Plastiche: alcune cifre per capire meglio l'impatto

https://www.plasticseurope.org/it/resources/publications/180 4-plastics-facts-2019

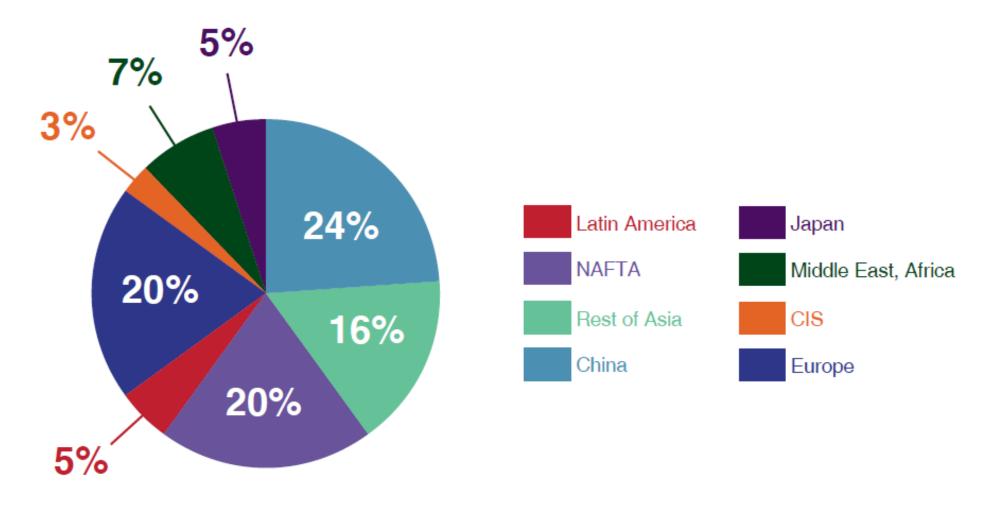


Plastics – the Facts 2019

An analysis of European plastics production, demand and waste data



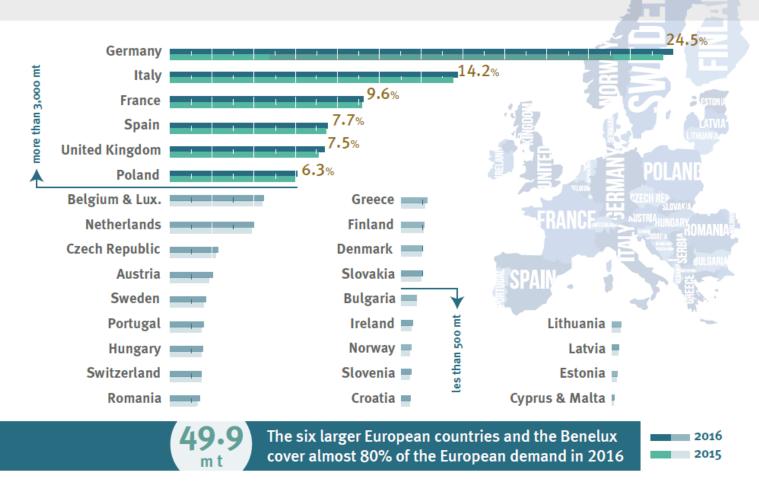
FIGURE 4: PLASTIC PRODUCTION PER REGION, 2012



Based on PlasticsEurope data [1]

Plastics EU converter demand per country

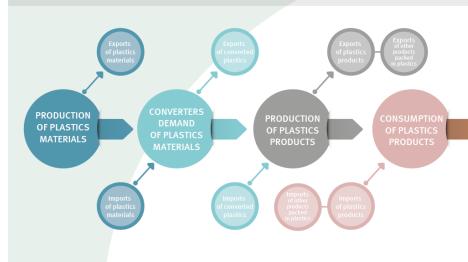
European plastic converter demand includes plastic materials (thermoplastics and polyurethanes) and other plastics (thermosets, adhesives, coatings and sealants). Does not include: PET fibers, PA fibers, PP fibers and polyacryls-fibers. Source: PlasticsEurope Market Research Group (PEMRG) and Conversio Market & Strategy GmbH (Consultic GmbH for 2015 data)



Life cycle of plastics

Understanding the life cycle of plastics products

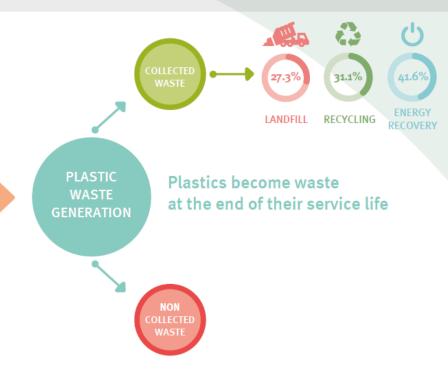
In order to understand the life cycle of plastics products it is important to understand that not all plastics products are the same and not all have the same service life. Some plastic products have a shelf life of less than one year, some others of more than 15 years and some have a lifespan of 50 years or even more.



Thus, from production to waste, different plastic products have different life cycles and this is why the volume of collected waste cannot match, in a single year, the volume of production or consumption.

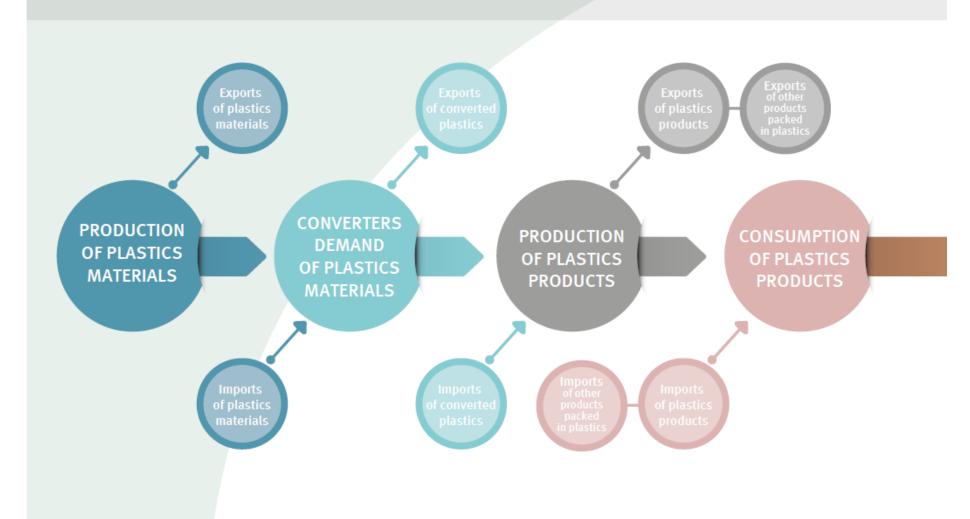
The service life of plastics products goes from less than 1 year to 50 years or more

LIFE SERVICE OF PLASTICS PRODUCTS



Understanding the life cycle of plastics products

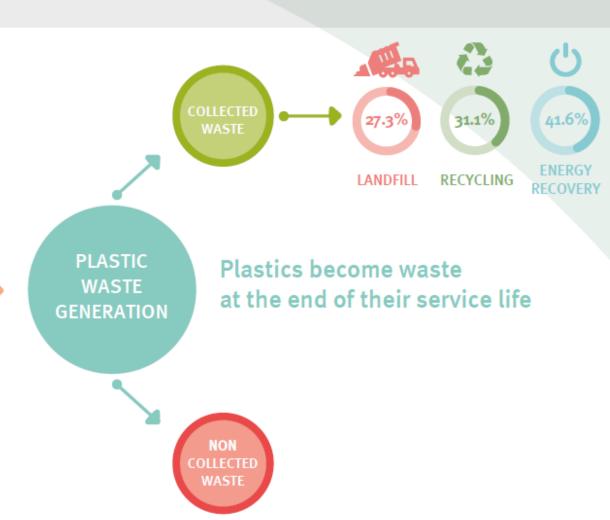
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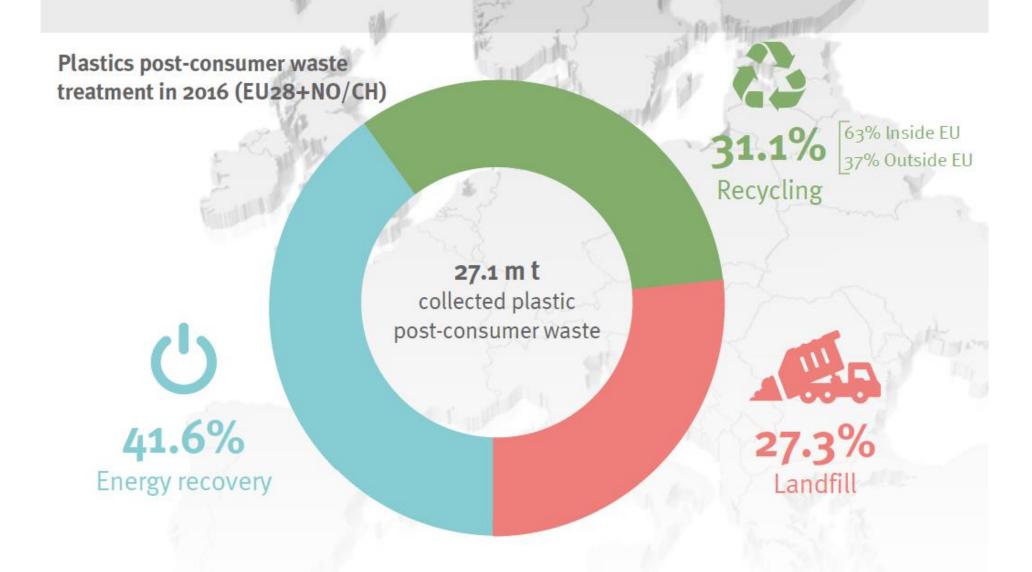
The service life of plastics products goes from less than 1 year to 50 years or more

