

POLÍMEROS

Julio Alberto Aguilar Schafer

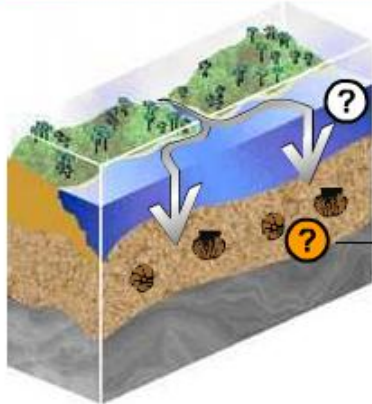
Formación geológica del petróleo

Formación del Petróleo



HOW ARE OIL AND GAS FORMED?

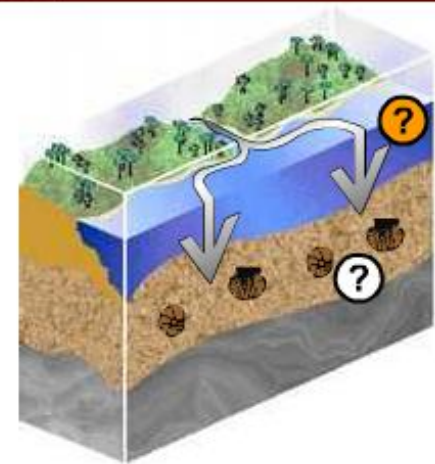
Oil and gas (hydrocarbons) are found in sedimentary basins.



Sedimentary rocks with organic material are source rocks. The "source" of hydrocarbons, they can be coal or shale.

HOW ARE OIL AND GAS FORMED?

and gas (hydrocarbons) found in sedimentary basins.



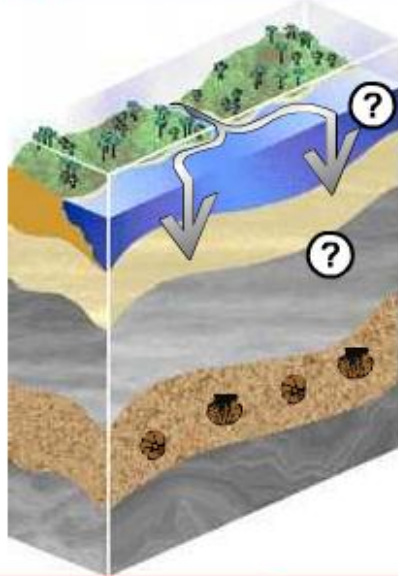
The type of organic material determines whether oil or gas is formed



HOW ARE OIL AND GAS FORMED?

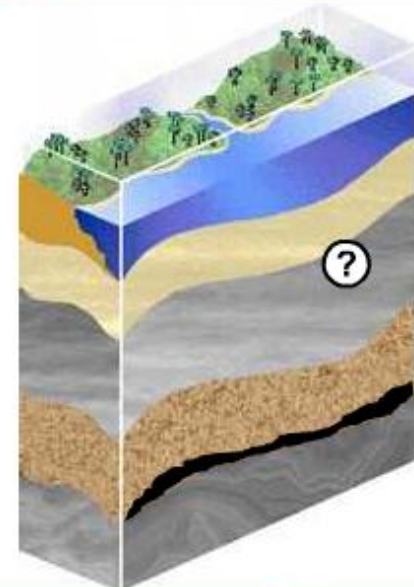
Oil and gas (hydrocarbons) are found in sedimentary basins.

Millions of years ago dead plants and microscopic animals (organic material) were deposited in sedimentary basins with mud, sand and other sediments. Over time the sediments were compacted and transformed into layers of sedimentary rock.



HOW ARE OIL AND GAS FORMED?

Oil and gas are formed by the thermal conversion of organic matter trapped in source rocks. This happens when source rocks are buried deeper and the temperature rises and pressure increases.

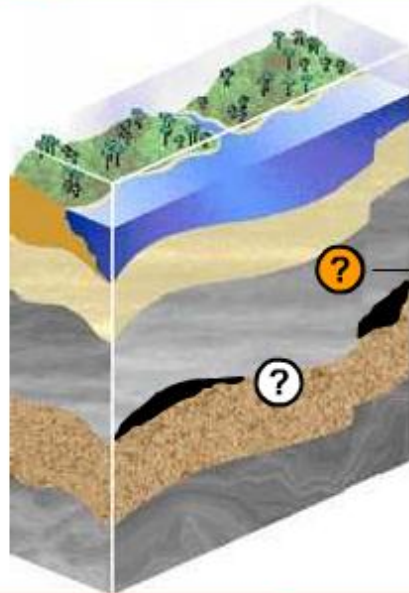




HOW ARE OIL AND GAS FORMED?

To form an oil or gas field, oil and gas must leave the source rocks and move through the pores and spaces of permeable rock or along minor fractures and fault lines.

If oil and gas are not trapped in a reservoir formation they will escape to the earth's surface as a seepage.

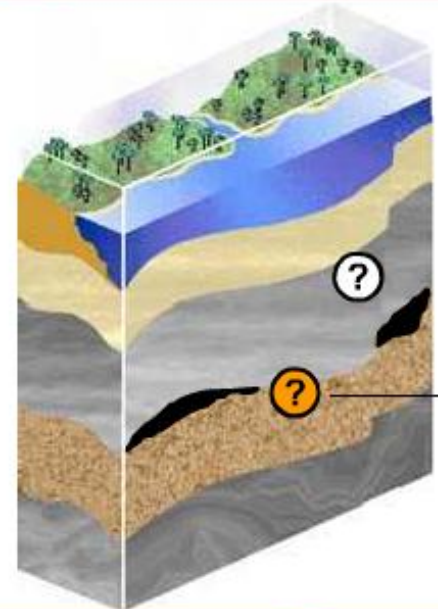


A fault traps oil & gas because it moves the rock layers to form a seal.

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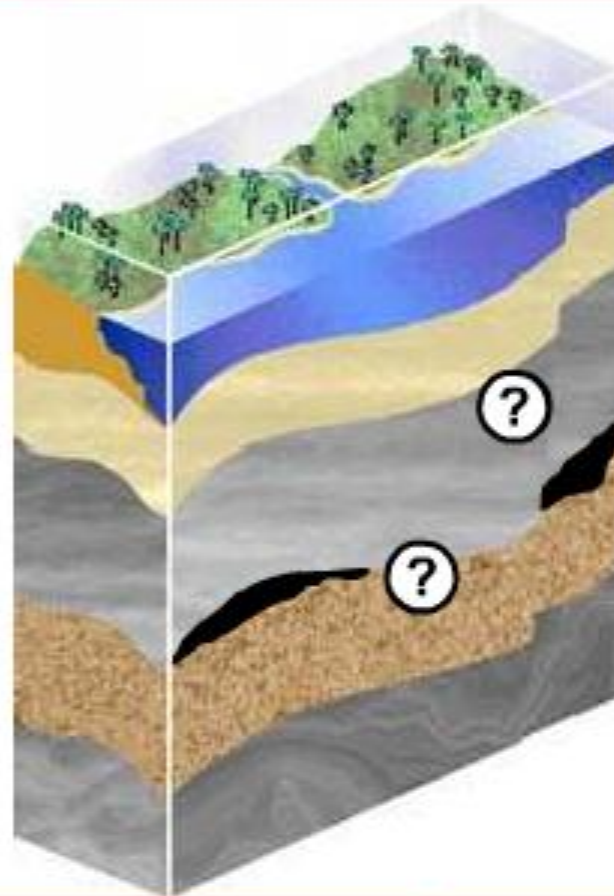
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HOW ARE OIL AND GAS FORMED?

Oil or gas fields are found in subsurface geological features called traps. For a trap to be effective it needs reservoir rock to hold the oil and gas and seal rocks to contain it.

Oil and gas are not a liquid lake but are contained in the reservoir rocks' pores.



Localización de Petróleo

Magnetometría aérea

AIRBORNE MEGATEM-GEOTEM TEM
by GEOTERREX-DIGHEM



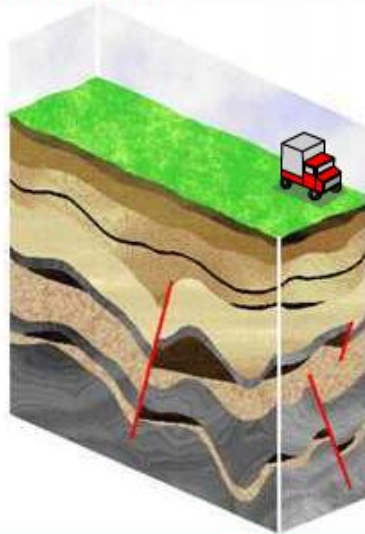
Localización de Petróleo



HOW ARE OIL AND GAS DEPOSITS LOCATED?

Scientists use geology and geophysics to "see" what is happening beneath the earth.

Seismic surveying is important. It can be performed on land and sea using different equipment. There are always three components:

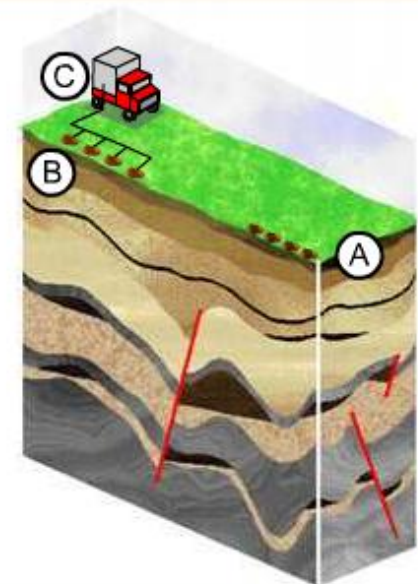


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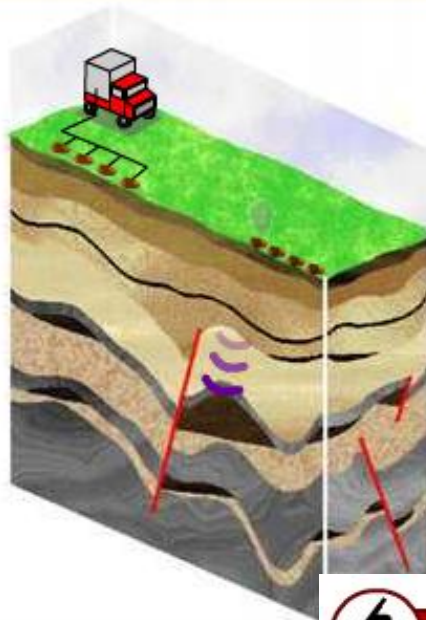
- A. a seismic source
- B. sensors
- C. recording equipment





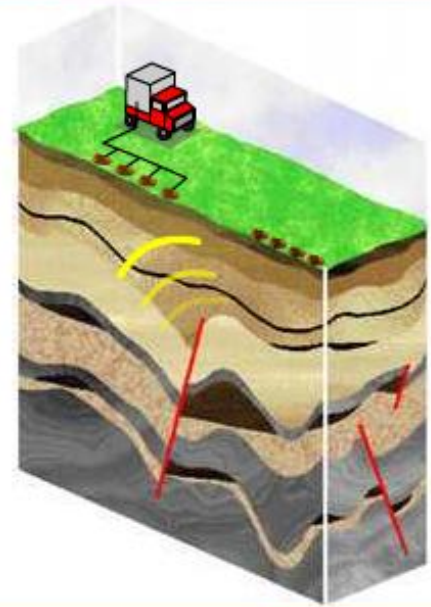
HOW ARE OIL AND GAS DEPOSITS LOCATED?

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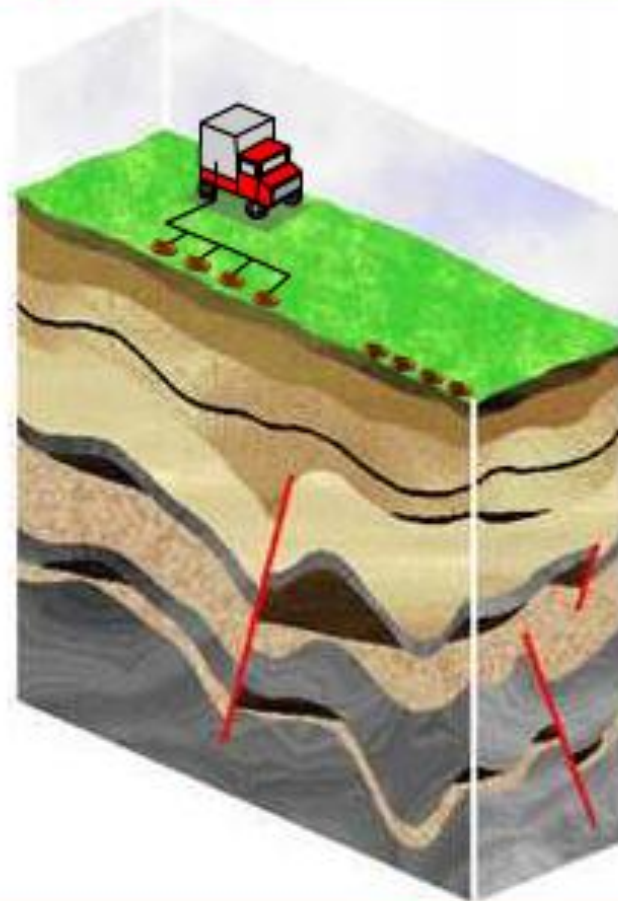


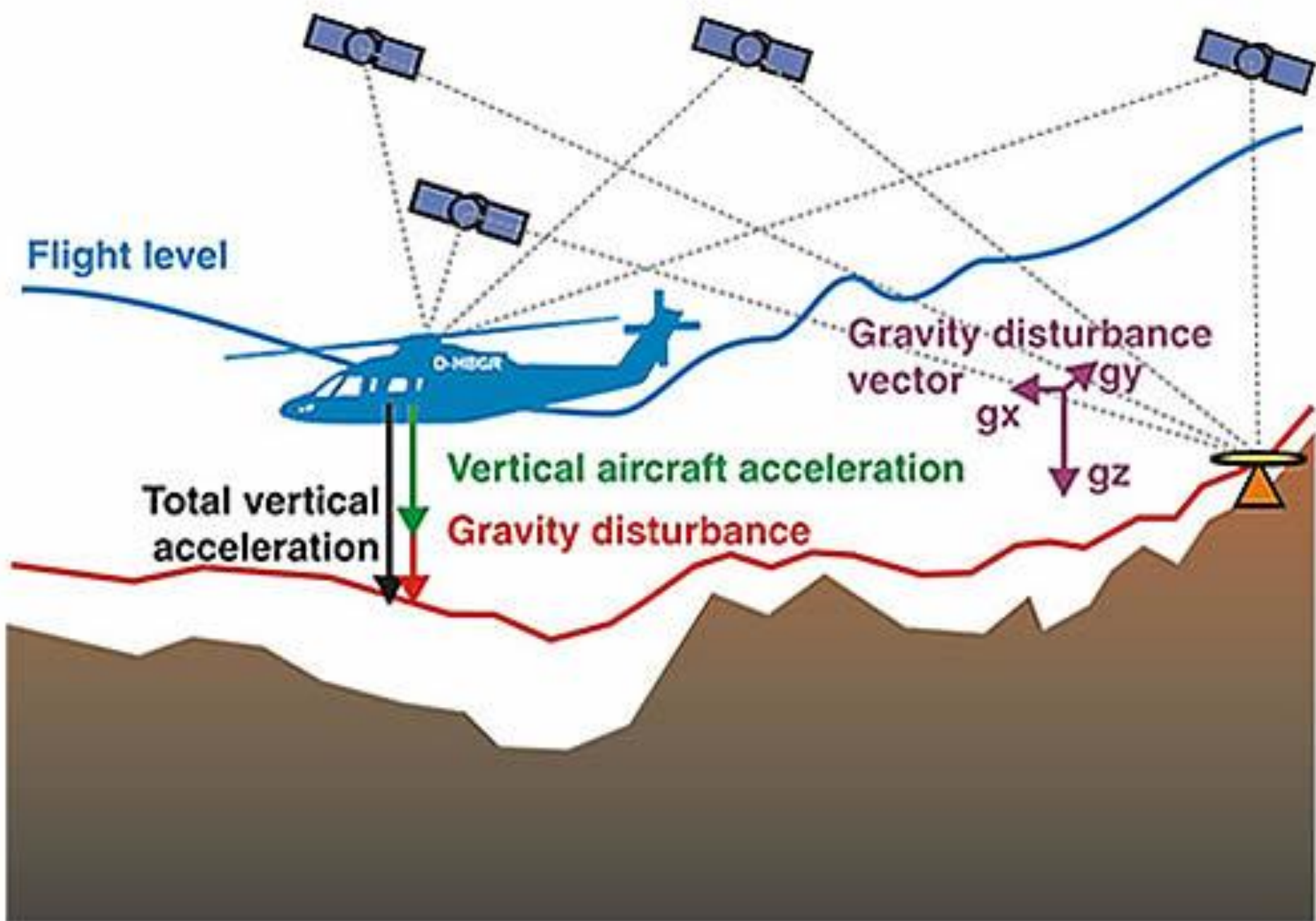


HOW ARE OIL AND GAS DEPOSITS LOCATED?

Scientists create a "picture" of the structure of the rock layers by measuring the time it takes energy waves to reach the surface.

Once the seismic data is collected it must be processed and carefully interpreted to decide whether further testing is needed or if exploration can begin.







Camión Vibrador



Perforadora
de mesa
giratoria y
brocas

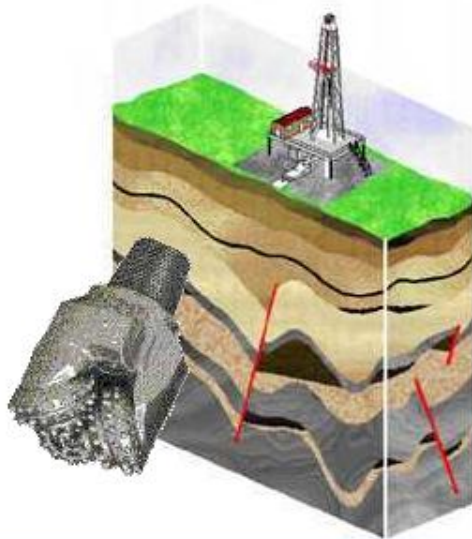
Cómo es descubierto



HOW ARE OIL AND GAS DISCOVERED?

The most common drilling technique is the rotary drilling system - a highly efficient mechanical system used on land and sea.

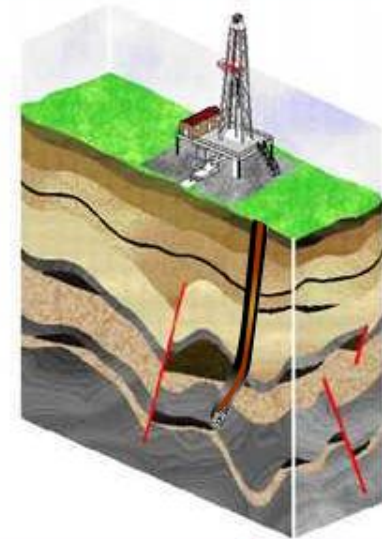
When seismic surveys and other data indicate that oil and/or gas could be present an exploration or 'wildcat' well will be drilled.



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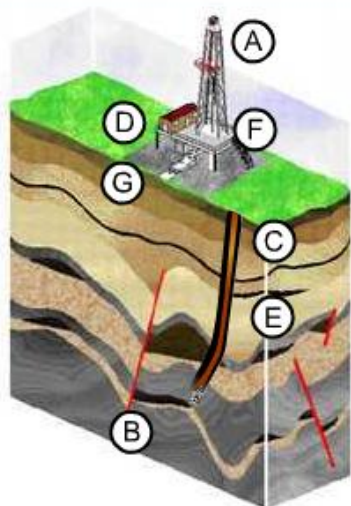




HOW ARE OIL AND GAS DISCOVERED?

A rotary drilling rig is made up of :

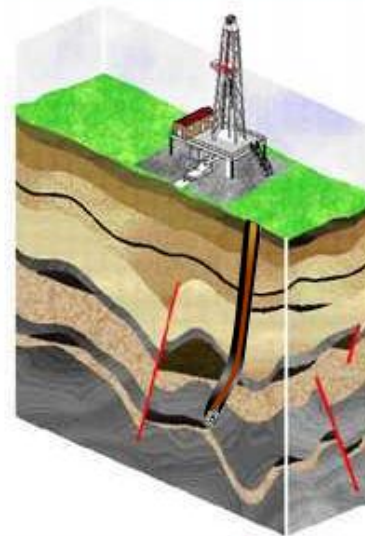
- A. Hoisting equipment
- B. Drilling bit
- C. Drill pipe
- D. Rotary equipment
- E. Mud circulating treating equipment
- F. Blowout prevention system
- G. Power source



HOW ARE OIL AND GAS DISCOVERED?

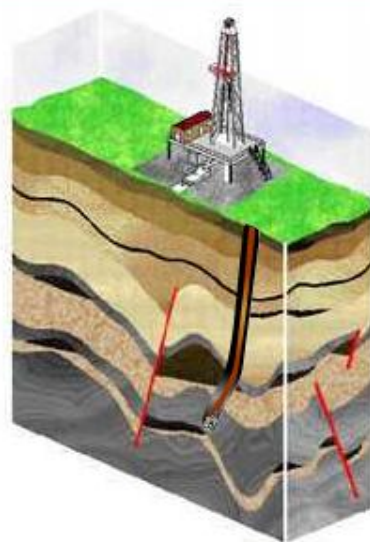
Drilling is a round-the-clock operation. A drilling supervisor leads a team of experts who work in shifts.

Wells are usually drilled vertically but can be drilled at an angle. This technique, called directional drilling, is used for a variety of reasons. A major advancement is horizontal drilling which drills along a reservoir and increases production.

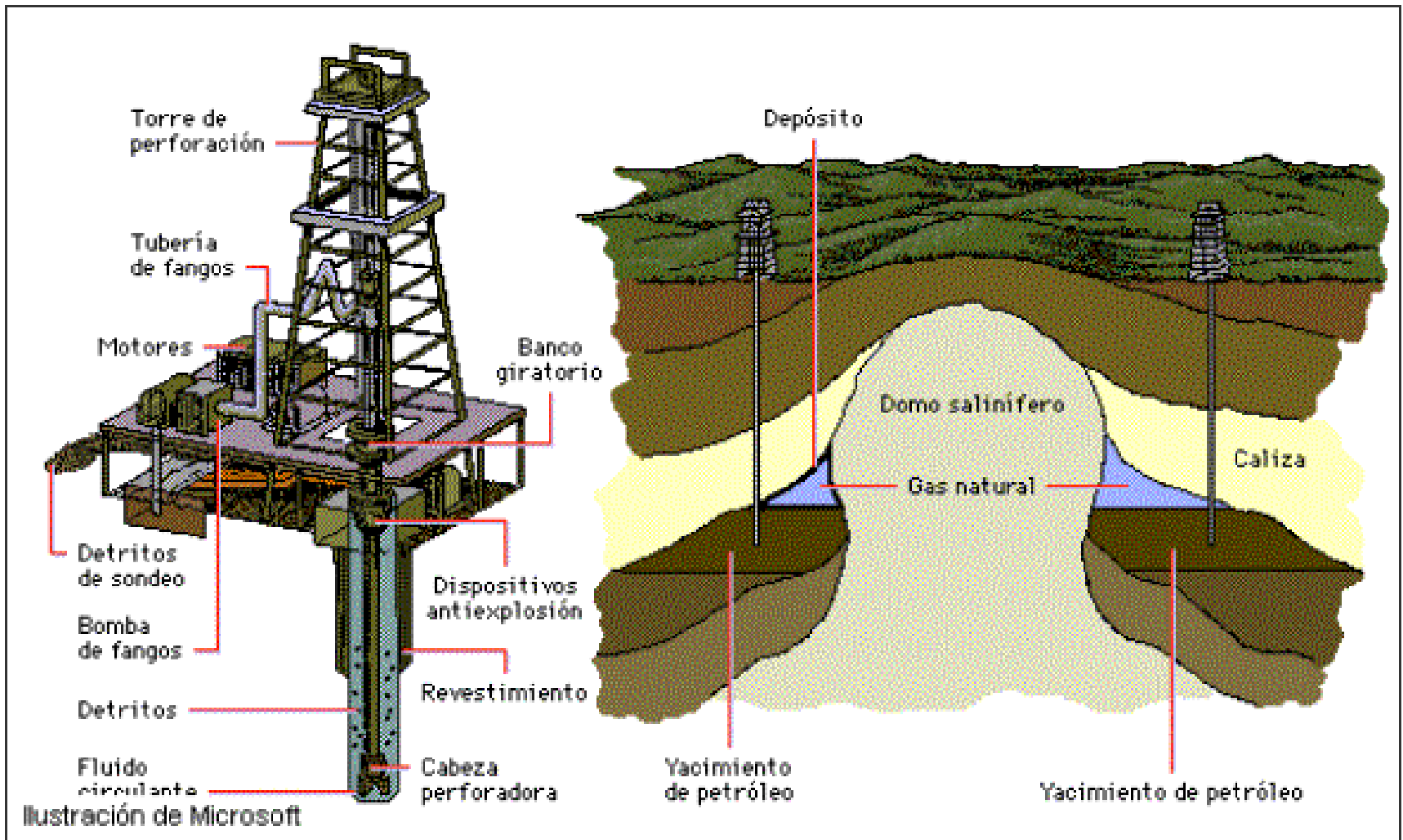


HOW ARE OIL AND GAS DISCOVERED?

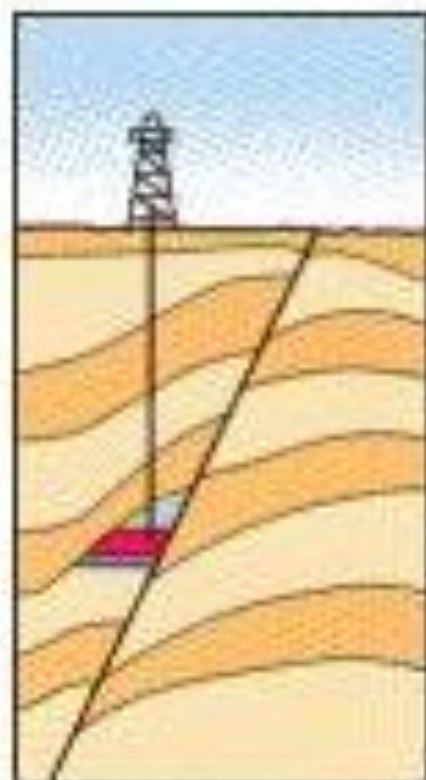
As the drill nears its final depth, engineers, geologists and management have to decide if there is enough oil and gas for the well to be completed or if it should be abandoned as a 'dry hole.'



