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Theoretical isotopic fractionation of magnesium between chlorophylls

Frédéric Moynier^{1,2} & Toshiyuki Fujii³

Magnesium is the metal at the center of all types of chlorophyll and is thus crucial to photosynthesis. When an element is involved in a biosynthetic pathway its isotopes are fractionated based on the difference of vibrational frequency between the different molecules. With the technical advance of multi-collectors plasma-mass-spectrometry and improvement in analytical precision, it has recently been found that two types of chlorophylls (*a* and *b*) are isotopically distinct. These results have very significant implications with regards to the use of Mg isotopes to understand the biosynthesis of chlorophyll. Here we present theoretical constraints on the origin of these isotopic fractionations through *ab initio* calculations. We present the fractionation factor for chlorophyll *a*, *b*, *d*, and *f*. We show that the natural isotopic variations among chlorophyll *a* and *b* are well explained by isotopic fractionation under equilibrium, which implies exchanges of Mg during the chlorophyll cycle. We predict that chlorophyll *d* and *f* should be isotopically fractionated compared to chlorophyll *a* and that this could be used in the future to understand the biosynthesis of these molecules.

Magnesium is a ubiquitous element in nature and has three naturally occurring stable isotopes, ²⁴Mg, ²⁵Mg and ²⁶Mg, with relative abundance of 78.99%, 10.00% and 11.01%, respectively. Magnesium also plays an essential role in a wide range of fundamental biological and non-biological reactions in the geological and biological system.

Chlorophylls are the green pigments, which allow phytoautotrophs to use light energy to produce organic matter by photosynthesis. Chlorophylls are formed of a Mg atom embedded in a porphyrin or chlorin ring^{1,2}. Chlorophylls can take several forms, with chlorophyll *a* the main pigment used by almost all oxygenic (oxygen-evolving) photosynthetic organisms. In addition to chlorophyll *a* (chl *a*), some organisms use accessory pigments: chlorophyll *b* (chl *b*) in green algae, and higher plant antennas, chlorophyll *c* (chl *c*) in certain algae, chlorophyll *d* (chl *d*) in some cyanobacteria and the recently discovered chlorophyll *f* (chl *f*) in some cyanobacteria³. In addition, there are a number of bacteriochlorophylls (BChl *a* through *g*) found in anoxygenic (non-oxygen evolving) photosynthetic bacteria^{2,4}.

Biological processes are widely recognized to have the ability to fractionate isotopes and create isotopic fractionation wherein the isotopic ratio of the product is different to that of the starting material (e.g. Zn, Fe, and Cu isotopic fractionation between plants and soils refs 5–10). Two studies found that Mg in cyanobacterial chl *a* is isotopically fractionated from the culture medium in which the cyanobacteria were grown and chl *a* and chl *b* have different isotopic compositions^{11,12}. They found that the ²⁶Mg/²⁴Mg of chl *a* was $0.43 \pm 0.15\%$ heavier than of chl *b*. These results have very important geochemical and biological implications: 1) Incorporating magnesium into chl *a* fractionates Mg isotopes and therefore the distribution of Mg isotopes in fossils may be used to search for the presence of fossil photosynthetic life and trace the origin of photosynthesis. 2) Studying the mechanism of isotopic fractionation by comparing the Mg isotopic composition measurements with *ab initio* calculations can help us understand the biosynthetic pathway of Mg during chlorophyll formation. In order to understand the origin of these fractionation Black *et al.*¹¹ performed *ab initio* calculation to estimate the theoretical isotopic fractionation between chl *a* and chl *b* and predicted an isotopic fractionation of 0.66‰ for the ²⁶Mg/²⁴Mg ratio (compared to 0.43 ± 0.15 in the natural samples).

Other chlorophyll structures than chl *a* and chl *b* exist, including the important red-shifted chl *d* and chl *f*. These pigments have not been studied using Mg isotopic methods yet.