LIFE SERVICE OF PLASTICS PRODUCTS



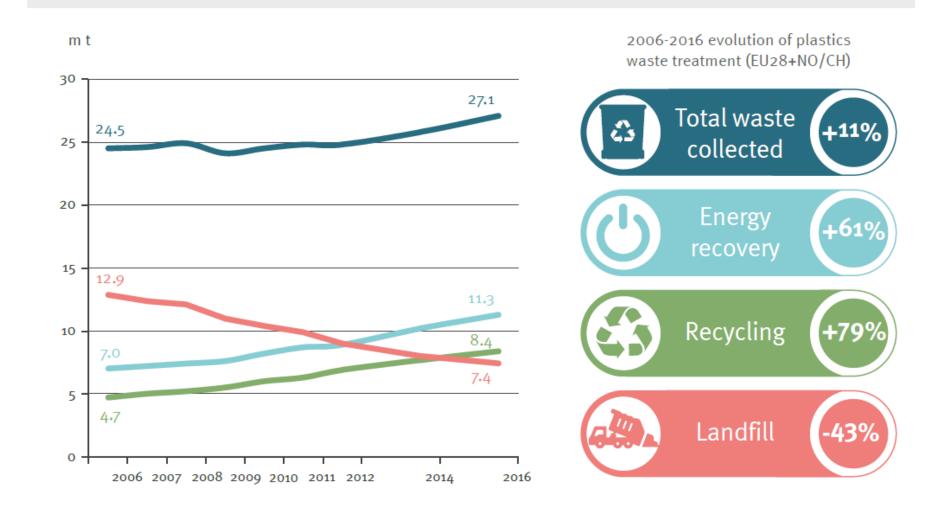
In 2016, for the first time, recycling overcame landfill

In 2016, 27.1 million tonnes of plastic waste were collected through official schemes in the EU28+NO/CH in order to be treated. And for the first time, more plastic waste was recycled than landfilled.



In ten years, plastic waste recycling has increased by almost 80%

From 2006 to 2016 the volumes of plastic waste collected for recycling increased by 79%, energy recovery increased by 61% and landfill decreased by 43%.



Plastic waste recovery is still very uneven in Europe

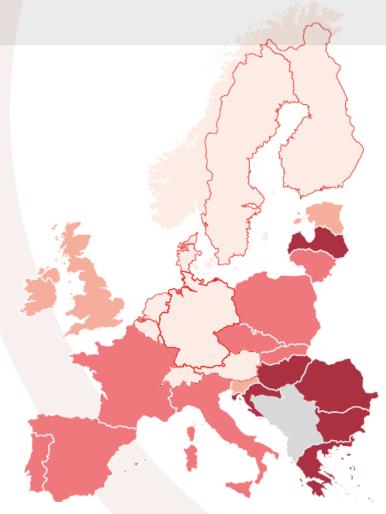
Although the total EU situation is improving, in many countries, landfill is still the first or second option of treatment for plastic post-consumer waste.

Source: Conversio Market & Strategy GmbH

Plastics post-consumer waste landfill rate across Europe

- 10% or less
- up to 30%
- up to 50%
- more than 50%
- Countries with landfill restrictions implemented

Plastics waste going to landfill in 2016



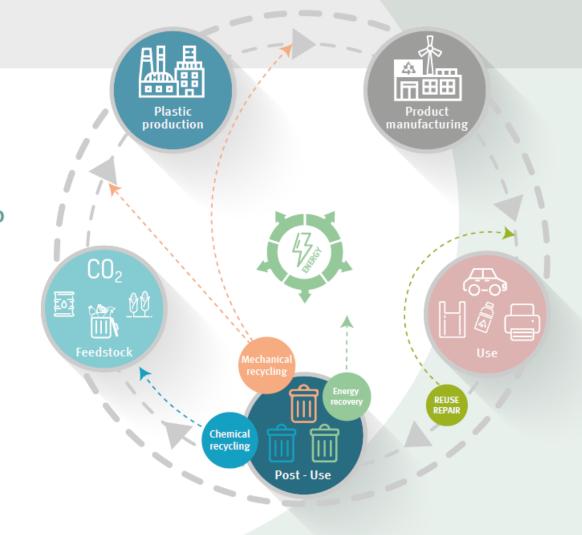
Full life cycle thinking is smart thinking

https://www.plasticseurope.org/it/resources/videos/403-full-

life-cycle-thinking-italian



At the end of their life, plastics are still very valuable resources that can be transformed into new feedstock or into energy.



THE CIRCULAR ECONOMY FOR PLASTICS

A European Overview



https://www.plasticseurope.org/it/resources/publications/189 9-circular-economy-plastics-european-overview Collecting and recycling plastics represents an answer to the problem and analysis by the Ellen MacArthur Foundation has shown that "replacing just 20% of single-use plastic packaging's with reusable alternatives offers opportunities for economic development worth at least 10 billion USD" [11].

Notably, in Europe, collection increased from 27.1 million tonnes in 2016 to 29.1 million tonnes in 2018. About 9.4 million tonnes (32.5%) of them were recycled inside or outside Europe while 24.9% ended up in landfills and the rest was incinerated [7].

Why recycling is not sufficient 1.

It must be underlined that plastics made from fossil fuels account for 20% of the total fossil oil consumption [11] and their manufacture, recycling and incineration are energy intensive and cause considerable greenhouse gases (GHG) emissions.

Analyses indicate that if plastic continues to be produced from fossil carbon sources, it will be responsible for a 15% of the maximum annual global carbon budget, needed to limit global warming to 2°C in 2050 [12].

Therefore, the use of **recycled materials**, when technically and legally feasible, is essential to **enable the dissociation of plastic production** from the exploitation of finite carbon sources. However, in a long-term perspective, it is necessary to boost a **transition to plastics obtained from renewable feedstock**, whenever robust evidences support their environmental and social benefits.

Why recycling is not sufficient 2.







Not all plastics/polymers can be recycled



La direttiva Europea SUP: single use plastics

(Atti legislativi)

L 155/1

DIRETTIVE

DIRETTIVA (UE) 2019/904 DEL PARLAMENTO EUROPEO E DEL CONSIGLIO

del 5 giugno 2019

sulla riduzione dell'incidenza di determinati prodotti di plastica sull'ambiente

(Testo rilevante ai fini del SEE)

Articolo 1

Obiettivi

Gli obiettivi della presente direttiva sono prevenire e ridurre l'incidenza di determinati prodotti di plastica sull'ambiente, in particolare l'ambiente acquatico, e sulla salute umana, nonché promuovere la transizione verso un'economia circolare con modelli imprenditoriali, prodotti e materiali innovativi e sostenibili, contribuendo in tal modo al corretto funzionamento del mercato interno.

TOP 10 SINGLE-USE PLASTIC ITEMS FOUND ON SEA SHORES

- 1 Drink bottles, caps and lids
- Cigarette butts
- **Cotton buds sticks**
- Crisp packets/sweet wrappers
- 5 Sanitary applications (sanitary towels, tampons etc.)
- 6 Plastic bags
- (7) Cutlery, straws and stirrers
- 8 Drinks cups and cup lids
- Balloons and balloon sticks
- (10) Food containers, including fast food packaging

Items in yellow could soon be banned as non-plastic alternatives are





WAYS TO REDUCE **YOUR SINGLE-USE PLASTICS** There are an estimated **5 TRILLION** pieces of plastic in the ocean worldwide, with **8 MILLION** metric tons added to the ocean each year*. Wildlife are dying at a rapid pace due to the ingestion of or entanglement in plastics. PLASTIC AND PLASTIC TRAVEL PRODUCE BAGS CLOTH BAGS BEVERAGE COFFEE LIDS **COFFEE MUG** CONTAINER YOU can help reduce waste by removing single-use plastics from your everyday activities. SINGLE-SERVE A POT OF COFFEE COFFEE PLUGS TRAVEL COFFEE PODS AND STIR STICKS COFFEE MUG DISPOSABLE MATCHES BALLOONS STRAW FREE #StopSingleUse



Plastic is responsible for around 10% of the generated total waste and composes 60-90% of the marine litter, mostly with food and beverage packaging, cigarette butts and bags. According to the United Nations Environment Programme (UNEP), 8 million tonnes of plastic are poured into the seas each year, an equivalent to a full garbage truck every minute [6][7].

