



HAL
open science

Unravelling the Role of Uncommon Hydrogen Bonds in Cyclodextrin Ferrociphenol Supramolecular Complexes: A Computational Modelling and Experimental Study

Pascal Pigeon, Feten Najlaoui, Michael James Mcglinchey, Juan Sanz García, Gérard Jaouen, Stéphane Gibaud

► To cite this version:

Pascal Pigeon, Feten Najlaoui, Michael James Mcglinchey, Juan Sanz García, Gérard Jaouen, et al.. Unravelling the Role of Uncommon Hydrogen Bonds in Cyclodextrin Ferrociphenol Supramolecular Complexes: A Computational Modelling and Experimental Study. *International Journal of Molecular Sciences*, 2023, Special Issue Cyclodextrins: Properties and Applications, 24 (15), pp.12288. 10.3390/ijms241512288 . hal-04176438

HAL Id: hal-04176438

<https://hal.science/hal-04176438>

Submitted on 3 Aug 2023

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Article

Unravelling the Role of Uncommon Hydrogen Bonds in Cyclodextrin Ferrociphenol Supramolecular Complexes: A Computational Modelling and Experimental Study

Pascal Pigeon ^{1,2,*} , Feten Najlaoui ³, Michael James McGlinchey ⁴, Juan Sanz García ⁵ , Gérard Jaouen ^{1,2} and Stéphane Gibaud ³

- ¹ Chimie ParisTech, PSL, 11 Rue Pierre et Marie Curie, CEDEX 05, 75231 Paris, France; gerard.jaouen@chimieparistech.psl.eu
- ² Institut Parisien de Chimie Moléculaire (IPCM)—UMR 8232, Sorbonne Université, 4 Place Jussieu, 75252 Paris, France
- ³ EA 3452/CITHEFOR, Faculté de Pharmacie, Université de Lorraine, 5 Rue Albert Lebrun, 54000 Nancy, France
- ⁴ UCD School of Chemistry, University College Dublin Belfield, D04 V1W8 Dublin, Ireland
- ⁵ 1MSME, Université Gustave Eiffel, CNRS, 5 Boulevard Descartes, 77454 Marne-la-Vallée, France; juan.sanz.garc@gmail.com
- * Correspondence: pascal.pigeon@chimieparistech.psl.eu or pascal.pigeon@sorbonne-universite.fr; Tel.: +33-14-427-6699

Abstract: We sought to determine the cyclodextrins (CDs) best suited to solubilize a patented succinimido-ferrociphenol (SuccFerr), a compound from the ferrociphenol family having powerful anticancer activity but low water solubility. Phase solubility experiments and computational modelling were carried out on various CDs. For the latter, several CD-SuccFerr complexes were built starting from combinations of one or two CD(s) where the methylation of CD oxygen atoms was systematically changed to end up with a database of ca. 13 k models. Modelling and phase solubility experiments seem to indicate the predominance of supramolecular assemblies of SuccFerr with two CDs and the superiority of randomly methylated β -cyclodextrins (RAME β CDs). In addition, modelling shows that there are several competing combinations of inserted moieties of SuccFerr. Furthermore, the models show that ferrocene can contribute to high stabilization by making atypical hydrogen bonds between Fe and the hydroxyl groups of CDs (single bond with one OH or clamp with two OH of the same glucose unit).

Keywords: cyclodextrin; ferrocene; modelling; phase solubility; database; atypical hydrogen bond



Citation: Pigeon, P.; Najlaoui, F.; McGlinchey, M.J.; Sanz García, J.; Jaouen, G.; Gibaud, S. Unravelling the Role of Uncommon Hydrogen Bonds in Cyclodextrin Ferrociphenol Supramolecular Complexes: A Computational Modelling and Experimental Study. *Int. J. Mol. Sci.* **2023**, *24*, 12288. <https://doi.org/10.3390/ijms241512288>

Academic Editors: Miguel A. Esteso and Carmen M. Romero

Received: 4 July 2023
Revised: 26 July 2023
Accepted: 28 July 2023
Published: 31 July 2023



Copyright: © 2023 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

1. Introduction

1.1. Cyclodextrins

Cyclodextrins (CDs) are cyclic oligosaccharides formed by glucose units bound together. Nowadays, they are used as excipients to improve the solubility of lipophilic compounds. Typical cyclodextrins contain a number of glucose units ranging from six to eight units in a ring and can be classified as α -cyclodextrin with six glucose subunits, β -cyclodextrin with seven glucose subunits (Figure 1) and γ -cyclodextrin with eight glucose subunits. Cyclodextrins are frequently transformed by chemical modification of the hydroxyl group, for example, to form alkyloxy derivatives (O-Me, O-Et, etc.) [1].

The most commercially viable strategies lead to the synthesis of partially and randomly alkylated cyclodextrins [2,3]. Chemically modified cyclodextrin derivatives available on the market are mixtures with different degrees of substitution [4]. These cyclodextrins can be used as excipients in many pharmaceutical formulations (e.g., tablets, aqueous parenteral solutions, eye drop solutions and nasal sprays) [5–8]. Although these randomly substituted cyclodextrins are widely used as solubilizers, they can form a wide range of complexes,

making the characterization of these mixtures quite challenging. In silico models of these mixtures are usually focused on one or very few well-defined structures.

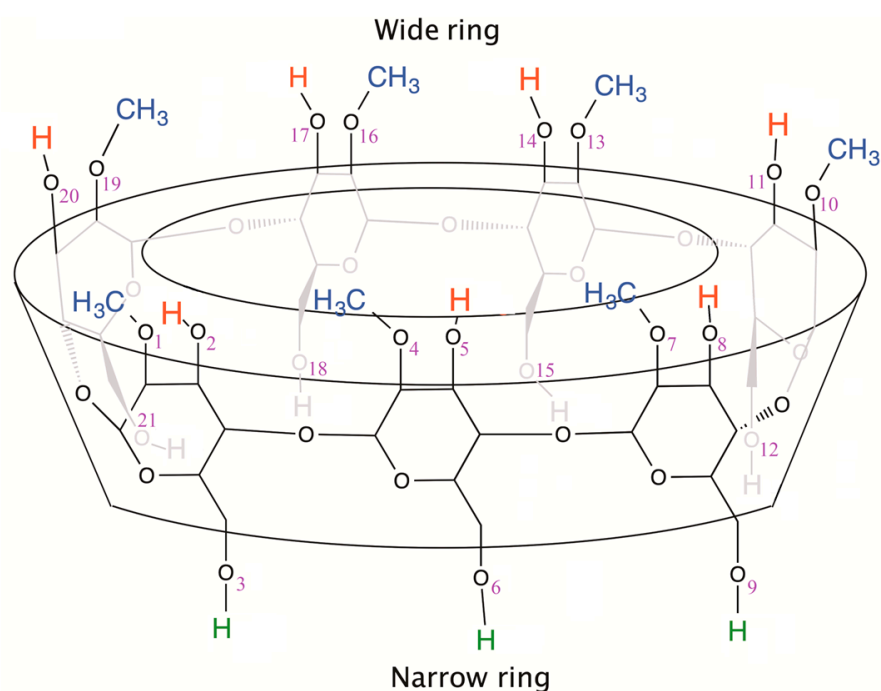


Figure 1. Representation of a β -cyclodextrin (CD, here 2-Me- β CD belonging to the category of well-defined CDs) with the oxygen atom numbering used for our modelling and database (glucose numbering is not displayed).

The term “well-defined” implies that all glucose units are identical. Figure 1 shows an example, 2-Me- β CD, in which all 2-positions (glucose numbering is not indicated) are methylated, and all 3- and 6-positions are unmethylated. CDs having at least two different glucose units are designated as undefined CDs in this work.

Although chemical modification of cyclodextrins might seem quite simple, it is in fact particularly challenging due to the presence of –OH groups in large numbers. More complex methods for selective modifications of cyclodextrins have been reviewed [9] and are currently used to prepare very specific varieties of them, such as the heptakis(2,3,6-tri-O-methyl)- β -cyclodextrin or the heptakis-(2,6-di-O-methyl)- β -cyclodextrin.

These cyclodextrins are much more expensive than the commercially available randomly methylated cyclodextrins (RAME β CDs), making their use very restricted. In fact, researchers are often led to focus on cheaper, randomly substituted cyclodextrins for animal studies and for clinical trials. A non-trivial question arises when modelling cyclodextrins: what is the difference between solutions prepared with a well-defined cyclodextrin and with a mixture of randomly substituted cyclodextrins? We already know that the association constant of randomly methylated cyclodextrins depends greatly on their degree of methylation [10].

1.2. Ferrociphenol SuccFerr

In this paper, we report that we have performed a series of experiments for formulation with CDs on a ferrociphenol derivative (ferrociphenol with an additional *p*-hydroxyphenyl group: *N*-{4-ferrocenyl-5,5-bis-(4-hydroxy-phenyl)-pent-4-enyl}succinimide; SuccFerr (or P722), Figure 2) recently developed in our research group. SuccFerr is the result of an optimization of the ferrociphenol family [11] whose members act by apoptosis or senescence [12], where the pharmacophore is the *trans* (ferrocene-double bond-*p*-phenol) motif [13] (in blue in Figure 2), and the main metabolite is a quinone-methide [14,15] able to attack proteins [16,17]. However, the attachment of a succinimidyl group decreased the IC₅₀ to around 40 nM on MDA-MB-231, A2780, A2780-Cis and K562 cell lines [18] thanks to

an intramolecular stabilization [19]. Moreover, recent research on glioblastomas has shown that SuccFerr has a surprising selectivity on patient-derived cell lines (PDCLs) [20]. This patented compound [21] is currently being tested for its anticancer activity, but a suitable formulation was needed to achieve a usable formulation because of its lipophilicity and poor solubility. SuccFerr was recently formulated with various nanocapsules in its diphenol form, or to increase its lipophilicity and the load in lipid nanocapsules, as monoacetate [22] or diacetate [22–24], with or without the unexpected formation of a gel [22]. It was recently formulated with randomly methylated cyclodextrins (RAME β CDs) and tested in vivo without noticeable adverse toxicity [25], but this last complexation has yet to be analyzed more finely by modelling and laboratory experiments.

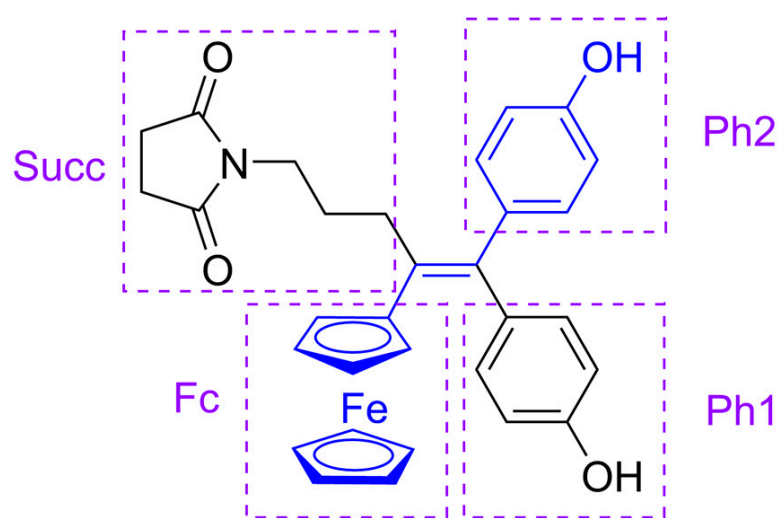


Figure 2. SuccFerr (the ferrociphenol motif is drawn in blue, and the moieties that can enter the cavity of the CDs are surrounded by purple rectangles with the abbreviations used in this work: the ferrocenyl group (Fc), phenol 1 (Ph1, *cis* to the Fc), phenol 2 (Ph2, *trans* to the Fc) and the succinimidylpropyl group (Succ).

2. Results

2.1. Experiments Prior to Modelling (Proof of Concept)

In our study, it was important to compare the modelling results with the laboratory solubilization tests. The complexing capacity is known to be very variable depending on the type of cyclodextrin, and it was therefore necessary to compare the results of solubilization of SuccFerr with various CDs.

The phase–solubility study was first described by Higuchi and Connors [26] to test the dissolving properties of the complexation.

In our experiment (Figure 3A), α -CD did not improve the solubility of SuccFerr, probably due to its small cavity size (internal diameter: 0.57 nm). In contrast, the methylated β -cyclodextrin (RAME β CD) showed significant solubilization properties already previously described for the phthalimide compound [27]. The internal diameter of this CD is 0.68 nm, and it is well adapted to complex aromatic and heterocyclic moieties [28].

The phase solubility diagram of SuccFerr in the presence of RAME β CD (see Figure 3A) exhibits a positive curvature, described as Ap-type by Higuchi and Connors [26]. This is observed if more than one CD can complex the drug corresponding to 1:2, 1:3, 1:4 (or more) stoichiometries. Our results confirm that RAME β CD can be used to prepare a solution of SuccFerr.

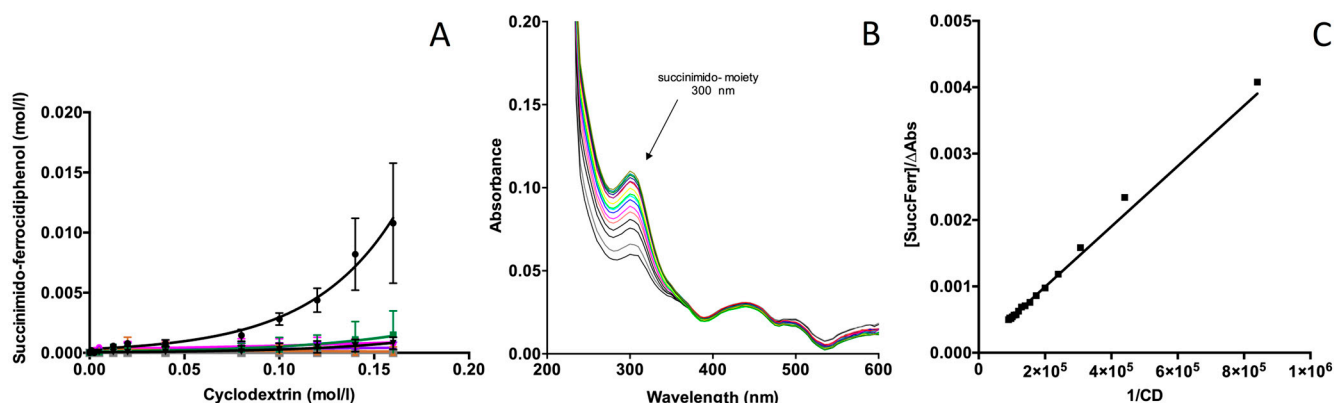


Figure 3. (A) Phase solubility diagram of SuccFerr in the presence of α CD (\blacksquare), RAME β CD (\bullet), DM β CD (\blacktriangledown), HP β CD (\blacksquare), HE β CD (\blacktriangle), SBE β CD (\blacklozenge), γ CD (\blacklozenge). (B) Spectra of SuccFerr (2.5×10^{-5} M). The control (i.e., lower curve) was obtained with 1% DMSO. The other curves were obtained by addition of various amounts of RAME β CD, exhibiting a hyperchromic effect at 300 nm. (C) Benesi–Hildebrand plots for the complexation of SuccFerr in RAME β CD ($\lambda = 300$ nm).

It is noteworthy that DM β CD does not similarly improve the solubility as RAME β CD does.

A second study was carried out to better understand the complexation mechanism. This experiment was performed to confirm the inclusion of the succinimidyl group in RAME β CD. The spectrum of pure SuccFerr exhibits an absorption maximum at 300 nm corresponding to the succinimidyl moiety (Figure 3B). The ferrocenyl moiety is known to exhibit a λ_{\max} at approximately 200 nm, whereas the λ_{\max} of phenols is found at approximately 270 nm. The addition of RAME β CD induces a hyperchromic effect at 300 nm (Figure 3B); in contrast, the inclusion of phenols and ferrocenyl moiety did not modify the spectrum at 254 nm and 270 nm.

The specific effect observed at 300 nm can be used to study the inclusion of the succinimidyl group using the method previously described for the phthalimide derivative [27] and derived from the Benesi–Hildebrand method. Consequently, we studied the variations in the absorption of SuccFerr as a function of the concentration of RAME β CD using the spectroscopic data. Each absorption value was taken from our spectra at 300 nm.

The corresponding theoretical equation is:

$$\frac{[\text{SuccFerr}]}{\Delta A} = \frac{1}{\epsilon_c \cdot l \cdot [\text{CD}_0] \cdot K_a} + \frac{1}{\epsilon_c \cdot l} \quad (1)$$

where

$$\Delta A = \frac{1}{\sum_{i=1}^n \frac{1}{\Delta A_i}} K_a = \frac{1}{\sum_{i=1}^n \frac{1}{K_i}}$$

This equation is similar to the Benesi–Hildebrand equations and is linear for $[\text{SuccFerr}]/\Delta A$ as a function of $1/[\text{CD}_0]$ (Figure 3C). Consequently, we can confirm that the linear curves obtained for double reciprocal curves of Benesi–Hildebrand correspond to a 1:1 apparent complexation with the succinimidyl group. In this equation, the absorbance coefficient and K_a can be defined as the harmonic mean of all the complexes if there is an inclusion of the succinimidyl moiety (see [27] for more details). $[\text{SuccFerr}]$ is the concentration of SuccFerr, $[\text{CD}_0]$ is the initial concentration of free cyclodextrin, ϵ_c is the change in the molar absorption coefficient after complexation, and l is the path length.

The curve obtained from experimental data (i.e., $[\text{SuccFerr}]/\Delta A$ as a function of $1/[\text{CD}_0]$) is presented in Figure 3C, and K_a was calculated ($K_a = 50.618 \pm 5.280 \text{ M}^{-1}$).

2.2. Modelling and Web Application

We used modelling (semiempirical PM3 quantum-mechanical method) to determinate the type of association SuccFerr could make with CDs. Two main types of series have

been produced. For the inclusion of SuccFerr into a single CD, eight possible systems were created (four moieties (Fc, Ph1, Ph2 and Succ, Figure 2) inserted by the narrow or the wide side of the CD, denoted 1-CD series S1 to S8). For the models with two CDs, only four series were made (2-CD series S1 to S4). These 12 trees have the same treatment in common. First, taking 1-CD models as an example, we calculated all the possible combinations of insertion (four moieties for SuccFerr \times two insertion sides per CD \times six possible well-defined CDs, see Tables S1 and S2 for 2-CD). Then, for each of the eight (moiety-CD side) combinations, we selected the model with the most negative ΔE (the energy difference between the supramolecular assembly and the separated molecules, while keeping the same geometry). The aim of this selection was to start with the lowest possible energy for each of these eight models that served as the G0 (generation 0) model for creating series/trees. Then, the methylations on all oxygen atoms (21 for 1-CD systems) were inverted, creating a G1 level with 21 models in the tree. The best model (lowest ΔE) was selected to create the G2 level of the tree in the same way, and so on.

We created a web application to handle the data of these 12 trees, which also provided statistics, but a need arose to obtain more data on our calculations. This is why a second program was created in C programming language. Its purpose was to verify the models (building error that would invalidate this work), to search for certain hydrogen bonds of interest, to count the average number of methyl groups around Fc and to generate the static webpages, copies of the dynamic webpages created by the web application. In the latter case, the goal of generating these static webpages was to dispense with the web application for simplified read-only consultation (to view a particular model or to navigate into the trees). These webpages can be downloaded as a compressed archive [29].

To facilitate a better understanding of this concept, the code of the PHP program, the code of the C program and the database (scheme and data) are provided in a data repository [30] (including the static webpages cited above). The web application (CDModelTree) was also referenced on Software Heritage via Hal [31] and is executable and testable (even with any other inserted molecule that readers of this article would like to test, with up to two CDs for inclusion, and without any need to adapt the code) using the XAMPP pack [32]. More details can be found in the supporting information.

The results of the calculations of these 12 series/trees are reported in Table 1.

Table 1. Comparison of all the 1-CD series and of four of the 24 possible 2-CD series (for each series/tree, average and best ΔE (in kJ/mol), generations reached, methylation rate (average Me per glucose unit), number of calculated samples and free CDs, ...), calculated with the semiempirical PM3 quantum-mechanical method.

Series	1-CD S1	1-CD S2	1-CD S3	1-CD S4	1-CD S5	1-CD S6	1-CD S7	1-CD S8
Moiety-side ^A	Fc NS	Fc WS	Ph1 NS	Ph1 WS	Ph2 NS	Ph2 WS	Succ NS	Succ WS
β CD G0 ^B	2-Me	3-Me	2-Me	3,6-diMe	2-Me	6-Me	2,3-diMe	3,6-diMe
ΔE G0 ^C	-122	-83	-111	-141	-48	-49	-161	-111
Average ΔE ^D	-154	-144	-127	-177	-68	-98	-187	-274
Best ΔE ^E	-194	-261	-142	-204	-127	-169	-219	-302
Max G/Gbest ^F	33/29	19/18	13/12	13/10	12/11	4/4	18/17	21/16
Average Me ^G	1.4	1.07	1.09	1.73	1.04	1.31	1.34	0.98
Me Domain ^H	0.6–2.4	0.3–2.0	0.6–1.6	1.1–2.3	0.6–1.3	1.0–1.6	0.6–2.3	0.3–2.1
Fe-H(CD) ^I	191(0) 1(0)	497(118) 145(0)	0(0) 0(0)	60(0) 0(0)	0(0) 0(0)	1(0) 0(0)	929(0) 0(0)	3414(1664) 2(0)
NbMeFc ^J	1.44 0.89	1.60 2.37	0.53 1.57	1.26 2.88	0.02 0.14	1.05 0.52	1.41 1.30	0.93 1.33
Samples ^K	2726	1994	571	326	187	79	1106	1903

Table 1. Cont.