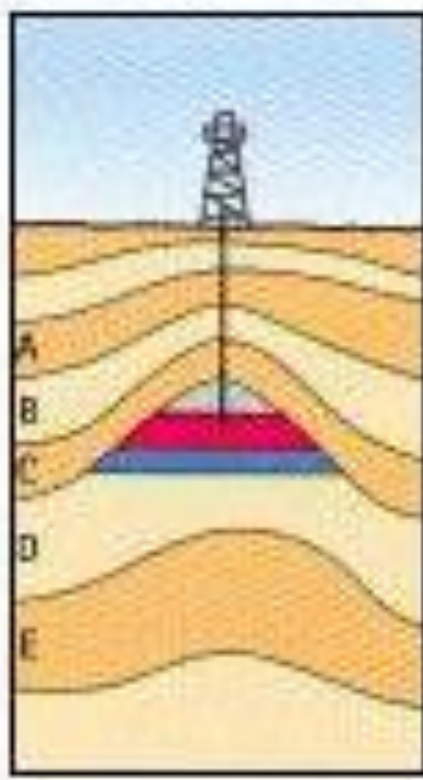
Extracción del petróleo



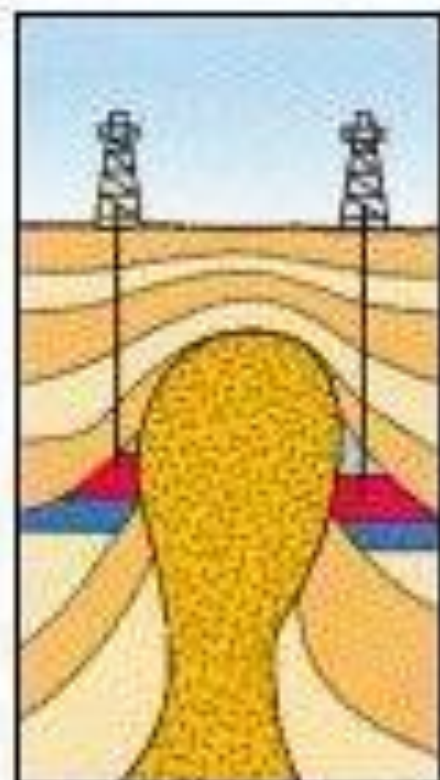
TRAMPAS MAS COMUNES



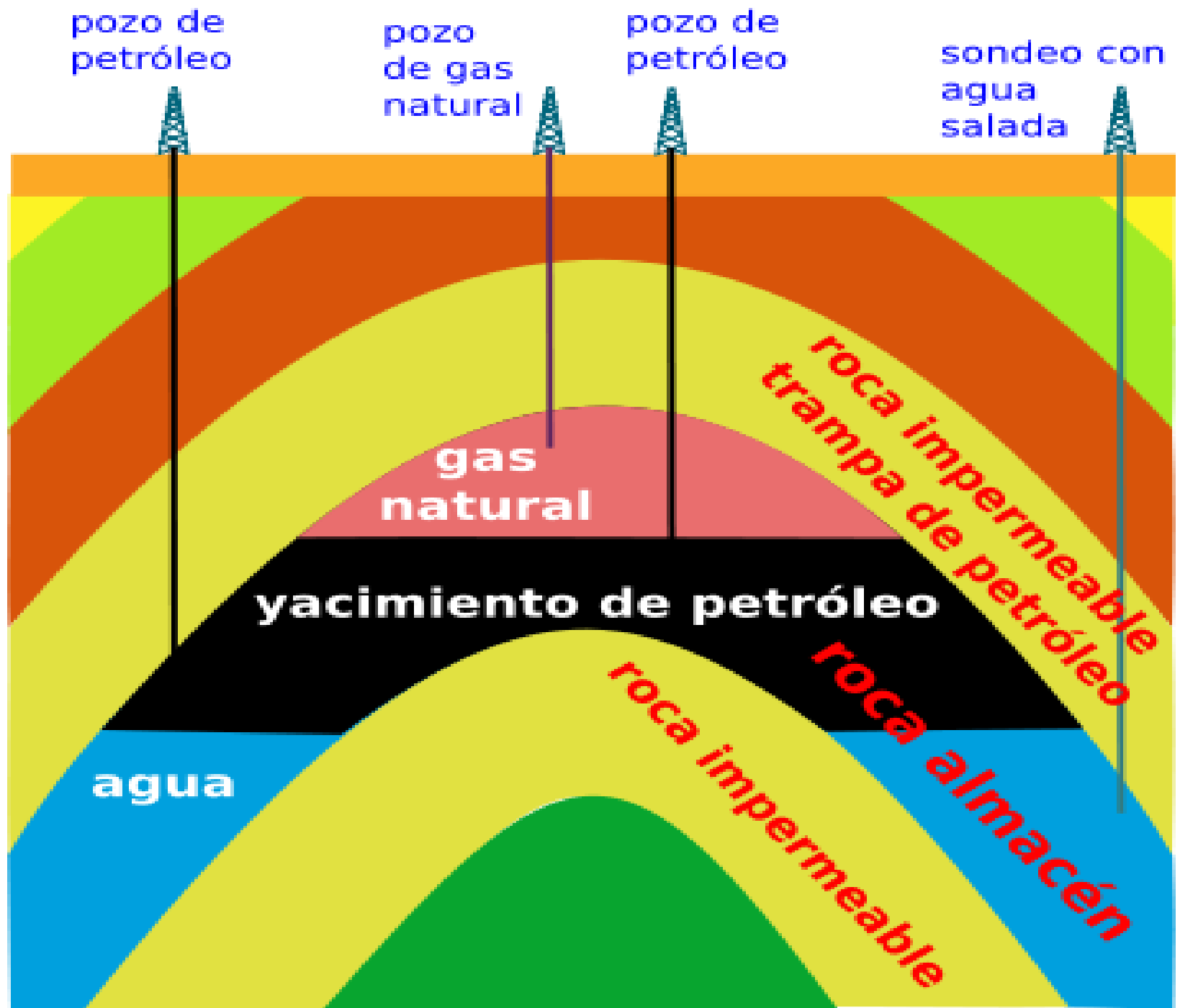
**TRAMPA
POR
FALLA**



**TRAMPA
POR
PLEGAMIENTO**



**TRAMPA
COMBINADA
(PLEGAMIENTO,
FALLA Y DOMO)**



TRAMPAS DEL PETROLEO



■	MARGA
■	CONGLOMERADO
■	ARENISCA
■	ARCILLA

■	CAUZA
■	DOMO DE SAL
■	FALLA
■	LENTES DE ARENA

■	ROCA ALTERADA
■	PUEGUES
■	PETROLEO - GAS

TRAMPAS DE PETRÓLEO

reservorio de
variación litográfica

reservorio de falla

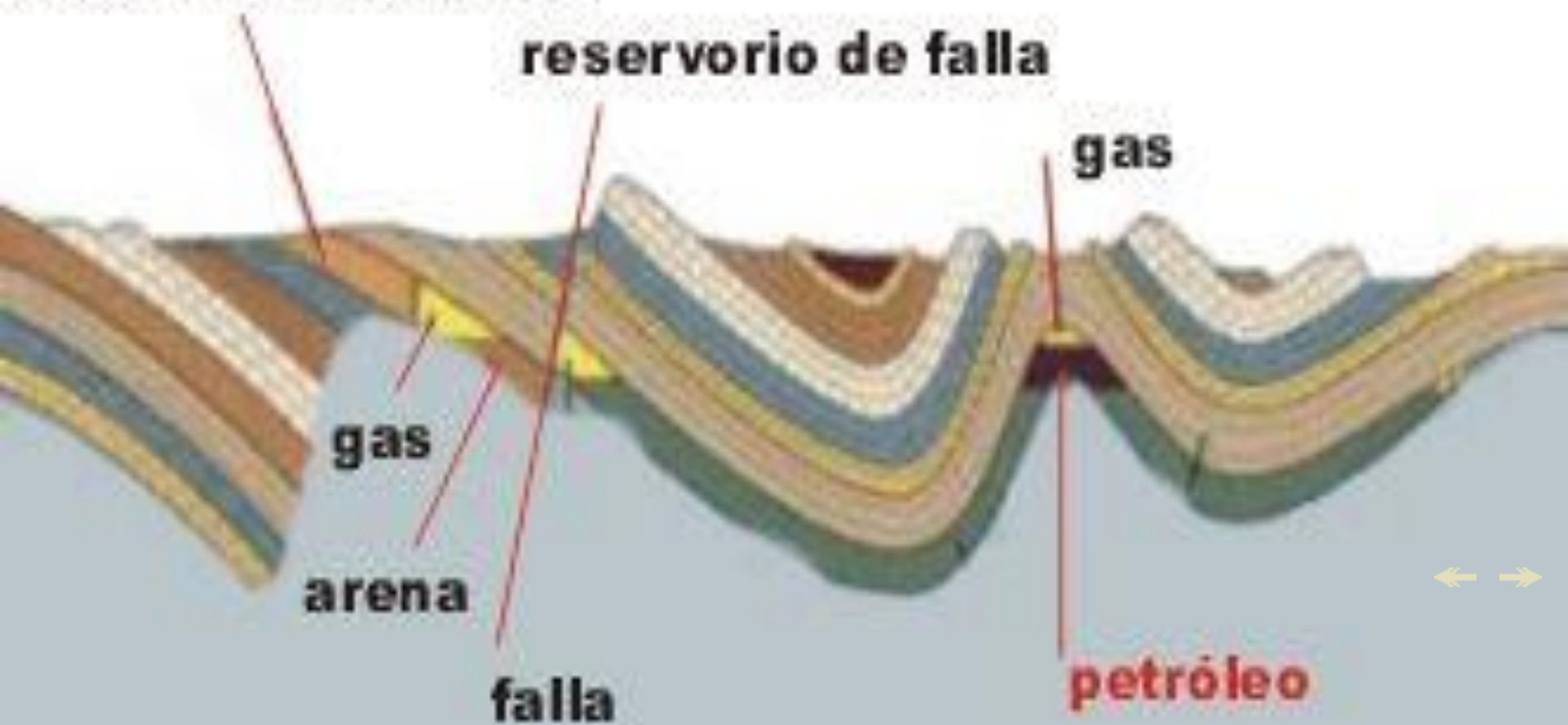
gas

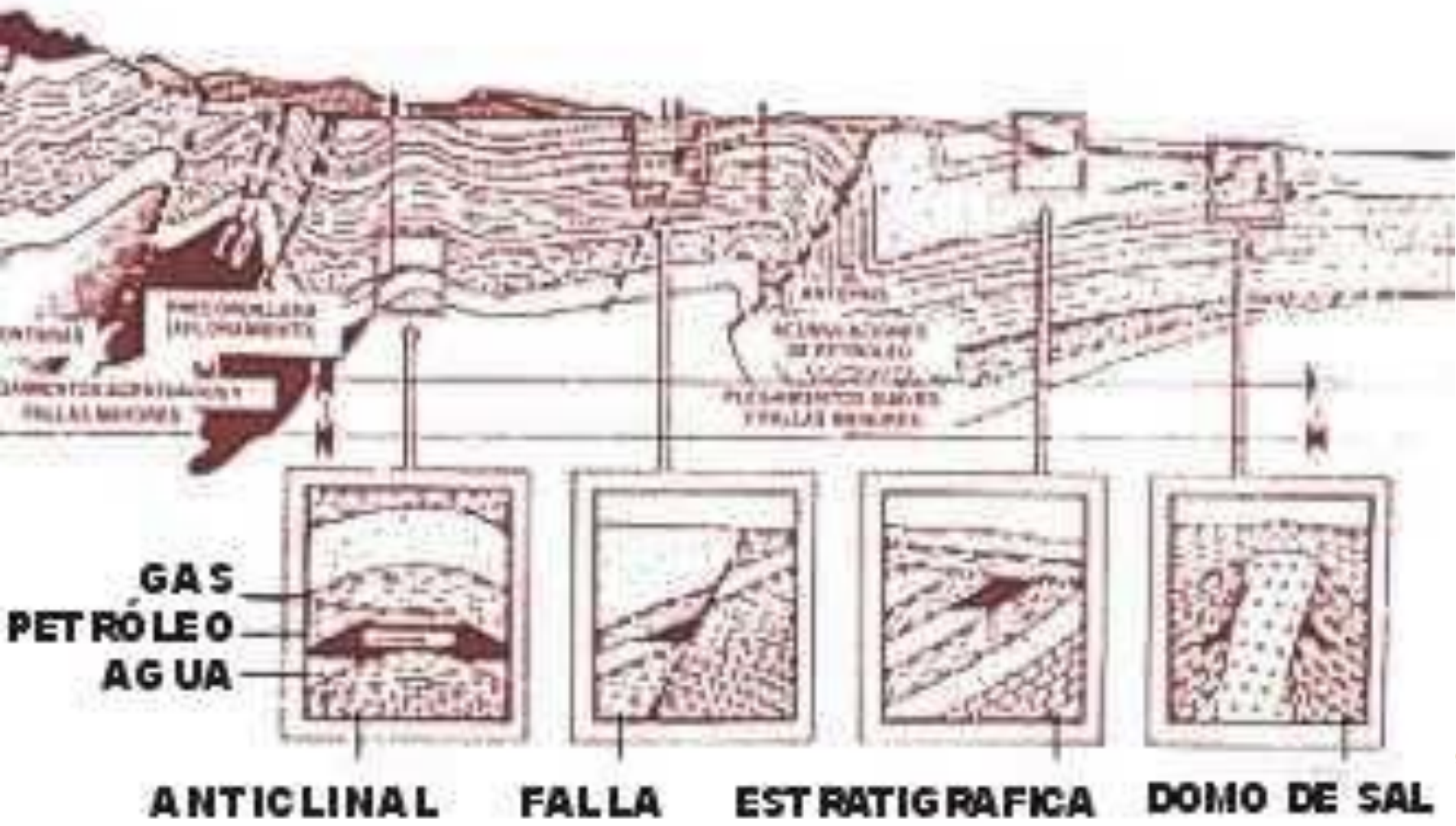
gas

arena

falla

petróleo





Perforación en plataforma submarina

Perforación en tierra



gas

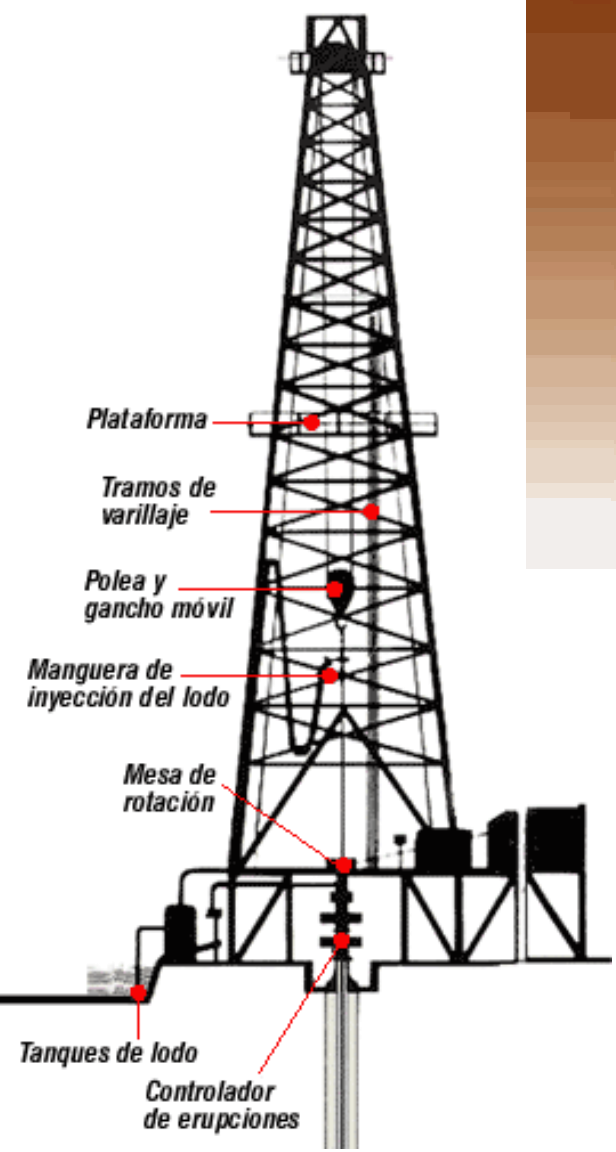
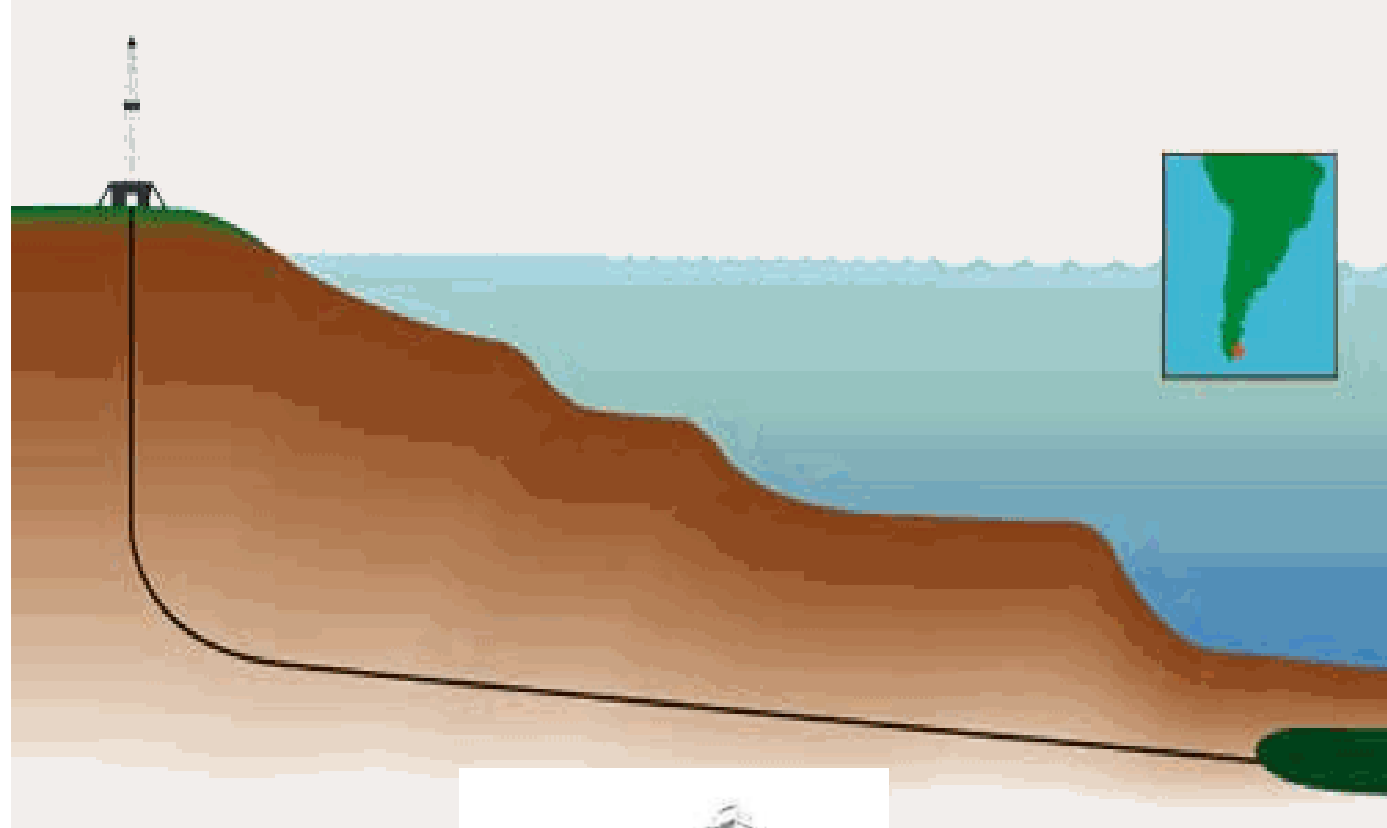
mezcla de hidrocarburos

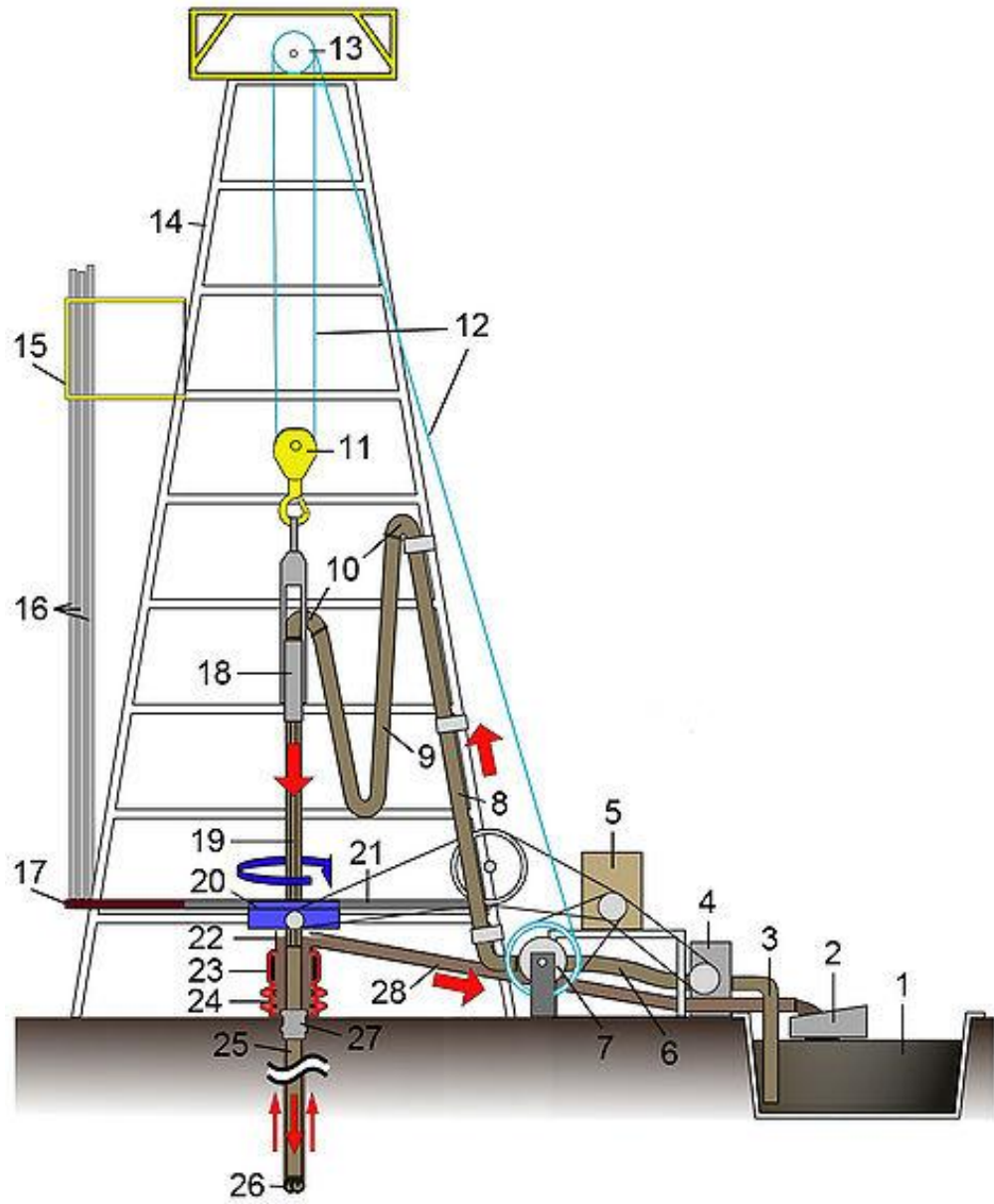
agua

gas

mezcla de hidrocarburos

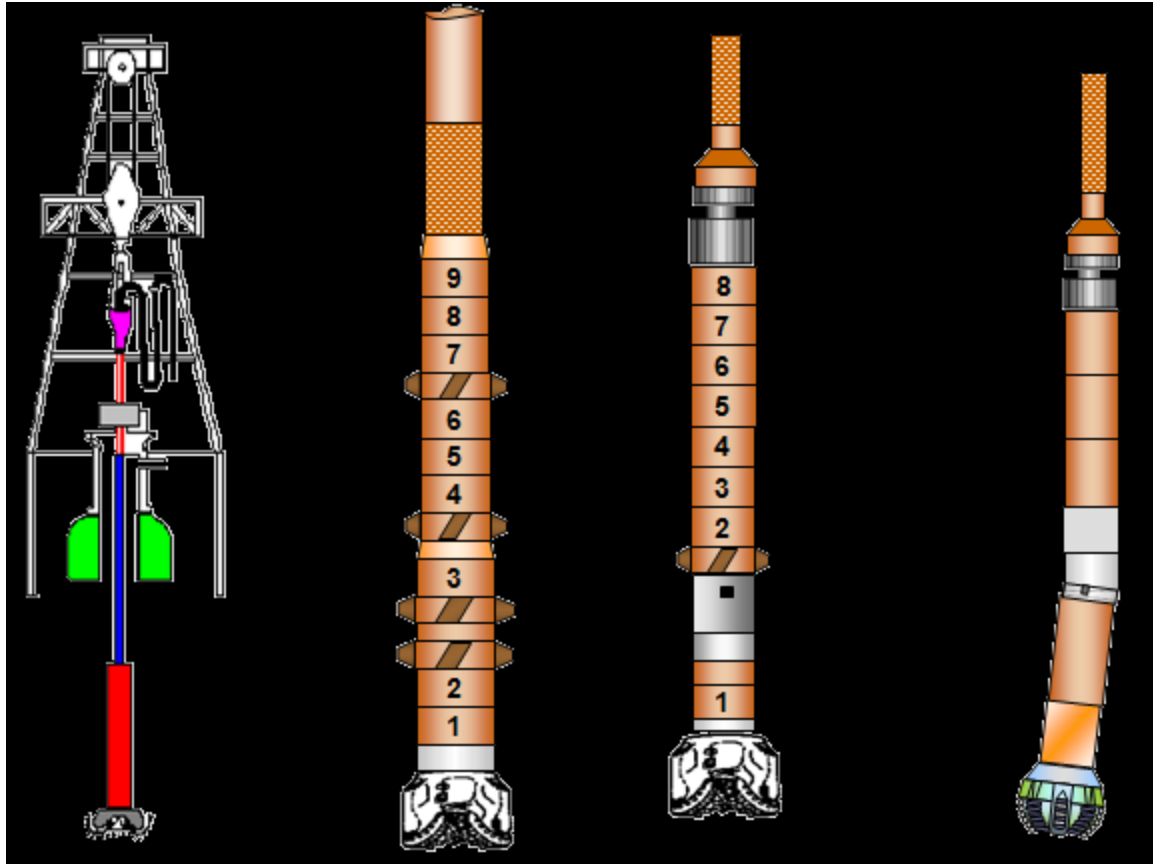
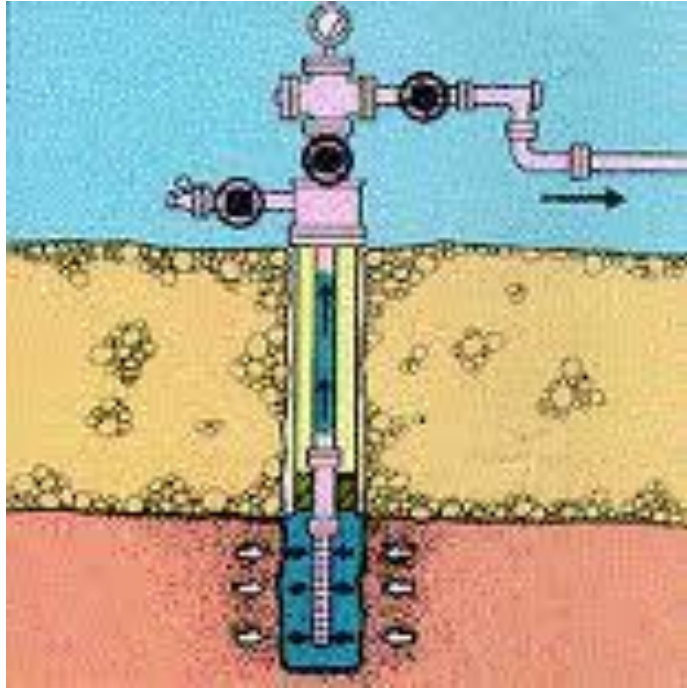
agua



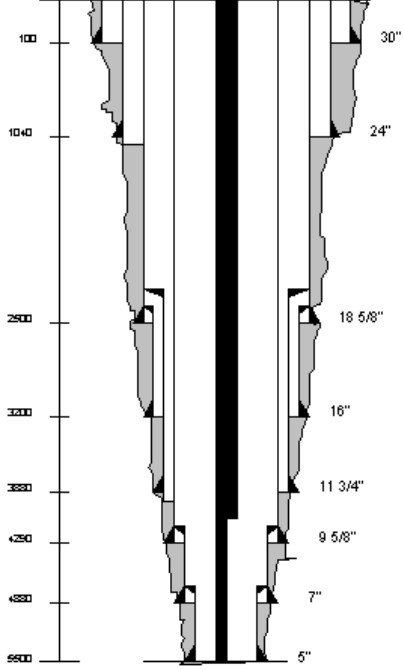


Identificación de la Fig. (Equipos usados en la Perforación de Pozos)

1. Tanque de lodo
2. Agitador de arcilla
3. Succión de lodo.
4. Bomba de lodo
5. Motor
6. Manguera de bomba
7. carrete de aparejo
8. Cañería de lodo
9. Manguerote
10. Cuello de ganso
11. Aparejo
12. Cable de aparejo
13. Bloque corona
14. Estructura
15. Piso del enganchador
16. Tiros
17. Rack
18. Conexión giratoria de lodo
19. Barra de perforación
20. mesa rotativa
21. Piso de perforación
22. Bell niple
23. Valvula BOP anular
24. Valvula BOP ciega de cañeria
25. Sarta de perforación
26. Trépano
27. Cabeza del casing
28. Línea de retorno del lodo.



Pozo Programado



Para mejorar el porcentaje de extracción (de 25% a 33% promedio) se utilizan dos procesos:

- Inyección de Agua:

Se bombea agua hacia el yacimiento para mantener o aumentar la presión. También aumenta el ritmo de producción.

- Inyección de Vapor:

Se emplea en depósitos de petróleo muy viscoso. Reduce la viscosidad.

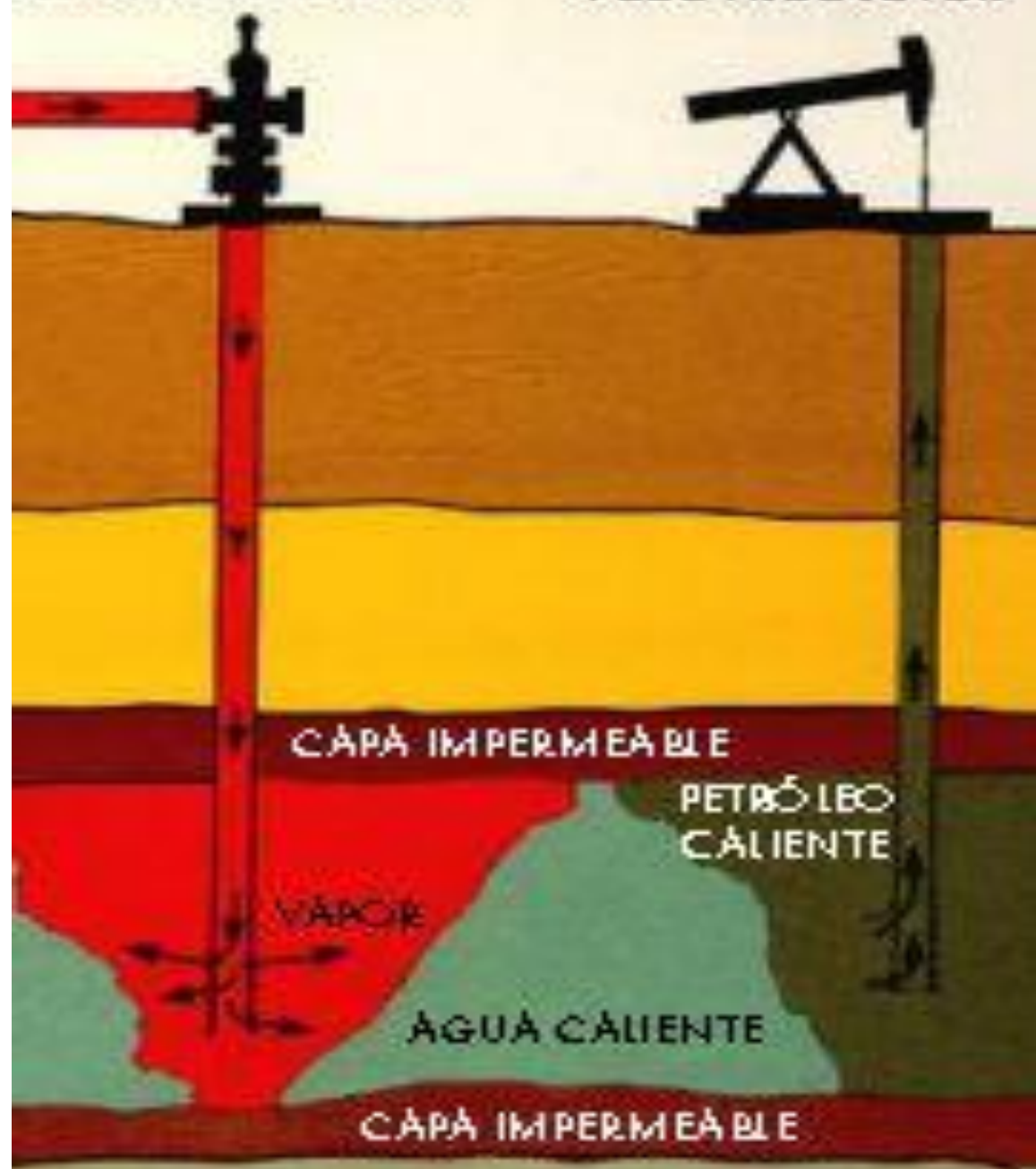


AP



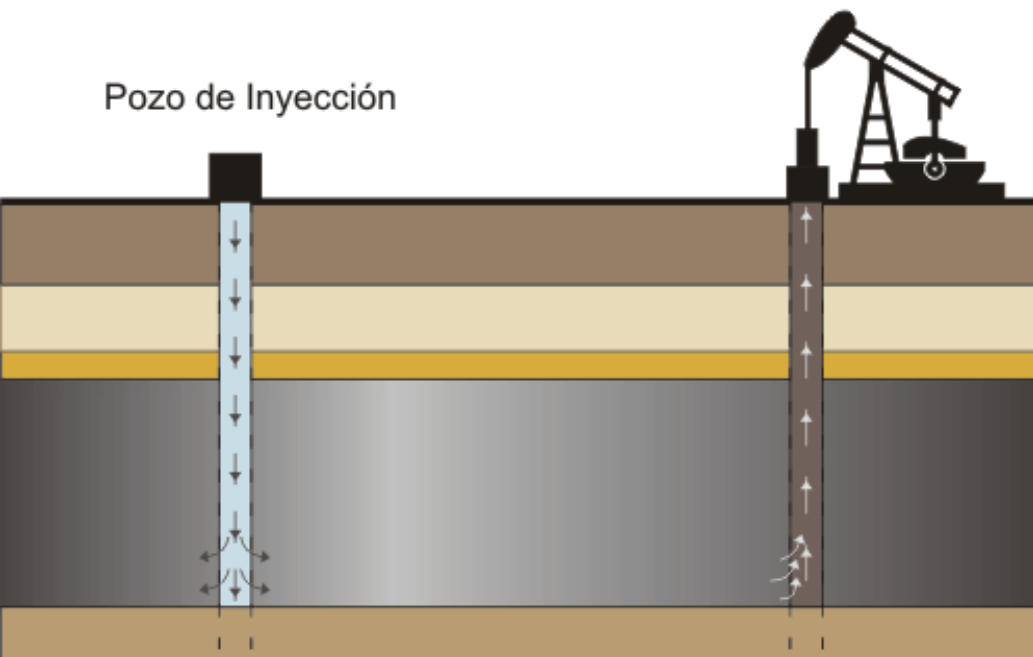
INYECTOR DE VAPOR

POZO PRODUCTOR



Pozo de Producción

Pozo de Inyección



Pozo de Producción

Pozo de Inyección

Dióxido de Carbono Inyección de Agua



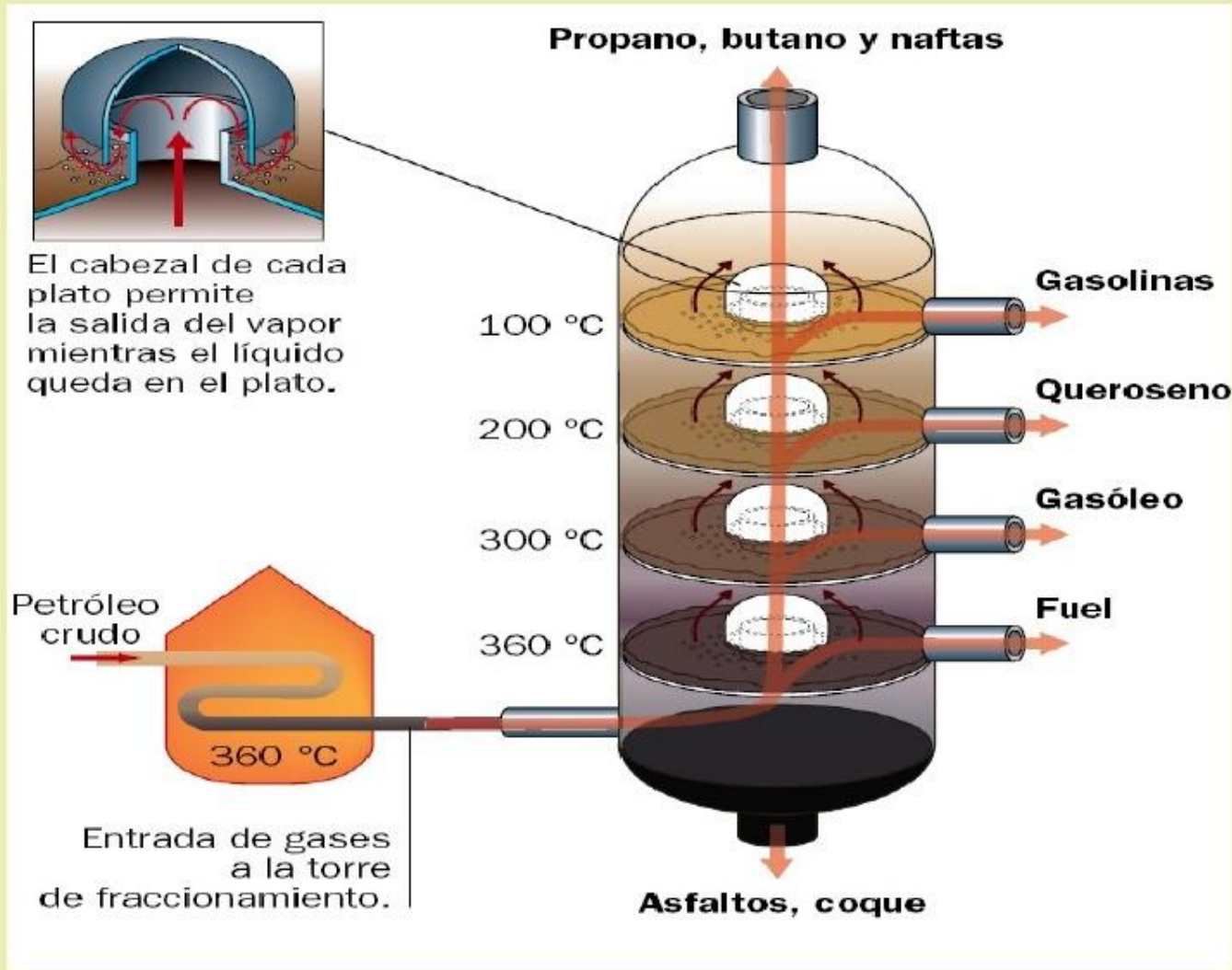


El oleoducto de Alaska (EEUU) transporta petróleo desde el campo de petróleo de la bahía de Prudhoe hasta los petroleros que atracan en el sur de Alaska. El oleoducto recorre 1.270 km de tierras deshabitadas y transporta hasta dos millones de barriles diarios de la costa ártica al golfo de Alaska.

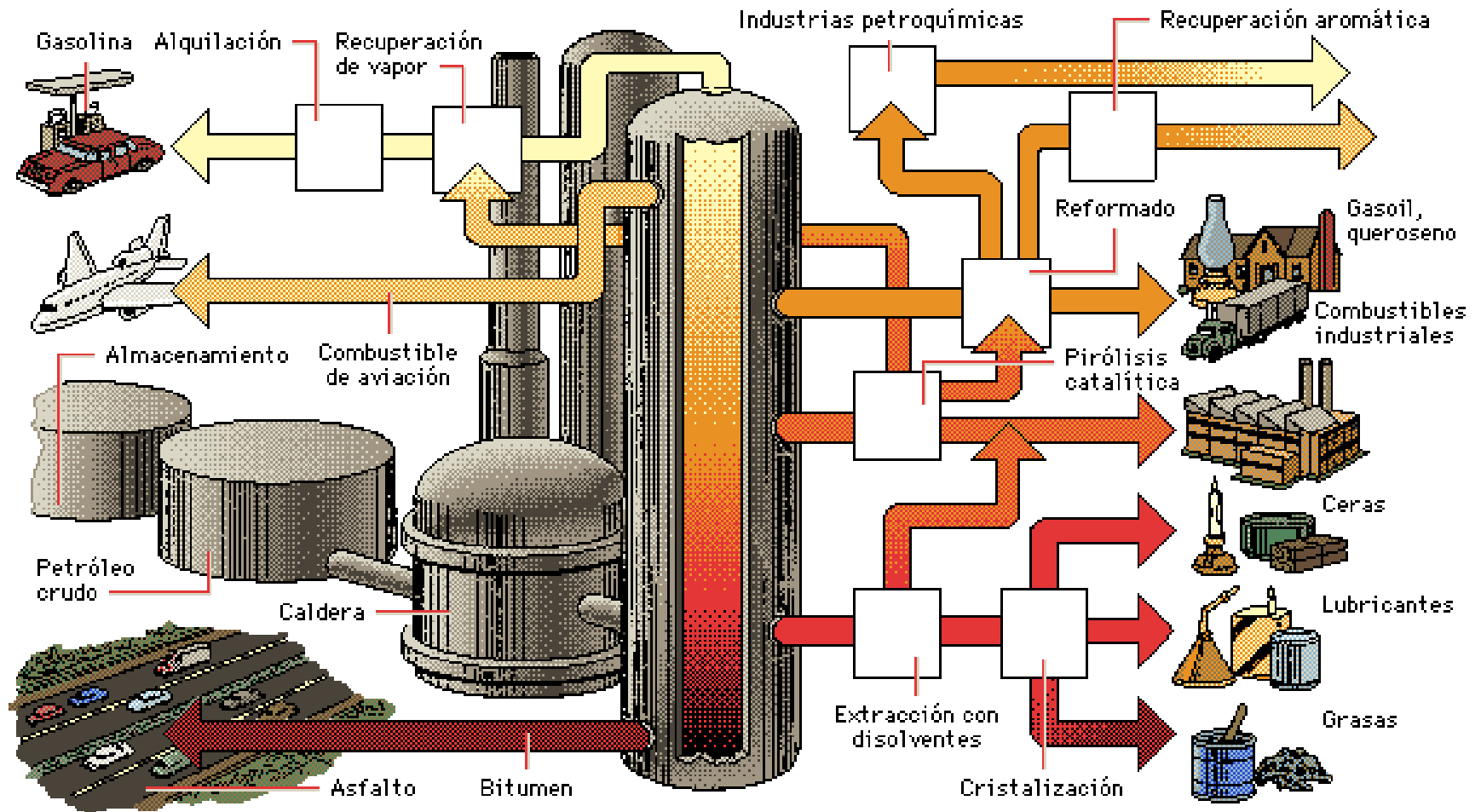
REFINACIÓN

- Esta operación sólo suministra productos en bruto, que deberán ser mejorados para su comercialización.
- Hoy, un barril de crudo produce 79,5 litros de gasolina, 11,5 litros de diesel, 34 litros de gasoil y destilados, 15 litros de lubricantes y 11,5 litros de residuos más pesados.

