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# Synthetic routes and fields of application of isohexides: comprehensive perspective of relevant industrial compounds

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Isohexides are a class of compounds that can be obtained from natural sources using simple processes, and have interesting properties. Among these, the intrinsic chirality of the rigid bicyclic structural cavity and the presence of two hydroxyl groups in positions 3 and 6, available for functional-group interconversion, appear the most interesting characteristics for real innovative applications. In this review, the strategies for the synthesis of isohexides from polysaccharides and biomasses of different complexities have been analyzed and the alternative processes to the traditional synthetic routes, in terms of sustainability, have been discussed. Secondly, the properties that make isosorbide the most suitable isohexide compound in terms of availability, stability and scalability for innovative production processes have been highlighted. The fields of application of these extremely interesting compounds have been investigated, proceeding with a discussion in which the areas of exploitation of isohexides are analyzed, in terms of their use as building blocks, chiral auxiliaries or constituents of chiral ionic liquids. Finally, the numerous applicative possibilities of isosorbide as a monomer or co-monomer of a wide range of polymeric materials of different natures, have been thoroughly investigated. This review represents a first investigation into the possible uses of isohexide-based compounds by analyzing the positive and negative aspects of their exploitation as ubiquitous alternatives to current oil-based intermediates. Certainly, future studies on the optimization of those production processes that occupy a large share of the chemical-product market will lead to the confirmation of the synthetic potential of compounds with an isohexidic structure.

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## Sustainability spotlight

Isohexides can be used in production processes and synthetic transformations as sustainable alternatives to those currently employed, derived from crude oil. In fact, these compounds adhere to at least 7 of the 12 principles of Green Chemistry. In particular, isosorbide can be extracted from renewable feedstocks, preventing the wastage of precious resources and incorporating atom economy, it can be used as a benign solvent leading to less hazardous syntheses, and it can be employed in catalytic quantities or as a chiral auxiliary in processes that traditionally require toxic organocatalysts. These are bicyclic compounds that can be isolated from sugar wastes that do not compete with the agri-food supply chain, namely low-cost materials that are capable of undergoing even complex transformations with harsh reaction conditions and can be used to produce highly performing polymeric materials. The reader is therefore exposed to the real possibility of ubiquitous application of compounds with an isohexidic structure in very different but fundamental areas of the new sustainable transition.

## 1. Introduction

In the United States alone, 400 million tons of sugar-based dry biomass are generated annually from the corn, soybean, flour, and sorghum production industries.<sup>1</sup> The complete replacement of fossil fuels with biomass fuels is difficult to achieve due to the enormous scale of fuel consumption and the market demand. For these reasons, biomass is currently more effectively used in the conversion to chemical feedstocks and chemical precursors.<sup>2</sup> However, the high market value of these

products sometimes requires the use of expensive unit operations for pre-treatment, processing, and separation, completely *ex novo* compared to traditional industrial precursor plants.

The growing interest in a circular economy and environmental sustainability has increased the search for renewable sources and their conversion into products, which include chemical reagents, energy resources and materials,<sup>3</sup> to replace the enormous demand for fossil fuels (Fig. 1).

Given the abundance of these materials and the lack of competition with the agri-food sector, during the current century a market segment dedicated exclusively to the valorization of waste biomass was successfully established. Therefore, research is evolving towards the formulation of integrated

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Fig. 1 US primary energy consumption by energy source in 2023; only 9% is renewable.<sup>3</sup>

processes that can lead to new products with high added-value, starting precisely from the mentioned biomasses.

Due to this promising background, many scientists from all over the world have dedicated their research to the extraction and separation of high-added-value components from wastes of different natures. This has opened new frontiers in different areas of the chemical industry, allowing components deriving from petrochemicals (used in already consolidated processes) to be replaced with interesting and more sustainable alternatives.<sup>1</sup>

In this context, agricultural raw materials come into play as a natural, renewable and sustainable resource to produce a large variety of biopolymers, biochemicals and bioenergy sources.<sup>4,5</sup> For example, cellulose can be obtained mainly from agricultural residues and is one of the largest sources of organic raw materials in the world. It is in fact considered a promising resource that can be transformed into sustainable fuels and chemicals.<sup>6</sup>

Since the 1990s, there has been growing research on the isolation, synthesis and innovative use of numerous sugar-based products.<sup>7,8</sup> As can be seen from the graph shown in Fig. 2, interest in this sector and the related scientific publications have witnessed an exceptional increase over the past decades.<sup>9</sup>

Alcohols derived from C<sub>5</sub> and C<sub>6</sub> sugars obtained from cellulose or hemicelluloses are considered as sustainable building blocks and important synthons in organic synthesis.<sup>10</sup> Among these, the hexitols are a class of polyols with six hydroxyl groups that can be obtained from the corresponding hexoses (Fig. 3).<sup>11</sup>

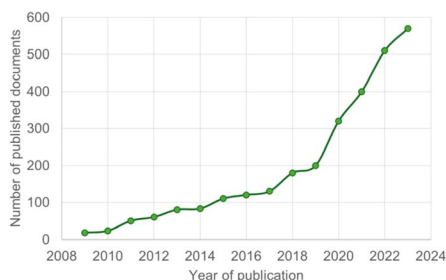


Fig. 2 Number of documents published on biomass valorisation per year; data from cited Scopus database.<sup>9</sup>



Fig. 3 Synthetic scheme to produce isohexide precursors starting from agro-food waste biomass.<sup>11</sup>

In fact, from plant-based biomasses rich in fibers (such as rice husk, wood pulp, banana peel, coffee husk, wheat straw, sisal fibers and sugarcane bagasse) it is possible to isolate cellulose microfibrils using recently consolidated processes.<sup>12</sup>

Crystalline structures that can be isolated from cellulose constitute some of the optimal precursors for the synthesis of hexose sugars from natural sources. The interconversion of functional groups starting from the OH present on the sugar-based carbon skeleton opens the door to different classes of compounds. Isohexides, for example, are compounds with a 6-carbon skeleton and 2 hydroxyl groups linked to different carbons, and are extremely interesting.