ID G0 ^L	1	2727	4721	5292	5618	5805	5884	6990
Free CDs ^M	2690	1877	550	305	166	57	1084	1872
Series	2-CD S1	2-CD S2	2-CD S3	2-CD S4				
Moiety-side 1 ^A	Fc NS	Fc NS	Fc NS	Succ NS				
Moiety-side 2 ^A	Succ WS	Ph2 WS	Ph1 WS	Ph1 WS				
β CD G0 ^B	3-Me	3-Me	2,3-diMe	2-Me				
ΔE G0 ^C	−278	−208	−203	−236				
Average ΔE ^D	−271	−293	−248	−283				
Best ΔE ^E	−330	−338	−296	−326				
Max G/Gbest ^F	22/19	15/15	12/11	27/26				
Average Me ^G	1.33, 1.36	1.03, 1.16	1.94, 1.88	0.97, 0.81				
Me Domain ^H	0.9–1.9	0.6–1.4	1.6–2.3	0.4–1.4				
Fe-H(CD) ^I	790(0) 58(0)	534(0) 24(0)	367(0) 48(0)	1175(0) 3(0)				
NbMeFc ^J	1.11 2.05	0.97 1.58	0.94 0.87	1.14 1.80				
Samples ^K	1138	642	449	1229				
ID G0 ^L	1	1139	1781	2230				
Free CDs ^M	495	253	195	451				

These data were extracted from the webpages generated by the PHP program and from the report files generated by the C program that can be consulted in the data repository [30]. (A) Moiety of SuccFerr inserted inside the cavity of the CD and entrance side: NS = narrow side, WS = wide side. (B) Well-defined CD used as the G0 starting system. For 2-CD systems, the same CD was used twice. (C) ΔE of the starting system G0 (Note A1). These values are the best ones of Tables S1 and S2. (D) Average ΔE of all the experiments (Note A2). (E) Best ΔE reached by the tree (Note A3). (F) Max G = higher generation reached by the tree (Note A4), Gbest = generation of the best experiment (Note A5). (G) Average Me = average methylation per glucose unit for systems better than G0 (for each of the two CDs for the 2-CD series, see details in SI). (H) Methylation-per-glucose-unit domains for systems better than G0 (for the 2-CD series, union of the overlapping domains of the two CDs) (see details in SI). (I) Code 'Fe-H(CD)' indicates, for the whole series, the total number of hydrogen bonds between Fe and a hydrogen atom of a CD for the best (first row) and worse (second row) systems, respectively (Notes A6, A7, A8). The values inside parenthesis indicate the total number of clamps with Fe. (J) Code 'NbMeFc' indicates the average number of Me close to the ferrocene moiety for the best (first row) and worse (second row) systems, respectively (Notes A7, A8, A9). (K) Number of calculated experiments for each series (Note A10). (L) ID (model number) of the G0 system. (M) Total number of distinct CDs, when freed from the supramolecular assemblage (Note A11).

3. Discussion

These calculations were carried out to determine which CDs should be the best fit for each moiety of SuccFerr. More particularly, these calculations were meant to serve, among other things, to determine the best suitable average methylation rate of these CDs.

Data collected in Table 1 and the representation of the variation in ΔE for the unique route going from G0 to the best model (called "main route", Figure S1) for each of the 12 series/trees (Figure 4) show the superiority of the 2-CD systems, even at the G0 level (i.e., with well-defined CDs), which is predictable because there are more stabilizing interactions. Among the G0 systems of 1-CD, series 7 (insertion of Succ via the narrow side), which was the best, is exceeded in these mutation trees, at G1 by series S8 (insertion of Succ from the wide side) and even at G6 by series 2 (insertion of Fc from the wide side). This shows that no conclusion can be made based on well-defined CDs, but also that undefined CDs can sometimes perform better, even though the RAME β CD mixture also contains CDs that are less efficient than well-defined CDs. At any rate, this behavior of 1-CD S8 and S2 merits further study.

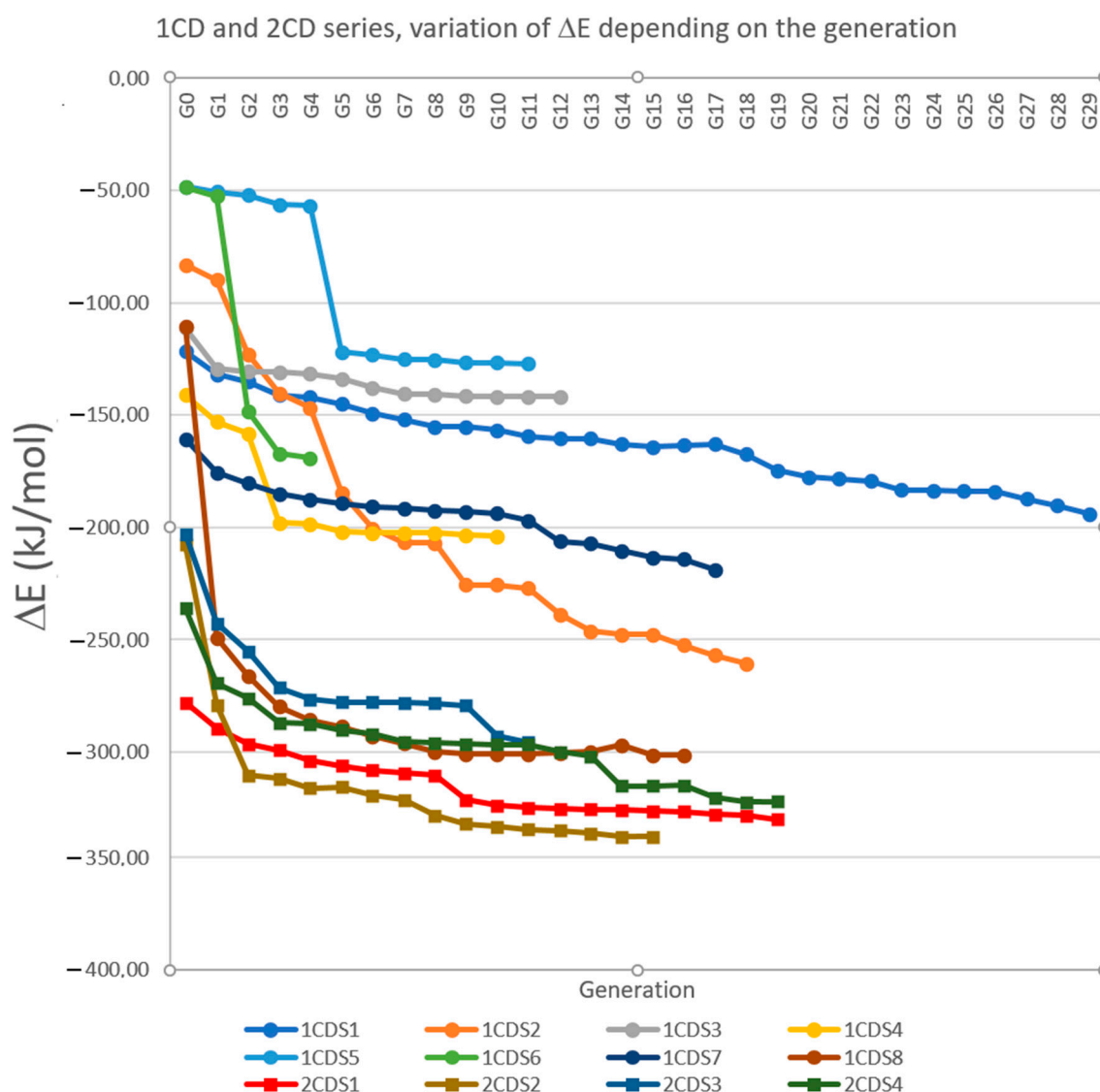


Figure 4. For the eight 1CD series (S1 to S8) and the four 2CD series (S1 to S4), unique route in the trees, starting from G0 and ending when reaching the best model of the series with display of the variation in ΔE (kJ/mol) at each generation. Some values can be found in Table 1: the starting ΔE (ΔE G0), the ending ΔE (ΔE of the best model: best ΔE), and the generation of the best model (final generation: Gbest). The PHP code to retrieve the data needed to fill this graphic from the database is explained in SI.

3.1. Observed Contribution of the Iron Atom to Stability

Although Table 1 and Figure 4 show a superiority of 2-CD over 1-CD (lower ΔE for G0, lower average ΔE , best lower ΔE of series), the 1-CD series S8 experiment shows very different ΔE from other 1-CD series and is in competition with the ΔE of the 2-CD. The webpage displaying the G0 model of 1-CD S8 and its first-generation descendants (G1) can be consulted to see the data (using the web application by typing its ID number (6990) into the form, or more simply displaying the static saved version “1-CD-webpage-ID-6990.html” file, available inside the repository [30]). In the list of the G1 descendants, experiment ID 7010 shows the strongest variation in ΔE (between itself and its parent, $\Delta\Delta E = -139$ kJ/mol) of all the 8892 calculated 1-CD experiments during the demethylation of the oxygen atom O₂₀. This brusque variation in ΔE can be seen in Figure 4 at G1. In addition, the generations derived from this experience all show a significant destabilization during the re-methylation of this atom (as in model ID 7169) but also of O₁₉ (for example,

ID 7017). These anomalies in the data can be easily detected by SQL queries ordering the experiments by their 'deltadelta' field (representing $\Delta\Delta E$ in the database, Note A12). It is worth mentioning that thanks to these anomalies in the data, two atypical hydrogen bonds between hydrogens atoms from O₁₉ and O₂₀ and the central ferrocene iron atom have been detected. This is characterized by the formation of a clamp ($-\text{O}_{19}-\text{H} \dots \text{Fe} \dots \text{H}-\text{O}_{20}-$), as can be seen with the best model of this series (Figure S5a). This type of bond, but intramolecular and without a clamp, has been reported with NH groups by an XRD analysis of a single crystal [33]. However, to the best of our knowledge, this is the first time that it has been described by modelling in supramolecular systems involving ferrocene and a cyclodextrin forming intermolecular Fe-H hydrogen bonds. This phenomenon is responsible for this decrease in energy, but also for its increase when the clamp is broken by methylating O₁₉ or O₂₀, both of which belong to the same anhydroglucose unit (positions 2 and 3 in the glucose numbering system). Another clamp ($-\text{O}_{10}-\text{H} \dots \text{Fe} \dots \text{H}-\text{O}_{11}-$, Figure S6) was also discovered in the 1-CD series S2, again associated with a strong positive value of 'deltadelta' ($\Delta\Delta E$) when O₁₀ or O₁₁ is methylated (for example, ID 3743 for O₁₀ and ID 3744 for O₁₁, seen when displaying the webpage of their common parent, model ID 3738). Unlike 1CD S8, where two Fe-H bonds (clamp) are formed at the same time, which explains the steep decrease in ΔE , the formation of the clamp for 1CD S2 is done gradually, as can be seen in Figure 4 (slower variation of ΔE , due to different stages passing through weaker Fe-H bonds formations). Even though both O₁₀ and O₁₁ were non-methylated since G1, this clamp is only formed at G9 due to the favorable influence of the modification of methylations managed by the web application. In a related way, the greatest increases in ΔE are not all due to the direct breaking of the clamp, but often also by indirect cleavage caused by steric hindrance of the latter during the methylation of an oxygen atom close to it, as methylation of O₁ for series S8 (ID 7011 and ID 7030, having $\Delta\Delta E > 176$ kJ/mol). In 2-CD systems, this phenomenon also exists, but the connection is single, or of an asymmetrical clamp type (one strong/short and one weak/long), probably because of the greater steric effect. Analyzing the geometries of all the 2-CD models, thanks to the C program (that does the same for 1-CD models) reading each of the XYZ files, we can confirm that no strong symmetrical clamp exists with Fe in any of these models, even though some simple bonds were found and are important for stability (Note A13).

Analyzing the geometries of the full dataset (XYZ files, using the C program) proved to be extremely useful in understanding the effect of the atypical and classic hydrogen bonds on the stability of supramolecular assemblies. We have systematically checked, for all the models, the existence, or not, of seven possible intermolecular hydrogen bonds. The maximum distance was set to 2 Å, and the results can be found in the provided Hbonds-1CD.txt [34] and Hbonds-2CD.txt [35] report files for 1-CD and 2-CD models, respectively. For SuccFerr, the atoms involved in these bonds are the iron atom, the two phenols (-O- and -H independently) and the two carbonyls (=O) of the imide group. For CDs, these are the hydroxyls (-O- and -H independently), the ethers and the anomeric group (-O-). For each series, the models were separated into two groups (worse models or best models, Note 8). The comparison of the number of each type of hydrogen bond found in these two groups showed that the effect is important especially when the iron atom is involved, even more so when it is implicated into a clamp (asymmetrical clamps (one bond > 2 Å and one bond < 2 Å) have been ignored). For phenols, the clamps only exist for the 2-CD series, whereas clamps involving the C=O groups were detected for both the 1-CD and 2-CD series. However, these two types of clamps do not carry as much weight in stability as those involving Fe (Note A13). To explain why, in the 2-CD models, no strong clamp involving the iron atom was detected, we can compare the 1-CD S8 (Figure 5) and 2-CD S1 series (which corresponds to the 1-CD S8 series with an additional CD in which Fc is inserted, Figure 6). We see on the models that this second CD impedes the approach of iron to two OH groups of the CD including Succ, as in the 1-CD S8 series, and instead makes a single bond with the iron atom (Figure 6).

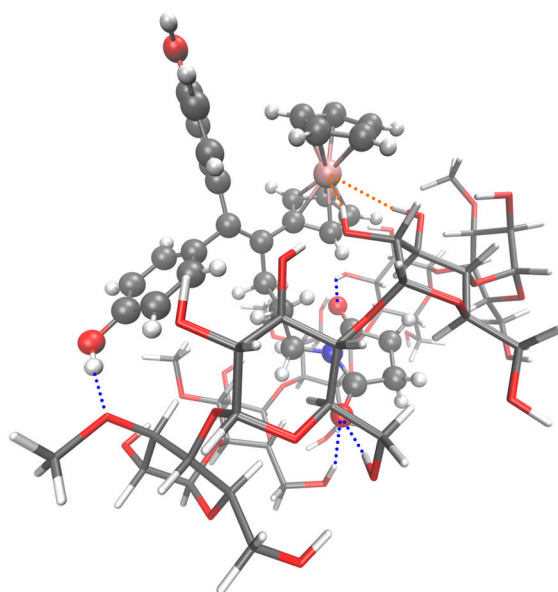


Figure 5. Molecular model of the best 1-CD system (series S8, ID 7759, Succ entering by the wide side of the CD) computed with Gaussian 16 at the DFT level of theory. SuccFerr is represented with a “ball-and-stick” model, and the CD is represented with a “licorice” model. Orange dashed lines (clamp between the iron atom and O₁₉-H and O₂₀-H) represent the two atypical hydrogen bonds between the iron atom and the CD. Another clamp between one of the C=O group and O₁₂-H and O₁₈-H is also visible, as bonds between the phenol group (Ph2) and O₁₄-Me and between the second C=O of imide and O₅-H (blue dashed lines).

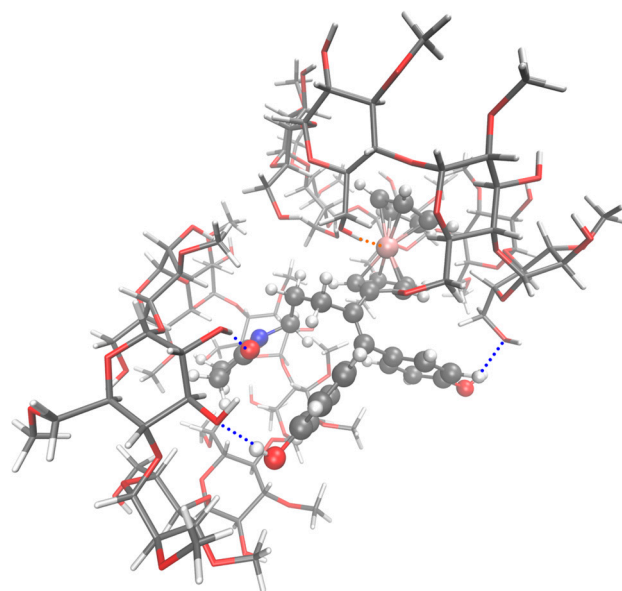


Figure 6. Molecular model of the best 2-CD series S1 system (ID 586, Succ entering by the wide side of CD1 and Fc entering by the narrow side of CD2) computed at the semiempirical PM3 level of theory. SuccFerr is represented with a “ball-and-stick” model, and CD1 and CD2 are represented with a “licorice” model. Orange dashed line is the hydrogen bond between the iron atom and O₃₆-H of CD2.

The lengths of the Fe-H bonds found with PM3 seeming surprisingly short (≈ 1.95 Å), the best 1-CD series 8 model was then computed in DFT (see computational details). The latter confirms the formation of this clamp with the iron atom (Figure 5), with bond lengths of 2.77 and 2.69 Å for O₁₉-H ... Fe and O₂₀-H ... Fe, respectively (the corresponding angles

are 159° and 166°). The hydrogen bond acceptors' and donors' distances are 3.69 and 3.64 Å for O₁₉,Fe and O₂₀,Fe. The calculation of the ferrocene + methanol models in water confirms the presence of Fe-H bonds in DFT in aqueous medium (in the compressed archive DFT.zip [36]). A further Natural Bond Orbitals (NBO) calculation has allowed us to estimate the stabilizing energy ($E^{(2)}$, see Table 2) of the atypical H-bonds found between the Fe from the SuccFerr compound and the alcohols from the CD. Adding all the NBO (donor–acceptor) energy contributions (see Table S4 in SI for further details) results in a total stabilisation energy of 86.45 kJ/mol (20.66 kcal/mol) and 119.24 kJ/mol (28.50 kcal/mol) for O₁₉-H...Fe and O₂₀-H...Fe, respectively. It is worth noting that in such particular supramolecular contacts, we observe not only a weak LP(Fe)- σ^* (H-O) intermolecular interaction (3.80 kJ/mol) comparable to that which can be seen in common H-bond interactions, but we also observe σ (H-O)-LP*(Fe) stabilizing contributions, as well as quite uncommon strong (56.77 kJ/mol) LP*(Fe)-Rydberg (H) contributions among some other minor contributions.

Table 2. Second-order perturbation stabilization energy contributions to the atypical H-bond Fe-HO interactions. NBO labels: LP = lone pair, LP* = antibonding lone pair, CR = core, σ^* =Antibonding sigma bond.

NBO Donor	NBO Acceptor	$E^{(2)}$ (kJ/mol)
Fe ---- H_O19		
LP _{All} (Fe)	Rydberg _{All} (H)	1.13
LP _{All} (Fe)	σ^* _{All} (H-O)	3.80
LP* _{All} (Fe)	Rydberg _{All} (H)	56.77
LP* _{All} (Fe)	Rydberg _{All} (O)	9.23
LP* _{All} (Fe)	σ^* _{All} (H-O)	4.48
σ _{All} (H-O)	LP* _{All} (Fe)	9.83
σ _{All} (H-O)	Rydberg _{All} (Fe)	0.29
LP _{All} (O)	LP* _{All} (Fe)	0.92
Fe ---- H_O20		
CR _{All} (Fe)	Rydberg _{All} (H)	0.21
LP _{All} (Fe)	Rydberg _{All} (H)	3.14
LP _{All} (Fe)	σ^* _{All} (H-O)	2.90
LP* _{All} (Fe)	Rydberg _{All} (H)	98.40
LP* _{All} (Fe)	σ^* _{All} (H-O)	4.43
σ _{All} (H-O)	LP* _{All} (Fe)	8.61
σ _{All} (H-O)	Rydberg _{All} (Fe)	0.50
LP _{All} (O)	LP* _{All} (Fe)	1.05

The PM3 and DFT methods gave comparable results (see Table 3 for the lengths and angles of some hydrogen bonds for the best model ID 7759).

Table 3. Lengths and angles of some hydrogen bonds for model 2CD S8 ID 7759 calculated by PM3 and DFT.

Type of Hydrogen Bond	Length PM3 (Å)	Length DFT (Å)	Angle PM3 (°)	Angle DFT (°)
O ₁₉ -H...Fe	1.933	2.774	167	159
O ₂₀ -H...Fe	1.896	2.694	177	166
C=O(1)...H(CD)	1.910	2.010	148	128
C=O(2)...H(CD) (first)	1.838	1.864	166	164
C=O(2)...H(CD) (second)	1.858	2.087	162	163
H(Ph2)...O(CD)	1.843	1.751	167	173
H...O intra CD (1)	1.863	1.874	163	173
H...O intra CD (2)	1.825	1.815	168	175
H...O intra CD (3)	1.862	1.868	172	164

Table 3. *Cont.*

Type of Hydrogen Bond	Length PM3 (Å)	Length DFT (Å)	Angle PM3 (°)	Angle DFT (°)
H...O intra CD (4)	1.816	1.836	174	175
H...O intra CD (5)	2.474	2.294	108	111
H...O intra CD (6)	1.862	1.868	172	164

O₁₉-H...Fe: first hydrogen bond of the clamp with Fe, O₂₀-H...Fe: second hydrogen bond of the clamp with Fe, C=O(1)...H(CD): hydrogen bond between the first CO of imide and H of the CD, C=O(2)...H(CD): hydrogen bond between the second CO of imide and H of the CD (first and second hydrogen bond of the clamp), H(Ph2)...O(CD): hydrogen bond between H of phenol 2 and O of CD, H...O intra CD (1 to 6): some selected intra-CD hydrogen bonds.

3.2. Contribution of Methyl Groups to Stability

One might think that since the ferrocenyl group is nonpolar, as are the phenyl groups, it could therefore interact better with methyl groups and prefer highly methylated CDs, or at least have many proximate methyl groups, as postulated by Buriez [37]. However, this is not reflected in our modelling for Fc, because the number of neighboring methyl groups is generally rather low, as shown in Figure 5. To understand the effect of the Fc-Me proximity on the stability, a systematic analysis has been performed by the C program (see code 'NbMeFc' in Table 1 and notes A7, A8, A9). We can see that the average number of these proximities are quite limited, and that there is no clear difference between best and worse models (except for 1CD S8 and S2 series, where some methyls should be removed to form the beneficial clamp with the iron).

Finding the best blend would not be of interest, as statistically, it only exists in small amounts in the RAME β CD mix and is closely followed by many other assemblies in terms of stability; moreover, it would be idealistic to seek to synthesize the complex corresponding CDs, merely to dissolve SuccFerr better. Thus, it is better to consider macroscopic values, such as the average degree of methylation of the ideal mixture. By "ideal mix", we mean that one should include in this calculation, for each series, only CDs participating in improved assemblies, that is, those for which $\Delta E < \Delta EG_0$. These average states of methylation are calculated by the PHP program and are given in Table 1. However, relying only on these average methylations is restrictive, as it is found that the ideal range for each run (lowest and highest methylation rate of all improved CDs in any given run) is large. We can therefore deduce that SuccFerr can adapt to RAME β CDs with very various methylation rates. The most important parameter is not this rate, but the distribution of CDs (in particular, the isomers), a parameter impossible to control during the synthesis of RAME β CD, or difficult to assess by analysis. In conclusion, it has been shown in this work that the solubility of SuccFerr with methylated CDs depends on many variables, and it is no simple task to determine the best methylation rate for the CDs.

