

NMRlipids Databank: Overlay Databank of Lipid Membrane Simulations Arising from Open Collaboration

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ABSTRACT

We present a databank of lipid bilayer simulations from the NMRlipids open collaboration project.

1 Introduction

The importance of sharing MD simulation data following the FAIR principles¹ has been widely recognized^{2–9}, and databanks are emerging^{9–16}. The relevance of quality evaluation of simulation trajectories in databanks regarding technical details of simulations and accuracy of the underlying physical description of the system (force field) has become evident^{3,7,10} and such quality evaluation has in some cases also been implemented^{10,12}. However, straightforward quality comparisons between individual simulations or force fields within these databanks remain challenging. While importance of such databanks for MD simulations is widely recognized^{2–9} and different kinds of approaches are emerging^{9–16}, generally accepted protocols and best practices are still under active development.

Here we present a solution for lipid bilayers based on overlay databank structure illustrated in Fig. 1. The concept of overlay databank is developed here to solve the practical challenges in generating databanks of MD simulation data enabling flexible analyses over large sets of simulation data, but it potentially used for wide range of situations, particularly when storage of raw data requires significant resources and final outcomes or best practices are not yet clear, overlay databank approach lowers the barrier to start without compromising the long term stability or scalability.

The practical relevance of the NMRlipids databank is exemplified by automatic quality evaluation and ranking of large amount of MD simulation data, data-driven analysis detecting correlations between properties of model cell membranes and analyses of rare phenomena that are beyond the scope of standard MD simulation studies. The NMRlipids databank provides new tools for researchers in wide range of fields in academia and industry from cell membrane biology to lipid nanoparticle formulations and data-driven computational chemistry and machine learning.

2 Results

2.1 Design of the NMRlipids databank

The key idea of the overlay databank is that the storage of raw data in layer 1 is distributed in publicly available repositories or other servers with long term stability and permanent links such as digital object identifiers. The core of the databank, layer 2, contains only information on the location and content of the raw data, thereby not requiring large resources to handle and maintain. This lowers the barrier for starting such databank as well as for long term storage. The NMRlipids databank is essentially a git repository containing information on the location of raw data and its indexing with universal naming convention. In addition to all computers where the databank is developed and used, the NMRlipids databank git is stored to Zenodo server, thereby enabling a very cost effective long term storage for the databank. The databank can be used in layer 3 by accessing the raw data and information stored in the databank by employing the universal naming conventions for atoms, molecules and simulation details. The applications can be linked to the core databank (layer 2) without actually including them, thereby enabling flexible extension of the data without compromising the simplicity and lightness of the core databank.

Currently the databank is composed of approximately 500 trajectories with the total length of approximately 231 microseconds of which most are contributed for the previous publications from the NMRlipids open collaboration^{17–20}. The distribution of lipids, force fields, length of the trajectories and available binary mixtures are shown in Fig. 1.

