

Towards a molecular level understanding of active sites for catalysis and the interactions with solvents in zirconium based MOFs at operating conditions

Moleculair inzicht in de actieve sites voor katalyse en de interactie met solventen in zirconium gebaseerde Metaal Organische roosters bij operationele condities

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List of Abbreviations

AIMD	<i>Ab Initio</i> Molecular Dynamics
AIM	Atomic layer deposition in MOF
BDC	Benzene-1,4-dicarboxylate
BTC	Benzene-1,3,5-tricarboxylic acid
BO	Born–Oppenheimer
BOMD	Born–Oppenheimer Molecular Dynamics
CMM	Center for Molecular Modeling
CPMD	Car–Parrinello Molecular Dynamics
CN	Coordination Number
CV	Collective Variable
DFT	Density Functional Theory
FES	Free Energy Surface
FFA	Free Fatty Acids
FTIR	Fourier-Transform Infrared
GCMC	Grand Canonical Monte Carlo
GGA	Generalized Gradient Approximation
HF	Hartree–Fock
HKUST	Hong Kong University of Science and Technology
HPC	High-Performance Computing
IR	Infrared
IRMOF	Isoreticular MOF
LDA	Local Density Approximation
LSDA	Local Spin Density Approximation
M–L	Metal–linker coordination
MC	Monte Carlo
MD	Molecular Dynamics
MIL	Matériaux de l’Institut Lavoisier
MOF	Metal–Organic Framework
MPV	Meerwein–Ponndorf–Verley
MTD	Metadynamics
MTK	Martyna–Tobias–Klein
NEB	Nudged elastic band
NU	Northwestern University
PBC	Periodic Boundary Conditions
PCM	Polarizable Continuum Model
PCP	Porous Coordination Polymer
PHVA	Partial Hessian Vibrational Analysis
PES	Potential Energy Surface

PSE	Post–Synthetic Exchange
PSLE	Post–Synthetic Ligand Exchange
PSM	Post–Synthetic Modification
PXRD	Powder X–Ray Diffraction
QM	Quantum Mechanics
RE	Replica Exchange
RDF	Radial Distribution Function
RPA	Random Phase Approximation
SBU	Secondary Building Unit
SCF	Self–Consistent Field
SE	Single Excitation
SXRD	Single–Crystal X–Ray Diffraction
TFA	Trifluoroacetic Acid
TGA	Thermogravimetric Analysis
TPS	Transition Path Sampling
TS	Transition State
TST	Transition State Theory
UiO	Universitetet i Oslo (University of Oslo)
US	Umbrella Sampling
WHAM	Weighted Histogram Analysis Method
XRD	X–Ray Diffraction
ZPE	Zero Point Energy

List of Symbols

\ddagger	Transition state
ΔE^\ddagger	Electronic energy barrier
ΔH^\ddagger	Enthalpy barrier
ΔG^\ddagger	Free energy barrier
B_0	Bulk modulus
E	Electronic energy
E_{ZPE}	Zero-point vibrational energy
F	Helmholtz free energy
G	Gibbs free energy
\hat{H}	Hamiltonian operator
H	Enthalpy
h	Planck constant
k	Rate coefficient unimolecular reaction
k_B	Boltzmann constant
N	Number of particles
N_A	Avogadro constant
N_{dof}	Number of degrees of freedom
p	Pressure
P	Product
Q	Overall partition function
Q_{trans}	Translational partition function
$Q_{rot,ext}$	Rotational partition function
Q_{vib}	Vibrational partition function
q_X	Partition function of species X
R	Reactant
R	Universal gas constant
r_{ij}	Distance between atoms i and j
S	Entropy
T	Temperature
U	Internal energy
V	Volume
V_0	Equilibrium Volume
ε_0	Electronic energy
ν_i	Vibrational frequency
ρ	Density

Samenvatting

Nanoporeuze materialen worden intensief bestudeerd de laatste decennia omwille van hun veelzijdigheid aan toepassingen in tal van sectoren. Dit toepassingspotentieel heeft vooral te maken met hun groot intern oppervlak en porie–volume, waardoor ze uitermate geschikt zijn binnen de heterogene katalyse en adsorptie toepassingen. Een groot scala van nanoporeuze materialen zijn momenteel beschikbaar zoals zeolieten en actieve kool die op grootschalige basis worden gebruikt binnen de petrochemie, geneeskunde en milieutoepassingen.

Recentelijk werden nieuwe generaties van nanoporeuze materialen ontwikkeld waarvan het bouwconcept gebaseerd is op bouwstenen die modulair kunnen worden samengebracht. Tot deze klasse behoren onder meer metaal–organische roosters (MOFs) en covalente organische roosters (COFs). Vooral MOFs zijn op korte tijd geëvolueerd tot één van de meest onderzochte materialen en dit omwille van hun grote variabiliteit in bouwconcept, wat het mogelijk maakt materialen te ontwerpen voor specifieke toepassingen. Hoewel ze initieel vooral werden onderzocht voor adsorptietoepassingen, wordt het toepassingspalet volop uitgebreid de laatste jaren. Op dit moment zijn reeds talrijke succesvolle voorbeelden beschikbaar van MOFs binnen het domein van de heterogene katalyse.

Het uniek bouwconcept berust op het samengaan van anorganische bouwblokken met multitopische organische linkers via coördinatie bindingen. Op die manier worden 1D-, 2D- of 3D- kristallijne netten gevormd. Door het feit dat men kan uitgaan van een groot aantal anorganische bouwblokken die op hun beurt kunnen gecombineerd worden met tal van organische linkers, kunnen een bijzonder groot aantal materialen worden gesynthetiseerd. Inderdaad tot op vandaag werden er meer dan 10000 materialen experimenteel gemaakt. De grote uitdaging bestaat er echter in om de materialen zo te ontwerpen zodat ze optimaal geschikt zijn voor een bepaalde toepassing. Naast deze grote verscheidenheid gegenereerd door de grote combinatoriek van bouwblokken, kunnen de materialen ook nog verder worden gefunctionaliseerd worden door middel van post–synthetische modificaties. Het mag duidelijk zijn dat metaal organische roosters omwille van bovenstaande aspecten bijzonder veel potentieel vertonen voor ontwerp en design naar specifieke toepassingen toe.

In tegenstelling echter tot zeolieten, de welbekende familie van nanoporeuze anorganische materialen die momenteel op grote schaal worden gebruikt binnen tal van industriële toepassingen, zijn ze echter minder stabiel. Dit heeft te maken met de aanwezigheid van de metaal–ligand binding die inherent zwakker is dan de

silicium–oxide binding die aan de basis ligt van zeolieten. Echter zeolieten kunnen niet gefabriceerd worden met een even grote verscheidenheid en lenen zich in die zin minder tot molecuair ontwerp voor specifieke toepassing.

In deze thesis worden metaal organische roosters onderzocht voor toepassingen binnen de katalyse. Zoals reeds gesteld, bieden metaal organische roosters de mogelijkheid om poreuze materialen te ontwerpen met het doel om optimale chemische conversies met gewenste productselectiviteit en opbrengst te bekomen. Naast deze specifieke eigenschappen, behoren ze uiteraard tot het domein van de heterogene katalyse, wat maakt dat ze toelaten meer duurzame processen te ontwerpen gezien inherent product separatie of toxicisch afval minder is in vergelijking met processen waarbij homogene katalysatoren worden gebruikt.

Een groot vraagstuk bestaat erin te begrijpen hoe actieve sites voor katalyse eruit zien en hoe zij kunnen worden gegenereerd en gemoduleerd. Al vrij vlug na de vlug na de initiële synthese van de eerste metaal organische roosters werd duidelijk dat deze materialen inherent een grote mate van wanorde kunnen bezitten en structurele defecten. Initieel werd dit vooral als een nadeel gepercipieerd, echter binnen de huidige wetenschappelijke context wordt de aanwezigheid van defecten en hun mogelijkheid tot modulatie geëxploiteerd voor specifieke toepassingen. Dit is met name belangrijk voor de creatie van actieve sites voor katalytische reacties. Binnen deze thesis zullen ondermeer materialen worden onderzocht die indien perfect opgebouwd enkel volledig gecoördineerde metaalsites bevatten en in die zin open metaal sites ontbreken waaraan katalytische reacties kunnen plaatsvinden. Het structureel inbouwen van defecten biedt de mogelijkheid om de materialen te activeren voor katalytische toepassingen. Verder kunnen defecten er ook voor zorgen dat de poriegrootte wordt aangepast bijvoorbeeld door structureel ontbreken van anorganische bouwbladen in het materiaalskelet. De grote uitdaging bestaat erin om de aanwezigheid van defecten te controleren en ook hun molecuair gedrag te begrijpen.

De experimentele karakterisatie van deze materialen tot op molecuair niveau is bijzonder uitdagend mede door de complexiteit van hun structuur en de aanwezigheid van structurele defecten. De aanwezigheid van wanorde maakt het niet triviaal voor experimentatoren om direct inzicht te verkrijgen in de moleculaire aard van de actieve sites en structuur–eigenschappenrelaties op te stellen. In dit opzicht zijn moleculaire modelleringstudies in nauwe samenwerking met experimentatoren van essentieel belang om het gedrag van MOFs te begrijpen en te voorspellen. Binnen het domein van de moleculaire modellering en de toepassing op nanoporeuze materialen heeft een enorme evolutie plaatsgegrepen de afgelopen tientallen jaren. Met behulp van computationele technieken is het nu mogelijk om MOFs te bestuderen op de nanoschaal. In die zin kunnen computersimulaties aangewend worden om experimentele waarnemingen te verklaren en in sommige gevallen kan gepoogd worden zelfs te voorspellen wat aangewezen materialen zijn voor bepaalde toepassingen of hoe reactieomstandigheden kunnen worden aangepast om een betere selectiviteit, activiteit te bekomen. Men moet zich echter realiseren dat dit laatste een bijzonder grote uitdaging blijft voor huidige modelleringstechnieken, gezien het zeer moeilijk is om theoretische modellen op te

stellen die met voldoende nauwkeurigheid experimentele omstandigheden kunnen nabootsen.

Een groot scala van moleculaire modelleringstechnieken zijn momenteel voorhanden om nanoporeuze materialen te bestuderen. Tal van keuzes dienen gemaakt te worden die een afweging zijn tussen enerzijds de haalbare computationele kost en anderzijds de accuraatheid waarmee het materiaal en het proces wordt beschreven. Idealiter benaderen moleculaire modellen zo goed als mogelijk de experimentele en reële condities. Afhankelijk van welke aspecten men wenst te bestuderen kunnen verschillende benaderingsmethoden worden toegepast. Indien men bijvoorbeeld geïnteresseerd is in fysische eigenschappen zoals fasetransformaties, kan het volstaan om de chemische interacties op benaderende wijze te beschrijven aan de hand van krachtvelden waardoor meer atomen kunnen worden in rekening gebracht worden. Echter voor de katalyse, is het essentieel om de chemische bindingen voldoende accuraat te beschrijven teneinde de actieve site en het reactiemechanisme correct in kaart te brengen. Dergelijke meer accurate methoden die gebaseerd zijn op een expliciete beschrijving van de elektronische structuur, vergen een hogere computationele kost waardoor de modelsystemen in grootte moeten beperkt worden. Concreet betekent dit het aantal atomen van het systeem beperkt blijft tot een paar honderdtal atomen. Een grote uitdaging bestaat erin de topologie van het materiaal correct in kaart te brengen en ook rekening te houden met procescondities zoals een realistische temperatuur, gasdruk of aanwezigheid van solventen in de poriën van het materiaal. De huidige computationele technieken zijn dermate geëvolueerd dat dergelijke beschrijving van een chemische reactie op procescondities binnen het bereik komt. Dit heeft niet alleen te maken met de opkomst van heel krachtige High Performance computers maar ook met de ontwikkeling van ingenieuze theoretische algoritmes die toelaten om grote moleculaire systemen accuraat te berekenen.

Binnen deze thesis werden zirconium materialen bestudeerd voor katalytische toepassingen. Meer in het bijzonder werd een uitgebreid onderzoek verricht van de actieve binnen het zeer stabiel UiO-66 materiaal. Er werd specifiek geopteerd om dit materiaal in detail te onderzoeken, gezien het een van de meest onderzochte materialen is voor verscheidene toepassingen en dit omwille van de zeer hoge stabiliteit binnen de familie van de metaal organische roosters. De grote stabiliteit vindt zijn oorsprong in de opbouw van het materiaal, namelijk zirconium oxide clusters worden geconnecteerd met tereftalaat linkers en dit met een heel hoge graad van connectiviteit. Elke anorganische bouwblok wordt geconnecteerd met 12 organische linkers in het defectvrije materiaal. Dit maakt dat het materiaal bijzonder robuust is in verscheidene chemische omstandigheden en thermische condities, waardoor het volop kan worden ingezet voor tal van katalytische toepassingen. Het defectvrije materiaal bevat geen open metaal sites en is in die zin niet geschikt voor katalytische toepassingen. Lange tijd was het onduidelijk hoe de actieve sites werden gecreëerd binnen dit materiaal. Door middel van geavanceerd fundamenteel wetenschappelijk experimenteel en theoretisch onderzoek, is het duidelijk geworden dat de structurele aanwezigheid van defecten aan de basis ligt van actieve sites voor katalyse. Zo kunnen ondermeer linkers ontbreken, maar aangezien het materiaal inherent een bijzonder hoge graad van connectiviteit vertoont, brengt

dit de stabiliteit niet in het gedrang. Dit materiaal met zijn veelzijdigheid van eigenschappen en mogelijkheid tot modulatie vormt een ideaal platform voor dit doctoraatsonderzoek.

Binnen dit doctoraatsonderzoek werden actieve sites voor katalyse in het UiO–66 materiaal bestudeerd met een groot scala van computationele technieken teneinde een begrip te krijgen van het reactief gedrag op procescondities. Er werd zowel gebruik gemaakt van statische technieken waarbij slechts enkele punten op het potentieel energie oppervlak werden gelokaliseerd als moleculaire dynamica technieken waarbij het gedrag in functie van de tijd op reële temperaturen en in reële omstandigheden van reactanten in aanwezigheid van solventen werd gevolgd. Het potentieel energie oppervlak werd systematisch beschreven met behulp van dichtheidsfunctionaaltheorie. Hierdoor wordt de elektronische structuur van het materiaal expliciet beschreven met een haalbare computationele kost. Initieel werd voor sommige reacties gebruik gemaakt van uitgebreide clustermodellen om een eerste inzicht te krijgen in de lokale omgeving van de katalytisch actieve site. Nadien werd systematisch overgestapt op moleculaire modellen waarbij de topologie van het materiaal met inbegrip van de periodieke randvoorwaarden correct in kaart werd gebracht. Het computationeel werk werd in grote mate verricht in samenwerking met verscheidene experimentele groepen. Verder werd ook samengewerkt met internationale theoretische partners om de kennis aangaande gevanceerde moleculaire dynamica technieken te versterken.

Als startpunt van het doctoraatsonderzoek werd de Fischer esterificatie bestudeerd, welke een Lewis zuur gekatalyseerde reactie is en waarvoor een gunstig effect werd waargenomen in het experiment door de aanwezigheid van water. Initieel was het onduidelijk wat het effect was van water op de actieve site. Dankzij het toepassen van moleculaire modelleringstechnieken, werd duidelijk dat de actieve site niet alleen bestaat uit een Lewis zure site maar dat ook de aanwezigheid van Brønsted zure sites essentieel is voor optimaal functioneren van de actieve site. De aanwezigheid van waterige solventen heeft een gunstige invloed op de reactie aangezien het de mobiliteit van protonen faciliteert, waardoor een soort van dynamische aciditeit van het materiaal wordt gegenereerd.

Startende van deze specifieke case studie, werd het duidelijk dat het gebruik van statische methoden, niet altijd voldoende is voor het bestuderen van katalytische reacties in deze materialen op procescondities, gezien in de realiteit het materiaal en de actieve site een sterk dynamisch gedrag kunnen vertonen. In groot deel van de thesis werd derhalve gebruik gemaakt van meer complexere modellen waarbij het solvent dat opgesloten is in de poriën van het materiaal explicet in rekening wordt gebracht op realistische experimentele condities. Door het gebruik van dergelijke gevanceerde moleculaire modelleringstechnieken werd duidelijk dat protische solventen het gedrag van het materiaal in grote mate kunnen moduleren door continu te interacteren met de onder gecoördineerde zirconium clusters. De aanwezigheid van een dergelijke solvent laat verder toe om geladen intermediairen te stabiliseren. Een onderbouwd inzicht werd bekomen in de manier hoe defecten in deze materialen kunnen worden gevormd en gemoduleerd. Door toepassing van gevanceerd moleculaire dynamicatechnieken, werd het duidelijk

dat de organische linkers een zekere graad van mobiliteit kunnen vertonen. Zij kunnen tijdelijk loskomen van de anorganische cluster, connecteren met een andere deel van de anorganisch bouwblok en dit terwijl de structurele integriteit van het materiaal behouden blijft. Dergelijke dynamisch gedrag van linkers blijkt tevens aan de basis te liggen van de postsynthetische modificatie van het UiO–66 materiaal. In samenwerking met experimentele partners werd dit proces nader bekeken. Een uniek inzicht werd bekomen in de moleculaire processen die aan de grondslag liggen van postsynthetische modificatieroutes.

In een volgende fase werd nagegaan in hoeverre het waargenomen dynamisch gedrag van het UiO–66 materiaal met hoge connectiviteit, kon worden veralgemeend naar andere zirconium gebaseerde materialen. In dit opzicht werd het MOF-808 materiaal onderzocht, wat opgebouwd is uit gelijkaardige zirconium oxide clusters maar waarbij de connectiviteit met organische linkers lager is. Het materiaal bezit een groot porievolume waardoor het actief wordt onderzocht binnen het domein van de katalyse. Het werd duidelijk dat het waargenomen gedrag voor UiO–66 niet rechtstreeks kan worden doorgetrokken naar andere zirconium gebaseerde materialen. De lagere connectiviteit zorgt voor een aantal fundamentele verschillen met betrekking tot stabiliteit. Verdere studies in dit domein naar andere potentieel interessante zirconium materialen bieden een interessant perspectief voor toekomstige katalytische toepassingen.

Samenvattend, werden in deze thesis een groot scala van computationele technieken aangewend om de eigenschappen van actieve sites voor katalyse te begrijpen op moleculaire schaal in zirconium gebaseerde materialen. Gezien in deze hoog geconnecteerde materialen de actieve sites worden gegeneerd door de creatie van structurele defecten, is het belang van solventen in de poriën van het materiaal van uitermate belang. Dit alles echter levert een nanogestrukteerd platform voor katalyse dat bijzonder complex is maar ook een groot scala van mogelijkheid biedt voor verder exploitatie van katalytische reacties. De modelleringstechnieken die nodig zijn om dergelijk complex systeem te modelleren zijn zeer geavanceerd en inzicht kan enkel bekomen worden door complementaire technieken in te zetten. De bekomen resultaten benadrukken het belang van geavanceerde modelleringstechnieken om het katalytisch gedrag van dergelijke complexe materialen te ontrafelen op moleculaire schaal.

Summary

In the past decades, nanoporous materials have become an intense field of study due to their numerous potential applications. This has to do mainly to their extremely high surface area and pore volume, that make them suitable in heterogeneous catalysts and adsorption applications. At present time, a wide range of nanoporous materials are available, such as zeolites and activated carbon, widely used in petrochemistry, medicine and environmental applications.

Recently, new generations nanoporous materials have been explored which rely on the concept of reticular chemistry and building blocks, such as Metal Organic Frameworks (MOFs) and Covalent Organic Frameworks (COFs). In particular, MOFs have rapidly become one of the most deeply investigated classes of materials due to their great variability in design, that allows the synthesis of materials for specific applications. Although they were initially investigated for applications as adsorbants, their range of applications has been enormously expanded in recent years. Currently, many successful examples of MOFs are available in the field of heterogeneous catalysis.

The unique construction concept of MOFs is based on the combination of inorganic building blocks with multtopic organic linkers via coordination bonds. This way, 1-, 2- or 3-D crystalline structures can be formed. A large number of inorganic building blocks can be combined with numerous organic linkers, allowing to construct a plethora of different materials. Indeed, up to today more than 10,000 MOFs have been synthesized experimentally. The big challenge lies in designing material that are optimally suited for a specific application. In addition to the large combination of building blocks, the materials can be further functionalized through post-synthetic modifications. Because of the above aspects, MOFs have a great potential to be designed to target a specific application.

However, in contrast to zeolites, the well-known family of inorganic nanoporous materials that are widely used in many industrial applications, MOFs are less stable. This is due to the presence of metal-ligand bonds that are inherently weaker than the Si-O bonds that lie at the basis of zeolites. However, zeolites cannot be manufactured with an equally large variety of structures and in this sense are less suitable for molecular design to target specific applications.

In this thesis, MOFs have been investigated for applications within catalysis. As already stated, MOFs offer the possibility to design porous materials to maximize the outcome of a given reaction, in terms of yield and selectivity for a desired prod-

uct. Moreover, as heterogeneous catalysts, they can bring numerous advantages to industry. As opposed to homogeneous catalysts, they offer the possibility to design more sustainable processes, with less toxic waste and easy product separation.

A major issue lies in the understanding of how active sites for catalysis look like and how they can be generated and modulated. Soon after the synthesis of the first MOFs, it became clear that these materials inherently possess a high degree of disorder and structural defects. If at first this was seen as a drawback, within the current scientific context it has become clear that the presence of defects and their modulation can be exploited for specific applications. This is particularly important for the creation of active sites for catalytic reactions. Some MOFs, if perfectly constructed only contain fully coordinated metal sites and in this sense are missing open metal sites where reactions can take place. The structural incorporation of defects offers the possibility to activate the materials for catalytic applications. Furthermore, defects can also increase the pore size, for instance if inorganic building blocks are missing. The big challenge lies in investigating the presence of defects and understanding their molecular behavior.

The experimental characterization of these materials at the molecular level is particularly challenging due to their structural complexity and the presence of structural defects. The presence of disorder makes it not trivial for experimentalists to have direct insight on the molecular nature of the active sites and to understand structure–property relationships. In this sense, modeling studies in close synergy with experimental research are of utmost importance to understand and predict MOF behavior.

The past decades have seen an enormous evolution within the domain of molecular modeling and its application to nanoporous materials. Using computational techniques, it is now possible to study MOFs at the nanoscale. In this sense, computer simulations can be used to explain experimental observations and in some cases even attempt to predict what are the appropriate materials for certain applications, or how reaction conditions can be adjusted to achieve better selectivity and activity. It must be realized, however, that the latter remains a major challenge for current modeling techniques, since it is very difficult to develop theoretical models that can simulate experimental conditions with sufficient accuracy.

Nowadays, a rich toolbox of molecular modeling techniques is available for the study of nanoporous materials which can be applied to MOFs. When studying such complex materials, there is always a trade-off between computational cost and the accuracy in the description of processes and material. Ideally, molecular models describe experimental conditions as closely as possible. Depending on which aspects one wants to focus on, different approaches can be used. For physical properties such as phase transformations, it may suffice to describe the chemical interactions approximately using force fields, so that more atoms can be taken into account in the model. However for catalysis it is essential to accurately describe the chemical bonds to correctly map active sites and reaction mechanisms. Such accurate methods, which are based on an explicit description of the electronic structure, are computationally more expensive, and the model systems must be

limited in size. In concrete terms, this means that the number of atoms in the system is limited to a few hundred atoms. It poses a major challenge to correctly map the topology of the material and at the same time take into account operating conditions, such as realistic pressure, temperature and presence of guest molecules. Current computational techniques have evolved to such an extent that a description of chemical reactions at operating conditions is now within reach. This is not only due to the growth in computational power of High Performance Computing facilities, but also to the development of ingenious theoretical algorithms that allow to calculate large molecular systems accurately.

Within this thesis, zirconium MOFs were examined for catalytic applications. More in particular, an extensive study was conducted on the active sites within the extremely stable UiO–66 material. UiO–66 was specifically selected as it is one of the most investigated MOFs for various applications, and because of its high structural stability within the MOF family. The exceptional stability originates from the structure of the material, in which zirconium–oxide bricks are bridged by terephthalate linkers with a high structural connectivity. In the defect-free material, each inorganic brick is connected to 12 linkers. This makes the material extremely robust and able to withstand the chemical and thermal conditions of numerous catalytic applications. The defect-free UiO–66 does not contain open metal sites and is therefore not suitable for catalytic applications. For a long time it was not understood how the active sites were created within this material. By means of advanced experimental and theoretical research, it has become clear that the presence of structural defects is responsible for the creation of active sites for catalysis. The highly connected material allows for the presence of crystal defects such as missing linkers, without compromising its stability. This material, with its versatility of properties and possibility of modulation forms an ideal platform for this doctoral research.

Within this doctoral research, the active sites in the UiO–66 material were studied with different advanced modeling techniques in order to gain an understanding of the reactive behavior at operating conditions. Static techniques were used, where only a few points on the potential energy surface are investigated, as well as molecular dynamics techniques, where the behavior was followed at realistic temperatures and operating conditions in presence of solvents. The potential energy surface was systematically described using density functional theory, which allows explicitly describe the electronic structure of the material with a feasible computational cost. For some reactions, extended cluster models were initially used to gain insight into the local environment of the active site. We then made use of periodic boundary conditions to correctly map the topology of the material. The computational work was largely carried out in collaboration with various experimental groups. Furthermore, there was also collaboration with international theoretical partners to strengthen the knowledge on advanced molecular dynamics techniques.

As a starting point for the doctoral research, the Lewis–catalyzed Fischer esterification was studied, for which a beneficial effect of the presence of water was observed experimentally. Initially, it was unclear what the effect of water was

on the active site. By use of molecular modeling techniques, it became clear that the active sites are not only composed of Lewis acid sites, but also Brønsted sites are needed for the optimal functioning of the catalyst in this reaction. The presence of aqueous solvent has a beneficial effect on the reaction, facilitating the mobility of protons and generating a sort of dynamic acidity in the material.

Starting from this specific case study, it became clear that static methods are not always sufficient to study the active sites in these materials at operating conditions, since the material can exhibit strong dynamic behavior. In the rest of this thesis, therefore, more complex models were used in which the solvent confined in the pores of the material is explicitly taken into account at realistic experimental conditions. By means of such advanced molecular modeling techniques, it became clear that protic solvents can modulate the behavior of the material to a large extent by continuously interacting with the under coordinated zirconium clusters. The presence of such a solvent also makes it possible to stabilize charged reaction intermediates. A substantial insight was obtained into how defects in these materials can be formed and modulated. By applying advanced molecular dynamics techniques, it became clear that the organic linkers can show a certain degree of mobility. They can temporarily decoordinate from the inorganic cluster and connect with another part of the inorganic building block, while maintaining the structural integrity of the material. Such dynamic behavior of linkers seems to lie at the basis of the post-synthetic modification of the UiO-66 material. This process was further examined in collaboration with experimental partners. A unique insight was obtained into the molecular processes that form the basis of post-synthetic modification routes.

In a subsequent phase, it was investigated to what extent the observed dynamic behavior of the UiO-66 material, with high connectivity, could be generalized to other zirconium-based materials. In this respect, the MOF-808 material was investigated, composed by similar zirconium–oxide clusters, but with lower structural connectivity. The material has a large pore volume and for this reason it is actively investigated in catalysis. It became clear that the behavior observed for UiO-66 cannot be directly extended to other zirconium- based materials. The lower connectivity provides fundamental differences with regard to stability. Further studies in this domain into other potentially interesting zirconium materials offer an interesting perspective for future catalytic applications.

In summary, a rich computational toolbox was used in this thesis to gain molecular understanding on the properties of active sites for catalysis in zirconium based material. As the active sites in these highly connected materials are generated by creation of structural defects, the role of solvent confined in the pores is extremely important. This provides a platform for catalysis that is particularly complex, but also offers many possibilities for further exploitation of catalytic reactions. The modeling techniques required to model such a complex system are very advanced and insight can only be obtained by using complementary techniques. The results obtained emphasize the importance of advanced modeling techniques to unravel the catalytic behavior of such complex materials at the molecular scale.

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Part I

**Towards a molecular level
understanding of active sites
for catalysis and the
interactions with solvents in
zirconium based MOFs at
operating conditions**

1

Introduction

Catalysis, from Ancient Greek *κατά* (katá = “down”) and *λύω* (lúō = “loosen”), is at the heart of almost every industrially relevant chemical process. A catalyst intervenes in the reaction, allowing chemical species to come in contact and react with a specific mechanism, without being consumed. Outstanding catalysts do not only increase the rate of a given reaction, enabling processes that would not happen spontaneously, but can even be selective towards specific end products. Catalysts can be divided in homogeneous and heterogeneous, according to the phase where they are located with respect to the reactants. Homogeneous catalysts share the same phase with the reactants, and can be for instance molecules dissolved in a solvent, such as with organometallic compounds. These type of catalysts are used in many industrial processes, but have the drawback of being difficult to separate from the products, and in many cases separation is the most costly step in the catalytic cycle. For instance, in the case of biodiesel production, the recovery and purification of the products accounts for 60-80% of the whole cost [1].

Heterogeneous catalysts, on the other hand, are located in a different phase with respect to the reactants, and for this reason they have the advantage of being easily separable from the reaction products. Especially green chemistry can widely benefit from heterogeneous catalysts, lowering the production costs and providing a viable alternative to petroleum [2]. Even if the name heterogeneous does not refer to a specific phase, industrially relevant heterogeneous catalysts are often in the solid phase. In order to exert their function, these materials must possess specific

active sites that are easy to reach for the reactants, where these can adsorb, react, desorb, and ultimately diffuse back in the bulk to leave space for a new cycle. For these processes to occur, the number of active sites and the area of contact between the two phases (or surface area) must be sufficiently high, and reactants and products must be able to easily diffuse in the material.