Marine species and humans are being harmed since the plastic waste enters the human food chain through fish consumption, [8] while the rapid spread of microplastics has made this problem even more alarming [9]. Because it is not effective to remove plastic waste and microplastics once they have entered the sea, plastic pollution needs to be tackled at its source [10].

The hidden natural capital cost of fossil-based plastics in numbers

A clear understanding of the environmental degradation and resource depletion connected to plastics must rely on a quantitative and transparent accounting of their impact on natural capital.

The term **Natural Capital** [13] describes "Earth's natural assets, including soil, air, water, and living things, existing as complex ecosystems, as well as the related ecosystem services that human societies need in order to survive and thrive".

Economic activities depend on these resources and services; however, the latter are often not factored into corporate accounting, and national accounts currently do not fully take their contribution into consideration.





In 2014, UNEP published a study focused on the evaluation of the natural capital costs of plastics, namely the environmental and social impacts caused by the use of plastic expressed in monetary terms to reflect the scale of the caused damage [6].

The study converted physical quantities of plastic into monetary values, using environmental or natural capital valuation techniques [14]. These techniques estimate the value of environmental goods or services in the absence of a market price and aggregate them into a single figure.

As an example, by calculating the amount of GHG caused by plastic production it is possible to ascribe a monetary value on each tonne of GHG in relation to its impact on climate change. Similarly, plastic waste incineration is associated to air pollution, which can be expressed in monetary terms, thus reflecting the scale of damage caused.

On this basis, the UNEP study estimated that the total natural capital cost of the plastic used in the consumer goods industry is above **US\$75 billion** per year.

Such approach translates physical impacts into a monetary figure, which expresses the potential value that companies would have to internalise if they were held accountable for their impacts.

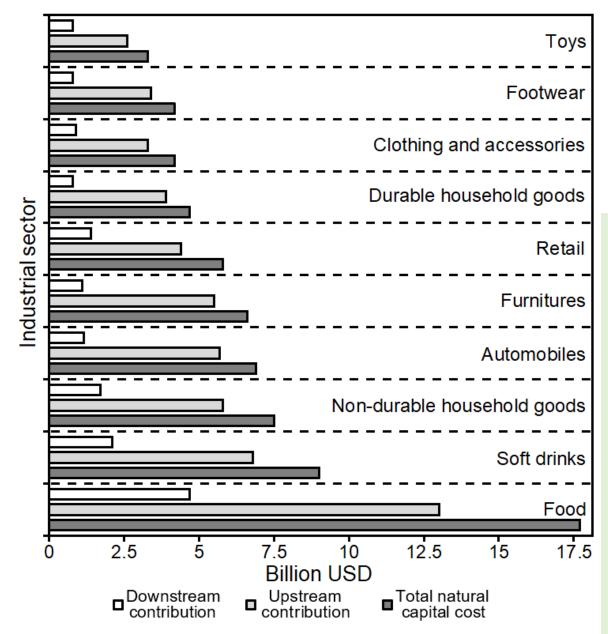
Upstream and downstream impacts

When considering that most types of plastic are petroleum-based products, one significant outcome of the UNEP analysis is that, across all sectors, over 75% of the known and quantifiable impacts associated with plastic usage are located in the upstream portion of the supply chain.

'Upstream' refers to "impacts generated from the extraction of raw materials to the manufacturing of plastic feedstock",

'downstream' refers to "impacts generated once the consumer has discarded the product".

For example, the downstream impact of consumer electronics is only 17% of its total impact due to established recycling initiatives, whereas the tobacco sector has the largest downstream impact (29%) of its total impact due to the littering of cigarette butts.



Total capital cost for some of the most relevant industrial sectors contributing to the plastic problem. The calculated costs arise from the accounting of the tonnage of plastic used in the selected key consumer goods sector (based on its expenditure), which gave the plastic intensity.

The natural capital cost for a certain sector is the **natural capital intensity** multiplied by the aggregate revenues and it expresses "the dependency of a certain sector on the natural capital".

The plastic intensity in each sector reflects different contributions of the three main categories of plastic usage:

- i) plastic used in products;
- ii) ii) plastic used as packaging;
- iii) iii) plastic used by suppliers (such as bags containing fertilizer used by farmers supplying the food sector).

This methodological approach made evident that, for the food, soft drink, retail and personal products sectors, the whole contribution comes from the packaging.

On average, the total natural capital cost of plastic use is 52% of the total economic cost. When only considering the upstream natural capital cost as a percentage of plastic prices, the potential cost increase is 44% on average. This means that, if the upstream impacts of plastic were taken into account and fully paid by businesses, the price of plastic would be 44% higher on average.

With current knowledge, the analysis indicates that, across consumer goods sectors, over 30% of the natural capital costs originates from GHG released in the upstream supply chain. The most significant downstream impact is marine pollution, which has a natural capital cost of at least \$13bn, and includes economic losses incurred by fisheries and tourism as well as time spent cleaning up beaches.

The environmental impacts associated with plastic use was calculated using lifecycle analysis techniques, using official databases as the US Toxic Release Inventory [15] and Plastics Europe eco-profiles [16]. The impact of additives leachate from plastics was also accounted, since there is a growing concerns on their impacts on human health and the environment [17,18]. Additives are added to plastic during their manufacturing to improve their mechanical and thermal properties and the study calculated the amount of additives per type of plastic based on a report of the Organisation for Economic Co-operation and Development [19]. According to the same report, the annual leaching rate of additives is 0.16% per year, which means that it would take 625 years for 100% of the additives to be released from the plastics. These data were used to quantify and, ultimately, valuate the toxic impact of plastics additives.

Overall, the disclosing of the natural capital costs of plastics highlighted "the urgent need for businesses to measure, manage and disclose information on their annual use and disposal of plastic, as many companies already do with carbon emissions" [6].





Companies need a single tool that measures environmental impacts in an integrated way together with other business issues. This is where **natural capital valuation** comes in. The technique enables companies to put a financial value on a range of impacts, including plastic, so environmental management can be fully embedded within the business.

This research then analyses the exposure of companies to these risks and opportunities by expressing quantities of plastic used as a natural capital cost. The results show that the total natural capital cost of plastic used in the consumer goods industry is over \$75bn per year. Broken down by sector, food companies are by far the largest contributor to this cost, responsible for 23% of the total natural capital cost (see figure 1). The results also show each sector's natural capital intensity – or its natural capital cost per \$1m of annual revenue. The toy sector has by far the highest intensity, at 3.9% of revenue.