4. Materials and Methods

4.1. Complexation Studies in Aqueous Phase

4.1.1. Phase Solubility Studies of SuccFerr Complexation

SuccFerr complexation with various CDs was evaluated using the phase-solubility method [26]. A suspension of a large excess of SuccFerr (30 mg) in 2 mL of aqueous solutions of the appropriate CD (concentrations ranging from 0.125 to 160 mM, pH adjusted to 7) was stirred in screw-capped amber vials for 24 h on a rock-and-roller agitator at 25 °C. Preliminary experiments indicated that equilibrium was reached after this 24 h stirring period. Each suspension was then centrifuged at 9000 g for 10 min and diluted from 1/10 to 1/1000 with acetonitrile, and the amount of dissolved SuccFerr was assessed by HPLC at 286 nm. Phase solubility curves (i.e., solubility of SuccFerr as a function of the CD concentration) were drawn for each CD.

4.1.2. UV–Vis Experiments—Benesi–Hildebrand Method

First, UV–vis spectra (Cary 50 spectrophotometer, Varian, Les Ulis, France) were obtained for a concentration of 2.5×10^{-5} M of SuccFerr and various concentrations of CDs (0 to 1.1×10^{-5} M). RAME β CD or HP β CD were added, and the SuccFerr concentration was kept at 2.5×10^{-5} M. All experiments were performed in 1% (*v/v*) DMSO/water mixture (the first dissolution of SuccFerr was performed in pure DMSO) to avoid precipitation, and the measurement was taken after 10 min to account for the kinetics of complex formation. Then, the mole-ratio titration method was used to calculate the values of the apparent binding constant (K_a) [38–40]. These experiments were performed at various wavelengths obtained from the spectra (low concentration of CD). Another series of experiments was performed with higher CD concentrations (2×10^{-6} – 25×10^{-5} M, or high concentration of CD) to confirm the binding constants.

4.2. Computer Methods

4.2.1. Software

Our experiments generated a large amount of data. We chose to create our own web application using the XAMPP pack (web server + database + PHP, version 8.2.0 was used) [32], which allowed us to design the very specific scheme required to store the data generated by our algorithms. We used this system to store our 13,261 models of inclusion of SuccFerr into one or two CD(s). The web browser serves as an interface between the scientist and our web application (entering data into a web form or displaying it). The PHP application acts as an intermediary with the MariaDB database, using SQL queries (see some examples in Appendix A), to store or access the data or even to generate statistics automatically.

The C program (to retrieve from the data repository [30]) was edited, compiled and executed using Code::Blocks version 20.03 64 bit [41]. It analyzes the XYZ files (checking for errors), creates the static webpages and performs additional statistics. More details can be found in SI.

4.2.2. Computational Details

All the SuccFerr- β -cyclodextrins complexes were computed using the program Spartan14 (Wavefunction, Irvine, CA, USA), performing a molecular mechanics optimization with the Merck molecular force field (MMFF) and a subsequent minimization with the semiempirical PM3 quantum-mechanical method. Starting from the best PM3 optimized geometry, a minimization was carried out using the Polarizable Continuum Model (PCM) in water. Density Functional Theory (DFT) was employed to perform this calculation using the ω B97XD functional with the diffuse-augmented polarization valence-triple- ζ (6-311G(d,p)) basis set including a set of p polarization functions for the hydrogen atoms and a set of d polarization functions for the second-row elements. For the iron atom, the uncontracted triple- ζ quality LANL08 basis set with an effective core potential (including 10 core electrons) was used in these calculations. This calculation was performed with the Gaussian 16 quantum package (Gaussian Inc., Wallingford, Connecticut, USA). On top of these DFT calculations, further NBO analyses were performed using the NBO version 3 integrated in Gaussian [42].

The PM3 method was used to determine the affinity of SuccFerr with the CD (ΔE , Equation (2) or Equation (3)):

$$\text{for 1-CD: } \Delta E = E_{(\text{SuccFerr}+\text{CD})} - E_{\text{SuccFerr}} - E_{\text{CD1}} \quad (2)$$

$$\text{for 2-CD: } \Delta E = E_{(\text{SuccFerr}+\text{CD1}+\text{CD2})} - E_{\text{SuccFerr}} - E_{\text{CD1}} - E_{\text{CD2}} \quad (3)$$

The energy of each element was calculated (energy method, for the ground state, no modification of geometry that should be the same as in the assemblage). The heat of formation (in kJ/mol) of each element and of the assemblage were copied from Spartan

and pasted into the form of the webpage of the web application that calculated ΔE , and this was saved in the database.

5. Conclusions

This work has shown that the RAME β CD mixture seems more suited than well-defined CDs to form stable associations with SuccFerr, and that the most probable assemblages are SuccFerr systems with two CDs, probably with a certain proportion of 1-CD series S8. The quantification results agree with the modelling showing that there are 2-CD assemblies. The modelling also revealed the special behavior of ferrocene, which forms atypical hydrogen bonds (one or two in the form of a clamp) between its iron atom and the hydrogen atoms of the hydroxyl groups of CDs. This result is particularly important because it has been possible, for the first time, to the best of our knowledge, to model the atypical hydrogen bonds between CDs and an anticancer molecule based on nonpolar ferrocene. These interactions are, in fact, responsible for the high affinity observed between the CDs and this molecule. Methyl groups also provide stability or instability depending on their position. For example, two OH groups at position 2 and 3 of the same glucose unit are not a sufficient parameter to allow a clamp with Fe, since some Me and OH groups should be placed at the right places elsewhere (6-Me- β CD in Table S1 does not have this clamp despite having seven glucose units with OH in position 2 and 3). Our method using trees of modifications of methylation permitted us to reach these particular configurations, and the discovery of these atypical bonds is due to the proper functioning of the experimental web application and to the automatic analysis of the model files by the C program.

Supplementary Materials: The following supporting information can be downloaded at: <https://www.mdpi.com/article/10.3390/ijms241512288/s1>.

Author Contributions: Conceptualization: P.P.; Data curation: P.P. and S.G.; Formal analysis: P.P., F.N. and S.G.; Funding acquisition: G.J. and S.G.; Investigation, P.P., F.N. and S.G.; Methodology, P.P., F.N. and S.G.; Project administration: S.G.; Resources: P.P.; Software: P.P.; Supervision: S.G.; Validation P.P., S.G. and F.N.; Visualization: P.P. and J.S.G.; Writing—original draft: P.P., M.J.M. and S.G.; Writing—review and editing: P.P., M.J.M., G.J., S.G. and J.S.G. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by Agence Nationale de la Recherche, ANR PRCE NaTeMOc no. ANR-19-CE18-0022-01. The APC was funded by the startup FEROSCAN (G. Jaouen is member of this startup).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: A part of the data supporting this article is available within the article and Supplementary Materials (Tables S1–S4 and more detailed descriptions of the program and examples of SQL queries adapted to our database). The main part of the data is publicly accessible in the ‘Recherche Data Gouv’ repository (space of Sorbonne Université) at <https://doi.org/10.57745/CBUPP3> (accessed on 22 April 2023) [30]. This data repository contains the database in various formats (CSV, SQL (preferred to import the database [43]), JSON and XML), 13361 models in XYZ format, 28 original Spartan files, the S8 DFT geometry in XYZ format, the files created by analysis of the models by the C program, the C program itself and the PHP program. See the README.txt file for details [44]. The PHP program was also referenced on Software Heritage via Hal [31].

Acknowledgments: The DFT simulations on Gaussian 16 were performed on the SACADO MeSU platform at Sorbonne Université. The authors thank Héloïse Dossman for assistance on MeSU.

Conflicts of Interest: The authors declare no conflict of interest.

Appendix A

Notes in the text and useful SQL requests to manage the database (replace 's' by the wanted series and 'calculscd' (name of the table of the database for 1-CD systems) by 'calculs2cd' for 2-CD systems).

Note A1. To recover the ΔE of a starting G0 experiment of series s: `SELECT deltaE FROM calculscd WHERE series='s' AND generation='0' LIMIT 1.`

Note A2. To calculate the average ΔE of a certain series s: `SELECT AVG(deltaE) FROM calculscd WHERE series='s'.`

Note A3. To calculate the best ΔE of a certain series s: `SELECT MIN(deltaE) FROM calculscd WHERE series='s'.`

Note A4. To recover the maximum generation of a tree of series s reached: `SELECT MAX(generation) FROM calculscd WHERE series='s'.`

Note A5. To recover the generation of the best experiment in a tree of series s: `SELECT generation FROM calculscd WHERE series='s' AND deltaE=(SELECT MIN(deltaE) FROM calculscd WHERE series='s').`

Note A6. The limit distance to determinate whether there is a hydrogen bond with Fe was set to 2 Å.

Note A7. These parameters were calculated by the C program that generated the raw data inside files 'Hbonds-1CD.txt' (1-CD models [34]), 'Hbonds-2CD.txt' (2-CD models [35]) and the formatted statistics inside webpages 'TableofHbondsbySeries-1CD.html' (1-CD models) and 'TableofHbondsbySeries-2CD.html' (2-CD models). These files are furnished inside the repository [30].

Note A8. To determinate whether a model belongs to the best or the worse part of the series, the limit ΔE was calculated as the average of the extreme ΔE values for this series.

Note A9. To count the average number of Me close to ferrocene: All Me were searched into the XYZ files (C atom connected to 3 H), the Me was counted as close to ferrocene if the distance between its C atom and the Fe atom is less than 7 Å, and the total number was divided by the number of models (for both best and worse models).

Note A10. To calculate the number of experiments of a certain series s: `SELECT COUNT(*) FROM calculscd WHERE series='s'.`

Note A11. To find the number of free CDs of a series (when freed from supramolecular assembly, see more details in SI: "Algorithm to calculate binarymin" and Figure S3).

Note A12. To find the higher negative variations in ΔE ($\Delta\Delta E$) by sorting the experiments based on 'deltadelta': `SELECT * FROM calculscd ORDER BY deltaxdelta.` The same for the higher positive variations of ΔE : `SELECT * FROM calculscd ORDER BY deltaxdelta DESC.`

Note A13. See static webpages "TableofHbondsbySeries-1CD.html" and "TableofHbondsbySeries-2CD.html", generated by the C program inside the compressed archive 'webpages.zip' [29], or download the whole dataset [30].

References

1. Wimmer, T. Cyclodextrins. In *Ullmann's Encyclopedia of Industrial Chemistry*, 5th ed.; Bohner, M., Ullmann, F., Eds.; Wiley-VCH: Weinheim, Germany, 2003.
2. Bakó, P.; Fenichel, L.; Töke, L.; Szente, L.; Szejtli, J. Methylation of cyclodextrins by phase-transfer catalysis. *J. Incl. Phenom. Mol. Recogn. Chem.* **1994**, *18*, 307–314. [[CrossRef](#)]
3. Szejtli, J. *Cyclodextrins and Their Inclusion Complexes*; Akadémiai Kiadó: Budapest, Hungary, 1982.
4. Arun, R.; Ashok, K.C.K.; Sravanthi, V.V.N.S.S. Cyclodextrins as Drug Carrier Molecule: A Review. *Sci. Pharm.* **2008**, *76*, 567–598. [[CrossRef](#)]
5. Conceicao, J.; Adeoye, O.; Cabral-Marques, H.M.; Sousa Lobo, J.M. Hydroxypropyl- β -Cyclodextrin and β -Cyclodextrin as Tablet Fillers for Direct Compression. *AAPS PharmSciTech.* **2018**, *19*, 2710–2718. [[CrossRef](#)] [[PubMed](#)]
6. Hafner, V.; Czock, D.; Burhenne, J.; Riedel, K.-D.; Bommer, J.; Mikus, G.; Machleidt, C.; Weinreich, T.; Haefeli, W.E. Pharmacokinetics of Sulfobutylether-Beta-Cyclodextrin and Voriconazole in Patients with End-Stage Renal Failure during Treatment with Two Hemodialysis Systems and Hemodiafiltration. *Antimicrob. Agents Chemother.* **2010**, *54*, 2596–2602. [[CrossRef](#)] [[PubMed](#)]
7. Johannsdottir, S.; Jansook, P.; Stefansson, E.; Loftsson, T. Development of a cyclodextrin-based aqueous cyclosporin A eye drop formulations. *Int. J. Pharm.* **2015**, *493*, 86–95. [[CrossRef](#)] [[PubMed](#)]

8. Yang, P.; Li, Y.; Li, W.; Zhang, H.; Gao, J.; Sun, J.; Yin, X.; Zheng, A. Preparation and evaluation of carfentanil nasal spray employing cyclodextrin inclusion technology. *Drug Dev. Ind. Pharm.* **2018**, *44*, 953–960. [[CrossRef](#)] [[PubMed](#)]
9. Khan, A.R.; Forgo, P.; Stine, K.J.; D'Souza, V.T. Methods for Selective Modifications of Cyclodextrins. *Chem. Rev.* **1998**, *98*, 1977–1996. [[CrossRef](#)]
10. Legrand, F.-X.; Sauthier, M.; Flahaut, C.; Hachani, J.; Elfakir, C.; Fourmentin, S.; Tilloy, S.; Monflier, E. Aqueous hydroformylation reaction mediated by randomly methylated β -cyclodextrin: How substitution degree influences catalytic activity and selectivity. *J. Mol. Catal. A Chem.* **2009**, *303*, 72–77. [[CrossRef](#)]
11. Vessières, A.; Top, S.; Pigeon, P.; Hillard, E.A.; Boubekour, L.; Spera, D.; Jaouen, G. Modification of the Estrogenic Properties of Diphenols by the Incorporation of Ferrocene. Generation of Antiproliferative Effects in Vitro. *J. Med. Chem.* **2005**, *48*, 3937–3940. [[CrossRef](#)]
12. Bruyère, C.; Mathieu, V.; Vessières, A.; Pigeon, P.; Top, S.; Jaouen, G.; Kiss, R. Ferrocifen derivatives that induce senescence in cancer cells: Selected examples. *J. Inorg. Biochem.* **2014**, *141*, 144–151. [[CrossRef](#)]
13. Tonolo, F.; Salmain, M.; Scalcon, V.; Top, S.; Pigeon, P.; Folda, A.; Caron, B.; McGlinchey, M.J.; Toillon, R.A.; Bindoli, A.; et al. Small Structural Differences Between Two Ferrocenyl Diphenols Determine Large Discrepancies of Reactivity and Biological Effects. *ChemMedChem* **2019**, *14*, 1717–1726. [[CrossRef](#)]
14. Wang, Y.; Pigeon, P.; Top, S.; McGlinchey, M.J.; Jaouen, G. Organometallic Antitumor Compounds: Ferrocifens as Precursors to Quinone Methides. *Angew. Chem. Int. Ed. Engl.* **2015**, *54*, 10230–10233. [[CrossRef](#)]
15. Richard, M.-A.; Hamels, D.; Pigeon, P.; Top, S.; Dansette, P.M.; Lee, H.Z.S.; Vessières, A.; Mansuy, D.; Jaouen, G. Oxidative Metabolism of Ferrocene Analogues of Tamoxifen: Characterization and Antiproliferative Activities of the Metabolites. *ChemMedChem* **2015**, *10*, 981–990. [[CrossRef](#)] [[PubMed](#)]
16. Wang, Y.; Richard, M.-A.; Top, S.; Dansette, P.M.; Pigeon, P.; Vessières, A.; Mansuy, D.; Jaouen, G. Ferrocenyl Quinone Methide-Thiol Adducts as New Antiproliferative Agents: Synthesis, Metabolic Formation from Ferrociphenols, and Oxidative Transformation. *Angew. Chem. Int. Ed. Engl.* **2016**, *128*, 10587–10590. [[CrossRef](#)]
17. Citta, A.; Folda, A.; Bindoli, A.; Pigeon, P.; Top, S.; Vessières, A.; Salmain, M.; Jaouen, G.; Rigobello, M.P. Evidence for Targeting Thioredoxin Reductases with Ferrocenyl Quinone Methides. A Possible Molecular Basis for the Antiproliferative Effect of Hydroxyferrocifens on Cancer Cells. *J. Med. Chem.* **2014**, *57*, 8849–8859. [[CrossRef](#)]
18. Pigeon, P.; Wang, Y.; Top, S.; Najlaoui, F.; Garcia Alvarez, M.; Bignon, J.; McGlinchey, M.J.; Jaouen, G. A new series of succinimido-ferrociphenols and related heterocyclic species induce strong antiproliferative effects, especially against ovarian cancer cells resistant to cisplatin. *J. Med. Chem.* **2017**, *60*, 8358–8368. [[CrossRef](#)] [[PubMed](#)]
19. Wang, Y.; Pigeon, P.; Top, S.; García, J.; Troufflard, C.; Ciofini, I.; McGlinchey, M.J.; Jaouen, G. Atypical lone pair- π interaction with quinone methides in a series of imido-ferrociphenol anticancer drug candidates. *Angew. Chem. Int. Ed. Engl.* **2019**, *58*, 8421–8425. [[CrossRef](#)]
20. Vessières, A.; Quissac, E.; Lemaire, N.; Alentorn, A.; Domeracka, P.; Pigeon, P.; Sanson, M.; Idbaih, A.; Verreault, M. Heterogeneity of Response to Iron-Based Metallodrugs in Glioblastoma Is Associated with Differences in Chemical Structures and Driven by FAS Expression Dynamics and Transcriptomic Subtypes. *Int. J. Mol. Sci.* **2021**, *19*, 10404. [[CrossRef](#)]
21. Jaouen, G.; Pigeon, P.; Top, S. Metallocene Derivatives with Anticancer Activity. Patent WO2015063201(A1), 7 May 2015.
22. Idlas, P.; Lepeltier, E.; Bastiat, G.; Pigeon, P.; McGlinchey, M.J.; Lautram, N.; Vessières, A.; Jaouen, G.; Passirani, C. Physicochemical Characterization of Ferrocifen Lipid Nanocapsules: Customized Drug Delivery Systems Guided by the Molecular Structure. *Langmuir* **2023**, *39*, 1885–1896. [[CrossRef](#)]
23. Najlaoui, F.; Pigeon, P.; Aroui, S.; Pezet, M.; Sancey, L.; Marrakchi, N.; Rhouma, A.; Jaouen, G.; De Waard, M.; Busser, B.; et al. Anticancer properties of lipid and poly(ϵ -caprolactone) nanocapsules loaded with ferrocenyl-tamoxifen derivatives. *J. Pharm. Pharmacol.* **2018**, *70*, 1474–1484. [[CrossRef](#)]
24. Idlas, P.; Ladacyia, A.; Némati, F.; Lepeltier, E.; Pigeon, P.; Jaouen, G.; Decaudin, D.; Passirani, C. Ferrocifen Stealth LNCs and Conventional Chemotherapy: A Promising Combination against Multidrug-Resistant Ovarian Adenocarcinoma. *Int. J. Pharm.* **2022**, *626*, 122164. [[CrossRef](#)] [[PubMed](#)]
25. Najlaoui, F.; Busser, B.; Taiwe, G.S.; Pigeon, P.; Sturm, N.; Giovannini, D.; Marrakchi, N.; Rhouma, A.; Jaouen, G.; Gibaud, S.; et al. Succinimido-Ferrociphenol Complexed with Cyclodextrins Inhibits Glioblastoma Tumor Growth In Vitro and In Vivo without Noticeable Adverse Toxicity. *Molecules* **2022**, *14*, 4651. [[CrossRef](#)] [[PubMed](#)]
26. Higuchi, T.; Connors, K. Phase Solubility Techniques. *Adv. Anal. Chem. Instrum.* **1965**, *4*, 117–212.
27. Najlaoui, F.; Pigeon, P.; Abdelkafi, Z.; Leclerc, S.; Durand, P.; El Ayeb, M.; Marrakchi, N.; Rhouma, A.; Jaouen, G.; Gibaud, S. Phthalimido-ferrociphenol cyclodextrin complexes: Characterization and anticancer activity. *Int. J. Pharm.* **2015**, *491*, 323–334. [[CrossRef](#)]
28. Del Valle, E.M.M. Cyclodextrins and their uses: A review. *Process Biochem.* **2004**, *39*, 1033–1046. [[CrossRef](#)]
29. Recherche Data Gouv (to Download File 'webpages.zip' and All Webpages within). Available online: <https://entrepot.recherche.data.gouv.fr/api/access/datafile/:persistentId?persistentId=doi:10.57745/COGDRQ> (accessed on 22 April 2023).
30. Pigeon, P. *Manipulation of a Cyclodextrin-Ferrociphenol Supramolecular Models Database Using a Web Application*; Recherche Data Gouv, V2; Recherche Data Gouv, Ministère de l'enseignement Supérieur et de la Recherche: Paris, France, 2023. [[CrossRef](#)]

31. Pigeon, P. *CDModelTree (V1): Manipulation of a Cyclodextrin-Ferrociphenol Supramolecular Models Database Using a PHP Web Application*; Software Heritage, INRIA: Le Chesnay-Rocquencourt, France, 2022. Available online: <https://hal.science/hal-03991394> (accessed on 22 April 2023).
32. XAMPP. Available online: <https://www.apachefriends.org> (accessed on 22 April 2023).
33. Förster, C.; Veit, P.; Ksenofontov, V.; Heinz, K. Diferrocenyl tosyl hydrazone with an ultrastrong NH \cdots Fe hydrogen bond as double click switch. *Chem. Commun.* **2015**, *51*, 1514–1516. [CrossRef]
34. Recherche Data Gouv (to Download File 'Hbonds-1CD.txt'). Available online: <https://entrepot.recherche.data.gouv.fr/api/access/datafile/:persistentId?persistentId=doi:10.57745/DQ8QXL> (accessed on 22 April 2023).
35. Recherche Data Gouv (to Download File 'Hbonds-2CD.txt'). Available online: <https://entrepot.recherche.data.gouv.fr/api/access/datafile/:persistentId?persistentId=doi:10.57745/WK5WE0> (accessed on 22 April 2023).
36. Recherche Data Gouv (to Download File 'DFT.zip' and All XYZ Models for DFT within). Available online: <https://entrepot.recherche.data.gouv.fr/api/access/datafile/:persistentId?persistentId=doi:10.57745/G2KRYZ> (accessed on 22 April 2023).
37. Buriez, O.; Heldt, J.M.; Labbé, E.; Vessières, A.; Jaouen, G.; Amatore, C. Reactivity and antiproliferative activity of ferrocenyl-tamoxifen adducts with cyclodextrins against hormone-independent breast-cancer cell lines. *Chem. Eur. J.* **2008**, *14*, 8195–8203. [CrossRef]
38. Benesi, H.A.; Hildebrand, J.H. A Spectrophotometric Investigation of the Interaction of Iodine with Aromatic Hydrocarbons. *J. Am. Chem. Soc.* **1949**, *71*, 2703–2707. [CrossRef]
39. Dotsikas, Y.; Kontopanou, E.; Allagiannis, C.; Loukas, Y.L. Interaction of 6-p-toluidinyl naphthalene-2-sulphonate with β -cyclodextrin. *J. Pharm. Biomed. Anal.* **2000**, *23*, 997–1003. [CrossRef]
40. Yoe, J.H.; Jones, A.L. Colorimetric Determination of Iron with Disodium-1,2-dihydroxybenzene-3,5-disulfonate. *Ind. Eng. Chem. Anal. Ed.* **1944**, *16*, 111–115. [CrossRef]
41. Code::Blocks. Available online: <https://www.codeblocks.org> (accessed on 22 April 2023).
42. Foster, J.P.; Weinhold, F. Natural hybrid orbitals. *J. Am. Chem. Soc.* **1980**, *102*, 7211–7218. [CrossRef]
43. Recherche Data Gouv (to Download File 'pascal.sql'). Available online: <https://entrepot.recherche.data.gouv.fr/api/access/datafile/:persistentId?persistentId=doi:10.57745/38MYXM> (accessed on 22 April 2023).
44. Recherche Data Gouv (to Download File 'README.txt'). Available online: <https://entrepot.recherche.data.gouv.fr/file.xhtml?persistentId=doi:10.57745/MDGGKE&version=2.0> (accessed on 22 April 2023).