Here we report the molecular orbitals of a large variety of Mg-chlorophyll species (chlorophylls *a*, *b*, *d*, *f*) to obtain the reduced partition function ratio (RPF) of isotopologues. We then use these data to test whether the Mg isotopic fractionation observed between chl *a* and chl *b* is due to chemical exchange of Mg and predict

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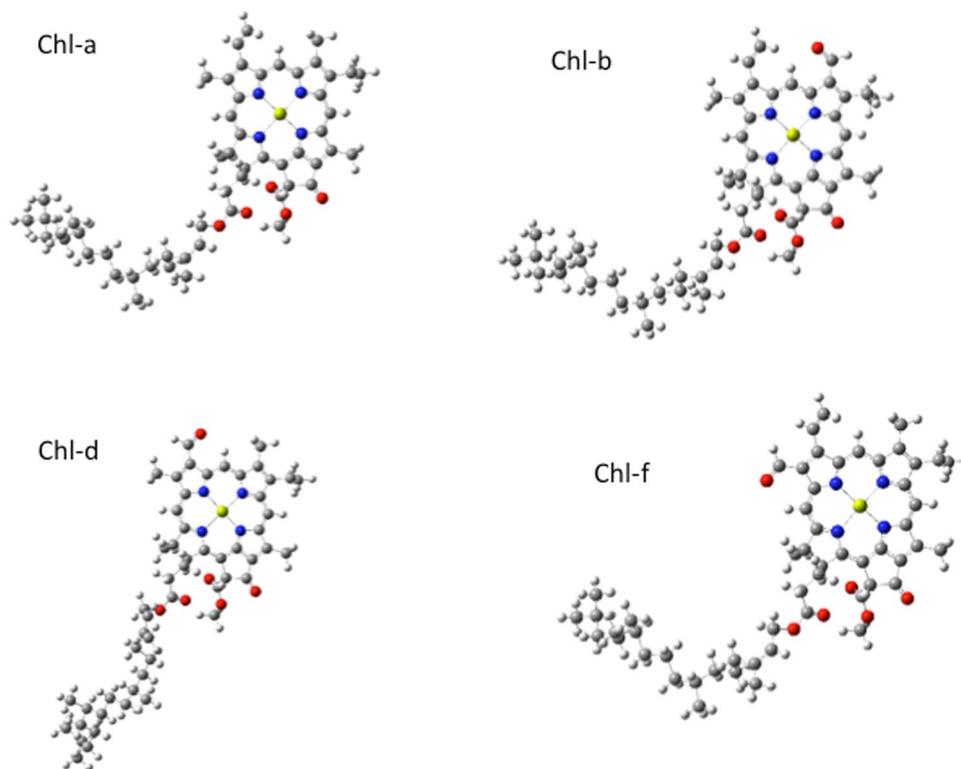


Figure 1. Optimized geometry for chlorophyll (a,b,d, and f).

whether such effect should affect chl *d* and chl *f*. Finally, we propose that by combining our calculations with natural Mg isotopic data would permit to better understand the biosynthetic formation of chlorophyll.

Methods

Magnesium has only three stable isotopes, ^{24}Mg , ^{25}Mg and ^{26}Mg . The isotopic composition of Mg is usually presented using the δ per mill (‰) notation defined as:

$$\delta^x\text{Mg} = \left[\frac{({}^x\text{Mg}/{}^{24}\text{Mg})_{\text{Samples}}}{({}^x\text{Mg}/{}^{24}\text{Mg})_{\text{Standard}}} - 1 \right] \times 1,000 \quad (1)$$

with $x = 25$ or 26 . All the data are reported relative to the same standard (DSM-3; Black, 2006 #4698).

And we define the difference of $\delta^x\text{Mg}$ between two species X and Y a $\Delta^x\text{Mg}_{\text{X-Y}}$, with $x = 25$ or 26 as:

$$\Delta^x\text{Mg}(X - Y) = \delta^x\text{Mg}(X) - \delta^x\text{Mg}(Y) \quad (2)$$

Computational method. Orbital geometries, vibrational frequencies, Gibbs free energies of aqueous Mg species are computed using density functional theory (DFT) as implemented by the Gaussian03 code¹³. The DFT method employed is a hybrid density functional consisting of Becke's three-parameter non-local hybrid exchange potential (B3)¹⁴ with Lee-Yang-and Parr (LYP) non-local functionals. The 6-311 + G(d,p) basis set, which is an all-electron basis set, will be chosen for H, C, O, and Mg. For the solvation effect, CPCM continuum solvation method (CPCM: conductor-like polarizable continuum model) is used.