All the above mentioned properties can be found in nanoporous materials, which possess pores with a diameter of < 100 nm, and are extensively used in the field of heterogeneous catalysis. According to the classification by the International Union of Pure and Applied Chemistry (IUPAC) [3], nanoporous materials can be differentiated on the basis of their pore size into microporous (pore size < 2 nm), mesoporous (pore size between 2 and 50 nm) and macroporous (pore size > 50 nm). Their pore structure provides them with an exceptionally high surface area, that for some materials can reach values larger than $7000\text{ m}^2/\text{g}$ [4, 5]. This facilitates diffusion of reactants inside the material, and allows shape selectivity to give specific products. Well-known classes of nanoporous materials include zeolites and metal organic frameworks (MOFs) [6]. Zeolites were one of first heterogeneous catalysts to be industrially exploited, and are the workhorses of petrochemistry, among other applications. They are extremely robust due to their inorganic nature and can withstand harsh reaction conditions. In contrast, MOFs are novel hybrid materials that are less robust, but extremely versatile. They possess even higher porosity, diverse composition, high metal content and tunable organic functions, and for these reasons they have a high catalytic potential.

In general, the study of solid heterogeneous catalysts is not trivial, as the exact concentration and nature of the active sites is often unknown. It is not always clear, in fact, where exactly the active sites may be located in the material, and how they interact with molecules. Moreover, these materials are far from perfect, and are often the defects which are tremendously important to induce the catalytic properties. In this sense, molecular modeling offers a complementary platform, not only to understand the catalytic function, but also to determine structure–activity relationships and to design structures to target specific applications [7].

1.1 Metal Organic Frameworks

MOFs are one of the most intriguing class of materials of current science. These materials, first called 'porous coordination polymers' (PCPs) were discovered in the late 50's, but only at the end of the past century with the works of Robson [8, 9], Kitagawa [10, 11], Yaghi [12] and Férey [13], the scientific community started to understand their full potential. At first the main focus of MOF research was in the discovery of new structures. However, in the last few decades, the field has seen an incredible explosion in scientific and industrial interest, with new applications being continuously explored [14]. MOFs are hybrid nanoporous materials that are composed by metal or metal–oxo clusters connected by multtopic organic

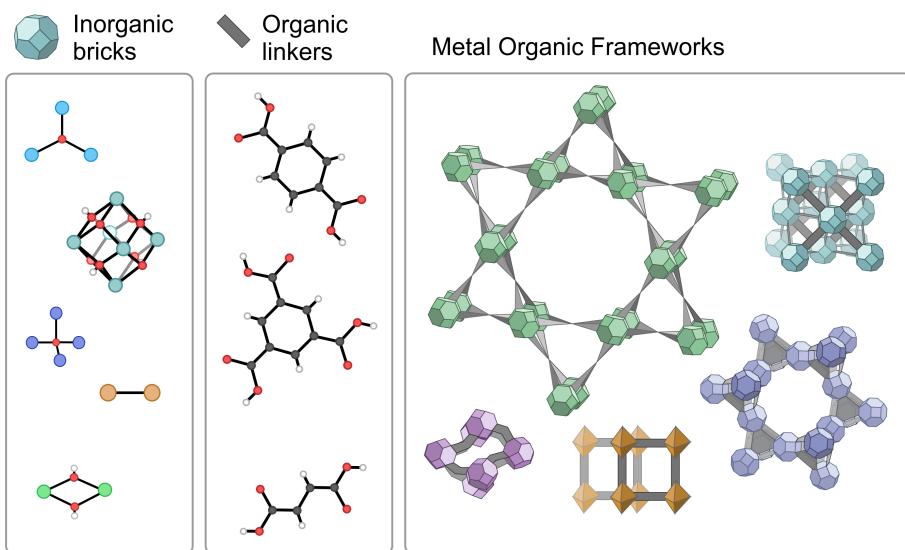


Figure 1.1: Schematic representation of the building block design in Metal-Organic Frameworks (MOFs). Left panel: some examples of inorganic bricks; middle panel: some examples of organic linkers; right panel: some MOFs with different topologies, pore size and shape

linkers, to form multidimensional porous structures. Compared to the already well-known zeolites, MOFs can be constructed without templating agents, with a far greater number of metals and with an exceptional structural diversity. In fact, their particular building block design (Figure 1.1), that makes use of secondary building units (SBUs), allows the creation of an almost infinite number of crystalline structures with different topology and chemical composition.

The coordination bonds which lie at the basis of MOF structures are weaker than in other heterogeneous catalyst such as zeolites. This on the one hand makes them less robust, but on the other allows facile structural modifications which are impossible to do on zeolites. In principle, the nature of the SBUs and their association can be finetuned [15], allowing control of properties such as pore shape and size, functionalization, or surface area. The response to chemical and physical stimuli can this way be easily modulated [16, 17]. One of the most intriguing concepts is the isoreticular synthesis, by which inorganic or organic SBUs can be replaced by topologically identical (or similar) building blocks. This way, starting from a given MOF precursor, whole families of MOF materials can be synthesized spanning a range of pore size and functionality. For instance, the pore size can be significantly increased up to the mesoporous range by using longer isoreticular linkers, such as in the IRMOF series, based on the MOF-5 precursor [18].

The tunability of MOF structures, along with their high crystallinity, metal

content and porosity, are very appealing for application in different industrially relevant fields, such as catalysis, gas storage and separation, drug delivery or sensing, as displayed in Figure 1.2. More specific applications are being further explored, such as warfare agents decomposition, magnetic applications, or membrane separation [14]. At present moment, the main limitation for large scale application lies in the structural stability.

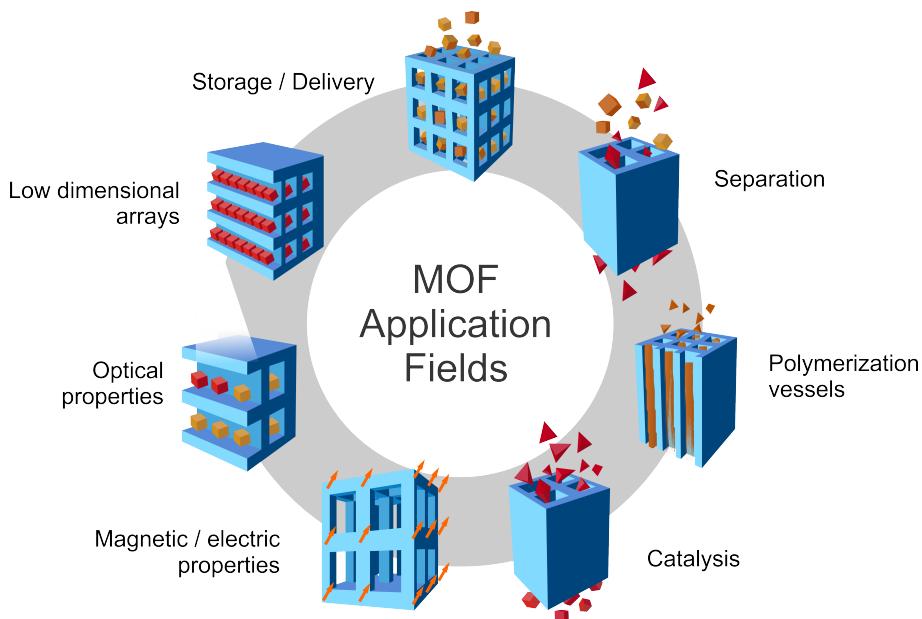


Figure 1.2: Schematic representation of some of the numerous MOF applications.

The study of MOFs has seen a continuous evolution in the past decades. Following the nomenclature proposed by Kitagawa, we can differentiate between three generations of MOFs [19]. First generation MOFs are defined as having a guest–molecules supported pore system that collapses when these are evacuated. For this reason, this first generation of MOFs found very limited use for practical applications. Those of second generation are more robust and have permanent porosity, that is retained even in absence of guest molecules. These materials show high potential for catalysis and other applications and are the main object of this dissertation. Finally, third generation MOFs are characterized by flexible pores that can reversibly change shape with the presence of guest molecules, or upon certain stimuli, such as temperature or pressure. Matsuka *et al.* [20] identified five types of response mechanisms in MOFs, denoted as shrinking, expanding, reshaping, swelling and gate opening or closing. A perspective on the types of stimuli that can induce responsive in MOFs has been reported in the work of Coudert [21].

Defects and active sites

MOFs are crystalline materials that possess structural disorder, such as vacancies. If at first this was seen as a drawback, limiting stability, it has been accepted that they can play a key role in the performance of the material. Defect-containing MOFs and defect engineering have become an active field in MOF research, representing an additional way to finetune and enhance the material properties. Following the classification proposed by Sholl *et al.* [22], defective sites in MOFs can be either point vacancies (such as missing linkers or clusters) or extended ones. Fang *et al.* [23] further divided extended defects into dislocations, planar defects, and micro- and mesoscale volume defects.

At low defect concentration, a random distribution of isolated point defects can be found, whereas at higher concentrations, clustering could occur [24] if the presence of a vacancy is influenced by other vacancies in proximity. The complexity arising from structural disorder makes the study of defects a current challenge in MOF research. In order to control their effect on certain properties, it is important to have precise information on their location, type and dispersion. Common experimental strategies for the characterization of defects in MOFs involve electron and fluorescence microscopy, Raman, infrared and X-ray spectroscopy, powder and single-crystal X-ray diffraction [23]. Complementary to experiment, molecular modeling provides an important tool to obtain atomic scale resolution on the defect sites and understand how they impact specific properties.

Particularly for catalysis and adsorption, vacancies in the material lead to two main effects: 1) an increase in porosity and mass transport, and 2) the presence of undercoordinated metal atoms, introducing highly desired Lewis acid sites. In this sense, provided the structures are stable at reaction conditions (i.e. no leaking is observed), MOFs are true single-site heterogeneous catalysts, that contain well-defined active sites which are inherent part of the framework [25]. We particularly refer to the importance of catalysis in MOF research, shown by the number of publications on the topic (Figure 1.3). According to the classification proposed by Rogge *et al.* [26], we can distinguish between three types of single-site catalysts in nanoporous materials, in which active sites for catalysis can arise from coordinatively unsaturated metals (type I), metal atoms embedded in porphyrin-based ligands (type II) or reactive functional groups (type III).

Coordinatively unsaturated metals are Lewis acid sites that can be present in the framework such as in HKUST-1 [27], or can be introduced by intentional creation of defects, such as in the stable UiO-66 MOF [28]. Another procedure that can lead to open metal sites without generating vacancies is the dehydration upon thermal treatment. To date, a plethora of MOFs have been synthesized containing open metal sites that possess different Lewis acidity. Moreover, their acidic properties can further be tuned by functionalization of linkers and bricks.

This way, it is in principle possible to engineer a material for a target catalytic application.

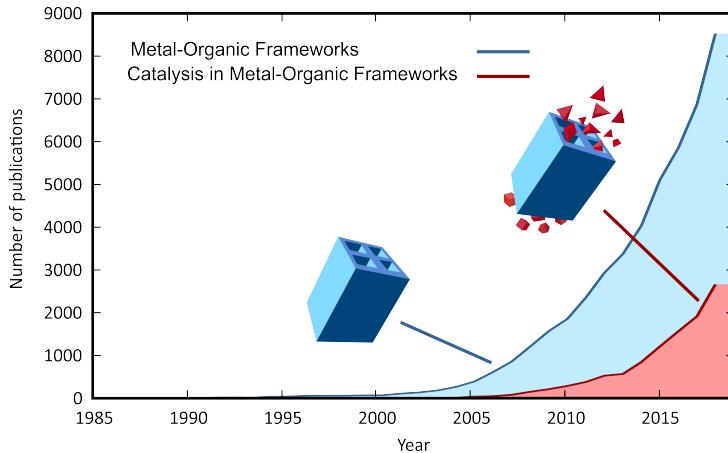


Figure 1.3: Number of publications on MOFs and catalysis in MOFs in years 1985-2018, showing the importance of catalysis in MOFs as indicated by Web of Science

Post-synthetic modification

When it is not possible to introduce functionalities with direct synthesis, post-synthetic modification (PSM) [29, 30] has become a well established procedure that allows the preparation of MOF materials with specific chemical composition. Via this technique, it is possible to modify the crystal after the synthesis, allowing to finetune the properties of the material. Moreover, multi-functional sites can be integrated at the same time in the crystals [31]. PSM strategies include encapsulation of guest molecules or nanoparticles in the pores, modifications of the linkers without breaking the metal-ligand bond, or post synthetic exchange (PSE) of linkers and metals, where building blocks are dynamically exchanged. PSE can also involve terminal ligands, or modulators, that do not contribute in connecting different bricks. This way, building blocks can be exchanged in a dynamic way. Moreover, they can also be eliminated from the framework to create vacancies, provided the material retains its crystallinity. For this reason, PSM has been used as an efficient strategy to introduce defective sites in MOFs. As one may expect, sufficient structural stability under the PSM conditions is a necessary prerequisite in order to apply PSM techniques [32].

Stability

In general, to function at operating conditions and to withstand modifications, materials need to retain their mechanical, thermal and chemical stability at those

conditions [33]. For instance, mechanical stability is needed when compressing MOF in pellets or other shapes for industrial processes [34]. Catalysis often also requires thermal stability, as the materials must be able to resist harsher conditions for certain processes such as in petrochemistry. Finally, chemical stability is crucial for many applications, such as drug delivery, molecular separation, or catalysis [35].

Unfortunately, the M–L coordination bond that makes MOFs so tunable is also regarded as one of their main drawbacks [36–38], as it is responsible for the lower structural stability when compared to already established nanoporous catalysts such as zeolites. For example, the first synthesized MOFs such as Cu_2^+ trimesate HKUST-1, or MOF-5, composed by Zn_2^+ clusters and BDC linkers were degraded by water even at mild conditions [39–42]. Moreover, mechanical, thermal and chemical stabilities can further decrease in the presence of defects. Very few MOFs show stability towards water and at different pH conditions [43], in particular the family of Zr-based MOFs.

Zr-based MOFs

Recently, a class of outstandingly robust MOFs have been synthesized [44]. Zr-based MOFs [45] which exploit the robustness of the Zr–O bond, show an unprecedented stability and are at present time one of the most studied classes of MOFs (Figure 1.4). Moreover, zirconium is an ubiquitous metal that is present in biological systems and has low toxicity, as well as limited cost. This makes Zr-based MOFs particularly promising for applications in catalysis, gas sorption, and drug delivery. An overview of the plethora of possible structures that can be synthesized with different inorganic SBUs can be found in a recent review by Bai *et al.* [45].

The vast majority of these materials is characterized by Zr(IV) atoms in a high coordination state (Figure 1.4), which interact strongly with the oxygens of carboxylate linkers of various topology. These MOFs are characterized by a $\text{Zr}_6\text{O}_4(\text{OH})_4$ cluster in which each of the 6 zirconium atoms is connected to 4 oxygen atoms (two μ_3 -OH and two μ_3 -O), each of which is in turn connected to three zirconium atoms, forming a polyhedron. Each zirconium atom can form 4 other bonds with ligands, accommodating up to 24 metal–ligand bonds per cluster. Every zirconium atom can therefore form a total of 8 coordination bonds oriented in a square–antiprismatic geometry, yielding a rich range of possible structures that can be synthesized with connectivity ranging from 12, as in UiO-66 [46], to 6 which is found in MOF-808 [44] (see Figure 1.4). The dual Lewis acid/base nature of the Zr–carboxylate bonds, along with the high metal oxidation state, gives rise to strong interactions between the SBUs, thus allowing processes such as PSM without compromising the stability of the structures. The high degree of connectivity between inorganic and organic SBUs lies at the origin of the high structural stability found in Zr-based MOFs [43, 45]. Open metal sites in these

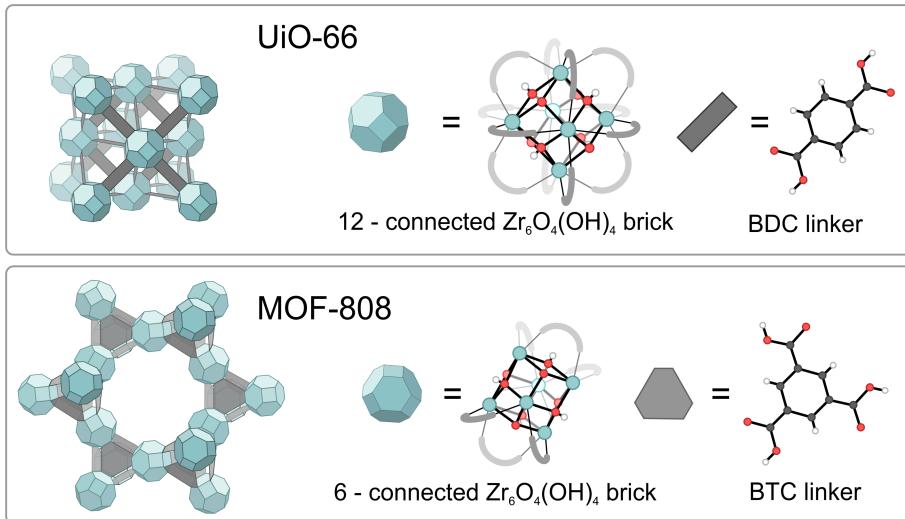


Figure 1.4: Structure of **UiO-66** (top) and **MOF-808** (bottom), the two Zr-MOFs investigated in this work of thesis.

materials can be present if the brick connectivity is lower than 12. This can be an inherent property of the structure, such as in the 8-fold connected NU-1000 [47], or can be induced by introduction of structural defects. Zirconium atoms that remain undercoordinated are Lewis acid sites where reactants can adsorb and that can function as catalytic centers.

UiO-66

The first material belonging to this class of Zr-based MOFs is the Zr–therephthalate based **UiO-66**, and was first synthesized at the Universiteit i Oslo (UiO) by Lillerud and coworkers [46]. The **UiO-66** material, shown in Figure 1.4 (top), is one of the most studied MOFs thus far and was an inspiration of other MOFs within this family. **UiO-66** is characterized by an extremely high connectivity that gives rise to an exceptional structural stability. In this material, each Zr_6 SBU is connected to 12 terephthalate (or benzenedicarboxylate (BDC)) linkers forming a cubic close packed structure with a space group $\text{F}\bar{m}3\text{m}$, No. 225. In this structure there are two different cavities of octahedral and tetrahedral shape, with window sizes of 10 Å and 25 Å, respectively. Each octahedral cage shares triangular windows with eight tetrahedral cages. This results in an extremely robust material which is stable up to 648 K and in a broad range of protic and aprotic solvents and pH conditions. Moreover, the $\text{Zr}_6\text{O}_4(\text{OH})_4$ bricks can be reversibly dehydrated upon thermal treatment at temperatures between 523 and 573 K. Up to two water molecules can be formed this way, yielding a Zr_6O_6 brick, where the zirconium atoms have a coordination of 7 [48]. Lowering of zirconium coordination by

dehydration can be a strategy to create open metal sites, along with the creation of defects.

A whole family of isoreticular MOFs can be derived from UiO–66 by using linkers of different size, spanning from fumaric acid [49] up to terphenyldicarboxylic acid [50], allowing to significantly tune pore size and surface area. Interpenetrated MOFs with UiO–66 topology have been also reported if longer linkers are used [51], however with a decrease in surface area. Moreover, different functional groups can be appended to the phenyl rings, such as bromo, amino, nitro, or naphthalene. Garibay and Cohen showed that UiO–66–NH₂ can be further modified to yield new functionalized frameworks [32]. Also the inorganic SBUs can be modified, for instance introducing titanium or hafnium [52]. Moreover, PSM can lead to creation of defects and open metal sites, as will be shown in **PAPER IV**. The exceptional thermal and chemical stability of UiO–66, along with its high connectivity, allows all these modifications of the structure, and for this reason, this material is often considered as a perfect MOF archetype, where new techniques can be tested.

Defects on UiO–66

The high connectivity of the UiO–66 material allows the presence of structural vacancies. It has been generally accepted that the material can contain defects in the form of missing linkers or clusters (Figure 1.5). These defects naturally originate during synthesis, as shown by experimental results, such as symmetry-forbidden reflections in the PXRD pattern, metal-linker ratio obtained by thermogravimetric analysis (TGA), higher than expected surface area, appearance of O–H stretching bands in the FTIR spectrum etc. [48,53]. Moreover, it was later discovered that the number of defective sites can easily be tuned by adapting the synthesis conditions, such as temperature and type of modulator [54, 55].

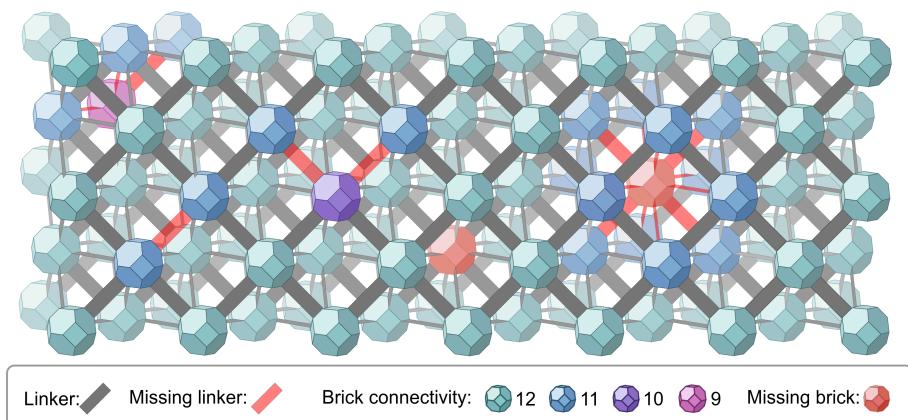


Figure 1.5: Representation of the UiO–66 material with missing linkers and clusters displayed in red. In this dissertation, we mainly focus on missing linkers.

Defects can influence the material properties to a large extent. The beneficial role of defects in UiO–66 has been explored in many applications such as gas storage and separation [54,56], sensing [57], drug delivery [58] and catalysis [26,59]. The physical properties of defective UiO–66 differ according to the number of defects and their location, as has been extensively studied both theoretically and experimentally [24, 60–62]. A decrease in the connectivity in the structure will naturally lead to a decrease in stability of the material. However, the extremely high connectivity of UiO–66 allows the presence of both missing linkers and clusters without loss of crystallinity. In this context, Rogge *et al.* investigated the influence of all possible configurations of one to two linker vacancies on the stability of UiO–66. The equilibrium volume is not affected by such vacancies, however properties such as bulk modulus and loss-of-crystallinity pressure, which in turn influence the stability, are affected by the number and configuration of missing linkers [60]. De Vos *et al.* [61] investigated the electronic properties for all possible configuration of defective UiO–66 with up to three missing linkers, showing that some configurations are energetically more stable than others, and that the number and position of missing linkers affect the band gap of the material. From these studies performed on small unit cells, it is already clear that the number of possible defective configurations can be extremely high, and it is still unclear to what extent such point defects are disordered in the material. A regular distribution of point defects that involve missing clusters can lead to different phases in the material, from *fcu* (non-defective), to *bcu* (missing linkers with open channels), *reo* (missing clusters) to *scu* (missing linkers and missing clusters). Missing clusters could in principle increase the catalytic activity of the material more than missing linkers due to the larger pore size. Recently, Cliffe *et al.* [24] showed in a combined theoretical-experimental work that missing clusters on UiO–66(Hf) were correlated and formed nanodomains in the material, characterized by a *reo* phase. The presence of missing clusters or missing linkers can be affected by the choice of modulator [63]. These reports shed light into the complex nature of defects in nanoporous materials.

Active sites for catalysis in UiO–66

One of the breakthroughs of MOF research was the discovery that UiO–66 could contain a high number of open metal sites. As reported earlier, active sites for catalysis in UiO–66 and other Zr–MOFs are present when the zirconium connectivity is decreased from its equilibrium value of 8.

A first way to obtain coordinatively unsaturated zirconium sites is by creation of defects, in the form of missing linkers or clusters. The inherent vacancies in UiO–66 bring unsaturated zirconium Lewis acid sites [28,53,54,59,64] and at the same time enhance the accessibility of the reactants to the active sites as the pore volume increases. The synthesis of defective UiO–66 can be tuned via modulators such as formic acid or trifluoroacetic acid (TFA), that are competing with BDC linkers in binding to the inorganic SBU. Vermoortele *et al.* first showed in a dual

computational–experimental study that the Lewis catalyzed cyclization of citronellal requires missing linkers [65]. For Meerwein reduction, another Lewis catalyzed reaction, the catalytic activity of UiO–66 could be significantly increased by making use of TFA, that introduced a large number of linker vacancies. Additionally, the non-modulated material that contained only a small number of defects showed nearly no catalytic activity [59]. Such relationship between catalytic activity and amount of defects has been reported for different Lewis catalyzed reactions giving clear evidence that the catalytic centers are located on the defective sites [59,65].

The molecular characterization of the defective sites on UiO–66 has been an ongoing research interest in MOF literature. Particular focus has been given on the species adsorbed on the defective site and on the coordination environment of zirconium (Figure 1.6). Early experimental work by Lillerud and coworkers [66] reported that when linker vacancies were present, XRD characterization of the material showed two types of Zr–bonded oxygen atoms. The first type was BDC oxygen, and the other was associated to water species coordinated to the zirconium on the defect site. Later from SXRD measurements, the group of Yaghi [67] proposed that the two Zr–bonded oxygens belonged to physisorbed water molecules and that the charge compensating species was a hydroxyl group. However, following reports did not confirm such configuration.

Molecular simulations were essential to shed light into the detailed molecular structure of the defective brick. Ling and Slater [68] performed MD simulations starting from the configuration proposed by Yaghi and showed a progressive stabilization towards another structure where the two adjacent zirconium atoms were coordinated to a physisorbed water molecule and a hydroxyl group, as shown in Figure 1.6. These sites have been confirmed by a comprehensive computational study of different adsorption possibilities of up to three water molecules on defective UiO–66 performed by Vandichel *et al.* [69] and by simulations performed in this thesis. Such water and hydroxy species are strongly interacting with the zirconium atoms and are sources of Brønsted sites, as indicated in Figure 1.6. It was remarkable to discover that some catalytic processes are not catalyzed by undercoordinated Lewis acid sites, but need also Brønsted sites arising from defect–coordinating species, as will be shown in **PAPER I**. Apart from the interactions of water with the active sites, they may also have a strong influence on the proton conductivity of the material. Serre and coworkers reported that a hydrogen bonded network that spans the octahedral and tetrahedral cages is responsible for the high proton conductivity showed by UiO–66 at high temperatures [62]. Kitagawa *et al.* showed the positive role of defects in proton conductivity in the UiO–66 material, as defective sites provide both larger pores and zirconium undercoordinated sites where water species are adsorbed and can function as proton donors [70]. Farha and coworkers identified three types of acidic protons in the material by means of potentiometric titration: $\mu_3\text{--OH}$ belonging to the inorganic SBU, and two arising from subsequent deprotonation of the physisorbed water and hydroxyl group on

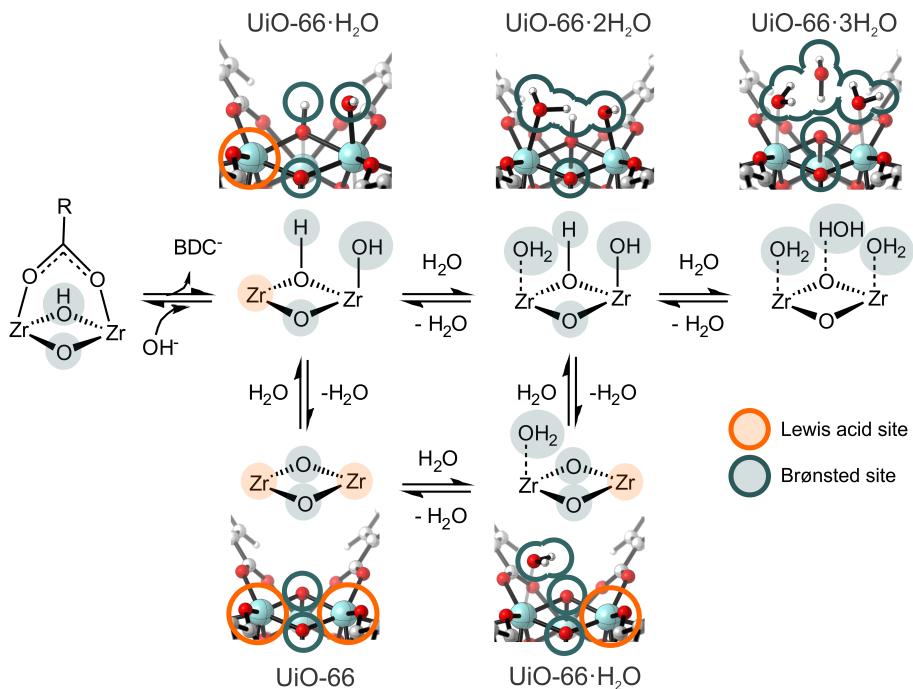


Figure 1.6: Lewis and Brønsted sites that are created when a linker is removed from the UiO-66 zirconium brick, with different number of coordinated water molecules.

the defective site. They made use of the acidic protons belonging to physisorbed water for a quantification of defects in the material as alternative to TGA [71]. The mechanism associated to the change in topology of the hydroxyl groups on the inorganic SBU has been investigated by Yang *et al.* [72]. The interaction between defect–coordinating water molecules and zirconium atoms is strong, and UiO–66 can be partially hydrolyzed [73] while retaining its stability, as will be further discussed in this dissertation.

The defect–coordinating species can be removed by thermal activation at $T > 423$ K (more information on this process is given in **PAPER I**). The more loosely bonded physisorbed water is the first to leave the defective site, followed by the chemisorbed water. The dehydrated active site obtained this way is missing a μ_3 –OH proton and has two 7-fold coordinated zirconium open metal sites as shown in Figure 1.6 (bottom left).

The second activation process by which open metal sites can be created on UiO–66 is the reversible dehydration of the $Zr_6O_4(OH)_4$ cornerstone to obtain Zr_6O_6 by removal of up to two water molecules performed at a temperature range between 523 and 573 K [48]. In this case also 7-fold coordinated zirconium atoms are created and may take the role as Lewis active sites, however the increase in pore size due to linker vacancies does not occur, and the active sites are less accessible than in the defective material. For instance for the citronellal cyclization [65], in absence of defects, linkers should still be partially decoordinated to make space for the reactants to adsorb on the zirconium atoms, therefore the process is less likely to occur.