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.

Supporting information

Unravelling the role of uncommon hydrogen bonds in cyclodextrin ferrociphenol supramolecular complexes: a computational modelling and experimental study

Pascal Pigeon ^{1,2,*}, Feten Najlaoui ³, Michael J. McGlinchey ⁴, Juan Sanz García ⁵, Gérard Jaouen ^{1,2} and Stéphane Gibaud ³

¹ PSL, Chimie ParisTech, 11 rue Pierre et Marie Curie, Paris F75231 Paris cedex 05, France

² Sorbonne Université, Institut Parisien de Chimie Moléculaire (IPCM) – UMR 8232, 4 place Jussieu, 75252 Paris Cedex 05, France; pascal.pigeon@sorbonne-universite.fr

³ Université de Lorraine, EA 3452/CITHEFOR, 5 rue Albert Lebrun (Faculté de Pharmacie), F-54000 Nancy, France

⁴ UCD School of Chemistry, University College Dublin Belfield, Dublin 4, Ireland

⁵ IMSME, Université Gustave Eiffel, CNRS, 5 Boulevard Descartes 77454 Marne-la-Vallée, France

Links:

All files needed for this work can be found into the data repository at <https://doi.org/10.57745/CBUPP3>. This includes the PHP program, the C program, the database, the static webpages, and other files (follow the link above and read the README.txt file for more details).

The PHP program (named CDMoDelTree) was also deposited into a forge (Software Heritage) using Hal:

Pascal Pigeon. CDMoDelTree (V1): Manipulation of a cyclodextrin-ferrociphenol supramolecular models database using a PHP web application. 2022, `{swh:1:dir:5c05be4fc1a1a2d4ca16bc48323231bb28379dec;origin=https://hal.archives-ouvertes.fr/hal-03991394;visit=swh:1:snp:438a3325327d8bef3a099271997a0dbdeca32da2;anchor=swh:1:rel:4ec1d7207a58cc63963137f2e79030f04cfdec96;path=/}`. `{hal-03991394}`

SUMMARY:	page
Modelling (tables S1 and S2)	3
Classification trees for models of SuccFerr- β CD assemblies (Fig. S1-S2)	7
Web application	16
Database and SQL	18
The PHP program and the webpages (table S3)	27
Installation of XAMPP, of the database and of the PHP software	29
Useful algorithms (Fig. S3)	30
Case of 1-CD series 9 and 10	39
Figure S4. Algorithm of the two PHP programmes	41
Examples without XAMPP (static webpages)	42
C program to control the models	42
Case of the best models of 1CD series 8 and series 2 (fig. S5-S6)	49
Table S4. Second-order perturbation stabilization energies for the NBOs	51
Description of the trees of modelling and of the web application to manage them	54

Modelling

Molecular Modelling calculations with the semiempirical PM3 quantum-mechanical method were performed using the 64-bit program Spartan 14 version 1.1.8 (Wavefunction Co., Irvine, CA, USA) with a Dell Precision T5810 PC (equipped with an Intel® Xeon® CPU E5-1630 v3 at 3.70 GHz, 4 cores, 8 logical CPU, 16 Gb RAM memory, 64 bits Windows 10). DFT calculations were performed using Spartan 20 on a workstation or Gaussian 09 on the supercomputer MeSU at Sorbonne Université, Paris (<https://sacado.sorbonne-universite.fr/mesu/>).

Each time a model was modified, a geometry optimization was carried out using the Merck molecular force field (MMFF) method followed by the semiempirical PM3 quantum-mechanical method. The semiempirical PM3 quantum-mechanical method was also used to determine the affinity of SuccFerr with the CD. This allowed one to evaluate the affinity of each of the 4 substituents of the alkene double bond for each side of each CD. In this SI, wdMβCD is an abbreviation of well-defined Methyl βCD (all glucose units are identical).

Typical method for inclusion of SuccFerr into well-defined βCDs

Each of the six tested CDs (mono-methylated: 2-Me-βCD, 3-Me-βCD, 6-Me-βCD and dimethylated: 2,3-diMe-βCD, 2,6-diMe-βCD and 3,6-diMe-βCD) were built separately and then optimized as described above. SuccFerr and the CD were copied and pasted into the same new window with forced inclusion of one of the four moieties of the double bond of SuccFerr approximately on the axis of the CD. Each moiety of the SuccFerr molecule, i.e., ferrocenyl (Fc), succinimidylpropyl (Succ), phenol #1 -*cis* with the ferrocenyl (Ph1) - and phenol #2 (Ph2) was included on each side of the CD (wide and narrow side) and built in separated files.

Typical method for inclusion of SuccFerr into two well-defined βCDs

To limit the number of possibilities the two wdMβCDs were identical (the six previous wdMβCDs, even for the special experiment CDxCD that gave two identical CDs as the best G0, see below) and only four combinations were calculated (Fc-Ph₁, Fc-Ph₂, Fc-Succ, Ph₁-Succ). However, the four possible combinations of inclusion were calculated (wide side – wide side, narrow side – narrow side, wide side – narrow side, narrow side – wide side). Files of previous experiments with one CD (CD1) on Fc or Ph₁ were duplicated, and in each copy, a second CD (CD2) was inserted

by copying and pasting the CD1 and placing by forced inclusion one of the remaining 3 substituents (Ph1, Ph2 or Succ for Fc and Succ for Ph1) with an eventual flipping (wide side or narrow side). The affinity required calculation of the energies of the reactions: SuccFerr + CD1 + CD2 \rightarrow molecular assembly ($\Delta E = E_{\text{SuccFerr-CD1-CD2}} - E_{\text{CD1}} - E_{\text{CD2}} - E_{\text{SuccFerr}}$).

Table S1. Calculated 1-CD systems (series S1 to S8) for inclusion into narrow or wide side of each of the moieties of SuccFerr into well-defined CDs with monomethylated CDs (2-Me- β CD, 3-Me- β CD, 6-Me- β CD) and dimethylated CDs (2,3-diMe- β CD, 2,6-diMe- β CD, 3,6-diMe- β CD). ΔE in kJ/mol.

Series 1, narrow side, Fc ¹	ΔE	Series 2, wide side, Fc	ΔE
2-Me-βCD ⁵	- 122	2-Me- β CD	- 58
3-Me- β CD	- 97	3-Me-βCD	- 83
6-Me- β CD	- 76	6-Me- β CD	- 43
2,3-diMe- β CD	- 96	2,3-diMe- β CD	- 53
2,6-diMe- β CD	- 73	2,6-diMe- β CD	- 52
3,6-diMe- β CD	- 68	3,6-diMe- β CD	- 75
Series 3, narrow side, Ph1 ²	ΔE	Series 4, wide side, Ph1	ΔE
2-Me-βCD	- 111	2-Me- β CD	- 52
3-Me- β CD	- 34	3-Me- β CD	- 47
6-Me- β CD	- 46	6-Me- β CD	- 53
2,3-diMe- β CD	- 42	2,3-diMe- β CD	- 41
2,6-diMe- β CD	- 72	2,6-diMe- β CD	- 40
3,6-diMe- β CD	- 38	3,6-diMe-βCD	- 141
Series 5, narrow side, Ph2 ³	ΔE	Series 6, wide side, Ph2	ΔE
2-Me-βCD	- 48	2-Me- β CD	- 37

3-Me- β CD	- 36	3-Me- β CD	- 44
6-Me- β CD	- 21	6-Me-βCD	- 49
2,3-diMe- β CD	- 34	2,3-diMe- β CD	- 40
2,6-diMe- β CD	- 38	2,6-diMe- β CD	- 28
3,6-diMe- β CD	- 43	3,6-diMe- β CD	- 31
Series 7, narrow side, Succ ⁴	ΔE	Series 8, wide side, Succ	ΔE
2-Me- β CD	- 61	2-Me- β CD	- 53
3-Me- β CD	- 40	3-Me- β CD	- 81
6-Me- β CD	- 41	6-Me- β CD	- 50
2,3-diMe-βCD	- 161	2,3-diMe- β CD	- 51
2,6-diMe- β CD	- 74	2,6-diMe- β CD	- 40
3,6-diMe- β CD	- 45	3,6-diMe-βCD	- 111

¹ Fc: ferrocene, ² Ph1: phenol 1 (*cis* to Fc), ³ Ph2: phenol 2 (*trans* to Fc), ⁴ Succ: succinimidylpropyl, ⁵ The best system and value for each series is displayed in bold and is the starting systems (G0) for calculation for diversely methylated cyclodextrins calculations.

Table S2. Calculated 2-CD systems: Inclusion of 2 moieties of SuccFerr into the wide side (WS) or into the narrow side (NS) of monomethylated wdM β CDs (2-Me- β CD, 3-Me- β CD, 6-Me- β CD) or dimethylated wdM β CDs (2,3-diMe- β CD, 2,6-diMe- β CD, 3,6-diMe- β CD). ΔE in kJ/mol.

Monomethylated		Dimethylated	
2-Me- β CD	ΔE	2,3-diMe- β CD	ΔE
Fc(WS)+Ph1(WS) ¹	- 132	Fc(WS)+Ph1(WS)	- 104
Fc(WS)+Ph2(WS)	- 150	Fc(WS)+Ph2(WS)	- 122
Fc(WS)+succ(WS)	- 118	Fc(WS)+succ(WS)	- 112
Ph1(WS)+succ(WS)	- 116	Ph1(WS)+succ(WS)	- 93

Fc(NS)+Ph1(NS)	- 176	Fc(NS)+Ph1(NS)	- 163
Fc(NS)+Ph2(NS)	- 148	Fc(NS)+Ph2(NS)	- 99
Fc(NS)+succ(NS)	- 182	Fc(NS)+succ(NS)	- 150
Ph1(NS)+succ(NS)	- 142	Ph1(NS)+succ(NS)	- 179
Fc(WS)+Ph1(NS)	- 113	Fc(WS)+Ph1(NS)	- 109
Fc(WS)+Ph2(NS)	- 126	Fc(WS)+Ph2(NS)	- 90
Fc(WS)+succ(NS)	- 114	Fc(WS)+succ(NS)	- 99
Ph1(WS)+succ(NS)	- 246²	Ph1(WS)+succ(NS)	- 120
Fc(NS)+Ph1(WS)	- 150	Fc(NS)+Ph1(WS)	- 203
Fc(NS)+Ph2(WS)	- 162	Fc(NS)+Ph2(WS)	- 173
Fc(NS)+succ(WS)	- 184	Fc(NS)+succ(WS)	- 78
Ph1(NS)+succ(WS)	- 178	Ph1(NS)+succ(WS)	- 77

3-Me-βCD	ΔE	2,6-diMe-βCD	ΔE
Fc(WS)+Ph1(WS)	- 131	Fc(WS)+Ph1(WS)	- 100
Fc(WS)+Ph2(WS)	- 140	Fc(WS)+Ph2(WS)	- 97
Fc(WS)+succ(WS)	- 111	Fc(WS)+succ(WS)	- 77
Ph1(WS)+succ(WS)	- 96	Ph1(WS)+succ(WS)	- 111
Fc(NS)+Ph1(NS)	- 150	Fc(NS)+Ph1(NS)	- 112
Fc(NS)+Ph2(NS)	- 187	Fc(NS)+Ph2(NS)	- 82
Fc(NS)+succ(NS)	- 89	Fc(NS)+succ(NS)	- 79
Ph1(NS)+succ(NS)	- 145	Ph1(NS)+succ(NS)	- 89
Fc(WS)+Ph1(NS)	- 122	Fc(WS)+Ph1(NS)	- 87
Fc(WS)+Ph2(NS)	- 110	Fc(WS)+Ph2(NS)	- 70
Fc(WS)+succ(NS)	- 124	Fc(WS)+succ(NS)	- 84
Ph1(WS)+succ(NS)	- 132	Ph1(WS)+succ(NS)	- 87
Fc(NS)+Ph1(WS)	- 133	Fc(NS)+Ph1(WS)	- 114
Fc(NS)+Ph2(WS)	- 208	Fc(NS)+Ph2(WS)	- 118
Fc(NS)+succ(WS)	- 278	Fc(NS)+succ(WS)	- 106
Ph1(NS)+succ(WS)	- 76	Ph1(NS)+succ(WS)	- 112

6-Me-βCD	ΔE	3,6-diMe-βCD	ΔE
----------	----	--------------	----

Fc(WS)+Ph1(WS)	- 141	Fc(WS)+Ph1(WS)	- 118
Fc(WS)+Ph2(WS)	- 119	Fc(WS)+Ph2(WS)	- 135
Fc(WS)+succ(WS)	- 116	Fc(WS)+succ(WS)	- 118
Ph1(WS)+succ(WS)	- 129	Ph1(WS)+succ(WS)	- 104
Fc(NS)+Ph1(NS)	- 130	Fc(NS)+Ph1(NS)	- 129
Fc(NS)+Ph2(NS)	- 143	Fc(NS)+Ph2(NS)	- 94
Fc(NS)+succ(NS)	- 120	Fc(NS)+succ(NS)	- 91
Ph1(NS)+succ(NS)	- 103	Ph1(NS)+succ(NS)	- 94
Fc(WS)+Ph1(NS)	- 131	Fc(WS)+Ph1(NS)	- 78
Fc(WS)+Ph2(NS)	- 87	Fc(WS)+Ph2(NS)	- 108
Fc(WS)+succ(NS)	- 93	Fc(WS)+succ(NS)	- 101
Ph1(WS)+succ(NS)	- 123	Ph1(WS)+succ(NS)	- 73
Fc(NS)+Ph1(WS)	- 143	Fc(NS)+Ph1(WS)	- 108
Fc(NS)+Ph2(WS)	- 170	Fc(NS)+Ph2(WS)	- 123
Fc(NS)+succ(WS)	- 106	Fc(NS)+succ(WS)	- 98
Ph1(NS)+succ(WS)	- 119	Ph1(NS)+succ(WS)	- 87

¹ Fc: ferrocene, Ph1: phenol 1 (*cis* to Fc), Ph2: phenol 2 (*trans* to Fc), Succ: succinimidylpropyl. Calculated combinations are WS-WS, NS-NS, WS-NS or NS-WS, but excluding all Ph1-Ph2 and Ph2-Succ combinations.² The best system and value for each series is displayed in bold and is the starting systems (G0) for calculation for diversely methylated cyclodextrins calculations.

Classification trees for models of SuccFerr- β CD assemblies

General background:

The information related to our SuccFerr-cyclodextrin complex models generate a large amount of data that needs to be saved reliably, but also to be readily consulted and even used in statistics or other calculations. The ideal method is therefore to save the data in a database which makes it possible to retrieve the desired information from thousands of other records by sorting and selecting using simple queries. The free pack XAMPP (version 8.2.0 was used) can be used to manage this very large scientific database. It contains a web server program (Apache), a database (MariaDB) supporting the SQL query language, a web system for manually entering SQL queries (phpMyAdmin) and a programming language interpreter (PHP). Access to the server through a web browser (whichever one is used, and whatever the computer's operating system) allows

information to be entered and displayed in HTML format, but also to make decisions automatically (PHP program). In addition to this database, all models are available in XYZ format satisfying the Findability, Accessibility, Interoperability and Reusability (FAIR) approach and additional treatments and statistics on them were carried out using a handmade C program. Together with other recognized technologies, such as ioChem-BD, AiiDa, ASE or Flask, these new packages can help in the exploitation and management of large amounts of computational data. More details follow.

Method:

In this work, we show how our web application was used to classify a series of cyclodextrin models (here limited to β CDs) according to their methylation state. These cyclodextrins serve to solubilize SuccFerr (Figure 2 of main text), which is very active against cancer cells but poorly soluble in water.

We have used the semiempirical PM3 quantum-mechanical molecular modelling method (abbreviated as “PM3”) to determine which methylated β cyclodextrins are the ablest to make a stable supramolecular assembly in order to solubilize SuccFerr. We calculated the models of SuccFerr with one or two cyclodextrins (1-CD or 2-CD systems), using different types of β CD. Assembly with 3 CDs was ruled out early on as models showed poor inclusion of “moieties” in the CD cavity, and instead the formation of aggregate-like assemblies. Considering the central double bond of SuccFerr, we can envisage four molecular fragments, designated here as “moieties” (shown in Figure 2 of main text, surrounded by purple rectangles) likely to enter the CD cavity: the ferrocenyl group (Fc), phenol 1 (Ph1, *cis* to the Fc), phenol 2 (Ph2, *trans* to the Fc) and the succinimidylpropyl group (Succ). Each moiety can access the cavity of the CD either from the wide side or from the narrow side. For 1 CD systems, this gives a total of 8 possible combinations, which we label as 1-CD series 1 to 8. For 2-CD systems, there are 24 possible series (6 combinations of 2 moieties times 4 combinations of wide and narrow sides). When one of these four moieties of SuccFerr is encapsulated within a CD, all 21 oxygen atoms become non-equivalent. We have therefore numbered (O1 to O21) each of the oxygen atoms that can be (de)methylated in order to identify them unambiguously not only in our models, but also in our database (Figure 1 of main text). A specific structure can be found in this database using a complex SQL query with the appropriate index for each oxygen atom (value of each O_i is even numbers for unmethylated oxygen atoms and odd numbers for methylated ones). Based on the glucose numbering system, oxygens at position 2 are given a number $3n + 1$, those at the positions 3 are

listed as $3n + 2$, and those at positions 6 are labelled as $3n + 3$, with n ($0 \leq n \leq 6$) being the anhydroglucose unit number. For simplicity, this number is not referenced in this article, and the first unit ($n = 0$) was chosen arbitrarily for each series. The SQL query to search for all supramolecular assemblies involving a specific CD, referred as “free CD”, is simpler and consists of finding all records containing a certain number. This number is a binary conversion of the CD configuration (unmethylated = 0, methylated = 1), retaining the smaller of the seven possible numbers from now on “binarymin” (depending on the starting anhydroglucose unit number, see details in SI: “Algorithm to calculate binarymin” and Figure S3). The number of possible systems for each of these 8 series becomes $2^{21} = 2,097,152$ combinations (2^{42} for 2-CD systems); evidently, these are impossible to calculate with limited resources in a reasonable time. Thus, we have devised a simple, robust, and sound algorithm to reduce the number of combinations to be computed (see “undefined-CD” section). This algorithm is based on the stability of the CD-SuccFerr complex. The computed stability requires calculation of the energies of the SuccFerr-CD(s) molecular assembly, of each CD, and of SuccFerr, the latter two in the conformations they had in the molecular assemblies to give the ΔE (energy variations, equation 2 or 3) of the reactions: SuccFerr + CD₁ [+ CD₂] → molecular assembly. Note that equation 3 also takes into account the eventual interactions between the two CDs, which makes it possible to predict a more negative ΔE than that of 1-CD systems.

$$\text{Eq. for 1-CD: } \Delta E = E_{(\text{SuccFerr}+\text{CD})} - E_{\text{SuccFerr}} - E_{\text{CD1}}$$

$$\text{Eq. for 2-CD: } \Delta E = E_{(\text{SuccFerr}+\text{CD1}+\text{CD2})} - E_{\text{SuccFerr}} - E_{\text{CD1}} - E_{\text{CD2}}$$

We started with well-defined CDs (see definition above) as a basic model, in order not only to compare them with undefined CDs but also to create the latter by modifying the states of methylation.

- Well-defined CDs; Having no idea which types of methylation would work best for each combination, the 8 1-CD series were each comprised of 6 well-defined CDs: three mono-methylated combinations: 2-Me- β CD, 3-Me- β CD, and 6-Me- β CD (i.e., on all the oxygen atoms at positions 2, 3 or 6, respectively, of their 7 anhydroglucose units, i.e., all glucose units are identical), and three dimethylated combinations: 2,3-diMe- β CD, 2,6-diMe- β CD (DM β CD) and 3,6-diMe- β CD (Table S1). The same calculations were made with the 2-CD assemblies (reduced to 16 of the 24 possible, eliminating models with bad inclusion characteristics, Table S2).

- Undefined CDs; The well-defined CDs were then used as the basic model on which to create the undefined CDs, that is, those with any combination of methylations. First, so as to start from already relatively stable models, the best one from each of the 8 series of the previous 1-CD models (i.e., having the most negative ΔE) was chosen (Table S1). For the 2-CD models, only the 4 best series (2-CD series 1 to 4, taken from Table S2) were studied because calculation times would be too long. When creating methylation modification trees (resembling a tree of DNA / RNA / protein mutations but without the deletions and insertions), these 12 basic systems were classified as generation zero (G0). By analogy, we will use the term "mutation" for each change in the methylation pattern, even if this nomenclature is perhaps non-standard. Initially, we are only allowed to continue mutations into the next generation (Gn + 2) if ΔE has decreased between Gn and Gn + 1 (rule 1). The series 2-CD S4 was exempted of this rule 1 to analyze its effect and utility (the remaining very best model without descendant of the whole series was systematically used to continue in the next generation, even if its ΔE has increased). Of course, it is only interesting when this model has a ΔE more negative than that of any model with a decreasing ΔE .

Rule 1: (applied to any series but 2-CD S4)

If($\Delta E(G_{n+1}) < \Delta E(G_n)$) then

Allow to continue with descendants Gn+2 within this branch (creation of new ramifications, one for all descendants)

else

stop (creation of new ramifications Gn + 2 are not allowed starting from this experiment of generation Gn + 1)

For the 8 series 1-CD and 4 series 2-CD, 21 descendants of G0 (42 for 2-CD systems) were created, each corresponding to a reversal of the methylation of the G0 system onto a different oxygen atom, creating the first generation of mutations (G1). The best blend (most negative ΔE) of the 21/42 G1 offspring in each series was in turn used as a template for the creation of a new mutation (G2). At the start of the branch, the first mutations (G1, G2, G3, etc.) were retained: rule 2: no modification of the methylation of an oxygen atom several times in the same chain / route of modifications. The series 2-CD S4 was also exempted of this rule 2 for the same purpose of studying the effect of this rule.