For the structure of chlorophyll a, b, d and f we used an optimized geometry shown in the Fig. 1 (data are available as supplementary materials).

Results and Discussion

The vibrational modes ν_1 , ν_2 , and ν_3 of hexaaqua complexes are the fundamental intramolecular vibration modes. The validity of vibrational frequencies and atomic distances of Mg-O of hydrated Mg^{2+} ion computed is discussed in detail in Schott *et al.*¹⁵. The calculated vibrational frequencies of $\text{Mg}(\text{H}_2\text{O})_6^{2+}$ are underestimated compared with the literature values determined by Raman and IR spectrometry. Setting 12 H_2O molecules at the second coordination sphere brought the calculation results closer to the literature values. The use of conductor-like polarizable continuum model (CPCM) of solvation for $\text{Mg}(\text{H}_2\text{O})_6^{2+}$ increases vibrational frequencies, but the effect is smaller than setting water molecules in the second coordination sphere.

Calculating how physicochemical properties vary with hydration of the species is a common strategy for examining the accuracy of theoretical calculations of aqueous species^{16,17}. In the theoretical study on the hydration enthalpy of Fe^{2+} and Fe^{3+} , Li *et al.*¹⁸ tested a small cluster model of 6 H_2O molecules as the first coordination

Hydration number n	Method/Basis set	$\Delta H_{\text{hyd}}^{\circ}$ / kJ/mol	Reference
6	O3LYP/6-31G(d)	-1893	This work
6	B3LYP/6-31G(d)	-1954	This work
18	O3LYP/6-31G(d)	-2047	This work
18	B3LYP/6-31G(d)	-2280	This work
—		-1990	Rosseinsky, 1965
—		-1921	Smith, 1977
—		-1931	Marcus, 1985

Table 1. Hydration enthalpy of $\text{Mg}(\text{H}_2\text{O})_n^{2+}$.

sphere and a large cluster model of 12 additional H_2O molecules as the second coordination sphere. For Fe^{3+} , the large cluster model brought the calculated data closer to the experimental results. For Fe^{2+} , both the small and the large cluster models reproduced the experimental results. Similarly, the hydration enthalpies of Ni^{2+} and Zn^{2+} were appropriately reproduced by using both the small and the large cluster models. Magnesium ion is a divalent one. The small cluster model may be applicable.

The hydration enthalpy of $\text{Mg}(\text{H}_2\text{O})_n^{2+}$ ($n = 6$ or 18) was also examined. In order to relate the calculated quantities to the experimental hydration enthalpy ($\Delta H_{\text{hyd}}^{\circ}$) at 298 K, the following correction terms were considered¹⁸,

$$-\Delta H_{\text{hyd}}^{\circ} = -\Delta E_{\text{b}} + \Delta E_{\text{sol}} + n\Delta H_{\text{vap}} + \Delta nRT - \Delta E(\text{Cp}) - \Delta E_{\text{zp}} + \Delta E_{\text{rel}} + \Delta E_{\text{geom}} \quad (3)$$

ΔE_{b} is the total binding energy of the gas phase cluster $[\text{Mg}(\text{H}_2\text{O})_n]$. ΔE_{sol} represents the solvation free energy of $\text{Mg}(\text{H}_2\text{O})_n^{2+}$. This term, which contains the entropic contribution to the solvation free energy for the continuum dielectric part, was neglected. ΔH_{vap} is the heat of vaporization of water, which is 10.50 kcal/mol¹⁹. The $\Delta E(\text{Cp})$ term arises from the difference in heat capacity of the components of the system, which corresponds to a small correction of ~ 1 kcal/mol at 298 K¹⁸. ΔE_{zp} is the difference in vibrational zero-point energy in forming clusters, ΔE_{rel} is a correction due to relativistic effects for metal centers, and ΔE_{geom} is a correction due to geometry relaxation for H_2O during the formation of the clusters. The magnitude of $-\Delta E_{\text{zp}} + \Delta E_{\text{rel}} + \Delta E_{\text{geom}}$ was reported to be negligibly small (a few kcal/mol)¹⁸, and hence these corrections were not included in our calculation. The calculated value of $\Delta H_{\text{hyd}}^{\circ}$ is shown in Table 1. The hydration enthalpy of metal cations has been determined by thermochemical methods^{20–22}, and literature values are shown for comparison. Setting 12 H_2O around the small cluster $\text{Mg}(\text{H}_2\text{O})_6^{2+}$ resulted in the increase of $\Delta H_{\text{hyd}}^{\circ}$. For the method, O3LYP, large cluster model showed similar value compared with literature data, while B3LYP showed that small cluster model showed similar value compared with literature data. For divalent cations, even small cluster models like $\text{Mg}(\text{H}_2\text{O})_6^{2+}$ can reproduce $\Delta H_{\text{hyd}}^{\circ}$.