Therefore, the different configurations of the stereogenic centers define the structural divergences of the isohexidic spatial isomers and their further chemical modification. These differences will be highlighted throughout the manuscript and the important asymmetric reactivity of the two OH groups will be especially emphasized.

In this review, the synthetic strategies for isohexides will be thoroughly discussed, focusing the reader's attention on the comparison between the processes traditionally used in industry or at the laboratory scale and the possible paths recently identified as greener alternatives.

Subsequently, the possible application of isohexides in the organic synthesis of discrete molecules, such as chiral auxiliaries and macromolecular materials with a polymeric structure, will be discussed, demonstrating the suitability of the use of compounds with an isohexidic structure in various industrial sectors.

### 1.1. Structure of isohexides

The term "isohexides" refers to a class of diol compounds whose structures are characterized by the presence of two tetrahydrofuran rings fused with a *cis*-type junction.

Isosorbide (1,4:3,6-dianhydrosorbitol **1**), isomannide (1,4:3,6-dianhydromannitol **2**) and isoidide (1,4:3,6-dianhydroiditol **3**), are three of the main isomers in this category of natural sugar-based compounds. They are distinguished by their different stereochemistries of the hydroxyl groups in position 3 and in position 6: the two OH groups of isosorbide **1** are *endo-exo*, while in isomannide **2** and in isoidide **3** they are

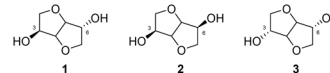


Fig. 4 Chemical structures of isohexides: isosorbide **1**, isomannide **2**, and isoidide **3**.



oriented *endo-endo* and *exo-exo*, respectively.<sup>13</sup> Their chemical structures are shown in Fig. 4.

The type of junction present between the two rings gives a concave vaulted structure to the isohexides, with a dihedral angle between the two tetrahydrofuran rings of about 120°. As can be observed in Fig. 5, in isosorbide **1**, the concavity of the structure offers the possibility of forming intramolecular hydrogen bonds between the hydrogen of the hydroxyl group in position 3 and the oxygen of the adjacent ring.<sup>4</sup> The two hydroxyl groups occupy different stereochemical surroundings; furthermore, we can deduce that they also show different degrees of reactivity.

In isomannide, the hydroxyl groups have the same stereochemistry, thus making **2** a highly symmetric derivative with the possibility of forming two intramolecular hydrogen bonds, thanks to the *endo* arrangement of both hydroxyl groups facing the inside of the chiral cavity. In isoidide **3**, no intramolecular hydrogen bond can occur due to the orientation of the two OH groups towards the outside of the cavity.

## 1.2. Properties of isohexides

Regardless of the *endo* or *exo* position of the hydroxyl groups, for the same carbon skeleton the reactivity of the two substituents in **3** and **6** is affected by the nearby chemical environment.

The structural difference between the three isohexides can be also predicted from the wavelengths at which the maxima of the different peaks relating to the stretching of the O–H bonds can be observed *via* IR spectroscopy (Fig. 6).<sup>15,16</sup>

Even the literature <sup>1</sup>H-NMR spectra<sup>15,16</sup> of isohexides demonstrate the non-equivalence of the above-mentioned groups, if all other assumptions can be considered equal. The data and deductions shown in Table 1 have been obtained from theoretical simulations, while at the experimental level these differences are rarely observed through the most common techniques of characterization of organic compounds, performed under standard conditions.

The structural difference between the three isomers clearly implies differences in their chemical–physical properties, such as their melting temperature (Table 2) and the reactivity of the hydroxyl groups.

As discussed before, the difference in melting temperature ranges can be attributed to the presence or absence of intramolecular hydrogen bonds. Considering the steric effects and the establishment of hydrogen bonds, isomannide **2** with the two *endo* hydroxyl groups is certainly the least reactive isomer, and isoidide **3** with the two *exo* groups is more reactive than isosorbide **1**. Therefore, it could be deduced that isoidide **3** is the most attractive isomer on the market, but, unfortunately, it



Fig. 5 Three-dimensional structures of isohexides with possible hydrogen bonds shown.<sup>4</sup>

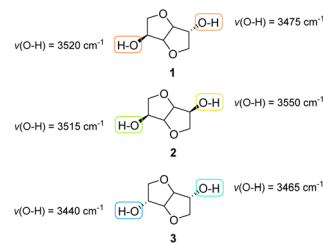


Fig. 6 Open-source FT-IR simulation data for O–H stretching of isohexides.<sup>14</sup>

is the rarest in nature and remains the most expensive for many industrial applications.

For the above-mentioned reasons, research attention in the field of isohexides derived from renewable sources and their applications is mainly directed towards the *endo-exo* isosorbide **1** and in recent years has witnessed a notable increase. The interest in isosorbide lies both in the possibility of selective derivatization of the OH groups and in the presence of the cavity-like structure. This cavity constitutes a true chiral confined system, within which molecules of appropriate size and stereochemistry can be hosted or converted, thus favouring enantioselection.

Meanwhile, the selective derivatization of the hydroxyl groups of isomannide **2** is possible only through the control of the stoichiometry of the reagents. Based on a statistical approach, a 2 : 1 ratio between isomannide and the derivatizing agent is usually employed, which often leads to the formation of a mixture of mono-derivatized product, disubstituted derivative and unreacted reagent.

On the other hand, the non-equivalence of the OH groups of isosorbide **1** enables its selective functionalization and hence the production of mono-substituted and non-symmetric composites with great industrial relevance.