FIGURE 1: TOTAL NATURAL CAPITAL COST AND INTENSITY OF SELECTED SECTORS

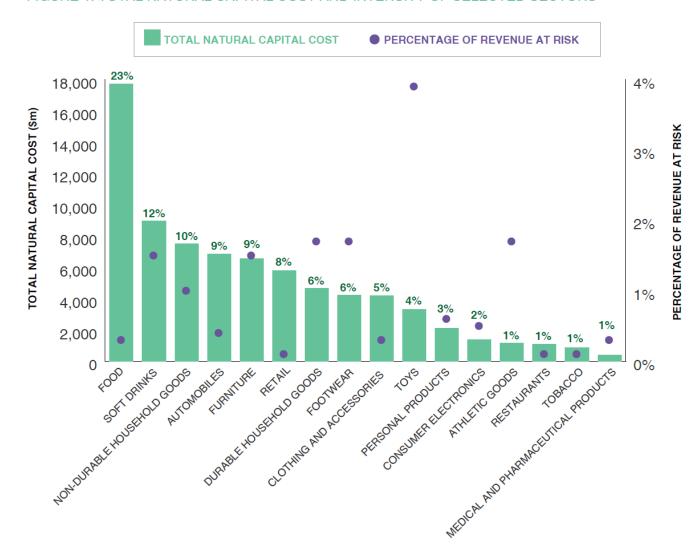


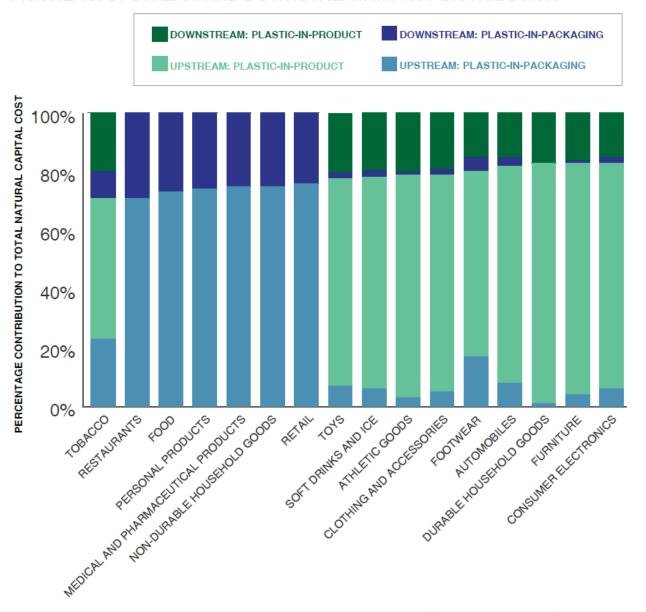
TABLE 3: TOTAL NATURAL CAPITAL COST (\$) AND NATURAL CAPITAL INTENSITY

SECTOR	TOTAL NATURAL CAPITAL COST (\$M)	TOTAL NATURAL CAPITAL COST (\$M) PER SERVICE YEAR
Food	17,700	8,900
Non-durable household goods	7,500	7,100
Soft drinks	9,000	4,500
Retail	5,800	2,900
Footwear	4,200	2,700
Tobacco	900	2,200
Personal products	2,100	1,000
Clothing and accessories	4,200	1,500
Furniture	6,600	1000
Durable household goods	4,700	800
Toys	3,300	700
Consumer electronics	1,400	600
Restaurants	1,100	500
Automobiles	6,900	500
Athletic goods	1,100	200
Medical and pharmaceutical products	400	200

TABLE 2: PLASTIC INTENSITY PER SECTOR (TONNES PER \$1m REVENUE)

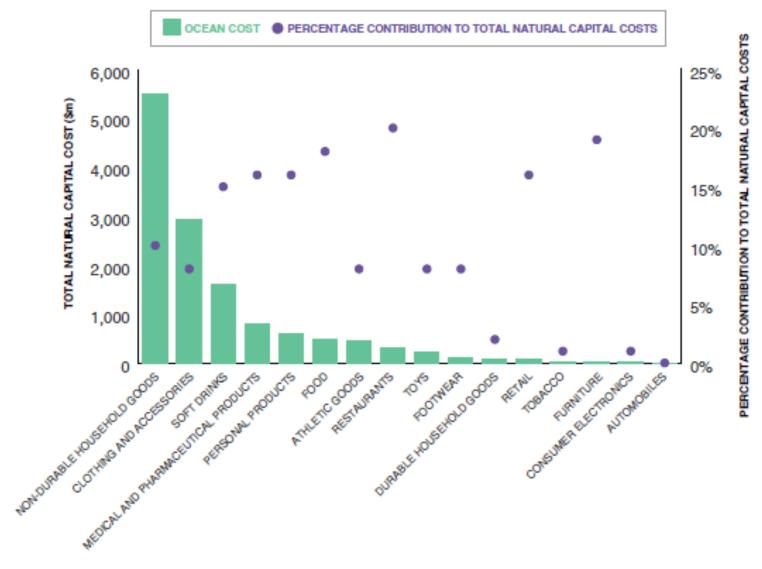
SECTOR	PLASTIC-IN-PACKAGING	PLASTIC-IN-PRODUCT	PLASTIC-IN-SUPPLY-CHAIN	TOTAL
Toys	2.9	37.5	7.6	48.0
Soft drinks	14.9	-	19.7	34.6
Furniture	0.8	15.2	10.9	26.9
Durable household goods	1.6	16.2	7.2	25.0
Footwear	3.6	13.8	6.8	24.2
Athletic goods	0.8	16.7	4.0	21.5
Personal products	5.9	not estimated	10.0	15.9
Non-durable household goods	1.0	9.7	3.8	14.4
Automobiles	0.0	4.5	5.3	9.9
Food	3.2	-	6.3	9.5
Consumer electronics	0.4	4.7	3.6	8.7
Clothing and accessories	0.2	3.3	4.7	8.2
Medical and pharmaceutical products	3.1	-	3.5	6.6
Restaurants	1.2	-	3.2	4.4
Tobacco	0.3	0.7	2.5	3.5
Retail	0.5	-	1.5	2.1

FIGURE 10: UPSTREAM AND DOWNSTREAM IMPACT DISTRIBUTION



Trucost calculations derived from, but not limited to, World Bank [7]; PlasticsEurope [8]; Eurostat [9], and the US EPA [10] datasets (full set of references and methodology available in appendices 3 and 4 of this report)

FIGURE 13: TOTAL NATURAL CAPITAL COST OF PLASTIC IN MARINE ECOSYSTEMS (\$)



Total natural capital costs correspond approximately to over 80 million tonnes of plastic. Trucost calculations derived from, but not limited to, World Bank [7]; PlasticsEurope [8]; Eurostat [9], and the US EPA [10] datasets (full set of references and methodology available in appendices 3 and 4 of this report)

Sectors with the highest supply-chain-plastic intensity in relation to the total plastic intensity are the retail (75%), restaurants (72%), tobacco (70%) and food (66%) sectors – meaning that companies operating in these sectors may need to pay attention to plastic used in their supply chains. The retail and restaurant sectors are located further down the supply chain compared to others. Their low direct intensity is because it only includes packaging added to the product in the shop, such as carrier bags. This explains the low intensity of the sector's direct operations and proportionately larger plastic-in-supply-chain intensity. The retail, restaurants, tobacco and food sectors are also significant users of agricultural commodities in their supply chain, which has been recognized as a plastic intensive sector. As highlighted by Plastics Europe, agriculture contributed 4.2% to the overall plastic demand in Europe in 2012.1

FIGURE 7: PLASTIC INTENSITY PER SECTOR (TONNES PER \$1M REVENUE)

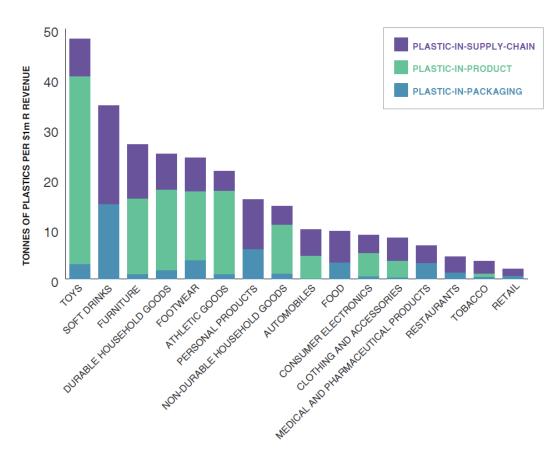
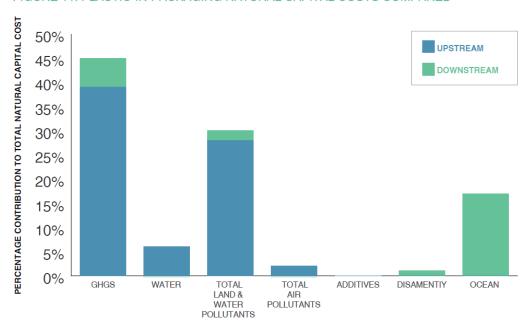
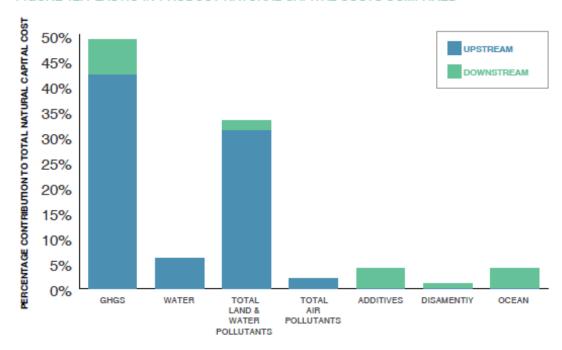


FIGURE 11: PLASTIC-IN-PACKAGING NATURAL CAPITAL COSTS COMPARED



Trucost calculations derived from, but not limited to, World Bank [7]; PlasticsEurope [8]; Eurostat [9], and the US EPA [10] datasets (full set of references and methodology available in appendices 3 and 4 of this report)

FIGURE 12: PLASTIC-IN-PRODUCT NATURAL CAPITAL COSTS COMPARED



Trucost calculations derived from, but not limited to, World Bank [7]; PlasticsEurope [8]; Eurostat [9], and the US EPA [10] datasets (full set of references and methodology available in appendices 3 and 4 of this report)

Albis Plastic GmbH (D, www.albis.com)