All the data are reported in Table 2 and Fig. 2 for some of the most relevant molecules discussed. The absolute values of our calculated $\ln\beta$ for $\text{Mg}(\text{H}_2\text{O})_6^{2+}$, $\text{Mg}(\text{H}_2\text{O})_{18}^{2+}$, chl *a* and chl *b* are in fairly good agreement with literature data^{11, 23, 24} while we are using different program packages: PQS ver. 3.3. for Black *et al.*¹¹ vs Gaussian09 in the present study and Schott *et al.*¹⁵.

While the relative difference between the $\ln\beta^{26/24}\text{Mg}$ of chl *b* and chl *a* (Table 3) that we obtain (0.34) is slightly lower than in Black *et al.*¹¹ (0.67); it is closer than the experimental data (0.43). The accuracy of our calculation using Gaussian09 for Mg isotopologues was tested by comparing our calculations with experimental data for a large set of aqueous species (sulphides, citrates, EDTA, oxalates, hydroxides; see ref. 23).

The calculated theoretical isotopic fractionation between the four chlorophylls species considered here (chl *a*, chl *d* and chl *f* compared to chl *b*, see Table 3) predicts resolvable isotopic variations between the different species of chlorophylls. The Mg isotopic composition of the chl *f* should be very close to chl *b* with $\Delta^{26}\text{Mg}(\text{chl } f\text{-chl } b) = 0.09$, chl *d* is slightly heavier, $\Delta^{26}\text{Mg}(\text{chl } d\text{-chl } b) = 0.18$ and finally chl *a* is the isotopically heaviest $\Delta^{26}\text{Mg}(\text{chl } a\text{-chl } b) = 0.34$.

Our results suggest that each type of chlorophyll should exhibit a distinct Mg isotopic composition. It is therefore possible to test whether Mg were exchanged at equilibrium via the Mg isotopic composition of the different chlorophyll. Following this approach, Black *et al.*¹¹ proposed that Mg were exchanged at equilibrium between chl *a* and chl *b* based on the similarity between the measured and calculated isotopic composition, suggesting that the timescale of Mg exchanges were similar to the lifetime of the chlorophylls. Our new results show even better agreements with the measurements further suggesting that Mg were exchanged during the lifetime of the chlorophyll.

It is known that the relative abundance of chl *a* and chl *b* in plants or green algae changes as a function of the light conditions²⁵ suggesting that the biosynthesis of chl *a* and chl *b* are closely linked. Tanaka *et al.*²⁶ isolated the gene encoding for the formation of chl *b* by oxygenation of chl *a* and proposed a chlorophyll cycle with an intermediary molecule, 7-hydroxymethyl. From their model, Mg is not involved in the cycle and, therefore, chl *a* and chl *b* should have similar isotopic composition. The fact that the Mg isotopes are fractionated between isolated chl *a* and chl *b* and follow our theoretical prediction confirms that the biosynthesis of chl *a* and chl *b* are closely linked and implies the Mg is also exchanged during the chlorophyll cycle. Our results imply that the biochemical pathway of the synthesis of chl *a* and chl *b* is more complex than presently modeled.