## 2. Synthesis of isohexides

### 2.1. From biogenic polysaccharides

Polysaccharides are a class of macromolecules that are widespread in nature among animals, plants, microorganisms and

Table 1 <sup>1</sup>H-NMR simulation data for isohexides<sup>17</sup>

Compound	$\delta$ [ppm]	H	Multiplicity	$J$ [Hz]
<b>1</b>	3.65–3.95	8H	dd	7.0, 5.9
	3.81		dd	12.9, 6.8
	3.87		dd	12.9, 4.7
	3.88		ddd	6.8, 5.9, 4.7
	3.65–4.00	8H	dd	7.3, 6.7
<b>2</b>	3.77		dd	10.4, 6.1
	3.90		ddd	6.7, 6.1, 5.5
	3.93		dd	10.4, 5.5
	3.66	2H	dd	7.0, 3.8
	<b>3</b>	3.82–4.00	6H	ddd
3.91			dd	10.5, 7.8
3.93			dd	10.5, 6.8



Table 2 Melting temperature of isohexides

Compound	Melting temperature [°C]
1	60–63
2	80–85
3	50–56

algae. These polymeric materials are commonly characterized by high molecular weight and complex structural features that determine their biological functions.<sup>18</sup>

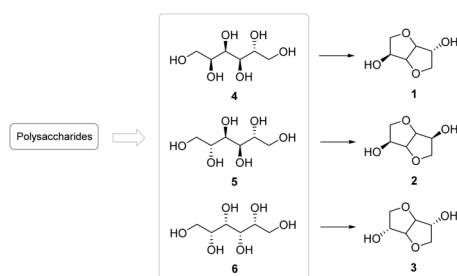
The performance of these functions can be crucial for the survival, growth and health development of the organism that synthesizes them or that exploits their metabolism. In recent years, there have been increasing attempts to study natural biomaterials based on polysaccharides. The enhanced properties of polysaccharides have been extended to various applications, including biomedical engineering, human drug delivery, food packaging, biofuel isolation from biomasses, contaminated wastewater treatment and even textile fiber production.<sup>19</sup>

Due to their biodegradable and biocompatible characters, bioactive polysaccharides have been largely used in hexitol production processes, according to the synthetic scheme shown in Fig. 7.

As previously mentioned in the Introduction (Section 1), hexitols are C<sub>6</sub> polyols: organic compounds with various hydroxyl groups present in the structure with a skeleton composed of six carbon atoms. Therefore, starting from polysaccharides it is possible to isolate hexitols, thus obtaining the precursors for isohexides directly from sustainable sources.

In fact, isohexides have been known since 1880 and their synthesis commonly involved the dehydration of the corresponding hexitols (sorbitol, mannitol and iditol) in an acid environment and in the presence of a solvent.

**2.1.1. Synthesis of isosorbide from sorbitol.** In the 1990s, the industrial synthesis of isosorbide involved a homogeneous batch process at atmospheric pressure, catalyzed by sulfuric acid and with water as a solvent medium.<sup>20</sup> The main drawbacks of the process, shown in Fig. 8, included the need for separation of the products by neutralization, the elimination of sulfur compounds, the required distillation stage (energy-demanding for each production batch and causing the acid catalyst to

Fig. 7 Scheme of isohexide production from hexitols.<sup>19</sup>Fig. 8 Isosorbide synthesis starting from sorbitol, using an acidic aqueous medium and traditional reaction conditions.<sup>20</sup>

become inert) and finally the recovery of the product with a yield that did not exceed 65% by weight.

The desire for the combination of Green Chemistry principles and industrial progress has led to the development of new production methods for this isohexidic intermediate. For example, in 2013 an innovative catalytic dehydration strategy of sorbitol **4** to isosorbide was proposed,<sup>21</sup> which employed microwaves (MWs) as a heating method and Amberlyst 35 as an acid catalyst (Fig. 9).

This research led to optimization of the reaction conditions, efficient recovery of energy and an increase in isosorbide yield. Following the principles of atomic efficiency and a good compromise between yield and purity of the product, this strategy is still nowadays the most widely used to produce isosorbide from hexitols.

**2.1.2. Synthesis of isomannide from mannitol.** As seen for isosorbide, isomannide **2** can also be produced from the respective hexitol, *i.e.*, mannitol. As an example, the data relating to the conditions used in the synthesis of isomannide starting from mannitol **5** and the relative yields are reported in Fig. 10 and Table 3, respectively.

The main disadvantages of these synthetic methods included the requirement of solvents, the commonly unsatisfactory yield achieved, the obtaining of a large amount of resinous distillation residues and the long reaction times often required, reaching over 85 hours for complete substrate-to-product conversion.

The issues, also highlighted in Section 1.1.1, exclude the possibility of preparing both isosorbide **1** and isomannide **2** on a large scale with these methods. In addition, the cost of the starting material (for example pure mannitol and sorbitol) is considerably high compared to the normal cost of the precursors used in the chemical industry to produce intermediates intended for commodities.

Subsequently, an innovative synthetic route for isohexides was proposed, where the authors obtained high yields of isomannide with a selective reaction, in the absence of solvent, by treating the crystalline hexitol with gaseous hydrogen halide (preferably HCl<sub>(g)</sub>), as shown Fig. 11. To date, these are the most used dehydration reaction conditions at the laboratory level to produce **1** and **2** from hexitols **4** and **5**, respectively.