- Agrana (A, www.agrana.at)
- Amynova Polymers GmbH (D, www.amynova.com)
- Arichemie GmbH (D, www.arichemie.com)
- Arkema SA (F, www.arkema.com/en/)
- BASF SE (D, www.basf.com/en.html)
- Bayer MaterialScience AG (D, www.bayermaterialscience.de)
- BioAmber (CA, www.bio-amber.com/)
- Bio-On srl (I, www.bio-on.itwww.minerv.it)
- Biomater Ltda.(BR, www.biomater.com.br)
- Biomer (D, www.biomer.de)
- BIOP Biopolymer Technologies AG (D, www.biop.eu)
- Biotec Biologische Naturverpackungen GmbH & Co. KG (D, www.biotec.de)
- BioTec Environmental (USA, www.bio-tec.com)
- Borregaard (N, <u>www.borregaard.com</u>)
- Braskem (BR, www.braskem.com.br/)
- Cardia Bioplastics (AU, www.cardiabioplastics.com)
- Cereplast (USA, <u>www.cereplast.com</u>)
- Cargill Inc. (USA, <u>www.cargill.com/</u>)
- Clarifoil (UK, <u>www.clarifoil.com</u>)
- DaniMer Scientific, LLC (USA, <u>www.danimer.com</u>)
- Dow Chemical (D, www.plastics.dow.com)
- Dow Wolff Cellulosics (D, www.dowwolff.com)
- DuPont (USA, www.dupont.com/Plastics)
- Eastman Chemical GmbH (D, www.eastman.com)
- Fkur Kunststoff GmbH (D, www.fkur.com)
- Futerro (B, www.futerro.com/)
- Gehr GmbH (D, www.gehr.de)
- Genomatica (USA, <u>www.genomatica.com/</u>)
- Goodfellow Cambridge Ltd. (UK, <u>www.goodfellow.com</u>)
- HallStar (USA, www.hallstar.com)
- Horn & Bauer GmbH & Co. KG (D, www.horn-bauer.de)
- Huhtamaki Deutschland GmbH & Co. KG (D, www.huhtamaki.com)
- Innovia Films Ltd. (UK, <u>www.innoviafilms.com/</u>)
- Ire Chemical Ltd. (K, www.irechem.co.kr)
- Jinhui Zhaolong High Tech (CN, www.ecoworld.jinhuigroup.com)
- Kaneka Corp. (JP, www.kaneka.co.jp/kaneka-e/)
- Kareline Oy Ltd (FIN, www.kareline.fi)
- Kingfa (CN, www.kingfa.net/)
- Koninklijke DSM N.V. (NL, www.dsm.com/corporate/home.html)
- Lanxess AG (D. lanxess.com/en/corporate/home/)
- Limagrain Cereales Ingredients (F, www.lci.limagrain.com/)
- Masterbatch Winter Herstellungs und Vertriebs GmbH (D, www.masterbatch-winter.de/)

Main players in the bioplastic sector

- Mazzucchelli 1849 SPA (I, www.mazzucchelli1849.it)
- Meredian Inc. (USA, www.meredianinc.com/)
- Metabolix Inc. (USA, www.metabolix.com/)
- Mitsubishi Chemical USA Inc (USA, www.mitsubishichemical.com)
- Mitsui Chemical Europe GmbH (D, www.mitsuichem.com)
- NAPAC Schweiz AG (CH, www.napac.ch)
- Natureplast (F, www.natureplast.eu)
- NatureWorks LLC (USA, www.natureworksllc.com/)
- Novamont SpA (I, www.novamont.com)
- Novomer (USA, www.novomer.com/)
- Ofotec Folien GmbH (D, www.ofotec-folien.de)
- Perstorp AB (S, www.perstorpcaprolactones.com)
- Peter Holland BV (NL, www.peterholland.nl)
- Plantic Technologies Ltd (AU, www.plantic.com.au)
- Polymer Chemie GmbH (D, www.polymer-chemie.de)
- Polyone Corp. (USA, www.polyone.com/en-us/Pages/default.aspx)
- Procter & Gamble (D, www.pg.com)
- Purac (NL, www.purac.com)
- Radici Plastic GmbH & Co. KG (D, www.radiciplastics.de)
- Rhein Chemie Rheinau GmbH (Lanxess Group, D, www.rheinchemie.com/)
- Rodenburg Biopolymers B.V. (NL, www.biopolymers.nl)
- Shanghai Disoxidation Macromolecule Materials Co., Ltd (CN, www.dmmsh.com)
- Showa Denko (JP, www.shp.co.jp)
- So.F.teR. Spa (I, www.softergroup.com/)
- Solvay SA (B, www.solvay.com/)
- Sukano AG (CH, www.sukano.com)
- SwissGel AG (CH, <u>www.swissgel.ch</u>)
- Tate & Lyle PLC (UK, <u>www.tateandlyle.com/Pages/default.aspx</u>)
- Telles (USA, www.mirelplastics.com)
- Tecnaro GmbH (D, www.tecnaro.de)
- Tianan Enmat (CN, www.tianan-enmat.com)
- Thantawan Industry PLC (BioFoammat Division, T, www.biofoammat.com)
- Toray Industries Inc. (JP, <u>www.toray.com/</u>)
- Toyobo (JP, www.toyobo-global.com/),
- Unitika Ltd. (JP, www.unitika.co.jp/terramac)
- Vegeplast S.A.S. (F, www.vegeplast.com/uk)
- Ventura AG Kunststofftechnik (CH, www.ventura-ag.ch)
- Wentus Kunststoff GmbH (D, www.wentus.de)
- W.W. Textile Co. Ltd. (CN, www.2wtextile.com)

Table 2.Naturally biosynthesized biopolymers and their chemically modified derivatives.

Chemical classification	Polymer	Properties and applications	Ref.
Polyisoprene (terpenes)	Natural rubber	Waterproof items, engineering applications in antiseismic buildings or offshore installations for oil extraction,	
	Starch based polymers; thermoplastic starch -TS	Component of biodegradable and biocompostable plastics.	[39]
Polysaccharides	Cellulose based polymers: Cellulose acetate Cellulose nitrate Acetylphthalylcellulose	Applications in textiles, cigarette filters, surface coatings, ink additive, photographic negatives, motion picture film, microfilm, microfiche, membranes for water desalinization. Chemical modifications decrease the biodegradation of cellulose although derivatives are attacked by both aerobic and anaerobic microorganisms.	[40, 41]
	Chitosan	Obtained from deacetylation of chitin. Biodegradable, non-toxic, bacteriostatic and fungistatic with wide application in the pharmaceutical field. Industrially applied as carrier for enzyme immobilization.	[42]
Polyphenols	Lignin based polymers	Because of its aromatic and phenolic components, lignin itself is used in polymer blends as compatibilizer, plasticizer, hydrophobizing agent or as a natural antioxidant in active packaging. Employed in flame retardants, optical modifiers, stabilizers. Lignin-based polyols, reacted with diisocyanates, are used as drop-in replacement of fossil polyols in polyurethane foams for their flame-retardant properties.	[43-45]

Table 3. Bio-engineered polymers bio-synthesized by microorganisms and plants.

Chemical classification	Polymer	Properties and applications	Ref.
Polyesters	Polyhydroxy alkanoates - PHAs: poly(3-hydroxybutyrate) and poly(3-hydroxybutyrate -co-3-hydroxy-hexanoate)	Biodegradable and compostable. Chain length determines the flexibility of PHA: short chain butyrate provides rigidity, with T _m of 160°C, whereas longer carbon chains confers T _m below 145°C. Sensitivity to thermal degradation makes its processing challenging. Fields of application include agriculture, packaging, biomedical sector.	[46]
	Polymalic acid	Linear anionic polyester composed of L-malic acid monomers, with potential applications as drug carriers, surgical suture, and biodegradable plastics.	[47]
Polyamides	Poly-γ-glutamic acid - PGA	Water-soluble, anionic, biodegradable, edible. Applications in foods, pharmaceuticals, healthcare, cosmetics, water treatment, curable adhesives.	[48, 49]

★ Not biodegradable

Table 4. Bio-based synthetic polymers obtained from bio-based monomers or a combination of bio- and fossil-based monomers.