Functionalization of UiO–66

Reactions on UiO–66 can proceed exploiting purely the Lewis acidity of under-coordinated zirconium atoms, or can also make use of Brønsted sites located on the defect sites. The acidic and basic properties of such groups can be further influenced by linker functionalization [74–76]. Functionalization, which can be done via PSM, represents a strategy to further finetune the properties of the material. For instance, the presence of electron–donating substituents showed an increase in catalytic activity for jasminaldehyde condensation [77]. A positive effect of electron–withdrawing groups was found by the same authors for Lewis catalyzed citronellal cyclization [65]. The beneficial role of amino groups was reported as well by Timofeeva *et al.* [78] for acetylation of benzaldehyde, as well as by Cirujano *et al.* [79, 80] for esterification. The electron donating effect of amino groups would in principle lead to a decrease in the Lewis acidity of the defective zirconium atoms, lowering the reaction rate. Therefore, a change in mechanism in the presence of $BDC-NH_2$ was hypothesized. However, Hajek *et al.* further studied aldol condensation in a computational work [81], and reported a beneficial, but

passive role of amino groups, which is further confirmed in **PAPER I** for Fischer esterification.

Active sites on MOF-808

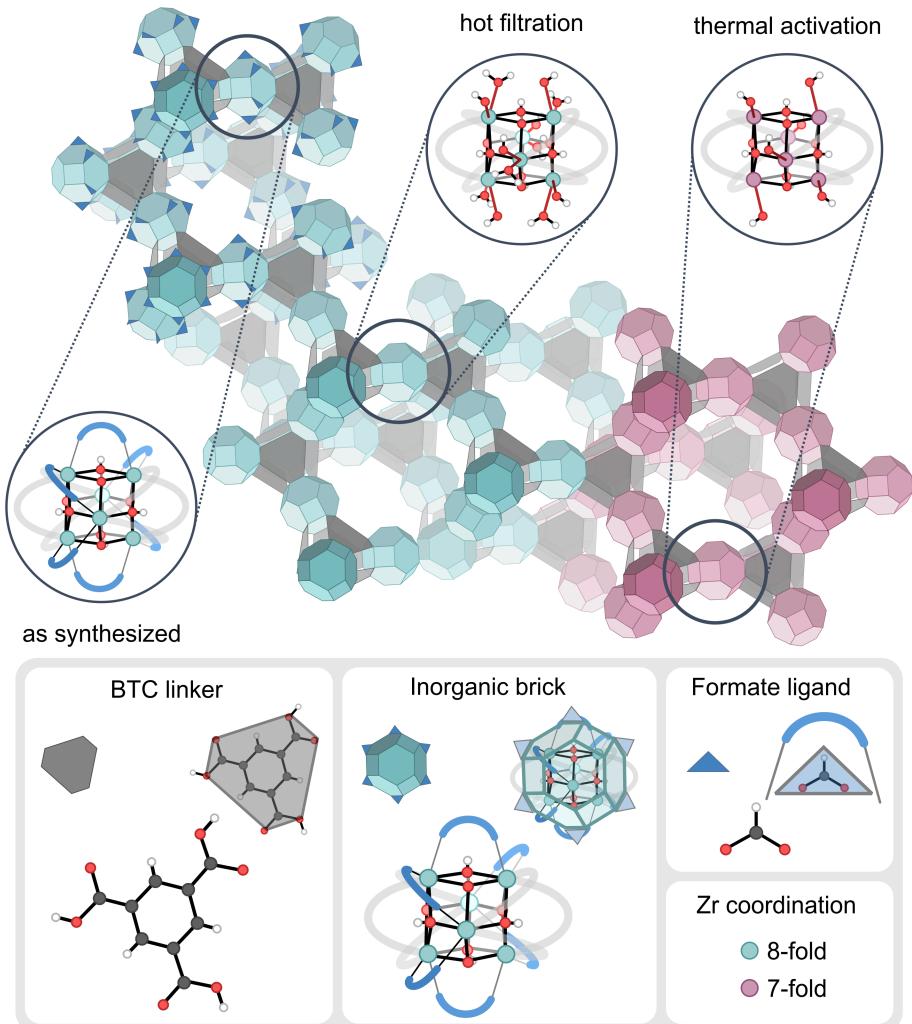


Figure 1.7: Schematic representation of as synthesized and upon activation MOF-808 structures

So far it was shown how active sites can be introduced on UiO-66 by linker or brick removal and dehydration. In the Zr-based MOF family, other less connected materials such as NU-1000 and MOF-808 contain inherent vacancies due to their topology. Although less robust than UiO-66, they possess a higher number of active sites and larger pores and for this reason they are regarded as promising

heterogeneous catalysts. Moreover, their large pore size allows for interesting modifications such as atomic layer deposition in MOFs (ALD) [47]. MOF-808, formed by the inorganic $Zr_6O_4(OH)_4$ cornerstone and trimesate (BTC) linkers, represent the least connected MOF in the Zr-MOF family [44], and its brick can be regarded as an extreme case of the defective UiO-66 (Figure 1.7). MOF-808 is a promising catalyst that is still less studied than UiO-66, but has a lot of potential for possible applications. For instance, it shows a higher catalytic activity for certain Lewis catalyzed reactions, such as Meerwein-Ponndorf-Verley (MPV) reduction [82, 83]. Moreover, MOF-808 represents the first evidence of a superacid in MOFs, showing superacidity after post-synthetic treatment with sulfuric acid [84].

The rigid BTC tritopic linkers stabilize the structure and yield exceptionally wide channels which are required for the diffusion of substrates. The octahedral crystals contain 6 BTC linkers per inorganic node and are characterized by a *spn* topology. In the material, each zirconium atom is bonded to four oxygens from the inorganic brick and two oxygens from the BTC linkers. The other two bonds that are needed to reach the total coordination of 8 are provided by species present in solution that do not contribute to the framework connectivity.

In the as-synthesized material (Figure 1.7), the zirconium atoms are capped with the relative mobile formate modulator. The material at this stage does not contain catalytically active sites and must be activated by post-synthetic treatment. By hot filtration, the formate groups can be eliminated and replaced by solvent water and hydroxyl groups [82, 83]. In the catalytically active material, each brick is connected to six BTC linkers, six water molecules and six hydroxyl groups, and possesses a high number of potential Brønsted sites. Klet *et al.* reported the presence of four different types of protons in the material [71], each yielding a different acidity. The acidity of these protons can influence reactions, and its characterization represent a current challenge. From this structure, Lewis acid sites can be created by thermal activation, similarly as in UiO-66. In this case, hydroxyl groups remain connected to the zirconium atoms and the material possesses both Lewis and Brønsted sites that can be used for reactions. Moreover, the hydroxyl groups could further be used as anchors to incorporate new features in the material.

1.2 Outline and goal of the thesis

In this PhD dissertation, state of the art molecular modeling techniques were used to unravel at the nanoscale the molecular nature of defects and active sites in two zirconium MOFs: UiO-66 and MOF-808. From a purely experimental basis it is very hard to follow processes such as activation, reactions or formation of defects at the molecular level. The comprehensive insight obtained from molecular modeling can offer precious understanding in these processes at the nanoscale

level. Furthermore it is the ambition to model the described processes at operating conditions, thus at realistic conditions of temperature, pressure and guest loadings in the pores of the material. This can ultimately allow to fine-tune the active sites to target specific applications. Processes taking place in MOFs usually occur at mild conditions in the presence of solvent confined in the pores therefore, multilevel modeling techniques are required to accurately describe both, the material and reaction environment. Moreover, the processes happening at the molecular level are very dynamic and can involve rare molecular events. All these complex dynamics need to be mapped appropriately. It is shown that enhanced sampling methods used in this thesis are crucial to reveal the strong interactions of protic solvent with the active sites modulating their nature and actively participating in chemical transformations. The insight obtained in this PhD into the solvent assisted ligand exchange mechanism shed light on the role of solvent in modulating the properties of the active sites at operating conditions, which influence structural stability and create active sites for catalysis. The research was performed in close collaboration with experimental and theoretical partners. The results obtained in this thesis rationalize the experimental findings and shed light on the molecular processes that occur around the active site, which are crucial to engineer MOFs for future applications. This PhD thesis is organized as follows:

- In Chapter 2, a condensed theoretical overview of current molecular modeling techniques within MOF research is given. Particular attention is drawn on how these techniques can be applied to obtain insight into structural and catalytic properties at operating conditions.
- In Chapter 3, the main results of this PhD thesis are summarized. The links between theory and experiment are highlighted throughout the chapter. All results are the result of fruitful experimental and theoretical collaborations and have been published in international peer-reviewed journals.
- In Chapter 4, the main conclusions of this thesis work, as well as perspectives on future research are given.

2

Modeling metal organic frameworks

The understanding of catalytic processes in MOFs is a very challenging task. MOFs are materials of complex nature, and reactive processes in these materials are elusive and difficult to track on a purely experimental basis. Molecular simulations offer an alternative approach that starts from the construction of models that can explain, complement, and predict experiments. With growing computational power, computational models can aim at giving a more and more accurate description of materials at operating conditions, narrowing the gap between theory and experiment. Often, the structural properties and chemical transformations that take place on the active sites need to be investigated using a combination of multiple computational techniques, and the problems need to be tackled from different angles. In this chapter, an overview of the different computational methods that can be used to study reactive processes in MOFs will be given.

2.1 Framework topology

A crucial decision when performing simulations lies in the choice of the model system, and what should be included in it. When choosing a model to represent the system under study, there is always a fine balance between accuracy and computational cost. On the one hand, it is crucial to use a computational model that captures all the relevant properties of the material and mimics the experimental structure as closely as possible. On the other hand, it is often convenient to use

approximations in order to obtain a description of the processes at a larger scale. The focus in this work are the active sites that can be used for catalysis, therefore an accurate electronic description of this region of the material is imperative. Nevertheless, the activity of these sites for chemical reactions can also be influenced by other factors, such as the pore size or functionalization. Therefore, to describe active sites in MOFs and other nanoporous materials, which can have rather large unit cells and non-periodic structural defects, the first question that needs to be asked is how to account for periodicity or molecular environment. Two conceptually different approaches can be used as shown below.

Extended cluster model

A very computationally efficient approach consists in neglecting periodicity and extracting a finite cluster of atoms from the periodic structure. This cluster model, displayed in Figure 2.1, contains the active sites and their surroundings but consists in a limited number of atoms, allowing to substantially decrease the computational cost. This allows both a more accurate treatment of the electronic structure, and a screening of different possible geometrical configurations of adsorbates, which is useful in the search for transition states (TS). Moreover, very efficient TS searching algorithms have been developed for such systems in Gaussian [85], the most widely used code for cluster calculations and that we used in the framework of this thesis. When extracting a cluster, particular attention has to be drawn to the termination of bonds and the charge compensation, that have to be done in the most realistic way. The rest of the crystal structure does not surround the external cluster atoms. Some of these atoms need to be fixed in order to mimic the periodic environment and prevent nonphysical deformations that would affect an estimation of the entropy [87].

Cluster calculations are an excellent way to benchmark and perform a first qualitative screening of reactions and possible configurations and have been for long the standard computational tool when studying reaction in nanoporous materials. However, confinement effects and possible structural rearrangements are neglected in this model, and for certain reactive processes they could play an important role. The influence of solvent in the pores can also be crucial for the outcome of a reaction and cannot be explicitly studied by cluster models. Periodic calculations resolve this shortcoming, and as computational power grows, the heterogeneous catalysis community is shifting towards these more expensive, however more accurate models.

Periodic model

Periodic models, that fully take into account the periodicity of the crystal, enable to describe the whole topology of the framework. These calculations make use of periodic boundary conditions (PBC), which allow to simulate bulk phases with a

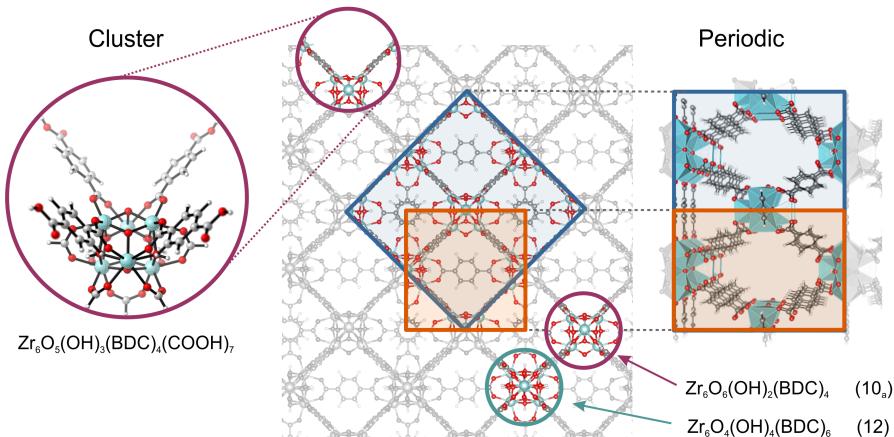


Figure 2.1: Left: extended cluster model cut from the periodic structure of UiO-66. The cluster contains the active site, the brick and the linkers in the closest proximity to the active site. Right: representation of the unit cells containing the defect. In blue, the conventional 4-brick unit cell, in orange, the 2-brick unit cell used for the calculations. The two different bricks are highlighted in orange. The 10-fold coordinated brick has two terephthalate linkers missing. Adapted from ref. [86] with permission of the American Chemical Society.

limited number of atoms. In this model, the unit cell is replicated infinite times in each direction. When one atom disappears from one side of the unit cell it will reappear on the opposite side and each atom interacts with its neighbors in the same unit cell but also in the adjacent ones. Spurious interactions between the atoms can be avoided by applying a *minimum image convention*, for which each atom interacts with its nearest neighbor or periodic image. In the case of long range interactions, such as the electrostatic ones, other techniques need to be used, such as Ewald summation [88], where the potential is divided into a short range contribution, calculated in real space, and a long range contribution, calculated in reciprocal space using a Fourier transform.

In the case of UiO-66, the conventional unit cell [46] contains 4 zirconium bricks (Figure 2.2, blue). In the calculations of this thesis, BDC linkers have been removed from the unit cell to introduce defects which are active sites in catalysis. Different amounts of missing linkers with different topologies have been considered 2.2. An interesting topology is the one denoted as type 6 in the work of Rogge et al. [26] that is characterized by a channel which offers good perspectives for the diffusion of guest molecules. This unit cell (displayed in blue in Figure 2.2) can be reduced by symmetry to a 2-brick unit cell (in orange, Figure 2.1) which offers the best compromise between accuracy and computational cost. This reduced unit cell is used in most of the calculations performed in this thesis. In **PAPER I** we

show that for catalytic purposes, the free energy differences calculated for the same chemical processes on the two unit cells is negligible, therefore the 2–brick unit cell represents a good model system. When modeling disorder, however, periodicity decreases and bigger unit cells have to be taken into account. For this reason, in **PAPER IV**, we studied defective 4–bricks unit cell with missing linkers ranging from zero to three. Moreover, in some cases it may be necessary to use supercells to avoid spurious interactions between adjacent cells.

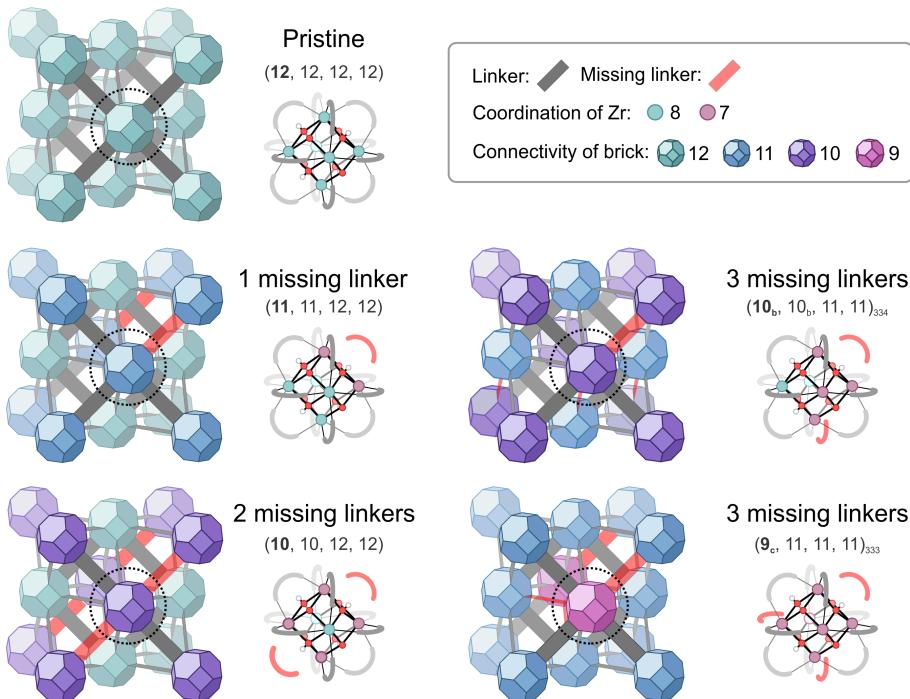


Figure 2.2: Representation of the different ways to remove up to three linkers from a 4–brick UiO-66 unit cell. Adapted from Ref. [89] with permission of the American Chemical Society.

2.2 Theoretical methods

Electronic energy methods

A key concept involved in the study of any chemical or physical transformation is the knowledge of the potential energy surface (PES), which is a function of the coordinates of all the atoms of the system. The PES is always the reference quantity in our simulations and every atomic configuration can be represented as a point in this hypersurface, with a given value of potential energy (Figure 2.3).

Ideally, by calculating the value of the PES for each atomic configuration we can obtain all information on the system and on the transformations that can occur. However, the complexity of this surface escalates quickly with the number of atoms, and the sampling of its relevant regions represents the main challenge of molecular simulations. The information gained by exploring the PES is tightly connected to the experimental observables. Statistical physics acts like a bridge between the microscopic insight that is gained through molecular simulations and the macroscopic properties which are measured experimentally. In principle, all macroscopic properties of a system can be derived from its wavefunction. To calculate it, the stationary Schrödinger equation is solved:

$$\hat{H} |\psi\rangle = E |\psi\rangle$$

where ψ is the wavefunction, \hat{H} is the Hamiltonian of the system, and E is the total energy. The resolution of this equation is at the heart of computational chemistry and will in principle provide the exact description of matter, but it is nevertheless extremely difficult to solve for most of the electron systems. The presence of electron–electron interactions makes it a many-body problem which is very hard to solve. Therefore, a series of approximations need to be made to solve this problem for systems with a high degree of complexity.

Born–Oppenheimer approximation

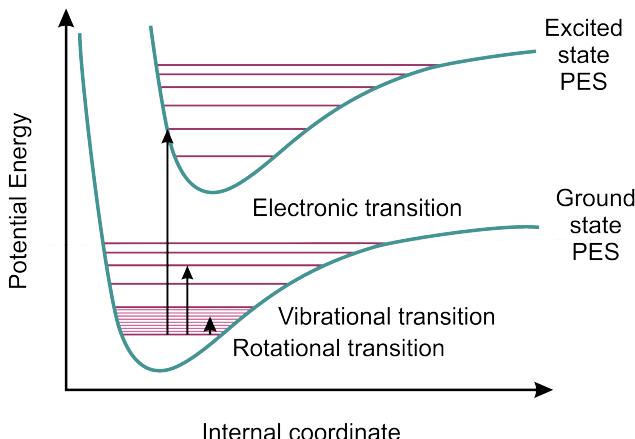


Figure 2.3: The two lowest PES in the BO approximation for a diatomic molecule. In blue, the electronic PES for the ground state and first electronic excited state (UV–Vis transitions). In orange, the vibrational energy levels (IR transitions), in grey the rotational levels (microwave transitions).

For all calculations performed in this work, we rely on the so-called Born–Oppenheimer (BO) approximation [90]. In this treatment, nuclei are considered

as classical points which move in the potential energy surface generated by the electrons (Figure 2.3). This way, to each nuclear configuration a corresponding electronic energy can be assigned, and nuclear coordinates enter in the Schrödinger equation only as parameters, allowing to construct a BO surface, or PES. This approximation holds since nuclei are much slower than electrons, therefore the motion of electrons is instantaneous from the nuclei point of view. This approximation does not always hold, especially when dealing with light nuclei such as hydrogen. In these cases, nuclear quantum effects can have an impact on the measured properties [91]. In most cases, the electronic ground state is also not interacting with the higher electronic states because of the high energy difference. In the BO approximation, the electronic energy levels are also considered fully separated and do not interact with each other. For this reason, the approximation is also called adiabatic approximation. Additional interactions have to be considered when two surfaces lie close to each other, for instance in the neighborhood of conical intersections.

Force Fields

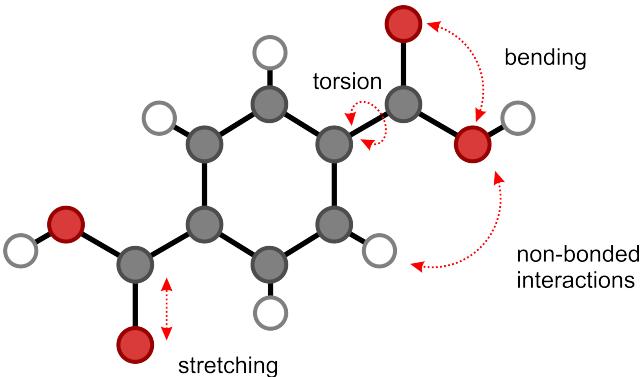


Figure 2.4: Representation of some of the molecular modes taken into account in a generic force field model.

The simplest way to describe interactions between atoms which determine the PES is the so-called “balls and springs” model. In this treatment, all interactions are represented by interatomic classical potentials which are parametrized to reproduce the results of more accurate quantum mechanical calculations. In this work, generic force field calculations have been used in some cases to give preliminary input structures for more costly *ab initio* calculations, through which the description of chemical transformations is possible. Force fields are often constituted by harmonic potentials (Figure 2.4), and the relative parameters are generated from reference experimental or *ab initio* data, with packages such as QuickFF [92]. Force fields can be successfully applied to describe physical transformations but cannot be employed in case of chemical reactions where bonds may be formed and

broken. Reactive force fields, such as ReaxFF [93] are currently being developed, but their application in complex heterogeneous reactions is still an ongoing chemical challenge and is out of the scope of this work. For this reason, the description of reactive processes needs a more advanced treatment, where electronic distributions are explicitly taken into account.

Density Functional Theory

Density functional theory (DFT) has become the method of choice for the study of chemical systems, due to its good trade-off between computational cost and accuracy of the obtained results. DFT began in the 1920's with the work of Thomas and Fermi [94, 95], but it was only in the '60s that it became a complete and accurate theory, shown in the work of Kohn, Hohenberg and Sham [96]. The fundamental property that DFT describes is electron density as opposed to many body electron wavefunctions, which allows to reduce enormously the number of variables in the case of complex systems. Two fundamental theorems by Hohenberg and Kohn state that there is a unique relation between electronic density and total wavefunction, therefore the ground state density allows us to determine all properties of the system. Moreover, the ground state density can be obtained from a minimization of the total energy functional with a variational method by solving the so-called Kohn–Sham equations. The global minimum value of the functional determines the exact ground state of the system. This way it is possible to obtain the total energy of the system and the forces which act on the atoms, two quantities which are needed in all the simulations performed in this work.

To minimize the energy, Kohn and Sham [97] introduced a method which replaces the many-body problem with an auxiliary system of non-interacting particles, allowing a fast solution of the eigenvalue problem. What needs to be added in this treatment is an additional functional which describes exchange and correlation. Nowadays one of the greatest challenges in DFT consists in the search for an accurate expression for the exchange–correlation functional. The simplest method is known as Local Density Approximation (LDA) initially proposed by Kohn and Sham [97] and can also be adapted to include spin in the Local Spin Density Approximation (LSDA) [98]. A more refined method is the Generalized Gradient Approximation (GGA) which involves the calculation of the gradient of electron density and includes functionals such as B88 [99], LYP [100] and PBE [101, 102], used in this thesis. More recent functionals are the so-called hybrid functionals, which include the Hartree–Fock (HF) exchange, such as B3LYP [99, 100, 103], which is a combination of B88, LYP and LDA with HF, and PBE0 [104], which mixes PBE with HF. These functionals can give a more accurate electronic description of the system but are computationally very expensive. As compromise between accuracy and computational cost is a geometry optimization with PBE, and a single point calculation to refine the energies with B3LYP, as performed in **PAPER I**. This approach, denoted as B3LYP//PBE, allows to avoid the high computational cost

associated to B3LYP geometry optimizations, especially in periodic calculations. A comparison of different combinations of functionals has been reported by Hajek *et al.* [81] for aldol condensation in cluster models. This work shows that PBE geometry optimizations are sufficiently accurate, and provide energy barriers that are comparable to B3LYP for this kind of systems.

Dispersion interactions

In this thesis we often encounter noncovalent interactions which need to be treated with high accuracy, such as the adsorption of guest molecules on the zirconium Lewis acid sites or interaction between solvent molecules. One of the challenges of DFT methods is the description of long range dispersive interactions such as London forces, which are commonly referred to as van der Waals interactions. These interactions are due to many particle electron correlation effects which are present also in absence of charges and can have a significant impact on the noncovalent interaction energy. To tackle this problem, various dispersion schemes have been proposed. One of the most used is currently the Grimme–D3 method [105], where a damped $-C_6R^{-6}$ function is added to the DFT functional. Recently, more advanced dispersion schemes have been developed, such as the many body dispersion scheme [106], or the one of Tkatchenko and collaborators [107], although for the systems we are studying not many benchmarks of these new methods have been performed so far. Wieme *et al.* [108] recently compared different dispersion methods with respect to RPA + SE (random phase approximation with single-excitation effects calculations) in reproducing the breathing behavior of the flexible MIL-53(Al) MOF. The influence of difference functionals and dispersion schemes was also tested by Hajek *et al.* for the adsorption of pentene on the H-ZSM-5 zeolite [109].

Geometry optimization

In order to obtain molecular structures that have physical significance and their relative energies, the arrangement of the atoms needs to be optimized. There are generally two types of molecular structures that we need to find in our simulations, the equilibrium geometries, which correspond to *minima* of the PES, and the transition state geometries, which correspond to first order saddle points, as displayed in Figure 2.5. These points are characterized by null first derivatives of the energy (the total forces acting on each atom are sufficiently close to zero), all positive second derivatives for local *minima* and one negative second derivative for first order saddle points, which correspond to transition states.

The geometrical optimization of reactants and products consists in a minimization of the energy along the nuclear coordinates. Often the starting point is the experimental structure which can be obtained from diffraction data. In the most used codes several minimization methods are implemented, each characterized by a different computational cost and robustness, such as steepest descent, conjugated

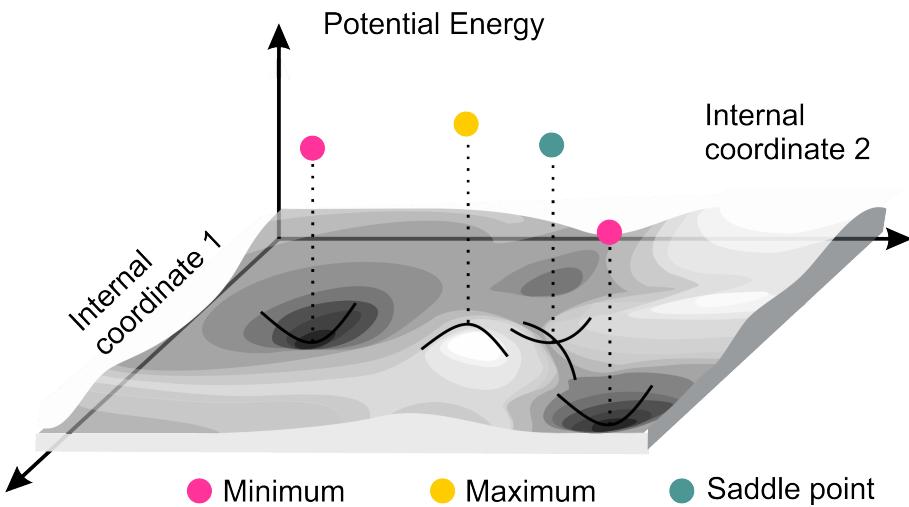


Figure 2.5: Schematic representation of the potential energy surface and the stationary points.

gradient or simulated annealing. The algorithms will find local *minima*, and do not guarantee that the system will be in a global *minimum*, therefore the minimizations must start from a sufficiently good guess.

The search for transition states is far from trivial and often requires an iterative procedure involving different methods and requiring a good knowledge about the system and chemical process under study. In the calculations performed in this thesis, we often start from an equilibrium structure and as a first guess, we adapt the bond lengths and angles to be close to the transition state with a molecular editor such as Zeobuilder [110]. These bond lengths are then fixed, and the rest of the structure is reoptimized. The Hessian of this partly optimized structure needs to be then computed, and the vibrational modes analyzed, to check which (if any) negative frequency corresponds to the transition state. The system can be optimized again without the constraint using an improved dimer method along a selected eigenvector [111] which is followed by the optimization with a quasi-Newton method [112]. In some difficult cases, the TS search can be initiated on simpler cluster models and optimized in a code in which methods are usually implemented to directly find the TS structure. This way, a first guess of the TS geometry might be obtained and further transferred to the periodic model. In both transition states and *minima*, if there are superfluous negative frequencies, these need to be removed. Often, it is sufficient to minimize the energy along that vibrational mode. Single point energy calculations can be performed for different values of the displacement along that mode, and the lowest point in energy can be used as starting point for a subsequent minimization of all the coordinates.