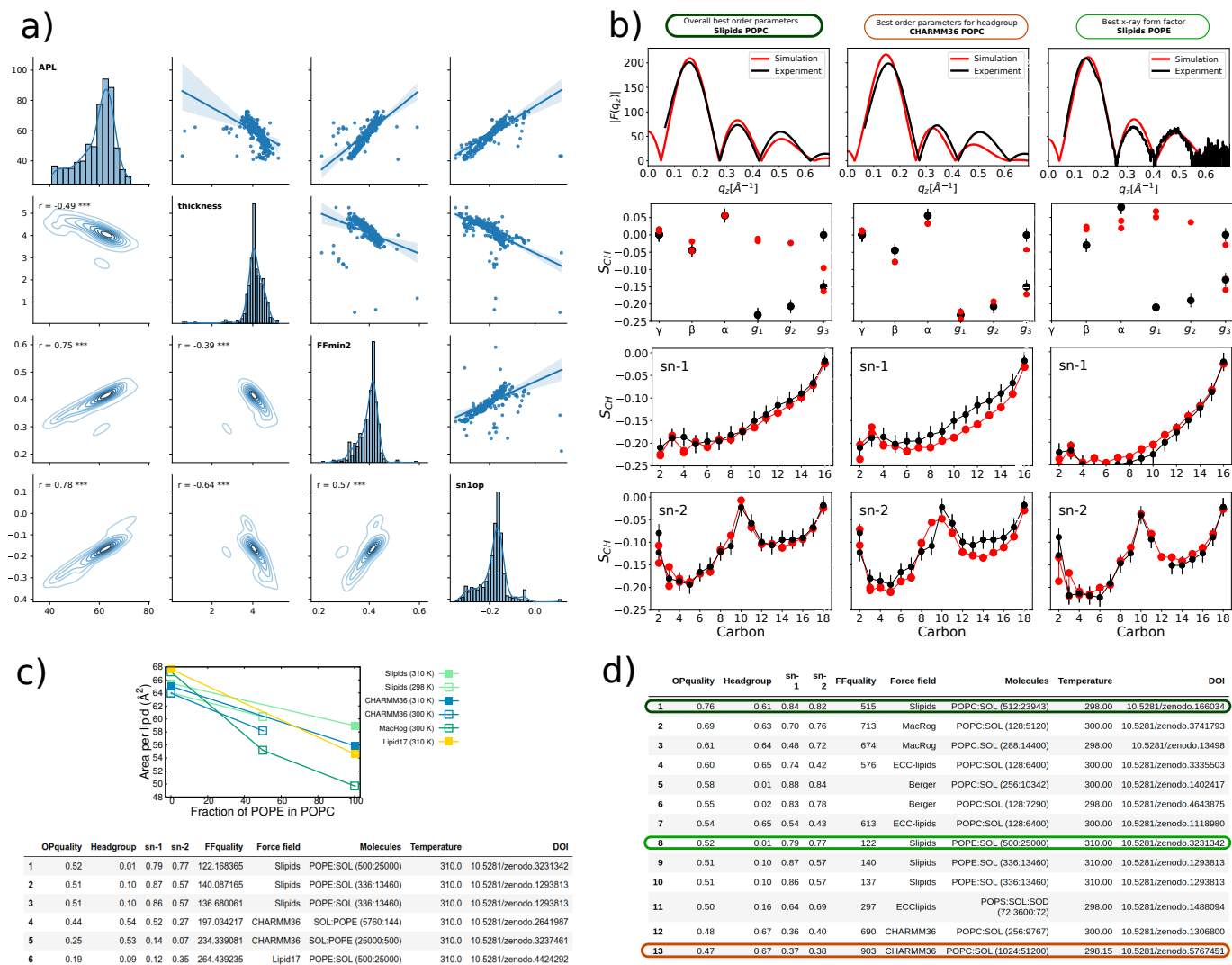


Figure 2. a) Correlations between membrane area per lipid, thickness, second minima of form factor and average order parameter of the *sn*-1 acyl chain extracted from the NMRlipids databank. All Pearson correlation coefficients shown in bottom left corner have p-value below 0.001. b) Evaluation against experimental data exemplified for the simulation giving the best qualities in overall, for headgroup and x-ray scattering form factor. c) Change in area per lipid upon addition of POPE to POPC in different simulation extracted from the NMRlipids databank and quality evaluation table of POPE simulations. d) Quality evaluation table showing the best 13 simulations according to the overall order parameter quality.

factor qualities can be compared only between simulations evaluated against the same experimental dataset.

The power of NMRLipids quality metrics in selecting the best model for particular application is demonstrated in Fig. 2 c) where the area per lipids of POPC:POPE mixtures from different force fields are shown and quality of POPE simulation in different force fields are evaluated. Among the simulations available in the NMRLipids databank, the Slipids force field gives the best quality in terms of acyl chain order parameters and form factor, and predicts the largest area per molecule for POPE and smallest difference with POPC. In conclusion, simulations with Slipids is the most reliable force field for membrane packing in POPC:POPE mixtures although its glycerol backbone is not accurately modelled. Similar comparisons utilizing the preliminary data from the NMRLipids databank have concluded that Slipids is relatively good model also for mixtures of charged POPC:POPS membranes, although models with better counterion binding have higher quality, but CHARMM36 is the best to study differences in headgroup conformational ensembles between different lipid types. Understanding such complex picture of lipid bilayer MD simulation quality would not be possible without automatic quality evaluation of large sets of simulations enabled by the NMRLipids databank.

2.3 NMRLipids databank reveals correlations between membrane properties

The power of NMRLipids databank to find correlation between membrane properties is exemplified in Fig. ?? showing how water diffusion through and along the membrane correlate with different properties of systems. While NMRLipids databank enables similar analyses for any membrane properties, these are selected due to their relevance in the development of MRI imaging methods[?].