Polytimethylene terephthalate) - Polytester. Same properties as fossi-based PTT. Scarcey blodegradable. Semi crystalline thermoplastic, easily moliced or thermoformed and spun into fibres. Polytethylene terephthalate)-PET - Polytester. Same properties as the fossi-based PET. High-performance plastic used for engineering applications, fibres, films, bottles. Polytester, Same properties as the fossi-based PET. High-performance plastic used for engineering applications, fibres, films, bottles. Polytethylene succinate) - PES - Polytethylene succinate) - PES - Polytethylene burcinate) - PES - Polytethylene burcinate) - PES - Polytethylene burcinate) - PES - Polytethylene furancel - PEF - Polytester. Not brodegradable. T. of 172° C. 7, of 37° C. 30° Cod oxygen barrier and elongation properties. Used for film applications. Moderately blodegradable. To design a latenative to polytene furancel - PEF - Polytester. Not brodegradable. T. of 172° C. 7, of 37° C. 30° Cod oxygen barrier and elongation properties. Used for film applications. Moderately blodegradable. To design a latenative to polytester. The oxygen blodegradable. To design a latenative to polytester. The oxygen blodegradable. To design a latenative to polytester. The oxygen blodegradable. To design a latenative to polytester. The oxygen blodegradable. To design a latenative to polytester. The oxygen blodegradable. To design and polytester blodegradable. The polytester. The oxygen blodegradable. To design and polytester blodegradable. The polytester. The oxygen blodegradable to the polytester blodegradable. The polytester is a polytester blodegradabl		etic polymers obtained from bio-based monomers of a combination of bio- and lossif-based monomers.
POTY: Polytethylene succinate) - PBS Polytethylene succinate) - PEF Polytethylene furanoate) - PEF Polytethylene furanoate - PEF Polytethylene fur	Polymer	Properties and applications
Poly(ettylene succinate) - PBS Polyetetylene succinate) - PBS Polytetylene succinate) - PBS Polytetylene succinate) - PBS Polytetylene furnanoate - PES Polytetylene furnanoate - PES Polytetylene furnanoate - PTF Polytimetrylene furnanoate - PTF Polytetylene adjata-co-1,4- butylene terpetylhallate - PBP Polytetylene problemane - Pthan - Polytester - Not biodegradable. T _m of 172°C, T _m of 57°C, good oxygen barrier properties. Employed in light weighting packaging. Polytetylene pertylentialete - PBP Polytetylene problemane - PTF Polytetylene problemane - PTF Polytetylene problemane - PTF Polytetylene problemane - Pthan	PTT*	TGood tensile and flexural strength, excellent flow and surface finish. Used in textiles and engineering applications (automotive parts, mobile phone housings.
Polytethylene succinate) - PEF Polytethylene succinate) - PEF Polytethylene succinate) - PEF Polytethylene furnoate) - PEF Polytene furnoate) - PEF Pol	Poly(ethylene terephthalate)-PET*	
Polyterhylene furanasel) - PEF Polysetser. Durable, good oxygen barrier. T., of 211 °C and T., of 89 °C. Suitable for packaging, in the food and beverage industry. Polytburylene furanasel) - PBF Polytonylene adpate-cer. 174 butylene adpate-cer. 144 butylene adpate-cer. 144 butylene adpate-cer. 145 butylene adpate-cer.	Poly(1,4-butylene succinate) - PBS	an alternative to polyolefins.
Polyfurfmethylene furanaste) - PTF Polyester. Not biodegradable. T., of 17°C, god owygen barrier properties. Employed in light weighting packaging. Polyfurfmethylene furanaste) - PBF Polyester. T., of 17°C, T., of 47°C, pod 40°C, bottomial replacer of PET and PBF. Polyfurfmethylene and polyester resins - Upcertee ware according the percentage of unsaturated diacid (e.g. itaconic acid) and the curing procedure. Applied in waterborne UV-curable coatings for wood and flooring industry. Polyester. Biodegradable. Used in blends with PLA and fibers due to low thermo-mechanical properties. Obtained from fossil feedstock or bio-terefalic acid wood and flooring industry. Polyester industry. Polyester. The of 17°C option of PET and PBF of Polyester. Biodegradable. Used in blends with PLA and fibers due to low thermo-mechanical properties. Obtained from fossil feedstock or bio-terefalic acid of wood and flooring industry. Polyester industry. Polyester. The of 17°C option of PET and PBF overties a coording the percentage of unsaturated diacid (e.g. itaconic acid) and the curing procedure. Applied in waterborne UV-curable coatings for wood and flooring industry. Polyester		Moderately biodegradable. Good oxygen barrier and elongation properties. Used for film applications.
Polyturylene furanoate) - PBF Polytarylene adipate-0-14- butylene terephthalate) - PBAT Unsaturated polyester resins - UPR Polytarylene adipate-0-14- butylene terephthalate) - PBAT Unsaturated polyester resins - UPR Polytarylene adipate-0-14- polytarylene - PELA Polytarylene - PE-0- (trom bio-esthanol) Polytarylene - PP- Polytarylene -	Poly(ethylene furanoate) - PEF	🎀 Polyester. Durable, good oxygen barrier. T _m of 211 °C and T _g of 86 °C. Suitable for packaging, in the food and beverage industry.
Polyfi.4-butylene adipate-co-1.4 butylene terephthalate) - PBAT Unsaturated polyester risins - UPR Poly(L-lactide) - PLLA Polyester. PECA Polyester. PECA Polyester. PECA Polyester. PECA Polyester. December 2 Polyester. December 2 Polyester. December 2 Polyester. December 2 Polyester. December 3 Polyester. December 3 Polyester. December 3 Polyester. December 3 Polyester. Thermoplastic. Processable by extrusion, injection modding, blow molding. Degradable by hydroysis rather than microbial attack. Industrially compositable. Crystallinity can be controlled by co-polymerization of selected ratios of L- to D-stereoisomers of lactic acid or lactic acid or lactic de. Mechanical, thermal and barrier properties using the properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive material properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive material properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive material properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications range from water management to cable coating, food contact products and automotive properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications and management to cable coating, food contact products and automotive properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications, properties as compared to PA 6.6. PA 4.10 has low moisture uptable. Applications	Poly(trimethylene furanoate) - PTF	¥Polyester. Not biodegradable. T _m of 172 °C, T _g of 57 °C, good oxygen barrier properties. Employed in light weighting packaging.
Unsaturated polyester resins - Unsaturated polyester resins - Unsaturated polyester resins - UPR Unsaturated polyester resins - UPR Poly(L-lactide) - PLLA Poly(L-lactide) - PLLA Polyamides containing four carbons - 4C PAs: 4; 4.6 and 4.10 Polyamides containing four carbons - 4C PAs: 4; 4.6 and 4.10 Polyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyyonyl chloride - PVC* Polyethylene - PE* (from bio-ethanol) Polyamides containing four carbons - 4C PAs: 4; 4.6 and 4.10 Polyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyyonyl chloride - PVC* Polyethylene - PE* (from bio-ethanol) Polymethylene - PP* Polymethyl methacrylate) - PMMA* Poly(laction) - PPM Polymethyl methacrylate) - PMMA* Polymethyl methacrylate) - PMMA* Not biodegradable: Good resistance to hot water and polar solvents but poorly resistant to aromatic and alignia devices and bulletproof vests. Polyurethanes - PURs Polymethyl methacrylate) - PMMA* Not biodegradable: Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties or molecular sieve adsorbernts medical material used as ployed a polyour polymers. Action of the produced through the resonance of the polymers with shock absorption. Ozone and thermal resistant properties. Poly(lurfuryl alcohol) - PFA Acrylonitrie butadiene styrene - Acrylonitrie butadiene styrene - Age polymers and a sew properties as the fossil P. Not biodegradable in a material used as component for electrochemical and electrical devices (bod contacting applications) and in packaging. Poly(methyl alcohol) - PFA Acrylonitrie butadiene styrene - Age polymers and as a component for electrochemical and electrochemical and electrochemical and electrochemical and electrochemical and electrochemica	Poly(butylene furanoate) - PBF	Polyester. T _m of 172 °C, T _g of 44 °C. Potential replacer of PET and PBT.
wood and flooring industry. Poly(L-lactide) -PLLA Polygester. Thermoplastic. Processable by extrusion, injection molding, blow molding. Degradable by hydrolysis rather than microbial attack. Industrially compostable. Crystallinity can be controlled by co-polymerization of selected ratios of L- to D-stereoisomers of lactic acid or lactide. Mechanical, thermal and barrier properties justify applications in food packaging. Used for medical applications and drug delivery because of its biocompatibility. Not biodegradable. 4C PAs match properties of fossil-based PAs 6 and 6.6, such as thermal durability and mechanical strength, with a Tm above 250°C. All 4C PAs have higher dielectric strength and higher retention of tensile properties as compared to PA 6.6. PA 4.10 has low moisture uptake. Applications range from water management to cable coating, food contact products and automotive. Polyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyymid chloride – PVC* Polyethylene – PE* (from bio-ethanol) Polyethylene – PE* (from bio-ethanol) Polygerian – PP Polymethyl methacrylate) – PMMA* Polymethyl methacrylate) – PMMA* Polymethyl methacrylate) – PMMA* Not biodegradable and poorly chemically degradable. Same properties as tosil-based PVC. Used in construction profile applications, bottles and non-food packaging. When made more flexible by the addition of plasticiers, it is used in electrical cable insulation, imitation leather, flooring and as rubber replacer. Polyolefin. Same properties as the fossil PP. Not biodegradable, recyclable through dedicated infrastructures. Thermoplastic, High Density PE (more propylene delement on construction sector. Low Density Polyethylene is used in packaging. Ultrahigh Molecular Weight Polyethylene has applications in construction sector. Low Density Polyethylene is used in packaging. Ultrahigh Molecular Weight Polyethylene has applications in construction sector. Low Density Polyethylene is used in packaging. Ultrahigh Molecular Weight Polyethylene has polyetiations.	butylene terephthalate) - PBAT	