Cell optimization

In the case of periodic systems, not only the structure, but also the unit cell needs to be optimized. This is not trivial, as when using a finite plane wave basis set the number of plane waves depends on the volume of the unit cell. If the volume changes during the optimization, artificial forces which go under the name of Pulay stress can arise. This would require many iterations to optimize the volume. In this thesis, another approach was used [113] which relies on an equation of state fit. For a given volume, for instance taken from experimental data, the unit cell is optimized. Then a set of equally spaced different volumes is defined and for each of these points the geometry and unit cell parameters are optimized. This way it is possible to construct an energy–volume curve, which for a rigid system can be fitted with a Birch–Murnaghan equation of state [114, 115], allowing to extract the volume V_0 which corresponds to the *minimum* electronic energy E_0 .

$$E(V) = E_0 + \frac{9V_0B_0}{16} \left\{ \left[\left(\frac{V_0}{V} \right)^{\frac{2}{3}} - 1 \right]^3 B'_0 + \left[\left(\frac{V_0}{V} \right)^{\frac{2}{3}} - 1 \right]^2 \left[6 - 4 \left(\frac{V_0}{V} \right)^{\frac{2}{3}} \right] \right\}$$

Where B_0 and B'_0 are the bulk modulus and its derivative. A new structure is then generated at this given volume and coordinates and unit cell parameters are optimized again.

Molecular vibrations

As seen in the previous paragraph, for many purposes in this thesis we need to calculate the second order derivatives (Hessian matrix) of the PES, which are associated to molecular vibrations. First of all, the Hessian gives us information about the curvature of the surface and the nature of the stationary points encountered during the minimization. The second order derivatives are obtained by displacing the atoms in the three directions and calculating the energies, then the Hessian is diagonalized to determine the eigenvectors that correspond to the vibrational motions. From the Hessian we can calculate the vibrational frequencies, which open the door to a lot more information on the system than a single point calculation. As a matter of fact, single point calculations are performed at 0 K, but even at this temperature nuclei vibrate around their equilibrium positions, and this movements are responsible for vibrational entropy. We can approximate these motions with those of harmonic oscillators, by using the vibrational frequencies constructed from the Hessian.

These frequencies can then be used to estimate the value of the vibrational entropy at finite temperatures, as will be explained later. In the calculations performed in this thesis, due to computational limits, a partial Hessian approach (PHVA) was used when dealing with reactions, as implemented in the TAMkin toolkit [116]. The quantity that needs to be derived from these calculations is the change in free energy, which mainly depends on the parts of the system that

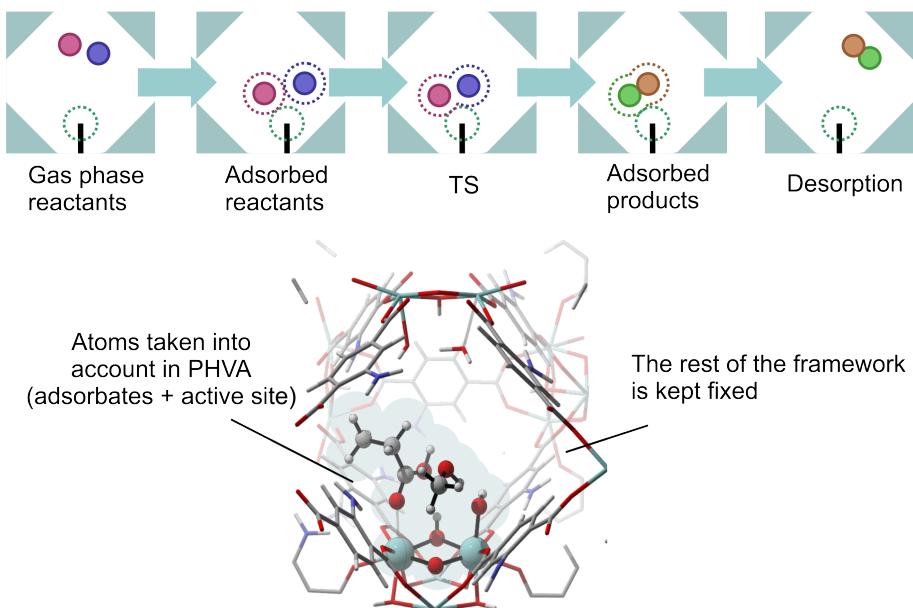


Figure 2.6: Representation of the atoms taken into account in the Partial Hessian Vibrational Analysis (PHVA) approach. Top: a schematic representation of a reactive process in nanoporous material, bottom: a snapshot from the static calculations where the atoms of the active site and the adsorbates are highlighted

change during the reaction, in the case of a heterogeneous catalyst the active site and the adsorbed reactants. Therefore, restricting the entropy calculations only to this part of the system is a good approximation that allows to decrease enormously the computational cost [117]. This approach has been used in the calculation of the free energy barriers for the Fischer esterification on UiO–66 (**PAPER I**), where the atoms taken into account were the adsorbed reactants and four atoms of the active sites in their immediate proximity, as displayed in Figure 2.6.

In many cases, negative or low frequencies can be present, especially when involving large systems such as species adsorbed in MOFs [87]. Negative frequencies with large absolute value ($\nu < -50 \text{ cm}^{-1}$) can arise if the system is not fully optimized, meaning that the energy can be further minimized along this coordinate. In this case, a series of geometries can be extracted along the coordinate corresponding to the negative eigenvalue. For each of these structures, the electronic energy can be obtained with a single point calculation. The geometry corresponding to the lowest energy along the vibrational mode can be then further optimized to find a lower energy minimum. This procedure can be iteratively performed for each of the negative frequencies. In case of frequencies approaching zero, the harmonic oscillator model would break down. Therefore, low frequency modes, or so-called soft modes ($|\nu| < |50 \text{ cm}^{-1}|$) are artificially replaced by a frequency of 50 cm^{-1} [87]. In this sense, PHVA approach allows to avoid considering all spurious low-frequency modes arising from lattice vibrations of the framework and only limit to those arising from the reactants.

2.3 Free energy

The central thermodynamic quantity that determines the outcome of a reaction is the free energy change associated to the process. In general, a chemical system will undergo changes in a direction that minimizes its free energy, until an equilibrium is reached. Knowing the difference in free energy between reactants and products allows us to know the equilibrium constant for a given reaction. The Gibbs Free energy can be decomposed in an enthalpic and an entropic contribution, that can be evaluated from the simulations knowing the molecular partition functions. Initially, the total internal energy U has to be obtained from the electronic energy ε_0 , the zero-point vibrational energy E_{ZPE} and the molecular partition function Q at constant number of particles n and volume V :

$$U = U_0 + RT^2 \left(\frac{\partial \ln Q}{\partial T} \right)_{n,V}$$

$$U_0 = \varepsilon_0 + E_{ZPE}$$

Where R is the gas constant, equal to the product $N_A \cdot k_B$ between Avogadro's number and Boltzmann constant. The molecular partition function Q can be split

in its translational, rotational, and vibrational components:

$$Q = Q_{trans} Q_{rot,ext} Q_{vib}$$

The enthalpy H corresponds to the total energy plus the work associated to the change in volume.

$$H = U + p_0 V$$

The entropy S can be directly obtained from the partition function:

$$S = R \ln Q + RT \left(\frac{\partial \ln Q}{\partial T} \right)_{n,V}$$

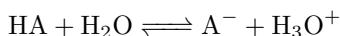
Finally, the Gibbs free energy G will be:

$$G = H - TS = U_0 + p_0 V - RT \ln Q$$

Where in the case of non-interacting particles $p_0 V = RT$ following the ideal gas law. We can therefore define a free energy surface (FES) that is function of coordinates of the system, and can be derived from the PES.

Equilibrium

As explained above, there is a tight connection between free energy and equilibrium concentrations in chemical reactions. As an example, an equilibrium which is of utmost importance in chemistry is the acidic dissociation of species in aqueous solution:



The acidic dissociation constant (K_a) is an important equilibrium constant in chemistry and is equal to the ratio between the concentration of products and reactants when the reaction reaches the equilibrium. It is often reported with its negative decimal logarithm as pK_a .

$$K_a = \frac{[\text{A}^-][\text{H}^+]}{[\text{HA}]}$$

The equilibrium constant is equal to the Gibbs free energy change from reactants to products.

$$\Delta G = RT \ln K_a$$

$$pK_a = \frac{\ln 10}{RT} \Delta G$$

Therefore, knowing the Gibbs free energy difference between reactants and products, we can have important information about the equilibrium composition of a chemical system. However, kinetic factors can sometimes play a major role, and equilibrium cannot always be easily reached.

Transition state theory

A chemical reaction is a process that through rearrangement of the atoms transforms one stable state into another. Every elementary reaction can be represented as a minimum energy path connecting two *minima* along the FES. Furthermore, along this reaction path the existence of a saddle can be postulated, which is the highest point in energy that needs to be crossed to go to the product state. The saddle point is typically called transition state or activated complex and it is the basis for the transition state theory (TST) developed by Eyring in the 1930's, one of the most successful chemical theories which allows to explain reaction rates of elementary chemical reactions. The assumption of the theory is that there is a *quasi*-equilibrium between reactants and activated complex, and the rate constant can be obtained by the size of the energy barrier and by the frequency at which the system can cross the barrier. This is possible because the barrier acts as a bottleneck in the reaction, its crossing is a rare event and all the kinetics depends only on it. For a unimolecular reaction, the rate constant can be derived from the partition functions of reactant, TS and their energy difference:

$$k(T) = \frac{k_B T}{h} \frac{q_{TS,\ddagger}}{q_R} e^{-\frac{\Delta E^\ddagger}{k_B T}}$$

Where k_B is the Boltzmann constant, h is the Planck constant, q_R and $q_{TS,\ddagger}$ are the molecular partition functions of reactants and activated complex for all coordinates except the reaction coordinate, evaluated from the zero-point vibrational level.

$$q_{vib,i} = \prod_{i=1}^{N_{dof}} \frac{1}{1 - e^{-\frac{h\nu_i}{k_B T}}}$$

Where N_{dof} is the number of vibrational degrees of freedom of the system. The energy difference ΔE^\ddagger includes electronic energy and zero-point vibrational energy difference at 0 K:

$$\Delta E^\ddagger = E_0^{TS} - E_0^R + \Delta E_{0,vib}$$

$$\Delta E_{0,vib} = \sum_{i=0}^{N_{dof}-1} \frac{h\nu_i^{TS,\ddagger}}{2} - \sum_{i=0}^{N_{dof}} \frac{h\nu_i^R}{2}$$

A distinction has to be made between intrinsic (ΔG_\ddagger) and apparent (ΔG_{app}) energy barrier, indicated in 2.7. ΔG_\ddagger , or intrinsic barrier, corresponds to the reference state in which all reactants are adsorbed on the material, and to a unimolecular rate equation. These are the barriers commonly reported in theoretical works. ΔG_{app} is the apparent activation energy measured in experiments, and corresponds to a reference state in which reactants are not yet adsorbed. In this case, a multimolecular rate equation has to be used. To properly compare the two activation energies, a good estimate of the adsorption energy of reactants is needed [118].

TST has some limitations, and may fail in the case of labile intermediates, when nuclei deviate from a classical behavior, or at high temperatures. For a given reaction, in fact, there will be many paths characterized by different barriers, and at low temperature only the lowest one will be likely to be crossed. When the kinetic energy is high enough, many other paths will be activated. The transition state will occupy a larger region of the PES, and it will not be possible to derive entropy from the vibrational partition functions.

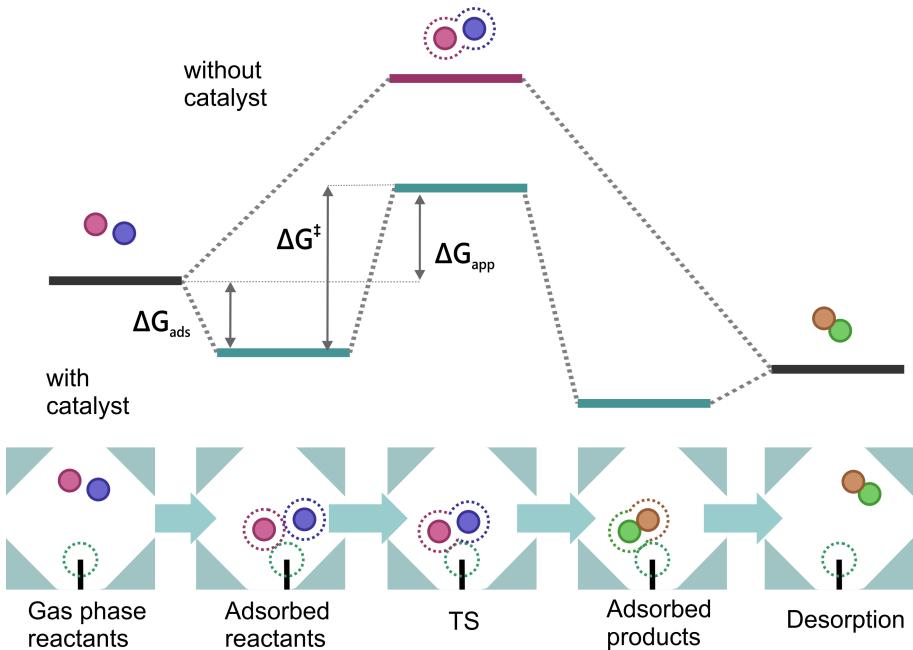


Figure 2.7: 1D free energy profiles for a given reaction with and without catalyst, indicating the adsorbed initial, final states and the localized transition state, schematically illustrated in below. The adsorption free energy, intrinsic and apparent barriers are obtained by static calculations and indicated by ΔG_{ads} , ΔG^\ddagger , and ΔG_{app} , respectively;

2.4 Exploring the free energy surface

Static calculations, where molecular vibrations are approximated using harmonic oscillators, can fail to give an accurate representation of the entropy when there is a high configurational freedom. When the FES is flat with respect to $k_B T$, the system at equilibrium can evolve in a larger region of the PES and move along more than one *minimum*. In this case, vibrational frequencies are anharmonic and it is not possible to represent the system by approximating around one single *minimum* (Figure 2.8). Therefore, static calculations are not always sufficient in

describing the system at operating conditions. In this view, molecular dynamics (MD) techniques, which follow the time evolution of the system, can resolve this shortcoming.

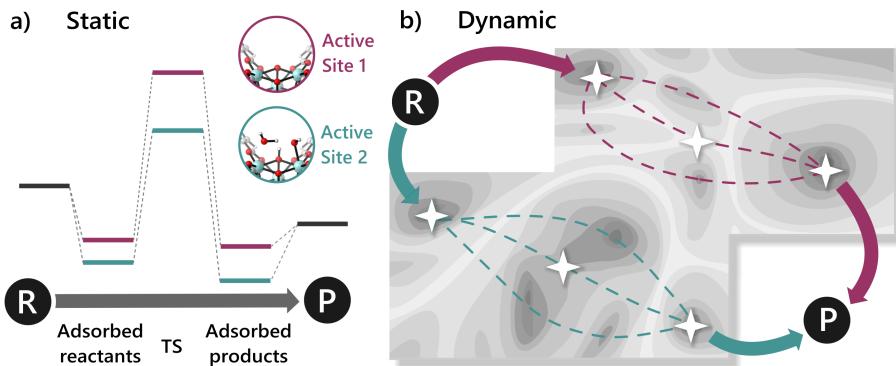


Figure 2.8: a) 1D free energy profiles for a given reaction on two different active sites in UiO–66 (insets), indicating the adsorbed initial and final states and the localized transition state for this reaction; (b) possible 2D representation of the given reaction on the two active sites as obtained using advanced dynamic techniques, indicating the three critical points on the potential energy surface. Adapted from ref. [26] with permission of the Royal Society of Chemistry.

Ab initio Molecular Dynamics

From MD simulations, thermodynamic properties such as free energy can be obtained taking into account a whole region of the PES instead of a single point. This is based on the ergodic theorem, that in one of its formulations states that the time average of equilibrium properties is equal to the ensemble average, in the limit of a sufficient long simulation.

MD simulations are based on solving Newton's equations of motion:

$$M_i \ddot{\mathbf{R}}_i = \mathbf{F}_i = -\nabla_i V$$

where M_i and \mathbf{R}_i are the mass of a given nucleus and its coordinates, \mathbf{F}_i the forces that act on it, which correspond to the gradient $\nabla_i V$ of the PES. There are many ways to calculate these quantities and to integrate the equations of motion, and at present time, chemists and physicists can choose between a plethora of MD techniques which span a whole range of complexity, accuracy and computational cost. In the calculations performed in this thesis, potential energy and forces on the PES are calculated from first principles by means of DFT to account for the full dynamic behavior of the material by ab initio molecular dynamics (AIMD). The calculation of electronic properties which define the PES is decoupled

from the propagation of nuclear motions, in a method called Born–Oppenheimer Molecular Dynamics (BOMD). Other famous AIMD methods, which differ by how the calculations of electronic potential and the equation of motion are combined, are the Car–Parrinello MD (CPMD) [119], where a fictitious electronic kinetic energy is added to the lagrangian, or the Ehrenfest MD, based on the namesake theorem [120, 121].

The first MD calculations were performed in the microcanonical (NVE) ensemble, where total energy, number of particles and volume are fixed. However, in experiments it is often the temperature that is fixed, not the energy. In general, the choice of the ensemble depends on the thermodynamic quantities that need to be determined. Nowadays there are many thermodynamic ensembles in which the simulation can be performed, an overview of which has been given by Rogge *et al.* [122]. The most convenient for a comparison with experiments are the canonical (NVT), with fixed number of molecules, volume and temperature, or the isothermal–isobaric (NpT), with fixed number of molecules and temperature, but where the volume can fluctuate. In order to have a fixed average temperature, some control of the kinetic energy of the atoms is needed. Various thermostats, which differ in terms of speed and robustness, are implemented in every MD code. In this thesis, Nose’–Hoover thermostat was used, where the system is connected to a heat bath. The pressure is also controlled in simulations by means of a barostat. The most commonly used is the one developed by Martyna, Tobias, and Klein (MTK) [123].

Radial distribution functions

From MD trajectories, different macroscopic properties can be derived. An important structural property is the radial distribution function (RDF) or pair correlation function $g(r)$, that describes the probability density between specific atoms as a function of the distance, normalized with respect to a probability distribution of a homogeneous gas with the same density. The RDF can be calculated from the following expression:

$$g_{ij}(r) = \frac{dn_{ij}(r)}{4\pi r^2 dr \rho_i}$$

where dn_{ij}/dr represent the number of atoms of type j within a certain distance of the atoms of type i , and $\rho_i = V/N_i$ represent the density of the homogeneous distribution. The integral of the RDF can also give valuable insight into the number of atom pairs at a given distance, such as what is the average coordination of a certain chemical species.

Vibrational density of states

Information on the time evolution of certain quantities in the system can be obtained by calculating time autocorrelation functions. The correlation of a certain

variable with itself is equal to:

$$\langle X(0) : X(t) \rangle = \lim_{N \rightarrow \infty} \frac{1}{T} \int_0^T X(t) X(t + \tau) d\tau$$

Autocorrelation functions offer a measure of the response of the system to a given stimulus, which is function of the density of states. The response of a system to a perturbation is equal to the power spectrum of the autocorrelation function of the fluctuations of the involved observable [124]. For instance, by calculating the power spectrum of the atomic velocities autocorrelation function, we can obtain the vibrational density of states, as done in **PAPER IV**. These spectra contain all the dynamic and anharmonic information that is neglected in the static frequency calculations.

Towards modeling at operating conditions

With the growth in computational time, the new challenge is constituted by modeling the system at operating conditions. To describe the active sites at reaction conditions, the computational model should reflect the realistic conditions during the reaction, and its complex reaction environment. Within the challenge of *operando* modeling, Grajcar *et al.* [125] gave a recent review on the various computational methods applied to heterogeneous catalysis, as shown in Figure 2.9.

Many chemical reactions in MOFs are performed at mild conditions involving the presence of a solvent. In order to move closer to modeling the system at operating conditions, the solvent in the pores can also be taken into account. This adds a lot of degrees of freedom to the system, and for this reason often an implicit description of the solvent is done, such as in the Polarizable Continuum Model (PCM) [126]. In the case of the work in this thesis, however, it is necessary to fully model the solvent molecules, as they are actively involved in proton transfers. To do so, the number of solvent molecules that can fit in the unit cell needs to be estimated. Monte Carlo method (MC) is an alternative approach to MD to explore the PES for complex systems. It was initially developed for the calculation of multidimensional integrals and is nowadays largely used in chemistry, especially when dealing with adsorption. In the framework of this thesis, it has been applied in the Grand Canonical ensemble (μVT , fixed chemical potential, volume and temperature) to determine the number of solvent molecules that could fill the pores of the material at realistic conditions [127]. The estimation of this number is not trivial, and depends on many factors such as the defect topology, or the choice of force field model used, as reported by Vandenbrande *et al.* [128].

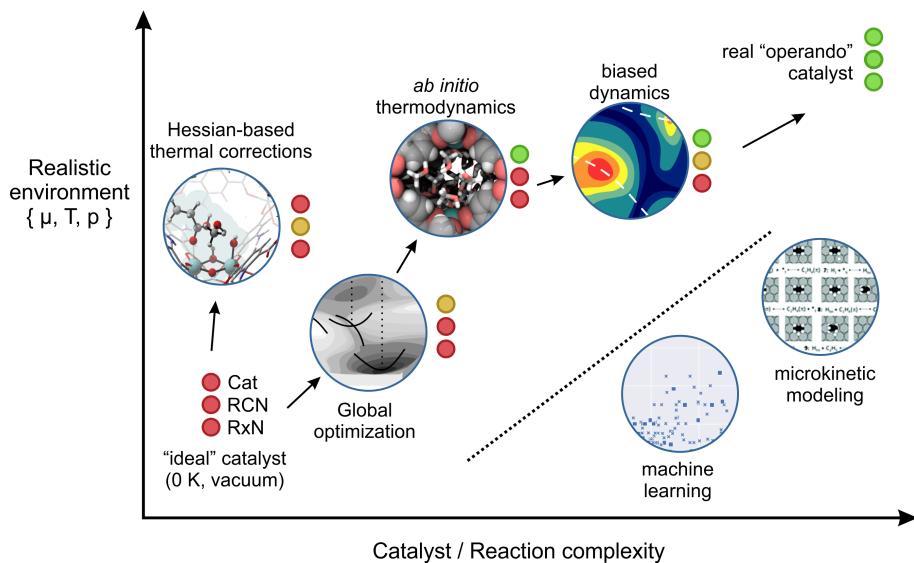


Figure 2.9: Schematic of the various computational methods applied to heterogeneous catalysis, which lie between an idealized ultra-high vacuum/0 K model and a realistic, *operando* model. The traffic light key depicts the quality of each method with respect to catalyst model complexity (Cat), reaction coordinate accuracy (RCN) and reaction network complexity (RxN). Adapted from ref. [125] with permission of the Royal Society of Chemistry.

Enhanced sampling MD methods to describe activated processes

MD simulations can offer valuable insights into the behavior of a chemical system at equilibrium conditions. From these simulations, many properties can be extracted, such as free energy differences, equilibrium geometries, vibrational spectra, diffusion coefficients, structural parameters etc. Configurations associated to higher (or lower) values of potential energy will be sampled for shorter (or longer) times, and in principle, if a certain process is sufficiently sampled, based on the ergodic theorem we can know its equilibrium constant, and in turn the free energy barrier associated to it. In general, to have reliable free energy for a given process, three components are needed: (i) an appropriate description of the potential energy, (ii) a sampling protocol that allows to generate an ensemble of configurations that is representative for the process, and (iii) the right method to estimate the relevant free energy differences [129]. Point (i) was already tackled in previous sections, and herein we focus on the last two points.

Chemical reactions, where bonds are broken and formed, are generally rare events that will not be sampled with a regular exploration of the PES. If the free energy barrier is high compared to $k_B T$, the probability that such event would spontaneously occur during the simulation time is practically none. This is especially true for complex molecular systems, in which processes occur in different time scales and that can only be simulated for a limited amount of time. For this reason, different enhanced sampling techniques have been developed to enhance the sampling of low probability regions of the free energy landscape [130–137]. Recently, enhanced sampling techniques have been successfully applied in heterogeneous catalysis to study processes at operating conditions [118, 138–143].

There are two main classes of methods that can be used to explore rare events on the free energy surface. The first class is characterized by methods that encompass all degrees of freedom and do not need any prior information on the free energy landscape between two points. Examples of these methods are Transition Path Sampling (TPS) [144] and Replica exchange (RE) [145]. The other class of techniques encompasses methods in which the sampling is enhanced along certain coordinates of the system, such as Umbrella Sampling (US) [146, 147] and Metadynamics (MTD) [131], shown in Figure 2.10.

Choice of collective variable

The PES is a highly dimensional surface, defined by the positions of all atoms in the system. However, often a reactive process can be described by few important coordinates called “collective variables” (CVs), which are projections of the high dimensional space. In fact, what can be considered a chemical configuration is an ensemble of microstates that are different in terms of absolute coordinates of each atom, but all contribute to the same macrostate. In some cases, a simple CV

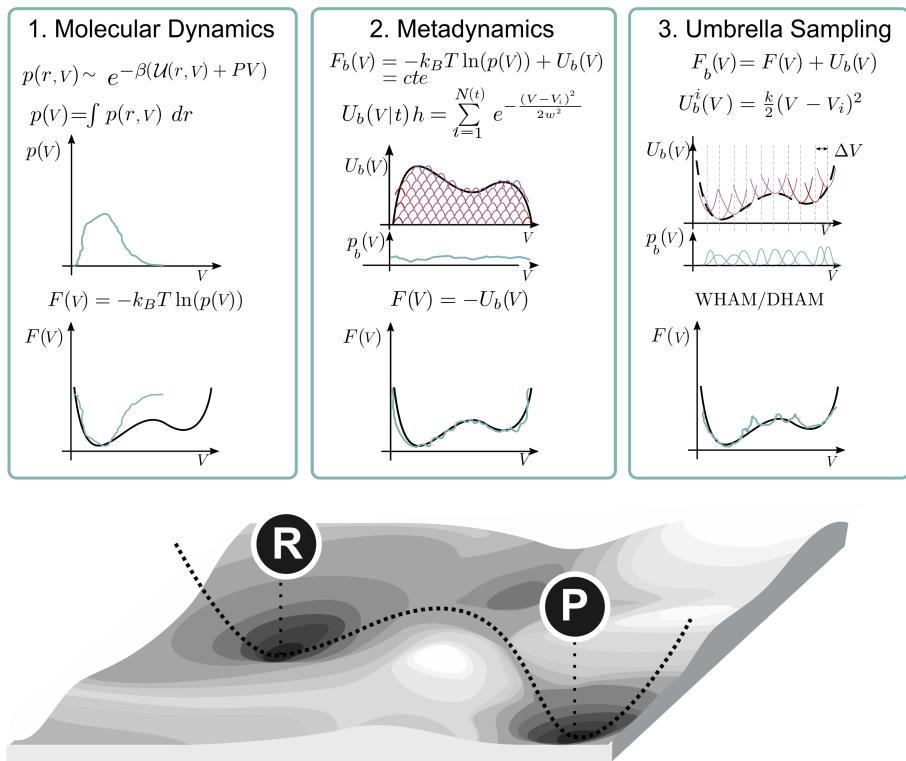


Figure 2.10: Schematic representation of different MD techniques that can be used to explore the PES. Adapted from Ref. [129] with permission of the American Chemical Society

can represent the reaction coordinate for the process, but often the choice is not trivial [148]. In general, choosing the right collective variable is crucial to describe the correct process. Often geometric parameters are used, such as distances, angles, dihedrals, or combinations of the latter. When studying certain processes, a switch function such as a coordination number (CN) derived from distance information is often the preferred choice. CNs represent a smart choice compared to distances, because for each pair of atoms, the coordination is zero for distances higher than a certain threshold.

$$\text{CN} = \sum_{i,j} \frac{1 - (r_{ij}/r_0)^{nn}}{1 - (r_{ij}/r_0)^{nd}}$$

Where CN is defined between two sets of atoms i and j , r_{ij} corresponds to the distance between atoms i and j , and r_0 is a threshold distance. The exponential parameters nn and nd define how sharply the function behaves around the value r_0 . In processes such as in the ones of **PAPER V**, we considered the coordination between a zirconium atom and the oxygen atoms of the solvent water. Such collective variable allows to consider the possibility that different water molecules could decoordinate and recoordinate, defining the same macrostate for a given value of the CN. We show that such processes cannot be studied by employing distances as CVs.

Metadynamics

Metadynamics (MTD) is a popular enhanced sampling method, used to overcome barriers in the free energy landscape, which was first proposed by Laio and Parrinello [131, 149]. During MTD, a history dependent bias potential is added in the form of gaussian hills to the PES along a certain CV. This way, potential energy is added to already visited states, allowing to escape local *minima* and explore different regions of the PES. When all states are sampled with equal probability, the free energy profile along the biased CV can be obtained from the added bias potential. As the bias is added, the system is able to evolve along all the other degrees of freedom. For this reason, this method can also provide insight into the mechanisms that occur during the exploration of the different regions of the CV, as has been done in **PAPER V**.

Umbrella sampling

In the umbrella sampling method, a series of biased MD simulations are performed along the chosen CV. In each simulation, a bias potential is added to constrain the system to adopt a specific value of the CV, while it can evolve along all the other degrees of freedom. This way, the whole range of the CV can be explored. As each simulation is independent, this method is highly parallelizable. The free energy profile can then estimated by using different methods to combine the information obtained from the set of simulations, such as the weighted histogram analysis (WHAM) or the multistate Bennett acceptance ratio [146, 150]. Even if the CV is

constrained in each simulation, this method can also offer valuable insight into the evolution of the system at different values of the CV, as has been done in **PAPER III**.

Computational details

In this thesis, cluster calculations were performed with Gaussian '09 [85]. Static periodic calculations were done with the Vienna Ab Initio Simulation Package (VASP) [151–155]. Molecular dynamics simulations were performed with CP2K [156]. GCMC calculations were performed with the RASPA code [157]. Enhanced sampling was applied by means of the PLUMED code [158] along with CP2K. For what concerns calculation time, an extended cluster calculation takes less than 1 node day. A static periodic optimization with frequency calculations on the 2-brick UiO-66 unit cell takes approximately 20–25 node days. A 50 ps CP2K calculation on the same unit cell including solvent takes about 150 node days. The simulations were run on the TIER1 Flemish supercomputer centrum, with access granted on a project basis, for a total of about 20000 node days used for the calculations in this thesis (Appendix B). The computational time indicated refers to nodes consisting of 2x14-core Intel Xeon processors, with 128 GiB RAM.