3 Discussion

Quality measure and automatic quality evaluation of lipid bilayer MD simulations introduced in the NMRLipids databank enables rapid ranking of available simulation models against experimental NMR and x-ray scattering data. This provides a tool for researchers to rapidly evaluate the credibility of MD simulations of their own and reported by other groups. Because such tool and quality measure has not been available, this will elaborate the quality of published MD simulations of lipid bilayers and reduce potentially misleading results[?]. The power of NMRLipids databank to select the best models for particular applications has been demonstrated for PC/PE lipid mixtures (Fig. 2), PC/PS lipid mixtures[?], and lipid headgroups²⁰.

The increasing amount of MD simulation data with programmatic access in the NMRLipids databank opens up possibilities for wide range of applications utilizing the large set of accessible data. Extend of the data in the NMRLipids databank in terms of quantity (e.g., simulation length and number of conformations), content (e.g, lipid compositions and ion concentrations) and quality enables analyses that are not possible to conduct from MD simulation data produced by a single research group. Applications of the NMRLipids databank to understand how diffusion of water through and along cellular membrane depends on its physical properties are demonstrated in Fig. ?. Permeation of water through membranes resembles the permeation of also other hydrophilic molecules, such as drugs, and detailed understanding of water dynamics through and along membranes is potentially useful for the development of MRI imaging methods²⁵. These examples demonstrate the practical applications of NMRLipids databank on problems in the biological and biomedical sciences.

Building accessible databanks of molecular dynamics simulation data has been challenging due to the required long term support for hardware and software maintenance. In the overlay model used in the NMRLipids databank, the demand of hardware can be distributed and open collaboration model reduces the risk for ending software maintenance. Furthermore, the open collaboration model used in the NMRLipids project credits contributors by offering authorship in published articles, thereby creating an incentive for contributions. This model could be extended also to other fields where similar barriers to establish publicly accessible databanks exist. Emerging applications of machine learning are increasing the impact of such databanks. For example, the existing Protein Databank (PDB)[?] containing experimentally determined protein structures with programmatic access has enabled the development of machine learning based tools building on the data collected in the databank over the years[?]. NMRLipids and other databanks with open programmatic access have potential to lead similar unforeseen applications in the future.

4 Methods

4.1 Structure of the databank

Structure of the NMRLipids databank is illustrated in Fig. 3. The required input information to create an entry into the NMRLipids databank are listed in table 1. While the raw simulation data is not directly stored in the NMRLipids databank, permanent links from where the raw data can be accessed have to be given and are then stored in the README.yaml files at <https://github.com/NMRLipids/Databank/tree/main/Data/Simulations>. These files contain all the essential information listed in table 1 on each simulation entry that are needed for further applications. The raw MD simulation data can locate in any stable publicly available repository, although all the current data locates in Zenodo www.zenodo.org.

key	description	type
DOI	DOI from where the raw data is found	user given (compulsory)
SOFTWARE	Software used to run the simulation (e.g. Gromacs, Amber, NAMD, etc.)	
TRJ	Name of the trajectory file found from DOI	
TPR	Name of the topology file found from DOI (trp file in the case of Gromacs)	
PREEQTIME	Pre-equilibrate time simulated before the uploaded trajectory in nanoseconds. ¹	
TIMELEFTOUT	Equilibration period in the uploaded trajectory that should be discarded in analyses. ²	
COMPOSITION	Molecules names used in the simulation and corresponding mapping files (see section 4.2)	
DIR_WRK	Temporary working directory in your local computer.	
UNITEDATOM_DICT	Information for constructing hydrogens for united atom simulations, empty for all atom simulations	
TYPEOFSYSTEM	Lipid bilayer or something else	
PUBLICATION	Give reference to a publication(s) related to the data.	User given (optional)
AUTHORS_CONTACT	Name and email of the main author(s) of the data.	
SYSTEM	System description on free text format	
SOFTWARE_VERSION	Version of the used software	
FF	Name of the used force field	
FF_SOURCE	Source of the force field parameters, e.g, CHARMM-GUI, webpage, citation to a publication, etc.	
FF_DATE	Date when force field parameters were accessed on the gives source (day/month/year).	
FFmolename	Molecule specific force field information, e.g., water model with FFSOL and sodium parameters with FFSOD.	
CPT	Name of the Gromacs checkpoint file.	
LOG	Name of the Gromacs log file.	
TOP	Name of the Gromacs top file.	automatically extracted data.
GRO	Name of the Gromacs gro file.	
TRAJECTORY_SIZE	Size of the trajectory file in bytes	
TRJLENGTH	Lenght of the trajectory (ps).	
TEMPERATURE	Temperature of the simulation.	
NUMBER_OF_ATOMS	Number of atoms in the simulation.	
DATEOFRUNNIG	Date when added into the databank	
EXPERIMENT	Potentially connected experimental data	
COMPOSITION	Numbers of lipid molecules (NPOPC, NPOPG, etc.) per membrane leaflet are calculated by determining on which side of the center of mass of the membrane the center of mass of the head group of each lipid molecule is located. Numbers of other molecules such as solvent and ions (NSOL, NPOT, NSOD, etc.) are read from the topology file.	