Rule 2: (applied to any series but 2-CD S4)

nbOH = 21 (1-CD models) or 42 (2-CD models)

for ($i = 1; i \leq \text{nbOH}; i++$) \leftarrow outer loop going through all oxygen atoms of current system, of generation G_n , to be modified

for ($j = 1; j < G_n; j++$) \leftarrow inner loop going through all previous generations G_j of this route, starting from G_1

if(O_i of ancestor G_j was modified) then

mark O_i as non-modifiable (will be ignored)

if(O_i is not marked) then

allow inversion of methylation

The goal of fixing these mutations is a logical choice: If they have had a positive effect, it is advisable not to cancel them in the following mutations, or in any case not soon after. However, when many methylations have been altered, the complex interaction can cause a mutation that was useful early on in advancing the stability of the SuccFerr-CD assembly, only to become troublesome later on. This has been verified several times in this study. In this case it is necessary, at a certain stage, mainly at the end of the road, to authorize this mutation at will (bypass of rule 2). In this study, the "end of the road" (dead end) will be how we describe a chain of modifications leading to a generation where none of its systems have seen their ΔE drop, which leads to a dead end (locally, a model not allowed by rule 1 is also a type of dead end). Bypass of rule 2 is applied to a certain parent experiment chosen by the user and concerns only its direct descendants $G_n + 1$, without changing the mode of operation of the rest of the tree. It brings up, for this parent experiment, other descendants to be calculated, which can unblock the road for a few more generations, as has been verified in this work. However, when even the bypass of rule 2 cannot prevent the end of the route, it may also be interesting to remove rule 1 (bypass of rule 1), although this only seems to make sense for better systems whatever branch they are in, and not only the current one. The exception is 2-CD S4 where rules 1 and 2 were not applied from the start, as stated above, for comparison of the two methods (with or without rules). Indeed, with a strict application of rule 1, a good system, but going through a bad stage, would never be reached, or by making a big detour. For example, the very best system # 3029 (unique value of its key 'ID') belonging to series 4 (the only series where rule 2 is systematically bypassed) of 2-CD systems, has undergone a chain of mutations starting from the G_0 system: $G_0 - O_{37} + O_2 - O_{31} - O_{40} + O_{18} -$

$O_4 - O_2 + O_{36} - O_{16} + O_{48} + O_{44} - O_{43} - O_{36} - O_{44} + O_{42} + O_{16} - O_7 + O_{44} - O_{42} + O_{36} - O_{48} + O_2 - O_1 + O_7 + O_{48} - O_{19}$ (plus = methylation, minus = demethylation, of the indicated oxygen atom). The demethylation of O_{48} , carried out in G25 (ID = 2029), produced a rise of ΔE , and without the bypass of rule 1, this chain, which continues until G27 (end of the route), would have been blocked in G25. Likewise, we see a demethylation of O_{44} carried out in G14 (ID = 2630), while this atom had already been modified in G11 (ID = 2518), and that without bypassing rule 2, the chain would have stopped in G14. Moreover, we also see that this atom is modified a third time in G18, just like O_{48} (G10, G21 et G25) and O_{36} (G8, G13 et G20). This latest information confirms what was previously described, namely that a change, and its undoing a few generations later, can both lead to an improvement in the system.

Always, in order to find more good systems, it is unavoidable to create bifurcations (creation of a tree of mutations) since a single road will necessarily ends, even with bypasses. Going back to generations before the end of the road and creating a fork from the second / third best system respecting rule 1 creates parallel chains and therefore the branches of this tree of mutations. It should be noted that this does not apply to 2-CD S4, where the current very best system is chosen, respecting rules 1, rule 2 or none, being inside the current road or inside a parallel one. Some of these parallel chains can lead to the same configuration (same mutations but in a different order), which would create a duplicate in the tree which would no longer be a spanning tree. To avoid this, you need a system that checks, before creating an experiment, that there is no risk of producing a duplicate by checking whether it already exists in the database (rule 3: prohibit duplicates). This rule 3 is thus systematically applied in each road to avoid doing a mutation in $G_n + 2$ that would lead to the same model G_n , by cancelling the mutation done in $G_n + 1$.

Rule 3: (no bypass allowed)

for ($i = 1$; $i \leq$ All structures this series; $i++$)

 if(Structure_{*i*} already exists), then

 Structure_{*i*} is displayed for information

 else

 Structure_{*i*} is allowed to be created in database

To summarize, to determine if rules 1 and 2 are really useful, we applied two methods and compared their effect:

- Method I; For all 1-CD series and for the first three 2-CD series, we apply rules 1 and 2 at the beginning of the trees and decide to lift these rules (bypass) when the progression becomes slower (the number of dead ends becomes significant). This method therefore requires the user to decide on when to apply these bypasses. We must then decide to bypass rule 2, but also decide if we do it with, or without, simultaneously bypassing rule 1. This method is therefore less clear;
- Method II; For the 2-CD S4 series, we systematically bypass rules 1 and 2 from the start (G0). The user therefore follows the algorithm without having to decide. Finally, to produce this 2-CD S4 series, a final adjustment was made: To simplify, for the 2-CD S1 to S3 series, we took the same CD for both insertions. But there was no reason for this CD to be the best one for both moieties at the same time, and therefore for us to start from the model with the lowest possible ΔE (using the best model of Table S2, reported in Table 1). Therefore, we made the 6 x 6 possible combinations, with the help of a new PHP program (CDxCD.php, provided) and a new database table (CDxCD, provided) and chose the best model as G0 of our tree for S4. Ironically, a model with twice the same CD (2-Me-CD) gave the best result!

Of course, rule 3 remains mandatory for both methods.

Figure S1 shows examples on a fictitious tree and Figure S2 compares the computer flowchart of methods I and II:

1. Rule 1: The mutations indicated all correspond to decreases in ΔE except the $G0 + O_4 + O_6$ route where methylation on oxygen atom 6 causes an increase in ΔE and will not be allowed to continue in G3 (bypass possible if the user so desires, and preferably only when reaching the ends of the road leading to the best systems or if this model is the current very best for 2-CD S4);
2. Rule 2: The mutation ($G0 + O_4 - O_8 + O_5 - O_4$) modifies oxygen atom 4 a second time and will not be authorized (but bypass possible, especially useful at the end of the road). For 2-CD S4, bypass is systematic;
3. Rule 3: The $G0 + O_4 - O_8 + O_5 - O_9 - O_3$ mutation gives the same configuration as the $G0 + O_4 - O_8 - O_3 + O_5 - O_9$ route, so the -3 mutation will be blocked, and this route will stop on demethylation in 9 (bypass not authorized whatever is the series).

The simplified computer flowchart common to the two PHP programs which manages the trees of the 1-CD and 2-CD system is shown in Figure S4.

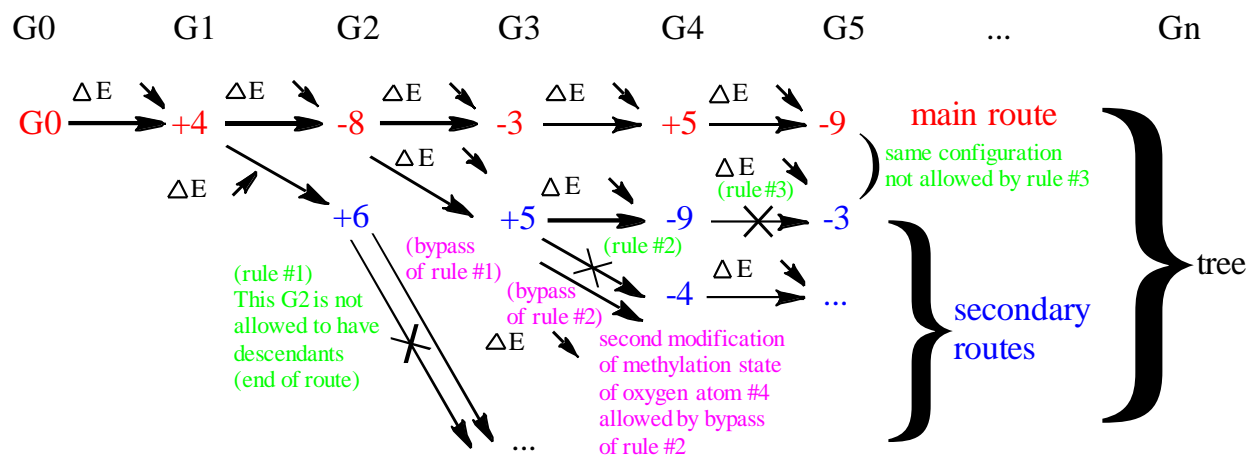


Figure S1. Example of a fictitious mutation tree with explanation of the rules. + indicates methylation on the oxygen atom of the number indicated and - demethylation. The main route/road is the one that goes through the best system of each generation and the secondary roads those that go through at least one system that is not the best of its generation. G0 (generation 0) represents the well-defined starting CD and G1, G2,... the following generations.

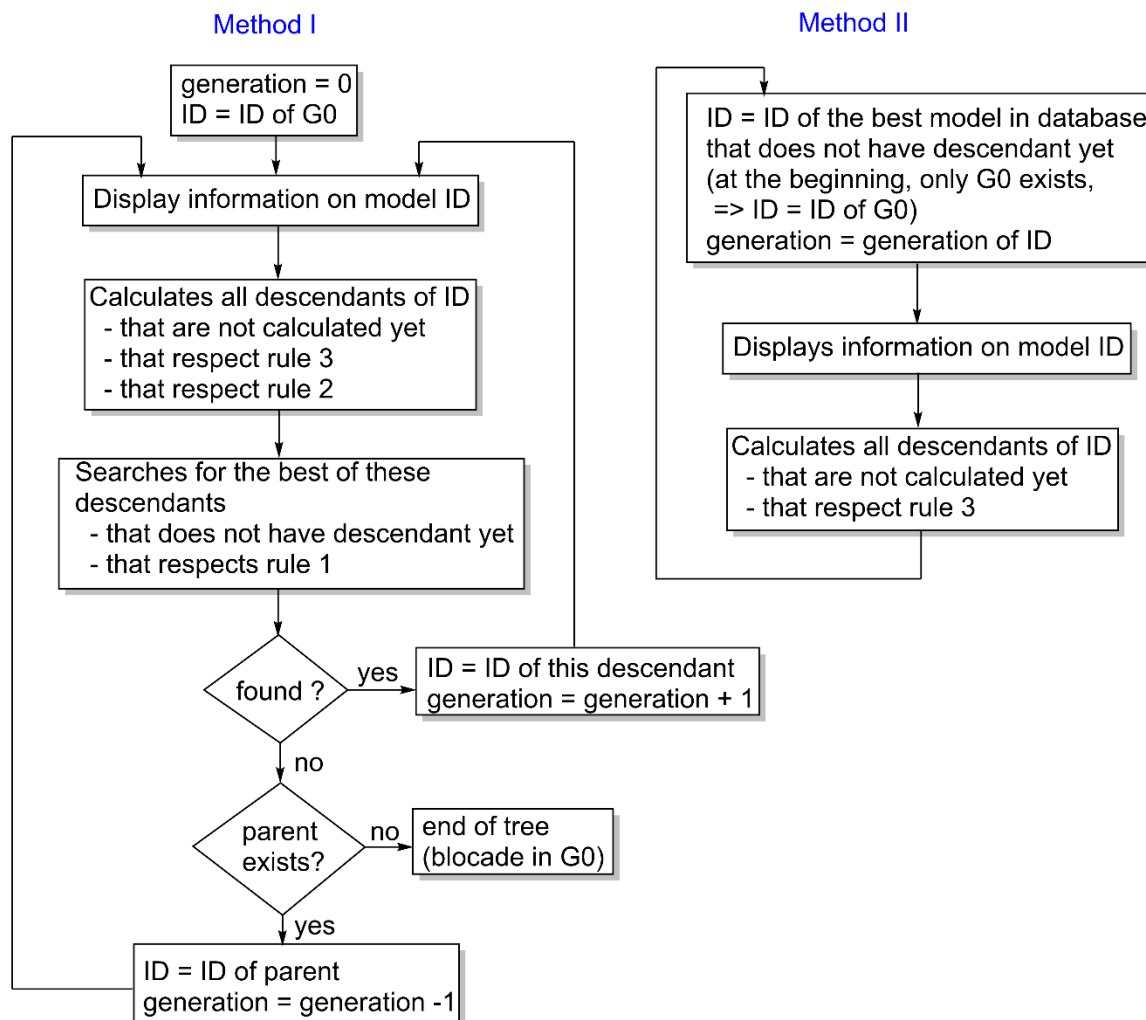


Figure S2. computer flowchart describing method I (left: rules 1 and 2 are respected) and method II (right: rule 1 and 2 are waived).

Comparing methods I and II

The study of the paths of mutations ('path' fields of the database) shows that with the bypass of rule 2, certain mutations are successfully cancelled sometimes only three generations later (for example, ID 7692 and 7759 for O16). We can therefore ask ourselves the question of the validity of rule 2, especially since its bypass is chosen more or less arbitrarily by the user, which introduces serendipity into the calculations and therefore confusion into the case where we want to apply this method for other systems. Similarly, rule 1 seems counter-productive. Indeed, when we arrive at an end of route in G_n , the method I that we used consists in downgrading by taking the second-best model respecting rule 1 of the previous generation (in G_{n-1} , if this model exists otherwise in G_{n-2} , ... see computer flowchart Figure S2 method I). This prolongs a parallel branch (a bifurcation with just one model up to now) in the tree, whereas a model not respecting this rule, in the current generation (G_n) may have a better affinity and not lead to the creation of a long new branch but to the extension of the current branch (tree less ramified and therefore less confusing). Method II not applying these two rules 1 and 2 with the 2-CD S4 series gives good results, but with a larger number of calculations (Figure S2 method II). This new simplified method II is therefore to be preferred over the old method I, even with more calculations (that also could lead to better models by serendipity).

Web application

To help handle the considerable number of experiments (9,767 for 1-CD calculations and 3,458 for 2-CD calculations) and corresponding data created by the ten series of experiments for 1-CD calculations and four series for 2-CD calculations, and to verify and respect the rules explained in section "Undefined CD" in the main text, an application was specially created with the XAMPP pack (PHP, MySQL/MariaDB and phpMyAdmin) (<https://www.apachefriends.org>). The pack should be installed (Windows, Linux, or OS X) to run the application and operate the database (see below).

Alternatively, to consult the results that are displayed into the dynamic webpages by the PHP program, static webpages, one for each model, were created by the C program (see

below). This C program permitted to avoid to manually save from the browser all pages one by one. The links operate off-line without XAMPP (they were adapted), but not the forms.

Description of the system:

More details can be found at <https://hal.science/hal-03991394> (then follow the link towards Software Heritage or directly use the link displayed in page 1 of this document).

- The data were stored into a database named 'pascal' operated by XAMPP (MariaDB, database engine: MyISAM, one table (calculscd) for the eight series of 1-CD calculations (+ 2 extra series S9 and S10, see below), one table (calculs2cd) for the four series of 2-CD calculations) and one table (cdxcd) for the special experiment to find the best model used as G0 for series 2-CD S4 (see above).
- Starting G0 systems for each series were either manually entered the table of the database using the phpMyAdmin software or simply created by a special option in the software by entering the methylation state of position 2, 3 and 6 for the seven anhydroglucose units. The other experiments, derived from them, were stored by the same software.
- The software was coded in PHP and was stored into three files (cdmodele.php to handle 1-CD calculations, cdmodele2cd.php for 2-CD calculations and CDxCD.php for the special experiment to find the best G0 for 2-CD S4, see below). It allows one to:
 - create a dynamic web page displaying the forms and the data of the trees into a browser (location: <http://localhost/cdmodele.php> for 1-CD calculations, <http://localhost/cdmodele2cd.php> for 2-CD calculations and <http://localhost/CDxCD.php?action=show> for special experiment). If the server (where the XAMPP pack is installed) and the client (where the browser operates) are not the same computer, "localhost" should be replaced by the IP address of the server,
 - receive the data sent from the forms (energies of the supramolecular assembly and of each of its components),
 - calculate ΔE according to the previous formula inside the main text of the article (eq. 2 for cdmodele.php, eq. 3 for cdmodele2cd.php and CDxCD.php),
 - store them into the table,

- allow continuation of experiments in the next generation for ameliorated systems only (rule 1) or force calculation of some next generations (bypass of rule 1 by direct access to them by typing their parent ID number in a form in the webpage),
 - create new allowed pending calculations into the database and keep track of them,
 - only allow calculations of systems not already treated by another branch (rule 3) and, if rule 2 activated (deactivated by clicking on a link), not corresponding to an already modified oxygen atom in the path from G0 to the present system,
 - calculate the statistics and automatically display them when displaying the G0 level of the tree of the current series (see static webpages 1-CD S1 to 1-CD S8 and 2-CD S1 to 2-CD S4 (.html) in compressed archive), display statistics for the eight series of 1-CD calculations into a table for comparison, ...
- The web page, used as an interface, allows one to display the forms:
- buttons for the creation of new pending experiments, if allowed, and direct access to a specific experiment for an eventual bypass of rule 1,
 - fields to input (copy/past from Spartan) calculated energies,
 - hypertext links (navigation forward and backward inside the trees, bypass of rule 2 or change of current series), ...

Database and SQL:

The structure of the database (tables `calculscd`, `calculs2cd` and `cdxcd`) is stored into the SQL file 'pascal.sql'. This file, that contains SQL queries to create the database with its data, should be used to import the database into the MariaDB system using the 'Import' option of phpMyAdmin (or other database management systems). This SQL file is the one that should preferably be used to recreate our database identically, because it includes the constraints on the keys, thanks to its system-independent SQL commands and is therefore more portable.

For 1-CD calculations, the unique table for the 10 series inside the database was created with this SQL command:

```
CREATE TABLE `calculscd` (  
  `ID` int(16) UNSIGNED NOT NULL,
```

```

`O1` tinyint(4) NOT NULL,
`O2` tinyint(4) NOT NULL,
`O3` tinyint(4) NOT NULL,
`O4` tinyint(4) NOT NULL,
`O5` tinyint(4) NOT NULL,
`O6` tinyint(4) NOT NULL,
`O7` tinyint(4) NOT NULL,
`O8` tinyint(4) NOT NULL,
`O9` tinyint(4) NOT NULL,
`O10` tinyint(4) NOT NULL,
`O11` tinyint(4) NOT NULL,
`O12` tinyint(4) NOT NULL,
`O13` tinyint(4) NOT NULL,
`O14` tinyint(4) NOT NULL,
`O15` tinyint(4) NOT NULL,
`O16` tinyint(4) NOT NULL,
`O17` tinyint(4) NOT NULL,
`O18` tinyint(4) NOT NULL,
`O19` tinyint(4) NOT NULL,
`O20` tinyint(4) NOT NULL,
`O21` tinyint(4) NOT NULL,
`changed` tinyint(4) NOT NULL,
`path` text NOT NULL,
`IDparent` int(16) UNSIGNED NOT NULL,
`deltaE` float NOT NULL,
`stop` tinyint(2) NOT NULL,
`series` smallint(6) NOT NULL,
`binarymin` int(16) UNSIGNED NOT NULL,
`generation` tinyint(2) UNSIGNED NOT NULL,
`deltadelta` float NOT NULL,
`mainseries` smallint(6) NOT NULL
) ENGINE=MyISAM DEFAULT CHARSET=latin1;

```

For 2-CD calculations, the unique table (calculs2cd) was created with this SQL command:

```
CREATE TABLE `calculs2cd` (  
  `ID` int(16) UNSIGNED NOT NULL,  
  `O1` tinyint(4) NOT NULL,  
  `O2` tinyint(4) NOT NULL,  
  `O3` tinyint(4) NOT NULL,  
  `O4` tinyint(4) NOT NULL,  
  `O5` tinyint(4) NOT NULL,  
  `O6` tinyint(4) NOT NULL,  
  `O7` tinyint(4) NOT NULL,  
  `O8` tinyint(4) NOT NULL,  
  `O9` tinyint(4) NOT NULL,  
  `O10` tinyint(4) NOT NULL,  
  `O11` tinyint(4) NOT NULL,  
  `O12` tinyint(4) NOT NULL,  
  `O13` tinyint(4) NOT NULL,  
  `O14` tinyint(4) NOT NULL,  
  `O15` tinyint(4) NOT NULL,  
  `O16` tinyint(4) NOT NULL,  
  `O17` tinyint(4) NOT NULL,  
  `O18` tinyint(4) NOT NULL,  
  `O19` tinyint(4) NOT NULL,  
  `O20` tinyint(4) NOT NULL,  
  `O21` tinyint(4) NOT NULL,  
  `O31` tinyint(4) NOT NULL,  
  `O32` tinyint(4) NOT NULL,  
  `O33` tinyint(4) NOT NULL,  
  `O34` tinyint(4) NOT NULL,  
  `O35` tinyint(4) NOT NULL,  
  `O36` tinyint(4) NOT NULL,  
  `O37` tinyint(4) NOT NULL,
```

```

`O38` tinyint(4) NOT NULL,
`O39` tinyint(4) NOT NULL,
`O40` tinyint(4) NOT NULL,
`O41` tinyint(4) NOT NULL,
`O42` tinyint(4) NOT NULL,
`O43` tinyint(4) NOT NULL,
`O44` tinyint(4) NOT NULL,
`O45` tinyint(4) NOT NULL,
`O46` tinyint(4) NOT NULL,
`O47` tinyint(4) NOT NULL,
`O48` tinyint(4) NOT NULL,
`O49` tinyint(4) NOT NULL,
`O50` tinyint(4) NOT NULL,
`O51` tinyint(4) NOT NULL,
`changed` tinyint(4) NOT NULL,
`path` text NOT NULL,
`IDparent` int(16) UNSIGNED NOT NULL,
`deltaE` float NOT NULL,
`stop` tinyint(2) NOT NULL,
`series` smallint(6) NOT NULL,
`binarymin1` int(16) UNSIGNED NOT NULL,
`binarymin2` int(16) UNSIGNED NOT NULL,
`generation` tinyint(2) UNSIGNED NOT NULL,
`deltadelta` float NOT NULL,
`mainseries` smallint(6) NOT NULL
) ENGINE=MyISAM DEFAULT CHARSET=latin1;

```

For the database to respond fast enough to requests, keys/index should be set on fields ‘ID’ (it is mandatory to have unique values for this field, so ‘ID’ was auto incremented by the database engine at each insertion of experiment) and ‘IDparent’ (mandatory for a large set of systems, because the database is slow without it).