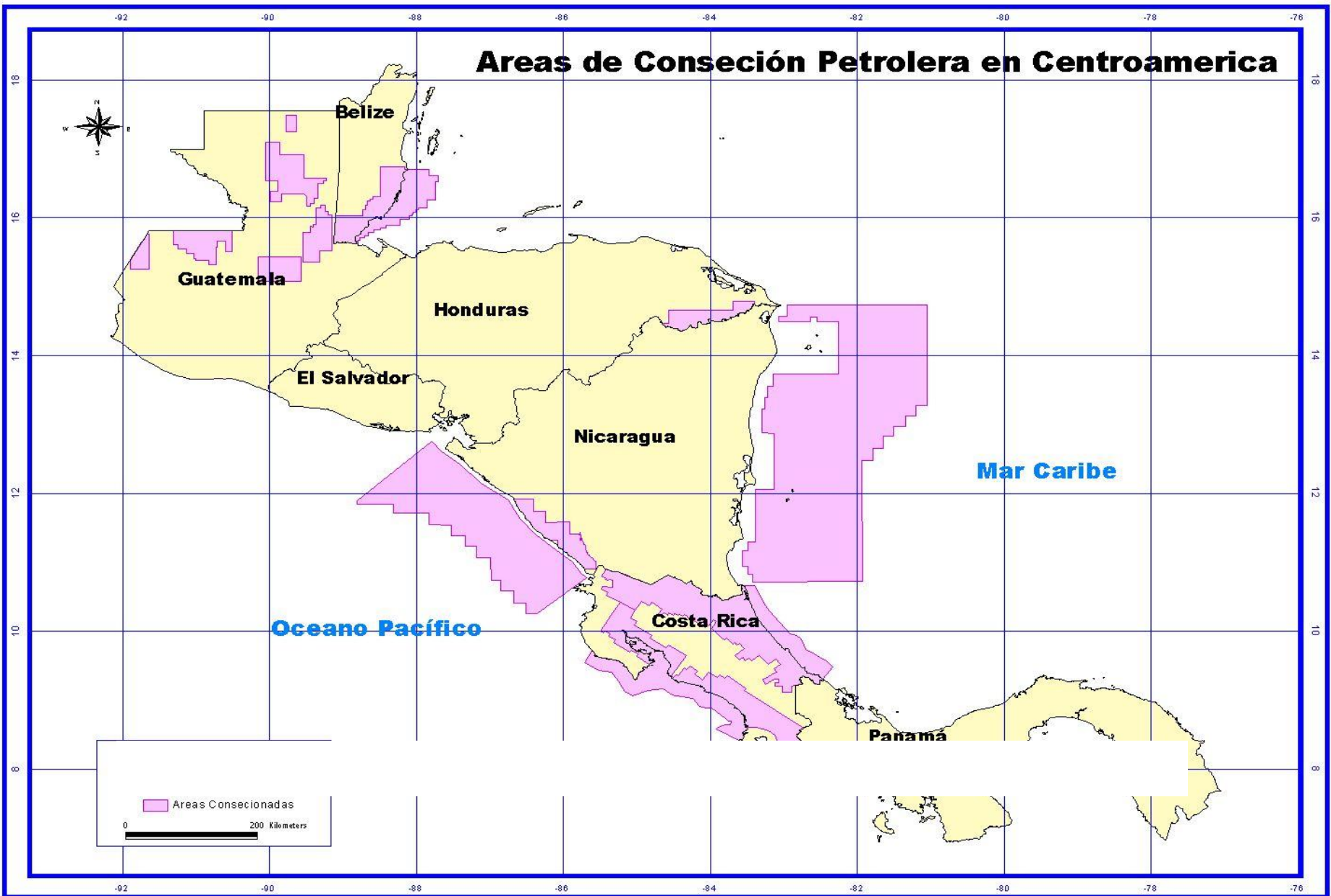
Destilación fraccionada



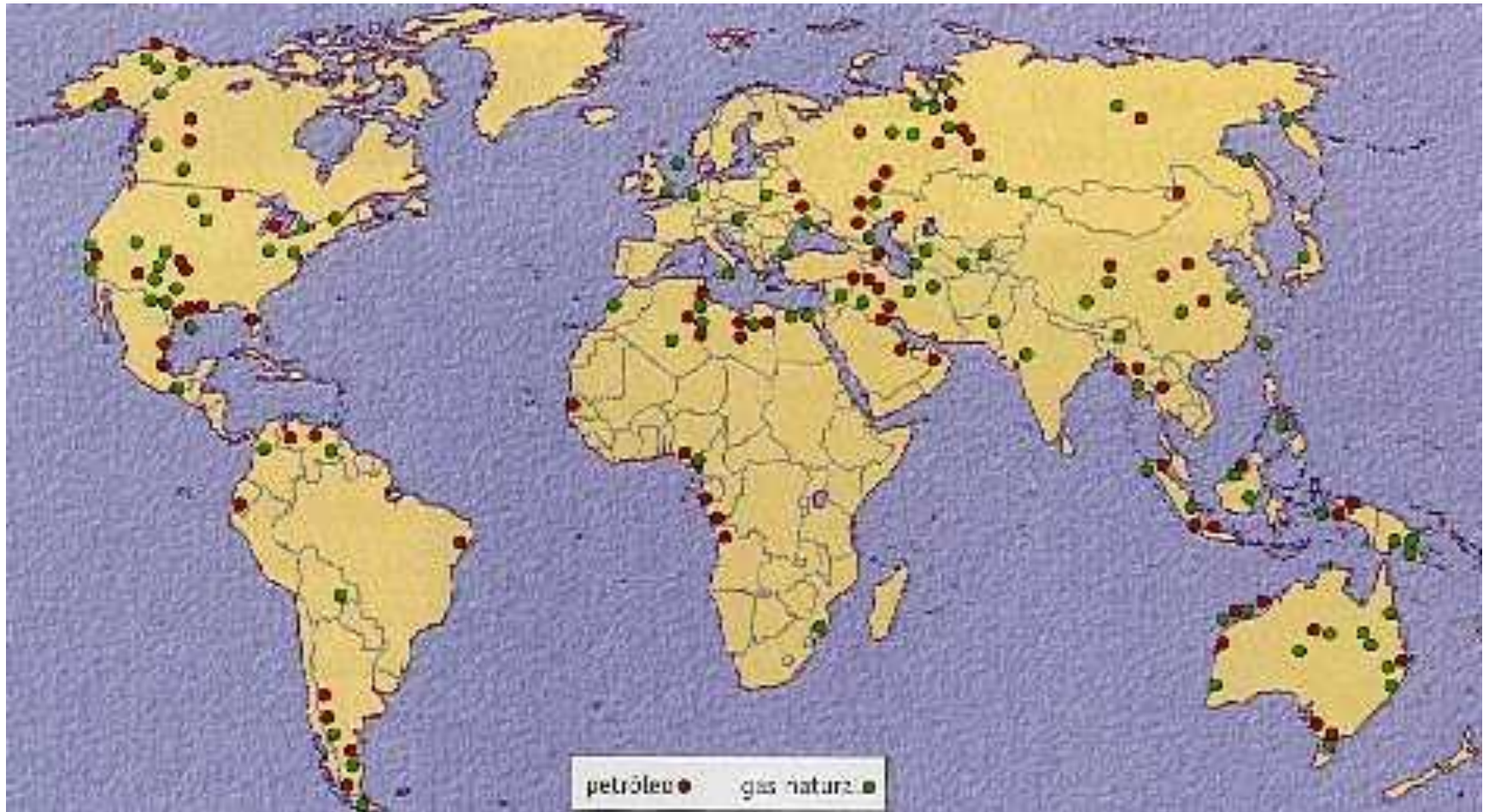
SISTEMA DE DESTILACIÓN DE HIDROCARBUROS



Áreas de Conseción Petrolera en Centroamérica



Países productores de petróleo

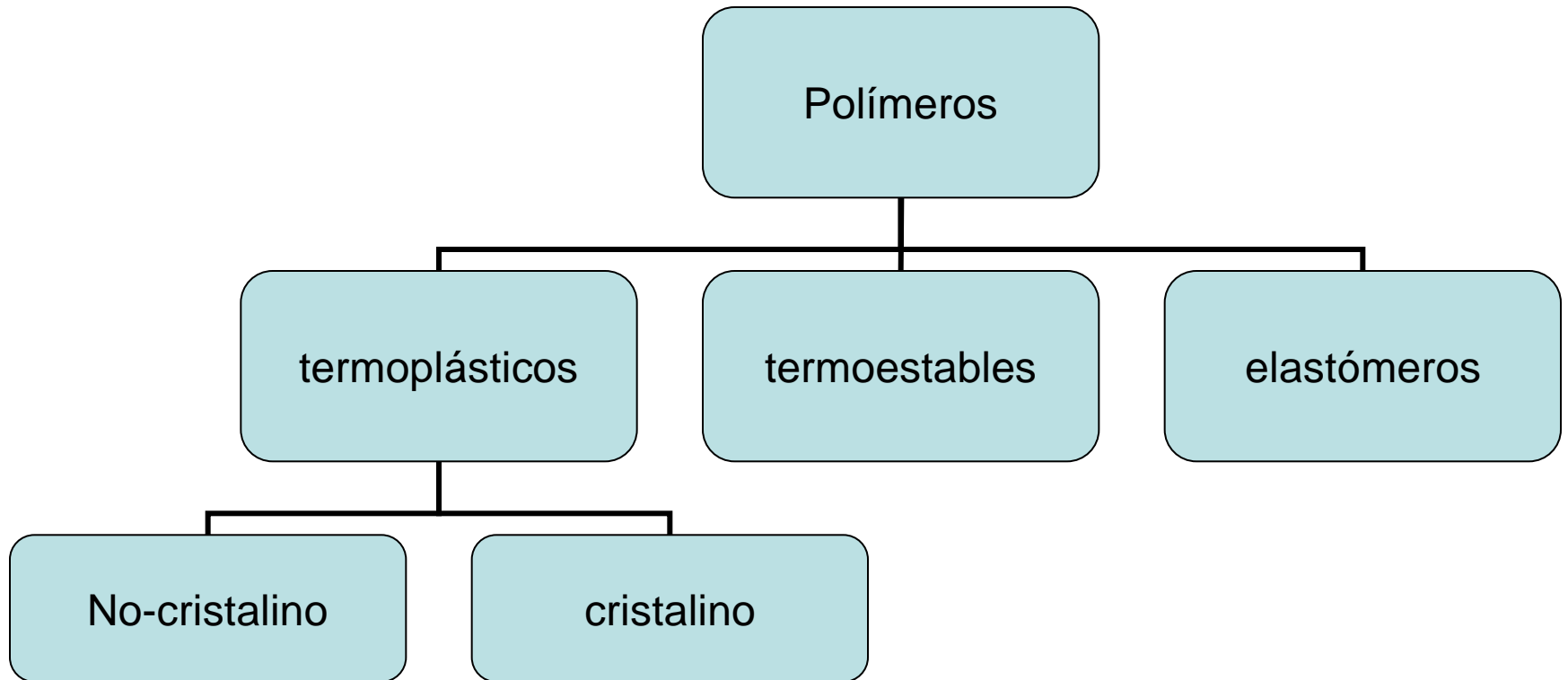


RESERVAS
DE
PETRÓLEO

Norteamérica	40
Asia	45
África	60
Europa	75
Suramérica	125
Irán	90
Irak	100
Kuwait	90
Arabia Saudí	245
Otros países del Oriente Próximo	120

billones de barriles de reserva

Clasificación de los polímeros

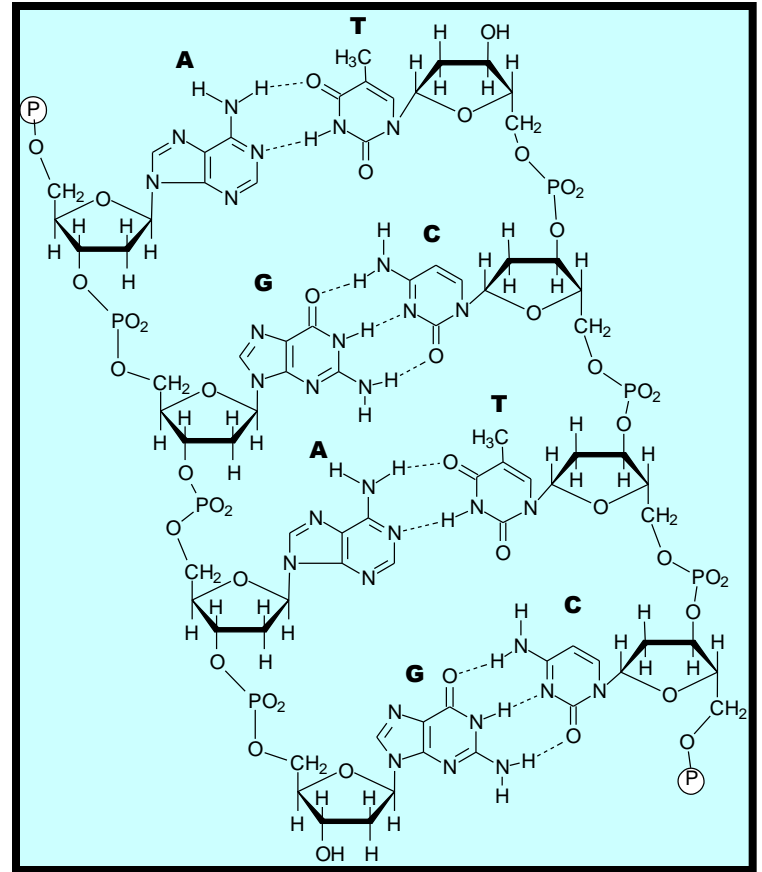
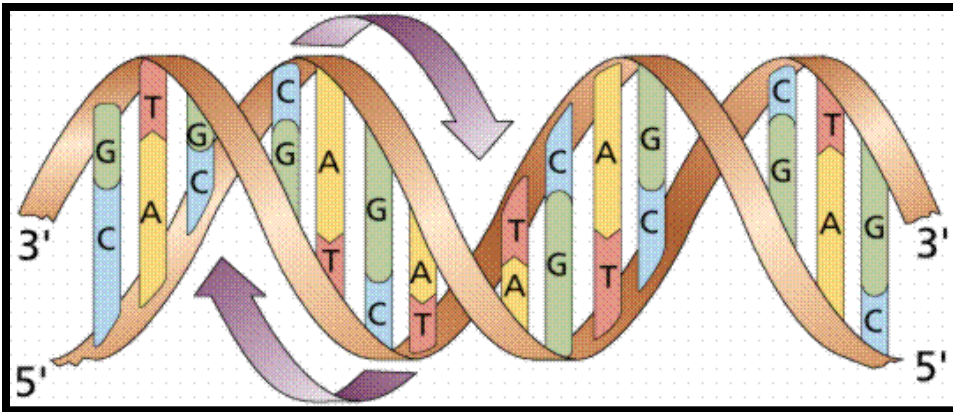
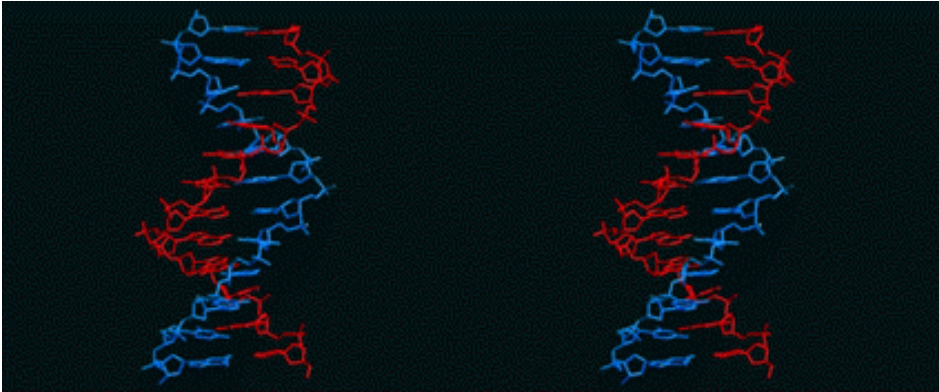


POLIMEROS SINTETICOS

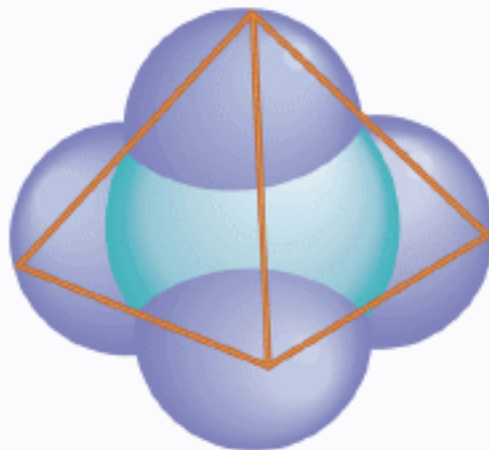
TIPOS DE POLÍMEROS

- **Termoplásticos:**
polietileno
- **Elastómeros:** **caucho**
- **Termoestables:**
baquelita
- **Fibras:** **poliéster**

POLIMEROS NATURALES: ADN



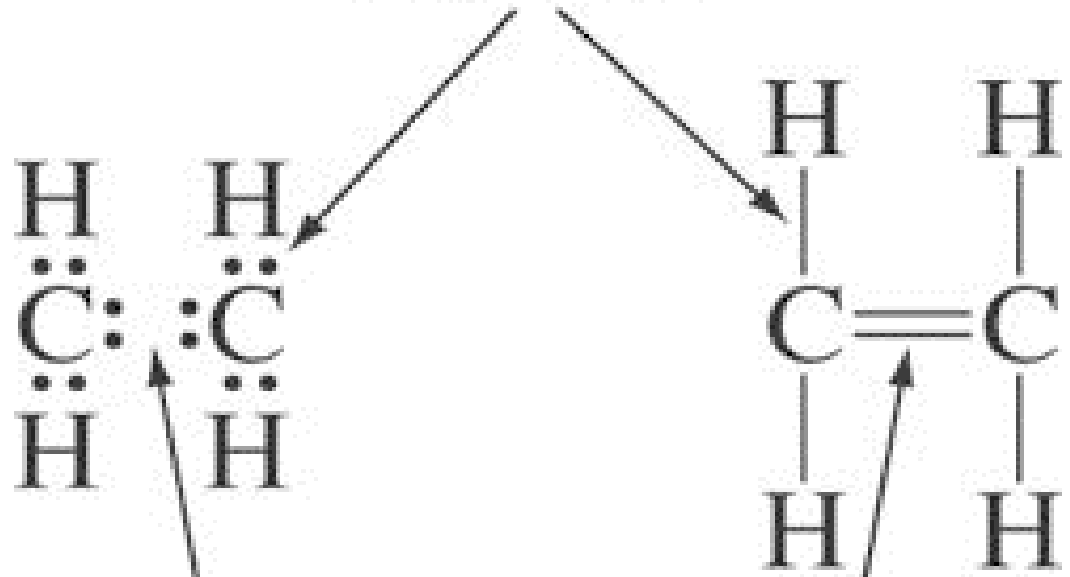
Unión covalente del metano



Hidrocarburo	Fórmula	Peso molecular	Densidad	T. de fusión
• Metano	CH ₄	16	gas	-182 °C
• Etano	C ₂ H ₆	30	gas	-183 °C
• Propano	C ₃ H ₈	44	gas	-190 °C
• Butano	C ₄ H ₁₀	58	gas	-138 °C
• Pentano	C ₅ H ₁₂	72	0,63	-130 °C
• Hexano	C ₆ H ₁₄	86	0,66	-95 °C
• Heptano	C ₇ H ₁₆	100	0,68	-91 °C
• Octano	C ₈ H ₁₈	114	0,70	-57 °C
• Nonano	C ₉ H ₂₀	128	0,72	-52 °C
• Decano	C ₁₀ H ₂₂	142	0,73	-30 °C
• Undecano	C ₁₁ H ₂₄	156	0,74	-25 °C
• Dodecano	C ₁₂ H ₂₆	170	0,75	-10 °C
• Pentadecano	C ₁₅ H ₃₂	212	0,77	10 °C
• Eicosano	C ₂₀ H ₄₂	283	0,79	37 °C
• Triacontano	C ₃₀ H ₆₂	423	0,78	66 °C
• Polietileno	C ₂₀₀₀ H ₄₀₀₂	28000	0,93	100 °C

Densidad y temperatura de fusión de hidrocarburos.

Enlaces
covalentes
sencillos

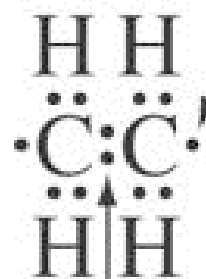


Enlaces
covalentes
dobles

a)

b)

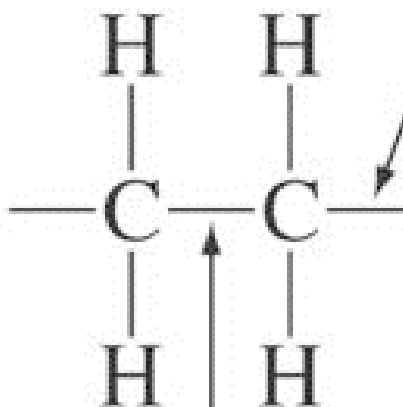
Electrón libre disponible
para enlace covalente



Enlace
covalente
sencillo

a)

Medio enlace covalente
o electrón libre

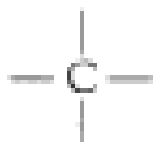


Enlace
covalente
sencillo

b)

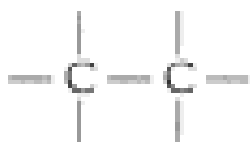
Nomenclatura de los compuestos orgánicos

Meth-1 Carbon



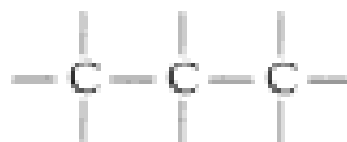
Methane

Eth-2 Carbons



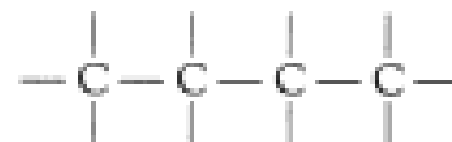
Ethane

Prop-3 Carbons



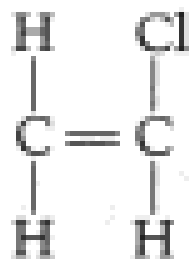
Propane

But-4 Carbons

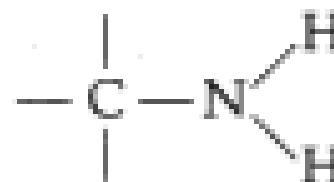


Butane

a) Cadenas saturadas

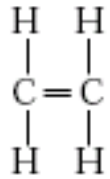


b) Cadena no saturada

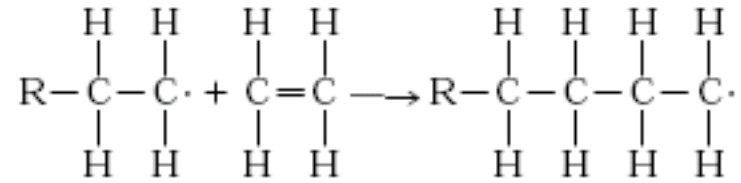
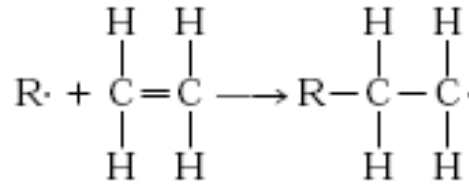


c) radical

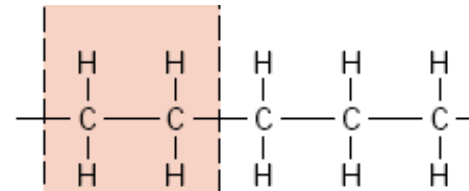
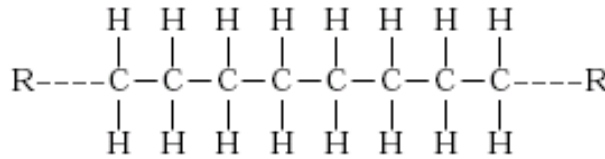
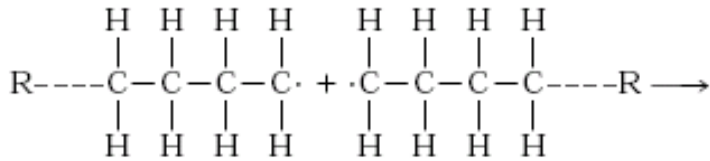
Síntesis del Polietileno (PE)



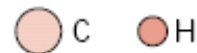
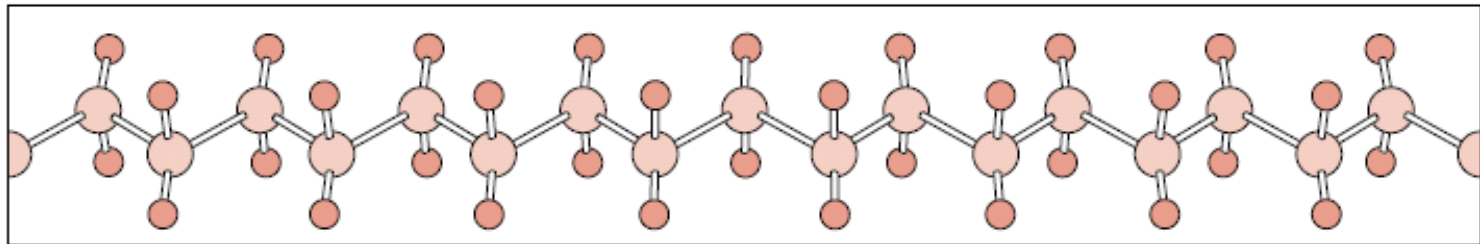
monómero

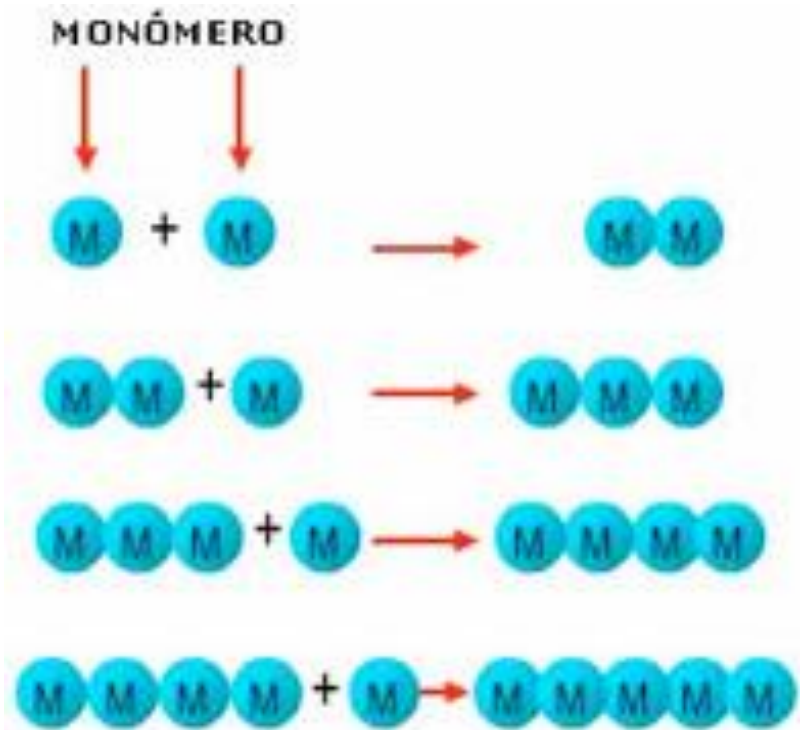


radical

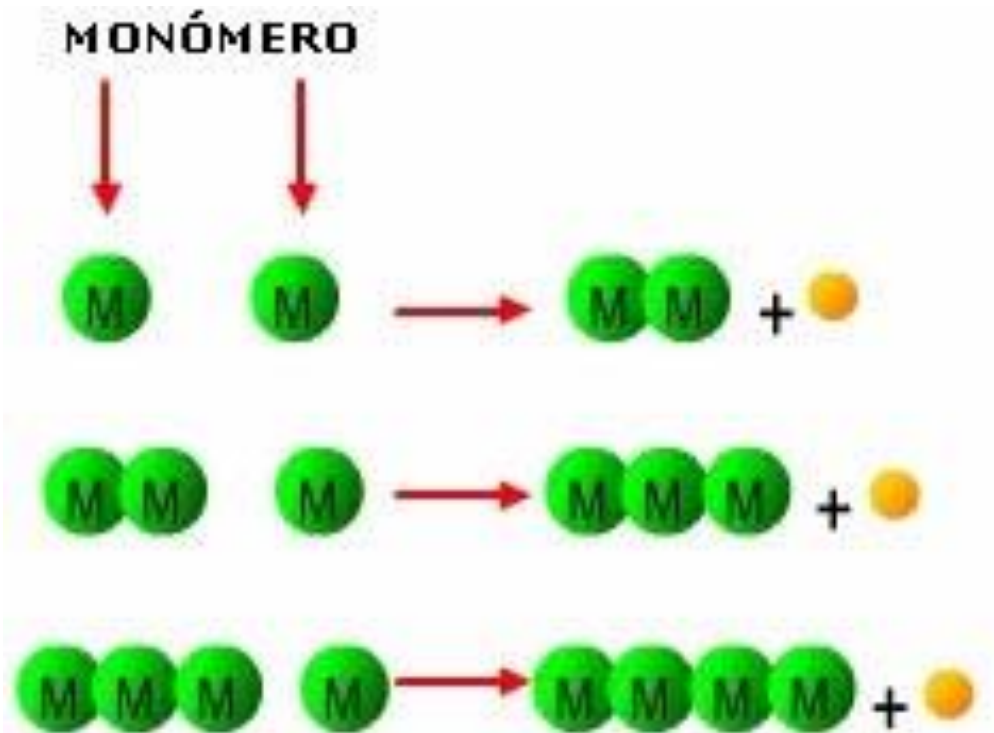


mero

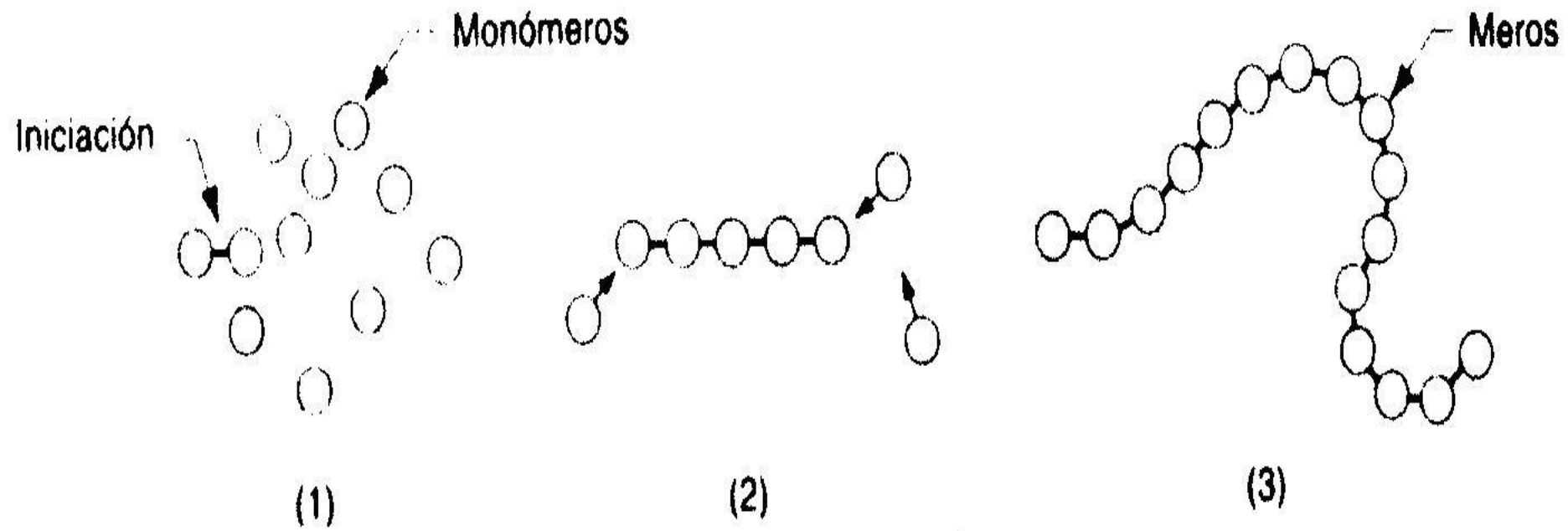




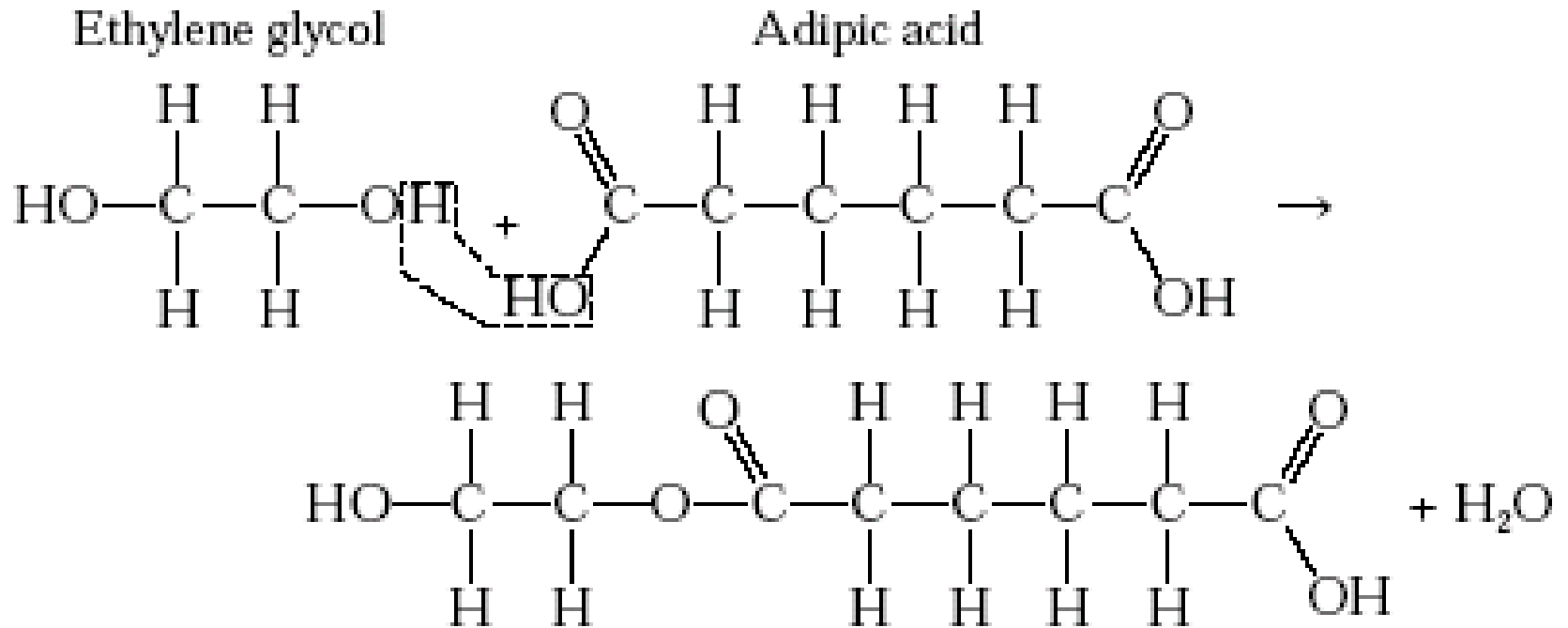
Esquema de polimerización por adición



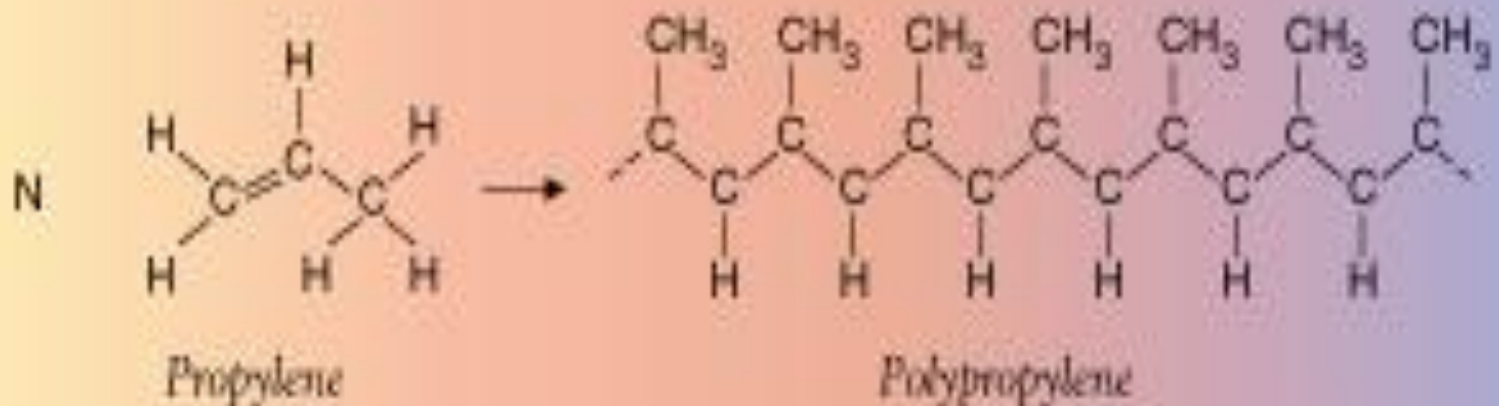
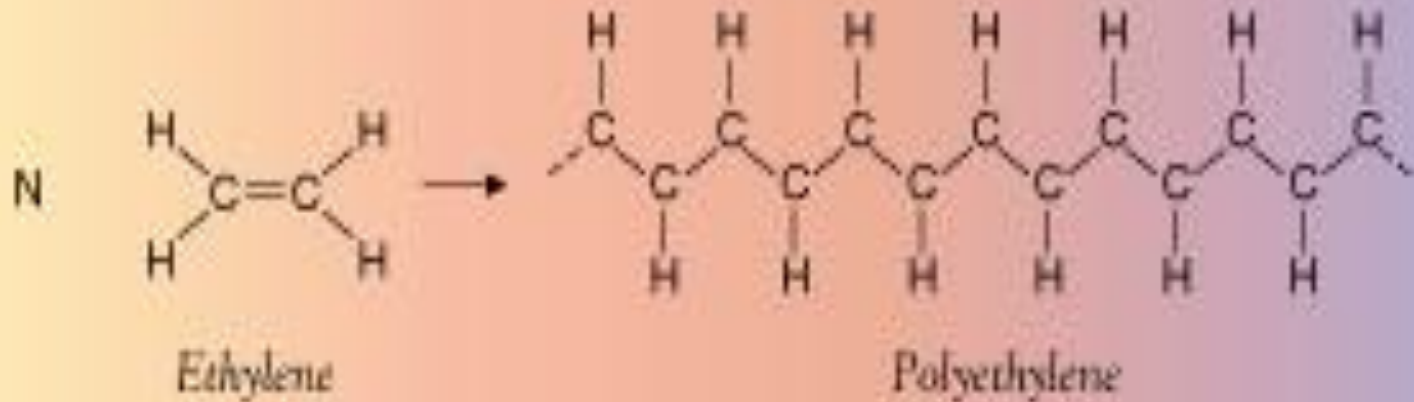
Esquema de polimerización por condensación



Polimerización por condensación



Polimerización del etileno y el propileno para obtener polietileno y polipropileno



