = \varphi_\sigma (1), \quad (D51b)
-2 \varphi_\sigma (0) - \varphi_{C_1} (1) + \varphi_\sigma (3) + \varphi_{C_1} (3) \quad (D51c)
\varphi_\sigma (2) - \varphi_{C_1} (2) + \varphi_\sigma (1) + \varphi_{C_1} (0) \quad (D51d)
\]

#### Case 2.1. \( n_T = 0 \).

\[
\theta_{T_1 T_1} = \frac{2 p_{T_1 T_1} \pi}{4}, p_{T_1 T_1} \in \{0, 1\}, \quad (D52a)
\varphi_T (1) + \varphi_T (0) = \theta_{\sigma T}, \quad (D52b)
2 \varphi_T (2) = \theta_{\sigma T}, \quad (D52c)
\varphi_T (3) = \theta_{\sigma T}, \quad (D52d)
\varphi_T (1) = \varphi_T (2) = \varphi_T (3), \quad (D52e)
2 \varphi_T (s) = \theta_T. \quad (D52f)
\]

#### Case 2.2. \( n_T = 1 \).

\[
\theta_{T_1 T_1} = \frac{2 p_{T_1 T_1} \pi}{4}, p_{T_1 T_1} \in \{0, 1\}, \quad (D53a)
-2 \varphi_{C_1} (0) = \theta_{C_1} T, \quad (D53b)
-2 \varphi_{C_1} (1) - \varphi_T (1) + \varphi_T (3) = \theta_{C_1} T, \quad (D53c)
-2 \varphi_{C_1} (2) - \varphi_T (2) + \varphi_T (1) = \theta_{C_1} T, \quad (D53d)
-2 \varphi_{C_1} (3) - \varphi_T (3) + \varphi_T (2) = \theta_{C_1} T, \quad (D53e)
-2 \varphi_\sigma (0) + \varphi_T (0) + \varphi_T (1) = \theta_\sigma T, \quad (D53f)
-2 \varphi_\sigma (1) + \varphi_T (1) + \varphi_T (0) = \theta_\sigma T, \quad (D53g)
-2 \varphi_\sigma (2) + 2 \varphi_T (2) = \theta_\sigma T, \quad (D53h)
-2 \varphi_\sigma (3) + 2 \varphi_T (3) = \theta_\sigma T. \quad (D53i)
\]

### b. Gauge fixing

We now gauge fix \( \varphi_\sigma (s) = \varphi_X (0, 0, 0; s), s = 0, 1, 2, 3 \), subject to the constraints listed above. First and foremost, we use the IGG freedom of \( G_{C_1} \) to fix \( \varphi_{C_1} (0) = 0 \). It follows from (D47a) that \( \theta_{C_1} = 0 \). Using (C2) and (D47b), we perform a sublattice dependent gauge transformation \( W_3 \)

\[
W_{0,3} = 1, W_1 = e^{-i \varphi_{C_1} (1) r_1}, W_2 = e^{-i \varphi_{C_1} (1) + \varphi_{C_1} (2) r_1},
\]

to fix \( \varphi_{C_1} (1, 2, 3) = 0 \).

#### Case 1. \( n_\sigma = 0 \). We use the IGG freedom of \( G_\sigma \) to fix \( \varphi_\sigma (2) = 0 \). (D48d) then implies \( \theta_\sigma = 0 \), and (D48e) subsequently implies \( \varphi_\sigma (3) = q_3 \pi, q_3 \in \{0, 1\} \). We perform a sublattice dependent gauge transformation \( W_0 = \exp [i \varphi_\sigma (0) r_1] \). \( W_{1,2,3} = 1 \) to fix \( \varphi_\sigma (0, 1) = 0 \), without affecting any previously fixed gauge. (D48f) then yields \( \theta_{\sigma C_1} = q_3 \pi \), so we rename \( q_3 \) as \( p_{\sigma C_1} \).

#### Case 1.1. \( n_T = 0 \). We use the IGG freedom of \( G_T \) to fix \( \varphi_T (0) = 0 \). By (D49a), we also have \( \varphi_T (1, 2, 3) = 0 \).

#### Case 1.2. \( n_T = 1 \). Eq. (D50a)-(D50d) yield \( \theta_{C_1 T} = 0 \) and

\[
- \varphi_T (1) + \varphi_T (3) = 0, \quad (D54a)
- \varphi_T (2) + \varphi_T (1) = 0, \quad (D54b)
- \varphi_T (3) + \varphi_T (2) = 0. \quad (D54c)
\]
or \( \varphi_T(1) = \varphi_T(2) = \varphi_T(3) \). Eq. (D50g) or (D50h) imply \( \theta_{\sigma_T} = 0 \), which together with (D50e) or (D50f) imply \( \varphi_T(0) = \varphi_T(1) \). We use the IGG freedom of \( G_T \) to fix \( \varphi_T(0) = 0 \), and it follows that \( \varphi_T(1, 2, 3) = 0 \).