In summary, this chapter gives an overview of the many methods that have been employed to understand the properties of MOFs. Reactive processes can be studied by using a plethora of different computational techniques that differ in cost and accuracy. Cluster calculations were used for a first estimate of the energies, whereas periodic models were used for all the remaining part of this thesis. The increase in computational time allowed to include more complexity in the model, and to explicitly take into account the role of solvent and structural modifications in the material. During the time frame of this thesis, we moved from a static description of the chemical events towards modeling at operating conditions that allows to better describe and predict real processes.

3

Major research results

This Chapter illustrates the main research results obtained in the framework of this thesis. The main goal of this work was the study of the nature of active sites on UiO–66 and MOF–808 upon activation processes, and how the solvent and the functionalization influenced their behavior. The role of active sites on defective UiO–66 was studied for Fischer esterification of free fatty acids (FFA), where it became clear that solvent played an unexpected active and beneficial role in the reaction mechanism. This study was inspired by experimental work performed by Cirujano *et al.* [79, 80]. Starting from this initial experimentally inspired problem, the level of complexity was increased in our modeling approach to obtain a fundamental understanding on the nature and creation of active sites as operating conditions. Different molecular modeling techniques based on static and dynamic methods have been applied to gain insight into the interaction between active sites and reactants in the material, as has been introduced in Chapter 2. Contrary to reactions in zeolites, processes in MOFs are often performed ad mild conditions in the presence of a solvent, which adds complexity to the model. So far, solvent in MOFs had been studied only with force–field approaches, but with the increase in computational power it became possible to include a full *ab initio* treatment of water and methanol solvent in the UiO–66 pores and to go towards an *operando* description of activation processes in MOFs, where realistic experimental conditions are taken into account. The most important scientific results will be highlighted in this Chapter. More details are to be found in the original articles, enclosed in Part II.

3.1 Activation processes in zirconium–MOFs

One of the main challenges in MOF research is the understanding of how active sites are created and how they impact the properties of the material. For this purpose, molecular modeling offers a platform that allows to study such activation processes and nature of active sites at the molecular level. In this sense, UiO–66, characterized by an exceptional stability, represents a perfect case study where different activation and PSM processes can take place without disrupting the stability of the structure. Initially, it was not clear how active sites were created in UiO–66, as the non-defective material contains only fully saturated sites. It was later understood from experimental and computational evidence [65] that defects, such as missing linkers, are necessary for the generation of active sites for catalysis.

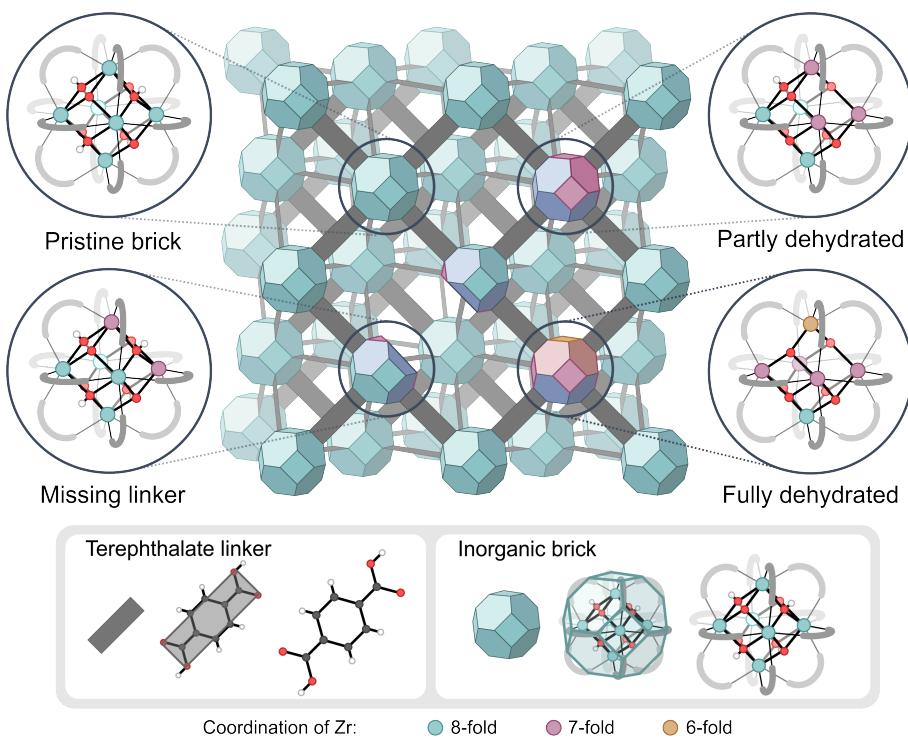


Figure 3.1: Schematic representation of the UiO-66 structure with possible configurations of the bricks that give rise to coordinatively unsaturated Zr atoms. The colors indicate the coordination of the Zr atoms. Adapted from ref. [159] with permission of the Royal Society of Chemistry.

Molecular structure of SBUs with missing linkers

In **PAPER I** we investigated the local defect topology upon removal of a linker on UiO-66 as starting point to investigate the catalytic role of defect coordinating species in the mechanism of Fischer esterification. This research was done in collaboration with the group of Dr. Francesc X. Llabrés i Xamena (Instituto de Tecnología Química, Universitat Politècnica de València), where they experimentally observed that the thermal treatment of the material had a strong effect on its catalytic behavior for Fischer esterification, but the reason was not well understood. When studying the catalytic behaviour of UiO-66, it is essential to understand the reactivity of the inorganic SBU and its molecular structure upon removal of a linker. On these sites, defect coordinating species can be adsorbed and have an impact on the catalytic properties of the MOF by introducing additional sites. Upon removal of one of the twelve negatively charged BDC linkers from the inorganic $Zr_6O_4(OH)_4$ SBU, a positive charge remains on the brick. Charge neutralization can be accomplished by either coordinating a negative ion such as hydroxyl group to one of the zirconium atoms, or by removing a proton from the brick. This latter case is characterized by two zirconium open metal sites, and is the type of active site that is obtained upon thermal treatment of the brick at $T > 423$ K. Initial studies on the catalytic properties of UiO-66 mainly pointed towards the Lewis sites as active sites for catalysis. However more recently it has been shown how Brønsted sites in the neighborhood of the Lewis sites may play also an active role for particular reactions [37, 64, 66, 68, 69, 71, 160, 161]. The presence of multiple sites makes it difficult to establish a simple structure–activity relation. In particular, it is important to understand from a mechanistic point of view how the presence of defect coordinating species may affect the catalytic activity on Zr-MOFs. The capping of these defect sites with various labile species was studied within the framework of **PAPER I** and the main results are shown in Figure 3.2.

The coordination of water species near the active sites can occur with different configuration, as schematically shown in Figure 3.2. In all structures the presence of possible Lewis and Brønsted sites which may play a role in reactions is highlighted. The simplest case taken as reference configuration is the unsaturated site obtained upon thermal activation, containing two μ_3 -oxygens bridging the zirconium atoms which may act as Brønsted sites. From this structure, the coordination of one physisorbed water molecule to one of the zirconium atoms is energetically favorable. A decrease in energy is observed when the molecule is deprotonated to the oxo atom as in configuration 1'. The chemisorption of this water molecule shields the Lewis character of the zirconium atom, but introduces an additional Brønsted site in proximity to the open metal site.

The most stable configurations are observed in presence of 2 or 3 adsorbed water molecules on the adjacent zirconium atoms. Two physisorbed water molecules of configuration 2' can be adsorbed to the two zirconium atoms, followed by an immediate dissociation of one of the two molecules into a hydroxyl group and a

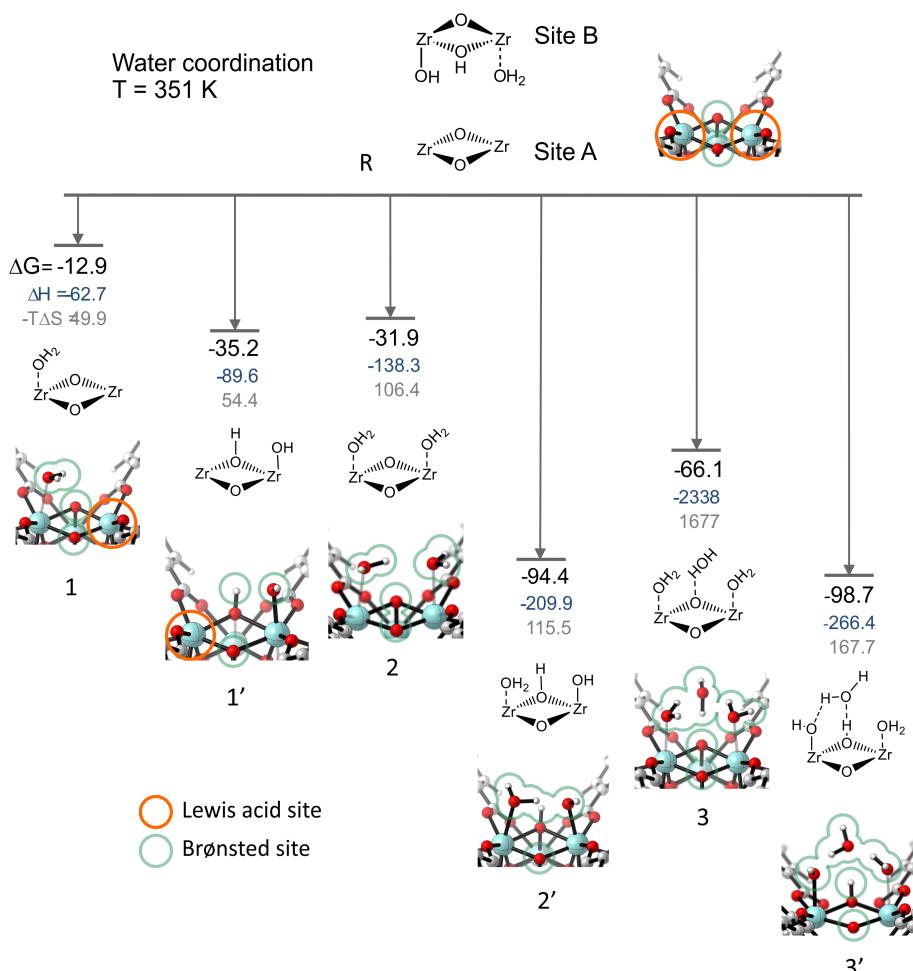


Figure 3.2: Coordination free energies at reaction temperature of 351 K of one, two and three water molecules at coordinatively unsaturated Zr-bricks in defective UiO-66 with respect to a water coordination free site (site R). The structure of the opposite site B corresponds with configuration 2' with two water molecules and consistently used in all periodic calculations considered in the figure. Free energies (in black) are given in kJ/mol, and their decomposition into enthalpic ΔH (blue) and entropic $-T\Delta S$ (grey) contributions. Energies are resulting from periodic calculations with PBE-D3 level of theory. In each configuration Lewis acid and Brønsted sites are indicated. Adapted from ref. [86] with permission of Elsevier.

proton on the adjacent μ_3 -oxygen. This configuration is characterized by a free energy difference of -94.4 kJ/mol at reaction temperature of 351 K compared to the dehydrated defective site. This is in agreement with previous SXRD results by Lillerud *et al.* [66], which show that in the defective material zirconium atoms are fully coordinated by oxygen. When starting from the thermally activated material that contains open Lewis sites, at standard condition, water present in the atmosphere will immediately coordinate to restore the 8-fold coordination of the zirconium atoms to give structure 2. This structure is consistent with the one proposed by the group of Farha [71] who identified three types of protons from potentiometric titration: μ_3 -OH, Zr-OH and Zr-OH₂.

A third bridging hydroxyl species as charge neutralizing species was proposed by Yaghi [67] from XRD data. They propose two physisorbed water molecules bridged by an OH⁻ counterion that is stabilized by a hydrogen-bond interaction with the neighboring μ_3 -OH of the brick. However, a recent study Ling and Slater [68] did not succeed in finding a corresponding minimum on the PES. In this work, we observe that the μ_3 -OH atom immediately deprotonates in proximity of such OH⁻ anion (configuration 3). Similarly as in the previous case, this structure can further stabilized by a deprotonation of one of the two physisorbed water molecules, to yield configuration 3' of Figure 3.2, in agreement with a previous report by Vandichel *et al.* [69]. Adsorption of reactants involved on the Fischer esterification reaction was also taken into account. Similar considerations on the deprotonation processes can be drawn when considering methanol instead of water as reported more in detail in **PAPER I** and **PAPER II**. The energies obtained clearly demonstrate that water molecules preferentially adsorb on the zirconium atoms and that a subtle balance between enthalpic stabilization and entropic penalty will determine the number of water molecules adsorbed on the inorganic brick. We here find that the inorganic brick will tend to stabilize itself with capping molecules to obtain the optimal coordination number of 8. However it must be emphasized that some of these capping species will have to be removed at the onset of the reaction to make place for the active reagents.

Nature of active sites for Fischer esterification explored by static simulation methods

Among the reactions that can be catalyzed by UiO-66, Fischer esterification, shown in Figure 3.3 is an important process in the production of biodiesel, a biofuel that is obtained from renewable sources, such as oils and animal fats [1, 2]. UiO-66 has been shown to be a stable and reusable Lewis catalyst with high conversion rate for the reaction [79, 80]. Experimental findings performed on both hydrated and dehydrated UiO-66 show that water has a beneficial role in the process, but a theoretical rationalization of the underlying causes was missing. Moreover, amino functionalization was shown to increase the reaction rate. To address these questions, the role of active sites on UiO-66 during the Fischer

esterification reaction was studied in **PAPER I**. Two possible lowest activated reaction pathways were identified for the hydrated and dehydrated active site.

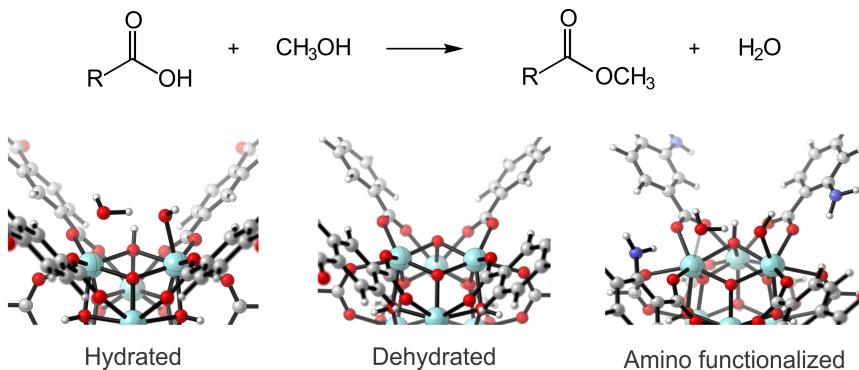
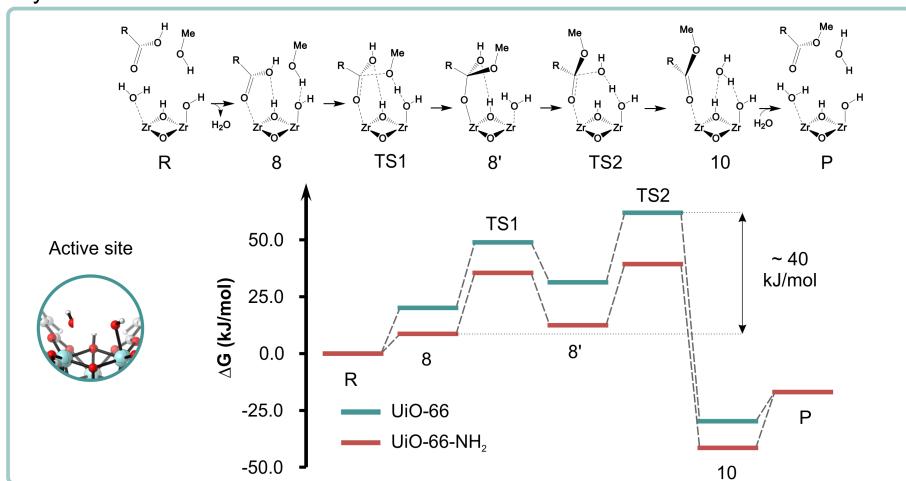


Figure 3.3: Top: Fischer esterification reaction; bottom: three different types of UiO-66 active sites.

The proposed reaction mechanism is shown in Figure 3.4 (top). From the reactant configuration, the physisorbed water molecule is displaced by the carboxylic acid that coordinates on the zirconium atom. The resulting configuration was identified after a series of static calculations probing possible geometries. In this configuration, the acid carbonyl group is bonded to the zirconium, and at the same time methanol is hydrogen bonded to the hydroxyl group coordinated to zirconium. The adsorption of the acid on the Lewis acid site gives to the carboxylic carbon a more electrophilic character, making it more prone to interact with the alcohol. The oxygen of methanol is at the same time made more nucleophilic due to the hydrogen–bonding interaction with the Brønsted basic site situated in close proximity. This favors the condensation between activated carboxylic carbon of the acid and methanol. Two TS are involved in the process, in which first a tetrahedral intermediate is formed, and then water is removed. In both, the hydroxyl group plays a role, first as proton acceptor, then as proton donor, while the acid maintains the bond with the carbonyl oxygen. The low energy barriers associated to the two TS are 28.9 and 30.6 kJ/mol at 351 K. The reaction can proceed both ways, until an equilibrium is reached. This mechanism is characterized by a dual participation of Brønsted and Lewis sites, and overlays with the experimental findings.

Upon thermal treatment at 423 K, UiO-66 loses the adsorbed solvent molecules without compromising the structure of the inorganic SBU, contrary to the dehydroxylation with release of two water molecule that takes part at $T > 523$ K [162]. In principle, these open metal sites should be more catalytically active, but a decrease in catalytic activity is experimentally observed. In the proposed mechanism, shown in Figure 3.4 (bottom), two Lewis acid sites, the zirconium atoms, and one Brønsted basic site, the μ_3 -oxygen, play an active role in the reaction. This mechanism is

Hydrated material



Dehydrated material

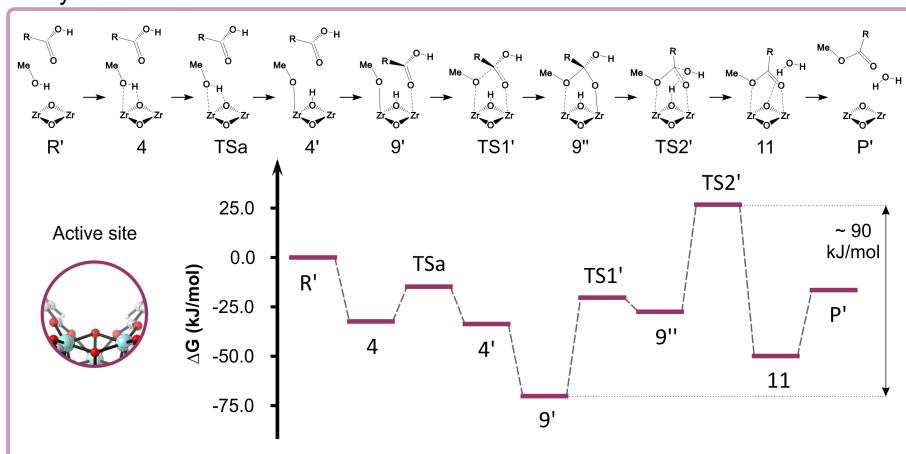


Figure 3.4: Mechanism and free energy profile for the esterification of propionic acid with methanol on a hydrated and defective UiO-66 material (blue), a hydrated defective UiO-66 material with amino functionalization of the BDC linkers (red), and on a dehydrated defective UiO-66 (black). Periodic calculations at B3LYP-D3//PBE-D3 level of theory, T=351 K. R corresponds with an empty frame with one linker defect and a pool with all reactants to guarantee mass balance. In P the defective Zr-brick is coordinated with two water molecules (configuration 2'). P' corresponds to the empty frame with the ester as final product and remaining water molecules in gas phase. Adapted from ref. [86] with permission of Elsevier.

characterized by three TS, in which: 1) methanol is deprotonated to the oxo atom, 2) an adduct is formed between the electrophilic carbon and the oxygen of methanol 3) the $\mu_3\text{-OH}$ group deprotonates to form water. This reaction is characterized by higher activation barriers (a total barrier of 90 kJ/mol at 351 K), which explain the lower catalytic activity of the dehydrated material.

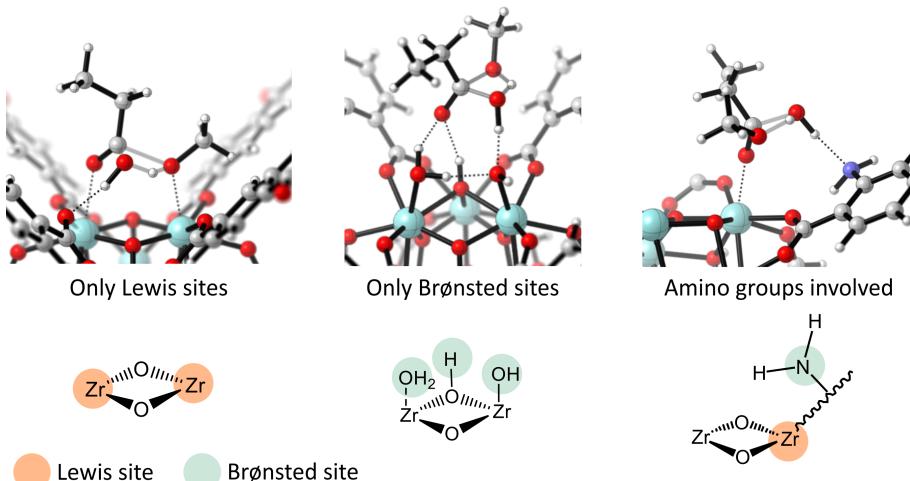


Figure 3.5: Transition states belonging to three cases where no suitable mechanism for Fischer esterification was found, suggesting that the reaction needs a concerted participation of Lewis and Brønsted sites

Other mechanisms that do not make use of either Lewis or Brønsted sites were investigated without success (Figure 3.5), as the energy barriers were too high to be likely to occur. Both proposed mechanisms are characterized by a dual participation of Lewis and Brønsted sites that work complementary to each other. The presence of acid and basic centers within molecular distances has been shown to be essential in the performance of the catalyst during Fischer esterification, as they cooperate in a concerted way during the chemical transformation. Most of the previous mechanistic studies on the UiO-66 material merely focused on the Lewis acidity of the undercoordinated sites, but it has become more and more clear that the bifunctional nature of the UiO-66 catalyst will play an important role in its future applications.

The proposed mechanisms overlay with experimental findings showing that the reaction rate decreases when the material undergoes thermal treatment, and that defective sites are the source of the catalytic activity. The effect of functionalization on the reaction barriers for this reaction will be further explored later in the thesis.

Active sites in larger defective unit cells

Understanding the nature of active sites on UiO–66 is not only important for reactions, but also for processes such as PSLE, where ligands are dynamically exchanged. For this reason, the adsorption of species present in the reaction environment of PSLE process was studied in the framework of **PAPER IV**. This work was done in collaboration with the group of Prof. Rob Ameloot (Centre for Surface Chemistry and Catalysis, University of Leuven), where experimental evidence was given of the active role of methanol in modulating the exchange of linkers and the number of missing linker defects during the process.

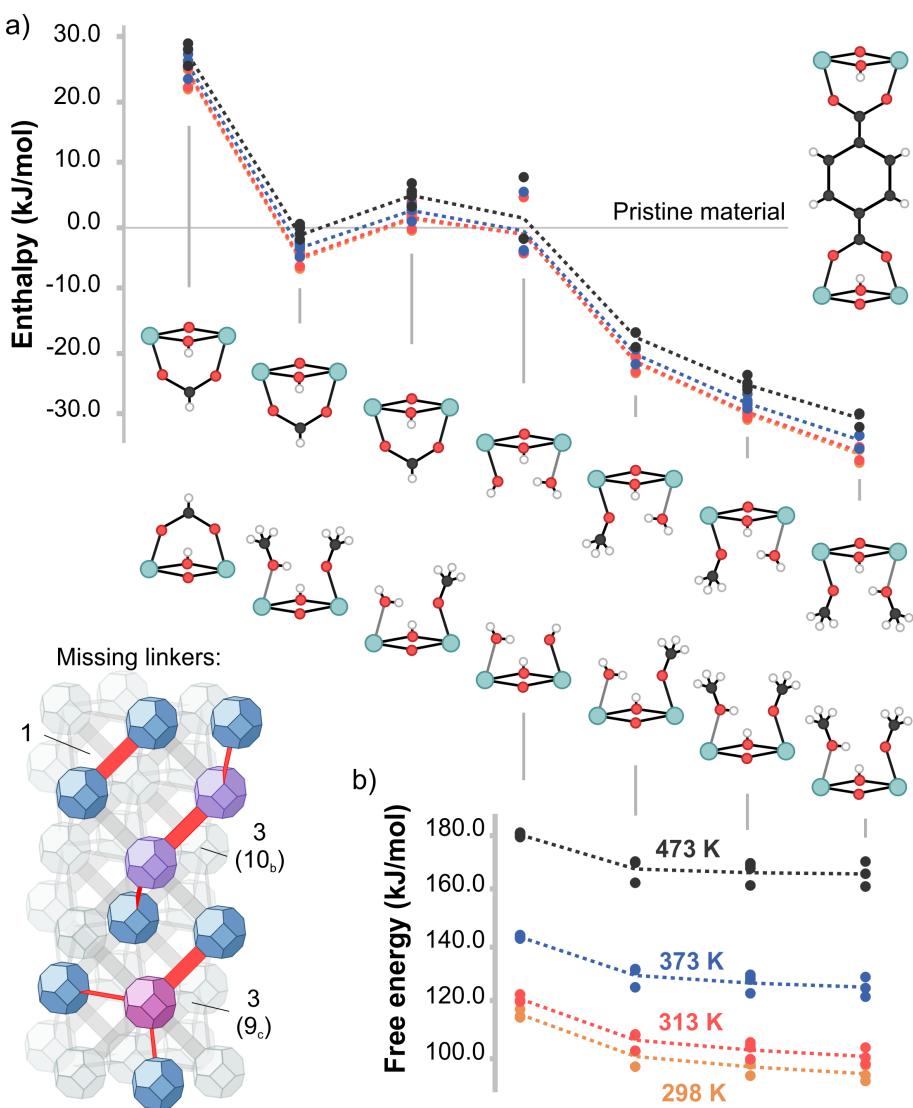


Figure 3.6: Energy diagrams for defective UiO-66 unit cells. Each dot represents a possible distribution of missing-linker defects (1 or 3 in total) within the unit cell; the connecting dotted line represents a weighted average. Values are normalized by the number of missing linkers in the unit cell. (a) Enthalpy difference between defect sites capped in different ways versus the non-defective material at $T = 298, 313, 373, 473\text{ K}$. (b) Temperature-dependence of the free energy difference of the defective structures indicated above (capped with $\text{H}_2\text{O}/\text{OH}^-$, $\text{H}_2\text{O}/\text{MeO}^-$ and MeOH/MeO^-) versus the non-defective structure. A representation of the clusters with different missing linker connectivities is also provided. Adapted from ref. [89] with permission of the American Chemical Society.

To rationalize the experimental findings, the first step was to see how different species present in solution would interact with the UiO-66 material at different defect concentration. To obtain this information, we made use of a larger unit cell, to be able to represent different amount of missing linker defects [60, 61, 63]. This increase in complexity of the model entails a higher computational cost. The defective structures taken into account in the modeling are obtained by removal of linkers from the conventional unit cell containing four inorganic $\text{Zr}_6(\text{O})_4(\text{OH})_4$ bricks [46]. Different numbers of missing linkers were modeled, to represent the different amount of defects in the experimental samples. A case with a low amount of defects is modeled by a unit cell with one missing linker. In this structure, two bricks are 12-fold coordinated, and two bricks are 11-fold coordinated, with an average of 11.5 linkers per brick. A second type of unit cell was taken with three missing linkers, corresponding to a higher amount of defects, with an average of 10.5 linkers per brick. De Vos et al. and Rogge et al. [60, 61] shown in their comprehensive studies that there are multiple topologically diverse possibilities to remove linkers from a 4-brick unit cell. In this work, we chose two possible, discrete cases to represent different coordination of the inorganic brick with this amount of defects. In a first unit cell with three missing linkers, two bricks are 10-fold coordinated and two are 11-fold coordinated. The other represents a more extreme case, with one 9-fold coordinated brick and three 11-fold coordinated bricks. The structures taken into account are denoted as $(10_b, 10_b, 11, 11)_{334}$ and $(9_c, 11, 11, 11)_{333}$ in the work of De Vos et al [61] and are reported in Figure 2.2.

The zirconium atoms on the defect sites have been capped by a variety of species which can be present in the framework during the activation process. In this sense, various combinations were possible by capping with water, hydroxo species, methanol, methoxide and formate. In our models, all zirconium atoms have been capped and are fully (8-fold) coordinated, being the most stable state at experimental conditions. On each defect site there is always a negatively charged species, to compensate for the removal of a carboxylate. When multiple species are adsorbed, configurations with different symmetries can arise, according to how the molecules are positioned on the defective sites. When possible, multiple possibilities for the positioning of these species were taken into account and modeled. The removal of a negative charge can be also compensated by removal of a proton, as obtained in the dehydrated material, with subsequent physisorption of two neutral species, but these configurations are higher in energy, as seen in **PAPER I**. The adsorption enthalpies for the different configurations are reported in Figure 3.6. Our results show a clear enthalpic preference for the methanol/methoxide pair in detriment of formate and water. The defective material synthesized with formate modulator is promptly attacked by methanol species present in solution. The obtained results are in line with a previous report [72] and highlight the preference of zirconium active sites for MeOH/MeO^- substitution as opposed to $\text{H}_2\text{O}/\text{OH}^-$, alternatively assumed as preferential charge balancing element for missing linker defects [67, 68]. No substantial difference in enthalpy is observed between the

unit cells with different amount of defects (energy in each defective structure represented by a dot in Figure 3.6). This is an indication that the thermodynamics that governs the process does not depend on the starting number of missing linkers. Such theoretical finding is in agreement with the experimental observations of **PAPER IV**, where it is shown that an equilibrium defect concentration is reached that does not depend on the initial defectivity.