Polímeros de adición

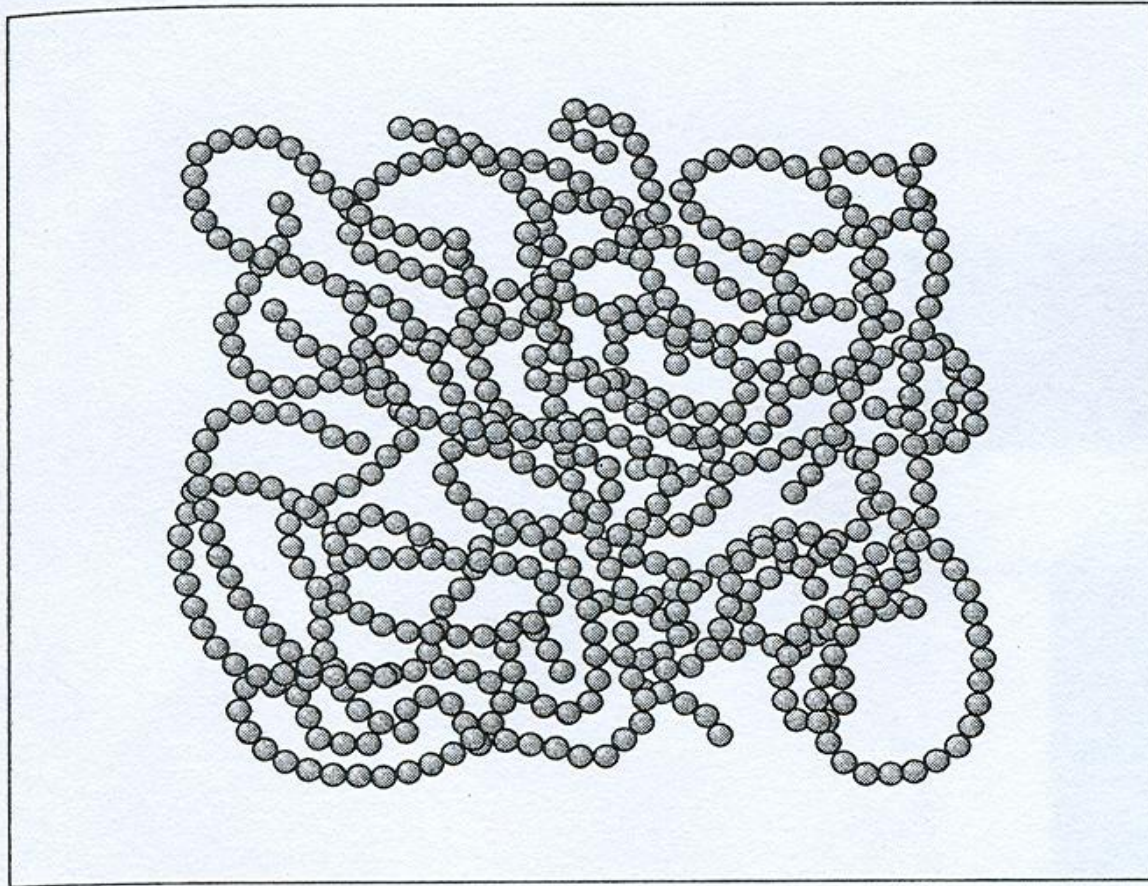
• MONÓMEROS

- Eteno
- Propeno
- cloroeteno
- tetraflúoreteno
- propenonitrilo
- butadieno
- fenileteno
- 2-clorobutadieno

• POLÍMEROS

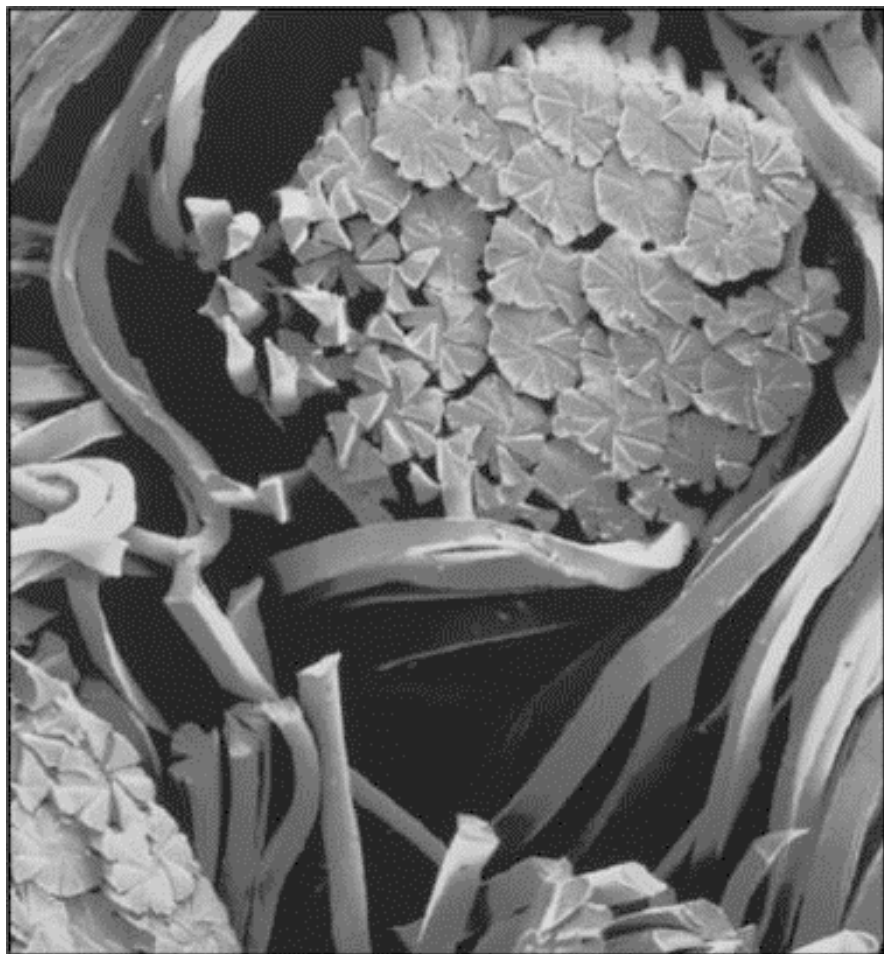
- ➡ – Polietileno
- ➡ – Polipropileno
- ➡ – policloruro de vinilo
- ➡ – teflón
- ➡ – poliacrilonitrilo
- ➡ – polibutadieno
- ➡ – poliestireno
- ➡ – neopreno

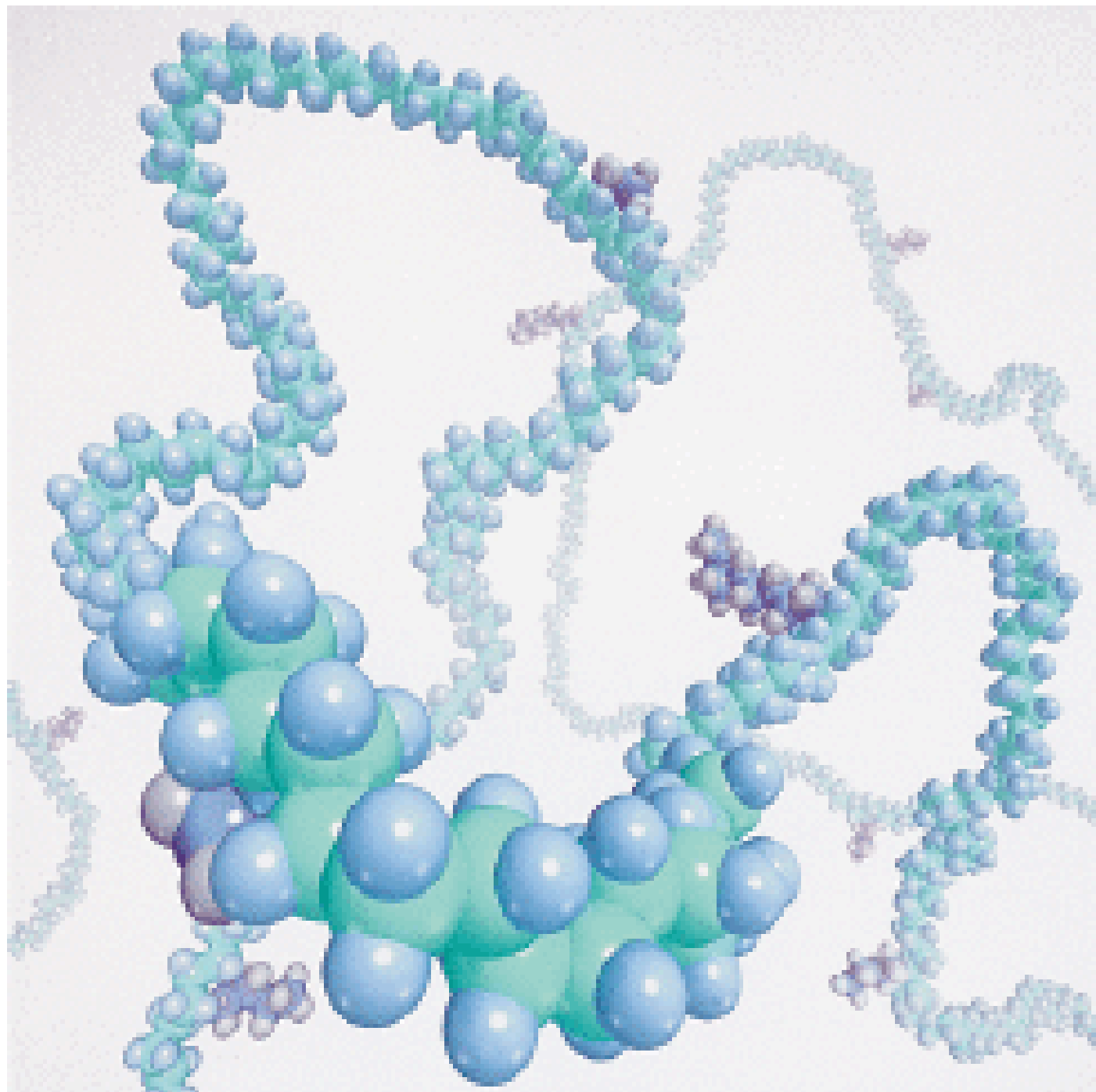
Cadenas de polímeros unidos por fuerzas de Van der Waals



(Según W. G. Moffatt, W. G. Pearsall y J. Wulff, «*The Structure and Properties of Materials*», vol. I, «*Structure*», Wiley, 1965, p. 104.)

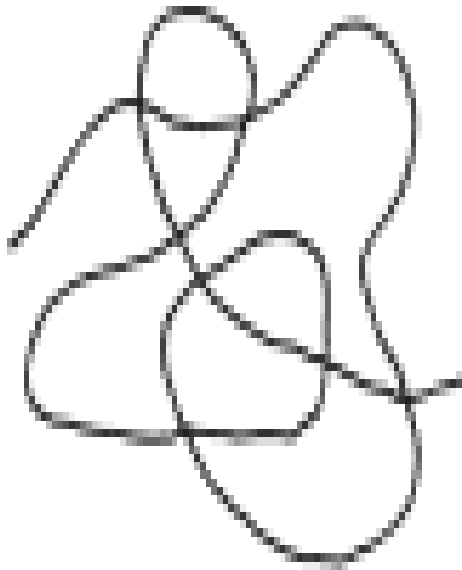
FIGURA 7.5. Representación esquemática de un polímero. Las esferas representan las unidades de la cadena del polímero de átomos no especificados.





Formas en que se pueden manifestar las cadenas de polímeros

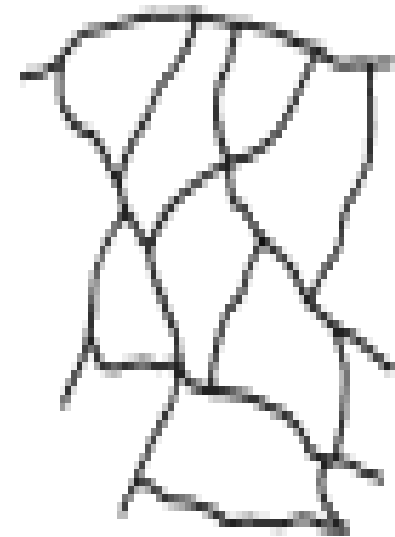
ALBERTO RAMÍREZ RAMÍREZ



Linear

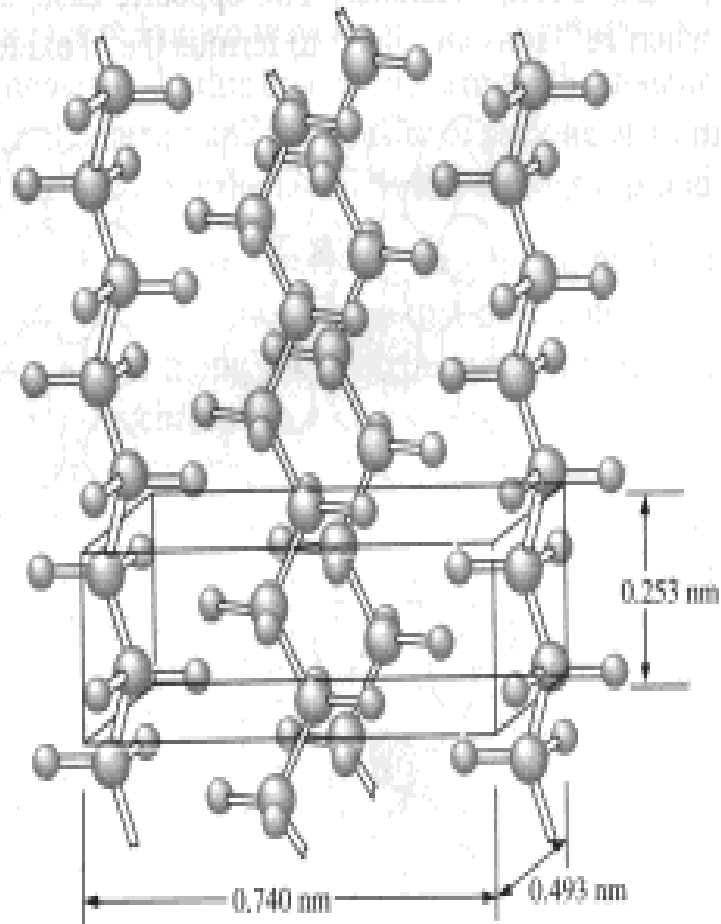


Branched



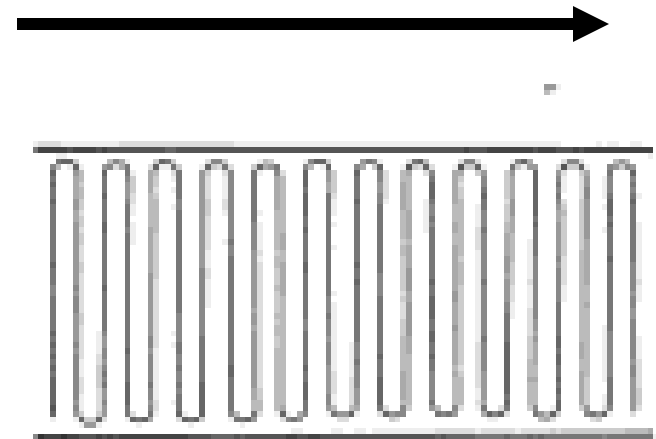
Network

Estructura cristalina del polietileno



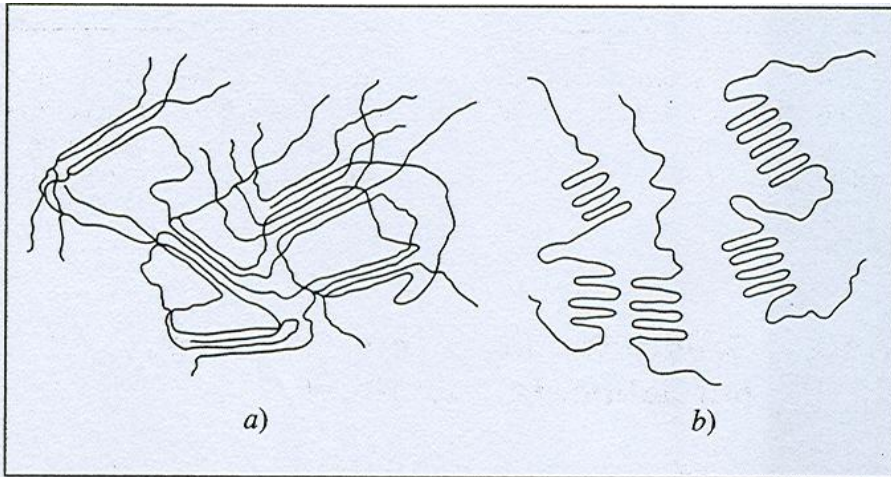
a) Alineación de la cadena cristalina del polietileno

Crecimiento del cristal



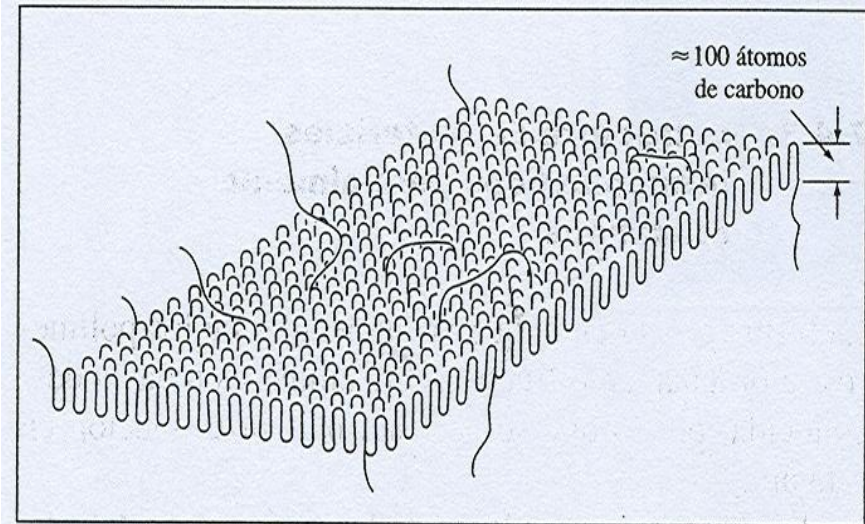
b) Ordenamiento de la cadena

Cadenas de polímeros de ordenaciones cristalinas



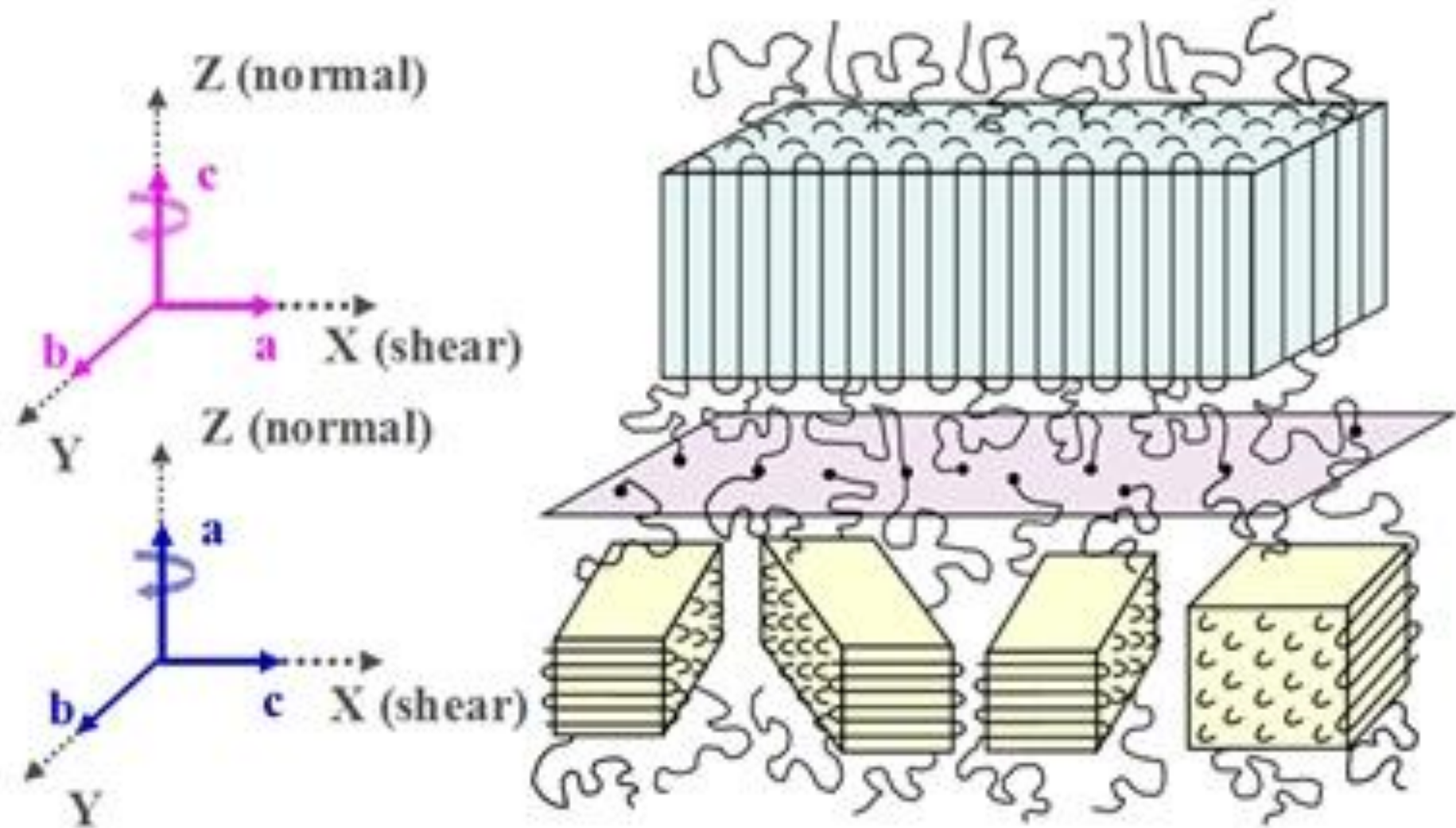
(Según F. Rodríguez, «Principles of Polymer Systems», 2.^a ed., McGraw-Hill, 1982, p. 42.)

FIGURA 7.16. Dos ordenaciones cristalinas sugeridas para materiales termoplásticos parcialmente cristalinos. a) Modelo de micela con flecos, y b) modelo de cadenas plegadas.



[Según R. L. Boysen, *Olefin Polymers (High-Pressure Polyethylene)*, en «Encyclopedia of Chemical Technology», vol. 16, Wiley, 1981, p. 405.]

FIGURA 7.17. Esquema de la estructura de cadena plegada de una lámina de polietileno de baja densidad.



Nivel de consumo de los principales polímeros


	<i>Use % of market</i>
Polyethylene, PE	32
Polypropylene, PP	11
Polystyrene, PS	9
Polyvinylchloride, PVC	<u>15</u>
	67

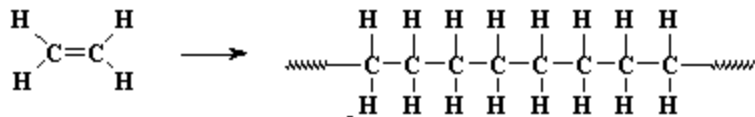
Algunos radicales substitutos del hidrogeno en la cadena

• H	-
• F	flouro
• Cl	cloro
• Br	bromo
• I	iodo
• CH ₃	methyl
• CH ₃ (CH ₂) _n	alkyl
• CN	cianuro
• NH ₂	amino
• N(CH ₃) ₂	dimethylamino
• NO ₂	nitro

Algunos monómeros para formar polímeros

TABLE 13.1 Summary of Important Polymers

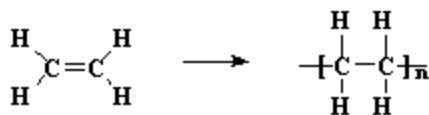
Group I. Thermoplastics		
Polymer	Percentage of Market	Monomer(s) Used
Polyethylene	32	$\begin{array}{c} \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{H} \\ \quad \\ \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{Cl} \end{array}$
Polyvinylchloride	15	
Polystyrene	9	$\begin{array}{c} \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{C}_6\text{H}_5 \end{array}$ <p>( is benzene, C₆H₆)</p>
Polypropylene	11	$\begin{array}{c} \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{CH}_3 \end{array}$
ABS	3	$\begin{array}{c} \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{C} = \text{N} \\ \quad \quad \quad \\ \text{H} \quad \text{H} \quad \text{H} \quad \text{H} \\ \quad \quad \quad \\ \text{C} - \text{C} - \text{C} = \text{C} \\ \quad \quad \quad \\ \text{H} \quad \text{H} \quad \text{H} \quad \text{H} \end{array}$
		<p>Acrylonitrile (graft)</p>
		<p>Butadiene (chain)</p>
		$\begin{array}{c} \text{H} \quad \text{H} \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{C}_6\text{H}_5 \end{array}$ <p>Styrene (graft)</p>
Acrylics (examples: polymethyl methacrylate, Lucite)	1	$\begin{array}{c} \text{H} \quad \text{CH}_3 \\ \quad \\ \text{C} = \text{C} \\ \quad \\ \text{H} \quad \text{C} = \text{O} \\ \\ \text{O} \\ \\ \text{CH}_3 \end{array}$



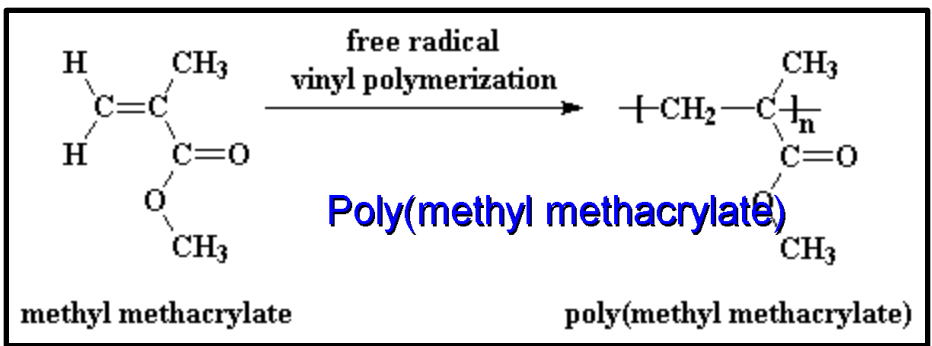
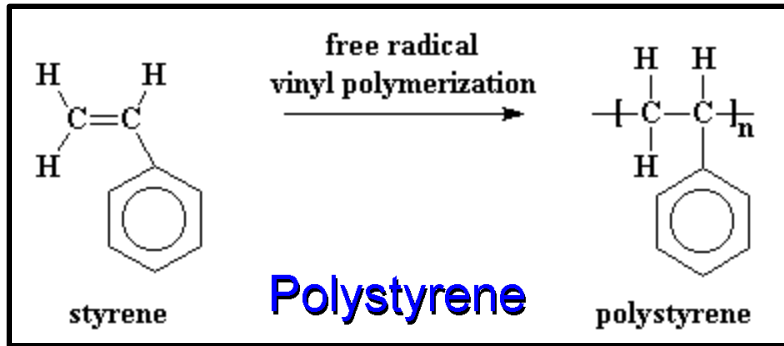
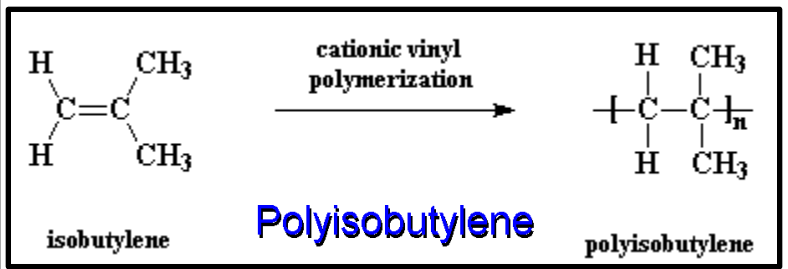
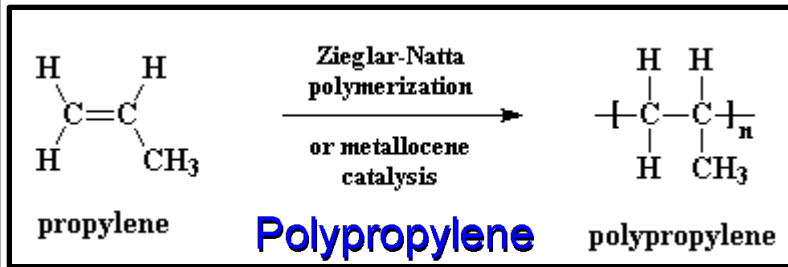
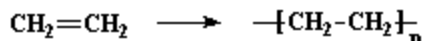
This can get tedious to draw, so we often use shorthand like this.

Polietileno

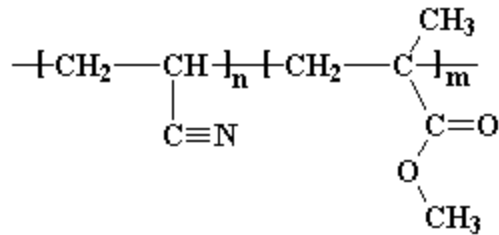
(Note: A line drawn between two atoms represents a pair of electrons shared by those atoms, which constitutes a chemical bond. Two lines represent two pairs of shared electrons, a double bond.)



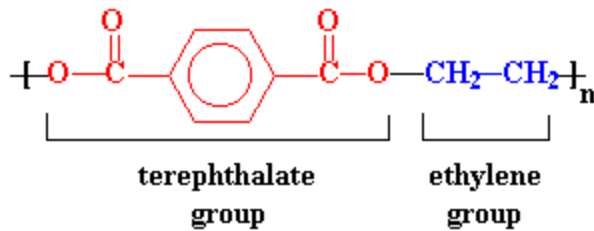
And when we're feeling really lazy we just draw it like this:



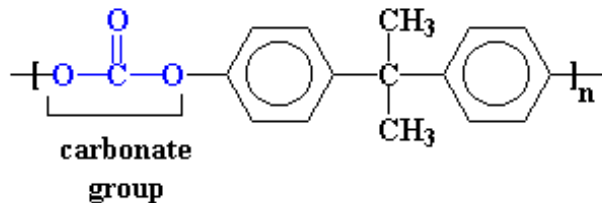
EJEMPLO DE POLIMEROS



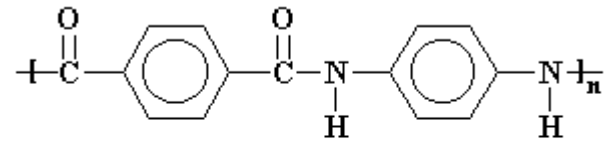
poly(acrylonitrile-*co*-methyl methacrylate)



Polyesters

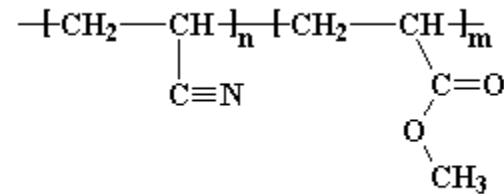


Polycarbonates

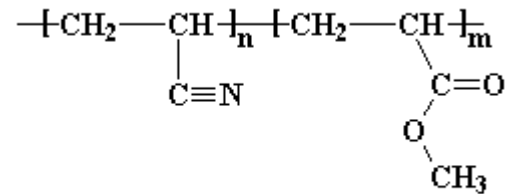


In Kevlar the aromatic groups are all linked into the backbone chain through the 1 and 4 positions. This is called *para*-linkage.

Aramids

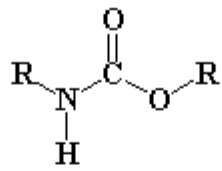


poly(acrylonitrile-*co*-methyl acrylate)

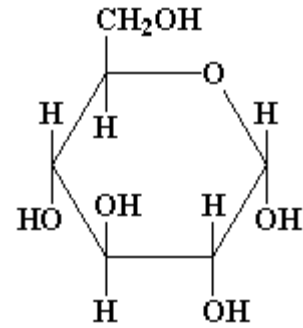


poly(acrylonitrile-*co*-methyl acrylate)

Polyacrylonitrile

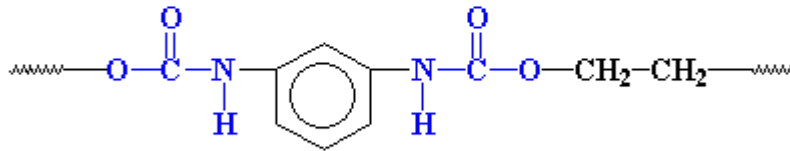


a urethane



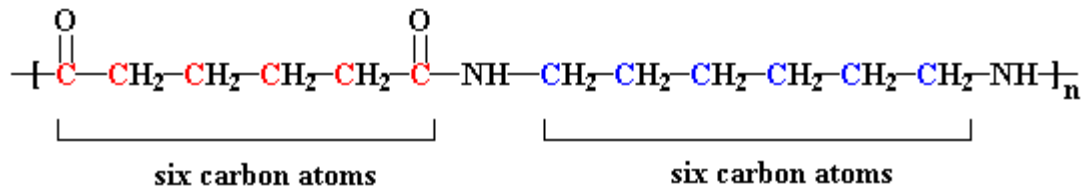
glucose

Cellulose



the urethane linkages
in a polyurethane

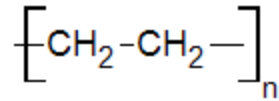
Polyurethanes



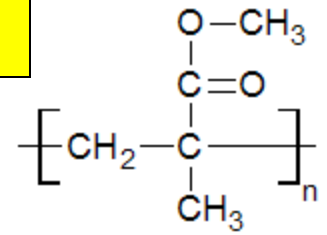
Nylons

EJEMPLO DE POLIMEROS

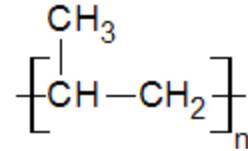
Polietileno



Polimetacrilato de metilo

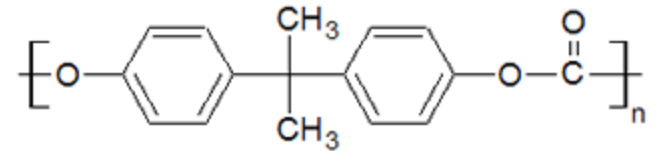
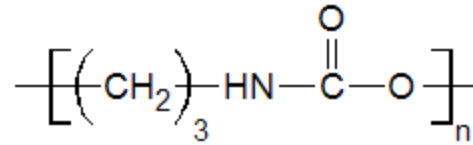


Polipropileno

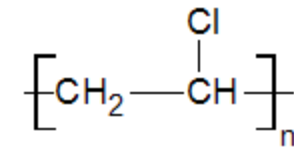


poli(4,4'-isopropilidendifenol carbonato (lexan)

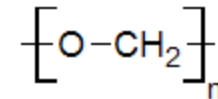
Poliuretano



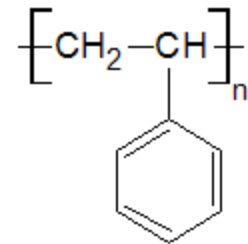
Cloruro de polivinilo
Poli(cloruro de vinilo)
(PVC)



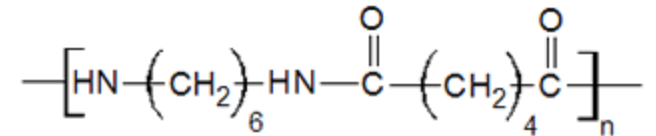
Polioximetileno



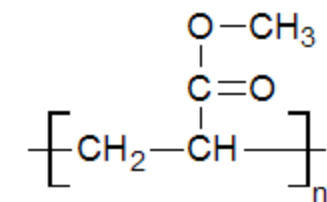
Poliestireno



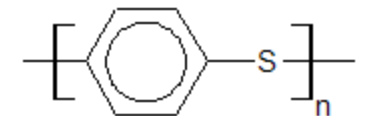
Nylon



Poliacrilato de metilo



Poli(sulfuro de fenileno)



Procesos de fabricación de polipropileno

Aplicaciones del polipropileno

- Autopartes
- Baldes, recipientes, botellas
- Muebles
- Juguetes
- Películas para envases de alimentos
- Fibras y filamentos
- Bolsas y bolsones
- Fondo de alfombras
- Pañales, toallas higiénicas, ropa
- Envases de pared delgada