**Case 2.** \( n_{\sigma} = 1 \). We perform a sublattice dependent gauge transformation \( W_0 = \exp[-i(\varphi_{\sigma}(2) - \varphi_{\sigma}(0))r_3] \), \( W_{1,2,3} = 1 \), such that \( \varphi_{\sigma}(0, 1) = \varphi_{\sigma}(2) \), without affecting any previously fixed gauge. Then, we use the IGG freedom of \( G_{\sigma} \) to fix \( \varphi_{\sigma}(2) = 0 \), and rename \( \varphi_{\sigma}(3) - \varphi_{\sigma}(2) \) as \( \varphi_{\sigma}(3) \). Eq. (D51c) implies \( \varphi_{\sigma}(3) = p_{\sigma c} \gamma, p_{\sigma c} \in \{0, 1\} \).

**Case 2.1.** \( n_T = 0 \). We use the IGG freedom of \( G_T \) to fix \( \varphi_T(1) = 0 \). It follows from (D52e) that \( \varphi_T(2, 3) = 0 \). We also have \( \theta_{\sigma_T} = 0 \) and \( \theta_T = 0 \) from (D52c) and (D52f) respectively. (D52b) then implies \( \varphi_T(0) = 0 \).

**Case 2.2.** \( n_T = 1 \). Eqs. (D53b)-(D53e) yield \( \theta_{C_1 T} = 0 \) and

\[-\varphi_T(1) + \varphi_T(3) = 0, \quad \varphi_T(2) + \varphi_T(1) = 0, \quad \varphi_T(3) + \varphi_T(2) = 0, \]

or \( \varphi_T(1) = \varphi_T(2) = \varphi_T(3) \). We use the IGG freedom of \( G_T \) to fix \( \varphi_T(1) = 0 \), and it follows that \( \varphi_T(2, 3) = 0 \). Eq. (D53h) or (D53i) imply \( \theta_{\sigma_T} = 0 \), and (D53j) or (D53g) subsequently imply \( \varphi_T(0) = 0 \).

We will exclude the solutions with \( n_T = 0 \) (Cases 1.1 and 2.1) because they have \( G_T(i) = 1 \) for all sites \( i \), which force \( u_{i j} = 0 \) for any pair of sites \( i \) and \( j \) by (10). These solutions, which lead to vanishing mean field ansatzes and thus a zero Hamiltonian, are unphysical. Let us count the remaining solutions. Each of 1.2 and 2.2 has three \( \mathbb{Z}_2 \) variables \( p_{T_1 T_2}, p_{T_2 T_2}, \) and \( p_{T_1 T_1} \). Therefore, we have in total \( 2 \times 2^3 = 16 \) gauge inequivalent solutions, i.e., 16 possible \( U(1) \) spin liquids. They are listed in Table I.

### 1. Relation to isotropic lattice

We first note that Ref. 48 (see also Ref. 47) have chosen a coordinate system in which the unit cell at the origin, \( (r_1, r_2, r_3) = (0, 0, 0) \), is associated with a down tetrahedron, whereas in our coordinate system it is associated with an up tetrahedron. However, it can be straightforwardly shown that if instead we choose a down tetrahedron at the origin and adopt the primitive translation vectors and sublattice labelings of Ref. 48, the expressions (4a)-(4e) of the action of the five space group generators remain invariant, so exactly the same PSG solutions (see Table I) for the breathing pyrochlore lattice will follow. This observation allows us to assume the coordinate system of Ref. 48 in this appendix and directly compare our PSG solutions to those of Ref. 48.

The sixfold rotonversion \( \mathcal{C}_6 = I C_3 \) and the twofold nonsymmetric screw \( S \) act on a generic site with coordinates \( (r_1, r_2, r_3; s) \) as [47, 48]

\[ \mathcal{C}_6 : (r_1, r_2, r_3; s) \longrightarrow (-r_3 - \delta_{s,3}, -r_1 - \delta_{s,1}, -r_2 - \delta_{s,2}; \mathcal{C}_6(s)), \quad (D56a) \]

\[ S : (r_1, r_2, r_3; s) \longrightarrow (-r_1 - \delta_{s,1}, -r_2 - \delta_{s,2}, r_1 + r_2 + r_3 + 1 - \delta_{s,0}; S(s)). \quad (D56b) \]