The effect of temperature on the free energy was also investigated on the relative stability of defective configurations with respect to the pristine unit cell, where the linker is not missing. The first observation is that configurations involving water and methanol are more stable at lower temperature, thus the formation of missing linker defect through ligand exchange is enthalpically driven and favored by lower temperatures. Secondly, at 473 K the energy of the non-defective framework is much lower than the defective cases, which explains why the synthesis of the non-defective UiO–66 is favorable at such high temperature [53].

The findings support the evolution of the ligand coordination on the UiO–66 material during the PSE process in methanol, which can successfully proceed at low temperatures. The PSE mechanism will be illustrated more in detail later in the chapter.

Activation by dehydration

The other process that can lead to activation of the UiO–66 material is the reversible dehydration of the brick performed at $T > 523$ K. The dehydration mechanism may have a decisive effect on certain catalytic reactions, where next to the Lewis acid site also the neighboring Brønsted base or acid site may take a cooperative role in the reaction mechanism. It is still unclear whether this lowering in the zirconium coordination without lowering of the brick connectivity is enough to induce catalytic activity. As shown for citronellal cyclization, missing linkers are required for the reaction to be catalyzed [65].

At elevated temperatures and low pressures, the zirconium core gets dehydrated and rearranged, and two water molecules are subsequently removed from the brick, as displayed in Figure 3.1. The fully dehydrated Zr_6O_6 brick contains undercoordinated sites [28, 48, 73, 162], with coordination of the zirconium atoms ranging from 8 to 6. The structure of UiO–66 is however preserved and the brick can easily be hydrated again with a reversible mechanism, giving evidence that the inorganic SBU can undergo dynamic processes. During these structural rearrangements, the presence of a infrared band related to hydroxy groups was observed by Nishida *et al.* [163] and by Shearer *et al.* [162] pointing towards structural rearrangements of the inorganic bricks. The mechanism of dehydration was first studied by Vandichel *et al.* by means of nudge elastic band calculations [69], showing the presence of loose hydroxyl groups, in agreement with former

experimental reports, as well as partially decoordinated linkers. These findings point towards an intrinsic mobility of the framework structure, that can accommodate rearrangements without disrupting its stability. However, the static study of this process poses serious limitations.

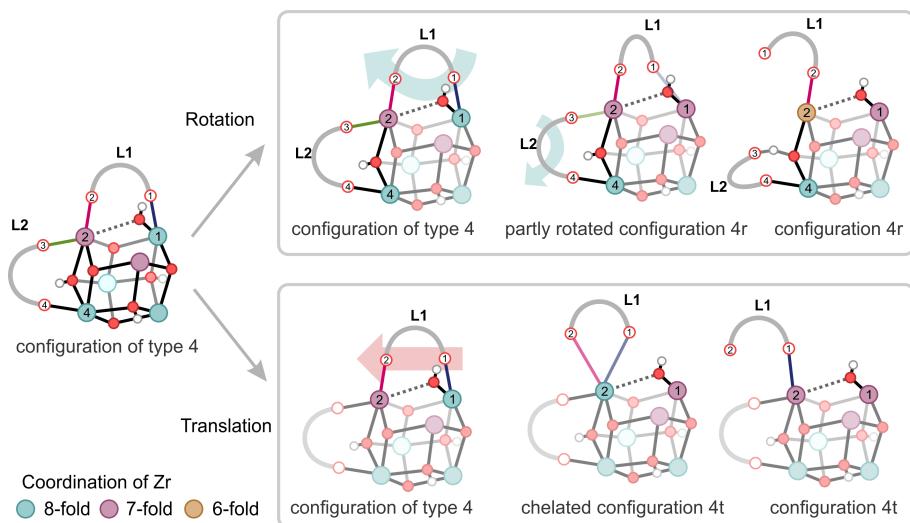


Figure 3.7: Umbrella sampling in two windows of $CV = 1.45$ and 1.51 , showing two distinct motions of the linkers. On the left column, a translation of the linker L1 generates a chelated structure and a subsequent shift in the carboxylic oxygen connected to Zr2 (configuration 4t). On the right column, a rotation of the linker L1 and a partial decoordination of linker L2 forming a hydrogen bond with an μ_3 -OH hydroxyl group is shown (configuration 4r). A proton transfer between the carboxylic oxygen O3 and the bridging μ_3 -O is also observed and is an indication of the occurrence of an intrinsic dynamic acidity. Colors indicate the coordination number of zirconium atoms. Adapted from ref. [159] with permission of the Royal Society of Chemistry.

To go beyond the static approach and follow the dehydration process *operando* and *in-situ*, in **PAPER III**, we follow on the fly the fast dynamic of the UiO-66 material at dehydration temperature of 573 K by means of umbrella sampling simulations. The results show an intrinsic dynamic behavior of the material, with open metal sites being created by continuous changes in the network connectivity due to labile M–L bonds. We identify two types of motions of the linkers, namely translation along the axis connecting the two adjacent zirconium atoms, and rotation along the linker axis connecting the two inorganic SBUs, as shown in Figure 3.7. At the same time, these motions are accompanied by a high mobility of hydroxy groups created by decoordination of the μ_3 -OH groups. These results show that linkers are more mobile than originally anticipated. The high connectivity

between the two SBUs of UiO–66 allows all these reversible rearrangements with activation barriers that can be easily accessible at experimental conditions.

These findings on the dynamic nature of UiO–66 were an eye-opener, as the structure of UiO–66 is regarded as one of the most stable and rigid among the whole MOF family. Herein, we discovered that the UiO–66 material possesses an intrinsic dynamic behavior, where the zirconium atoms have a degree of flexibility in changing their coordination number. This is connected to its high versatility in undergoing other processes where the coordination environment of the brick can change, such as in PSLE.

Thermal activation of MOF–808

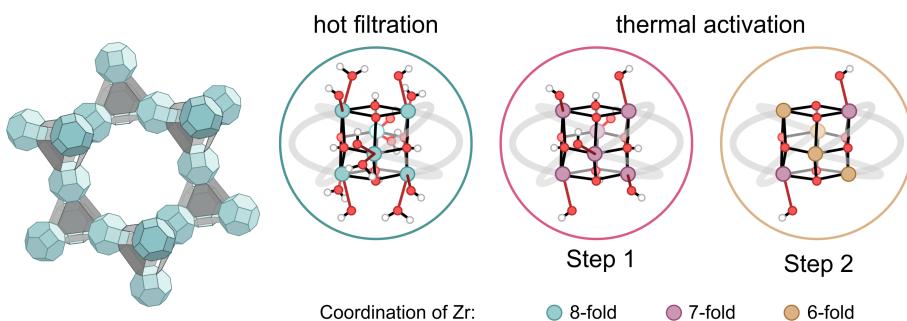


Figure 3.8: Schematic representation of as synthesized and upon activation MOF-808 structures

The active sites and coordination changes upon thermal activation have been studied in the case of MOF–808 in **PAPER VI**. MOF–808 shares the $\text{Zr}_6\text{O}_4\text{OH}_4$ brick with UiO–66, but the brick is connected to only six tritopic BTC linkers, making it the least connected MOF in the Zr-MOF family. The material possesses a high catalytic potential due to the intrinsic presence of defective sites of complex nature, and large pores [44]. For instance, it shows a higher catalytic activity than UiO–66 for MPV reduction [82, 83].

In this work, we wanted to see in how far the material can be activated by thermal treatment to obtain such Lewis acid sites, and whether the conclusions obtained for UiO–66 could be applied also to other MOFs possessing the same inorganic SBU. In particular, we were interested in the stability of the material upon these activation processes.

In the as synthesized material (Figure 3.8), the zirconium atoms that are not connected to BTC linkers are capped by formate groups. The material is activated by hot filtration, whereby each formate is replaced by water and hydroxyl group, for a total of six water molecules and six hydroxyl groups per inorganic SBU [82, 83].

Each pair of adjacent zirconium atoms has a similar configuration as the stable configuration 2 of Figure 3.2, but the presence of multiple Brønsted sites in close proximity gives rise to a more complex nature of the active sites. Experimentally, four different types of protons can be identified in this material [71], each yielding a different acidity.

By thermal activation, similarly to UiO–66, water can be removed from the active sites to open Lewis acid sites for catalysis. From each brick, six physisorbed water molecules can be decoordination. In a second stage, up to four of the hydroxyl groups could in principle be further decoordination by extracting a μ_3 -OH proton from the brick to form water, giving rise to mixed-coordination bricks with 6 and 7-fold coordinated zirconium atoms.

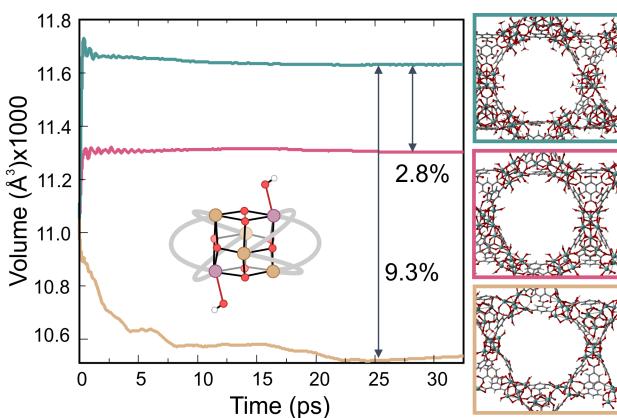


Figure 3.9: Change of volume in time of the three investigated structures with different Zr coordination

In **PAPER VI**, the behavior and stability of the material upon activation processes was investigated by means of a series of independent MD simulations at 300 K at variable unit cell parameters. We show that the dehydration of the inorganic brick, shown in Figure 3.8, substantially affects the stability of the structure. In the hydrated form, the material possesses a high number of Brønsted sites that show dynamic acidity in the form of proton transfer between water and hydroxyl groups that are located in close proximity. This may be important for proton conductivity. Upon thermal treatment, first the physisorbed water is removed leading to a homogeneous distribution of 7-fold coordinated zirconium atoms in the brick. Upon this dehydration, only a slight decrease in the unit cell volume is observed (Figure 3.9, pink curve), and the structure remains stable, even though it possesses a high amount of undercoordinated sites. Moreover, these Lewis sites are located in proximity to Brønsted sites arising from hydroxyl groups, making dehydrated MOF–808 a dual heterogeneous catalyst. Further removal of the hydroxyl groups together with the μ_3 -OH protons causes a collapse of the structure.

Due to the overall low structure connectivity the adsorbed hydroxyl groups have to be considered as inherent part of the framework composition and their removal results in the collapse of the material (Figure 3.9, yellow curve). In contrast to UiO–66, that can undergo reversible dehydration processes where the coordination of the zirconium atoms can be lowered to 6 without disrupting the structure, MOF–808 cannot sustain such decrease in coordination. The physical and chemical properties of UiO–66 cannot be easily extended to MOF–808, even though they share the same inorganic SBU. The lowest possible zirconium coordination is 7. These observations are especially significant in the context of potential applications in heterogeneous acid-base catalyzed reactions.

Linker functionalization on UiO–66

Linker functionalization can play an important role in MOF catalyzed reactions and in modulating the nature of the active sites. For instance, the acidic and basic properties of the active sites can be altered by the presence of electron–donating or electron–withdrawing groups and have an effect on the energy barriers on reactions, such as in Fischer esterification. MOFs can be functionalized by PSE procedures. On UiO–66, functionalization is particularly easy due to its intrinsic dynamic nature and its structural stability. The following results concern linker functionalization on UiO–66 and its effects on the catalytic activity of the material.

Post synthetic linker exchange

As reported in Chapter 1, functionalization in UiO–66 can be induced by PSLE, exploiting the robustness of the framework, which can easily undergo coordination changes. In **PAPER IV**, the mechanism of PSLE was investigated in a dual experimental–computational study performed in collaboration with the group of Prof. Rob Ameloot. The PSLE in UiO–66 that leads to functionalization of the BDC linkers with amino groups was performed in methanol at mild conditions (313 K). Results show that the initial amount of missing linkers did not have an effect on the final composition of the material, pointing towards low energy barriers for the exchange. Moreover, the process was accompanied by an initial lowering of the BET surface area, shown in Figure 3.10, that could be explained with a hindering of the pores by additional adsorbed species.

In light of these experimental observations and the computational findings reported in Figure 3.6, a hypothetical exchange mechanism was modeled starting from both pristine and defective material (Figure 3.11). In the proposed mechanism, linker exchange is initiated by coordination of methanol on the zirconium sites, causing a partial hydrolysis of one of the BDC linkers. This metastable state is characterized by a dangling linker which is stabilized by hydrogen bonds with other carboxylic oxygen atoms and the μ_3 –OH hydrogen from the brick, in a similar fashion as what observed in **PAPER III**. This structure causes a hindrance of the

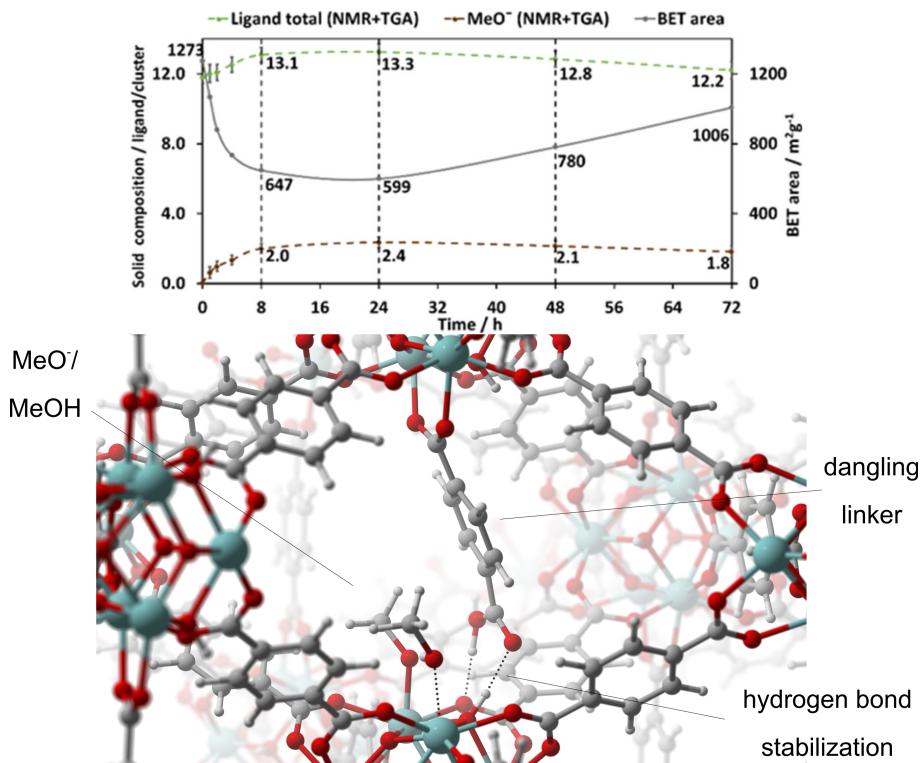
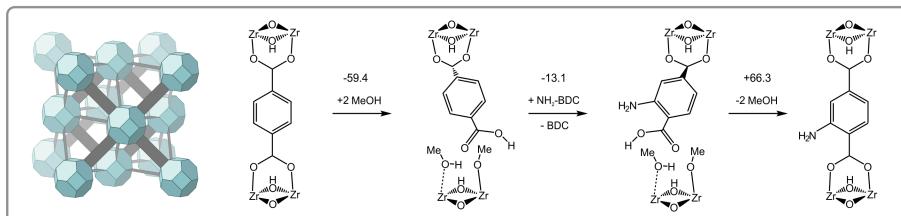


Figure 3.10: Porosity evolution during PSE of UiO-66 Top: MeO^- content in the MOF material during PSE (dashed brown) as determined by NMR and TGA. The total number of ligands per cluster is shown in dashed green. The evolution of BET areas measured for samples collected at selected instances is shown in gray. Bottom: Representation of an energy-optimized UiO-66 structure with both dangling linkers and MeO^-/MeOH pairs. Adapted from ref. [89] with permission of the American Chemical Society.

pore in agreement with the reduction in BET surface area, shown in Figure 3.10. From this configuration, exchange of mono-coordinated BDC linkers with NH₂-BDC quickly ensues, with partial coordination of the new linker. In a following step, methanol is desorbed from the active site. We postulate that after this step the linker can connect to the zirconium atoms, reestablishing the binding between the two bricks. This preferential adsorption of BDC-NH₂ is explained by the lowering of the enthalpy of about 7 kJ/mol per linker. When starting from a defective material (Figure 3.11), a similar mechanism can be proposed. In this second scenario, the synthesized material contains formate on both defective sides. Formate will be substituted by methanol present in solution, and the defect healing can proceed in a similar way as in the previous case. All these rearrangements proceed rather fast, as the final experimental composition of the material does not depend on the initial percentage of functionalized or missing linkers. During the process, the zirconium atoms rapidly change their coordination number, however, giving preference to 8-fold coordinated active sites that enhance the stability of the structure.

Non-defective material:



Defective material:

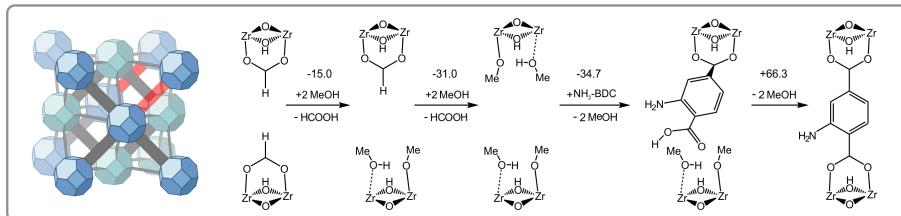


Figure 3.11: Proposed PSE mechanism in non-defective and defective UiO-66. MeOH facilitates ligand exchange through the creation and stabilization of defects. Enthalpy differences are given in kJ/mol at 313 K (PSE temperature). Adapted from ref. [89] with permission of the American Chemical Society.

In conclusion, we showed how molecular modeling can support experimental findings to give insight into the PSE process on UiO-66. The active role of MeOH solvent was revealed, in the creation and stabilization of dangling linker configurations. A fast exchange of these dangling linkers was also observed. We showed that the initial number of defects in the material does not have an effect on the final composition. This points towards low activation energies for the

rearrangements of linkers in the material and to a dynamic behavior of UiO–66. These static results were the onset for further work on the coordination changes of the zirconium atoms, which will be further explored later in this thesis.

Effect of linker functionalization on Fischer esterification in UiO–66

In order to understand the effect of the functionalization of the BDC linkers on the catalytic activity of defective UiO–66, the esterification reaction (Figure 3.4) was investigated also on the amino functionalized UiO–66-NH₂, as it showed higher catalytic activity. Amino functionalization on UiO–66 brings an electron donating group that would in principle lower the strength of the Lewis acid site. However, experimental insights into the Fischer esterification reaction showed that the amino functionalized UiO–66 was more catalytically active than its non-functionalized counterpart. For this reason, it was speculated that amino groups located in proximity to the defect sites would play an active role during the reaction. Our computational results showed that pathways that actively involve the amino groups are highly activated. We therefore studied the reaction following the same pathway that we propose for the hydrated brick (Figure 3.4) in the non-functionalized material.

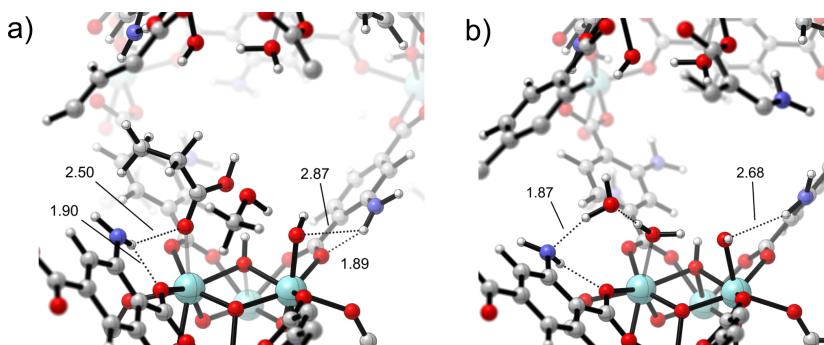


Figure 3.12: Network of hydrogen bonds between acid, methanol, hydroxyl group and amino groups. a) reactive complex 8, b) an additional water molecule present in solution. Key O–H and N–H distances are indicated in Å. Adapted from ref. [86] with permission of Elsevier.

The decrease in Lewis acidity upon amino functionalization is indeed confirmed by the increase of the Zr–O distances of the adsorbates [65]. However, for UiO–66-NH₂, stronger stabilization of the adsorbates is observed, as well as a slight decrease in the overall energy barrier of about 11 kJ/mol, which confirms the experimental findings [79, 80]. The amino groups, although not playing an active role in the reaction mechanism, indirectly modulate the properties of the Lewis and Brønsted sites. A stronger adsorption of the reactants is caused by the formation of a network of hydrogen bonds with the amino groups that cannot be observed

in the pristine material. Amino groups provide additional sites where solvent can form hydrogen bonds, as displayed in Figure 3.12. This indirect positive effect of amino groups on the catalytic properties was also observed by Hajek et al. for aldol condensation [28]. The effect of other functional groups was further analyzed (see Supplementary Material of **PAPER I**). Electron-withdrawing substituents such as $-NO_2$ did not decrease the energy barriers, although increasing the Lewis acidity of the metal. Once again, this confirms the dual Lewis/Brønsted character of the UiO-66 catalyst, that can be enhanced by the presence of additional sites within molecular distance.

3.2 Role of solvent: towards operating conditions

So far, a simple model to represent the active sites was used, where only the species immediately coordinated to the zirconium atoms were considered. Moreover most conclusions taken so far, were based on static calculations, relying on a discrete number of points on the PES, as shown in Chapter 2. When considering a more realistic reaction environment, solvent may be present in the pores of the material, which has an intrinsic dynamic nature and structure. In robust heterogeneous catalysts such as zeolites, processes take place at high temperatures and reactants are often in gas phase. However, in MOFs reaction conditions are usually milder, and liquid solvent is present. Moreover, coordination bonds are more labile and can be broken by interactions with the solvent, so it is important to consider all these possible interactions. A first level of complexity could be to take into account the solvent using microsolvation, adding a small amount of solvent molecules. However, the findings of the previous section point towards a complex nature of the active sites in the material, where solvent may play an active role in reactive processes and structural modifications. For instance, water was shown to have a beneficial effect on the catalytic performance of Fischer esterification by providing additional Brønsted sites as well as stabilization of intermediates through hydrogen bonding. Moreover, methanol solvent played an active role in solvent assisted PSLE, by allowing to exchange linkers and introduce or repair defective sites in the material in a dynamic way. Moreover, UiO-66 can be partially hydrolyzed [73] while retaining its stability.

Based on this knowledge, in **PAPER II** and **PAPER V** we went beyond the static approach and simple microsolvation approach by including an explicit treatment of the solvent in the pores of the UiO-66 material, as schematically indicated in Figure 3.13. This increase in complexity of the model allows to take into account the processes that can occur at operating conditions, and to represent the full solvent environment within the pores at realistic temperatures and pressures. The following results are work performed in a theoretical collaboration with the group of Evert Jan Meijer (Van 't Hoff Institute for Molecular Sciences, Universiteit Van Amsterdam).

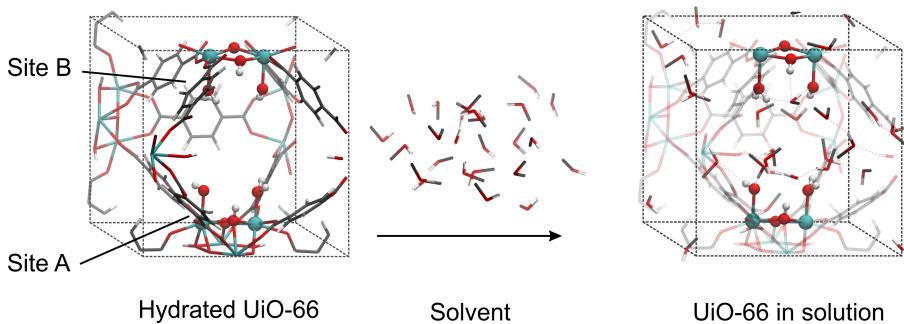


Figure 3.13: Schematic representation of solvent being inserted in the UiO–66 unit cell.
Adapted from ref. [164] with permission of Wiley.

Interaction between UiO–66 and confined solvent

The behavior of a liquid in a confined space, such as in the pores of the UiO–66 material, is substantially different from the bulk. Moreover, UiO–66 possesses both hydrophobic and hydrophilic regions, linkers and bricks, that can interact with a protic solvent in different ways. Molecular simulations allow to investigate how the material responds at the molecular level to the presence of solvent and how the solvent behaves when it is confined in the pores.

To understand how a protic solvent such as water and methanol interacts within the UiO–66 material, in **PAPER II** and **PAPER V**, solvent was inserted in the pores. GCMC simulations were used (see Chapter 2) to estimate the maximum uptake of solvent molecules in the defective hydrated UiO–66 unit cell and the starting molecular configurations. On the defective UiO–66 unit cell, a series of independent *ab initio* MD simulations were performed in which a full loading of methanol and two different loadings of water are included in the pores. In order to have a comparison, the empty material and the bulk solvent were also simulated.

Molecular simulations can offer precious structural insight into the interactions between solvent and material. In particular, vibrational density of states and radial distribution functions can provide information on the hydrogen bond structure and on the structural stability of the material (see Chapter 2). Moreover, unlike in experiments, such analysis from theoretical data can be performed for only part of the system under study, allowing to isolate specific interactions. In this case, for the sake of the analysis of these properties, the system can be divided in two components: material and solvent. We were interested in comparing the properties of the solvated material with the empty one, and the properties of the confined solvent with the bulk solvent. Figure 3.14 displays the vibrational density of states obtained from the power spectrum of the velocity autocorrelation function for the atoms of material and solvent.

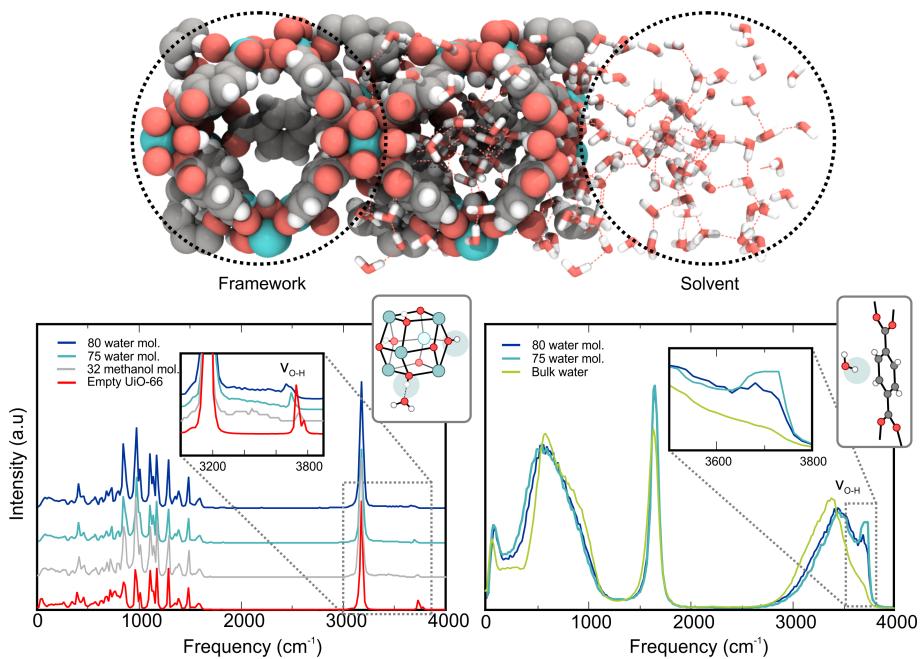


Figure 3.14: Top: schematic representation of the empty pore, pore with the solvent, and confined solvent without the material. Bottom: vibrational density of states obtained from the velocity autocorrelation function power spectra of selected atoms of the simulation. Bottom left: solvated material compared to the empty material. Bottom right: water in the pores compared to bulk water.