**2.1.3. Synthesis of isoidide from iditol.** Isoidide can also be synthesized by dehydration of the corresponding hexitol. In fact,

Fig. 9 Isosorbide synthesis starting from sorbitol, using Amberlyst 35 in acidic aq. medium and a MW heating system.<sup>21</sup>

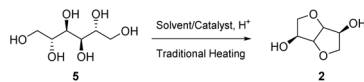


Fig. 10 Isomannide synthesis starting from mannitol, using different solvent–catalyst systems and traditional heating.<sup>20</sup>

Table 3 Reaction conditions and yields obtained in isomannide synthesis starting from mannitol

Solvent/Catalyst	2 yield [wt%]
Ethyl acetate/dioxane	35
Dichloroglycerol	36
Conc. hydrochloric acid	25–40

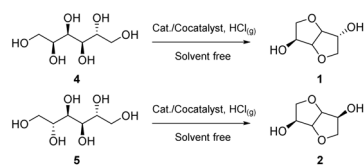


Fig. 11 Solvent-free isosorbide and isomannide synthesis starting from sorbitol and mannitol, using catalysts and gaseous hydrogen halide.<sup>20</sup>

3 can be produced starting from iditol **4**, a water-soluble C<sub>6</sub> sugar polyol that can be industrially obtained from biomasses such as agricultural wastes.

The reaction conditions shown in Fig. 12 are analogous to those traditionally used. Therefore, an acidic medium, in which the acid functionalities catalyze the dehydration reaction and consequent intramolecular closure of the bicyclic structure, was successfully employed.

**2.1.4. Synthesis of isoidide from isosorbide.** An interesting synthetic strategy proposed by Saska *et al.*<sup>22</sup> involved the synthesis of isosorbide through the epimerization of isosorbide. The term epimerization refers to a stereochemical process by which a diastereomer can be converted into another diastereomer by changing just one chiral center in the entire structure.<sup>23</sup>

The authors were able to propose a reaction mechanism that involved the two steps shown in Fig. 13: a catalytic dehydrogenation–hydrogenation reaction and a hydride abstraction/re-addition step. The first step included the use of Ru/C as a catalyst in 2-propanol, while the hydride abstraction was performed with Pb(OAc)<sub>2</sub> in acetic anhydride, leading to an overall isoidide purity with a value higher than 99%.

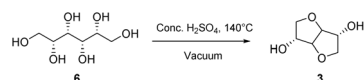


Fig. 12 Isoidide synthesis starting from iditol, using concentrate sulfuric acid and a traditional heating system.<sup>20</sup>



Fig. 13 Isoidide synthesis starting from isosorbide, using a two-step chemo-catalytic process.<sup>22</sup>

In fact, a few years earlier, the optimized conditions for the synthesis reaction of isoidide from isosorbide had been developed by the group of Le Nôtre *et al.*,<sup>15</sup> highlighting the potential applications of epimerization reactions of isohexides.

As demonstrated by the survey discussed in Section 1, isohexide **1** has the greatest applicative possibilities and the most abundant availability in nature. Therefore, the discussion will focus on the industrial and laboratory perspectives of using isosorbide as a building block in the applied fields of organic synthesis.

## 2.2. Synthesis of isosorbide from raw biomasses

A more complex but more abundant sugar source than simple biogenic polysaccharides is constituted by cellulose and lignocellulosic biomass. This type of biomass does not compete with the agri-food sector and often constitutes a waste volume that aggravates down-stream stages of industrial production processes of products such as paper, pulp, cellulose, *etc.*

To date, studies have been carried out on the possibility of obtaining isosorbide by means of a one-pot synthesis starting from cellulose or lignocellulosic biomasses. The synthesis shown in Fig. 14 involved the following cascade: (i) the hydrolysis of cellulose or starch to glucose **7**, (ii) the hydrogenation of glucose to sorbitol **4**, (iii) the dehydration of sorbitol to sorbitan **8**, and lastly (iv) the dehydration of sorbitan to isosorbide **1**.

The one-pot process required both an acidic environment and a hydrogenation catalyst. The acidity of the medium, the crystallinity and particle size of the cellulose, and the nature and therefore the origin of the raw materials, significantly influenced the yield of the isosorbide production process. The deactivation of the catalyst in the presence of lignin (present in most cellulosic raw material stocks) and its non-recyclability in an aqueous environment were considered the main problems of this proposed synthetic path.



Fig. 14 One-pot isosorbide synthesis starting from biomasses.<sup>15</sup>



In these studies, the possibility of formulating a sustainable and interesting way of designing a heterogeneous catalyst composed of a supported metal, designed to be stable in water and tolerant to the presence of lignin, was hypothesized. Carrying out this reaction in continuous flow using two alternating catalytic beds (suitable for the regeneration of the catalyst) could constitute an alternative solution for the synthesis of isosorbide from raw biomasses.

There was recent experimental evidence of a certain success for the synthesis of **1** starting from lignocellulosic biomass, which employed a heterogeneous ruthenium-based hydrogenation catalyst supported on carbon.<sup>24</sup> Unfortunately, the reaction conditions shown in Table 4 led to the synthesis of isosorbide with low yields that do not exceed 65% by weight, a value that can be traced back to the phenomenon of deactivation of the hydrogenation catalyst, which significantly lowers the product yield.

Research in this area is moving towards the design of a heterogeneous catalyst composed of a transition metal supported on a carrier with acid moieties, stable in water and tolerant to the presence of the lignin contained in the starting material. In this way, it would be possible to overcome the difficulties encountered to date in undertaking a synthesis with a very complex matrix, such as lignocellulosic biomasses.

Certainly, the choice of synthesizing dianhydrous hexitols starting from biomass of plant origin represents the most complex strategy at the laboratory level, but it could constitute a scalable innovation process suitable for the industrial production of important intermediates such as isosorbide.