When \( s = 0, 1, 2, 3 \), \( \mathcal{C}_6(s) = 0, 2, 3, 1 \) and \( S(s) = 3, 1, 2, 0 \). Recall that, for \( U(1) \) spin liquids, the gauge transformation \( G_X \) associated with a symmetry operator \( X \) has the specific form (13). The distinct \( U(1) \) spin liquids that respect both the \( Fd\overline{3}m \) space group of the regular pyrochlore lattice and the time reversal symmetry are characterized by the PSGs in (20a)-(20c), (21a)-(21e), and Table I of Ref. 48, which we quote below,

\[ \phi_T(1, r_1, r_2, r_3; s) = 0, n_{T_1} = 0, \quad (D57a) \]

\[ \phi_T(1, r_1, r_2, r_3; s) = -\chi_1 r_1, n_{T_2} = 0, \quad (D57b) \]

\[ \phi_T(1, r_1, r_2, r_3; s) = \chi_1 (r_1 - r_2), n_{T_1} = 0, \quad (D57c) \]

\[ \phi_{S_1}(r_1, r_1, r_2, r_3; s) = -\chi_1 r_1 (r_2 - r_3) - [2 + (\delta_{s,2} - \delta_{s,3})] \chi_1 r_1 + \delta_{s,2} \chi_1 r_1 + \phi_{S_6}(s), \quad (D57d) \]

\[ \phi_S(r_1, r_2, r_3; s) = \chi_1 \left[ \frac{r_1 r_3 + 1}{2} - \frac{r_2 (r_2 + 1)}{2} - r_1 r_2 \right] + (2 + \delta_{s,1} - \delta_{s,2}) \chi_1 r_3 + r_1 \]

\[ + (2 \delta_{s,1} - \delta_{s,2}) \chi_1 r_2 + \phi_S(s), \quad (D57e) \]

\[ \phi_T = 0, n_T = 1. \quad (D57f) \]

with

\[ \varphi_{\mathcal{C}_6}(0, 1, 2) = 0, \varphi_{\mathcal{C}_6}(3) = \chi_1, \]

\[ \varphi_S(0, 2) = 0, \varphi_S(1) = \chi_1, \varphi_S(3) = \chi_1, \]

\[ \chi_1, \chi_1 \mathcal{C}_6 S \in \{0, \pi\}. \quad (D58) \]

The two choices in each of \( n_{\mathcal{C}_6}, n_S, \chi_1, \) and \( \chi_1 \mathcal{C}_6 S \) gives rise to \( 2^4 = 16 \) \( U(1) \) spin liquids. Notice that every \( \chi_1 \) is an integer multiple of \( \pi \), which is invariant under a sign change.

The point group generators of the breathing pyrochlore lattice are related to the regular one by \( C_3 \equiv \mathbb{Z}_3 \) and \( \sigma = C_1 \Sigma C_3^{-1} \), where \( \Sigma \equiv SI \) is a reflection across the plane perpendicular to \( [110] \) (c.f. \( \sigma \)). For convenience of later calculations, we note that

\[ I : (r_1, r_2, r_3; s) \longrightarrow (-r_1 - \delta_{s,1}, -r_2 - \delta_{s,2}, -r_3 - \delta_{s,3}; s), \quad (D59a) \]

\[ \Sigma : (r_1, r_2, r_3; s) \longrightarrow (r_1, r_2, -r_1 - r_2 - r_3; \Sigma(s)), \quad (D59b) \]

where \( \Sigma(s) = 3, 1, 2, 0 \) for \( s = 0, 1, 2, 3 \). We also note that \( \mathcal{C}_6^3 = I \) [47, 48].