Table 1. Keys stored in the README.yaml files of simulations.

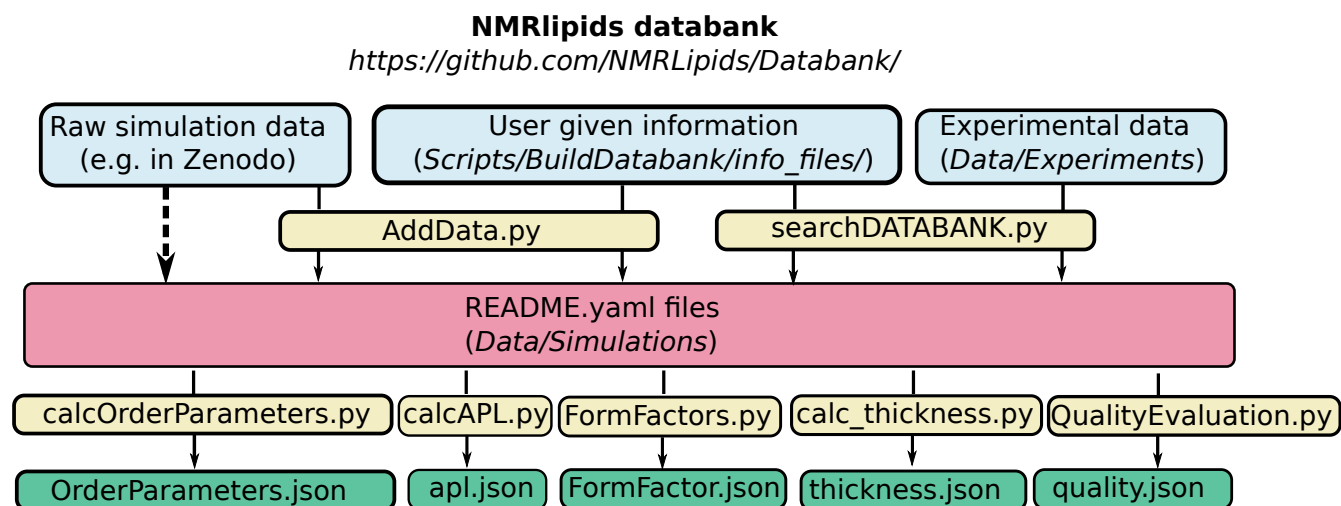


Figure 3. Structure of the NMRLipids databank. Manually added input data (blue boxes) includes basic information on the simulation, permanent links to the raw data, and experimental data if available. The databank entries (red box) and analysis results (green boxes), locating at <https://github.com/NMRLipids/Databank/tree/main/Data/Simulations> are automatically generated by the computer programs included in the NMRLipids databank (yellow boxes). Because raw data are not permanently stored but can be accessed based on the information in the databank, this connection is marked with the dashed line.

key	description
DOI	DOI of the publication related to the experimental data.
TEMPERATURE	Temperature of the experiment.
MOLAR_FRACTIONS	Dictionary of molar fractions of bilayer components
ION_CONCENTRATIONS	Dictionary of ion concentrations of the system
TOTAL_LIPID_CONCENTRATION	Total concentration of lipid components. If exact concentration is not known, but experiments are performed in excess water, 'full hydration' can be given.
COUNTER_IONS	Type of counter ions if present.

Table 2. Keys stored in the README.yaml files of experiments.