### 3. Isohexides as building blocks

In the chemical field, the term “building blocks” refers to a virtual molecular fragment or a real chemical compound that possesses reactive functional groups. Building blocks are used for the bottom-up assembly of molecular architectures such as nanoparticles, metal–organic structures, organic compounds with high molecular complexity and supra-molecular complexes.<sup>30</sup>

The selective reactivity of the two hydroxyl groups, the chirality, the controlled stereochemistry and the performance of its derivatives, lead to isosorbide being a promising “green fine chemical”. In fact, isohexides in general are considered fine chemicals as they are produced in limited quantities and at

a high market price and are therefore destined to be converted into value-added products.

#### 3.1. Chiral intermediates based on isohexides

Dianhydrous hexitolic derivatives have been successfully employed in asymmetric catalysis.<sup>31</sup> Asymmetric catalysis is the most widely used method for carrying out an asymmetric synthesis, *i.e.*, the conversion of a prochiral substrate into a chiral product with preferential formation of one of the possible enantiomers.<sup>32</sup> To perform a reaction within asymmetric catalysis, a chiral catalyst is commonly needed. Usually, a transition-metal complex coordinated by chiral ligands is employed and when it is formed of purely organic chiral ligands, the reaction is known as asymmetric organocatalysis.<sup>33</sup>

The efficiency of an enantioselective synthesis process can be very high, but each stage of the synthesis influences the final product yield; therefore, an industrial design suitable for the purpose will be required. This approach is particularly important in the field of pharmaceutical synthesis, since different enantiomers or diastereomers of a molecule often have different biological activities.<sup>34</sup>

Among the many possible synthetic intermediates that can be produced from isohexides, chiral derivatives are of vital importance. Given the exceptional properties and advantages mentioned above, the mono- and disubstituted derivatives of isosorbide will be mainly discussed.

At first glance, some of the isosorbide derivatives worth mentioning are reported in Fig. 15.

The synthesis of a wide spectrum of mono- and diesters, mono- and dinitrates, and mono- and ditosylates of isosorbide, such as compounds **9–13**, were reported by Bhat *et al.*,<sup>36</sup> expanding the chiral intermediates that can be obtained from biomass-derived isosorbide. For example, most of the amino derivatives of isosorbide are designed for the polymer field (*e.g.*,

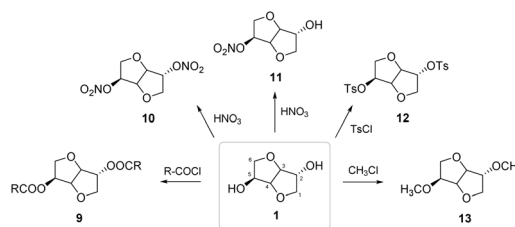


Fig. 15 Industrially relevant derivatives of isosorbide.<sup>35</sup>

Table 4 Reaction conditions and yield results for isosorbide synthesis starting from lignocellulosic biomass<sup>24</sup>

Dehydrating agent	<i>T</i> [°C]	H <sub>2</sub> [bar]	Time [h]	Ru/C [mg]	<b>1</b> yield [wt%]	Ref.
[HCl] = 0.1 M	215	60	0.5	20	50	25
[H <sub>2</sub> SO <sub>4</sub> ] = 0.5 M	220	40	2	40	50	26
[BPO <sub>4</sub> ] = 0.1 M	230	60	24	20	28	27
[H <sub>4</sub> SiW <sub>12</sub> O <sub>40</sub> ] = 55 M	160	50	7	100	13	28
[H <sub>4</sub> PW <sub>12</sub> O <sub>40</sub> ] = 55 M	160	50	7	100	17	28
[H <sub>4</sub> SiW <sub>12</sub> O <sub>40</sub> ] = 3 M	210	50	7	1000	65	29



polyurethanes and polyamides), as will be discussed in Section 3.3, but the role of isosorbide-based building blocks has attracted great curiosity, especially in the chemical industry for the asymmetric synthesis of enantiopure compounds.<sup>37</sup>

In fact, in recent years the call for a sustainable transition away from well-known polluting and hazardous reactions and an upgrade of the large-scale synthesis of intermediates, such as ligands and chiral auxiliaries, has found an outlet in asymmetric structures based on isosorbide derivatives.

For instance, Wu *et al.*<sup>38</sup> developed a novel isosorbide-based dinitrile synthesis procedure, leading to the production of compounds with one extra carbon atom for each hydroxyl group through isosorbide epimerization under mild and base-catalyzed reaction conditions (Fig. 16). The author proposed a detailed reaction mechanism through a structural analysis of the possible dinitrile isomers *via* DRX analysis<sup>39</sup> and DFT calculations,<sup>40</sup> demonstrating that different characteristic structural parameters of the chiral cavity, such as dihedral angles and torsion angles, can vary over a wide range and modify the outcome of the reaction.

The concept of structural rigidity of the bicyclic cavity was overcome by the authors, who proposed the possibility of less conservation of torsion and were consequently able to demonstrate access to an interesting class of building blocks in which the substituent orientation effects the yield of the extended C<sub>2</sub>/C<sub>5</sub> isohexidic compounds.<sup>38</sup>

Isohexidic derivatives have been used in asymmetric catalysis, as chiral metal ligands for the asymmetric reduction of alkenes, ketones and imines, and for the formation of C–C bonds *via* allylic substitution, alkylation and hydroformylation.<sup>31,41</sup>

They have also been employed as chiral auxiliaries in Diels–Alder reactions (Fig. 17), affording products with enantiomeric excesses of up to 99% for the desired product. The authors were able to fully demonstrate the efficiency of using isosorbide-based compounds as chiral auxiliaries, leading to comparable or even improved results with respect to the more traditional auxiliary compounds based on natural sources, such as camphor and menthol derivatives.