In this study we have expanded our calculations to the two most recently discovered chlorophylls, chl *d* and chl *f*. Both chl *d* and chl *f* are always found in association with chl *a*²⁷. Chlorophyll *d* and *f* are red-shifted chlorophyll

Species	Method/basic sets	$\ln\beta^{25/24}$ 298 K	$\ln\beta^{25/24}$ A, B ^a	$\ln\beta^{26/24}$ 298 K	$\ln\beta^{26/24}$ A, B ^a	Reference
Mg(H ₂ O) ₆ ²⁺	O3LYP/6-31G(d)	12.43	1.0425, 0.671	23.88	2.0041, 1.272	This study
Mg(H ₂ O) ₆ ²⁺	B3LYP/6-31G(d)	13.73	1.1448, 0.816	26.37	2.1998, 1.556	This study
Mg(H ₂ O) ₁₈ ²⁺	O3LYP/6-31G(d)	12.48	1.0470, 0.666	23.96	2.0104, 1.282	This study
Mg(H ₂ O) ₁₈ ²⁺	O3LYP/6-31G(d)	13.18	—	25.22	—	Black <i>et al.</i> ¹¹
Mg(H ₂ O) ₁₈ ²⁺	B3LYP/6-31G(d)	14.26	1.1881, 0.857	27.40	2.2852, 1.615	This study
Mg(H ₂ O) ₁₈ ²⁺	BP86/6-31G(d)	—	—	26.74	—	Rustad <i>et al.</i> ²⁴
Chlorophyll-a	O3LYP/6-31G(d)	14.61	1.2077, 0.979	28.07	2.3255, 1.828	This study
Chlorophyll-a	O3LYP/6-31G(d)	15.21	—	29.27	—	Black <i>et al.</i> ¹¹
Chlorophyll-a	B3LYP/6-31G(d)	15.14	1.2510, 1.022	29.08	2.4053, 1.944	This study
Chlorophyll-b	O3LYP/6-31G(d)	14.43	1.1949, 0.948	27.73	2.2995, 1.785	This study
Chlorophyll-b	O3LYP/6-31G(d)	14.88	—	28.64	—	Black <i>et al.</i> ¹¹
Chlorophyll-b	B3LYP/6-31G(d)	14.99	1.2397, 1.001	28.79	2.3830, 1.905	
Chlorophyll-d	O3LYP/6-31G(d)	14.53	1.2028, 0.956	27.91	2.3123, 1.822	This study
Chlorophyll-d	B3LYP/6-31G(d)	15.13	1.2509, 1.017	29.08	2.4042, 1.949	This study
Chlorophyll-f	O3LYP/6-31G(d)	14.48	1.1999, 0.937	27.82	2.3063, 1.795	This study
Chlorophyll-f	B3LYP/6-31G(d)	15.03	1.2427, 1.010	28.89	2.3913, 1.906	This study

Table 2. Logarithm of the reduced partition function, $\ln\beta$ (%), for the pair ²⁶Mg-²⁴Mg and ²⁵Mg-²⁴Mg of Mg(II) complexes. ^a $10^3 \ln\beta = 10^6 A T^{-2} + B$.

Species	Method	$\Delta^{25}\text{Mg}$	$\Delta^{26}\text{Mg}$	Reference
chl a vs. chl b	O3LYP/6-31G(d)	0.18	0.34	This study
chl a vs. chl b	O3LYP/6-31G(d)	0.33	0.63	Black <i>et al.</i> ¹¹
chl d vs. chl b	O3LYP/6-31G(d)	0.10	0.18	This study
chl f vs. chl b	O3LYP/6-31G(d)	0.05	0.09	This study

Table 3. Theoretical isotopic fractionation between different forms of chlorophylls and chlorophyll a for 298 K.