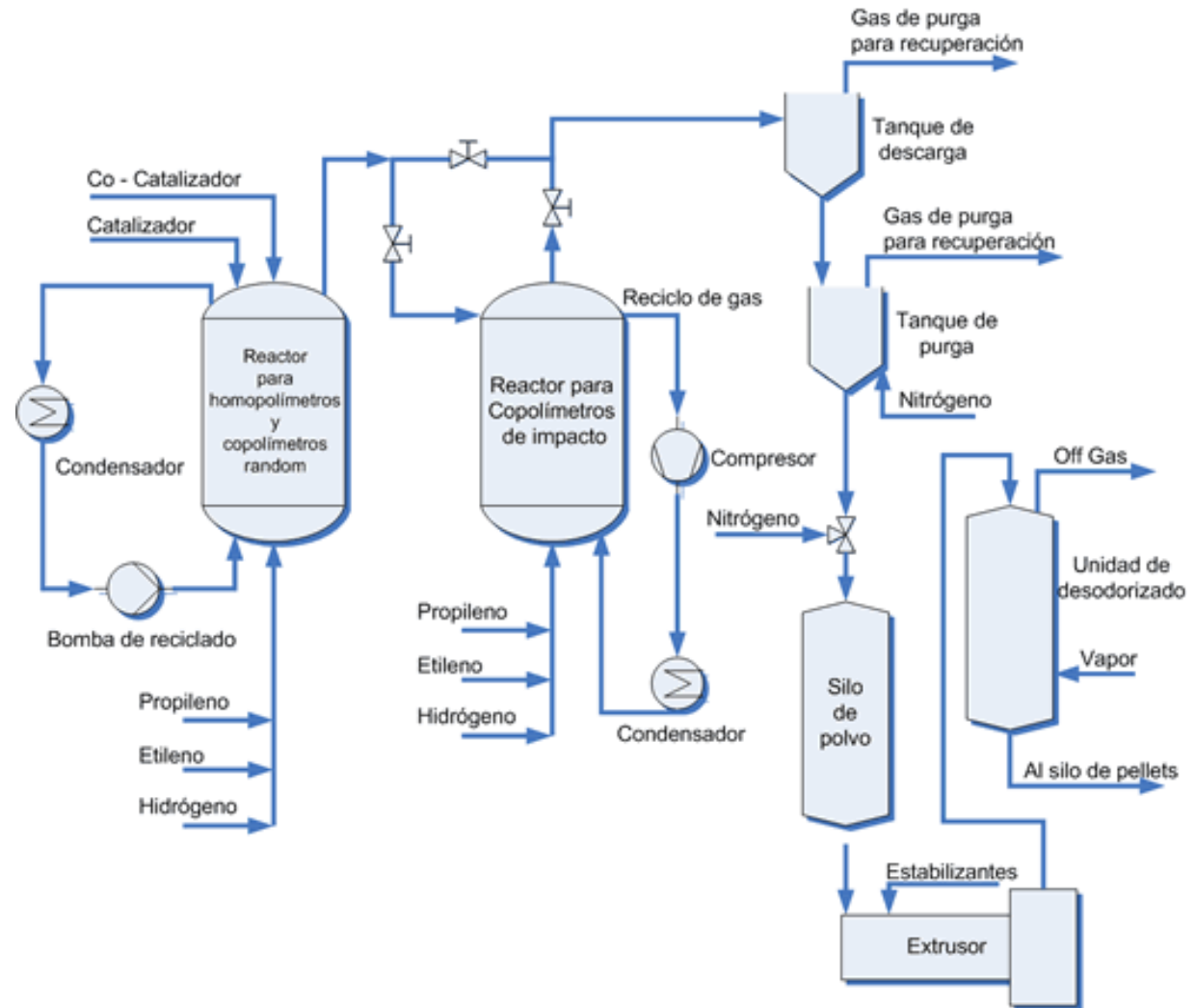
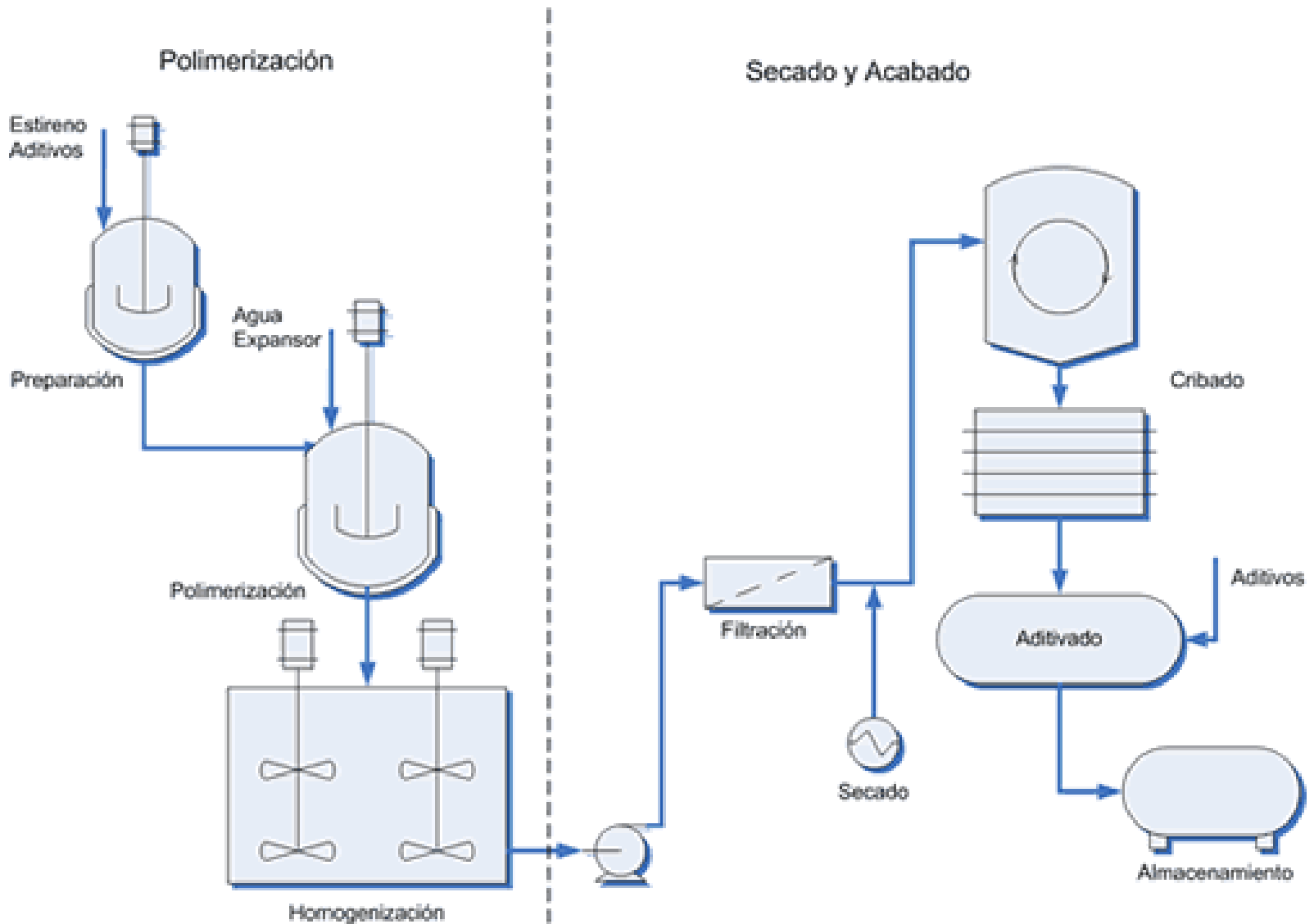
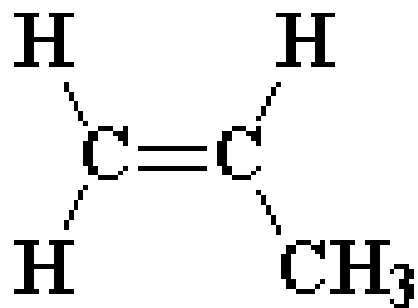


Diagrama de flujo de Polimerización



Isómeros del polipropileno

(misma composición química
pero diferentes característica física)

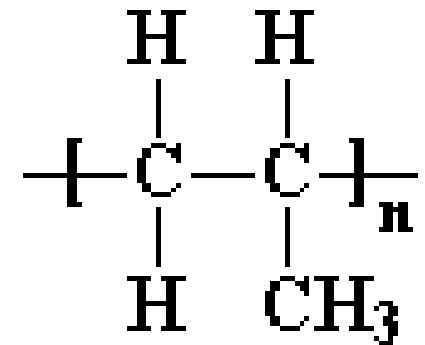


propylene

Ziegler-Natta
polymerization

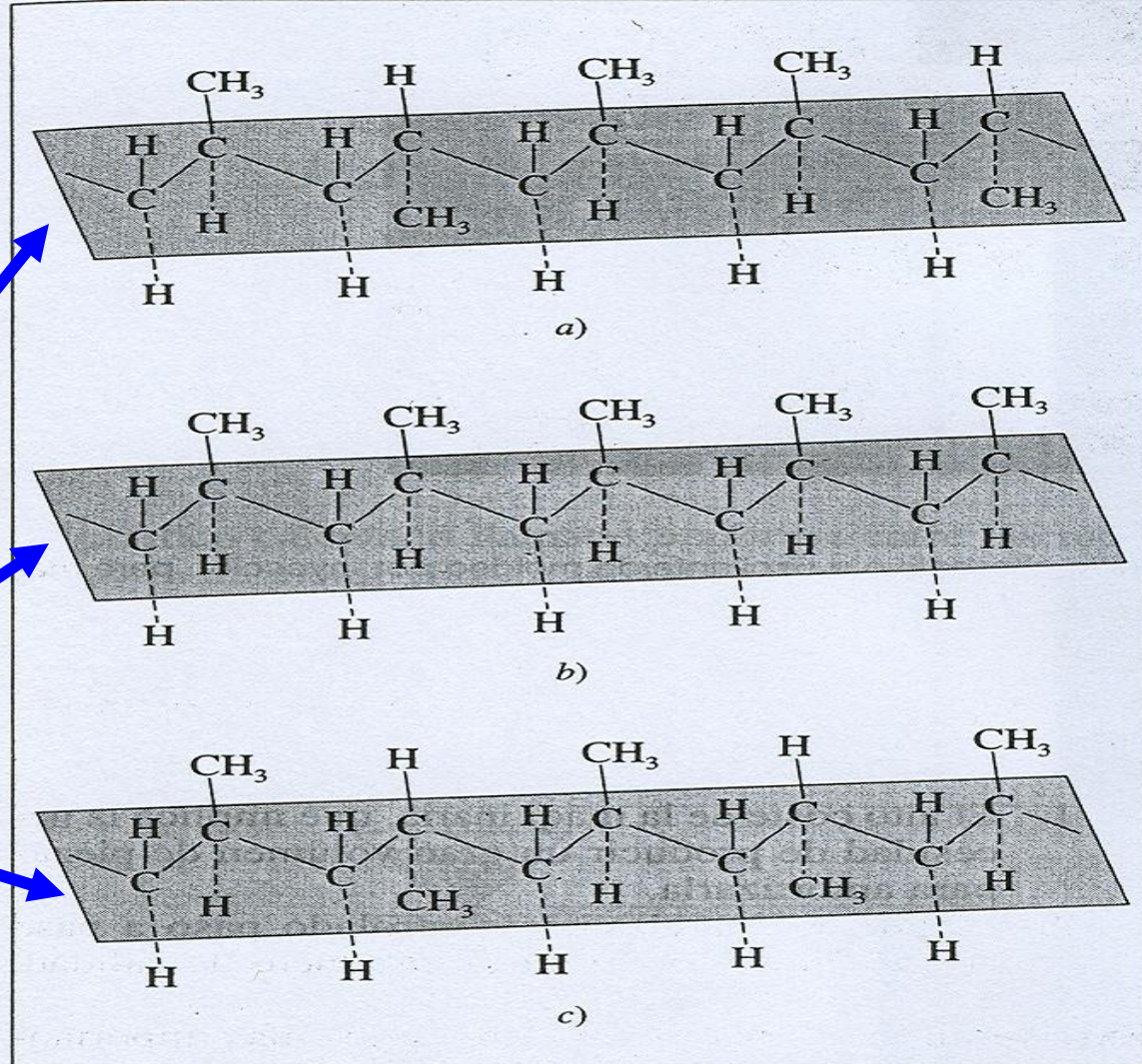


or metallocene
catalysis



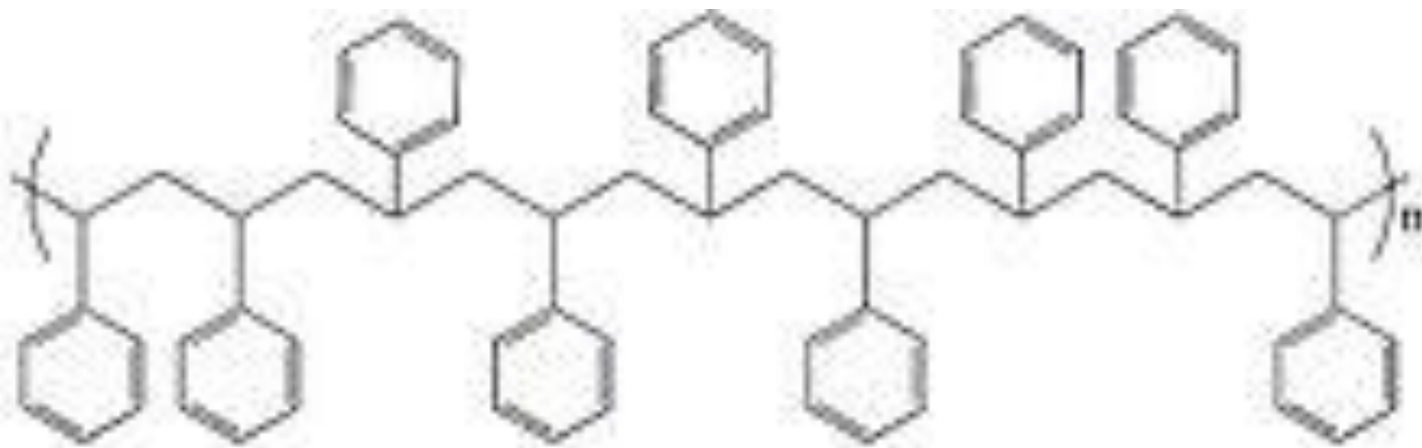
polypropylene

Isómeros:
atáctico
isotáctico
sindiotáctico

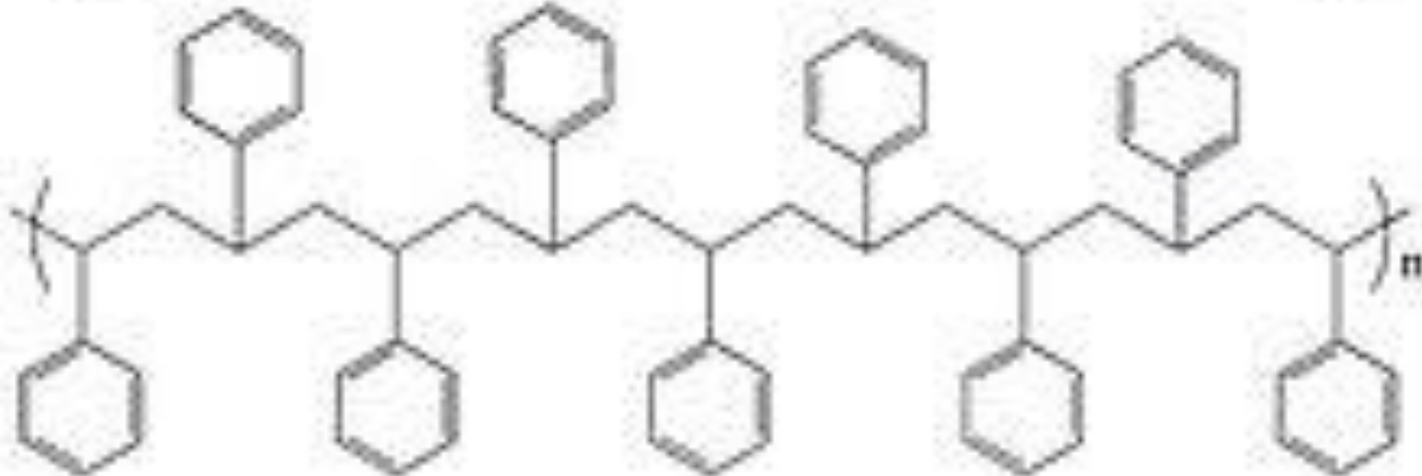


[Según G. Crespi y L. Luciani, *Olefin Polymers (Polyethylene)*, en «*Encyclopedia of Chemical Technology*», vol. 16, Wiley, 1982, p. 454.]

FIGURA 7.19 Estereoisómeros del polipropileno. a) Isómero atáctico en el que los grupos metilo sustituyentes se encuentran dispuestos aleatoriamente a uno u otro lado de la cadena principal; b) isómero isotáctico, en el que los grupos CH₃ sustituyentes se encuentran todos del mismo lado de la cadena principal, y c) isómero sindiotáctico en el cual los grupos CH₃ están ubicados de modo regular y alternado a uno y otro lado de la cadena principal.



Atáctico

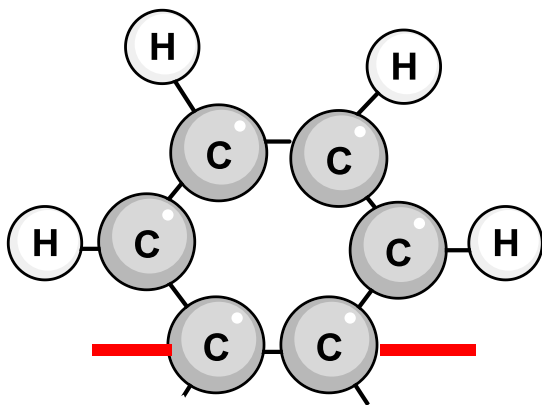


Sindiotáctico

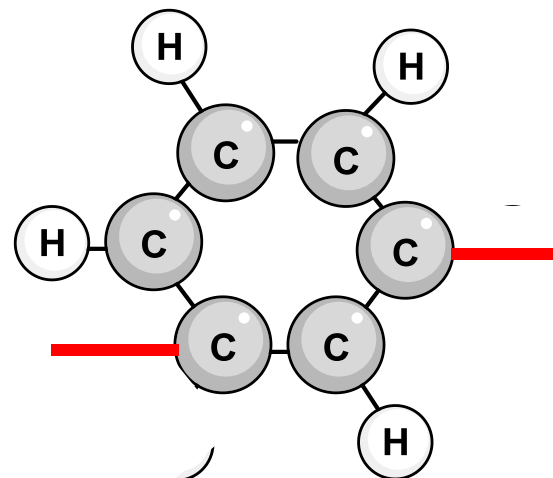


Isotáctico

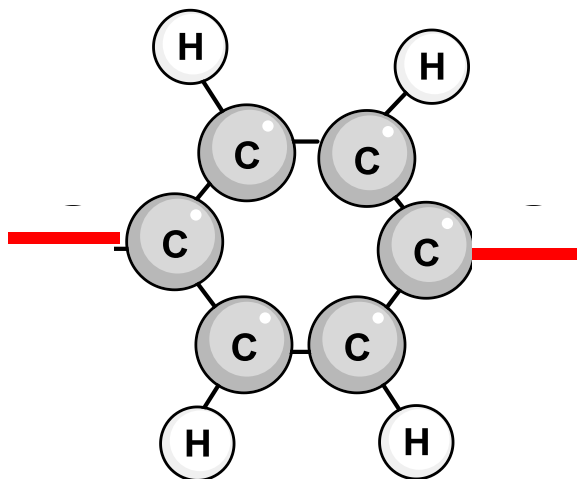
Isómeros de benceno



ORTO



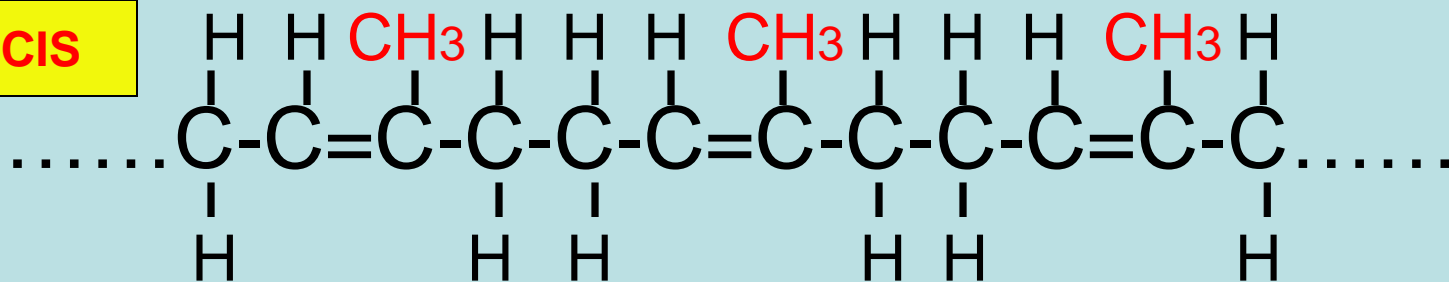
META



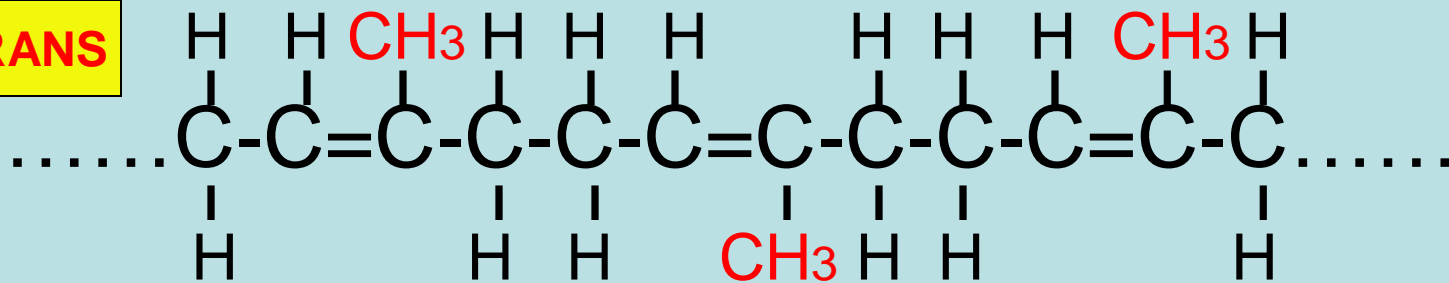
PARA

Isómeros

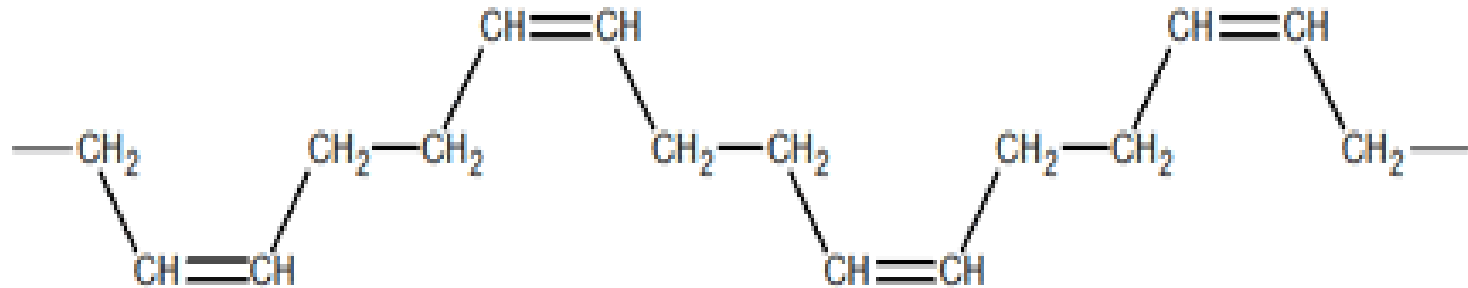
CIS



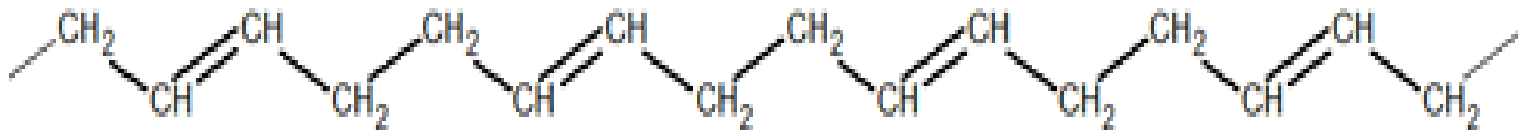
TRANS



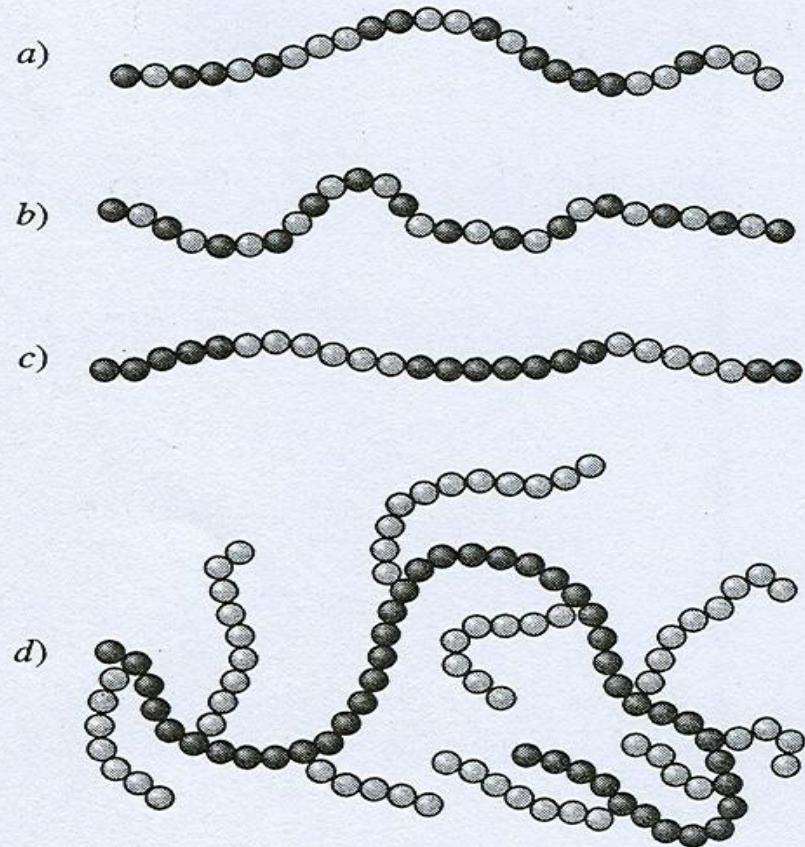
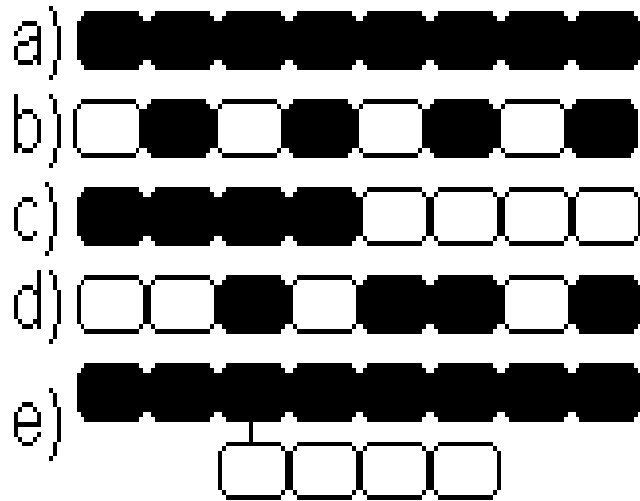
La primera se llama configuración *CIS* y así es el hule de la hevea y el de Guayule que se caracterizan por su flexibilidad y su elasticidad, la cual deben en parte a su estructura *CIS*, que es irregular y les impide cristalizar.



En cambio, la configuración *TRANS*, con gran regularidad estructural, está presente en el hule de gutapercha, que es cristalino, mucho menos elástico, con alta dureza, por lo que se lo emplea como cubierta de pelotas de golf.

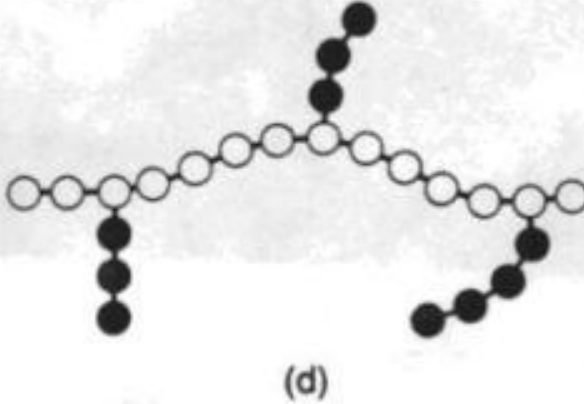
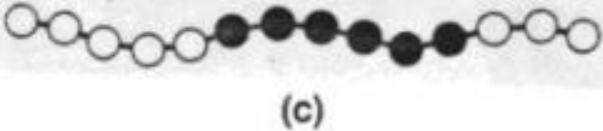
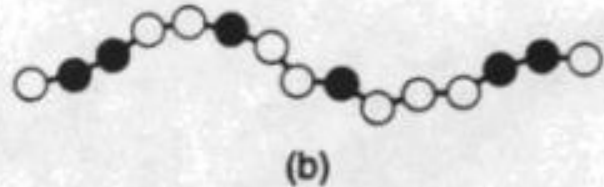
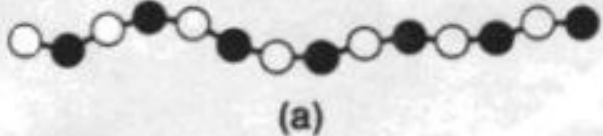
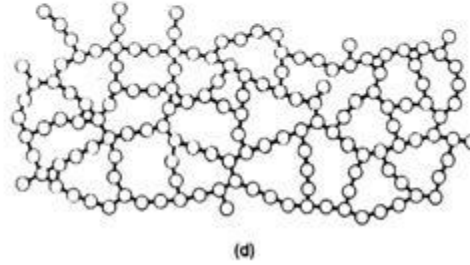
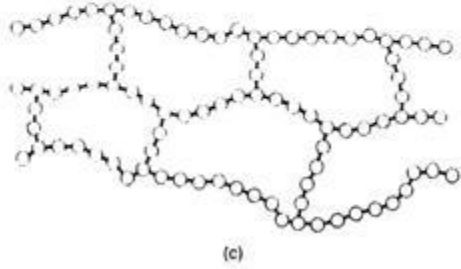
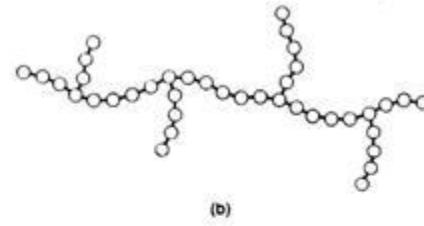
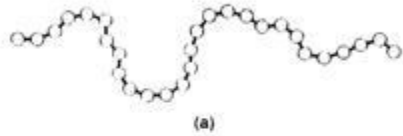


Copolímeros



(Según W. G. Moffatt, G. W. Pearsall y J. Wulff, «The Structure and Properties of Materials», vol. I, «Structure», Wiley, 1965, p. 108.)

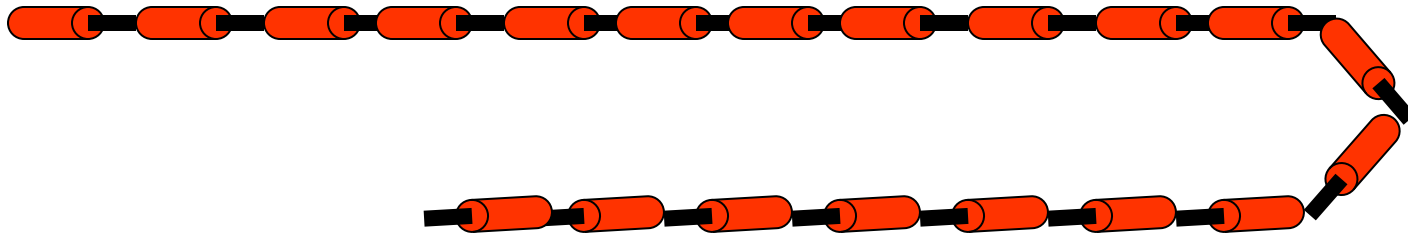
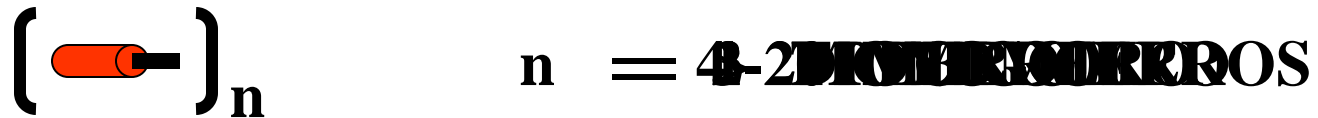
FIGURA 7.8. Ordenaciones de copolímeros. a) Copolímero en donde las distintas unidades están distribuidas al azar a lo largo de la cadena; b) las unidades están alternadas regularmente; c) un copolímero en bloque, y d) un copolímero de injerto.



POLIMEROS:

¿Que es un polímero?

Es una macromolécula formada por la unión de moléculas de menor tamaño que se conocen como monómeros.