In order to evaluate the quality of simulations, sets of C-H bond order parameters from NMR and from factors from x-ray scattering are included in the NMRLipids databank (<https://github.com/NMRLipids/Databank/tree/main/Data/experiments>). The required information for an experimental dataset are listed in table 2. A simulation is connected to a experimental data set if molar concentrations of all molecules are within ± 5 percentage units, charged lipids have the same counterions, and temperature is within ± 2 degrees. In such cases, a simulation and experimental data are paired by adding the path to the experimental data into the simulation README.yaml file.

Because README.yaml contains all the essential information on each simulation, arbitrary analyses can be automatically performed over all the simulations in the NMRLipids databank. In addition to the order parameters for each C-bonds and x-ray scattering form factors used in the quality evaluation, the NMRLipids databank contains the area per lipid and thickness calculated from all simulations in the databank. These results are stored in the same folders as the README.yaml files in <https://github.com/NMRLipids/Databank/tree/main/Data/Simulations>.

4.2 Molecule and atom naming convention

Because universal convention for lipid molecules and atoms therein has not been defined, the naming conventions vary between authors and force fields. To enable automatic analyses over large sets of simulation in the NMRLipids databank, we have defined unique naming conventions for lipid molecules and atoms. The abbreviations of molecule names used in the NMRLipids

¹For example, if you upload 100-200 ns part of total 200 ns simulation, this should value should be 100.

²For example, if you upload 0-200 ns part of total 200 ns simulation where the first 100 ns should be considered as an equilibration, this value should be 100.

databank are listed in table 3. The unique atom names for each molecule and corresponding names in each simulation are defined using mapping files introduced in the NMRLipids project (<https://nmrlipids.blogspot.com/2022/04/new-yaml-format-of-mapping-files.html>).

Molecule and atom names in each simulation are connected to the unique naming convention with the COMPOSITION dictionary in the README.yaml file. Upon addition of a new entry in the databank, the molecule names in the simulation corresponding the unique names and mapping files (available at https://github.com/NMRLipids/Databank/tree/main/Scripts/BuildDatabank/mapping_files) are defined in the dictionary. The numbers of molecules in the system are then automatically calculated by the NMRLipids databank codes (AddData.py in Fig. 3) and stored in the README.yaml together with other content of the COMPOSITION dictionary. This information can be then used to find each molecule and atom from each simulation in the analysis codes.

Abbreviation	Molecule name
POPC	1-palmitoyl-2-oleoyl-sn-glycero-3-phosphocholine
POPG	1-palmitoyl-2-oleoyl-sn-glycero-3-phosphoglycerol
POPS	1-palmitoyl-2-oleoyl-sn-glycero-3-phospho-L-serine
POPE	1-palmitoyl-2-oleoyl-sn-glycero-3-phosphoethanolamine
CHOL	cholesterol
DHDMAB	dihexadecyldimethylammonium
POT	potassium ion
SOD	sodium ion
CLA	chloride ion
CAL	calcium ion
SOL	water

Table 3. Abbreviations used in the databank

4.3 Quality evaluation

The quality of conformational ensembles of individual lipid molecules in simulations are evaluated using the C-H bond order parameters which can be directly compared with robust experimental data²¹. The order parameters are defined as

$$S_{CH} = \frac{1}{2} \langle 3 \cos^2 \theta - 1 \rangle, \quad (1)$$

where θ is the angle between the C-H bond and the membrane normal and angular brackets denote the ensemble average, i.e. average over all sampled configurations of all lipids in a simulation. Because conformational ensembles of individual lipids are independent in a simulation of a fluid lipid bilayer, the number of independent sample points for each C-H bond, n , equals the number of lipids in a simulation. Denoting the S_{CH} from Eq. 1 as the sample mean and s its variance calculated over individual lipids, $\frac{S_{CH} - \mu}{s/\sqrt{n}}$ has a Student's t-distribution with $n - 1$ degrees of freedom, where μ is the real mean of the order parameter. Because lipid bilayer simulations contain at least dozens of lipids, the Student's t-distribution could be safely approximated with a normal distribution. However, when calculating the probability for a simulation to locate within experimental order parameters, the normal distribution gives values below the numerical accuracy of computers for simulations values far from experiments. To avoid such numerical instability, we use the first order Student's t-distribution. Therefore, we estimate the probability for the simulation order parameter to locate within experimental error bars from equation

$$P = f\left(\frac{S_{CH} - (S_{exp} + \Delta S_{exp})}{s/\sqrt{n}}\right) - f\left(\frac{S_{CH} - (S_{exp} - \Delta S_{exp})}{s/\sqrt{n}}\right), \quad (2)$$

where $f(t)$ is the first order Student's t-distribution.