From (14a)-(14f), Table I, and from (D57a)-(D57f), and (D58), we see that \( G_T \) of both PSGs agree, while \( G_{T_1} \), \( G_{T_2} \), and \( G_{T_1} \) agree upon the identification of \( \theta_{T_1 T_2} \) with \( \chi_1 \). For the point group generators, let the gauge transformation parts of \( (G_{\mathcal{C}_6} \mathcal{C}_6^3)^4 = (G_{\mathcal{C}_6} \mathcal{C}_6^3 G)_S (G_{\mathcal{C}_6} \mathcal{C}_6^3)^{-1} \) by \( G_{\chi_1} \) and \( G_{\sigma} \) [which are expressed in terms of \( \phi_{S_{\chi_1}, \sigma} \) and \( n_{\chi_1, \sigma} \) according to (13)] respectively.
Let $i = (r_1, r_2, r_3; s)$. We start with
\[
\phi_{C_6}(i) = \phi_{C_6}(i) + \phi_{C_6}(C_6^{-1}(i)) + n_{C_6} \pi \\
= \chi_1 r_1(r_1 - r_2) + \chi_1^2 + \chi_3 + n_{C_6} \pi, \quad (D60)
\]
which further yields
\[
\phi'_{C_6}(i) = \phi_{C_6}(i) + \phi_{C_6}(C_6^{-2}(i)) \\
= -\chi_1 r_1(r_2 - r_3) + \chi_1 \delta s_1 + \delta s_2, n'_{C_6} = 0. \quad (D61)
\]
The uniform additive factor $n_{C_6} \pi$ in (D60) can be further dropped without penalty. We proceed to calculate
\[
\phi_F(i) = \phi_{C_6}(i) + \phi_{C_6}(C_6^{-1}(i)), \\
n_F = n_{C_6} \in \{0, 1\}, \quad (D62)
\]
which gives
\[
\phi_F(r_1, r_2, r_3; 0) = 0, \quad (D63a)
\phi_F(r_1, r_2, r_3; 1) = \chi_1 (r_2 - r_3 + 1), \quad (D63b)
\phi_F(r_1, r_2, r_3; 2) = \chi_3 (r_3 + 1), \quad (D63c)
\phi_F(r_1, r_2, r_3; 3) = \chi_1. \quad (D63d)
\]
Then, we calculate
\[
\phi_S(i) = \phi_S(i) + \phi_F(S^{-1}(i)), \\
= \chi_1 \left[ \frac{r_1(r_1 + 1)}{2} - \frac{r_2(r_2 + 1)}{2} - r_1 r_2 \right] \\
+ \chi_1 + \chi_{S_6} \delta s_1, \\
n_S = (n_S + n_F) \bmod 2 \in \{0, 1\}, \quad (D64)
\]
Finally, we calculate
\[
\phi'_S(i) = \phi_{C_5}(i) + \phi_S(C_5^{-1}(i)) - \phi_{C_5}^2 \quad (D65)
\]
We further add $\chi_{C_6} S - \chi_1$ to $\phi'_S(i)$ uniformly, and then perform a sublattice dependent gauge transformation $W_0 = \exp(i \chi_{C_6} S r_3), W_1 = \exp(i \chi_1 r_3), W_{2,3} = 1$ so that
\[
\phi'_{C_5}(r_1, r_2, r_3; s) = -\chi_1 r_1(r_2 - r_3), n'_{C_5} = 0, \quad (D66a)
\]
\[
\phi'_{C_5}(r_1, r_2, r_3; s) = -\chi_1 \left[ \frac{r_2(r_2 - 1)}{2} - \frac{r_3(r_3 - 1)}{2} + r_2 r_3 \right] \\
+ \chi_{C_6} \delta s_3, n'_{C_5} \in \{0, 1\}, \quad (D66b)
\]
without affecting other gauges.

We can now compare (14d) and (14e) with (D66a) and (D66b) respectively, and identify $\chi_{C_6} S$ with $p_{C_5}$ in Table I. We conclude that the 16 $U(1)$ spin liquids of the regular pyrochlore lattice [48] are continuously connected to the 8 $U(1)$ spin liquids of the breathing pyrochlore lattice with $\theta_{C_5} = 0$, in the fashion of a two-to-one mapping defined by $n_{C_5} = (n_{C_6} + n_S) \bmod 2$. We see that the additional symmetries of the regular pyrochlore lattice, on the one hand forces $\theta_{C_5} = 0$, on the other hand give rise to two $Z_2$ variables $n_{C_6}$ and $n_S$ instead of the sole $n_{C_5}$ in the breathing pyrochlore lattice. Therefore, while the number of $U(1)$ spin liquids are same in both lattices, they are not in a one-to-one correspondence.

Appendix E: Analytical solution of mean field theory

For the $U(1)_{\pi}$ state, diagonalizing $d_k$ yields the 16 eigenvalues $\epsilon_0(k) \{8\}, \epsilon_{++}(k) \{2\}, \epsilon_{+-}(k) \{2\}, \epsilon_{--}(k) \{2\}, \epsilon____(k) \{2\}$, where each curly bracket indicates the number of times that the corresponding eigenvalue appears,
\[
\epsilon_0(k) = -\chi_1 + \chi_2, \quad (E1a)
\epsilon_{++}(k) = \chi_1 + \chi_2 + \left[ 4 \chi_1^2 + 4 \chi_2^2 - 4 \chi_1 \chi_2 \right] \quad (E1b)
\]
\[
\epsilon_{--}(k) = \chi_1 + \chi_2 + \left[ 4 \chi_1 + 4 \chi_2^2 - 4 \chi_1 \chi_2 \right] \quad (E1c)
\]
and
\[
f(k) = -\cos(2k_1 - k_2) + \cos(k_2 - k_3) + \cos(k_3 - 2k_1) \quad (E2)
\]
Note that the function $f(k)$ defined here has nothing to do with that in Sec. IV B. We merely recycled the notation. $f(k)$ has maximum and minimum of 2 and $-6$ respectively.