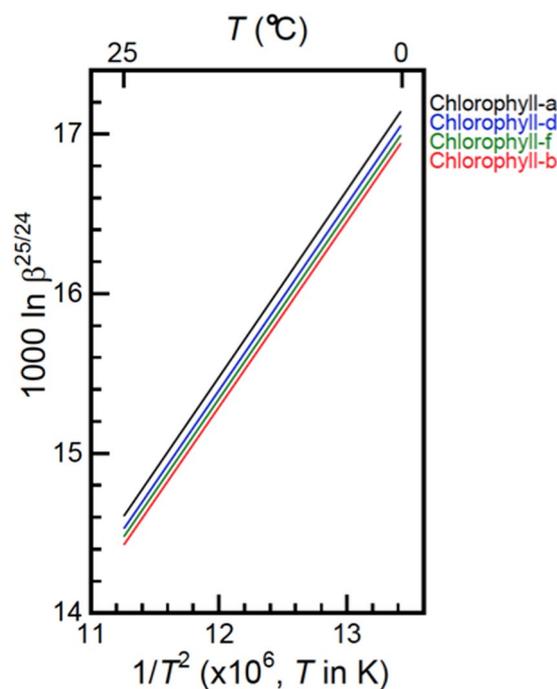


Figure 2. Temperature dependence of $\ln\beta$. The $\ln\beta^{25/24}$ values of chlorophyll a, b, d, and f calculated by using O3LYP/6-31G(d) are shown as linear functions of T^{-2} .

that can adsorb light with wavelength up to 760 nm while chl a and b do not absorb light <700 nm. These pigments have only been discovered in a limited set of organisms so far (e.g. *Acaryochloris Marina*, *Halomicronema Hongdechloris*) and are supposed to represent the consequence of adaptation of the micro-organisms to specific

ecological niches. The understanding of the biosynthesis of these novel pigments is still limited and while there is no consensus on the existence of chl *d*-chl *a* and/or chl *f*-chl *a* cycles there are hints that the biosynthesis of chl *d* and *f* are linked to chl *a*. Using spiked oxygen marking Schliep *et al.*²⁸ suggested that chl *d* was formed directly from chl *a* via oxygenase-type reactions but the enzymes responsible for the synthesis have yet to be discovered. Our new results on the two long wave-length absorbing chlorophylls exhibit distinct isotopic composition suggesting that it would also be possible to test the kinetic of Mg exchanged compared to the lifetime of chl *d* and chl *f*.

Isotopic fractionation of Mg at equilibrium between 4 types of chlorophylls (*a*, *b*, *d*, and *f*) was demonstrated theoretically. We show that our data are consistent with previous calculations and confirms that chl *a* and chl *b* are isotopically fractionated in plants following a thermodynamic equilibrium. This implies that during the synthesis of chl *b* and chl *a*, Mg is also exchanged. We expand these calculations to the two latest discovered pigments, chl *d* and chl *f* and show that their Mg isotopes should also be isotopically fractionated during equilibrium exchanges reactions.