polyamides containing four carbons - 4C PAs: 4'. 46. and 4.10' Polyamides containing four carbons - 4C PAs: 4'. 46. and 4.10' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. PAS: 6.10': 10.10': 11 and 12' Polyamides with longer chains. Polyethylene – PE* (from bio-eithanol) Polygethylene – PE* (from bio-eithanol) Polygethylene – PP* Polyotelfin. Same properties of fossil-based PE. Not biodegradable, recyclable through dedicated infrastructures. Thermoplastic. High Density PE (more propylene dene monomer – EPDM (synthetic rubber) Polyurethylene propylene dene monomer – EPDM (synthetic rubber) Polyurethanes -PURs Polyurethanes -PURs Polyurethanes -PURs Acrylonitrile butadiene styrene - ABS Polycar(lis superabol-beath polymers) Poly(urfuryl alcohol) - PFA Acrylonitrile butadiene styrene - ABS Polycar(lis superabol-beath polymers) - Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Denylore properties and controlled by co-polymers and far and chains and the packaging is used a polymers with a polyol. Microbial degradation in fuel lines in cars, offshore pipelines, gas distribution properties. Not biodegradable and poorly chemically degradable. Same properties as fossil-based PVC. Used in construction profile applications, bottles and non-food provided in fuel tectrical cable insulation, initiation leather, flooring and as rubber replacer. Polyolefin. Same properties as the fossil PP. Not biodegradable, non-polar. Partially crystalline finds with polyolefin Same properties with food		wood and flooring industry.
Polyamides with longer chains. PAs: 6.10; 10.10; 11 and 12 Polyoring choined – PVC* Polyethylene – PE* (from bio-ethanol) Polymethylene – PP* Polypropylene - PP* Polypropylene - PP Polyurethylene - PPMAs Ethylene propylene diene monomer – EPDM (synthetic rubber) Polyurethanes - PURs Polyurethanes - PURs Poly(urfuryl alcohol) - PFA Acrylonitrile butadiene styrene - ABS Polyacrylic superabsorbent polymers - PA-SA Polyoric superabsorbent poly(flatconic acid) - PIA Poly(itaconic acid)	Poly(L-lactide) -PLLA	compostable. Crystallinity can be controlled by co-polymerization of selected ratios of L- to D-stereoisomers of lactic acid or lactide. Mechanical, thermal and
PAs: 6.10; 10.10; 11 and 12 Polyvinyl chloride – PVC* Polyethylene – PE* (from bio-ethanol) Polypropylene - PP* Poly(methyl methacrylate) – PMMA* Not biodegradable and poorly chemically degradable. Same properties as fossil-based PVC. Used in construction profile applications, bottles and non-food packaging. When made more flexible by the addition of plasticizers, it is used in electrical cable insulation, imitation leather, flooring and as rubber replacer. Polyethylene – PE* (from bio-ethanol) Polypropylene - PP* Poly(methyl methacrylate) – PMMA* Not biodegradable. To be the form of the splications and in packaging. Poly(methyl methacrylate) – PMMA* Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Polyurethanes - PURs Poly(turfuryl alcohol) - PFA Acrylonitrile butadiene styrene - ABS Polyacrylic superabsorbent polymers - PA-SA Poly(itaconic acid) - PIA Poly(i		4C PAs have higher dielectric strength and higher retention of tensile properties as compared to PA 6.6. PA 4.10 has low moisture uptake. Applications range from water management to cable coating, food contact products and automotive.
Polyethylene – PE* (from bio-ethanol) Polypropylene - PP* Polyopropylene diene monomer - EPDM (synthetic rubber) Polyurethanes - PURs Polyopropylene diene monomer - EPDM (synthetic rubber) Polyurethanes - PURs Polyopropylene - PPROpylene propylene diene monomer - EPDM (synthetic rubber) Polyurethanes - PURs Polyopropylene diene monomer - EPDM (synthetic rubber) Polyopropylene diene monomer - EPDM (synthetic rubber of the cherical clothing, elastomers with shock absorption. Ozone and thermal resistant. Electrical insulation properties. Used also for automotive applications. Produced through the reaction of a diisocyanate with a polyol. Microbial degradation depends on the chemical structure. Often blended with polyethers to increase flexibility or extensibility. Used as de-halogenated flame retardant foams, paints, powder coatings, medical devices (blood contacting applications). Not biodegradable. Synthesized from bio-based furfuryl alcohol (FA) deriving from sugars. Used in the fabrication of nanoporous carbons structures for molecular si		Telectronics, sports equipment, furniture and automobile components.
crystalline) finds applications in construction sector. Low Density Polyethylene is used in packaging. Ultrahigh Molecular Weight Polyethylene has applications in medical devices and bulletproof vests. Polygropylene - PP* Polyolefin. Same properties as the fossil PP. Not biodegradable, non-polar. Partially crystalline thermoplastic with low density. Used in a large variety of applications and in packaging. Polymethyl methacrylate)—PMMA* Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Not biodegradable. Good resistance to hot water and polar solvents but poorly resistant to aromatic and aliphatic hydrocarbons. Chlorine-free synthetic rubber used for technical clothing, elastomers with shock absorption. Ozone and thermal resistant. Electrical insulation properties. Used also for automotive applications. Prolycurethanes -PURs Polyurethanes -PURs Poly(furfuryl alcohol) - PFA Acrylonitrile butadiene styrene - ABS Polyacrylic superabsorbent polymers - PA-SA Polygricaconic acid) - PIA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Deloy(itaconic acid) - PIA Deloy(itaconic acid) - PIA Poly(itaconic acid	Polyvinyl chloride – PVC*	🗫 packaging. When made more flexible by the addition of plasticizers, it is used in electrical cable insulation, imitation leather, flooring and as rubber replacer.
Poly(methyl methacrylate)—PMMA* Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Ethylene propylene diene monomer – EPDM (synthetic rubber) Polyurethanes -PURs Polyurethanes -PURs Poly(furfuryl alcohol) - PFA Acrylonitrile butadiene styrene- ABS Polyacrylic superabsorbent polymers - PA-SA Poly(furfuryl suconic acid) - PIA Poly(futaconic acid) - PIA Applications and in packaging. And biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties. Not biodegradable. Good resistance to hot water and polar solvents but poorly resistant to aromatic and aliphatic hydrocarbons. Chlorine-free synthetic rubber used for automotive applications. Produced through the reaction of a diisocyanate with a polyol. Microbial degradation depends on the chemical structure. Often blended with polyethers to increase flexibility or extensibility. Used as de-halogenated flame retardant foams, paints, powder coatings, medical devices (blood contacting applications). Not biodegradable. Synthesized from bio-based furfuryl alcohol (FA) deriving from sugars. Used in the fabrication of nanoporous carbons structures for molecular sieve adsorbents, membranes and as a component for electrochemical and electronic devices. Obtained from butadiene rubber dispersed in a matrix of styrene-acrylonitrile copolymer. Not biodegradable. Thermoplastic, used to make light, rigid, moulded solublisation. Find applications in personal disposable hygiene products, such dispers and sanitary napkins. Used as de-halogenated flame retardant foa		👱 crystalline) finds applications in construction sector. Low Density Polyethylene is used in packaging. Ultrahigh Molecular Weight Polyethylene has applications
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monomer – EPDM (synthetic rubber) Polyurethanes -PURs Polyurethanes -PURs Poly(furfuryl alcohol) - PFA Acrylonitrile butadiene styrene ABS Polyacrylic superabsorbent polymers - PA-SA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Wased for technical clothing, elastomers with shock absorption. Ozone and thermal resistant. Electrical insulation properties. Used also for automotive applications. Ozone and thermal resistant. Electrical insulation properties. Used also for automotive applications. Produced through the reaction of a diisocyanate with a polyol. Microbial degradation depends on the chemical structure. Often blended with polyethers to increase flexibility or extensibility. Used as de-halogenated flame retardant foams, paints, powder coatings, medical devices (blood contacting applications). Biodegradable polyurethane scaffolds have been used in tissue regeneration. Not biodegradable. Synthesized from bio-based furfuryl alcohol (FA) deriving from sugars. Used in the fabrication of nanoporous carbons structures for molecular sieve adsorbents, membranes and as a component for electrochemical and electronic devices. Obtained from butadiene rubber dispersed in a matrix of styrene-acrylonitrile copolymer. Not biodegradable. Thermoplastic, used to make light, rigid, moulded products such as pipes, automotive parts. Used also for its flame retardant properties. Its high swelling capacity is tuneable by controlling the degree of crosslinking. Its biodegradation in soil can be improved under conditions that maximize solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins. Due to the presence of a vinyl moiety, itaconic acid is structurally similar to acrylic and methacrylic acid, providing a suitable bio-based alternative to poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-	Poly(methyl methacrylate)-PMMA*	🕌 Not biodegradable. Lightweight material used as glass replacement in automotive for shatterproof and UV resistant properties.