The analysis proceeds very much along the same lines as in Sec. IV B, so we will only outline the steps and omit the details. Let us choose $\chi_1 > 0$ without loss of generality. We first assume that $\chi_2 > 0$. One can show that $\epsilon_{++}(k), \epsilon_{--}(k) > \epsilon_0(k)$ for all $k$, so the dispersing bands are well separated from, and higher in energy than, the flat bands throughout the Brillouin zone. Then, filling the lower half of the energy eigenstates, i.e., all the flat bands, the total energy of a system with $N \times N$ sites is given by
\[
E_S = \sum_k \left[ 16 \epsilon_0(k) + 24 \left( \frac{J_1}{4} \chi_1 + \frac{J_2}{4} \chi_2 \right) \right] \\
= \sum_k \left[ 16 \left( \frac{J_1}{4} \chi_1 + \frac{J_2}{4} \chi_2 \right) + 24 \left( \frac{J_1}{4} \chi_1 + \frac{J_2}{4} \chi_2 \right) \right]. \quad (E3)
\]
Next, we assume that $\chi_2 < 0$, and without loss of generality $|\chi_1| > |\chi_2|$. In this case, the dispersing bands are well separated from the flat bands, with the $\pm \chi_1 \chi_2$ bands lying above (below) the flat bands. One can show that, by filling half of the lower energy eigenstates, i.e., $2 -- \chi_2$ bands, $2 + -- \chi_2$ bands, and 12 of the flat bands, the total energy $E_A$ thus obtained is strictly
greater than $E_S$ in (E3). Therefore, we can exclude the case with $\chi_2 < 0$.

Finally, minimizing (E3) with respect to $\chi_1$ and $\chi_2$ yields $\chi_{1,2} = -1/3$ and the ground state energy per site $-(J_1 + J_2)/24$.

**Appendix F: Low energy effective field theory**

In this appendix, we provide details of the field theoretic treatment of the $U(1)_0$ state, from which we derive the low temperature heat capacity. Starting from (30a) and (30b), we integrate out the spinons to obtain an effective action (of the spin liquid, in which quadratically dispersing density fluctuations [65, 67, 74] so that it can be neglected.

We first calculate the diagram in Fig. 5c, which we call $\Pi^{\alpha}_{xx}(q)$. Let us choose a coordinate system in which $q$ aligns in the $z$ direction, so that $a_z = 0$ by the Coulomb gauge. For the transverse components of the gauge field, $\Pi^{\alpha}_{ij}$ is nonzero only when $ij = xx$ and $yy$,

$$\Pi^{\alpha}_{xx}(q) = \frac{-2}{m^2 \beta V} \sum_{kn} G(k + q) G(k) k_x^2$$

$$= \frac{-2}{m^2 \beta} \int_k \frac{1}{i(\omega_n + \nu_1) - \epsilon_{kq} + (\omega_n - \epsilon_k)^2},$$

(F3)

where the minus sign comes from the fermion loop, the factor of 2 is due to the two spin species, $G(k)$ is the free fermion Green function, $\int_k$ is the shorthand notation for $\int d^3k/(2\pi)^3$, and $\epsilon_k = |k|^2/2m$. The corresponding expression for $\Pi^{\alpha}_{yy}(q)$ is obtained with the replacement $x \rightarrow y$. Using the contour integral method [88, 95, 96] to perform the sum over $\omega_n$,

$$\Pi^{\alpha}_{yy}(q) = \frac{-2}{m^2} \int_k \frac{|f(\epsilon_{kq}) - f(\epsilon_k)|^2}{\epsilon_{kq} + (2k \cdot q + |q|^2)/2m},$$

$$= \frac{-2}{m^2} \int_k \frac{|f(\epsilon_{kq=2}) - f(\epsilon_{kq=2})|^2}{\epsilon_{kq} + (2k \cdot q/m)},$$

(F4)

where $f(\epsilon) = 1/[\exp(\beta \epsilon) + 1]$ is the Fermi Dirac distribution, and we have shifted $k$ by $-q/2$ in the second line to obtain a more symmetric expression. It is very difficult if not impossible to evaluate the integral, so we study the small $|q|$ limit. More precisely, we expand

$$f(\epsilon_{kq=2}) = \sum_{n=0}^{\infty} \frac{1}{n!} \frac{\partial^n f(\epsilon_k)}{\partial \epsilon_k^n} \left( \frac{k \cdot q + |q|^2}{2m} \right)^n$$

(F5)

with $|q|/\sqrt{mT} \ll 1$ as the small parameter. The inclusion of $T$ in defining the small parameter is important because each derivative of $f(\epsilon)$ with respect to $\epsilon$ brings down a factor of $1/T$, so we require higher powers of $|q|/\sqrt{mT}$ to be less important than lower powers to justify a truncation of the expansion (F6) at finite $n$. (There should also be a constraint $|k| \lesssim \sqrt{mT}$. Nevertheless, large momentum modes are thermally suppressed by the Fermi Dirac distribution function, so we
will take the upper limits of the \(|k|\) integrals below to infinity.)
Note that this is unlike the case of a spinon Fermi surface,
where one can expect to take \(q\) against the Fermi wavevector \(k_{F}\),
that is, taking \(|q|/k_{F} \ll 1\) as the small parameter, without a direct
comparison with \(T\), and use a zero temperature approximation
for \(f(e)\) which is effectively a step function \([74, 88]\).