References

- Blankenship, R. E. *Molecular Mechanisms of Photosynthesis*. (Blackwell, 2002).
- Grimm, B., J., P. R., Rudiger, W. & H., S. Chlorophylls and Bacteriochlorophylls: Biochemistry, Biophysics, Functions, and Applications. (Springer, 2006).
- Chen, M. *et al.* A Red-Shifted Chlorophyll. *Science* **329**, 1318–1319 (2010).
- Niedzwiedzki, D. M. & Blankenship, R. E. Singlet and Triplet Excited State Properties of Natural Chlorophylls and Bacteriochlorophylls. *Photosynthesis Research* **106**, 227–238 (2010).
- Viers, J. *et al.* Evidence of Zn isotopic fractionation in a soil-plant system of a pristine tropical watershed (Nsimi, Cameroon). *Chem. Geol.* **239**, 124–137 (2007).
- Moynier, F. *et al.* Isotopic fractionation and transport mechanisms of Zn in plants. *Chem. Geol.* **267**, 125–130, doi:10.1016/j.chemgeo.2008.09.017 (2009).
- Weinstein, C. *et al.* Cu isotopic fractionation in plants. *Chem. Geol.* (2011).
- Moynier, F., Fujii, T., Wang, K. & Foriel, J. Ab initio calculations of the Fe(II) and Fe(III) isotopic effects in citrates, nicotianamine, and phytosiderophore, and new Fe isotopic measurements in higher plants. *Comptes Rendus Geoscience* **345**, 230–240, doi:10.1016/j.crte.2013.05.003 (2013).
- Guelke-Stelling, M. & von Blanckenburg, F. Fe isotope fractionation caused by translocation of iron during growth of bean and oats as models of strategy I and II plants. *Plant soil* **352**, 217–231 (2012).
- Moynier, F., Vance, D., Fujii, T. & Savage, P. In Non-traditional stable isotopes Vol. 82 (eds F-Z Teng, J. Watkins, & N. Dauphas) 543–600 (Mineralogical Society of America, 2017).
- Black, J. R., Yin, Q. Z., Rustad, J. R. & Casey, W. H. Magnesium isotopic equilibrium in chlorophylls. *J Am Chem Soc* **129**, 8690, doi:10.1021/ja072573i (2007).
- Black, J. R., Casey, W. H. & Yin, Q. Z. An experimental study of magnesium-isotope fractionation in chlorophyll-a photosynthesis. *Geochim. Cosmochim. Acta* **70**, 4072–4079, doi:10.1016/j.gca.2006.06.010 (2006).
- Gaussian 09, Revision B.01, Gaussian, Inc.: Wallingford CT (2009).
- Becke, A. D. Density-functional thermochemistry. 3. *The role of exact exchange*. *J. Chem. Phys.* **98**, 5648–5652 (1993).
- Schott, J., Mavromatis, V., Fujii, T., Pearce, C. R. & Oelkers, E. The control of carbonate mineral Mg isotope composition by aqueous speciation: Theoretical and experimental modeling. *Chem. Geol.* **445**, 120–134 (2016).
- Fujii, T., Moynier, F., Blichert-Toft, J. & Albarede, F. Density functional theory estimation of isotope fractionation of Fe, Ni, Cu, and Zn among species relevant to geochemical and biological environments. *Geochim. Cosmochim. Acta* **140**, 553–576, doi:10.1016/j.gca.2014.05.051 (2014).
- Moynier, F. & Fujii, T. Calcium isotope fractionation between aqueous compounds relevant to low-temperature geochemistry, biology and medicine. *Scientific Reports* **7**, 44255, doi:10.1038/srep44255 (2017).
- Li, J., Fisher, C. L., Chen, J. L., Bashford, D. & Noodleman, L. Calculation of redox potentials and pKa values of hydrated transition metal cations by a combined density functional and continuum dielectric theory. *Inorg. Chem.* **35**, 4694–4702 (1996).
- Lide, D. R. *CRC Handbook of Chemistry and Physics*, 89th Ed., (CRC Press, 2008).
- Rosseinsky, D. R. Electrode potentials and hydration energies. Theories and correlations. *Chem. Rev.* **65**, 467–490 (1965).
- Smith, D. W. Ionic hydration enthalpies. *J. Chem. Edu* **54**, 540–542 (1977).
- Marcus, Y. *Ion Solvation*. (John Wiley & Sons, 1985).
- Schott, J., V., M., Fujii, T., Pearce, C. R. & Oelkers, E. The control of carbonate mineral Mg isotope composition by aqueous speciation: Theoretical and experimental modeling. *Chem. Geol.* **445**, 120–134 (2014).
- Rustad, J. R. *et al.* Isotopic fractionation of Mg²⁺ (aq), Ca²⁺ (aq), and Fe²⁺ (aq) with carbonate minerals. *Geochim. Cosmochim. Acta* **74**, 6301–6323, doi:10.1016/j.gca.2010.08.018 (2010).
- Björkman, O. *et al.* *Carnegie Inst Washington Yearb.* **1972**, 115–135 (1972).
- Tanaka, A. *et al.* Chlorophyll a oxygenase (CAO) is involved in chlorophyll b formation from chlorophyll a. *Proceedings of the National Academy of Sciences of the United States of America* **95**, 12719–12723 (1998).
- Li, Y. & Chen, M. *Functional Plant Biology* **42**, 493–501 (2015).
- Schliep, M., Crossett, B., Willows, R. D. & Chen, M. 18O Labelling of chlorophyll d in *Acaryochloris marina* reveals that chlorophyll a and molecular oxygen are precursors. *J. Biol. Chem.* **37**, 28450–28456 (2010).

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Author Contributions

F.M. and T.F. designed the project. T.F. performed the calculations. F.M. wrote the manuscript.

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