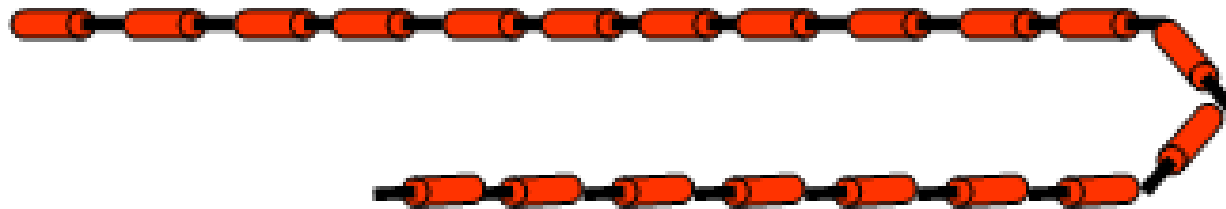




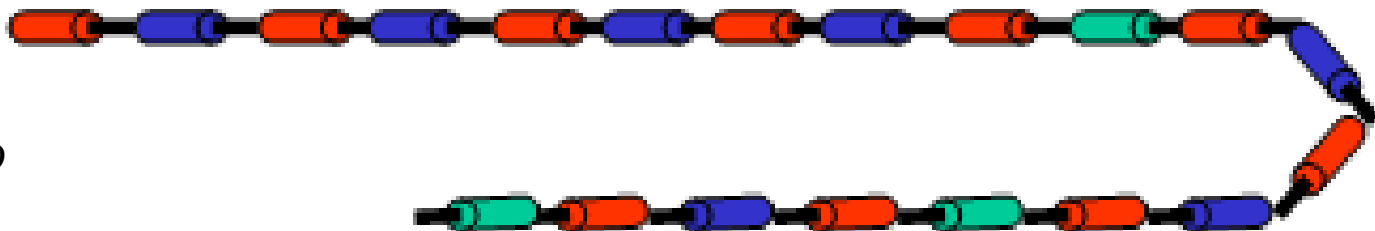
$n = 4 - 20$ OLIGOMEROS



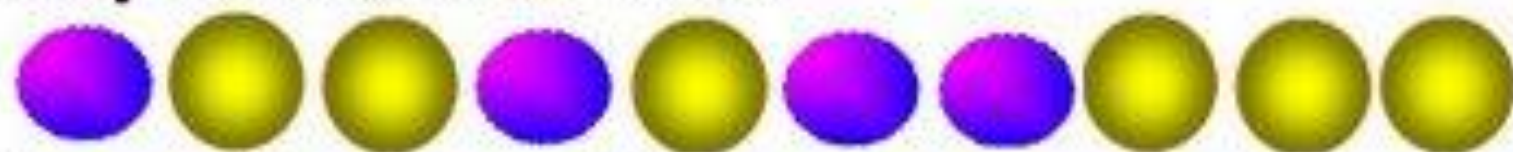
Homopolímero



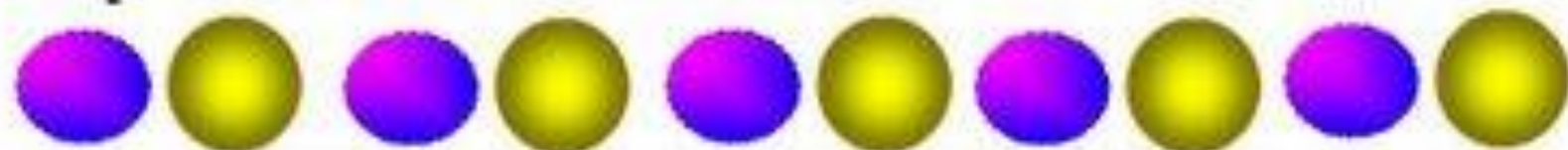
Copolímero



Copolímero al azar



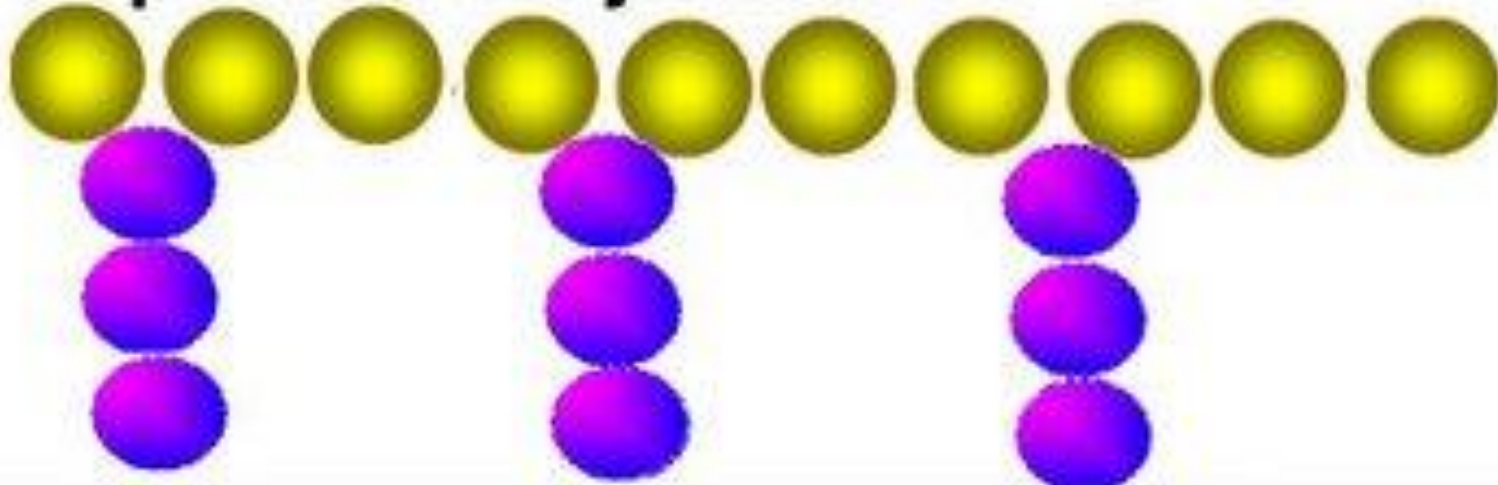
Copolímero alternado



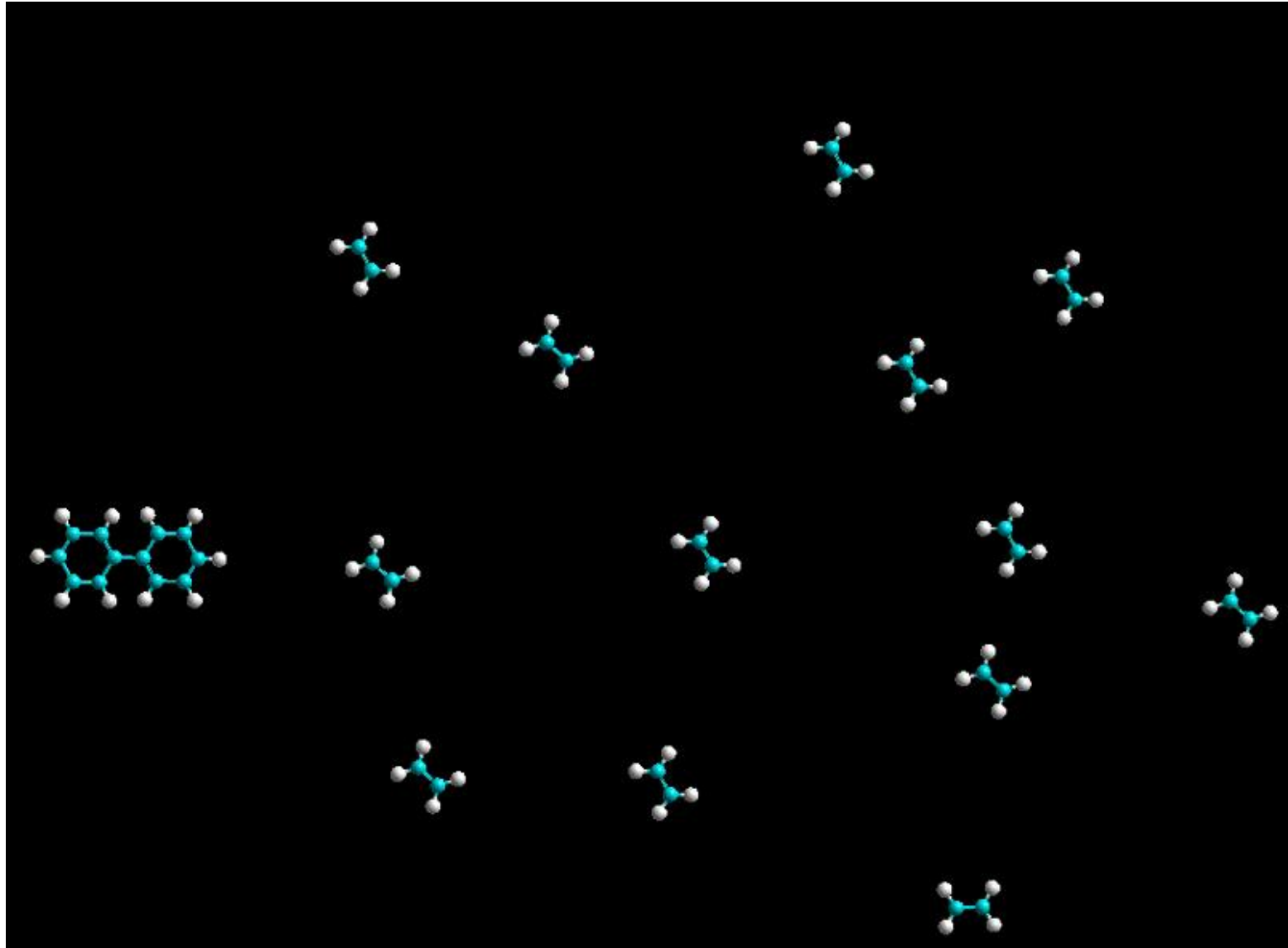
Copolímero en bloque



Copolímero injertado



POLIMERIZACIÓN



Copolímero

policloruro de vinilo-poliacetato de vinilo

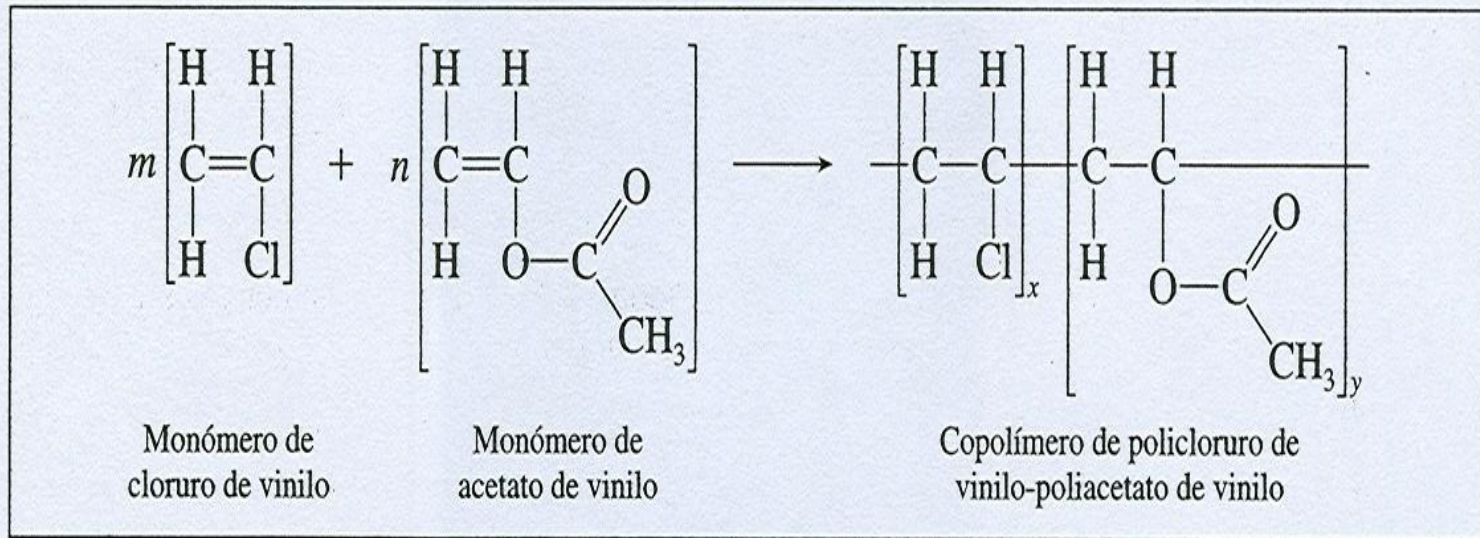
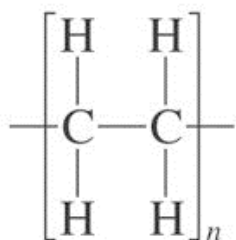
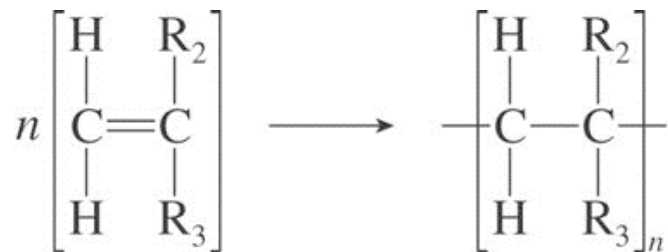
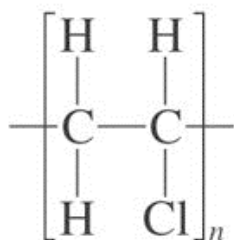


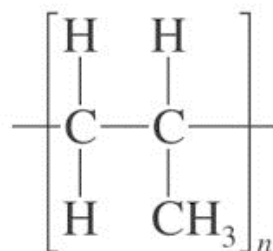
FIGURA 7.9. Reacción de polimerización generalizada de monómeros de cloruro de vinilo y acetato de vinilo para producir un copolímero de policloruro de vinilo-poliacetato de vinilo.



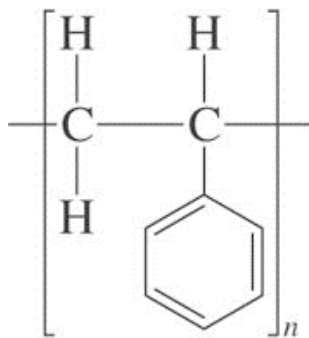
Polietileno
 pf: 110-137°C
 (230-278°F)



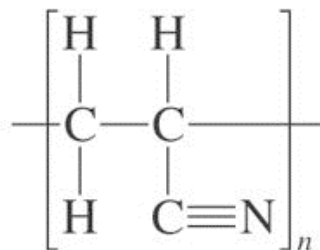
Policloruro de vinilo
 pf: ~204°C
 (~400°F)



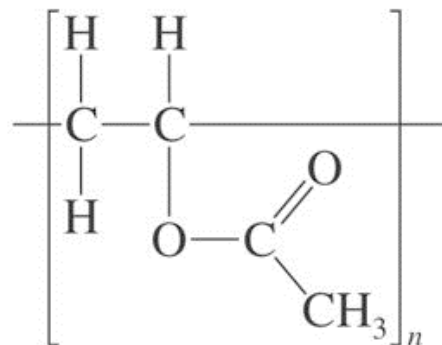
Polipropileno
 pf: 165-177°C
 (330-350°F)



Poliestireno
 pf: 150-243°C
 (330-470°F)

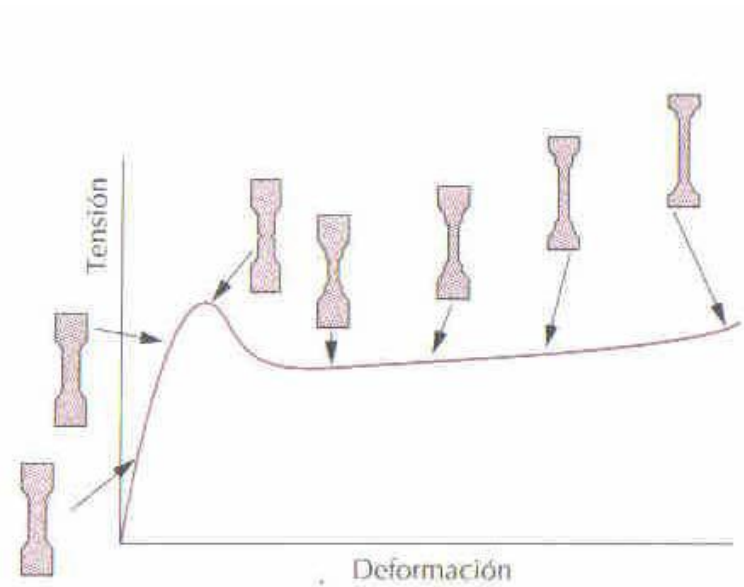
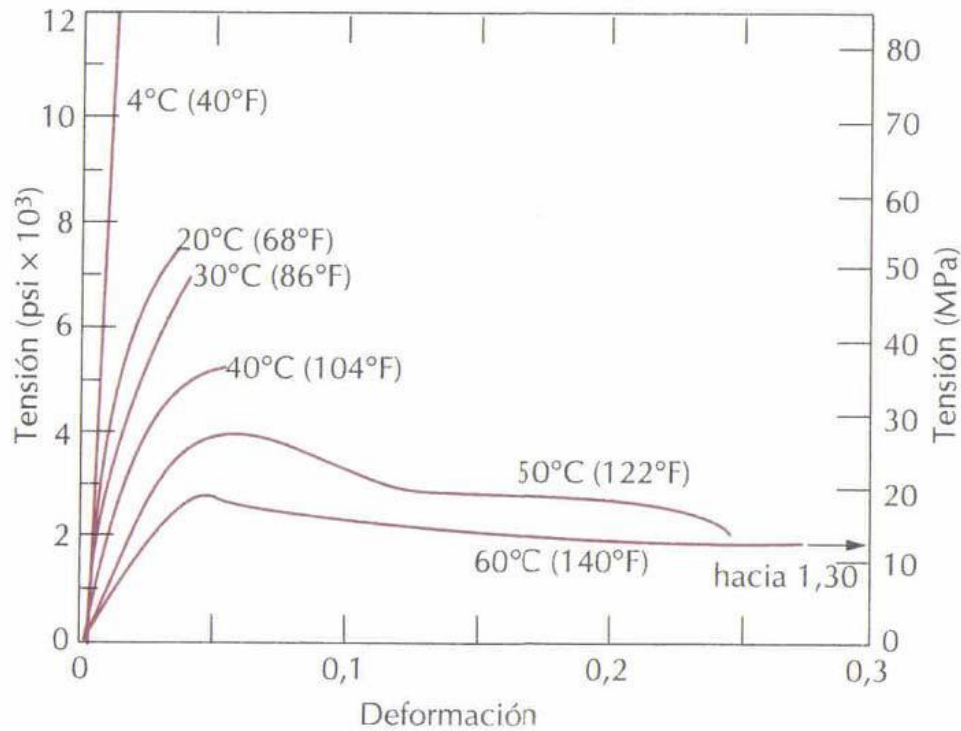


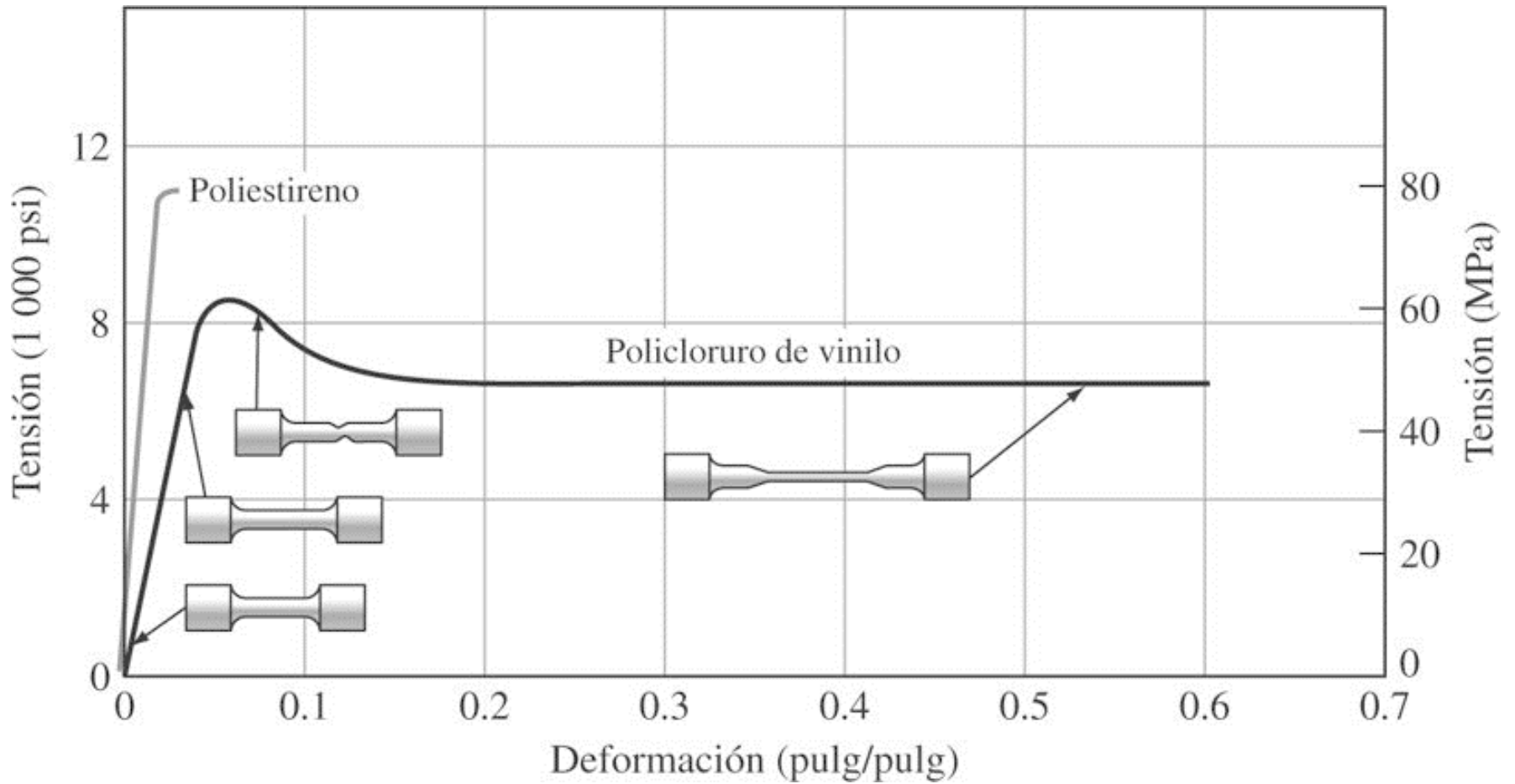
Poliacrilonitrilo
 (no se funde)



Poliacetato de vinilo
 pf: 177°C (350°F)

Tensión-deformación de polímeros





Datos de tensión-deformación para el termoplástico amorfo policloruro de vinilo (PVC) y el poliestireno (PS). Los dibujos presentan diversos modos de deformación de la muestra en varios puntos sobre la curva de esfuerzo-deformación.

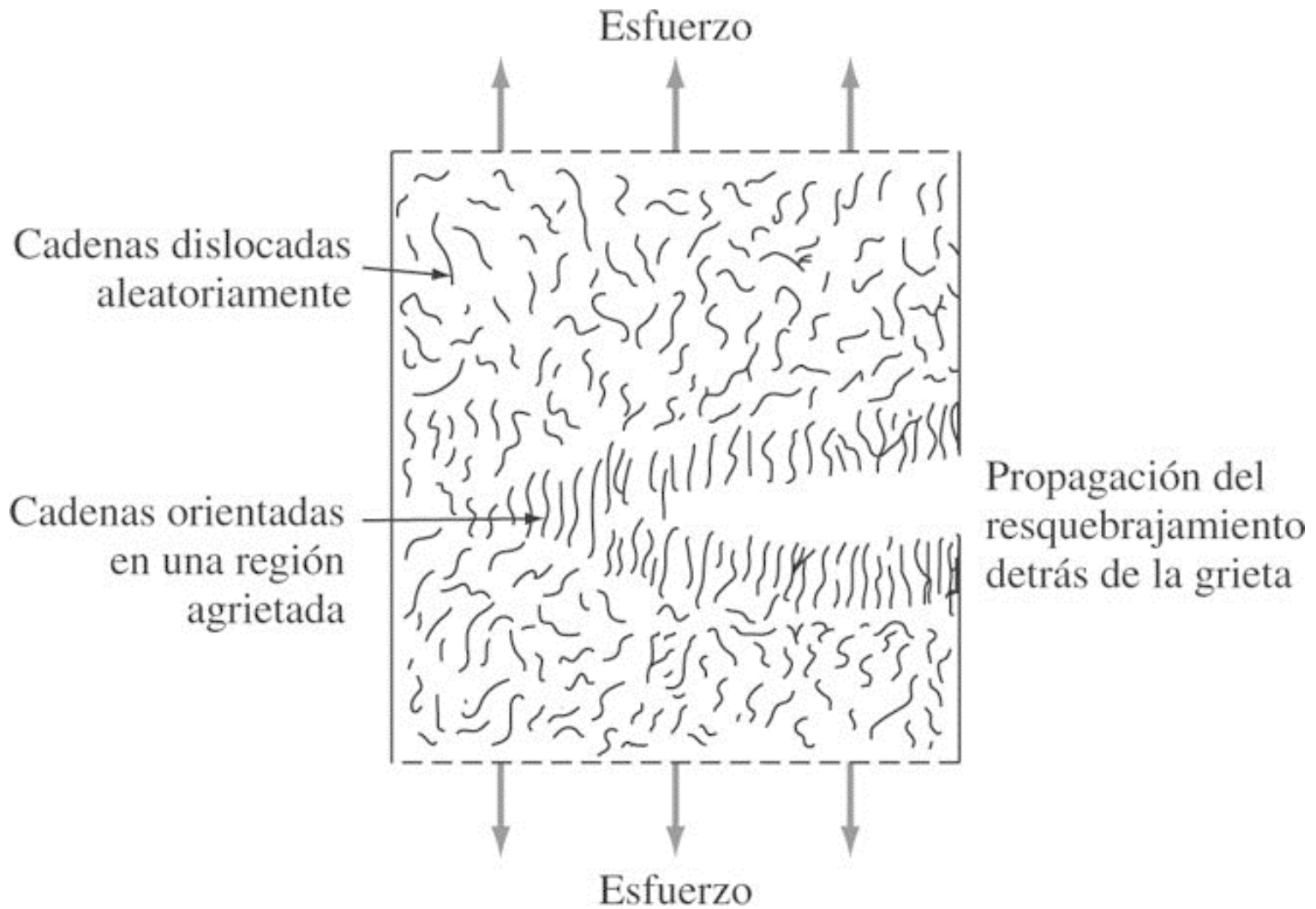
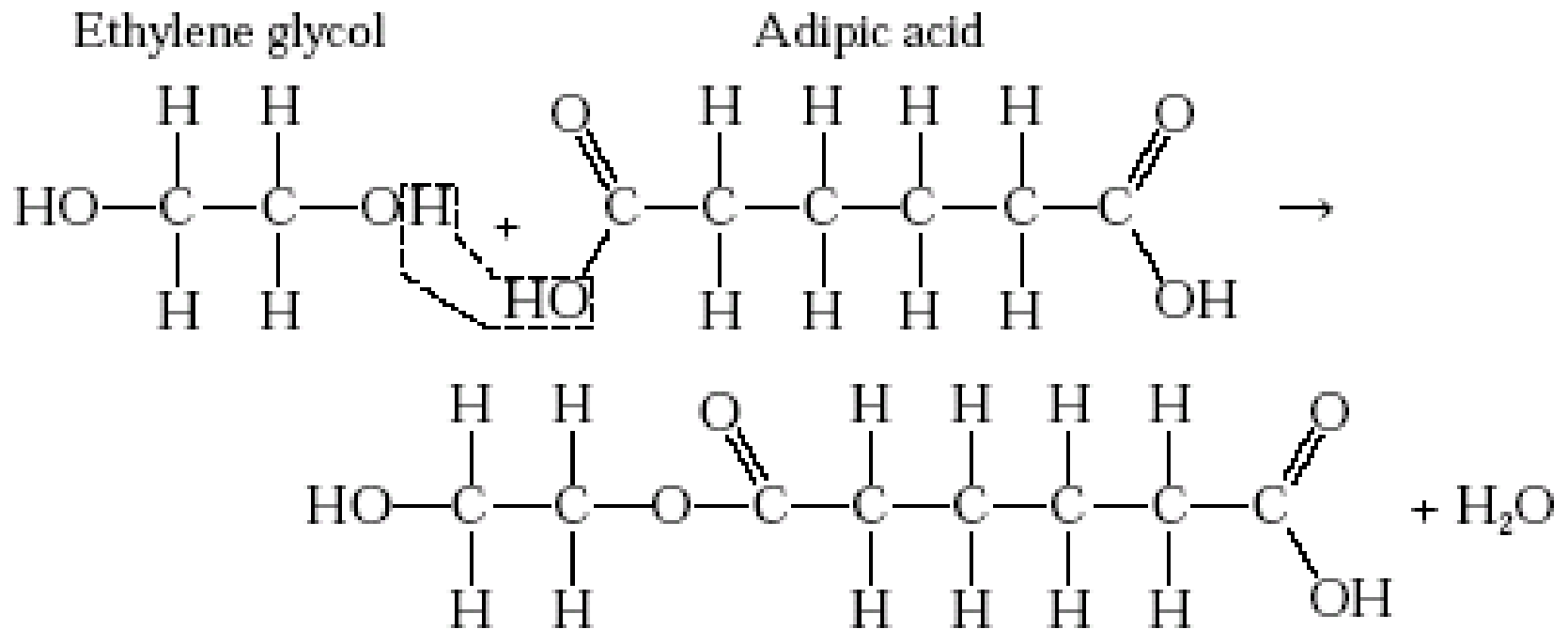
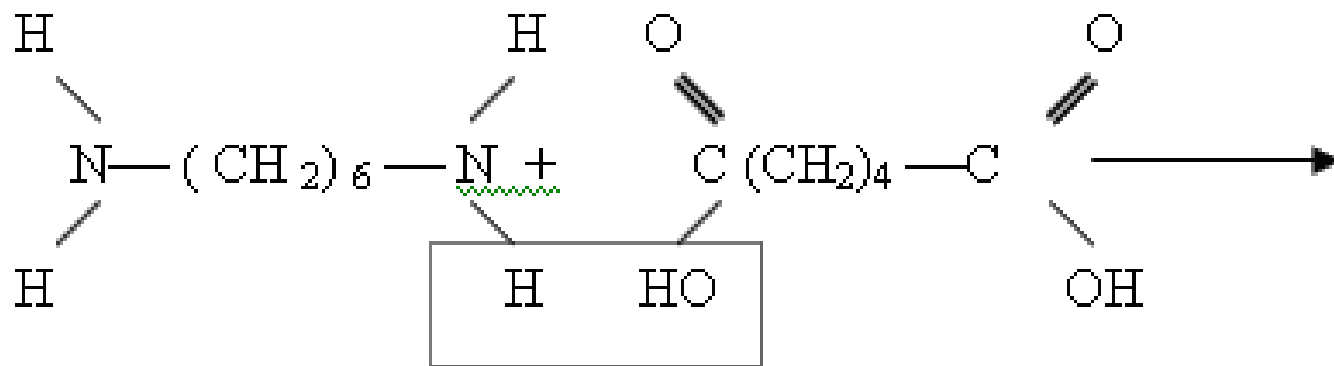


Ilustración esquemática de la estructura de un agrietamiento cercano al extremo de una fisura en un termoplástico vidriado.

COPOLÍMEROS

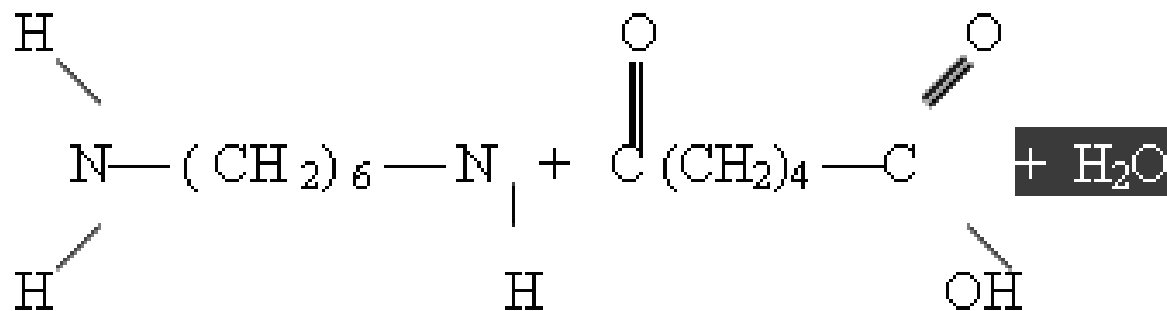
Polimerización por condensación





Hexametilendiamina

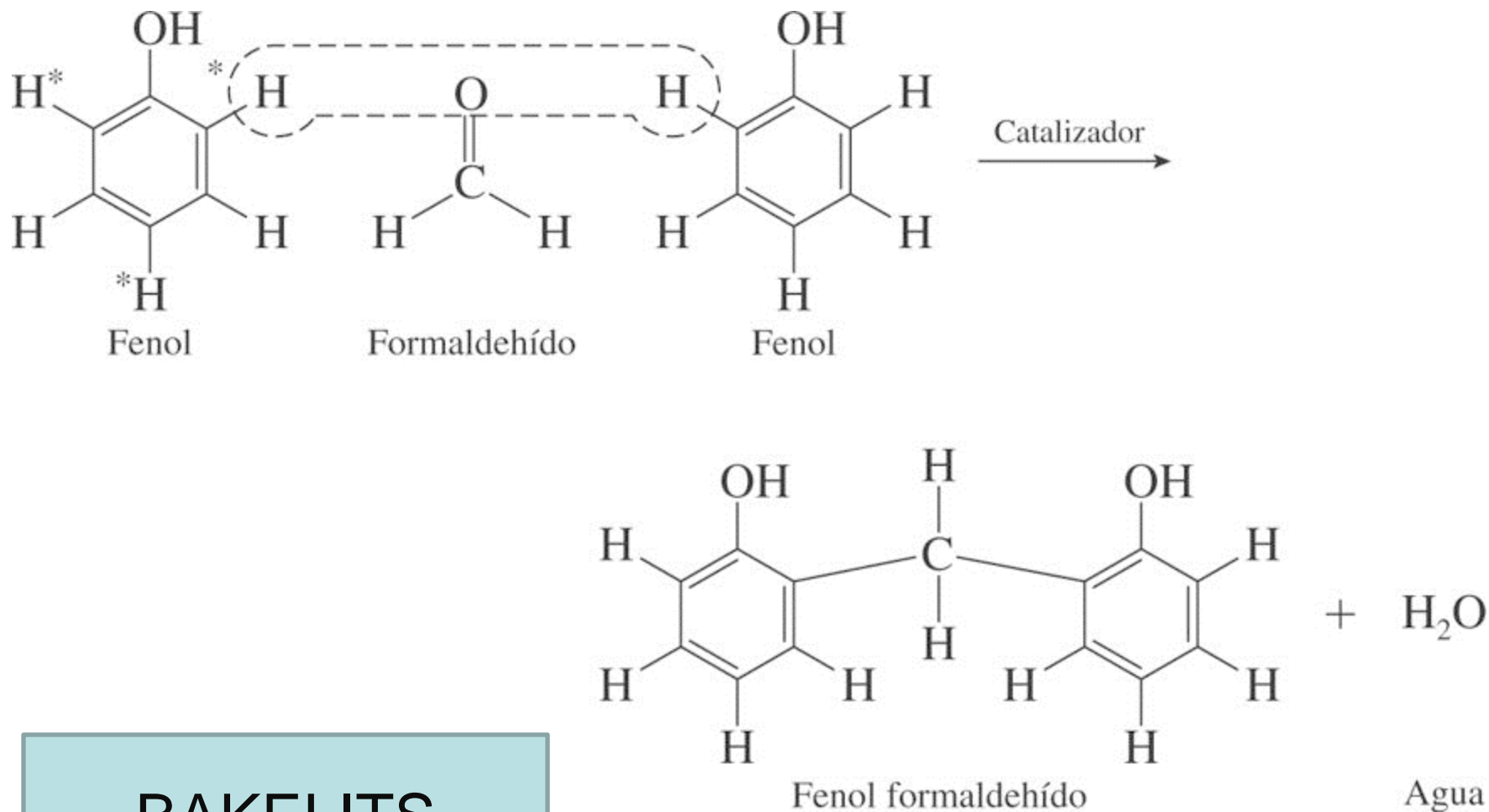
Ácido adípico



Hexametilendiamina

Agua

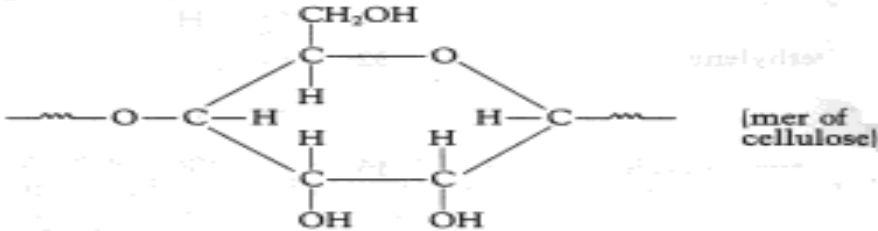
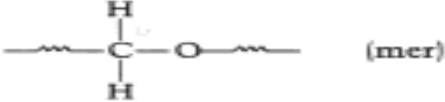

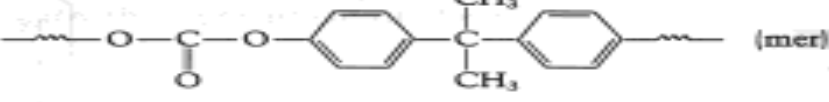
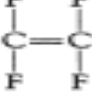
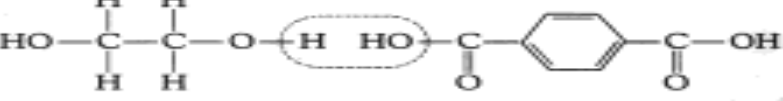
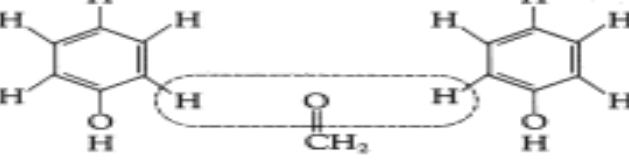
La correspondiente poliamida es el nylon 6,6



BAKELITS

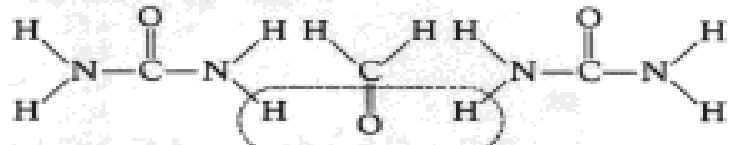
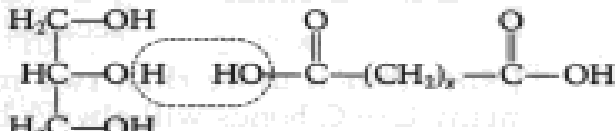
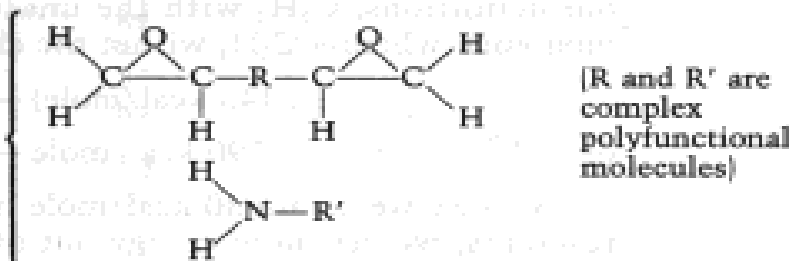
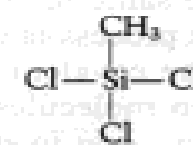
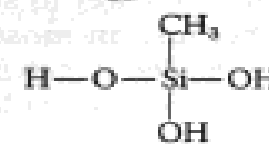
COPOLÍMEROS

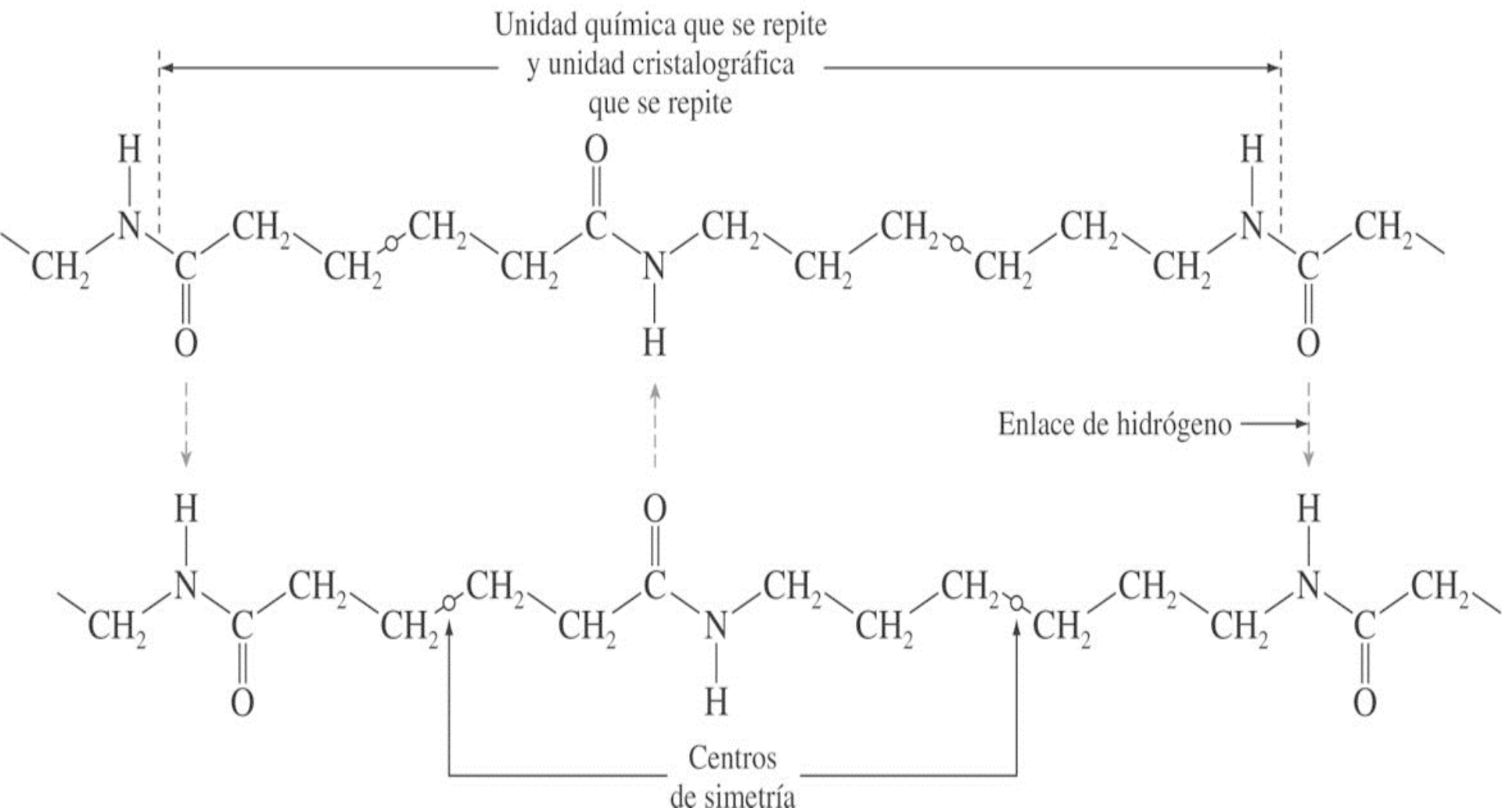
TABLE 13.1 Summary of Important Polymers (Continued)