These two options are set by the SQL command:

```
ALTER TABLE `calculscd` ADD UNIQUE KEY `ID` (`ID`) USING BTREE, ADD KEY `IDparent` (`IDparent`);
```

For the 'ID' field to be set in auto_increment mode:

```
ALTER TABLE `calculscd` CHANGE `ID` `ID` INT(16) UNSIGNED NOT NULL AUTO_INCREMENT;
```

The version in the pascal.sql file is a little different from the initial one (creation of an empty table), since it considers the already filled (with 9767 models, the next one to be created being 9768) table:

```
ALTER TABLE `calculscd` MODIFY `ID` int(16) UNSIGNED NOT NULL AUTO_INCREMENT, AUTO_INCREMENT=9768;
```

For the calculs2cd table, the SQL commands are:

```
ALTER TABLE `calculs2cd` ADD UNIQUE KEY `ID` (`ID`) USING BTREE, ADD KEY `IDparent` (`IDparent`);
```

```
ALTER TABLE `calculs2cd` CHANGE `ID` `ID` INT(16) UNSIGNED NOT NULL AUTO_INCREMENT;
```

Here too, we can see that the table is already filled (with 3458 models, the next one to be created being 3459) table:

```
ALTER TABLE `calculs2cd` MODIFY `ID` int(16) UNSIGNED NOT NULL AUTO_INCREMENT, AUTO_INCREMENT=3459;
```

There is no need to create a multicolumn index, as unicity constraint for example. Indeed, the PHP software verifies the constraints (rule 3) before creating experiments. Moreover, it would be impossible, since the Oi fields can take value 0, 1, 2, 3, 6 or 7 (the unicity should be based on the bit 0 of these fields (0 = unmethylated (even number), 1 = methylated(odd number)), not on the value itself.

Examples of useful SQL requests:

Calculation/recovery of some statistics or values by SQL requests for a particular series of experiments for 1-CD systems (some can be used for 2-CD systems, replacing the table 'calculscd' by 'calculs2cd'): The PHP software can use these SQL requests, but they also can be manually

used using phpMyAdmin (in this last case, the variable '\$series' should be replaced by 1 to 10 depending on the series we are interested in):

- The best ΔE value of a series: ***SELECT MIN(deltaE) FROM calculscd WHERE series='\$series'***
- The total number of calculated systems of a series: ***SELECT COUNT(*) FROM calculscd WHERE series='\$series'***
- The total number of changes of methylation state on a certain oxygen atom n (replace \$n by: [1..21] for methylation, [-1..-21] for demethylation) for all the experiments of a series: ***SELECT count(*) FROM calculscd WHERE changed='\$n' AND series='\$series'***
- The same, but only changes that improved ΔE (respecting rule 1 and 2 only): ***SELECT count(*) FROM calculscd WHERE changed='\$n' AND series='\$series' AND (stop='0' OR stop='2')***
- The total number of free CDs of systems that have a ΔE better than a certain value (\$v, in kJ/mol): ***SELECT COUNT(DISTINCT binarymin) FROM calculscd WHERE deltaE < '\$v' AND series='\$series'***
- Number of untreated sub-trees: ***SELECT count(*) FROM calculscd WHERE series='\$series' AND stop='0'***
- Total number of treated sub-trees: ***SELECT count(*) FROM calculscd WHERE series='\$series' AND (stop='2' OR stop='3')***
- Recovery of the generation of a certain experiment '\$ID': ***SELECT generation FROM calculscd WHERE ID='\$ID' LIMIT 1***
- Recovery of its ΔE : ***SELECT deltaE FROM calculscd WHERE ID='\$ID' LIMIT 1***
- Recovery of the number of systems that are allowed to continue for the next generation for a certain generation Gn (respecting rule 1 and 2 only): ***SELECT count(*) FROM calculscd WHERE generation='\$Gn' AND series='\$series' AND (stop='0' OR stop='2')***
- Recovery of the number of systems of a certain generation Gn: ***SELECT count(*) FROM calculscd WHERE generation='\$Gn' AND series='\$series'***
- To find the average positive variation of ΔE ($\Delta\Delta E$): ***SELECT AVG(deltadelta) FROM calculscd WHERE series='\$series' AND deltax > 0***
- To find the average negative variation of ΔE ($\Delta\Delta E$): ***SELECT AVG(deltadelta) FROM calculscd WHERE series='\$series' AND deltax < 0***

- With phpMyAdmin, ordering the experiments of series '\$series' by decreasing stability (i.e. starting from the lowest ΔE): ***SELECT ID,deltaE,stop FROM `calculscd` WHERE series='\$series' ORDER BY deltaE.*** The change of the 'stop' field value (with phpMyAdmin, by double-clicking on it) to 3, unlocks the bypass of rule 2 for its possible (allowed by rule 3) descendants. Then, typing the value of the 'ID' field into a form in the web page generated by the PHP software permits one to directly access to this experiment (without having to navigate into the tree, an even if not allowed by rule 1) to calculate its descendants (if not treated yet, and even if not allowed by rule 2).
- To calculate the average ΔE of a certain series: ***SELECT AVG(deltaE) FROM calculscd WHERE series='\$series'***
- To calculate the best ΔE of 1-CD systems (used as the limit to determinate the 2-CD systems that are better than any 1-CD system): ***SELECT MIN(deltaE) FROM calculscd***
- Then, to calculate the number of 2-CD systems that are better than any 1-CD system: ***SELECT COUNT(*) FROM calculs2cd WHERE deltaE<(SELECT MIN(deltaE) FROM calculscd)***

Database table design

The tables, which are attached to a database must contain all the identifying values of an experiment (one row represents an experiment). The columns represent the state information, the values of certain calculations, but also the links between the experiments which derive from one another and form a tree, this one consisting of parent-descendant couples attached to each other. Table S3 is part of the database table "calculscd" which contains information for 9767 1-CD system experiments. The 8 (+2) series / trees are stored in this table, the "series" field allowing one to select only the experiments concerning the desired series (tree).

- The 'ID' field represents the unique experiment number (key), assigned by the database management system when creating a new experience row in the table (the 'AUTO_INCREMENT' attribute added to the table after its creation by an SQL command 'ALTER TABLE' allows this uniqueness, see above).
- The "Parent ID" field indicates the parent experiment (its "ID" field) from which the current experiment is derived by modifying the methylation of one of its 21 oxygen atoms. It allows navigation in the tree going back to the previous generation ($G_n \rightarrow G_{n-1}$) thanks to the PHP

program and the hypertext links displayed in the web page it creates. A zero value indicates that this is the initial CD G0 (no parent).

- Conversely, searching for the value of the 'ID' field of a parent experience in the 'Parent ID' field of all the rows in the table allows you to find its descendants (downward navigation $G_n \rightarrow G_{n+1}$).

- The modified oxygen atom in the offspring is indicated in the 'changed' field, a positive value indicating methylation, and a negative value indicating demethylation.

- This change is reflected in the state attributes 'O1' through 'O21' which represent the methylation state of the corresponding oxygen atom. The values taken are: 0 = initially unmethylated and unmodified, 1 = initially methylated and unmodified, 2 = changed once to demethylated, 3 = changed once to methylated, 6 = unmethylated and 7 = methylated. Values 6 and 7 represent the last change when the possibility to change several times the methylation state of certain oxygen atoms was allowed and done by bypass of rule 2). For this reason, a unicity constraint in the table (unicity index set on O1... O21 + series) is useless, because inoperative to control respect of rule 3 (done by the PHP program, not by the database engine).

- The 'deltaE' attribute represents ΔE and 'deltadelta' its variation between itself and its parent ($\Delta\Delta E$). A calculated positive value indicates destabilization which sets the "stop" attribute to 1, indicating to the PHP program that the tree should not continue in this route / branch (rule 1). A negative value indicates a system improvement and "stop" is set to 0 to allow the PHP program to display a hyperlink in the web page to be able to calculate the descendants of the descendant. When at least one descendant has been created in the database, "stop" changes to 2 to indicate that this system is a parent. A value then changed to 3 for "stop" indicates that although we should not calculate some of its descendants (mutation already made in this route), we still allow it (bypass of rule 2).

- The 'binarymin' attribute corresponds to a very simple representation of a CD, a binary representation (but seen as a number in base 10, see algorithm below): the methylation of an oxygen atom corresponds to a value of 1 and a non-methylation (hydroxyl group) to 0. All these values represent the bits of the binary number. As a β CD is composed of 7 anhydroglucose units connected in a cycle, depending on the unit from which we start to count (but always from position 2 (glucose numbering system) of each unit and in the direction $2 \rightarrow 3 \rightarrow 6 \rightarrow 2 \rightarrow 3 \rightarrow 6 \rightarrow \dots$) we can calculate 7 binary numbers. The one selected is the smallest. Two experiments that have the same 'binarymin' attribute have in common the same CD that is placed differently around SuccFerr,

which makes it easier to search by SQL query than using the 21 fields 'O1' through 'O21'. A description of binarymin and how to calculate this number (algorithm and Figure S3) can be found below.

- 'mainseries' indicates the type of insertion (moieties of SuccFerr involved + insertion side of the CD) as 'series', but 'mainseries' permits any starting CD as G0 (i.e., permits to create several series 'series' with the same insertion type but with a different starting CD), where the starting CD was fixed with 'series' and is unique. Originally, S8, S9 and S10 were sub-series of 'mainseries'=8 (but disassociated because of denaturation of the model during modelling). Now 'mainseries'='series' for all the models (see below).

- The last attribute, not shown in Table S3 for clarity, is "path". It is a string of characters which indicates, in an understandable way for a person, the chain of mutations that has taken place since G0. For example, for experiment (ID) # 7011, 'path' = " 1CD succinimide into wide side of 3-6-diMe-CD: - O ₂₀ + O ₁", we see what the G0 system corresponds to and that it has then undergone two mutations: demethylation to O₂₀ (sign -) followed by methylation to O₁ (sign +). This representation makes it possible to see immediately the route followed (without having to reconstruct it with each "changed" field encountered starting from G0). It is displayed in the web page for each experiment, so that the formatting HTML tags are interpreted, and that "- O ₂₀ + O ₁" is displayed simply and cleanly "- O₂₀ + O₁".

The table 'calculs2cd' (2-CD systems) follows the same model but 'binarymin' is replaced by 'binarymin1' and 'binarymin2' (for each of the two CDs) and fields O31 to O51 are added to represent the 21 atoms of oxygen from the second CD. The numbering starts at 31 because it suffices to subtract 30 to find the numbering of the first CD and to guess the numbering position of the glucose.

Table S3. Partial representation (a few rows selected) of the 'calculscd' table (1-CD systems) concerning experiments in series S8. Values in kJ/mol for deltaE (ΔE of model ID) and deltadelta ($\Delta\Delta E$: ΔE of model ID - ΔE of parent model IDparent). Other fields (integers) do not have units.

ID	O1	O2	...	O19	O20	O21	changed	IDparent	deltaE	stop	series	binarymin	generation	deltadelta
6990	0	1	...	0	1	1	0	0	-110.51	2	8	1797558	0	0
...
7010	0	1	...	0	2	1	-20	6990	-249.138	2	8	1273270	1	-138.628
...
7029	0	1	...	0	2	1	-18	7010	-266.011	2	8	748980	2	-16.873
...
7039	0	1	...	0	2	1	-5	7029	-279.769	2	8	748852	3	-13.758
8882	0	1	...	0	2	1	-6	7029	-259.584	1	8	738742	3	6.427
...

For clarity, the fields 'O3' to 'O18' as well as the field 'path' are not shown. The beginning of the tree is visible: experiment 6990 corresponds to the starting wdM β CD (G0), experiments 7010 (G1), 7029 (G2), 7039 (G3) are the descendants of the main branch (G0 \rightarrow -20 \rightarrow -18 \rightarrow -5 \rightarrow ...) which continues in G4 (stop = 2). Experiment 8882 is the end (stop = 1) of a secondary route because ΔE (deltaE) has increased ($\Delta\Delta E$ (deltadelta) > 0).

The PHP program and the webpages

The role of the program written in PHP is to generate the web page describing part of the tree (a G_n generation experiment and all of its G_n + 1 descendants) by fetching the information to display in the database. It also manages user interactions with the page (click on a hypertext link or validate forms), calculates ΔE and updates the information in the database. SQL queries are created by the program from the data it owns and are represented as strings that are sent to the database manager by the PHP function "mysqli_query()". For the queries, the database sends back to the program a response in the form of a table which must then be traversed line by line (if several lines) by extracting the attributes (one or more columns) requested by the "SELECT" clause. For modifications to the table, the SQL commands "INSERT" (insertion of a new experiment/row) and "UPDATE" (update of one or more columns of an existing experiment/row) save the program data to the database.

The program is responsible for displaying in the web page the characteristics of an experiment (identification number ID and generation G_n) chosen by the user and, in a table, all its authorized

direct descendants. These are the $G_n + 1$ respecting rule 2 or authorized by bypass of rule 2, or already created (in the case of the bypass of rule 1, which is then no longer controlled for this descendant), otherwise they are not displayed. The program displays their status and possible action for them. In the case of a descendant authorized but not created, the program displays a form button to create it (will trigger an SQL "INSERT" command in the database with automatic attribution of a unique "ID" experiment number serving key). Then you have to use the Modelling software to build the model according to the instructions on the web page (parent serving as the starting model and its oxygen atom to be modified). The energies of formation of the supramolecular assembly and its constituents are then calculated, then the web form displayed for the experiments awaiting results is completed. The program receives the information from the form sent, calculates ΔE for this descendant $G_n + 1$ and updates the fields 'deltaE' (ΔE), 'deltadelta' ($\Delta\Delta E$: variation of ΔE) and 'stop' (authorization to continue or not). The page then displays the calculated ΔE and the possible actions on this descendant according to its 'stop' value:

- 'stop' = 0: display a hypertext link (because authorized by rule 1) to display the detail of this $G_n + 1$ descendant in a web page (which replaces the current web page), which also allows one to display and calculate all its descendants of generation $G_n + 2$ (those allowed by rule 2).
- 'stop' = 1: indicates that ΔE has increased for this $G_n + 1$ experiment so rule 1 prohibits calculating the $G_n + 2$ descendants of this $G_n + 1$ experiment. No hypertext link is displayed to the web page describing the details of the $G_n + 1$ experience and therefore its $G_n + 2$ descendants are inaccessible and therefore not calculable (end of the route).

Bypass of rule 1:

However, it is still possible to bypass the absence of a hyperlink to the description of a $G_n + 1$ descendant and to calculate the $G_n + 2$ descendants (bypassing rule 1 in a roundabout way). A form on the web page allows you to enter an experience number "ID" to go directly to its description page without any restriction. Then, the creation of a $G_n + 2$ experience will automatically perpetuate this link, even if rule 1 does not allow it.

Bypass of rule 2:

By default, the web page describing a Gn experiment, only displays its Gn + 1 descendants allowed by rule 2 (oxygen atoms not having already been modified in this route). To also display those which do not respect rule 2 (bypass of rule 2), you must set the 'stop' field of the parent experiment Gn to 3 (this mode only concerns the current experiment Gn and have an influence only on the display of its descendants Gn + 1). This can be done by clicking on a hyperlink at the bottom of the web page (or by changing this value with phpMyAdmin, see above).

Rule 3 being essential, the bypass is not possible, but the future descendants which do not respect it (already calculated by a parallel branch of the tree) are displayed with the other authorized descendants and with their ΔE for information. No link is displayed to go to their description, in order to keep a spanning tree: no cycle in the tree and horizontal movement prohibited). To go to this experiment, one has to go backward in the previous generations, up to a common ancestor, then has to go forward in the branch containing this experiment (it is also possible to simply type the experiment ID into the form for direct access).

Installation of XAMPP, of the database and of the PHP software

The code of the PHP programs is accessible for a better understanding. It was deposited into the data repository (<https://doi.org/10.57745/CBUPP3>) and on Software Heritage passing through Hal (<https://hal.science/hal-03991394>).

This software (consisting of several PHP files, one for 1-CD systems (cdmodele.php), one for 2-CD systems (cdmodele2cd.php), one “include file” for functions common to both previous files (functions.inc), and CDxCD.php to calculate the best combination of two CDs to use as G0 model for series 2-CD S4). The database is needed to access and navigate into the trees, and to see examples that are cited inside the paper as ‘ID ...’. An alternative method, if you just want to consult the data, is to use the static web pages (see below). After installation of the XAMPP pack, downloadable for free at URL <https://www.apachefriends.org> (version 8.2.0 was used in this work), download the pascal.sql file (<https://doi.org/10.57745/38MYXM>). Start the XAMPP control panel (if error message, launch it with administrator rights) then start ‘Apache’ and ‘MySQL’ with their “start” buttons (allow them with the firewall) and click “Admin” button for MySQL to launch phpMyAdmin (or use url <http://localhost/phpmyadmin/>). To install the “pascal” database, click on “home” icon to be sure to not be into another database, click “import” in the

menu, select the “pascal.sql” file on your disk, then click “Import”. The imported database “pascal” appears in the left panel.

Access the PHP software typing, into a browser, the url <http://localhost/cdmodele.php> for 1-CD systems, <http://localhost/cdmodele2cd.php> for 2-CD systems (<http://localhost/CDxCD.php?action=show> for the special method to find the best G0 for 2-CD S4 series). This software can be freely used and modified for academic research, but in case of publication please cite the present article. For help to adapt this software to another scientific problem, the author (P.P.) can be contacted.

Useful algorithms:

Notes of main text. Calculation of the average methylation-per-glucose-unit, methylation-per-glucose-unit domains (+ extra information: percentage of Me occupancy on the narrow sides and on the wide sides of the CDs), for all systems of each series that are better than their G0 (improved systems) and their corresponding free CDs (CDs without inclusion):

The PHP code is displayed in black, the comments in green and the SQL queries in blue.

```
for(i=1 ; i≤8 ; i++) //for each series 1-CD
    nMeseries=0; //overall number of Me for this series i (systems better than G0)
    deltaELevel0 ← "SELECT deltaE FROM calculscd WHERE series='$i' AND IDparent='0'
    LIMIT 1" //deltaE of G0 system (limit to determinate improved systems)
    ntotal ← "SELECT COUNT(*) FROM calculscd WHERE series='$i'" //overall number of
    systems for this series i
    nbestg0 ← "SELECT COUNT(*) FROM calculscd WHERE series='$i' AND
    deltaE<$deltaELevel0" //number of systems better than G0 for this series i
    nMemin=21 //number of minimum Me/CD for this series=maximum possible for now
    nMemax=0; //number of maximum Me/CD for this series=minimum possible for now
    nMenarrow=0; //number of Me found on narrow side
    nMewide=0; //number of Me found on wide side
    for(j = 1; j ≤ All structures of series i that are better than their G0 system*; j++)
        nMe=0 //number of Me for this CD = 0 for now
```

```

for(k = 1 ; k ≤ 21 ; k++) //all O atoms Ok of the CD (1-CD system = 21)
    if( parity of Ok is odd ) //1, 3 or 7 => Ok is methylated
        nMe++ //one more Me found for this CD
        if( k % 3 == 0 ) //numbers on narrow side: 3, 6, 9, 12, 15, 18, 21
            nMenarrow++ //one more Me found on narrow side
        else
            nMewide++ //one more Me found on wide side
    if(nMe>nMemax) nMemax=nMe; //max number of Me/CD for this series
    if(nMe<nMemin) nMemin=nMe; //min number of Me/CD for this series
    nMeseries+=nMe; //total Me for this series
    rateMe =nMeseries / 7 / nbestg0 //average rate of Me/anhydroglucose unit for the
    best systems of this series

nfreeCD ← "SELECT COUNT(DISTINCT binarymin) FROM calculscd WHERE
deltaE<'$deltaELevel0' AND series='$i'" //number of distinct free CDs**

Displays calculated/found information

```

* All structures of series i that are better than their G0 system: "SELECT * FROM calculscd WHERE deltaE<(SELECT deltaE FROM calculscd WHERE series='\$i' AND generation='0' LIMIT 1) AND series='\$i'"

** Free CDs: In a series, certain systems can correspond to the same CD being associated to the same moiety but with different placements (rotation around the moiety). To easily find the number of distinct CDs in the series, the binarymin value is used as an identifier for each CD.

Note: for the SQL queries (in blue), the PHP variables as \$i will be replaced by their values (1 to 8) before being sent to the database management system.

For 2-CD systems, we first must retrieve the list of union of CD1s and CD2s (CDs used as both CD1 and CD2 are counted as only one): *SELECT DISTINCT binarymin1 FROM calculs2cd where series='\$series' GROUP BY binarymin1 UNION SELECT DISTINCT binarymin2 FROM calculs2cd where series='\$series' GROUP BY binarymin2*. The number of lines founds by the SQL request is the researched number of free CDs.

PHP code to retrieve the data to fill Figure 4:

Code to be copied into the PHP file (cdmodele.php) for 1CD models (not included in the published version). 2CD models works the same (PHP file cdmodele2cd.php), but replace table “calculscd” by table “calculs2cd”, change the values in array \$stablebests and limit to 4 values, replace 9 by 5 in the two for loops).

```
$stablebests[1]=1553; //list of the best models for each of the 8 series
$stablebests[2]=3738;
$stablebests[3]=5051;
$stablebests[4]=5370;
$stablebests[5]=5732;
$stablebests[6]=5838;
$stablebests[7]=6040;
$stablebests[8]=7759;

$maxgeneration=0; //max generation to be displayed (for Figure 4 of the article)
for($laserie=1;$laserie<9;$laserie++) //for the 8 1CD series
{
    $ID2=$stablebests[$laserie]; //retrieves the current series
    do //searches from best model to the G0 model (decreasing generation)
    {
        $requete="SELECT IDparent,deltaE,generation FROM calculscd WHERE ID='$ID2'
LIMIT 1"; //for model $ID2, retrieves deltaE, the generation and its parent model (its ID)
        if($row=fctselect1X($requete,1,$id)) //for this handmade function, see reference
https://hal.science/hal-03991394 (or Software Heritage whose link is inside)
        {
            list($IDparent,$deltaE,$generation)=$row; //retrieves data from database.
            if($maxgeneration==0) $maxgeneration=$generation; //finds the higher generation to be
displayed in Figure 4.
            $tabledeltaE[$laserie][$generation]=$deltaE; //saves into array for latter.
        }
    }
}
```



```

else { print("<p>ID $ID2 not found !</p>"); exit(); } //should not be.
$ID2=$IDparent; //Continues with the parent to find its own parent.
}
while($generation>0); //loop until G0 is reached.
}
for($g=0;$g<=$maxgeneration;$g++) //print data ordered by generation.
{
print("<p>G$g"); //(print the generation: G0, G1, ...)
for($laserie=1;$laserie<9;$laserie++) //then by series
{
if(isset($tabledeltaE[$laserie][$g])) $d=$tabledeltaE[$laserie][$g]; else $d=0; //if no value
for deltaE, replaces by 0 (0 will be erased in Excel afterwards). Otherwise, copies the value.
print(";$d"); //writes the value.
}
print("</p>"); //closes the paragraph.
}

```

When the webpage finishes to display the table, copy/past it into a text file, open Excel and insert the content of the file, erase all 0 values, do the graphic.