Poly(furfuryl alcohol) - PFA Acrylonitrile butadiene styrene ABS Polyacrylic superabsorbent polymers - PA-SA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Increase flexibility or extensibility. Used as de-halogenated flame retardant foams, paints, powder coatings, medical devices (blood contacting applications). Biodegradable polyurethane scaffolds have been used in tissue regeneration. Not biodegradable. Synthesized from bio-based furfuryl alcohol (FA) deriving from sugars. Used in the fabrication of nanoporous carbons structures for molecular sieve adsorbents, membranes and as a component for electrochemical and electronic devices. Obtained from butadiene rubber dispersed in a matrix of styrene-acrylonitrile copolymer. Not biodegradable. Thermoplastic, used to make light, rigid, moulded products such as pipes, automotive parts. Used also for its flame retardant properties. Its high swelling capacity is tuneable by controlling the degree of crosslinking. Its biodegradation in soil can be improved under conditions that maximize solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins. Due to the presence of a vinyl moiety, itaconic acid is structurally similar to acrylic and methacrylic acid, providing a suitable bio-based alternative to poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-	monomer – EPDM (synthetic	🔆 used for technical clothing, elastomers with shock absorption. Ozone and thermal resistant. Electrical insulation properties. Used also for automotive
Acrylonitrile butadiene styrene - ABS Polyacrylic superabsorbent polymers - PA-SA Poly(itaconic acid) - PIA Molecular sieve adsorbents, membranes and as a component for electrochemical and electronic devices. Obtained from butadiene rubber dispersed in a matrix of styrene-acrylonitrile copolymer. Not biodegradable. Thermoplastic, used to make light, rigid, moulded products such as pipes, automotive parts. Used also for its flame retardant properties. Its high swelling capacity is tuneable by controlling the degree of crosslinking. Its biodegradation in soil can be improved under conditions that maximize solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins. Due to the presence of a vinyl moiety, itaconic acid is structurally similar to acrylic and methacrylic acid, providing a suitable bio-based alternative to poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-	Polyurethanes -PURs	increase flexibility or extensibility. Used as de-halogenated flame retardant foams, paints, powder coatings, medical devices (blood contacting applications).
ABS Polyacrylic superabsorbent polymers - PA-SA Poly(itaconic acid) - PIA Poly(itaconic acid) - PIA Polyacrylic such as pipes, automotive parts. Used also for its flame retardant properties. Its high swelling capacity is tuneable by controlling the degree of crosslinking. Its biodegradation in soil can be improved under conditions that maximize solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins. Due to the presence of a vinyl moiety, itaconic acid is structurally similar to acrylic and methacrylic acid, providing a suitable bio-based alternative to poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-		Not biodegradable. Synthesized from bio-based furfuryl alcohol (FA) deriving from sugars. Used in the fabrication of nanoporous carbons structures for molecular sieve adsorbents, membranes and as a component for electrochemical and electronic devices.
polymers - PA-SA solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins. Due to the presence of a vinyl moiety, itaconic acid is structurally similar to acrylic and methacrylic acid, providing a suitable bio-based alternative to poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-	ABS	🔭 products such as pipes, automotive parts. Used also for its flame retardant properties.
Poly(itaconic acid) - PIA poly(meth)acrylates via radical polymerization to yield poly(itaconic acid) (PIA). Applications include fibers, coatings, adhesives, thickeners, binders. As co-		solubilisation. Find applications in personal disposable hygiene products, such diapers and sanitary napkins.
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Building blocks and monomers	Structural evolution	Targeted performance	_
Aromatic lignin derivatives	2,4-, 2,5-, and 2,6-pyridinedicarboxylic acid obtained by re-routing the lignin degradation pathways of <i>Rhodococcus jostii</i> RHA1	New bio-based aromatic / apliphatic polyesters obtainable via enzymatic polycondensation with Mn around 14000 Da	
Ricinoleic acid	Confers biocidal activity to poly(hexamethylene succinate) modified at the chain ends. Imidazolium salt was anchored on C=C bond of ricinoleic acid to improve biocidal activity.	•	
Terpenes	Pinene transformed into pinocarvone, which contains a reactive exomethylene group exploitable for radical polymerization	High molecular weight plytherpenes with excellent thermal properties (Tg > 160 C). Polymerization of pinene would require low temperatures (-70°C) unviable for industrial purposes.)
	Branched chains of polyamide 4.	Moderation of rigidity. Increased MW. Improved mechanical properties without decreasing $T_{\rm m}$.	
Amides	$\{(4,40\text{-diyl}-\alpha\text{-truxillic} \text{ acid dimethyl ester}) 4,40\text{-diacetamido}-\alpha\text{-truxillamide}\},$ obtained from bio-based 4-aminophenylalanine, UV coupled with cinnamic acid	9	
Isosorbide	Confers rigidity	Increasing thermos and mechanical properties while preserving the biodegradability.	
Modified lactides for improved PLAs	Phenyl-substituted lactide synthesized by cyclic dimerization of bio-based mandelic acid to obtain mandelide (meso stereoisomer), which is polymerized <i>via</i> ring opening polymerization (ROP)	bulky side chains. $^{\circ}$ Polymandelide has T_{o} > 100°C and is less biodegradable than PLLA.	
violance lectices for improved 1 LAS	Norbornene-substituted lactide obtained by brominating the bio-based lactide. Elimination and Diels Alder reactions yield the norbornene lactide used in ring-opening metathesis polymerization.	Polymers have T _g > 190°C and narrow polydispersity.	_
Cyclic diols	Bio-based 1,4-cyclohexanedimethanol (CHDM) is obtainable from renewable terephthalic acid.	As co-monomer in polyesters of 2,5-furandicarboxylic acid-increases rigidity, confers mechanical properties comparable to PET and improves barrier properties. Its polycondensation requires temperature around 240-280 °C due to the high boiling point but such temperatures promote its decomposition. Mild enzymatic polycondensation overcomes this drawback.	r
Phenols	4-hydroxycinnamic acid (4HCA)	The aromatic ring confers liquid crystalline properties to polyesters. The biobased liquid crystal polymers exhibits remarkable properties (strength = 63 MPa, Young's modulus = 16 GPa, maximum softening temperature = 169 °C	3
Succinic acid derivatives	Polyesters obtained by co-polymerization of succinic acid with furan dicarboxylic acid (FDCA)	Modifying soft properties of linear poly(succinates)s by introducing aromatic furan moieties. The corresponding polyesters poly(butylene succinate-co-butylene furandicarboxylate)s (PBSF) have Mw from 39 000 to 89 000 g/mol and display excellent thermal stability. Their structure and properties can be tuned ranging from crystalline polymers with good tensile modulus (360-1800 MPa) and strength (20–35 MPa) to nearly amorphous polymer of low Tg and high elongation (~600%), so that they may find applications in thermoplastics as well as elastomers or impact modifiers.	-
Furan derivatives	Nucleophilic aromatic substitution polymerization of 2,5- bis(4- fluorobenzoyl)furan (BFBF) derived from FDCA and potassium salts of aromatic bisphenols	Bio-based poly(thioether ketone) (PEEK) with Tm >300 $^{\circ}$ C, comparable to fossil-based PEEK	
Itaconic acid derivatives	Functionalization of the unsaturated double bond of dimethylitaconate by thia-Michael addition reaction using 1-octanethiol.	Improve the stability of itaconic derivative monomers toward common conditions of polycondensation (high temperatures and metal-based catalysts)	1
	Post-polymerization modification of vinyl group of poly(itaconate) <i>via</i> Michael addition of primary amines.	Amine-triggered degradable materials; oligoesters displaying amine functionalities for biomolecules anchoring or covalent crosslinking.	
	Michael additions of proline, cysteine and other S-containing nucleophiles to vinyl moiety of poly(itaconate)s. Michael addition of C-nucleophiles (acetylacetone and dimethyl malonate) to	- Addition of pendants to polyester chain. Modifying polymer properties.	

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