Let us first examine the case when \(l \neq 0\). Expanding
the numerator in (F5) to the lowest nontrivial order in \(q\),
\[
\Pi_{\tau}^{\ell}(q) \approx -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q) m \times k^2}{x_{y}}

= -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q/m)^2}{x_{y}^2 + (k \cdot q/m)^2} k^2 \phi.
\]
(F7)

Since \(v_{f} \equiv 2\pi l T\) and \(k \cdot q/m \ll v_{f}\) for \(l \neq 0\), which allows us to approximate
\[
\Pi_{\tau}^{\ell}(q, v_{f}) \approx -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q)}{m v_{f}} \bigg[ k_x \bigg]^2

= -\frac{2}{m^4} \int_{0}^{2\pi} \frac{2\pi}{(2\pi)^3} \int_{0}^{2\pi} d\phi \cos^{2} \phi \int_{-1}^{1} d(\cos \theta) \sin^{2} \theta \cos^{2} \theta

\times \int_{0}^{\infty} d|k| \frac{|k|^{2} \partial f(e_{k})}{\partial e_{k}} \frac{|q|^{2}}{v_{f}^2} \int_{0}^{\infty} dx \frac{x e^{x^2}}{(e^{x^2} + 1)^2} \approx c_{3} T^{3/2} \phi.
\]
(F8)
The dimensionless \(x\) integral evaluates to 1.44 (to three significant figures).

On the other hand, the diagram in Fig. 5d, which we call \(\Pi^{0}\), is much easier to evaluate than \(\Pi^{\ell}\). As before, \(\Pi_{ij}^{0}\) is nonzero only when \(ij = xx\) and \(yy\). Contracting the fermion
lines forces \(q = q'\), so
\[
\Pi_{\tau}^{0}(q) = -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q)}{m v_{f}} \bigg[ k_x \bigg]^2

= -\frac{8\pi \sqrt{2m}}{2\pi^3} \int_{0}^{\infty} dx \frac{x e^{x^2}}{(e^{x^2} + 1)^2} \approx c_{2} T^{3/2} \phi.
\]
(F9)

which is independent of \(q\). The dimensionless \(x\) integral evaluates to 0.339 (to three significant figures). Adding up (F8)
and (F9) then yields (32c).

Next, we examine the case when \(l = 0\). Expanding
the numerator in (F5) to the lowest nontrivial order in \(q\),
\[
\Pi_{\tau}^{\ell}(q, v_{f}) = -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q) m \times k^2}{x_{y}}

\approx -\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{(k \cdot q/m)^2}{x_{y}^2 + (k \cdot q/m)^2} k^2 \phi.
\]
(F10)

To evaluate the terms in the square bracket, we note that
\[
\frac{\partial f(e)}{\partial e} = -\frac{\partial f(e)}{\partial T}.
\]
(F11)

which allows us to replace the derivative with respect to energy
by one with respect to temperature, and pull the temperature

The first derivative term evaluates to
\[
\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} |k|^2 \sin^{2} \cos^{2} \phi

= -\frac{4}{m^2} \int_{0}^{\infty} dx \frac{x e^{x^2}}{(e^{x^2} + 1)^2} = c_{2} T^{3/2} \phi.
\]

(F12)

which is same as (F9) apart from a minus sign. The second
derivative term evaluates to
\[
-\frac{2}{m^2} \int_{k} \frac{\partial f(e_{k})}{\partial e_{k}} \frac{|q|^2}{8m} |k|^2 \sin^{2} \cos^{2} \phi

= -\frac{|q|^2}{m} \int_{0}^{\infty} dx \frac{x e^{x^2}}{(e^{x^2} + 1)^2} \approx c_{2} T^{3/2} \phi.
\]

(F13)

The dimensionless \(x\) integral evaluates to 0.536 (to three significant figures). Similarly, the third derivative term evaluates to
\[
-\frac{2}{m^2} \int_{k} \frac{\partial^{3} f(e_{k})}{\partial e_{k}^3} \frac{|q|^2}{24m^2} |k|^4 \sin^{2} \cos^{2} \phi

= \frac{\sqrt{2/m}}{8\pi^2 T^{3/2} |q|^2} \int_{0}^{\infty} dx \frac{x e^{x^2}}{(e^{x^2} + 1)^2} \approx c_{2} T^{3/2} \phi.
\]

(F14)

On the other hand, \(\Pi^{0}(q)\) is still given by (F9), which
cancels out (F12). We are left with (F13) and (F14), and
adding them up yields (32b).