The error of $\Delta S_{exp} = 0.02$ is currently assumed for all experimental order parameters²¹. Because phospholipids sample their conformational ensemble within nanosecond timescale²⁶, all simulations in the databank would be sufficiently long to sample the realistic conformational phase of individual lipids. However, some force fields exhibit too slow dynamics which leads to large error bars in order parameter values²⁷. Because large error bars widen the Student's t-distribution in Eq. 2 thereby artificially increasing the probability to find the simulated value within experimental error bars, the order parameters with error bars larger than the experimental error 0.02 are not included in the quality evaluation.

The qualities of different fragments within each lipid type in a simulation, $P^{frag}[lipid]$, are then estimated by averaging the probabilities for individual C-H bond order parameters to locate within experimental errors according to Eq. 2, and dividing the

average with the percentage of order parameters for which the quality is available within the fragment, $p_{\text{frag}}[\text{lipid}]$,

$$p^{\text{frag}}[\text{lipid}] = \frac{\langle P[\text{lipid}] \rangle_{\text{frag}}}{p_{\text{frag}}[\text{lipid}]}, \quad (3)$$

where frag can be *sn*-1, *sn*-2, headgroup or total (all order parameters within a molecule). The overall quality of different fragments in a simulation are then defined as a molar fraction weighted average over different lipid components

$$P^{\text{frag}} = \sum_{\text{lipid}} \chi_{\text{lipid}} P^{\text{frag}}[\text{lipid}], \quad (4)$$

where χ_{lipid} is the molar fraction of a lipid in the bilayer.

Qualities of form factors were evaluated using the same approach as in SIMtoEXP program²⁸. Because experiments give form factors only in relative intensity scale, they were scaled to the simulation data using equation

$$k_e = \frac{\sum_{i=1}^{N_q} \frac{|F_s(q_i)| |F_e(q_i)|}{(\Delta F_e(q_i))^2}}{\sum_{i=1}^{N_q} \frac{|F_e(q_i)|^2}{(\Delta F_e(q_i))^2}}. \quad (5)$$

The quality of each form factor was then calculated from the equation

$$\chi^2 = \frac{\sqrt{\sum_{i=1}^{N_q} (|F_s(q_i)| - k_e |F_e(q_i)|)^2 / (\Delta F_e(q_i))^2}}{\sqrt{N_q - 1}}, \quad (6)$$

where F_s and F_e are form factors from a simulation and experiment, respectively, and summation goes over the experimentally available N_q points.

4.4 Analysing simulations in the NMRLipids databank

The README.yaml files contain all the essential information to perform arbitrary analyses of simulations in the NMRLipids databank, i.e., the permanent location of the original data and naming convention for all atoms and molecules in each system. In practise, the analyse codes contains a loop over all README.yaml files (i.e., simulations in the NMRLipids databank) which first downloads the raw simulation to a local computer and then uses the information about the atom and molecule naming conventions in README.yaml and mapping files to perform the desired analyses. For example, the code that calculates all C-H bond order parameters of all systems is available at <https://github.com/NMRLipids/Databank/blob/main/Scripts/AnalyzeDatabank/calcOrderParameters.py> and minimal example of a analysis code is available at <https://github.com/NMRLipids/Databank/blob/main/Scripts/AnalyzeDatabank/template.ipynb>.

While the order parameters, form factors, area per lipid and thickness are stored within the NMRLipids databank (<https://github.com/NMRLipids/Databank/tree/main/Data/Simulations>), further analyses can be conveniently stored in separate repositories with the same folder structure based on hash identities of trajectory and topology files. For example, results from further analyses performed here are stored in folders at <https://github.com/NMRLipids/DataBankManuscript/tree/main/Data>. Such organization of the data enables further upcycling of the analyzed data as similarly to the original NMRLipids databank repository.

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Author contributions statement

Must include all authors, identified by initials, for example: A.A. conceived the experiment(s), A.A. and B.A. conducted the experiment(s), C.A. and D.A. analysed the results. All authors reviewed the manuscript.

Additional information