Group I. Thermoplastics (Continued)		
Polymer	Percentage of Market	Monomer(s) Used
Cellulosics	<1	 <p>(mer of cellulose)</p>
Acetals	<1	 <p>(mer)</p>
Nylons	1	
Polycarbonates	<1	 <p>(mer)</p>
Fluoroplastics [example: polytetrafluoroethylene]	<1	
Polyester, thermoplastic type [example: polyethylene- terephthalate (dacron)]	2	
Group II. Thermosetting Polymers		
Phenolics (example: phenol formaldehyde)	6	

COPOLÍMEROS

TABLE 13.1 Summary of Important Polymers (Continued)

Polymer	Percentage of Market	Monomer(s) Used
<i>Group II. Thermosetting Polymers (Continued)</i>		
Amino resins (example: urea formaldehyde)	4	
Polyesters, thermoset type	3	
Epoxies	1	 <p style="text-align: right;">[R and R' are complex polyfunctional molecules]</p>
Polyurethane, also thermoplastic	4	$\text{OCN}-\text{R}-\text{NCO} + \text{HO}-\text{R}'-\text{OH}$ <p style="text-align: center;">(diisocyanate)</p> <p style="text-align: right;">[R and R' are complex polyfunctional molecules]</p>
Silicones	1	<div style="display: flex; align-items: center;"> <div style="font-size: 4em; margin-right: 10px;">{</div> <div style="margin-right: 20px;">  <p style="margin-left: 20px;">Trichlorosilane</p> </div> <div style="margin-right: 20px;">}</div> <div style="margin-right: 20px;">  <p style="margin-left: 20px;">Trihydroxy silane</p> </div> </div>



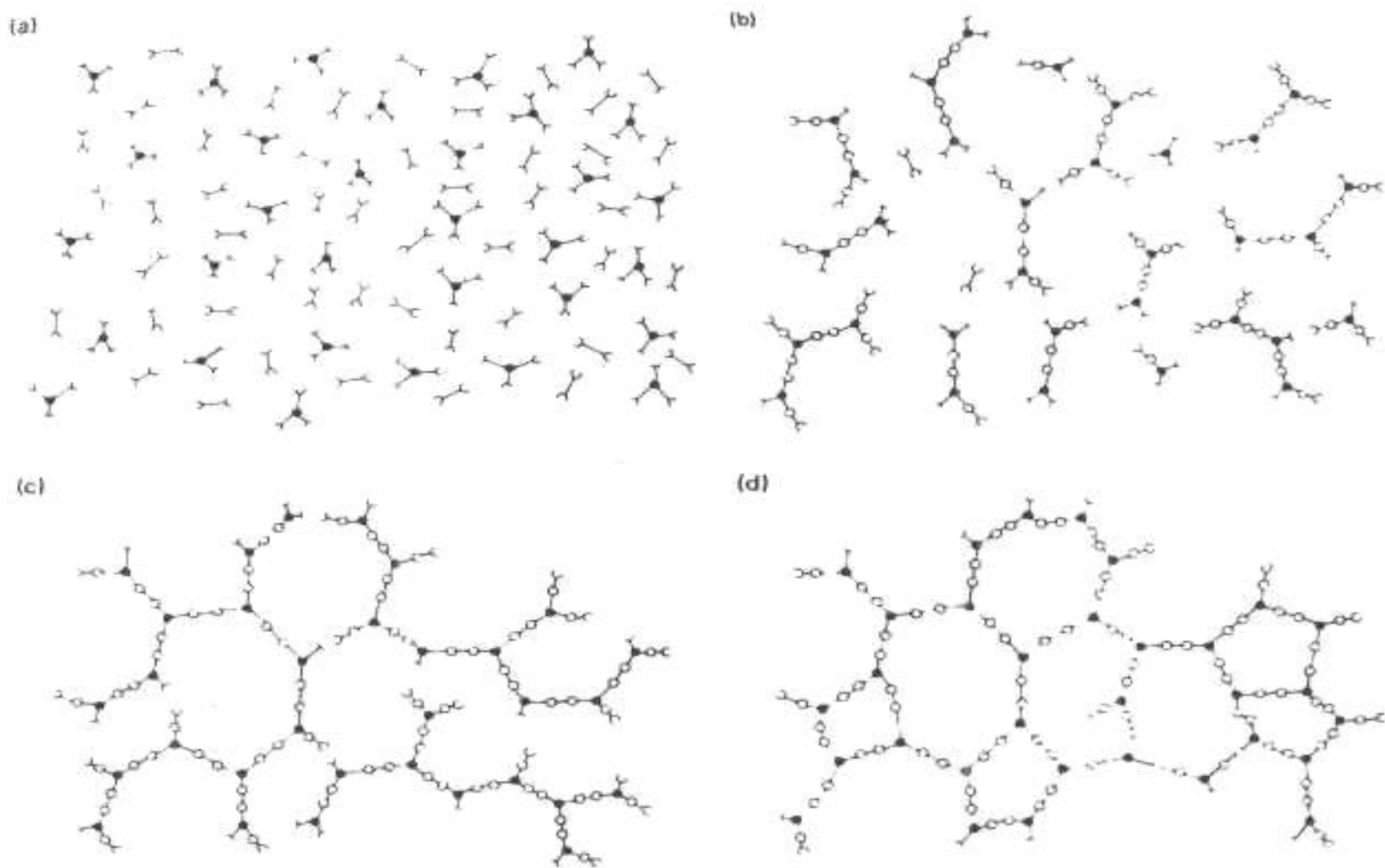
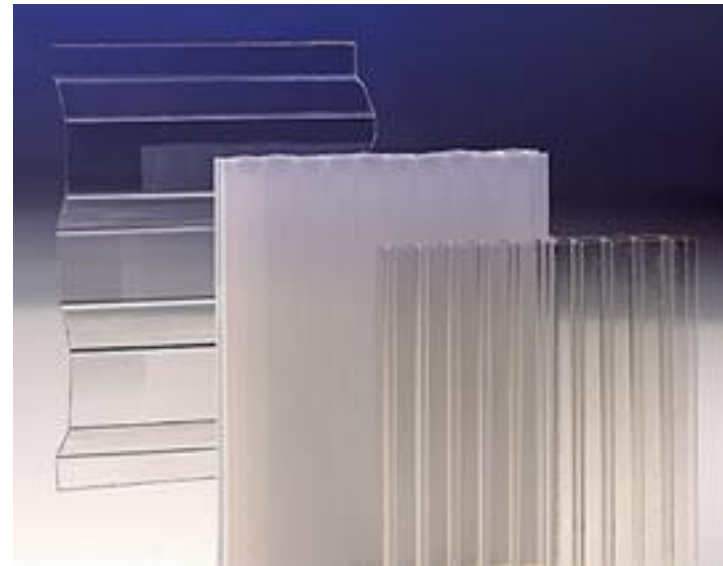
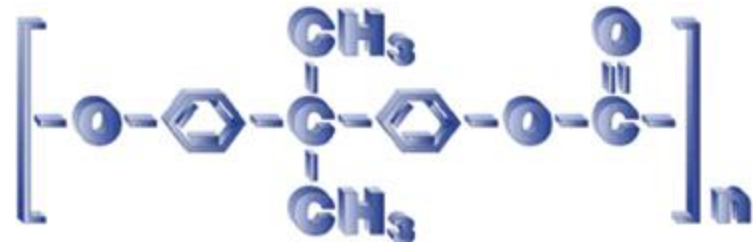
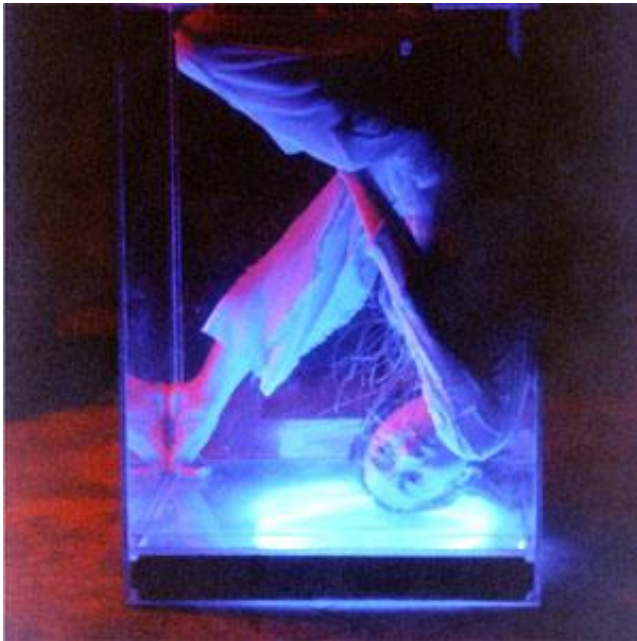
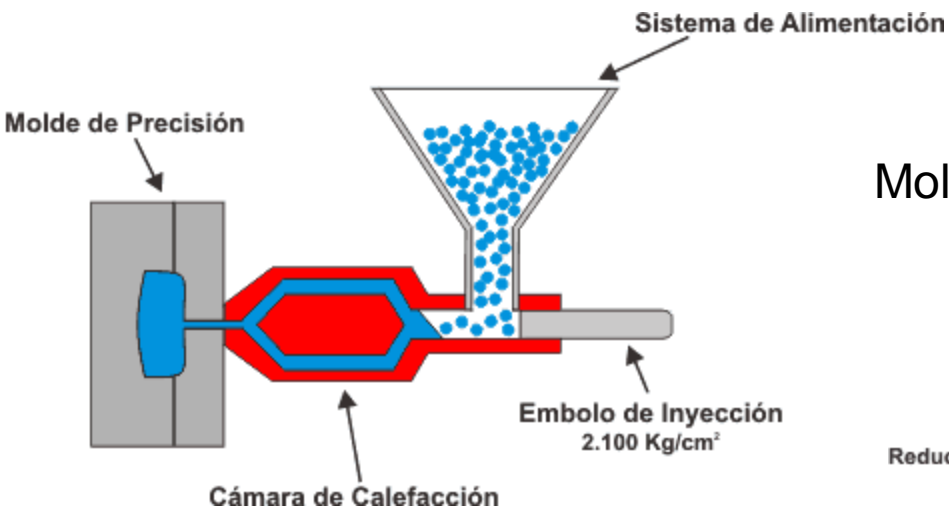


Figura 1.1 Representación en dos dimensiones del curado de un termoestable. (a) Monómeros. (b) Crecimiento lineal y ramificación por debajo del punto de gel. (c) Formación de un gel, reticulación incompleta. (d) Termoestable, curado total. (de Turi, 1981).

POLICARBONATO

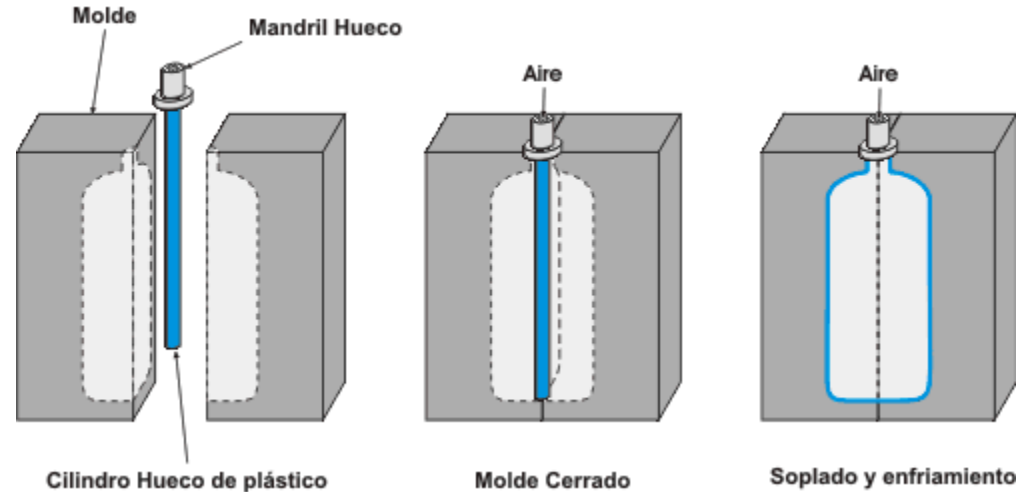
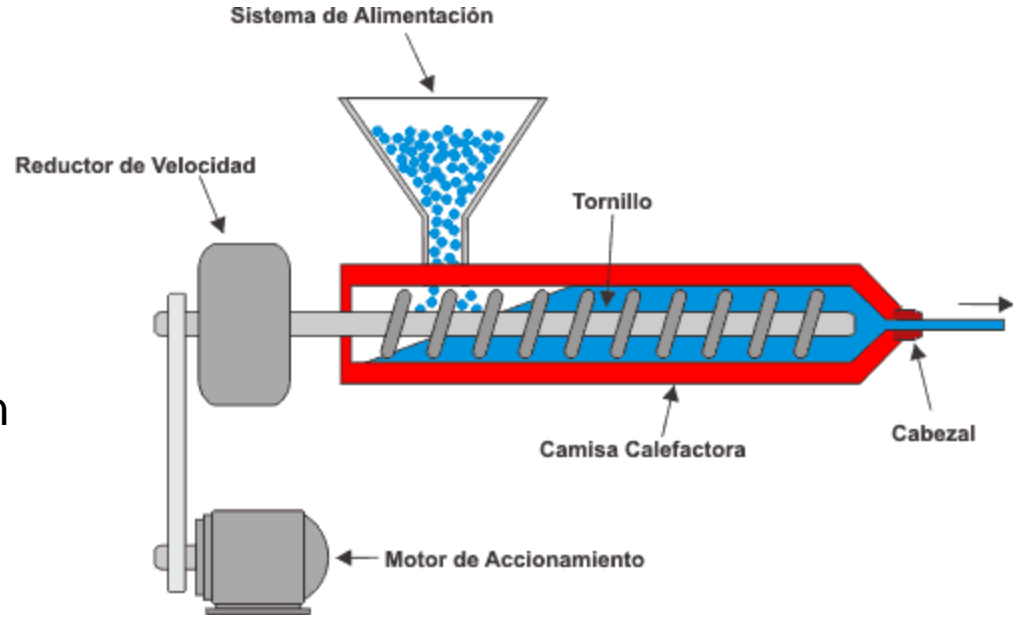
200 veces más resistente que el vidrio al impacto



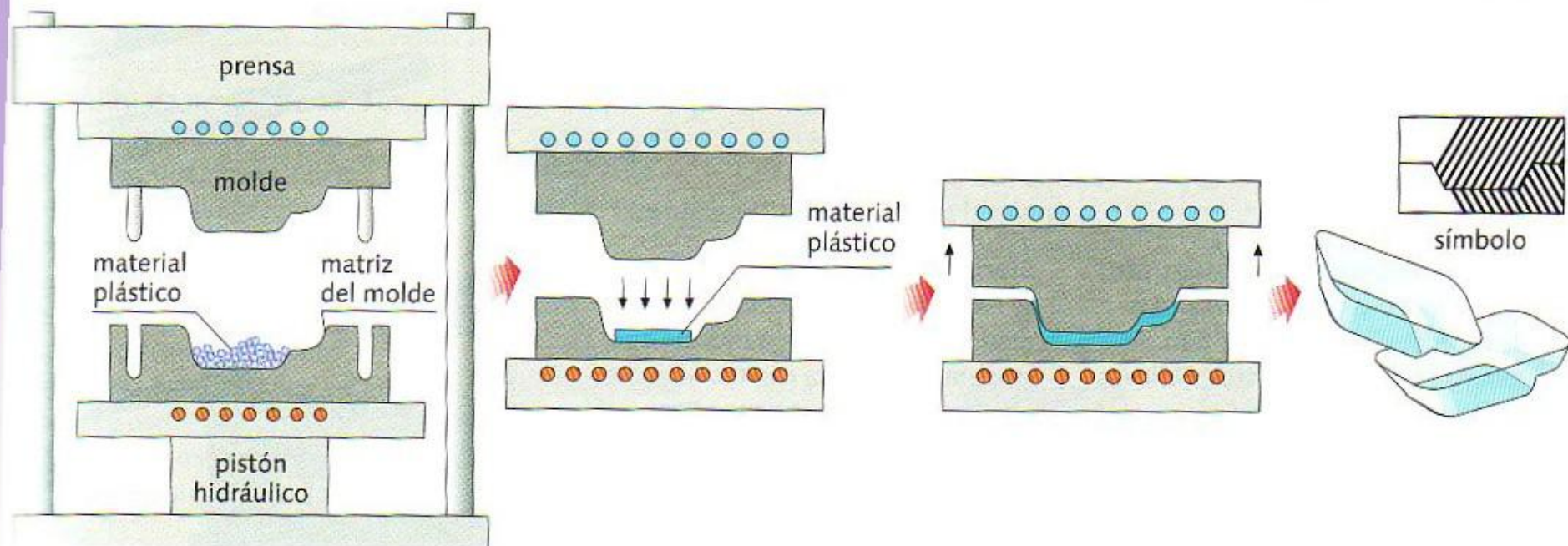
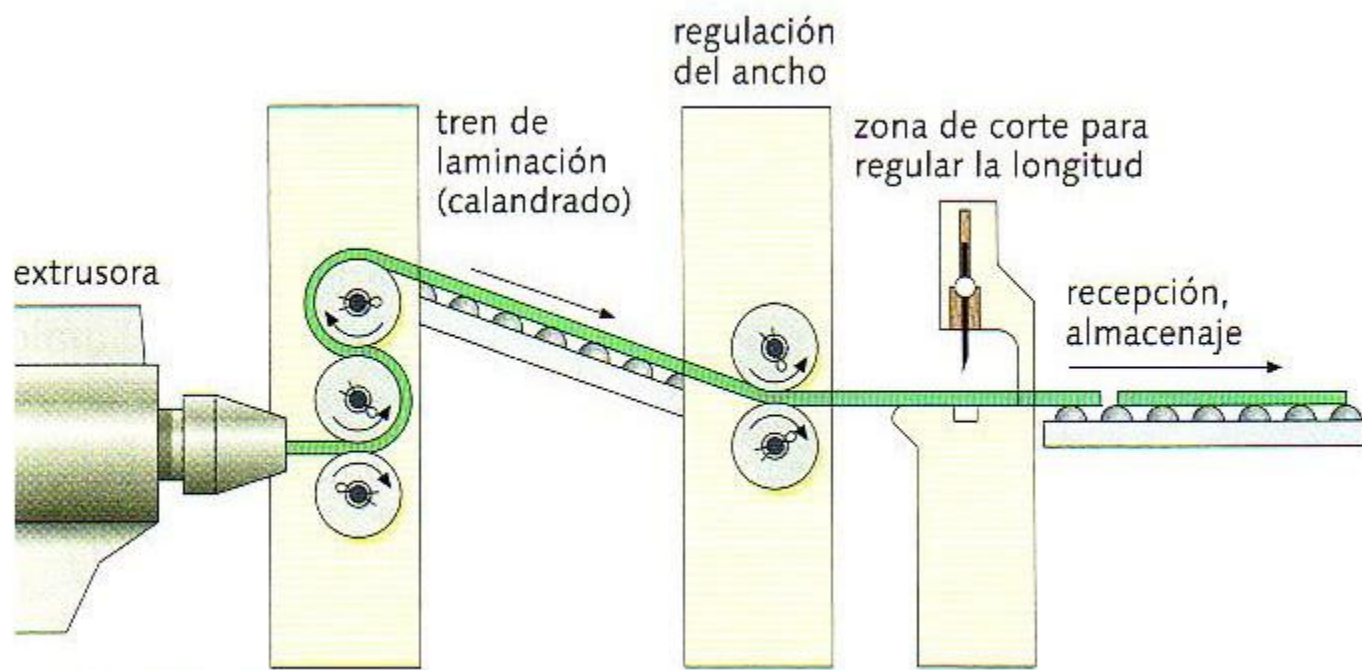


Moldeo por inyección

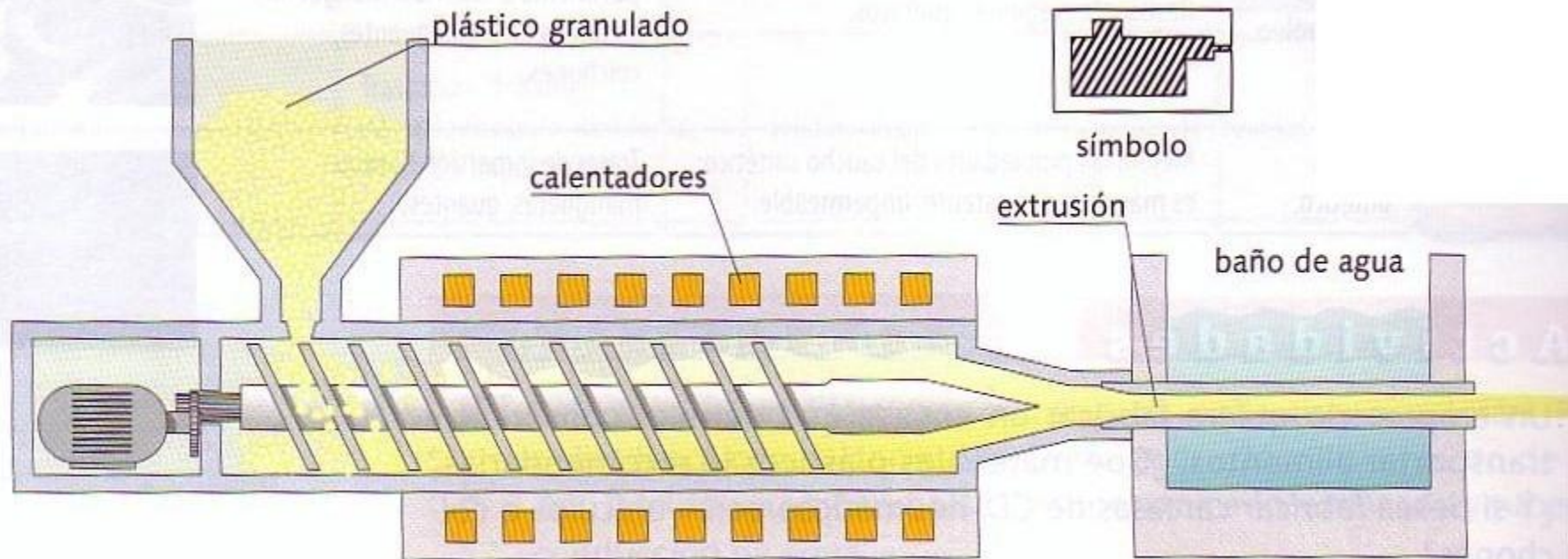
Moldeo por extrusión



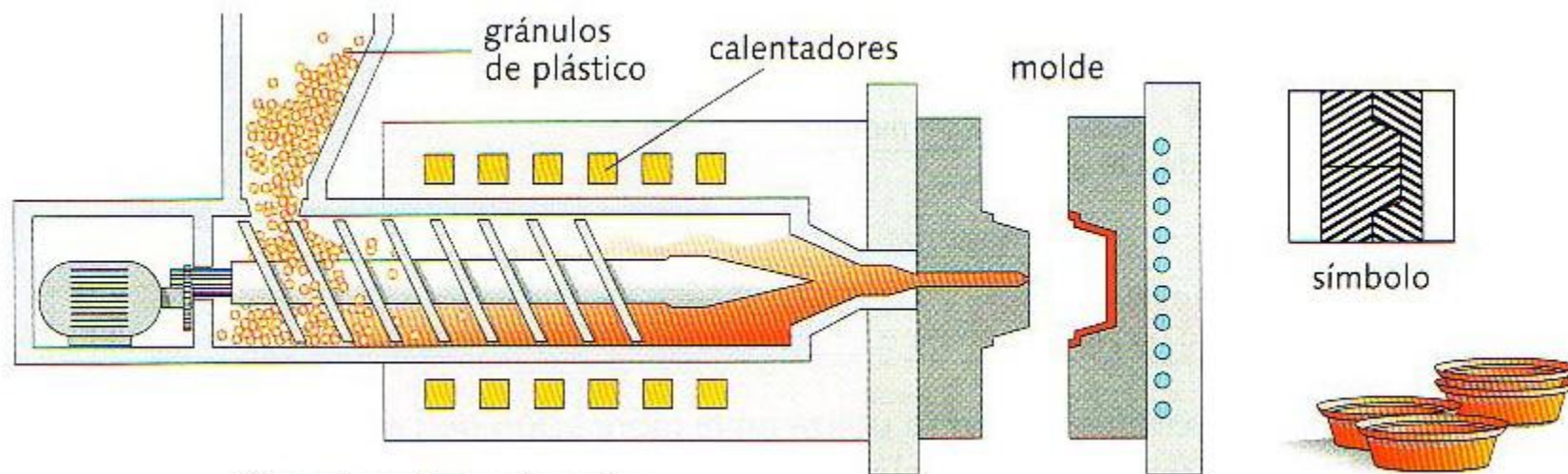
Moldeo por insuflación de aire



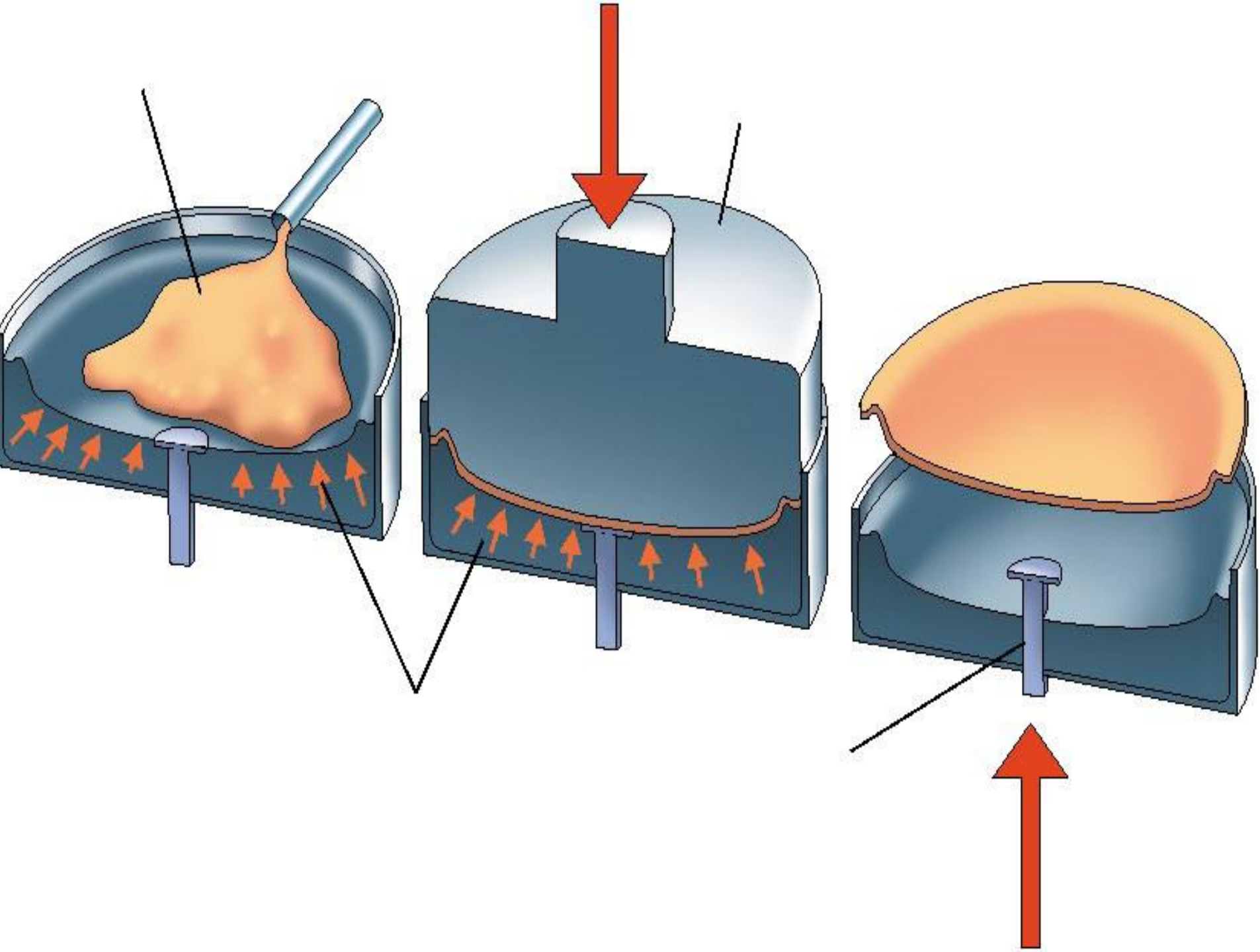
El moldeo por compresión se realiza en una máquina llamada prensa.

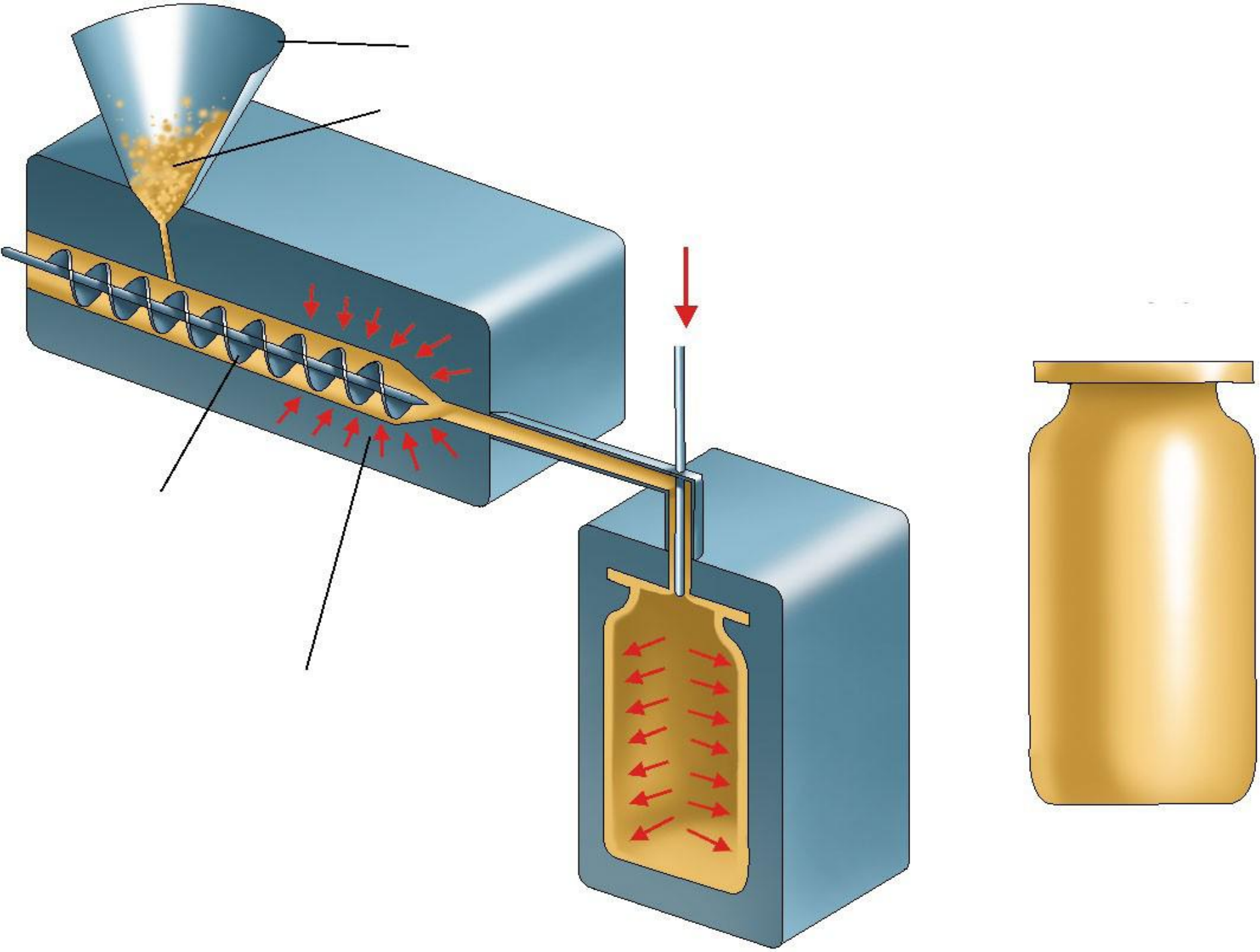


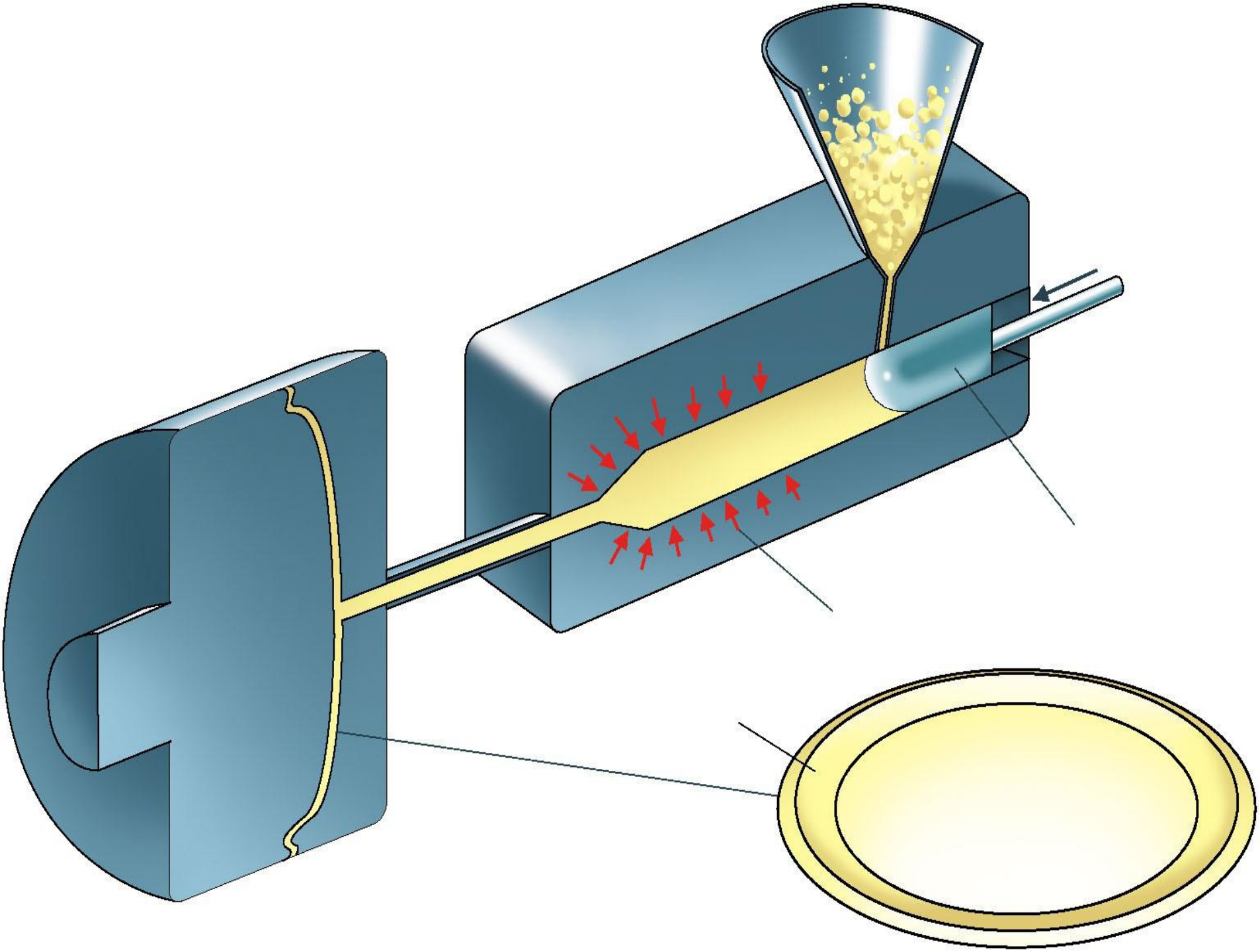
Técnica de extrusión.