We can then write down the effective Lagrangian of the \(U(1)\)
gauge field \((32a)\), the corresponding partition function \((33a)\),
and the free energy \((33b)\). We now calculate the free energy
contribution \((34b)\) from the \(l \neq 0\) modes, as follows. Changing
the summation over \(q\) to an integral, the free energy density
reads
\[
f_{\text{dyn}} \equiv \frac{F_{\text{dyn}}}{V} = T \sum_{l \neq 0} \int_{q} \ln \left( c_{2} T^{3/2} - c_{3} T^{5/2} \frac{|q|^2}{v_{f}^2} \right)

\approx T \sum_{l \neq 0} \int_{q} \ln \left( c_{2} T^{3/2} - c_{3} T^{5/2} \frac{|q|^2}{v_{f}^2} \right),
\]
(F15)

where \(c_{3}' \equiv c_{3}/c_{2}\). Since we have assumed \(|q| \ll \sqrt{mT}\) from
the beginning, the \(q\) integral should have an upper limit of
the form \(x_{0} \sqrt{mT}\) \([65]\), where \(x_{0} < 1\) is some small number,
whose precise value is not important for the current analysis.
Performing the integration over \(q\) yields
\[
f_{\text{dyn}} = \frac{8\pi T}{(2\pi)^3} \sum_{l \neq 0} \left[ \left( x_{0} \sqrt{mT} \right)^{3} \ln \left( c_{2} T^{3/2} - c_{3} T^{5/2} \frac{(x_{0} \sqrt{mT})^{5}}{5v_{f}^2} \right) \right].
\]
(F16)
The first term is independent of \( \nu_f \), so the summation over natural numbers yields a divergence, which can be regularized using the Riemann zeta function [97],

\[
\zeta(s) = \sum_{n=1}^{\infty} n^{-s} = \frac{1}{\Gamma(s)} \int_0^\infty dx \frac{x^{s-1}}{e^x - 1},
\]

where \( \Gamma(s) = \int_0^\infty dx x^{s-1} e^{-x} \)

is the gamma function. In particular, we use \( \zeta(0) = -1/2 \) and \( \zeta(2) = \pi^2/6 \) for the first and second terms in (F16) respectively, and we obtain

\[
f_{\text{dyn}} = -\frac{8\pi T}{(2\pi)^3} \left[ \frac{(x_0 \sqrt{mT})^3}{6} \ln(c_2T^{3/2}) + c_3' \left( \frac{(x_0 \sqrt{mT})^5}{120T} \right) \right].
\]

We then calculate the free energy contribution (34a) from the \( l = 0 \) mode. The free energy density reads

\[
f_{\text{sta}} = \frac{F_{\text{sta}}}{V} = T \left[ \ln \left( c_1 \sqrt{T} |q|^2 \right) \right] = \frac{4\pi T}{(2\pi)^3} \left[ \frac{(x_0 \sqrt{mT})^3}{3} \ln \left( c_1 x_0^3 mT^{3/2} \right) \right] \left( \frac{2}{3} \right).
\]

Adding up (F19) and (F20) yields

\[
f = f_{\text{sta}} + f_{\text{dy}} = \frac{(x_0 \sqrt{mT})^3}{6\pi^2} \left[ \ln \left( 0.0659 x_0^3 \right) - 0.0567 x_0^2 - \frac{2}{3} \right] T^{5/2}.
\]

With \( x_0 \leq 1 \), the square bracket in (F21) evaluates to a negative number. The volumetric heat capacity is readily obtained as

\[
\frac{C(T)}{V} = -T \frac{\partial^2 f}{\partial T^2} \sim T^{3/2}.
\]

Appendix G: Parton representation of Dzyaloshinskii-Moriya interaction

We first introduce the triplet hopping and pairing channels, \( \hat{E}_{ij} \) and \( \hat{D}_{ij} \) [78–81],

\[
\hat{E}_{ij} = \sum_{a\beta} f_{ia}(\sigma^a)\alpha f_{j\beta}, \quad \text{(G1a)}
\]

\[
\hat{D}_{ij} = \sum_{a\beta} f_{ia} i(\sigma^\gamma\sigma^a)\alpha f_{j\beta}. \quad \text{(G1b)}
\]

Let \( (\lambda, \mu, \nu) \) be a cyclic permutation of \( (x, y, z) \). The Dzyaloshinskii-Moriya interaction in (35) involves spin products in the combination

\[
\pm \frac{|D|}{\sqrt{2}} \left( s_\lambda^\mu s_\nu^\gamma - s_\nu^\mu s_\lambda^\gamma \right), \quad \text{(G2)}
\]

The relation between triplet hopping parameters \( E_{ij} \), which are purely imaginary if nonzero, in the \( U(1)_\mu \) state, \( s \) and \( t \) are the sublattice indices of sites \( s \) and \( t \) respectively. The corresponding relation for the \( U(1)_\nu \) state can be obtained from this table in conjunction with Fig. 2b.