Isosorbide has also been employed as a starting material for the synthesis of many novel chiral amino alcohols and diamines, as shown in Fig. 18. These derivatives have been synthesized in moderate to good yields (43–75%), depending on



Fig. 17 Diels–Alder reaction of an acrylate derived from *exo*-mono-benzylated isosorbide and cyclopentadiene.<sup>42</sup>

the RNH<sub>2</sub> reagents used. The above-mentioned study demonstrated the potential use of the obtained products as chiral ligands for asymmetric transfer hydrogenation (ATH) reactions of aromatic ketones catalyzed by ruthenium, providing chiral alcohols with enantiomeric excesses up to 91%.<sup>43</sup>

The exploitation of biomass-based starting materials can also be applied in the case of isosorbide itself and not only for its derivatives,<sup>44</sup> whose interconversion has appeared very attractive in recent years. In fact, the *endo*–*exo* spatial arrangement of the two hydroxyl groups in positions 3 and 6 of the isosorbide skeleton has been successfully converted to synthesize a wide range of organo-catalysts and chiral auxiliaries.<sup>45</sup>

Zullo *et al.*<sup>46</sup> were able to synthesize an amino alcohol with an isohexide structure by subjecting isosorbide to various reactive steps, including the innovative selective acetylation of the *endo*-OH group through a protection performed by a biocatalyst in heterogeneous form (Fig. 19).<sup>46</sup>

Starting from the amino alcohol in question, they were able to successfully use this compound as a chiral solvating agent in the enantio-discrimination of complex aromatic structures, demonstrating the possibility of using the functionality of asymmetric induction from a chiral pool.<sup>47</sup>

Recently, the Mitsunobu reaction<sup>48</sup> has also been successfully performed on isosorbide, leading to the synthesis of some bioactive intermediates with improved pharmacodynamic and pharmacokinetic properties.<sup>49</sup>

In fact, conjugates containing isohexidic moieties have been synthesized that confer improved solubility and permeability properties to the structure, compared to the use of bare scaffold compounds towards target compounds. Only a few of the possible examples of isosorbide-based compounds with high added-value in the field of enantioselective and asymmetric synthesis have been mentioned, but research has witnessed a large increase in possible chiral discriminants based on the strategies discussed in Section 3.1.

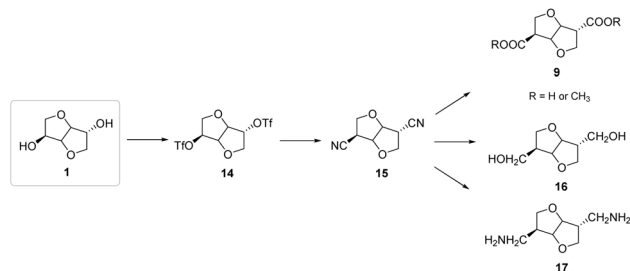


Fig. 16 Synthesis of 1-carbon-extended isosorbide-based compounds through epimerization reactions.<sup>38</sup>



Fig. 18 Synthesis of some amino derivatives of isosorbide.<sup>43</sup>





mass and high percentage of functionalization of the formed oligomers.<sup>13,59</sup> By the comparison of different studies, it emerged that the inclusion of isosorbide functionalities can lead to oligomers with a molar mass of 260–770 g mol<sup>-1</sup> with an increased impact and tensile strength.

The mentioned results were obtained despite the characteristic lower glass-transition temperature of diglycidyl ethers based on isosorbide compared to traditional epoxy resins that use BPA as a reagent.<sup>60</sup>

**3.3.2. Polyesters.** Considering its diol nature, isohexide **1** can also be used in the synthesis of polyesters. Polyesters are a class of polymers characterized by the presence of an ester bond (–CO–O–) in the main chain of the macromolecule and they can be obtained *via* the reaction of diols with diacids or diesters.<sup>61</sup> Isosorbide was successfully employed as a monomer in the production of thermoplastic materials, specifically in the synthesis of polyester.<sup>62–64</sup>

On an industrial scale, polyesters can be obtained from carboxylic acids or derivatives and diols using a two-stage polycondensation process that involves the release of a low-molecular-weight by-product. They can be either aromatic or aliphatic based on the nature of the monomers used in the production process.

Among the most common aromatic polyesters we can find poly(ethylene terephthalate) (PET), poly(butylene terephthalate) (PBT), poly(cyclohexylene dimethylene terephthalate) (PCT) and poly(ethylene naphthenate) (PEN), while among the aliphatic ones we can find poly(butylene succinate) (PBS), poly(ethylene succinate) (PES), poly(lactic acid) (PLA), poly( $\epsilon$ -caprolactone) (PCL), *etc.*<sup>61</sup>

Promising results have been obtained in the synthesis of a polyester based on the structure of PET, in which the ethylene glycol has been replaced with isosorbide. The polycondensation of isosorbide with terephthalic acid and its acid dichloride led to the production of poly(isosorbide terephthalate) (PIsT), as shown in Fig. 22.<sup>65</sup>

Given the versatility of the hydroxyl groups, **1** can also be converted into the respective diacids and/or diesters and used in the formulation of ester-based copolymers with innovative and interesting structures.<sup>66</sup> Recently, Wu *et al.*<sup>67</sup> were able to synthesize and thoroughly characterize a class of polyesters (shown in Fig. 23) that were completely based on the carbon skeleton of isosorbide, for both the diol and the diester components, named poly(isosorbide isosorbate) (PIsI).