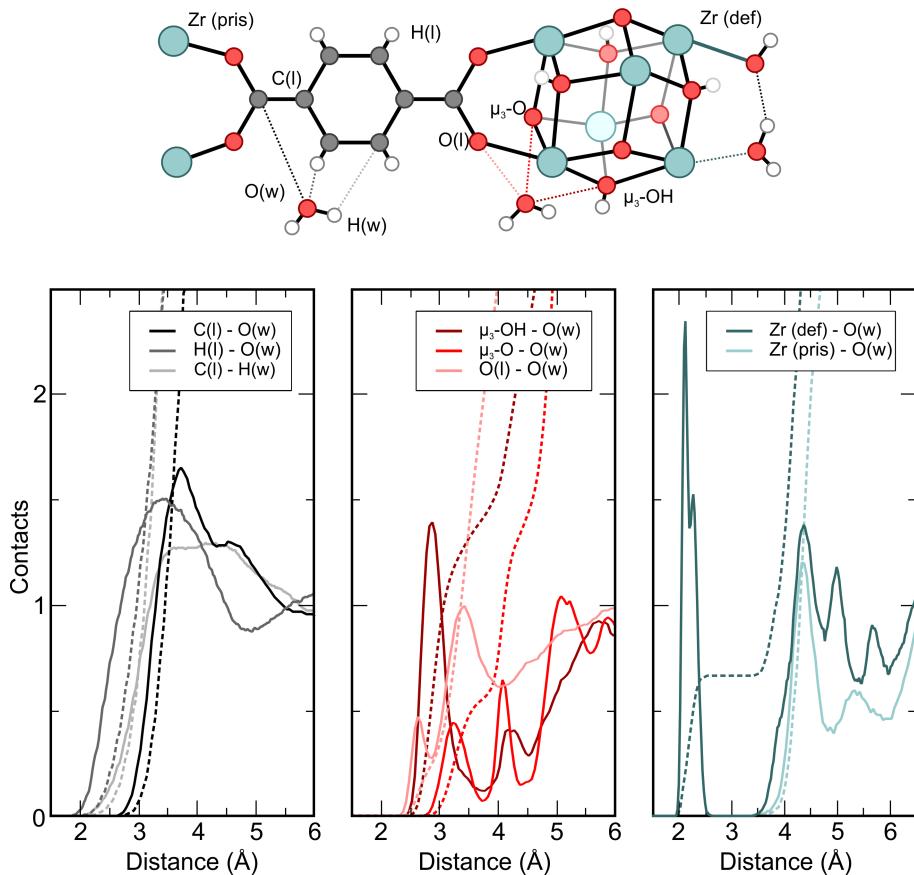


Figure 3.15: Radial distribution functions or pair correlation functions $g(r)$ between oxygen and hydrogen of water ($O(w)$, $H(w)$), and different atoms of the material obtained from the simulation with 80 water molecules in the unit cell. Full lines indicate the $g(r)$, dashed lines indicate its integral. Left panel: RDFs between water and linker carbons and hydrogens ($C(l)$, $H(l)$); middle panel: RDFs between water and oxygen atoms of the linkers and bricks ($O(l)$, μ_3 -OH, μ_3 -O); right panel: RDFs between water and zirconium atoms of defective and pristine bricks ($Zr(def)$, $Zr(pris)$).

The spectrum of the UiO–66 atoms under 3000 cm^{-1} shows no shifts in the peaks between empty and solvated pores, giving insight on the stability of the framework in protic solvents. For the OH stretching at 3750 cm^{-1} , however, there is a broadening when solvent is included. This is due to the strong interactions between μ_3 -OH hydrogens of the bricks and oxygens of solvent, that form hydrogen bonds. To further investigate this, we analysed the vibrational density of states for the solvent and compared it to the bulk solvent simulated in the same unit cell and at the same conditions. We can notice the appearance of a peak at $\nu > 3700\text{ cm}^{-1}$ in the confined solvent, which is due to O–H bonds that do not form hydrogen bonds. This peak is due to the water molecules whose hydrogen atoms are pointed towards the linkers and do not interact with them, in line with previous reports on hydrophobic confinement [165–167].

These results point towards a dual hydrophobic–hydrophilic interaction between solvent and material, where on the one hand, the solvent experiences a hydrophobic confinement due to the interaction with the linkers, on the other it binds strongly to the μ_3 -OH hydrogens of the bricks. To further gain insight into this behavior, we analyzed the RDFs for specific pairs of atoms of material and solvent. The results in case of water are displayed in Figure 3.15. The RDF between carbon atoms of the linkers and water clearly shows a hydrophobic confinement behavior, being nearly zero at distances below 3 Å. On the other hand, a strong interaction is observed between water and oxygen atoms belonging to bricks and linkers, showing formation of a network of hydrogen bonds. The coordination between zirconium atoms and water is also analyzed. In the defective brick, the first peak is due to the coordination between water and zirconium on the defect site.

From this data, we can conclude that solvent does not leave the active site during the simulation time, as the interaction is rather strong, in agreement with the previous static calculations and dynamic results on small models. These simulations give indication of the changes in behavior of the solvent upon confinement and the strong interactions around the bricks and in particular on the active site. The dynamic behavior of the solvent around the active sites will be examined from a mechanistic point of view in the next two sections.

Local structure of the solvent on the defective sites

The local interactions between solvent and defect sites play an important role in reactive processes. It is particularly important to assess the topology of such interactions and what type of phenomena can occur around the active sites at operating conditions. For instance, it is known that protons in defective UiO–66 show a dynamic acidity [68], that is reflected on the high proton conductivity measured experimentally measured in the defective material [70].

In **PAPER II**, we used a multilevel modeling approach to analyze the local behaviour of methanol solvent around the active sites from a mechanistic point of

view. We observe a breakage of the symmetry for the two active sites generated by linker removal. In an arbitrary way, we indicated the two different sites as site A and site B. The site denoted as Site A shows a trigonal network, similar to what reported in static calculations where three molecules were taken into account. This configuration is not broken during the simulation, giving evidence of its stability. Moreover, proton transfers in a similar fashion to what was reported by Ling and Slater [68] are observed between hydroxyl group and physisorbed water, with the bridging methanol molecule shuttling the proton from one site to the other (Figure 3.16). No proton transfers are observed from the $\mu_3\text{-OH}$ proton to the hydrogen–bonded methanol molecule, as this configuration is higher activated. Site B, however, shows a more complex evolution, with chains of hydrogen bonds connecting up to 5–6 methanol molecules and forming closed loops. This evolution is reported in Figure 3.17, where the number of molecules involved in these loops is followed during the simulation time. The system alternates between 4 to 6–membered rings that are anchored on the defect–coordinating water species and the $\mu_3\text{-OH}$ group.

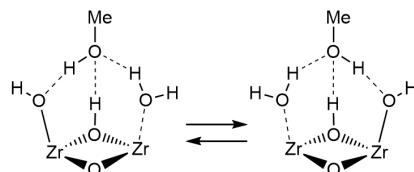


Figure 3.16: Dynamic Brønsted acidity in one of the structures established on the active site in defective UiO-66 and liquid methanol in the pores.

The behaviour is similar to what observed for bulk methanol [168–170], in which a mixture of chain and ring structures formed by six to eight methanol molecules was observed. In this case, the chains are shorter because of confinement, nevertheless, the pore size of defective UiO-66 allows these structures to form. Moreover, open chains of methanol molecules are observed that for part of the simulation connect the two active sites and can in principle provide a way to transfer protons from one site to the other. However, during the simulation time this event is not observed due to the barrier associated to the charge separation.

Processes involving proton transfers may also be important for reactions. Experimental reports show that the network of hydrogen bonds is responsible for a high proton conductivity observed at high temperatures [62]. Moreover, defective sites increase pore size and provide sites where water species are adsorbed and can function as proton donors [70]. To understand how proton transfers occur from a mechanistic point of view, proton mobility in the pores of the material was further analyzed. A charge displacement was induced by artificially removing a proton from the active site and inserting it in different positions in the pore of

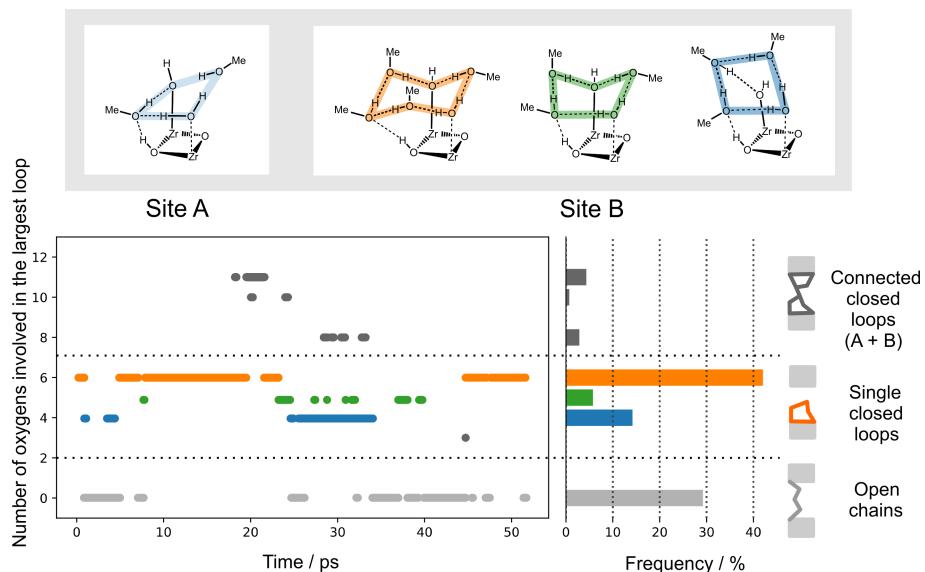


Figure 3.17: Top: Ring configurations observed at site A and site B originating from the interaction between the Zr-bonded hydroxo and water and the solvent molecules. Bottom: Appearance of the various structures during the simulation. The frequency of occurrence of the different structures is also reported. A threshold of 2.2 \AA for the donor-acceptor distance was chosen to determine a hydrogen bond and observations were smoothed over 0.5 ps. Adapted from ref. [164] with permission of Wiley.

the material (Figure 3.18) and the system response was followed. In the methanol solvent, the proton can be either stabilized by solvent molecules and maintain its position in the unit cell, or a Grotthuss charge transfer mechanism can occur in which the charge defect travels through a chain of hydrogen bonds, similarly as what observed by Morrone et al. [170]. In one of the simulations, the proton is transported towards the other active site (Figure 3.18). A configuration is obtained where the original site remains deprotonated and the other is protonated. This configuration is retained for the whole remaining simulation time and is stabilized by the presence of solvent molecules. The zirconium–oxygen distance in the protonated case increases, and the two water molecules physisorbed on the site are more mobile and prone to leave the active site, and a proton transfer could be the initiator to reactant exchanges on the active site. Static periodic calculations were performed to investigate the energy difference related to this charge separation which report a value of 89.4 kJ/mol difference in free energy. Such difference between static and dynamic picture gives indication of the positive role of a protic solvent in stabilizing charged configurations that can be reflected in the stabilization of charged intermediates during catalytic processes. These simulations shed light on the role of Brønsted sites in the material. The $\mu_3\text{OH}$ group is heavily involved in the stabilization of supramolecular structures around the active site, but does not deprotonate to the solvent, contrary to water and hydroxyl groups that show dynamic acidic behavior. These findings show the importance of a solvent beyond being a substrate in the reaction, as it can exchange protons, affect reaction mechanisms, and stabilize intermediates through its remarkable interactions with the active sites. In the next section we will see how solvent can be exchanged on the active sites and induce structural rearrangements in the material.

Activated processes related to dynamic changes in the zirconium coordination number

The interaction between solvent and zirconium atoms on the defective sites investigated in the previous sections is strong, and during regular MD simulations, the coordination is never broken. Previous simulations do not show any change in the zirconium coordination, as such processes are rare events characterized by an activation barrier that does not allow their sampling at normal conditions. However, Zr–O bond breaking is a fundamental event in many processes, such as in the exchange of solvent around the active sites, dehydration, defect formation or PSLE. For this reason, in **PAPER V**, we further focused on the dynamic changes in coordination of the zirconium atoms on the defective sites. We made use of enhanced sampling by means of a series of independent MTD simulations to investigate the coordination changes around the defective zirconium atoms in water solvent.

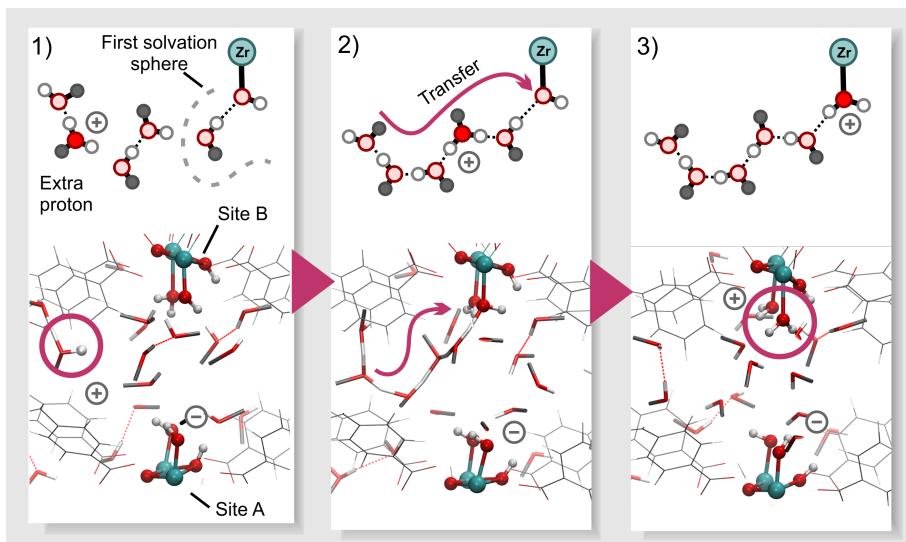


Figure 3.18: Three snapshots of the molecular dynamics simulation which starts from a deprotonated site A and a protonated methanol molecule, with corresponding schematic representation of the process (above). 1) starting structure with protonated solvent 2) a snake-like chain of hydrogen bonds is formed which leads a proton to site B 3) site B is protonated, while site A is missing a proton. Adapted from ref. [164] with permission of Wiley.

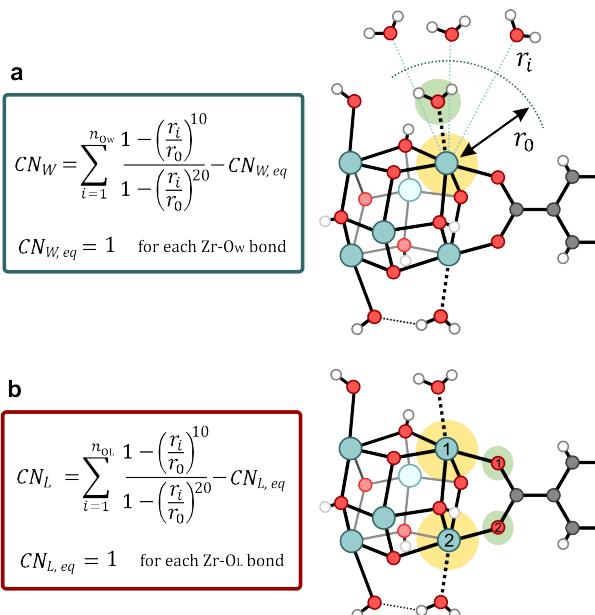


Figure 3.19: Coordination numbers used in the simulation a: coordination number CN_W between zirconium and all water oxygens. Also the linker that shows dynamic movement which induces changes in the zirconium coordination number is visualized. b: coordination number CN_L between each zirconium atom and linker oxygen atoms. n_{OW} and n_{OL} are the number of oxygen atoms considered in the two cases, r_i is the zirconium-oxygen distance, r_0 a cutoff distance of 2.9 Å. In yellow, the zirconium atoms considered in the CN. In green, the oxygen atoms that have a weight close to one and substantially different from zero in the summation.

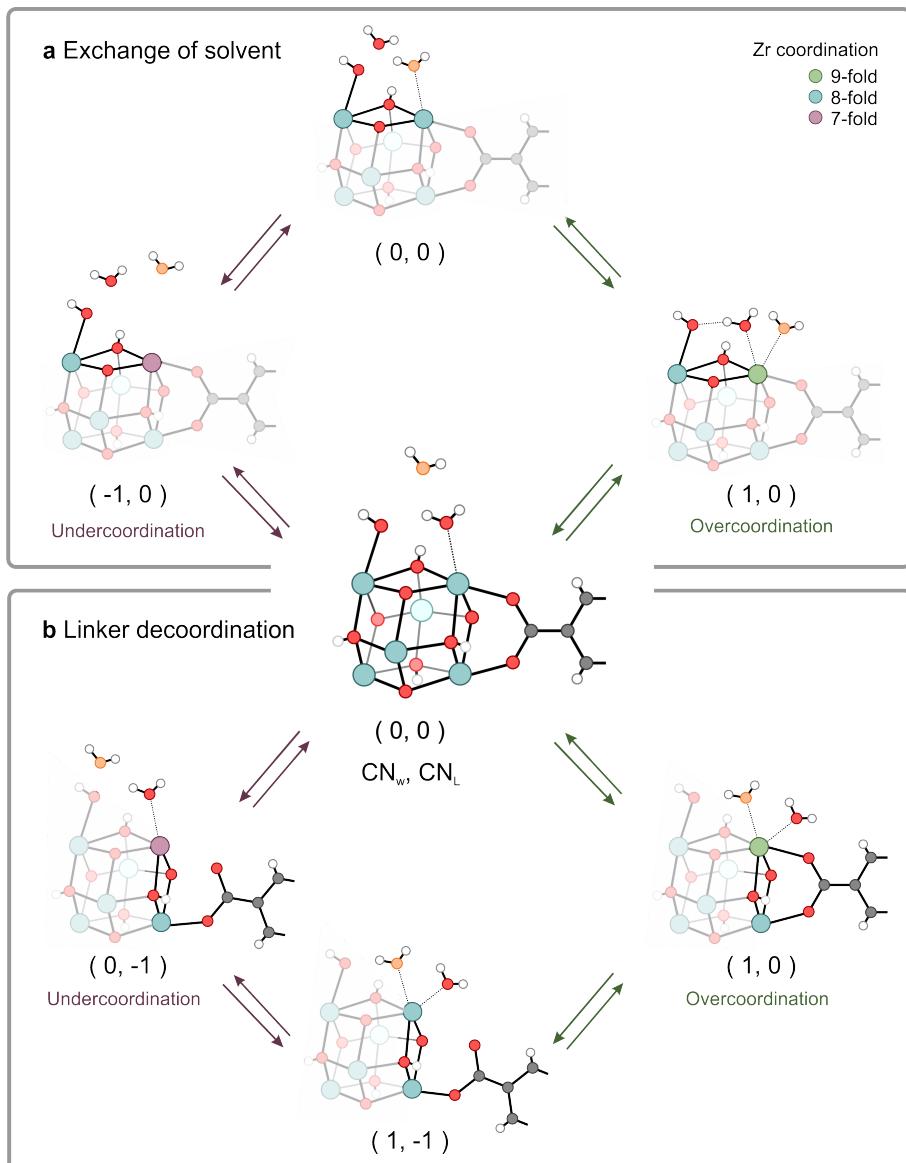


Figure 3.20: Coordination and value of the CV during (a) the exchange of solvent and (b) linker decoordination in the MTD simulations. top paths: stepwise pathway which goes through undercoordinated zirconium; bottom paths: concerted pathways which go through overcoordinated zirconium.

We particularly focus on the changes in coordination of zirconium with the adsorbed water molecules and with the linkers. Undercoordinated and overcoordinated zirconium states can be observed when a solvent is present. Undercoordination is crucial for catalysis, as the Lewis acid site needs to be open for the reactants to adsorb. Overcoordination, on the other hand, can be the trigger for a decoordination of the linker which is a key step in PSE.

To describe these two possible phenomena, two CVs were chosen, as displayed in Figure 3.19: a first one, denoted as CN_W , representing the coordination between zirconium and oxygen atoms of water, and a second one, denoted as CN_L , describing the coordination between zirconium and oxygen atoms of a defect–bridging linker. Both CVs refer to the difference between the coordination state and its equilibrium state. Therefore, starting from an equilibrium value of zero, when a bond is broken, the coordination will decrease by one, describing the undercoordinated case, or increase by one if an additional bond is formed, describing overcoordination. Using these two coordination numbers, it is possible to describe many events taking place around the brick, which are schematically reported in Figure 3.20. In the equilibrated structure, the state defined by the two CNs is (0,0), where the numbers refer to the values of CN_W and CN_L , respectively. Undercoordination and creation of open Lewis acid sites can occur by desorption of water or by breakage of Zr–linker bond to give a zirconium coordination number of 7. Overcoordination is observed when an additional water molecules coordinates to the zirconium atom increasing its coordination number to a value of 9.

In a first case study, the coordination CN_W between one of the defective zirconium atom and the water oxygens was biased. At the same time the CN_L between the same zirconium atom and the oxygen belonging to the defect bridging linker is monitored, as displayed in Figure 3.21. In this simulation, fluctuations of the coordination towards negative values are observed, where the open Lewis acid site is created. Via this undercoordinated pathway, solvent molecules can be exchange on the active site in a stepwise fashion. After this process has been sampled, the system evolves towards an overcoordination of the zirconium atom, to which an additional water molecule is coordinated. From this metastable configuration, the system evolves with structural rearrangements which are reported in the bottom of Figure 3.21. After the second water molecule has entered the coordination sphere of zirconium, it transfers a proton to the neighboring hydroxyl group and pushes the parent water molecule closer to the linker. The water molecule forms a hydrogen bond interaction with the linker which is in turn decoordinated from the zirconium atom, restoring the equilibrium coordination of 8 in the zirconium atom. After these events, the linker recoordinates to the zirconium, to decoordinate again few ps later. In the simulation, we observe a plethora of events such as undercoordination and overcoordination of zirconium, proton transfers, exchange of solvent molecules, and reversible linker decoordination. These fluctuations show how the role of solvent is crucial in inducing such coordination changes and in stabilizing partially

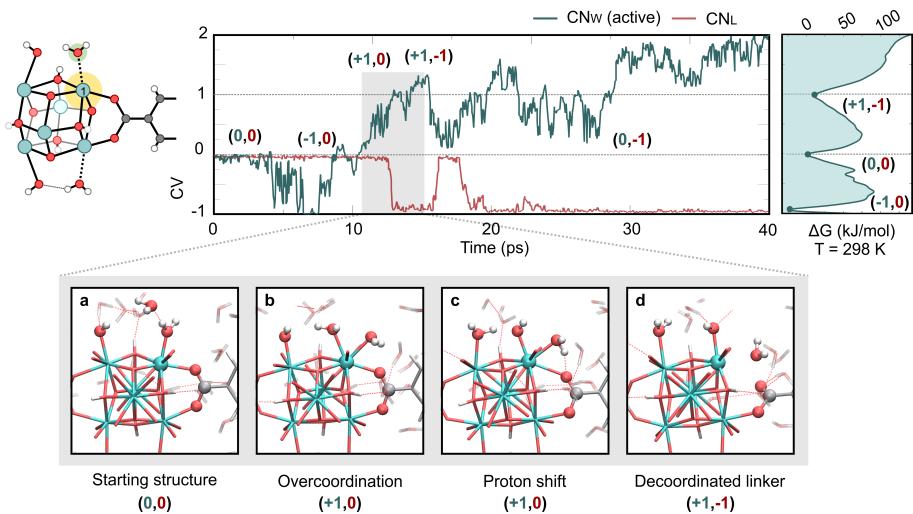


Figure 3.21: Top: evolution of the collective variable CN_W and free energy profile for the first MTD simulation along the CV representing the coordination between the zirconium atom Zr1, highlighted in yellow, and all water molecules (green curve). CN_L between Zr1 and the neighbouring defect-bridging linker oxygen (red curve) is also monitored; bottom: snapshots on the linker decoordination triggered by overcoordination

decoordinated linkers that can lead to structural rearrangements, while maintaining the stability of the material.

In a second case study, we biased the coordination CN_L between one linker and two defective zirconium atoms, and at the same time monitored the coordination CN_W between the latter atoms and solvent water. In this case, we are directly biasing the coordination between zirconium and the linker. The evolution of the collective variable is displayed in Figure 3.22, as well as snapshots that offer mechanistic insights on the different processes. During the simulation, different states are explored, in which the linker is partially decoordinated, can translate to give chelated structures, similarly to what observed in **PAPER III**, or be in a dangling state where both Zr–O bonds are broken, giving an indication of what was postulated in **PAPER IV**. In this case, the linker is decoordinated without intermediate role of solvent, and represents an alternate mechanism by which PSLE, defect creation, or even hydrolysis of the material could occur. This dangling linker scenario is stabilized by hydrogen bonding interactions with the solvent. The process would be completed if two water molecules would diffuse to the zirconium atom and restore the 8-fold coordination, while at the same time one of the two deprotonates to the linker. However this is not observed during the simulation time, as water does not have the time to diffuse, while hindered by the motions of the linker.

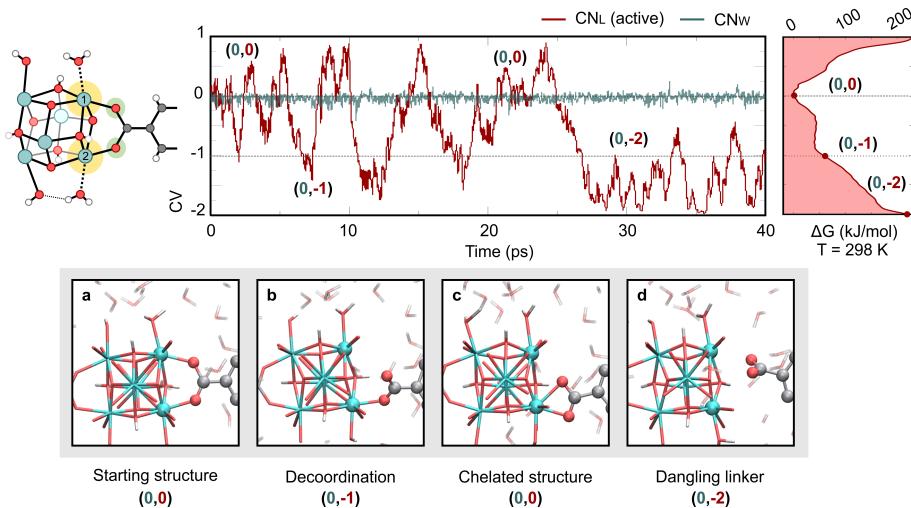


Figure 3.22: Top: free energy profile for the first MTD simulation along the CV representing the coordination between two zirconium atoms and a linker, middle: evolution of the collective variable, bottom: snapshots of the linker decoordination

By means of this series of simulations, we conclude that both solvent exchange around the active sites and structural rearrangements can either occur via a step-wise mechanism where zirconium atoms are undercoordinated or a concerted one, mediated by an overcoordination. The mobility of the linkers observed in **PAPER III** is herein observed without rearrangements of the bricks that are induced at dehydration conditions. In the presence of a solvent, such processes can undergo at much milder conditions, without disrupting the stability of the material. The role of solvent in allowing these processes is crucial, as it provides hydrogen bonds that can stabilize charged configurations, and can itself strongly interact with the zirconium atoms.

4

Conclusions and Perspectives

In this thesis, we shed light on the molecular characterization of the active sites in MOFs by means of different computational techniques. We focused in particular on zirconium based MOFs as they have been used extensively for catalysis. Molecular modeling can offer a rich toolbox to understand the properties of materials and how these properties can be tuned to target specific applications. This is especially true in MOF catalysis, where the active sites are often elusive and arise from disorder in the material that cannot be easily tracked experimentally. We especially focus on UiO-66, which is an example of an extremely stable MOF that due to its exceptional connectivity can be easily tuned and undergo modifications without losing its crystallinity. In close collaboration with experimental partners, we gained insights into the nature of active sites and the interaction of solvents with the inorganic brick, and on the role of complex events that can be observed when modeling processes at operating conditions. To unravel these processes, it was shown that a complementary modeling approach is mandatory. Within this approach, initial understanding of the local structure of the active sites is obtained from static calculations, and subsequently more advanced techniques are necessary to investigate realistic reaction environments. First principles MD simulations in presence of a realistic loading of solvent have been used, as well as enhanced sampling techniques that allow to study events that take place at activated conditions.

The high thermal, mechanical and chemical stability of UiO-66 allows the

material to undergo different activation processes. In this thesis, we focused on the decrease in the zirconium coordination caused by missing linker defects and dehydration. Changes in the zirconium coordination number are necessary to induce catalytic activity of the material.

At the start of this doctoral research work, we investigated the Fischer esterification in close synergy with the experimental group of Francesc Llabrés i Xamena of ITQ (**PAPER I**). This Lewis catalyzed reaction was modeled on UiO–66, initially focusing on the role of the hydration state of the defective sites of the material. We used a series of static calculations, first on small cluster models, then taking into account the periodic MOF structure, with thermal corrections. We show that in the most stable configuration, water is coordinated to the defective zirconium atoms, in line with previous reports. In the proposed reaction mechanism, the chemisorbed water acts as a Brønsted base site that actively takes part in the reaction, along with the zirconium Lewis acid site. It was an eye opener to discover that the catalytic centers are not restricted to Lewis sites in this material. An alternative mechanism was investigated on the dehydrated material, where the reaction proceeds without the assistance of water. We show that in this case, the inorganic brick is actively involved in the reaction, in which zirconium plays a role as Lewis acid site, while $\mu_3\text{-O}$ acts as Brønsted base. However, the beneficial role of water is missing, and the catalytic activity is remarkably decreased due to higher energy barriers, in agreement with the experimental findings. In both proposed mechanisms, UiO–66 acts as dual Lewis/Brønsted catalyst where reactions proceed with a remarkable interplay between the two catalytic centers. The acidic/basic character of protons and oxygens on the brick seems to have a major role, and could be further investigated by pKa calculations. Experimentally by potentiometric titration it was shown that in the defective material three types of protons are present, each possessing different acidity. Proton transfers are a fundamental step in many reactions, and the acidity of Brønsted sites can alter both reaction mechanism and yield. Further theoretical studies could allow to understand the dependency of acidity of various protons of zirconium materials on different conditions and their reactivity.

A remarkable property of UiO–66 is that it can be reversibly dehydrated upon thermal treatment. Dehydration also lowers the coordination number of zirconium up to 6, therefore introducing Lewis functionalities. In **PAPER III**, we followed on the fly the behavior of the material during the activation process of dehydration. We discovered that UiO–66 possesses an intrinsic dynamic nature by means of an exceptional linker mobility. We showed that linkers can decoordinate and recoordinate to the brick without disrupting the stability of the material.

Apart from UiO–66, a series of other zirconium based materials have been synthesized with larger pore sizes and a varying degree of zirconium coordination number. These materials are fully tested and exploited for applications, however it

remains an open question in how far concepts found in the stable UiO–66 are generic for other materials. To this end, some of the information obtained on UiO–66 can in principle be extended to other zirconium–MOFs possessing the same inorganic SBU. However, the behavior of UiO–66 cannot be easily generalized, as the exceptionally high connectivity and the local environment can play a major role in the stability and activity of the material. In **PAPER VI** we further investigated how MOF–808 responds to similar activation processes such as the UiO–66 dehydration. We show that the MOF–808 material possesses an exceptional amount of Brønsted sites that can be created upon hot filtration. By thermal activation, Lewis sites can be created in proximity of the Brønsted sites without disrupting the stability of the structure, offering great potential for future catalytic applications. However, we also observe that further dehydration generates a collapse of the framework and loss of crystallinity. In MOF–808 the coordination of zirconium cannot be decreased below 7 without causing a collapse of the structure.