Algorithm to calculate binarymin:

As stated above binarymin is a binary representation of the CD. Indeed, each oxygen atom (O_i with i ranging from 1 to 21 in 1-CD + 31 to 51 systems for 2-CD) can only take a binary state (methylated or not), like the bits which represent the smallest information of a computer (1 or 0). In addition, the oxygen atoms are arranged in a certain order (their numbers), as an integer encoded in the memory of a computer consists of a series of numbered bits (0, 1, 2, ...), each bit having a weight equal to 2^{number} ($2^0, 2^1, 2^2, \dots$). The value of the integer, converted to base 10, is the sum of each bit multiplied by its weight (so only bits with 1 count in this sum). By analogy, we therefore converted the configuration of a CD into a binary number. The reason for creating binarymin is to simplify the SQL query to find a particular CD. Indeed, an O_i can take 6 possible values (0, 1, 2, 3, 6 and 7), which makes the request complicated by considering the 21 O_i while the search on binarymin is simple and very fast (theoretically more again if we placed an index

on this field). This binarymin number is calculated only once, when creating the experiment in the database (inserted in the SQL query "INSERT INTO"). The binarymin2 field of the second CD of 2-CD systems is calculated similarly.

Depending on the anhydroglucose unit from which we start to count, and always starting from a position 2 (in the glucose numbering system, then 3, 6, then the following anhydroglucose unit) we can obtain 7 binarymin numbers different. We only keep the smallest. The algorithms of the two functions performing this calculation are shown below followed by the explanatory diagram.

//The first function converts the configuration of the CD to a binary number depending on the starting anhydroglucose unit (Shift). Array o contains the configuration (methylation state of all O atoms 1 to 21).

```
function ConvertCDtoBinaryNonUnique(o, shift)
    binary=0 //binary number to calculate
    multi=1 // → weight of the bit = 20 (=1 for starting bit 0)
    for(i=0 ; i<21 ; i++) //for each oxygen atom Oi (i.e., o[i])
        binary += multi * o[ ( ( 3 * shift + i ) %21 ) +1 ] //adds value of the bit x its weight
        multi *= 2; //then the weight become 21, 22, 23, ... for next bit
    return binary //send calculated binary
```

//The second function converts the configuration to a unique binary number by calling seven times function ConvertCDtoBinaryNonUnique(), each one starting from a different anhydroglucose unit (the smaller binary number on the 7 possibilities, will be saved in database afterwards). Array o contains the configuration (methylation state of all O atoms 1 to 21).

```
function ConvertCDtoBinaryUnique(o)
    BinaryMin = 2097152; // 221 = max+1
    for(i =1 ; i<22 ; i++) o[i]=o[i] & 1; //0, 2 or 6 →0, 1, 3 or 7 →1 (conversion to 0 or 1 only)
    for(i=0 ; i<7 ; i++) //for each starting anhydroglucose unit
        binary=ConvertCDtoBinaryNonUnique(o, i) //start counting from anhydroglucose
        unit i
        if(binary < BinaryMin) BinaryMin=binary //retains the smaller one
    return BinaryMin //and send it
```

For example, the best 1-CD series 8 experience (ID = 7759) has an official binarymin of 9346, which corresponds, following the order of the numbers, and starting at number 1, to an H-Me-H-H-H-H-Me-H-H-Me-H-H-Me-H-H-H-H-H-H string starting the count at O1. By converting the string into 0 (H) and 1 (Me), and by returning the string, to write the binary number in conventional notation (decreasing weight), we obtain 000000010010000010, that is to say 9346 in base 10 notation. The other combinations starting the count at O4, O7, O10, O13, O16, O19 (the other positions numbered 2 in glucose notation), give larger numbers, so 9346 is retained.

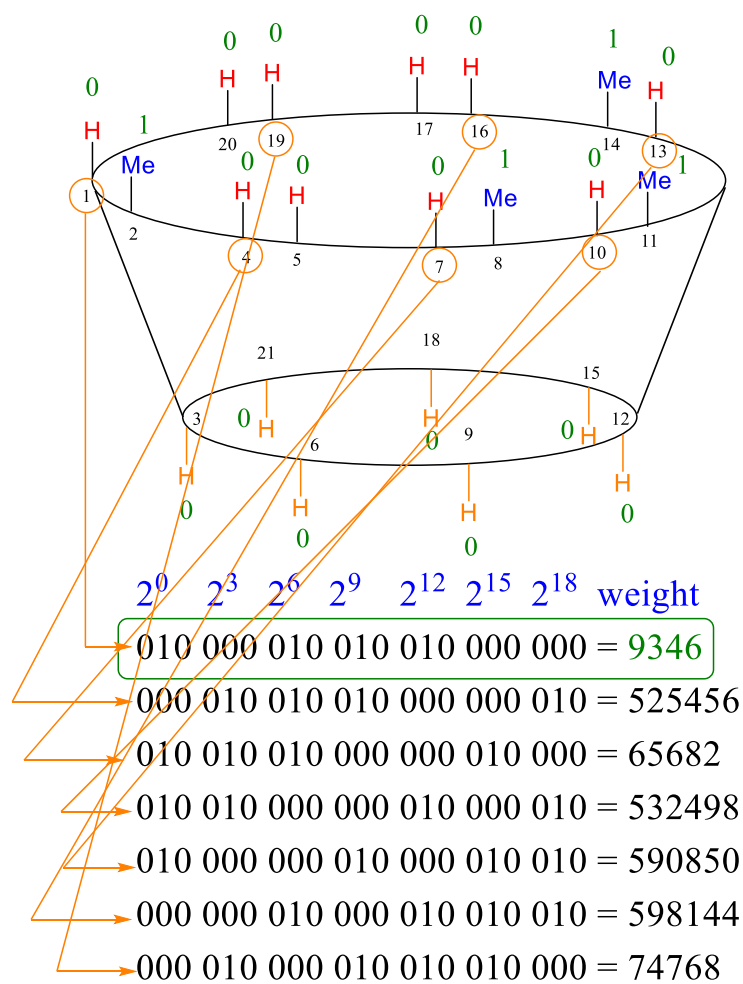


Figure S3. Determination of the binarymin code, the minimum of the 7 binary numbers calculated when starting from position 2 (glucose numbering system) of each of the 7 anhydroglucose units (orange circles). Binary numbers are represented from left to right to be consistent with the way of reading the CD but should be reversed in conventional binary notation.

With this binarymin number, it is easy to find the number of free CDs of a series 's' (when freed from supramolecular assembly, i.e., the number of distinct CDs):

- For 1-CD series (remark: DISTINCT eliminates doubletons): *SELECT COUNT(DISTINCT binarymin) FROM calculscd WHERE series='s'*.
- For 2-CD series: 1) First, we recover the list of total CDs by the intersection (UNION, that also eliminates doubletons) of CD1 and CD2: *SELECT DISTINCT binarymin1 FROM calculs2cd WHERE series='s' UNION SELECT DISTINCT binarymin2 FROM calculs2cd WHERE series='s'*, 2) then we look at the number of results the database displays.

To collect all the free CDs of the 12 series (1-CD and 2-CD) to determine in how many series they are involved, it is necessary to make the intersection of the free CDs of the 1-CD and those of 2-CD systems (by making the intersection of CD1 and CD2), by the SQL command:

SELECT DISTINCT binarymin FROM calculscd GROUP BY binarymin UNION SELECT DISTINCT binarymin1 FROM calculs2cd GROUP BY binarymin1 UNION SELECT DISTINCT binarymin2 FROM calculs2cd GROUP BY binarymin2.

Then, number of series a CD is implicated into (replace b by the researched binary value, and this for all the free CDs found above): *SELECT COUNT(DISTINCT series) FROM calculscd WHERE binarymin='b'* (for 1-CD) or *SELECT COUNT(DISTINCT series) FROM calculs2cd WHERE binarymin1='b' OR binarymin2='b'* (for 2-CD).

Insertion of a new experiment into the table 'calculscd' (called by the PHP software by its "handmade" fctinsert() function; the variable '\$num' receives the unique 'ID' value attributed by the database).: `$num=fctinsert("INSERT INTO calculscd VALUES ('0','$o[1]','$o[2]','$o[3]','$o[4]','$o[5]','$o[6]','$o[7]','$o[8]','$o[9]','$o[10]','$o[11]','$o[12]','$o[13]','$o[14]','$o[15]','$o[16]','$o[17]','$o[18]','$o[19]','$o[20]','$o[21]','$signe2$oxygenet','$path $signe O_{$oxygenet}','$IDparent','0','0','$series','$NumUniqueCD','$generation','0','$mainseries'))",0,$id).`

First parameter (ID): 0 indicates to the database to attribute a new and unique ID to the new record.

Second to 22th parameters (O₁ to O₂₁): \$o is the table that contains the values of all O atoms O_{1...21}.

Thus, \$o[i] represents oxygen atom number i into table \$o of the PHP program.

23th parameter (changed = \$signe2\$oxygenet): O atom (\$oxygenet is its number) that was changed with sign (\$signe2) – for demethylation and sign + for methylation.

24th parameter (path): “\$signe O_{\$oxygenet}” is appended to the end of the string ‘path’ to indicate in a comprehensive way, the O atom of the parent experiment that is changed in this descendant. The HTML tags < > are interpreted when displaying path in webpage (O atom number (\$oxygenet) is set to subscript).

25th parameter (IDparent'): ID of the parent experiment this experiment derivate from.

26th parameter (deltaE: ΔE): Set to 0 to indicate this value is to be calculated (pending calculation).

27th parameter (stop): No decision yet if the experiment can continue in next generation (waiting for calculation).

28th parameter (series). Series 1 to 10.

29th parameter (binarymin): Calculated and stored in PHP variable \$NumUniqueCD.

30th parameter (generation): generation was incremented by the PHP program (generation of its parent + 1).

31th parameter (deltadelta: ΔΔE): 0 for now (waiting for calculation).

32th parameter (mainseries): Permits to gather series that have the same inclusion type (same moiety + same inclusion side of the CD) but starting from a different CD for its G0 model. This is the field used (with O1 – O21) to search for an existing model to avoid creating a doubleton into the table of the database (rule 3). For details on its utility, see below (1-CD S9 and S10).

The database was first opened calling the PHP function `mysqli_connect()` (into include file `functions.inc` that was used by the three PHP files) that returns a value (stored by the PHP program into variable ‘\$id’, if the database was opened successfully) that serves to link PHP and the database it has opened. This variable \$id should be furnished with each query sent to the database. It should be noted that this variable \$id should not be confused with variable \$ID (variables are case-sensitive) that represent the number of a model (its ID field inside the database table).

All queries are sent by the PHP program to the database management system using the native “\$res=mysqli_query(\$id,\$requete);” command, where: - \$id is the handler for the database (see just above), - \$requete is a text variable containing the SQL query as a text. The result of the query is returned and stored into a variable (here \$res), whose value is ‘false’ if the query failed (SQL syntax error for example).

For illustration: The handmade PHP function `fcinsert(...)`, is used to send a 'INSERT INTO' SQL request to the database to save the new pending experiment by creating a new row (see above). This function uses the "`fcsql($requete,$stop,$id);`" command (where `fcsql` is also an handmade function, where `$requete` and `$id` were described above and `$stop` indicates if the PHP program should stop or not if the query failed). This last function calls the native `mysqli_query()` PHP function.

Example of update of several attributes of the experiment '`$ID`' in the table of the database:

When the calculation of the pending experiment was done by Spartan, and the energies were entered the form in the webpage, the PHP program receive the data, calculates `deltaE` (ΔE calculated and stored in PHP variable `$e`), and allows or not to continue in next generation ('`stop`'), depending on calculated '`deltadelta`' (variation of ΔE ($\Delta\Delta E$) calculated and stored in PHP variable `$deltadeltacal`):

UPDATE calculscd SET deltaE='\$e', stop='\$stop', deltaxdelta='\$deltadeltacal' WHERE ID='\$ID' LIMIT 1.

LIMIT 1 was added to the request to limit the research of the experiment in the database to only one (there is only one to be modified, so searching all the database when one was already found is a loss of time).

Case of 1-CD series 9 and 10

A question that arises using our method is: "Is this technique able to detect the best models?" As said in the article, we are not looking for the global minimum, because no complex CD will be synthesized to make this assembly. However, an evaluation of the effectiveness of the method is required to appreciate the convergence towards a local minimum close to the global minimum (which could theoretically only be found and proven by making a calculation on all the models, which is currently impossible). To help answer this question, we decided to make a modification to the method and the web application first used for 1-CD S1 to S8. Two new 1-CD series have been made (S9 and S10), but with the same inserts as for the S8 series (reminder: Succ inserted into the wide side of 3,6-diMe-CD). The goal was not only to see if we converge to a better minimum than S8 starting from a different point into the combination space, but also to find if we converge to some local minima already found by S8. Starting from different CDs (for S9, from 2-

Me-CD which is the exact inverse of 3,6-diMe-CD, and from 3-Me-CD for S10), the 2 series each formed a tree of mutations evolving in parallel, with only one exception: rule 3 (no duplicates) applies for S8 UNION S9 UNION S10 thanks to the creation of a new field in the database, "mainseries". For S8, S9 and S10, this field has been set to 8 (mainseries = series for others), indicating that the PHP program must now rely on this field, and not on the "series" field to detect duplicates before they are created. It is this rule that must make it possible to see the collision between these 3 series/trees and therefore to see if we converge well towards the same models. However, by analyzing these models, we realized that the Succ group was not well inserted into the CD, and worse, that after a few generations, it was the Fc group that was partially inserted automatically into the CD cavity (see ID 9191 for S9 and 9690 for S10), which corresponds more to S2 than to S8. The S9 and S10 series have therefore been distorted and have been disassociated from S8 in the database (they became independent, setting mainseries = series using a SQL query). They have also not been associated with S2 because some of their models are intermediate between S8 and S2 and moreover, the control of duplicates was not made with respect to S2. These two series have therefore not been included in Table 1 on main text, but their models are provided for information as for the other series in the form of XYZ files. We wanted to know more about the reason for this failure and noticed that from G0, Succ is almost not inserted into the cavity. The reason is due to hydrogen bonds between one (e.g., ID 8996) or two carbonyls (ID 8997) of the imide which anchor the succinimide group at the cavity entrance. However, methylation of one of these OH groups on the wide side of the CD, even though breaking the hydrogen bond, does not allow Succ to penetrate deep into the cavity because the carbonyl then finds another hydroxyl to replace it (for example ID 8909 – (+ O₈) → ID 8914 where O₈ is replaced by O₁₂). Forced manual insertion of Succ when modifying the model does not work around this problem; when calculating the geometry, Succ is still sticking out of the cavity. We have therefore demonstrated that this method can succeed or fail depending on the CD chosen for the G0 generation, and that it is preferable to start from the models with the lowest ΔE for G0, i.e., from the most suitable CD, as we instinctively did for the others before the S9 and S10 series. However, the method of starting from different points into the combination space, to browse larger space (but remaining reasonable), is validated, since it could permit to get around falling into a local minimum that hampers to reach a better minimum.

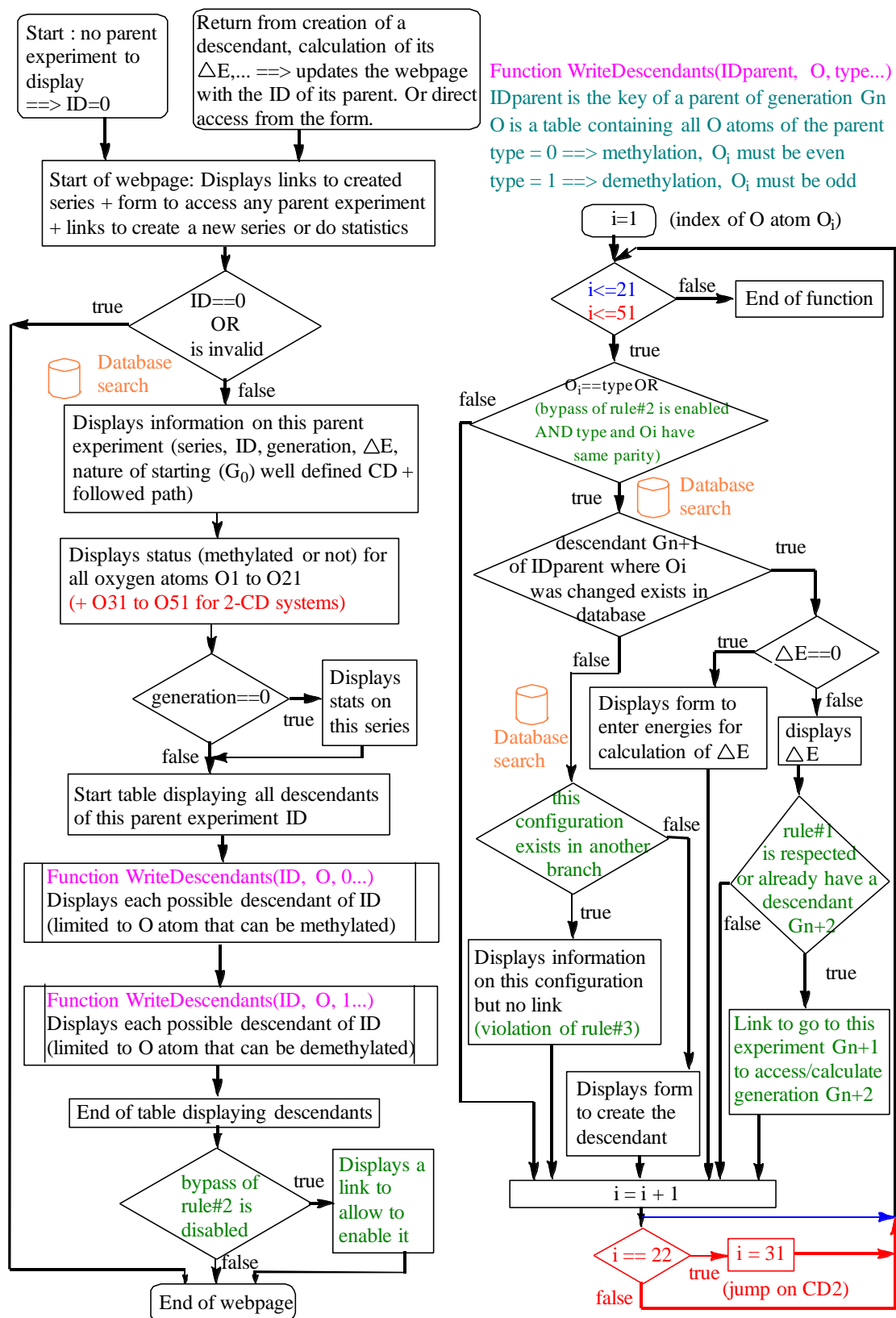


Figure S4. Algorithm of the two PHP programmes. Drawing in blue is specific to 1-CD systems and drawing in red to 2-CD systems. Effects of rules 1,2,3 and their bypasses are drawn in green.

Examples without XAMPP (static webpages)

To be able to see the results of the web application without installing XAMPP, all the corresponding static webpages were created by the C program (see below) and can be seen using a navigator. These webpages can be found into the data repository and into Software Heritage (links in the first page of this document) as a compressed archive. The archive should be downloaded, unzipped, and almost any webpage (preferably the first HTML file) can serve as an entry point to then reach any model. Indeed, the links work, permitting to navigate off-line into the models.

C program to control the models

This C program works on 1-CD series or on 2-CD series depending on a constant set in the program (“#define CD1OR2 1” to work on 1-CD series, “#define CD1OR2 2” for 2-CD series) before compiling it. It needs the corresponding database table to work. For this purpose, the tables were exported from phpMyAdmin as tab files (openable with a text editor, alternatively with Excel). In fact, these files were saved from phpMyAdmin as .csv files then renamed as .tab files (extension changed) to be compliant with the data repository system (to be visible on-line). The needed files to be opened by the C program are “calculscdprog.tab” for the “calculscd” table and “calculs2cdprog.tab” for the “calculs2cd” table. These files have tabulations as separator and the fields are not escaped by quotes. It should be noted that these tab files only differ from the “calculscd.csv” and “calculs2cd.csv” files (also in the repository) by their format (separators = comma, fields escaped by quotes), format requested by the repository for CSV files. These files make it easy to recreate the database regardless of the OS or database management system. But the C program can also read the .tab file, extracts its content, and fills a structure (typedef struct { ... } table;) whose instance is called “database”. Some of the SQL queries executed by the PHP program are simulated by the C program by accessing the fields of this structure (or an adapted copy where the Oi fields are simplified (0 or 1 only) to simplify the simulated SQL query).

The program also needs to have access to the folders where the models' files are saved. When a parent model was used to create its descendants in Spartan, a folder was created inside the same folder than the parent and all descendants were put inside, and so on for several generations. When the XYZ files were created from Spartan, these files were saved into the same folder as its original Spartan model file. The program can know and access to the corresponding folder and file of each model thanks to the "path" field of the table/.tab file. However, some information is lost in XYZ files, such as chemical bonds and atom labels. Nevertheless, these text files are easily openable by handmade software, which can recreate the lost chemical bonds according to the interatomic distances, and then to recreate the molecule by linking neighboring atoms and control if the files and the database match.

Although the PHP program could analyze these models, this procedure is not especially applicable because of the constraints applied to a web application, which thus shows its limits. A "standalone" program is therefore desirable, and we have carried out this audit program in the C language. The data can be easily retrieved, each line representing a model.