This review has demonstrated the possibility of optimizing the atom economy of polyester production processes and the sustainable derivation of the raw materials used to obtain polymeric products, with characteristics comparable to those currently on the market for that specific commercial use.<sup>68,69</sup>



Fig. 22 Isosorbide as a monomer for the synthesis of PET analogue PIsT.<sup>65</sup>

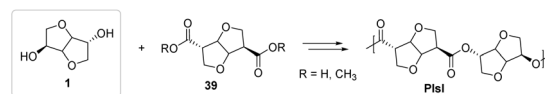


Fig. 23 Fully isosorbide-based polyesters.<sup>67</sup>

Still in the field of aliphatic polyesters, Wu *et al.*<sup>70</sup> have been able to carry out a CALB-catalyzed enzymatic polymerization (EP) of diethyl succinate, 1,4-butanediol (1,4-BDO) and isosorbide, obtaining copolyesters of poly(butyleneisosorbide-*co*-succinate) (PBIsS) in good yields. The polymeric materials produced by the authors (Fig. 24) showed a tunable temperature of glass transition and crystallinity in a wide range, which could make them suitable for new fields of applications, such as textile fibers or in engineering plastic materials.<sup>71,72</sup>

The most relevant scientific cases concerning the use of isosorbide derivatives in the innovative polymer industry will be discussed below, starting from aliphatic copolyesters based on **39**, the methyl diester of isosorbide.

As shown in Fig. 25, the methyl diester of isosorbide has been successfully employed as one of the two diacid reagents in a polycondensation process that included the employment of dimethyl adipate and 1,4-BDO.<sup>73</sup> The authors were able to fully demonstrate the production of novel systems with improved viscoelastic properties with an average molecular weight of approximately 24 000–47 600 g mol<sup>-1</sup>, as well as surprising self-adhesion capabilities and consequent industrial weldability.

Thiyagarajan *et al.*<sup>74</sup> reported the synthesis of AB-type homopolyesters based on *exo*-OH monomers of isohexides, with molecular weights comparable to those witnessed from the BB- or AA-type analogues.<sup>75</sup> With the same molecular weight of 2500 g mol<sup>-1</sup>, the thermal properties of the produced samples were found to be suitable for application at higher temperatures. Poly(isoidide-*co*-isoidide dicarboxylic acid) (PII) and poly(isosorbide-*co*-isoidide dicarboxylic acid) (PIsI) were the references, and the samples proposed by the authors consisted of a set of six compositions of isohexides and their respective methylcarboxylates.<sup>74</sup>

The various bifunctional derivatives of isosorbide, isomannide and isoidide also have high application potential in the polymer industry.<sup>76</sup> They were successfully included in the production of biobased polymers, solvents, fuel additives, nitrogenous-based plastics such as polyamides, polyimides and polyurethanes, and many others.<sup>13</sup>



Fig. 24 Fully aliphatic isosorbide-based copolyester *via* enzymatic polymerization using lipase.<sup>70</sup>





Fig. 25 Fully aliphatic isosorbide-based copolyester via melt polycondensation process.<sup>73</sup>

**3.3.3. Polyimides.** Polyimides (PIs) are polycondensation polymers characterized by the presence of an imide bond ( $-\text{CO}-\text{NR}-\text{CO}-$ ) and they are synthesized starting from dianhydrides and dianilines. PIs based on aromatic structures have shown a wider spectrum of application.<sup>77</sup>

Isohexidic functionalities were successfully inserted into the structures of partially biobased polyimide monomers by Thiyagarajan *et al.*<sup>78</sup> The presence of isosorbide in the repeating chain resulted in polymeric samples exhibiting higher mechanical stiffness and thermal stability comparable to their petrochemical counterparts. In fact, the innovation resided in their excellent optical transparency and optical activity properties, attributable to the chiral cavities of isosorbide.<sup>79</sup>

Poly(ether imide)-based polymers (PEIs) that exploit the structure of isosorbide as a co-monomer to impart special properties to the produced materials have been successfully synthesized and fully characterized (Fig. 26).<sup>80</sup> The authors were able to demonstrate that most of the samples derived from isosorbide possessed thermal properties (namely glass transition temperatures), tensile strengths and Young's moduli comparable to those of petroleum-based poly(ether imide)s.

**3.3.4. Polyurethanes.** Polyurethanes (PUs) are polymers characterized by the urethane group ( $\text{R}-\text{NH}-\text{CO}-\text{O}-$ ) and can be obtained by a polyaddition reaction, generally starting from a diol and a diisocyanate, even if in most cases the starting material is made up of polyfunctional monomers.<sup>81</sup>

Polyurethanes have a very diversified field of application, but are commonly used to produce both flexible and rigid foams.<sup>82</sup> An important structure–property correlation study of innovative polymers within the class of sustainable polyurethanes was carried out by the Kieber *et al.*<sup>83</sup> The authors provided an interesting screening of the possibility of employing different

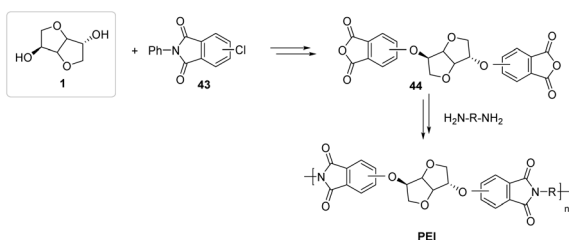


Fig. 26 Poly(ether imide)s (PEIs) synthesized starting from isosorbide-derived dianhydrides.<sup>80</sup>

stereochemical isomers of isohexides in the synthesis of polymeric materials with molecular weights in the range of 6–34 kDa.

The isosorbide-based PU in Fig. 27 showed the best results in terms of mechanical strength, higher density, reduced free volume and good thermal stability compared to the counterpart considered,<sup>84,85</sup> based on biomass-derived diols<sup>86</sup> of hydroxymethyl furfural (HMF).