| \( s \) | \( t \) | up tetrahedron | down tetrahedron |
|-------|------|----------------|------------------|
| 0     | 1    | \((0, E^z_0, E^\gamma_0, -E^\gamma_0)\) | \((0, -E^z_0, E^\gamma_0, E^\gamma_0)\) |
| 0     | 2    | \((-E^z_0, 0, E^\gamma_0)\) | \((-E^z_0, 0, E^\gamma_0)\) |
| 0     | 3    | \((E^z_0, 0, -E^\gamma_0, 0)\) | \((E^z_0, 0, -E^\gamma_0, 0)\) |
| 1     | 2    | \((-E^z_0, E^\gamma_0, 0)\) | \((E^z_0, E^\gamma_0, 0)\) |
| 2     | 3    | \((0, E^\gamma_0, E^z_0)\) | \((0, E^\gamma_0, E^z_0)\) |
| 3     | 1    | \((E^z_0, 0, E^\gamma_0)\) | \((E^z_0, 0, E^\gamma_0)\) |

which, with the parton representation of spins (5), can be expressed in terms of products of bond operators,

\[
\left( \frac{|D|}{\sqrt{2}} \right) \left[ (\hat{x}_{ij} \pm i\hat{E}_{ij}^3) (\hat{x}_{ij} \pm i\hat{E}_{ij}^3) \right] \left( \hat{\Delta}_{ij} \pm i\hat{D}_{ij} \right) + \hat{E}_{ij}^\mu \hat{E}_{ij}^\nu + \hat{E}_{ij}^\nu \hat{E}_{ij}^\mu + \hat{D}_{ij}^\mu \hat{D}_{ij}^\nu + \hat{D}_{ij}^\nu \hat{D}_{ij}^\mu , \quad \text{(G3)}
\]

where we have used the identity \( \hat{\Delta}_{ij}^3 \hat{\Delta}_{ij} = -\hat{E}_{ij}^3 \) (up to some constant) to ensure stability of the resulting mean field Hamiltonian [81]. From (G3), a mean field decoupling (or, more formally, a Hubbard Stratonovich transformation) \( \hat{O}^\dagger \hat{O} \approx (\hat{O}^\dagger) \hat{O} - (\hat{O}^\dagger)(\hat{O}) \) is now straightforward. For \( U(1) \) spin liquids, we set the pairing terms \( \Delta_{ij} \) and \( \hat{D}_{ij} \) to be zero.

Similar to (B4), the mean field Hamiltonian for the triplet channels can be written in terms of a trace [54],

\[
H_{\text{MF}}^{\text{triplet}} = \sum_{ij} \sum_{x,y,z} \text{Tr} \left[ \sigma^\mu \Psi_{ij}^\mu \Psi_{ij}^\mu \right] T_{ij} \quad \text{(G4)}
\]

where the \( 2 \times 2 \) matrix \( u^\mu_{ij} \) contains the coefficients of \( \hat{E}_{ij}^\mu \) and \( \hat{D}_{ij}^\mu \). For a space group element \( X \), PSG constrains the triplet ansatz via

\[
u^\mu_{x(i)x(j)} = \sum_v O^\nu_X X(i)u^\nu_{ij} G^\dagger_X (X(j)),
\]

where \( O_X \in SO(3) \) encodes the \( SU(2) \) spin rotation associated with \( X \) [54]. There are only two nontrivial \( O_X \) in our problem,

\[
O_{C_i} = \begin{pmatrix} 0 & 0 & 1 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix}, \quad O_\sigma = \begin{pmatrix} -1 & 0 & 0 \\ 0 & 0 & 1 \\ 0 & 1 & 0 \end{pmatrix}.
\]

On the other hand, for the time reversal symmetry,

\[
u^\mu_{ij} = -G_T(i)u^\mu_{ij} G_T(j). \quad \text{(G7)}
\]

An excellent account of the treatment of triplet ansatz can be found in Ref. 54, to which interested readers can refer for more information. See also Ref. 81. Let \( E^\gamma_0 \) (\( E^z_0 \)) be the \( y \) component of the triplet hopping parameter on the bond connecting
sublattices 0 and 1 on an up (down) tetrahedron. Omitting the details of the calculation, we summarize the relation between the triplet ansatze $E_{ij} = \{E_{ij}^x, E_{ij}^y, E_{ij}^z\}$ for the $U(1)_0$ state in Table II, which can be extended to the $U(1)_\pi$ state by enlarging the unit cell and multiplying by $\pm 1$ as prescribed in Fig. 2b. In both states, a nonzero $E_{ij}^\theta$ is purely imaginary due to time reversal symmetry. The relations between the singlet ansatze $\chi_{ij}$ remain the same as before.

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