The C program was created for several purposes:

- As stated above, for fast consultation of the trees of models displayed into the dynamic webpages by the PHP program, but without the need to use it, static webpages, one for each model, were furnished into the repository as clones of the dynamic webpages. Saving all pages one by one from the browser would have caused several issues: - It would have taken a long time, - Manually giving to the saved webpages a name related to their ID would have caused some errors, - The original links would have been unsuitable to operate off-line, hindering the navigation into the trees, or would have necessitated to manually adapt the links for each page. To create these static webpages, the behaviour of the C program was copied on the code of the PHP program but adapting the links.
- When using a parent model to create its descendants in Spartan, obviously errors are unavoidable. The errors can be: - changing the wrong oxygen atom or forgetting to change it, - accidentally erasing an atom (this unvalidated a whole series having more than 1000 models that had to be redone because the GO was wrong and the error has spread into descendants), - starting from the wrong parent, forgetting to convert some Spartan models into XYZ files... The C program having access to the

“path” field of the database, was able to determinate the path to the file, to look for its presence, to open it and to create a modified copy. The modifications are:

- Gathering all XYZ files into the same folder “XYZ” (the one that was compressed into an archive deposited into the repository), but each series being into its own folder “aCDSb” where “a” is the number of CDs (1 or 2) and “b” is the series.
- Changing the name of the copy to comply with the format “a-CD-Sb-ID-c.xyz” where “a” and “b” are those described above and c is the ID of the model.
- Fixing an error of Spartan that does not respect the format for XYZ files. Indeed, the two first lines are missing when saving XYZ files from Spartan 14 (not from Spartan 20). These two lines were added with the first line being the number of atoms into the model/XYZ file, and the second line being a facultative comment (that was filled with information about the model, based on its “path” field).
- Removing the two pseudo atoms “Lig” than Spartan adds to connect ligands to the metal atom, here Fe connected to two Cp (cyclopentadienyl), but that are confused with lithium by other modelling software.
- Accessory, reducing the size of the XYZ files by replacing several consecutive spaces by a unique tabulation.
- Finding the real name of the original XYZ file. Indeed, for high generation models, the path to the file became too long, provoking errors in Windows 10. The only possibility, if we want to keep the path to the folder, was to reduce the name of the original XYZ file to reduce the total path (folders path + file name). To do this, the beginning of the file was truncated. The C program needed to scan the last folder of the path to find the real name of the file finishing with "G0.xyz" for the G0 model, "+a.xyz" for models finished by a methylation or "-a.xyz" for models finished by a demethylation, where “a” is the modified oxygen atom.

Then, the program reads the new XYZ file and writes into a file (message.txt) all errors that were found. To check if the configuration of the CD(s) was correct, the “binarymin” of the XYZ file was calculated and compared to that of the database: Starting from the first found

anomeric carbon atom, the program progress from neighbour atom to the next one, checks the methylation state of the three important oxygen atoms on its way, and this for each glucose unit. It should be noted that this method does not guaranty to find all errors for the G1 models, because three groups of CDs have in common the same (free) CD (those having the inversion of methylation on the same number, in glucose numbering: 2, 3 and 6). Moreover, the number of atoms with a certain valence where also counted (H, O, C and N).

- Hydrogen bonds were found to be important for the stability of the supramolecular models, in particular hydrogen bonds implicating the iron atom of ferrocene. We used the C program to do extra statistics based on these hydrogen bonds but limited to intermolecular bonds (between the inserted molecule and a CD). To find these bonds, we fixed the limit to 2 Å (#define HBONDLIMIT2 4.0 in the C program is the square of this value, because with wanted to compare squares of distances to avoid using the square root function that is time consuming). For SuccFerr the atoms involved in these bonds are the iron atom, the two phenols (-O- and -H independently) and the two carbonyls (=O) of the imide group. For CDs these are the hydroxyls (-O- and -H independently), the ethers (-O-) and the anomeric group (-O-). The C program created files Hbonds-aCD.txt, where “a” is the number of CDs (1 or 2). Then, it has reread these files to make a webpage containing statistics on these bonds, including the found clamps (“TableofHbondsbySeries-aCD.html” where a is the same as above, inside the compressed archive where the other webpages are gathered).

Details on the C program and call to functions

The C program flows in this order:

- Program starts.
- Creates the structure “table” and creates an instance: “database”. It will contain all the fields of the MariaDB table (“calculscd” for 1-CD and “calculs2cd” for 2-CD models) plus complementary fields that are calculated on-demand by the PHP program but that will be calculated once by the C program and stored in “database”. Its size is set to 9800 (#define NLINESMAX 9800) for 1-CD or 2-CD models.
- Opens the "message.txt" file for writing. All found errors (if any) will be written inside.
- Opens "Hbonds-aCD.txt" for writing (where “a” is the number of CD). All H bonds of interest and Me (methyl groups) close to Fc (ferrocene) will be written inside.

- Calls “ReadCSVToTable()”:
 - Open the csv file ("calculscdprog.tab" or "calculs2cdprog.tab").
 - Reads each line of the file (one line = one model) and fills the database structure for each record “ID”. Since the table was ordered, ID = number of the record.
 - If constant “MAKEXYZFILES” was set to 1 (default, allows to create adapted XYZ files):
 - Calls function “CopyConvertedXYZFiles()” that reads, modifies and copies the new corresponding XYZ file for this experiment, knowing its ID and path.
 - Calls function “ChecksBinaryConfiguration()” that reads the new adapted XYZ file, calls function “VerifyCount()” that counts the number of atoms by type and valence, checks if incoherences exist into the XYZ file (writes errors in file "message.txt"), calls function “CalculationOfBinary()” that calculates the “binarymin” for the CD (then the “binarymin” for the second CD, for 2-CD models).
 - Checks for errors (if calculated “binarymin” for the XYZ file and in database are different) and if any, writes in file "message.txt".
 - If constant “MAKEHBONDSSTUDY” is set to 1, calls function “FindFe()” to search for the iron atom, calls function “FindThe2Carbonyls()” to find the two carbonyl groups (imide), calls function “FindThe2Phenols()” to find the two oxygen atoms and the two hydrogen atoms of the two phenol groups. Then calls function “FindHBonds()” for each type of hydrogen bond, that are written into file "Hbonds-1CD.txt" or "Hbonds-2CD.txt". Calls function “FindNumberOfMeCloseToFe(Fe)” to count the number of Me close to ferrocene (to Fe) and to write this information into file "Hbonds-1CD.txt" or "Hbonds-2CD.txt".
 - Determines how many series are inside the CSV file (store into variable “NumberOfSeries”) then closes it.
- Closes file "Hbonds-1CD.txt" or "Hbonds-2CD.txt".
- For each series, calls function “TreatTree()” to complete the database structure by some calculations (for values not in the original database but calculated on-demand by the PHP program):

- Determines, in “database”, the area where the models of this series are located (to increase the search speed by reducing this area).
 - Calls the recursive function “TreatTreeRecursive()” that browses this model and all its descendants (sub-branch starting from this model) to determine all fields not in the original database: “best” (number of descendant models that are better than this model), “pending” (number of models that are pending, i.e., whose ΔE was not calculated yet), “finished” (all descendants are blocked by rule 1 or not, dead end or not for this sub-branch), “max_generation” (the higher generation reached inside this sub-branch), “bestdeltaE” (the higher ΔE reached inside this sub-branch), “descendants” (the number of descendants this model has inside the sub-branch this model is the ancestor).
- If constant MAKEHBONDSSTUDY and constant MAKEXYZFILES are set to 1, creates a stat report on H bonds by reading file “Hbonds-1CD.txt” or “Hbonds-2CD.txt”.
- Opens file “Hbonds-1CD.txt” or “Hbonds-2CD.txt” for reading.
 - Calls function “CreateHBondsStatReport()” to create the report uniquely with data of the txt file.
 - Opens file "TableofHbondsbySeries-aCD.html" (“a” = number of CDs = 1 or 2) for the webpage containing the statistics about hydrogen bonds and Me close to Fc for each series. Starts to write the beginning of its HTML tag + information.
 - Finds each series inside the text file.
 - For each series, calls function “CalculatesLimitDeltaE()” to Find the minimum and maximum ΔE for this series and the average value (will be needed to find if a model should be classified as good or worse model).
 - For each series, reads each model, class it as good or worse models, counts the number of hydrogen bond for each type and the average number of Me close to Fc found. Writes the values into a table into the webpage, with a column for the good models and a column for the worse models.
 - Closes the HTML file.
 - Closes the text file.
- If constant MAKEHTMLFILES was set to 1 (creates the webpages for all the models)

- Calls function “AdaptedDatabase()” to create a copy of array “database” (dynamic allocation) with simplified fields O_i : 0 or 1 only. It permits to simplify the query that will imitate the SQL query to find if a model exists in database.
 - Calls function “CreateTheWebPages()”.
 - Determines, in “database”, the area where the models of this series are located (to increase the search speed by reducing this area).
 - For each series, Calls function “CreateOneWebPage()” that creates the webpage for the model whose ID is furnished in parameter. It opens the webpage, includes into the webpage the beginning that is common to all webpages, then the content that depends on the database. It calls function “WriteDescendants()” for the descendants of this model that were methylated, then for the descendants that were demethylated, then closes the webpage.
 - Free the memory allocated for the copy of “database”.
- Closes text file “message.txt”.
 - End of program

Case of the best models of 1CD series 8 and series 2 (figures S5 and S6)

Comparison of best model of 1CD series 8 calculated in PM3 (Spartan 14, Figure S5a) and in DFT (Spartan 20, Figure S5b):

Note: This DFT calculation (Figure S5b, ω B97X-D, 6-31G*, medium=water, option PCMRAD=FE~2.2, starting from the PM3 model) is different from Figure 4 (main text) that was calculated with the 6-311G* basis set and Gaussian 09, but gave comparable results.

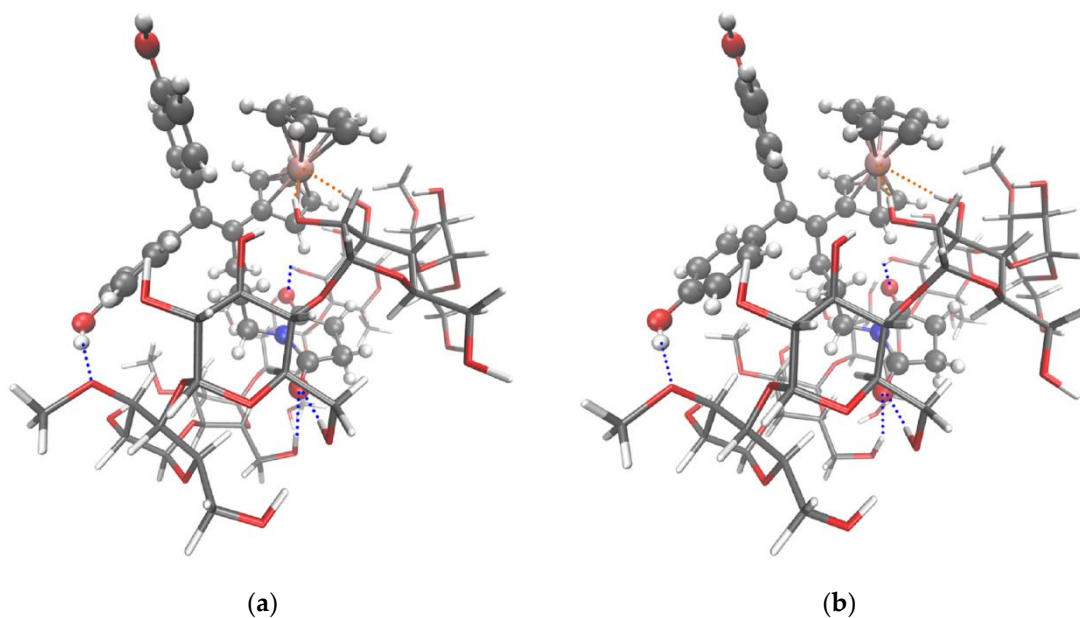


Figure S5. Molecular models of the best 1-CD system (series S8, ID 7759, Succ entering by the wide side of the CD). SuccFerr is represented with a “ball-and-stick” model and the CD is represented with a “licorice” model. Orange dashed lines (clamp between the iron atom and O₁₉-H and O₂₀-H) represent the two atypical hydrogen bonds between the iron atom and the CD. Another clamp between one of the C=O group and O₁₂-H and O₁₈-H is also visible, as bonds between the phenol group (Ph2) and O₁₄-Me and between the second C=O of imide and O₅-H (blue dashed lines). The numbering is not shown. **(a)** Computed with Spartan 14 at the semiempirical PM3 level of theory (i.e., in vacuum), with length of Fe-H bonds ≈ 1.9 Å and angles of O₁₉-H...Fe and of O₂₀-H...Fe $\approx 167^\circ$ and 177° . **(b)** Computed with Spartan 20 at the DFT level of theory, with length of Fe-H bonds ≈ 2.6 Å and angles of O₁₉-H...Fe and of O₂₀-H...Fe $\approx 165^\circ$ and 169° .

Best models of 1CD series 2 (PM3) (this calculation did not succeed in DFT):

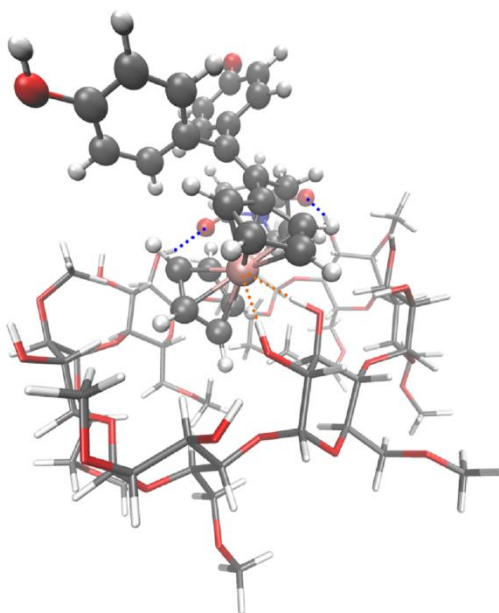


Figure S6. Molecular model of the best system of 1-CD series S2 (ID 3738, Fc included by wide side of the CD, $\Delta E = -261$ kJ/mol) computed at the semiempirical PM3 level of theory. SuccFerr is represented with a “ball-and-stick” model and the CD is represented with a “licorice” model. The 2 hydrogen bonds between the iron atom and the CD are represented by orange dashed lines (clamp between the iron atom and O₁₀-H and O₁₁-H, length ≈ 1.9 -2.0 Å). Hydrogen bonds also exist between each C=O of imide group and -O₁-H and -O₁₇-H (blue dashed lines).

Table S4. Second-order perturbation stabilization energies for the NBOs involving atoms taking part in the atypical H-bond Fe-HO interactions. NBO labels: LP = Lone pair, LP* = Antibonding lone pair, CR = Core.

Fe ----- H₂₁₀ – O₁₉	NBO Donor	NBO Acceptor	E⁽²⁾ (kJ/mol)
	LP (1) (Fe)	Rydberg (1) (H ₂₁₀)	0.92
	LP (1) (Fe)	Rydberg (4) (H ₂₁₀)	0.21
	LP (1) (Fe)	σ^* (1) (H ₂₁₀ – O ₁₉)	1.80
	LP (2) (Fe)	σ^* (1) (H ₂₁₀ – O ₁₉)	1.67
	LP (3) (Fe)	σ^* (1) (H ₂₁₀ – O ₁₉)	0.33
	LP* (6) (Fe)	Rydberg (1) (H ₂₁₀)	25.48
	LP* (6) (Fe)	Rydberg (2) (H ₂₁₀)	2.80
	LP* (6) (Fe)	Rydberg (3) (H ₂₁₀)	6.69
	LP* (6) (Fe)	Rydberg (4) (H ₂₁₀)	14.85
	LP* (6) (Fe)	Rydberg (5) (H ₂₁₀)	0.75
	LP* (7) (Fe)	Rydberg (2) (H ₂₁₀)	0.67
	LP* (7) (Fe)	Rydberg (4) (H ₂₁₀)	0.38
	LP* (8) (Fe)	Rydberg (1) (H ₂₁₀)	1.97
	LP* (8) (Fe)	Rydberg (3) (H ₂₁₀)	0.29
	LP* (8) (Fe)	Rydberg (4) (H ₂₁₀)	1.38
	LP* (9) (Fe)	Rydberg (5) (H ₂₁₀)	1.51
	LP* (6) (Fe)	Rydberg (1) (O ₁₉)	0.50
	LP* (6) (Fe)	Rydberg (3) (O ₁₉)	1.80
	LP* (6) (Fe)	Rydberg (4) (O ₁₉)	1.88
	LP* (6) (Fe)	Rydberg (7) (O ₁₉)	0.29
	LP* (7) (Fe)	Rydberg (1) (O ₁₉)	2.09

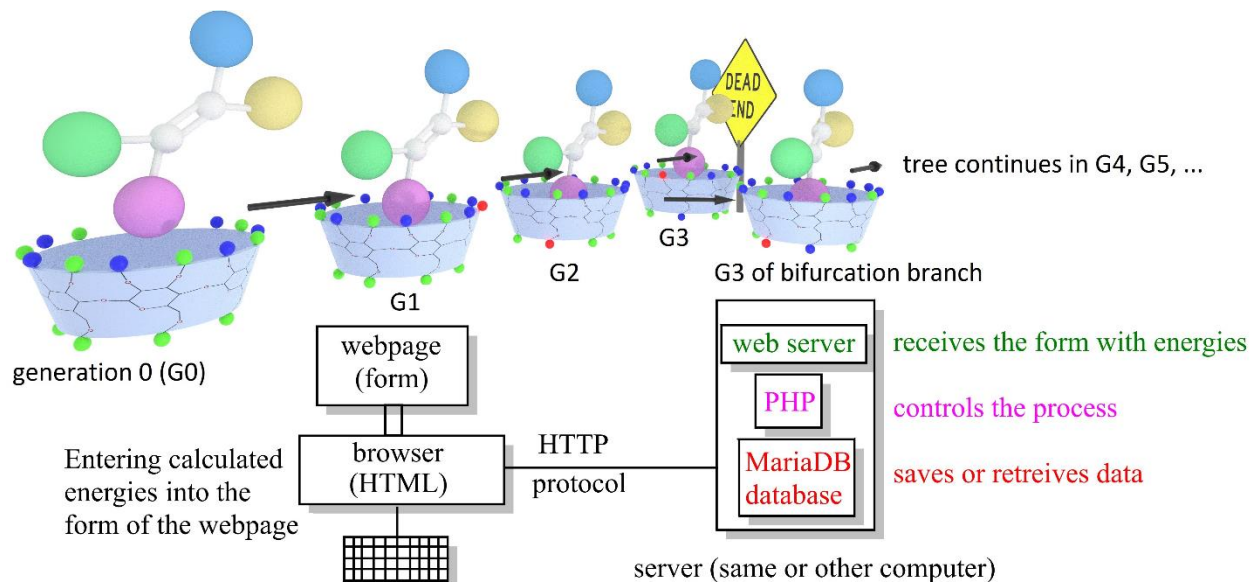
LP* (7) (Fe)	Rydberg (4) (O ₁₁₉)	1.71
LP* (8) (Fe)	Rydberg (3) (O ₁₉)	0.21
LP* (9) (Fe)	Rydberg (7) (O ₁₉)	0.75
LP* (8) (Fe)	σ^* (1) (H ₂₁₀ – O ₁₉)	4.48
σ (1) (H ₂₁₀ – O ₁₉)	LP* (6) (Fe)	7.49
σ (1) (H ₂₁₀ – O ₁₉)	LP* (7) (Fe)	0.96
σ (1) (H ₂₁₀ – O ₁₉)	LP* (8) (Fe)	1.17
σ (1) (H ₂₁₀ – O ₁₉)	LP* (9) (Fe)	0.21
σ (1) (H ₂₁₀ – O ₁₉)	Rydberg (9) (Fe)	0.29
LP (1) (O ₁₉)	LP* (7) (Fe)	0.25
LP (2) (O ₁₉)	LP* (6) (Fe)	0.67

Fe ----- H₁₉₃ – O₂₀

CR (1) (Fe)	Rydberg (4) (H ₁₉₃)	0.21
LP (1) (Fe)	Rydberg (1) (H ₁₉₃)	1.34
LP (1) (Fe)	Rydberg (3) (H ₁₉₃)	0.63
LP (1) (Fe)	Rydberg (4) (H ₁₉₃)	0.92
LP (3) (Fe)	Rydberg (3) (H ₁₉₃)	0.25
LP (1) (Fe)	σ^* (1) (H ₁₉₃ – O ₂₀)	1.30
LP (3) (Fe)	σ^* (1) (H ₁₉₃ – O ₂₀)	1.60
LP* (6) (Fe)	Rydberg (1) (H ₁₉₃)	23.68
LP* (6) (Fe)	Rydberg (3) (H ₁₉₃)	18.33
LP* (6) (Fe)	Rydberg (4) (H ₁₉₃)	38.99

LP* (6) (Fe)	Rydberg (5) (H ₁₉₃)	0.42
LP* (7) (Fe)	Rydberg (2) (H ₁₉₃)	1.63
LP* (8) (Fe)	Rydberg (1) (H ₁₉₃)	0.79
LP* (8) (Fe)	Rydberg (3) (H ₁₉₃)	0.84
LP* (8) (Fe)	Rydberg (4) (H ₁₉₃)	0.96
LP* (9) (Fe)	Rydberg (5) (H ₁₉₃)	12.76
LP* (8) (Fe)	σ^* (1) (H ₁₉₃ – O ₂₀)	4.43
σ (1) (H ₁₉₃ – O ₂₀)	LP* (6) (Fe)	6.23
σ (1) (H ₁₉₃ – O ₂₀)	LP* (8) (Fe)	2.38
σ (1) (H ₁₉₃ – O ₂₀)	Rydberg (1) (Fe)	0.25
σ (1) (H ₁₉₃ – O ₂₀)	Rydberg (11) (Fe)	0.25
LP (1) (O ₂₀)	LP* (7) (Fe)	0.21
LP (1) (O ₂₀)	LP* (8) (Fe)	0.21
LP (2) (O ₂₀)	LP* (6) (Fe)	0.63

Description of the trees of modelling and of the web application to manage them



Explanation: In a very schematic view, the molecule (SuccFerr, with its central double bond where are attached 4 moieties (Fc, Ph1, Ph2 and Succ)) is partially inserted into a CD by one of its moieties. The methylation state of the CD was changed at each generation, inverting all Me and H (blue and green spheres, on wide and narrow side of the CD, the current modification for this generation is a red sphere). The best G1 model (other G1 models are not shown for clarity) was used to do the same for several generations. To show that the model is more stable, its insertion into the CD is drawn deeper (even though this is not so simple!). If the model is less stable (shallow insertion), this branch is finished (dead end, here in G3, the worse model is drawn specially for illustration) and a better model of the same generation is used to continue the tree, or from a previous generation.

The server has the XAMPP pack installed (Apache web server, PHP interpreter, MariaDB database server). The client (same computer or another one) communicates with the server by HTTP protocol. The server sends the dynamic webpage (content based on the PHP program and on the content of the database) to the client that displays it using its web browser (compatible with any OS, even if the server and the client are different computers that have different OS).

The image was created with Blender (up) and with Chemdraw (down) and was assembled with Paint.