Another process that can influence the catalytic activity of the material is the introduction of linker functionalization, that can be done by PSE procedures. In **PAPER IV**, we studied the PSE mechanism by means of static periodic calculations at various defect composition and concentration. The work was performed in strong collaboration with the group of Rob Ameloot from KU Leuven. Experimental findings showed the active role of methanol during the process, which can be performed at mild conditions. From the computational and experimental insight, we postulate the presence of dangling linkers as intermediate states during the exchange, that are induced and stabilized by interaction with methanol. These structural rearrangements in the material lead to the replacement of BDC with BDC–NH₂. Moreover, the modified material is characterized by a specific concentration of defects that does not depend on the initial number of missing linkers or nodes. This gives insight on the low energy barriers associated to the process and on the role of methanol in facilitating the exchange.

Experimentally, the presence of amino functionalized linkers that can be introduced by PSE was shown to have a beneficial role in the catalytic performance of the material for different Lewis catalyzed reactions. However, the cause of this improvement was not well understood. For this reason, in **PAPER I** we further focused on the role of BDC–NH₂ functionalization on the catalysis of Fischer esterification. The electron-donating amino groups are shown to decrease the Lewis acidity of the metal centers, therefore should in principle lower the activity. However, we found that amino groups play a positive role in the reaction although not actively taking part in the mechanism. Stronger adsorptions of reactants and lower energy barriers are observed for the functionalized UiO–66 compared to the pristine material. Investigation of the reactants geometries and experimental results pointed towards an increase on water adsorption around the active site due to a stronger network of hydrogen bonds supported by the amino groups. This stabilization could play a positive role in facilitating specific reactant geometries

and supporting metastable intermediate configurations. Theoretical modeling of Fischer esterification performed in **PAPER I** shed light on the crucial role of water in the reaction and on the complex nature of the active sites and their interaction with protic solvents. These results already hint towards the impact of solvent species in the pores of the material on the reaction outcome.

Reactions in MOFs are often performed at mild conditions where reactants are in the liquid phase, which adds further complexity to the model. For this reason, the understanding of the role of solvent close to the active site cannot be neglected. Solvent can have an impact on the creation of active sites, be a substrate, determine the formation of particular isomers, influence the rate and selectivity, affect the reaction mechanism, or even activate/deactivate a specific reaction. Moreover, the M–L bonds in MOFs are more labile than in other heterogenous catalysts composed of covalent bonds. This makes MOFs more prone to structural rearrangements and modifications mediated by protic solvents. In order to move towards an operating description of the processes, it is crucial to understand the behavior of the solvent when confined in the pores of the material.

The role of solvent was investigated in **PAPER II** and **PAPER V**, in which a multilevel modeling approach was employed to study the behavior of defective UiO–66 in presence of a full loading of methanol and water. By means of GCMC simulations the pores of the material were filled with methanol or water solvent, to reproduce the temperature and pressure conditions during a reactive process. The confined solvent behaves differently from the bulk when confined into the pores of the material. Besides pores, UiO–66 possesses both hydrophobic and hydrophilic regions. By performing a series of MD simulations, in **PAPER V** we show that on the one hand, the linkers provide hydrophobic confinement, and on the other, solvent strongly interacts with the inorganic SBUs and in particular with the zirconium atoms on the defective sites.

We further investigated the possible interactions between solvent and active sites in the case of methanol from a mechanistic point of view (**PAPER II**). We observed a remarkable dynamic behavior of the solvent. Supramolecular structures stabilized by hydrogen bonds are formed around the active sites and evolve dynamically during the simulations. Within these active sites, a dynamic behavior of the protons is observed, as anticipated in **PAPER I**. Such proton transfers may play a substantial role in catalytic processes like Fischer esterification, or in structural rearrangements which occur during PSE. Moreover, we show how solvent can transfer protons within the pores and stabilize charged configurations in proximity of the active site. These findings point towards a positive role of protic solvents in reactions involving charged intermediates, that goes beyond simple solvation. These simulations shed light on how the active sites may be modulated via these dynamic interactions. However, regular MD simulations do not allow to track activated processes.

During regular MD simulations, the coordination number of zirconium atoms does not change. Nevertheless, activated processes often involve coordination changes of zirconium metal centers. In order to understand the dynamic interactions between solvent and material on these sites and explore the rare events that can take place, we relied on enhanced sampling simulations. We performed a series of independent MTD simulations where we explore the changes in the coordination of the zirconium at operating conditions. These simulations show that the total coordination number of zirconium can either decrease from 8 to 7 or increase from 8 to 9. Undercoordination is crucial for the creation of Lewis acid sites while overcoordination can trigger linker decoordination. In these two events we observe dynamic exchange of solvent around the active sites, as well as structural rearrangements which are characteristic of PSE processes. The strong interaction between solvent molecules and defective sites can lead to a dynamic behavior of the linker, similarly as what observed in **PAPER III**, but this time at milder conditions, by solvent mediation and without altering the brick. These findings shed light on the dynamic interplay between a protic solvent and the UiO–66 material at operating conditions during PSE. Moreover, one can postulate that these active sites may be dynamically opened for catalysis by temporary linker decoordination, giving a variable Lewis acidity to the metals and more opportunities for Lewis catalyzed reactions. The highly connected UiO–66 material allows all these structural rearrangements where a plethora of active sites that can work in synergy can be generated.

The findings reported in this thesis gave valuable insight into the complexity of the Zr–MOF catalysts, with UiO–66 as main case study. We investigated how the presence of disorder and the interactions with solvent lead to the creation of active sites of different nature that can work in a cooperative fashion during Lewis catalyzed reactions. There is an intriguing interplay between solvent and material, and much work is still to be done in this area of research which is in full exploration. The exceptional chemical versatility of MOFs can be exploited only if we gain enough understanding at the molecular level of their properties at operating conditions. The analysis of the relation between intrinsic properties and desired application have become an important research factor. The use of well-studied model systems such as UiO–66 is therefore crucial to connect theoretical and experimental findings and to understand the sources of errors in the models. The methods used in this thesis reveal molecular level insight into the activation of one of the most stable Metal-Organic Frameworks. They open perspectives to study and benchmark new materials, and further explore the role of active sites and disorder at operating conditions.

In the past years, computational research in MOFs has evolved from providing simple representations of model systems to more complex models that can represent the events that can occur at operating conditions. The growth in computational power will allow to access larger time and length scales, and to make models that

can represent processes at realistic conditions more accurately, especially in complex fields such as heterogeneous catalysis (Figure 4.1). As the modeling field matured, we went from a stage in which it was limited to the rationalization of experimental data to being able to work in synergy with experiments and make new predictions. The further step is the understanding of how computational modeling can help to design active sites and the screening of structural databases to target a specific property, that can then be validated by experiments. The multiple theoretical and experimental tools that have been developed so far complement each other on the investigation of complex phenomena. For this reason, the use of databases in which different computational and experimental results can be stored and accessed will be more and more important in the future of MOF research. Therefore, the publication of data cannot be limited to successful experiments, but has to be extended to failures, which also carry precious information. This way, with more data available, it will ultimately be possible to use statistical models to predict structures that match specific properties. The new theoretical predictions will be able to guide new experiments, that can in turn provide new data, and ultimately lead to materials and conditions that can target specific applications.

Molecular modeling has evolved substantially thanks to ingenious theoretical models and stronger computational power

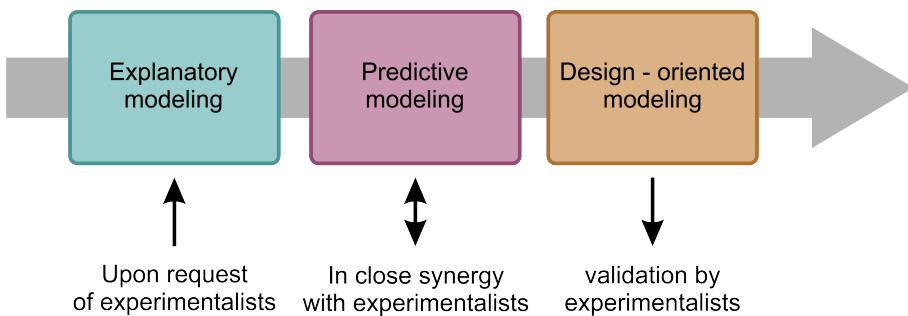


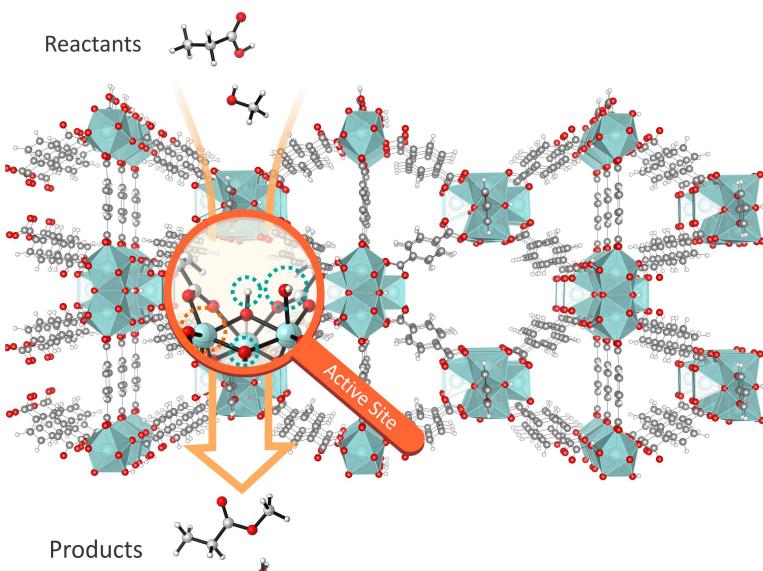
Figure 4.1: Different stages of the close relationship between modeling and experiments.

Part II

Published papers

Paper I

Nature of active sites on UiO–66 and beneficial influence of water in the catalysis of Fischer esterification



C. Caratelli, J. Hajek, F. G. Cirujano, M. Waroquier, F. X. Llabrés i Xamena, V. Van Speybroeck, *Journal of Catalysis*, **352**, 401–414 (2017)

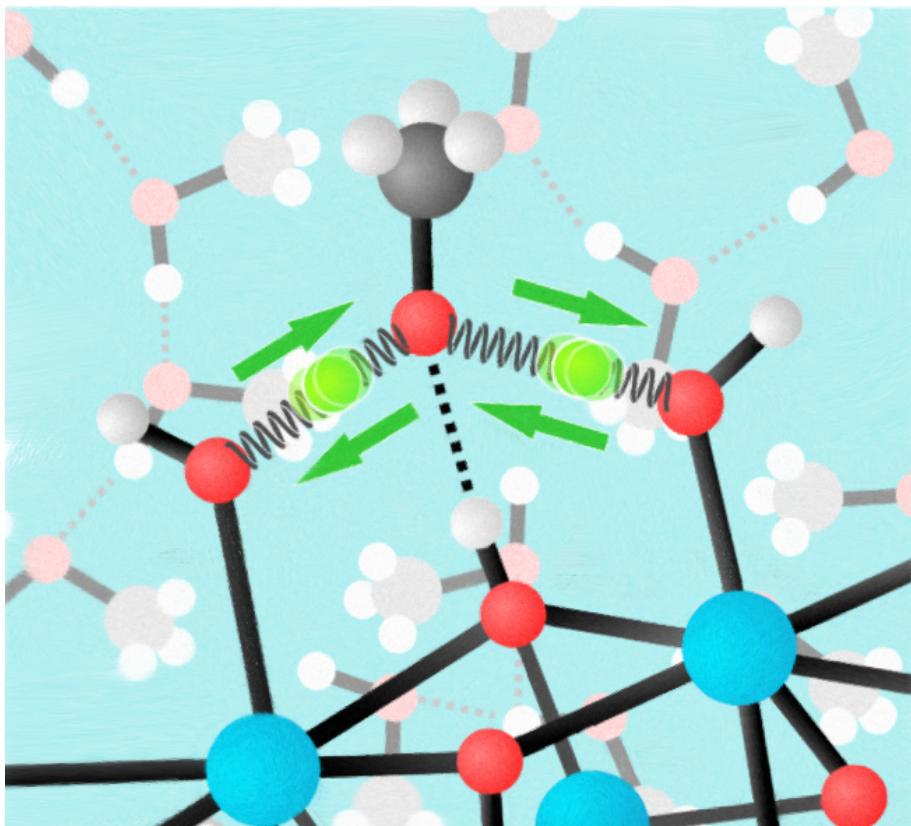
C. Caratelli performed all the calculations and prepared the manuscript.

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Paper II

Influence of a confined methanol solvent on the reactivity of active sites in UiO–66



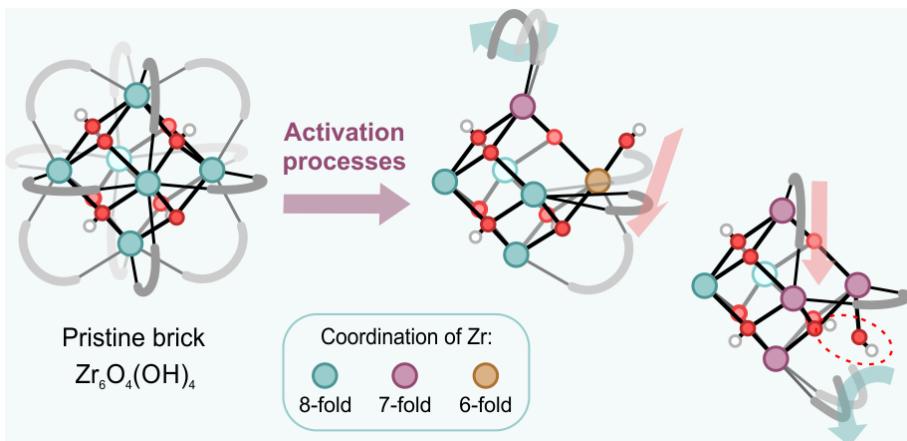
C. Caratelli, J. Hajek, S.M.J. Rogge, S. Vandenbrande, E.J. Meijer, M. Waroquier, V. Van Speybroeck, *ChemPhysChem*, **19**, 420–4290 (2018)

C. Caratelli performed all the calculations and prepared the manuscript.

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Paper III

On the intrinsic dynamic nature of the rigid UiO–66 metal–organic framework



J. Hajek, C. Caratelli, R. Demuynck, K. De Wispelaere, L. Vanduyfhuys, M. Waroquier, V. Van Speybroeck, *Chemical Science*, **8**, 2723–2732 (2018)

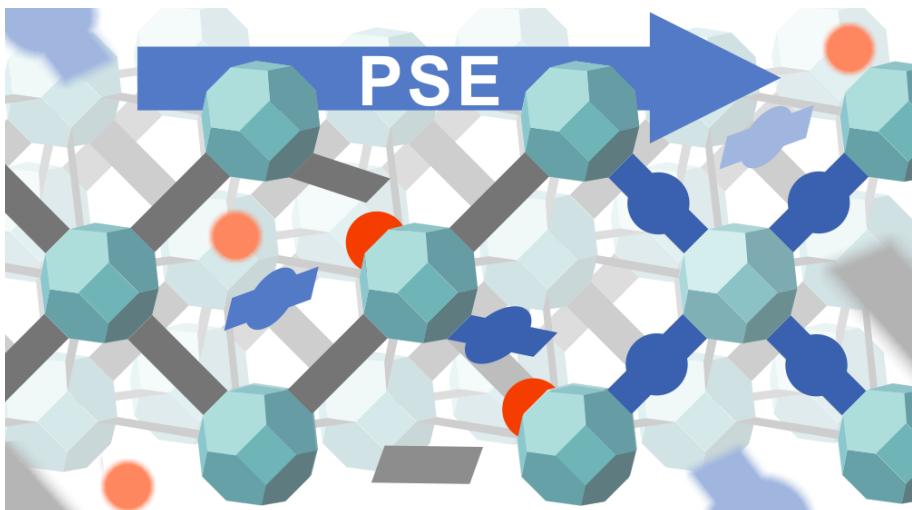
C. Caratelli was involved in the manuscript preparation.

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Paper IV

Active Role of Methanol in Post-Synthetic Linker Exchange in the Metal–Organic Framework UiO–66



J. Marreiros, C. Caratelli, J. Hajek, A. Krajnc, G. Fleury, B. Bueken, D. De Vos, G. Mali, M. Roeffaers, V. Van Speybroeck, R. Ameloot, *Chemistry of Materials*, **31**, 4, 1359–1369 (2019)

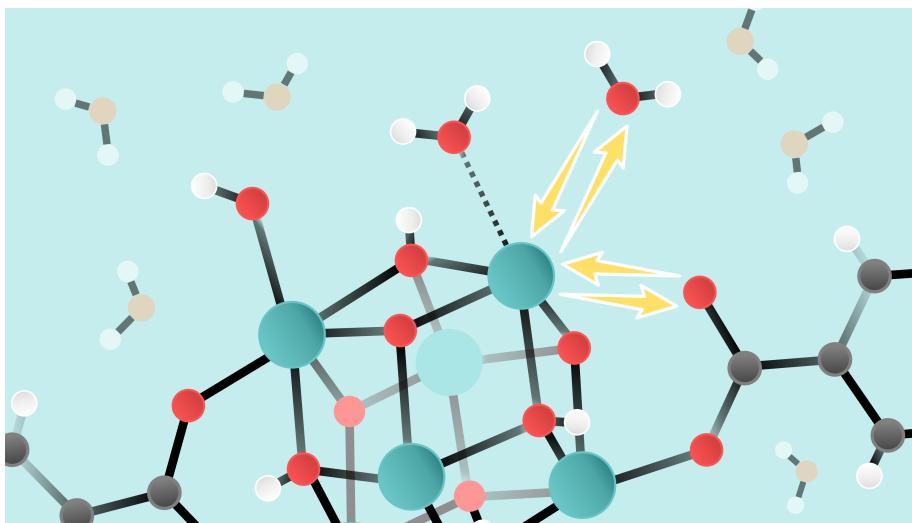
C. Caratelli performed the computational research and was involved in the manuscript preparation.

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Paper V

Dynamic interplay between defective UiO–66 and protic solvents in activated processes



C. Caratelli, J. Hajek, E.J. Meijer, M. Waroquier, V. Van Speybroeck, **Submitted**

C. Caratelli performed all the calculations and prepared the manuscript.

Paper VI

Investigating zirconium coordination changes in MOF-808 upon activation processes

In preparation

C. Caratelli performed part of the calculations and was strongly involved in the manuscript preparation.

A

Publication List

Updated May 2019

Publications in international peer-reviewed journals

1. M. Filez, C. Caratelli, M. Rivera-Torrente, F. Muniz-Miranda, M. Hoek, M. F.M. Altelaar, A. J.R. Heck, A.K Dutta Chowdhury, V. Van Speybroeck, B. M. Weckhuysen, *Molecular Nucleation Mechanism of Zeolitic Imidazolate Frameworks*, in preparation
2. J. Hajek, C. Caratelli, ... M. Waroquier, V. Van Speybroeck, *Investigating zirconium coordination changes in MOF-808 upon activation processes*, in preparation
3. C. Caratelli, J. Hajek, E.J. Meijer, B. Ensing, M. Waroquier, V. Van Speybroeck, *Dynamic interplay between defective UiO-66 and protic solvents in activated processes*, Submitted
4. J. Marreiros, C. Caratelli, J. Hajek, A. Krajnc, G. Fleury, B. Bueken, D. De Vos, G. Mali, M. Roeffaers, V. Van Speybroeck, R. Ameloot, *Active*

Role of Methanol in Post-Synthetic Linker Exchange in the Metal-Organic Framework UiO-66, Chemistry of Materials, **31** (4), 1359–1369 (2019)
 IF: 9.890

5. J. Hajek, C. Caratelli, R. Demuynck, K. De Wispelaere, L. Vanduyfhuys, M. Waroquier, V. Van Speybroeck, *On the intrinsic dynamic nature of the rigid UiO-66 metal-organic framework*, Chemical Science, **8**, 2723–2732 (2018)
 IF: 8.688
6. C. Caratelli, J. Hajek, S.M.J. Rogge, S. Vandenbrande, E.J. Meijer, M. Waroquier, V. Van Speybroeck, *Influence of a confined methanol solvent on the reactivity of active sites in UiO-66*, ChemPhysChem, **19**, 420–4290 (2018)
 IF: 3.075
7. C. Caratelli, J. Hajek, F. G. Cirujano, M. Waroquier, F. X. Llabrés i Xamena, V. Van Speybroeck, *Nature of active sites on UiO-66 and beneficial influence of water in the catalysis of Fischer esterification*, Journal of Catalysis, **352**, 401–414 (2017)
 IF: 6.844

Conference contributions

Oral presentations

1. Dynamic interplay between solvent and material in UiO-66 at catalytic conditions
C. Caratelli, J. Hajek, S. M.J. Rogge, M. Waroquier, V. Van Speybroeck
 Europacat 2019, Aachen, Germany, 18-23 August 2019
2. Dynamic interplay between defective UiO-66 and confined solvent: insights into a reaction environment at operating conditions
C. Caratelli, J. Hajek, S. M.J. Rogge, A. Lamaire, M. Waroquier, V. Van Speybroeck
 XXth Netherlands' Catalysis and Chemistry Conference (NCCC XX), Noordwijkerhout, The Netherlands, Mar 4–6 2019
3. Modeling nanoporous materials at the nanoscale: the role of high performance computing in materials science
C. Caratelli
 HPC-UGent User Meeting, Ghent, Belgium, Jan 28 2019
4. Post synthetic linker exchange of UiO-66: understanding the role of solvent and defects by time resolved characterization
J. Marreiros, C. Caratelli, J. Hajek, A. Krajnc, G. Fleury, B. Bueken, D. De Vos, V. Van Speybroeck, G. Mali, M. Roeffaers, R. Ameloot

IVth International Conference on Metal-Organic Frameworks and Open Framework Compounds (MOF 2018), Auckland, New Zealand, 9–13 December 2018

5. Investigating solvent effect in the acidity of UiO-66 metal organic framework for catalysis
C. Caratelli, J. Hajek, A. Tiwari, S. M.J. Rogge, B. Ensing, M. Waroquier, E.J. Meijer, V. Van Speybroeck
 IVth International Conference on Metal-Organic Frameworks and Open Framework Compounds (MOF 2018), Auckland, New Zealand, 9–13 December 2018
6. Towards a molecular level understanding of chemical and physical phenomena in metal-organic frameworks
J. Wieme, C. Caratelli, R. Demuynck, A. De Vos, J. Hajek, A. E. J. Hoffman, A. Lamaire, K. Lejaeghere, S. M.J. Rogge, S. Vandenbrande, L. Vanduyfhuys, M. Waroquier, V. Van Speybroeck
 Congrès français des MOFs, Paris, France, 16–18 May 2018
7. Influence of structural topology on the catalytic properties of Zr based MOFs: the case of UiO-66 and MOF-808
J. Hajek, C. Caratelli, M. Waroquier, V. Van Speybroeck
 XIXth Netherlands' Catalysis and Chemistry Conference (NCCC XIX), Noordwijkerhout, The Netherlands, 5–7 March 2018
8. Influence of a confined methanol solvent on the reactivity of active sites on UiO-66
C. Caratelli, J. Hajek, M. Waroquier, E.J. Meijer, V. Van Speybroeck
 2nd DEFNET School, Bochum, Germany, 18–21 September 2017
9. First principle study of active sites on UiO-66 for Fischer esterification
C. Caratelli, J. Hajek, F. G. Cirujano, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck
 Europacat 2017, Florence, Italy, 27–31 August 2017
10. First principle characterization of active sites on UiO-66 and their role in the catalysis of Fischer esterification
C. Caratelli, J. Hajek, F. G. Cirujano, M. Waroquier, F. X. Llabrés i Xamena, V. Van Speybroeck
 XVIIIth Netherlands' Catalysis and Chemistry Conference (NCCC XVIII), Noordwijkerhout, The Netherlands, 6–8 March 2017
11. Nature of active sites on UiO-66 and UiO-66-NH₂ in the catalysis of Fischer esterification
C. Caratelli, J. Hajek, G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck

Chemical Research in Flanders Symposium (CRF-1), Blankenberge, Belgium,
24–26 October 2016

12. Mechanistic study of Fischer esterification on UiO-66 and UiO-66-NH₂
C. Caratelli, J. Hajek, F.G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck
1st DEFNET School, Valencia, Spain, 21–24 June 2016

Poster presentations

1. How the connectivity of stable Zr-based MOFs affects the metal coordination and the nature of the active sites
J. Hajek, C. Caratelli, M. Waroquier, V. Van Speybroeck
MOFSIM 2019, Ghent, Belgium, Apr 10–12 2019
2. Dynamic creation of active sites on UiO-66 by interaction with protic solvents
C. Caratelli, J. Hajek, S. M.J. Rogge, M. Waroquier, B. Ensing, E.J. Meijer, V. Van Speybroeck
MOFSIM 2019, Ghent, Belgium, Apr 10–12 2019
3. Dynamic creation of active sites on UiO-66 by interaction with protic solvents
C. Caratelli, J. Hajek, S. M.J. Rogge, M. Waroquier, B. Ensing, E.J. Meijer, V. Van Speybroeck
1st KNCV-CTC symposium, Amsterdam, The Netherlands, Mar 26 2019
4. How the connectivity of stable Zr-based MOFs affects the metal coordination and the nature of the active sites
J. Hajek, C. Caratelli, M. Waroquier, V. Van Speybroeck
XXth Netherlands' Catalysis and Chemistry Conference (NCCC XX), Noordwijkerhout, The Netherlands, Mar 4–6 2019
5. Exploring the intrinsic dynamics of rigid Zr-based MOFs
J. Hajek, C. Caratelli, R. Demuynck, K. De Wispelaere, L. Vanduyfhuys, M. Waroquier, V. Van Speybroeck
IVth International Conference on Metal-Organic Frameworks and Open Framework Compounds (MOF 2018), Auckland, New Zealand, 9–13 December 2018
6. Modeling reactive processes in nanoporous materials: a look into the complexity
C. Caratelli, K. De Wispelaere, J. Hajek, P. Cnudde, S. M.J. Rogge, S. Vandenbrande, R. Demuynck, L. Vanduyfhuys, M. Waroquier, V. Van Speybroeck
Ghent, Belgium, Jun 1 2018
7. Investigating the outstanding dynamic behavior of protons on UiO-66 defective sites

C. Caratelli, J. Hajek, A. Tiwari, M. Waroquier, B. Ensing, E. Jan Meijer, V. Van Speybroeck

EuroMOF 2017, Delft, The Netherlands, Oct 29 -- Nov 1 2017

8. Post synthetic linker exchange of UiO-66: understanding the role of solvent and defects by time resolved characterization

J. Marreiros, C. Caratelli, J. Hajek, A. Krajnc, G. Fleury, V. Van Speybroeck, G. Mali, M. Roeffaers, R. Ameloot

EuroMOF 2017, Delft, The Netherlands, Oct 29 -- Nov 1 2017

9. Influence of a confined methanol solvent on the reactivity of active sites in UiO-66

C. Caratelli, J. Hajek, S. M.J. Rogge, S. Vandenbrande, E.J. Meijer, M. Waroquier, V. Van Speybroeck

XIXth Netherlands' Catalysis and Chemistry Conference (NCCC XIX), Nordanwijkerhout, The Netherlands, Mar 5–7 2018

10. Catalytic sites on UiO-66 for Fischer esterification

C. Caratelli, J. Hajek, G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck

MOLSIM 2017: Understanding Molecular Simulations, Amsterdam, The Netherlands, Jan 9–20 2017

11. Catalytic sites on UiO-66 for Fischer esterification

C. Caratelli, J. Hajek, G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck

Annual IAP Meeting IAP-PAI P7/05, Liège, Belgium, 12 October 2016

12. Catalytic role of UiO-66 and UiO-66-NH₂ in Fischer esterification: a mechanistic study

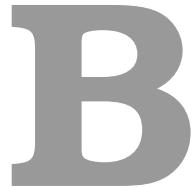
C. Caratelli, J. Hajek, G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck

3rd DEFNET workshop, Ghent, Belgium, 22–24 March 2016

13. Catalytic role of UiO-66 and UiO-66-NH₂ in Fischer esterification: a mechanistic study

C. Caratelli, J. Hajek, G. Cirujano, A. Corma, M. Waroquier, F.X. Llabrés i Xamena, V. Van Speybroeck

XVIIth Netherlands' Catalysis and Chemistry Conference (NCCC XVII), Nordanwijkerhout, The Netherlands, 7–9 March 2016



Granted computational Projects

Granted computational projects under the Vlaams Super Computing Centrum infrastructure

1. Operando study of post synthetic structural modifications in UiO-66
Mar 2019 - Nov 2019
4176 node days (Breniac)
2. Active sites on MOF-808 for industrially relevant MPV reduction
Nov 2018 - Jul 2019
4236 node days (Breniac)
3. Intrinsic dynamics of rigid Zr-based metal organic frameworks
Jul 2018 - Mar 2019
2766 node days (Breniac)
4. Investigating solvent effect in the acidity of UiO-66 metal organic framework for catalysis
Jul 2018 - Mar 2019
4764 node days (Breniac)
5. Probing the strength of basic sites on UiO-66 using pKa calculations
Jul 2017 - Jan 2017
4450 node days (Breniac)

6. *Ab initio* umbrella sampling simulations of dehydration of UiO-66
Nov 2016 - Apr 2017
3972 node days (Breniac)
7. Pilot Tier 1 project
Jul 2016
1000 node days (Breniac)
8. Investigating active sites in hydroxylated and dehydroxylated UiO-66 for
catalysis of Oppenauer-type oxidation
Feb 2016 - Oct 2016
2110 node days (Muk)

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