**3.3.5. Polyacetals.** An interesting strategy was developed by Rajput *et al.*<sup>87</sup> for the synthesis of a narrow range of diacetal isohexides, intended for producing chiral building blocks for the polymer industry, to be used in pharmaceuticals and medicine.<sup>87</sup>

Polyacetals are polymeric materials with simple bond repeating units ( $-\text{CH}_2-\text{O}-$ ) and are commonly used in the production of components for household appliances, gears, bearings and automotive parts, and components for biomedical devices.<sup>88</sup> They are used as commodities and, given the large market demand, the possibility of replacing oil-based components with more sustainable solutions appears to be an excellent starting point for the green transition of the economy in the context of polyacetals.

Isosorbide-based diacetals were subjected to acid-catalyzed polycondensation (Fig. 28) under mild conditions. The authors were able to synthesize innovative materials with good yields, appearance and intrinsic characteristics, such as a molecular weight of  $3200-27600 \text{ g mol}^{-1}$  for the isolated product. This publication led several groups of researchers to investigate the possibility of using isosorbide in this field.<sup>89,90</sup>

**3.3.6. Polyamides.** Another class of polymeric materials in which the typical functionalities of isohexides can be exploited to obtain innovative materials is constituted by polyamides (PAs).

PAs are macromolecular materials characterized by an amide bond ( $-\text{CO}-\text{NH}-$ ) in the main chain. In polymers, this bond is established by a polycondensation reaction of a  $-\text{COOH}$  terminal of a carboxylic acid and an  $-\text{NH}_2$  terminal of an amine.<sup>91</sup> The amide bond can resonate between two structures and can assume a partial double-bond character, thus giving the bond a planar geometry, a certain rigidity and a non-negligible Debye dipole moment. Commonly, these



Fig. 27 Polyurethane synthesis starting from biomass-derived monomers.<sup>85</sup>



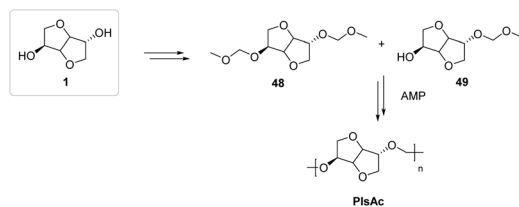


Fig. 28 Isosorbide-based polyacetals synthesized through acetal metathesis polymerization (AMP).<sup>87</sup>

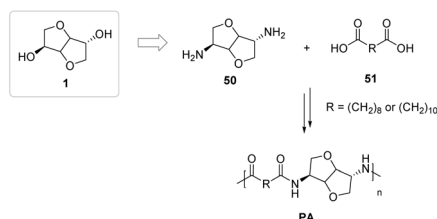


Fig. 29 Polyamide synthesis starting from isosorbide-based diamine and aliphatic natural dicarboxylic acids.<sup>95</sup>

characteristics allow macromolecules to be organized in *trans*-planar linear structures.

In addition, the spatial arrangement of the chains allows for effective overlap and the possibility of forming hydrogen bonds translates into a high degree of crystallinity that reaches values of up to 40–50%, and molecular weights of  $10^4 \text{ g mol}^{-1}$ .<sup>92</sup>

Jasinska-Walc *et al.*<sup>93</sup> were able to synthesize fully biobased polyamides from diamino-isosorbide and sebacic or brassylic acid,  $C_{10}$  and  $C_{13}$  aliphatic diacids derived from castor oil, as shown in Fig. 29. To date, all polyamides produced on an industrial scale from monomers derived from biomass have been based on monomers derived from castor oil.<sup>94</sup> A problem that has not yet been addressed is the competition with the food supply chain for the use of building blocks of natural origin to produce polymeric commodities. The use of isohexides such as isosorbide could constitute the green breakthrough so sought after by many researchers in the sector of macromolecular products.<sup>95</sup>

In Section 3, the current discussion on the sustainability of the monomer feedstock supply chain, and on the main potential application fields of isosorbide-based compounds in the polymeric materials industry, has been thoroughly highlighted.

## 4. Conclusions

In this review, a comprehensive survey of peculiar compounds with isohexidic structure was performed with a focus on isosorbide **1**, a dianhydro hexitol with two *endo-exo* OH groups in positions 3 and 6, respectively. The main properties that distinguish isohexides from any other reactant class have been addressed in Section 1, with particular attention to the correlations between the molecular structure and properties of the final product. In Section 2, it was discussed how isosorbide can be successfully synthesized *via* common dehydration of hexitols

or separated from vegetable waste biomasses that do not compete with the agri-food industry. Isosorbide's interesting characteristics, which can be exploited for the replacement of different diol compounds commonly derived from the crude oil sector, have been emphasized starting from Section 3. The singular structure of isosorbide is based on two tetrahydrofuran rings fused with a *cis*-type junction and the spatial arrangement of the two hydroxyl substituents allows this isohexide to be used for its intrinsic chiral property as a chiral auxiliary and/or as a reactant.

In the last part of the review, the reader can find in-depth bibliographic research with salient examples of uses of isosorbide in the synthesis of polymeric materials, with a high percentage of biobased components, thanks to its high reactivity and the possibility of forming intramolecular hydrogen bonds.

For instance, the use of isosorbide as a monomer, comonomer and precursor has been demonstrated in the production processes of different types of polymers, such as epoxy resins, polyesters, polyimides, polyurethanes, polyacetals, and polyamides with good to excellent yields and high average molecular weights.

Overall, this review has shown how and why isosorbide can be considered a compound of great industrial relevance in many chemical sectors. The reader will be able to recognize the potential exploitation of isosorbide in the field of asymmetric synthesis, the realization of chiral ionic liquids and the production of polymeric materials with high applicative potential. Following the author's personal interpretation of the cited literature, the differences that can be encountered when isosorbide is used as a reactant or auxiliary diol, instead of the traditional diols deriving from the petrochemical chain, will be clarified.

## Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

## Conflicts of interest

There are no conflicts of interest to declare.

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