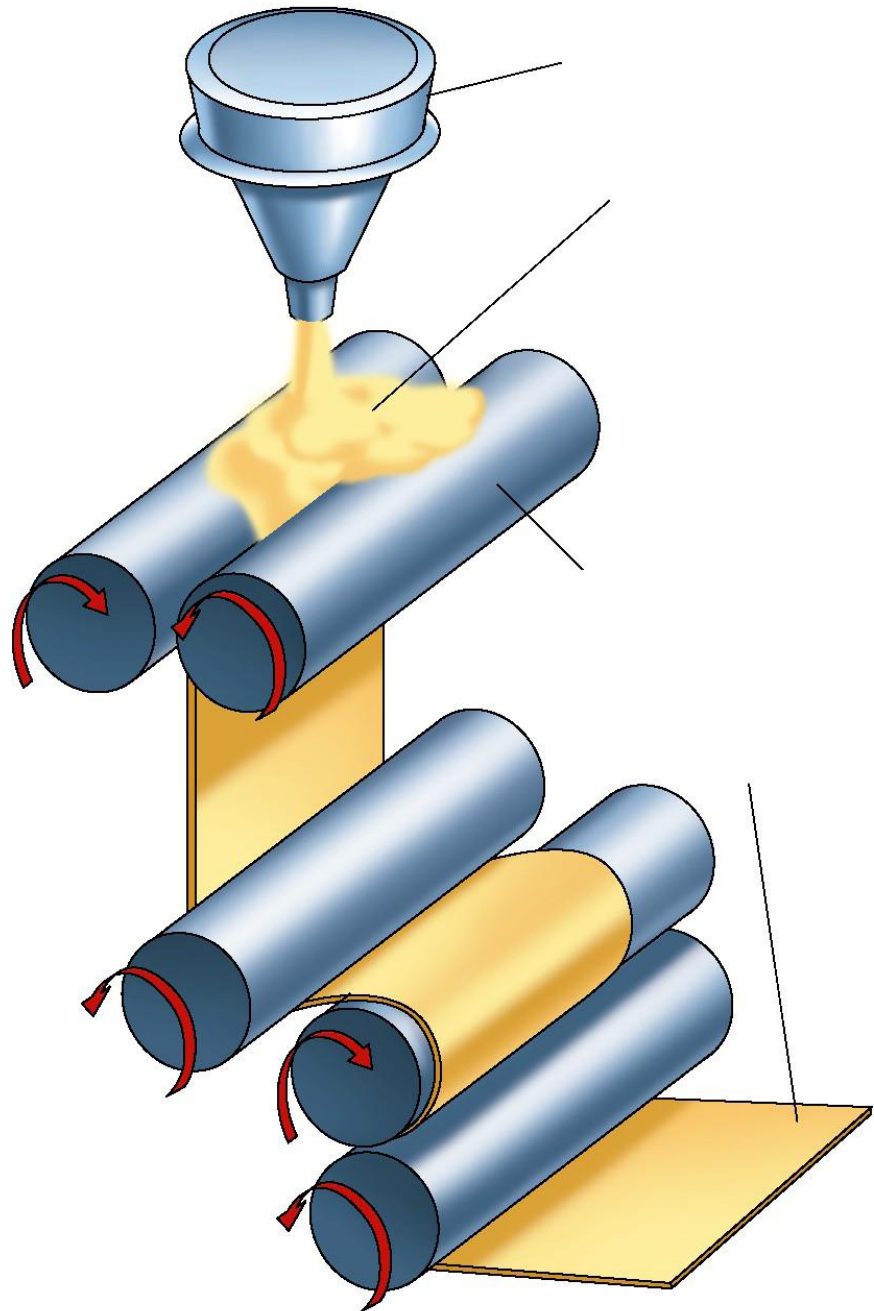


Técnica de moldeo por inyección.









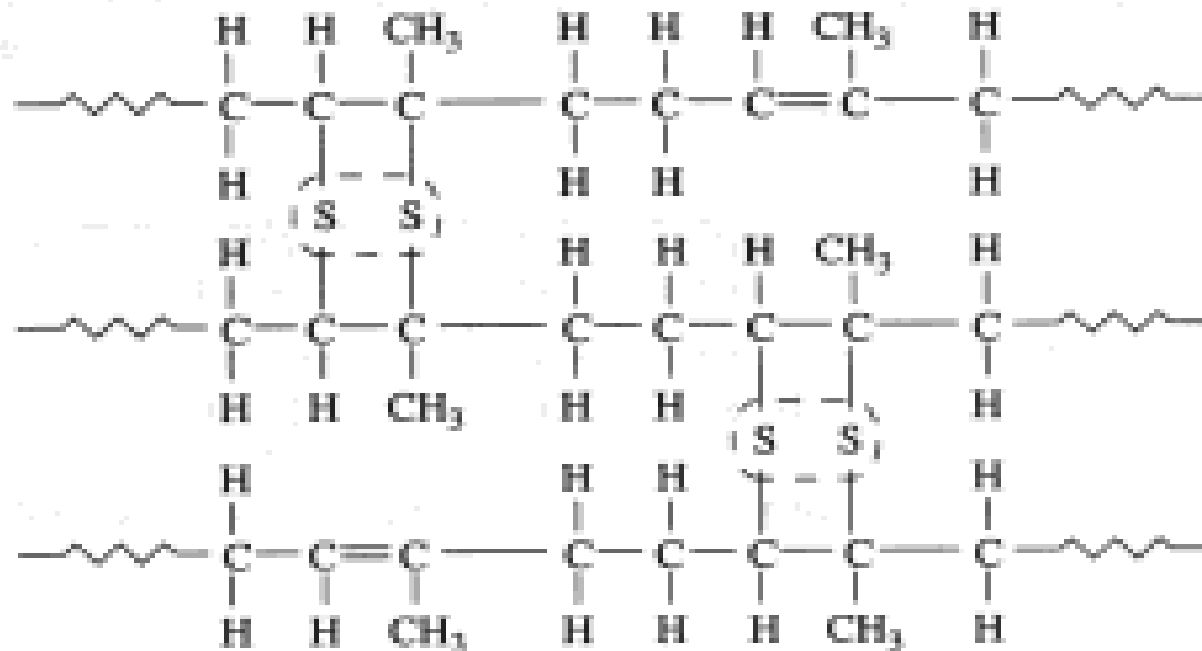
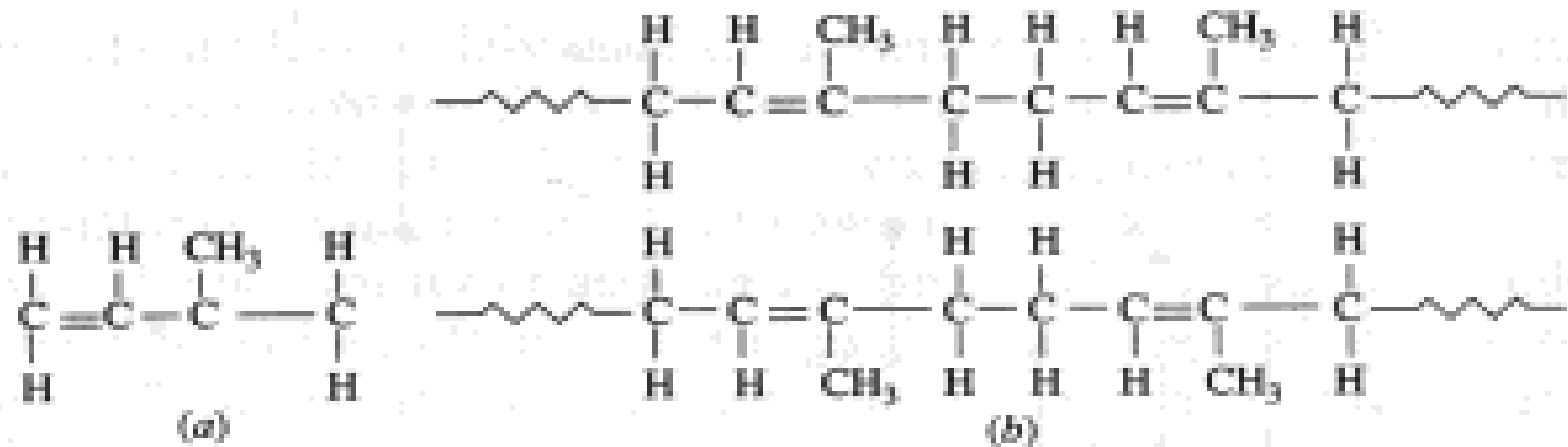
ELASTOMEROS

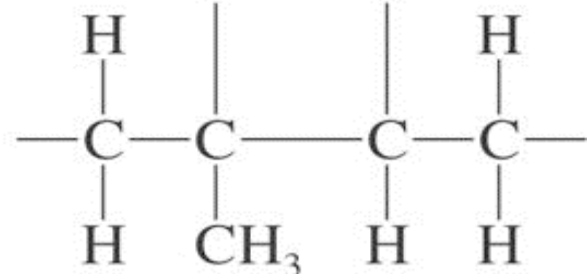
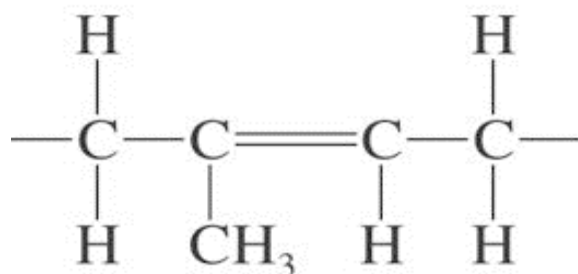
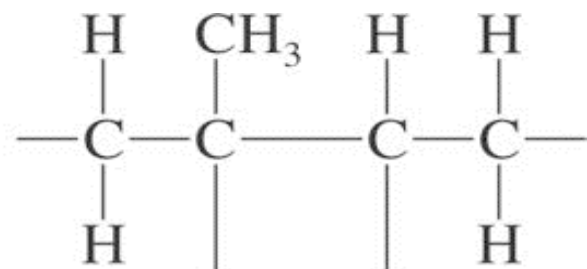
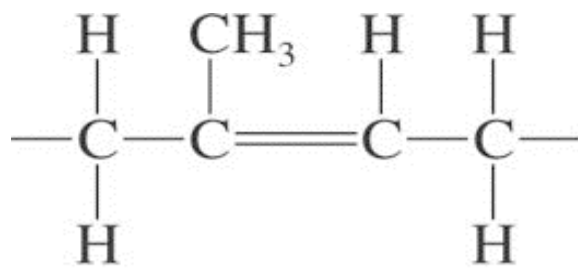
Vulcanizado de hule polisopreno (hule natural)



Extracción de látex de un [árbol](#); El látex se emplea en la producción de [goma](#)

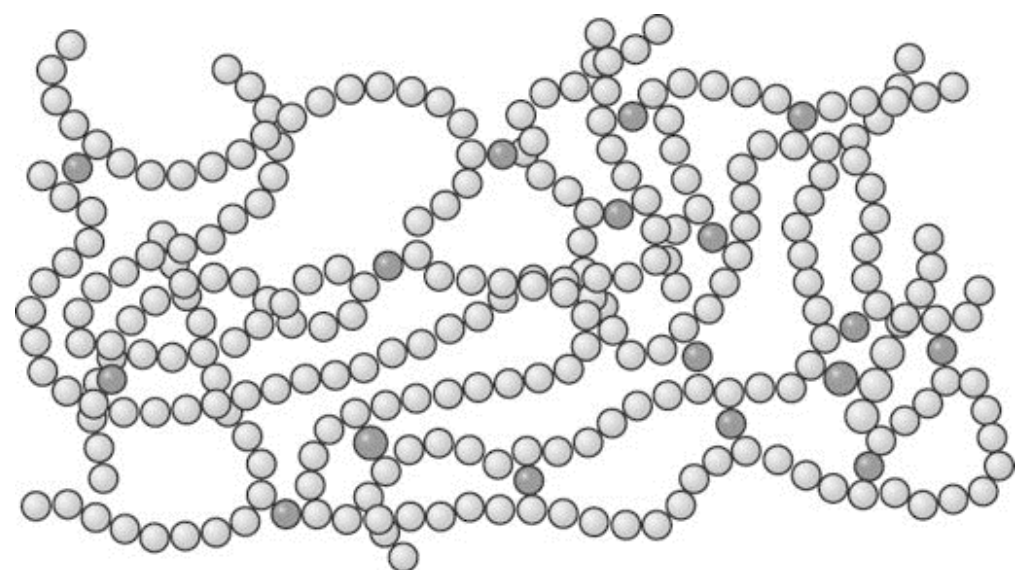
Vulcanizado de hule polisopreno (hule natural)



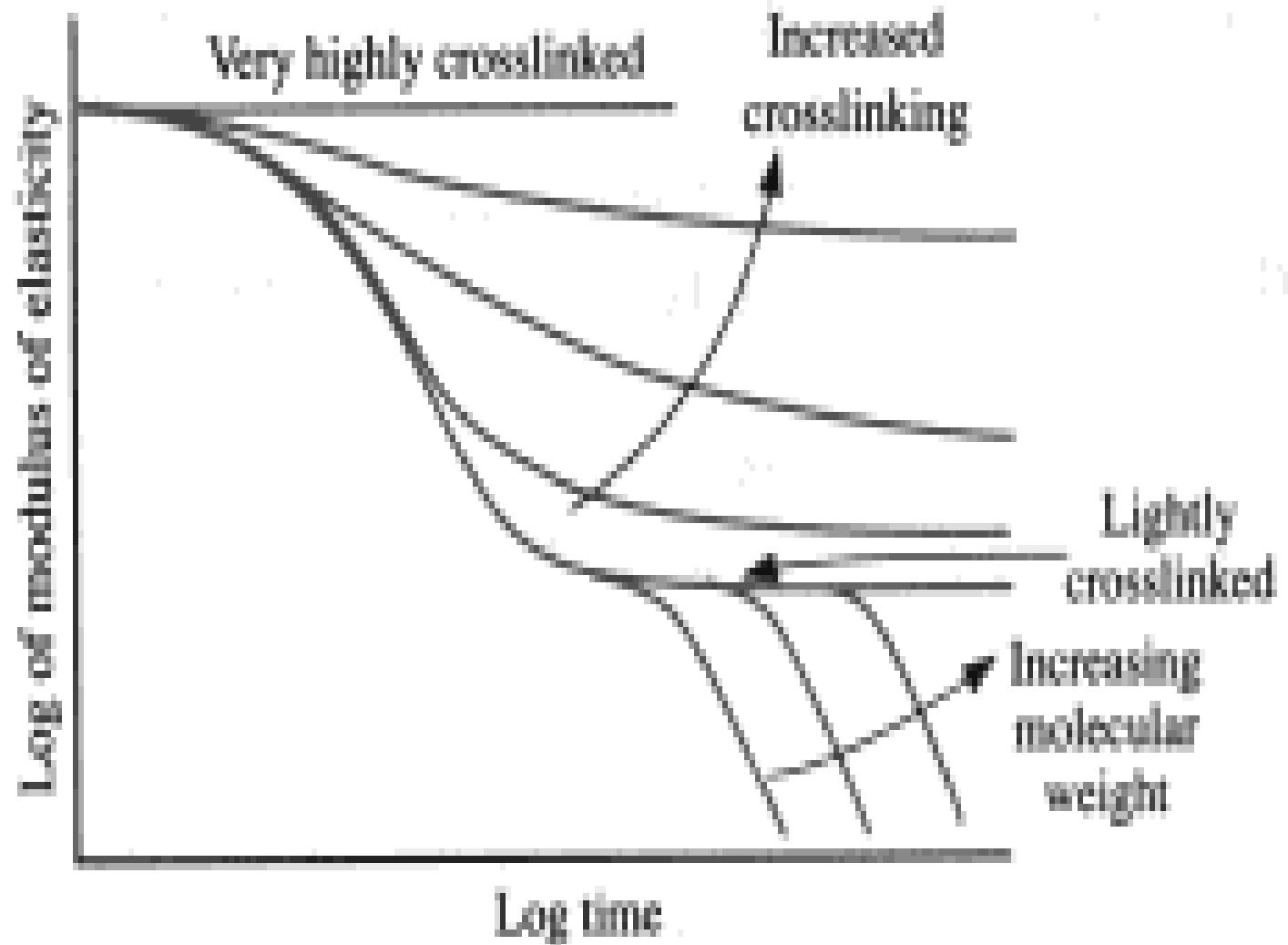


a)

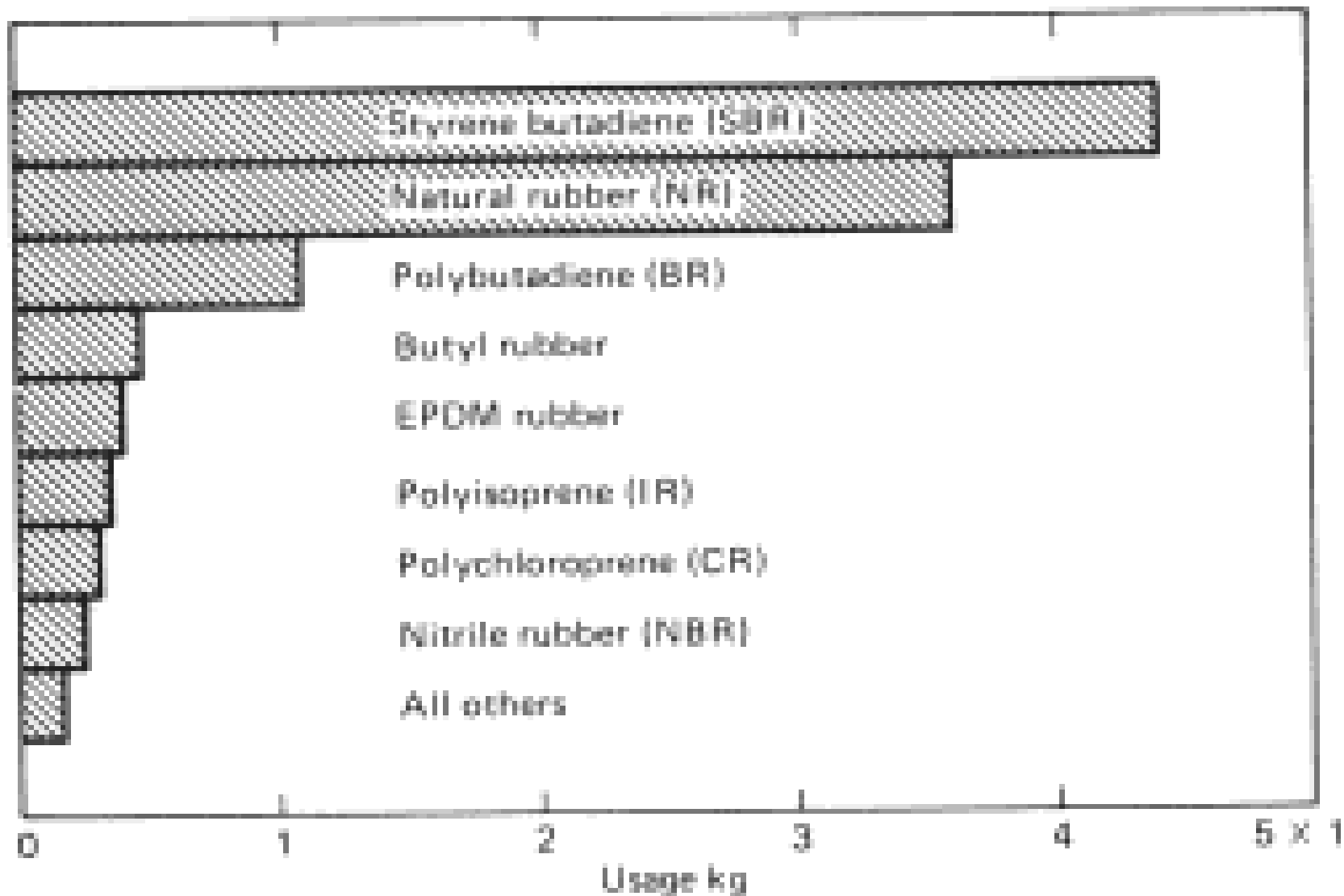
b)



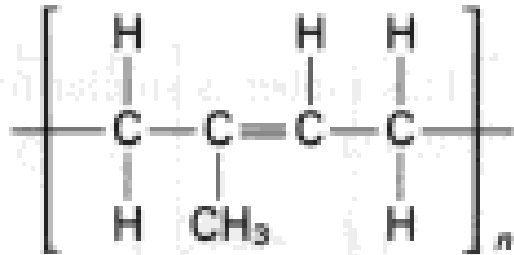
Efectos de elasticidad en modulos de hule



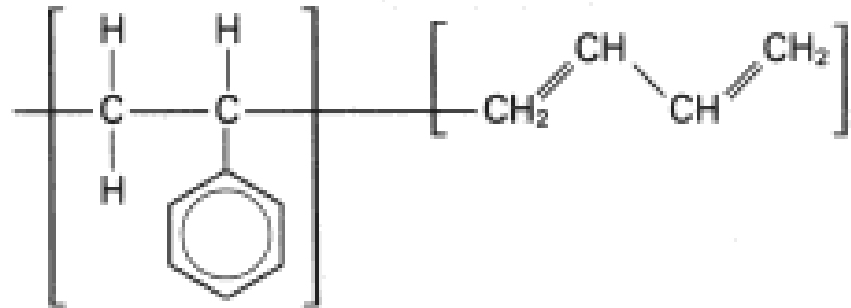
Consumo mundial de elastómeros



Monomeros elastoméricos más comunes



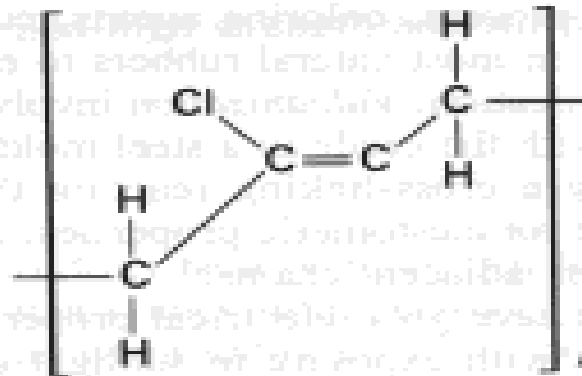
Polyisoprene (IR)



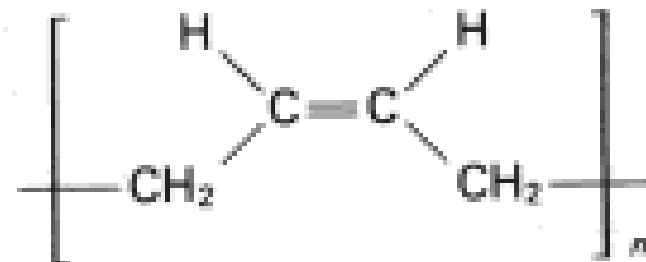
Polystyrene-1 part

(SBR)

Butadiene-3 parts

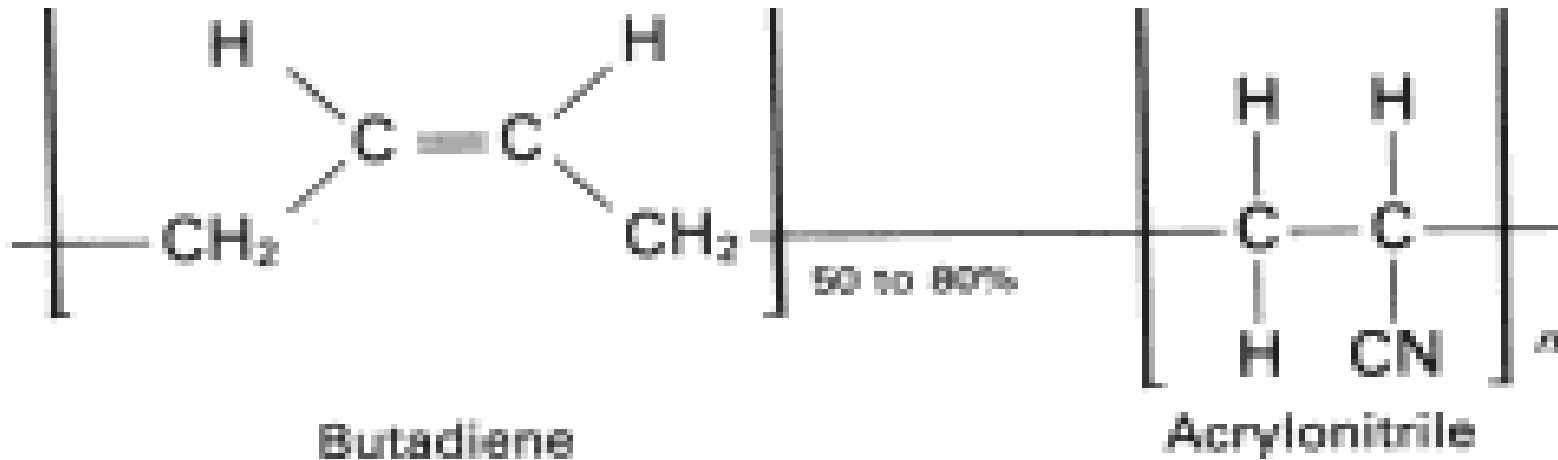


Polychloroprene (CR)



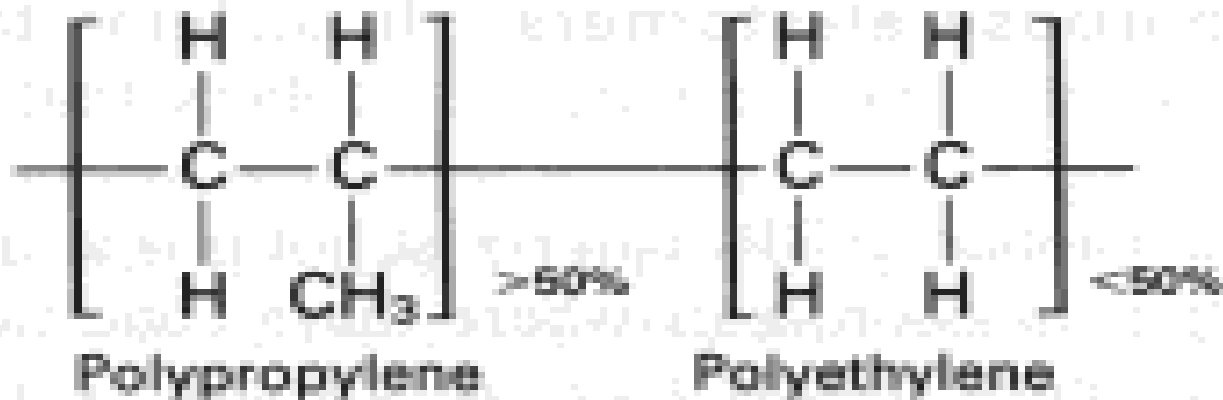
Polybutadiene (BR)

Monomeros elastoméricos más comunes



50 to 80%

Nitrile rubber (NBR)

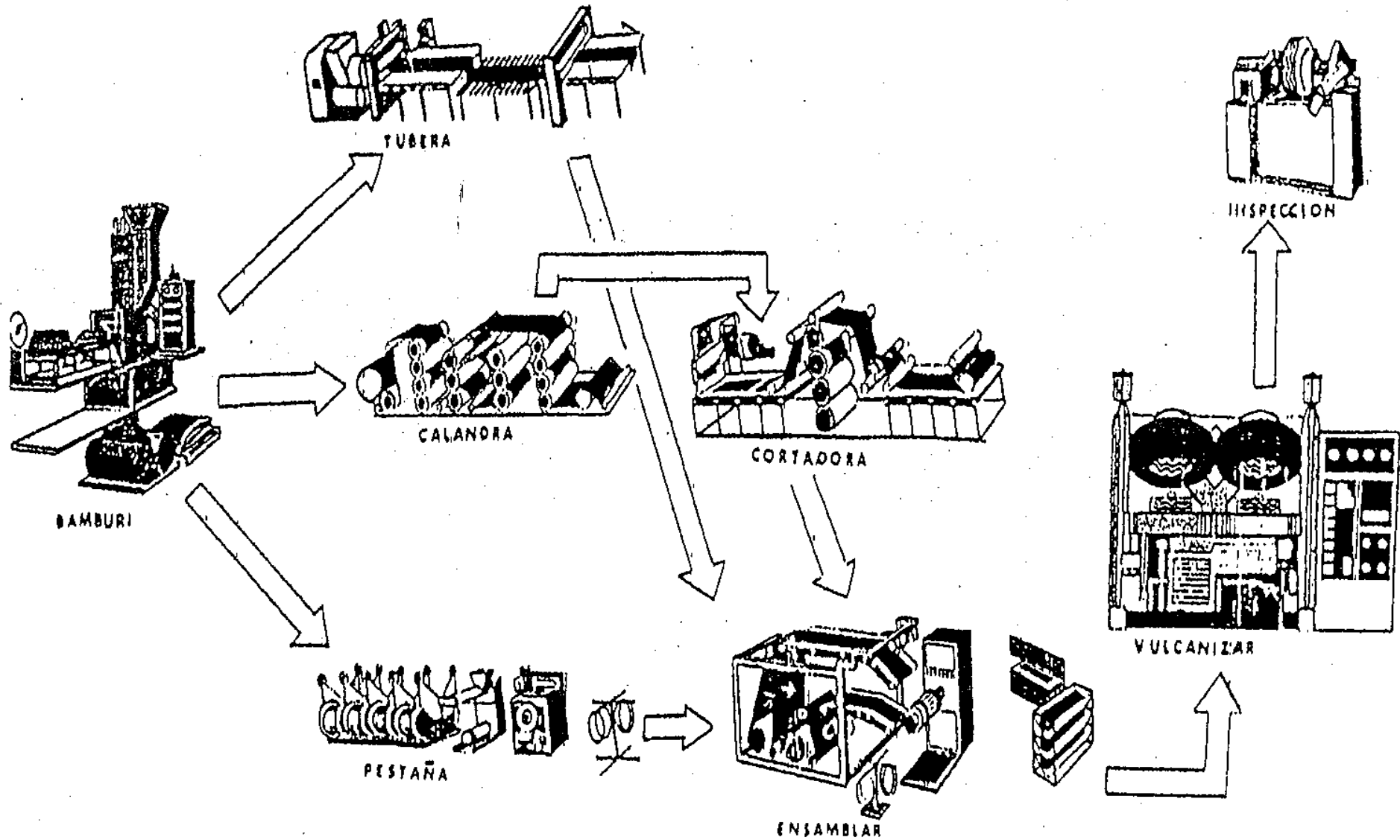


>50%

<50%

EPM rubber

PROCESO DE FABRICACIÓN DE NEUMÁTICOS



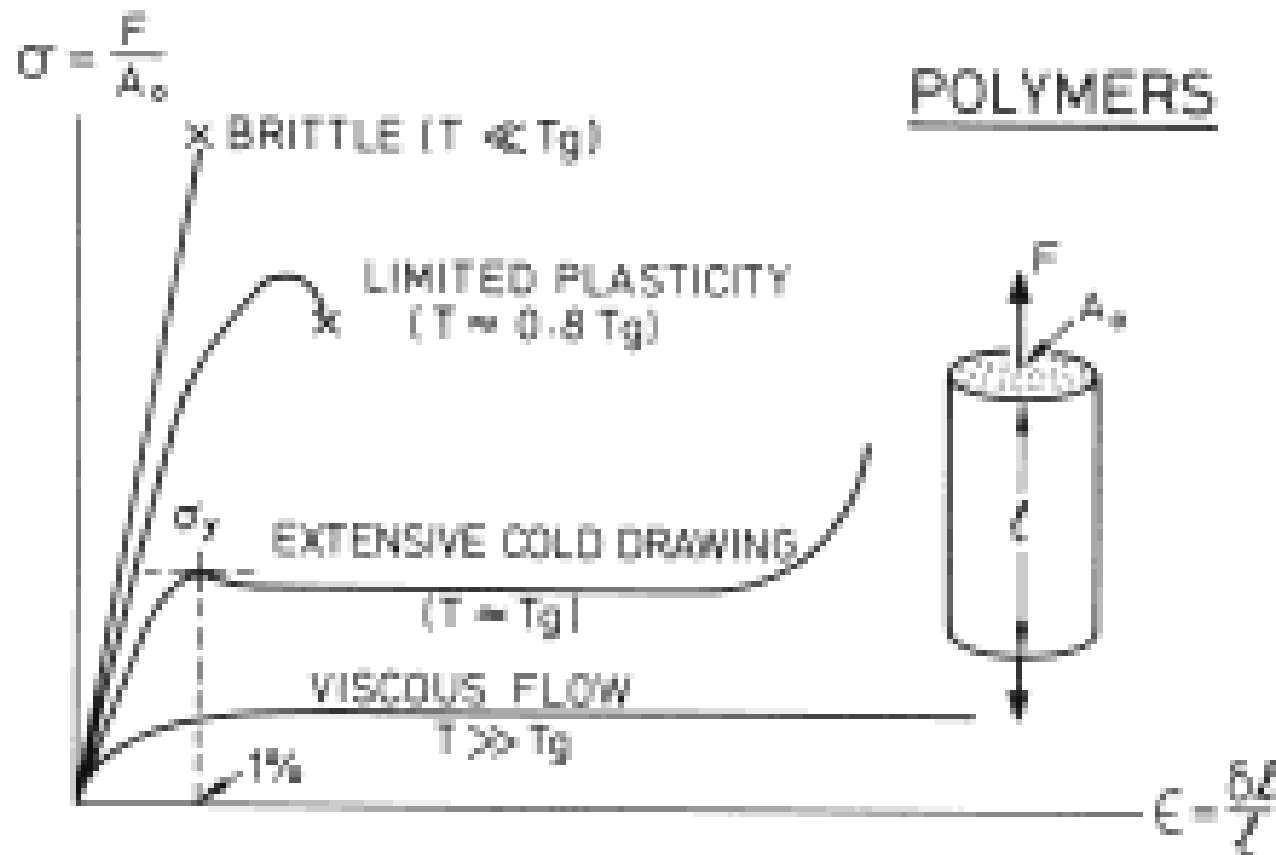
Datos de materiales elastomeros

TABLE 13.5 Properties of Different Rubbers (Elastomers)

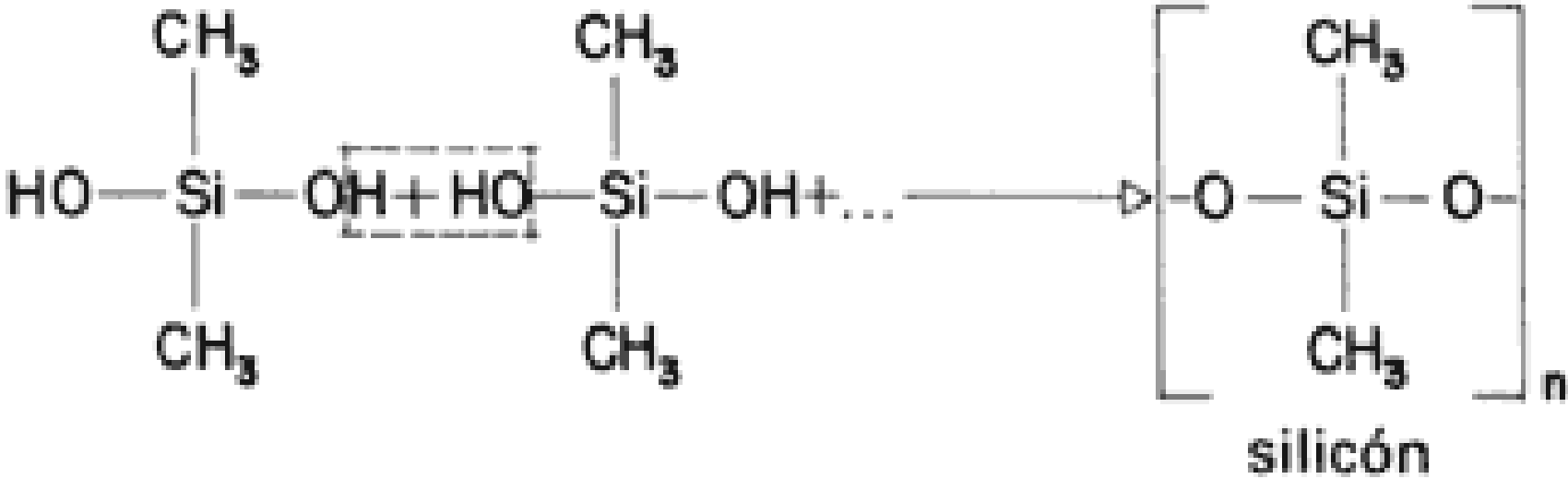
<i>Common Name</i>	<i>Chemical Name</i>	<i>Tensile Strength,* psi</i>	<i>Percent Elongation</i>	<i>Resistance to Oil, Gas</i>	<i>Useful Temperature Range, °F (°C)</i>
Natural rubber	<i>cis</i> -Polyisoprene	3,000	800	Poor	-60 to 180 (-51 to 82)
GR-S or Buna S	Butadiene- styrene copolymer	250	3,000	Poor	-60 to 180 (-51 to 82)
Isoprene	Polyisoprene	3,000	400	Poor	-60 to 180 (-51 to 82)
Nitrile or Buna N	Butadiene- acrylonitrile copolymer	700	400	Excellent	-60 to 300 (-51 to 149)
Neoprene (GR-M)	Polychloroprene	3,500	800	Good	-40 to 200 (-40 to 93)
Silicone	Polysiloxane	700	300	Poor	-178 to 600 (-117 to 315)
Urethane	Diisocyanate polyester	5,000	600	Excellent	-65 to 240 (-54 to 115)

*Multiply psi by 6.9×10^{-3} to obtain Mn/m^2 [MPa] or by 7.03×10^{-4} to obtain kg/mm^2 .

Curva de carga-deformación de los polímeros

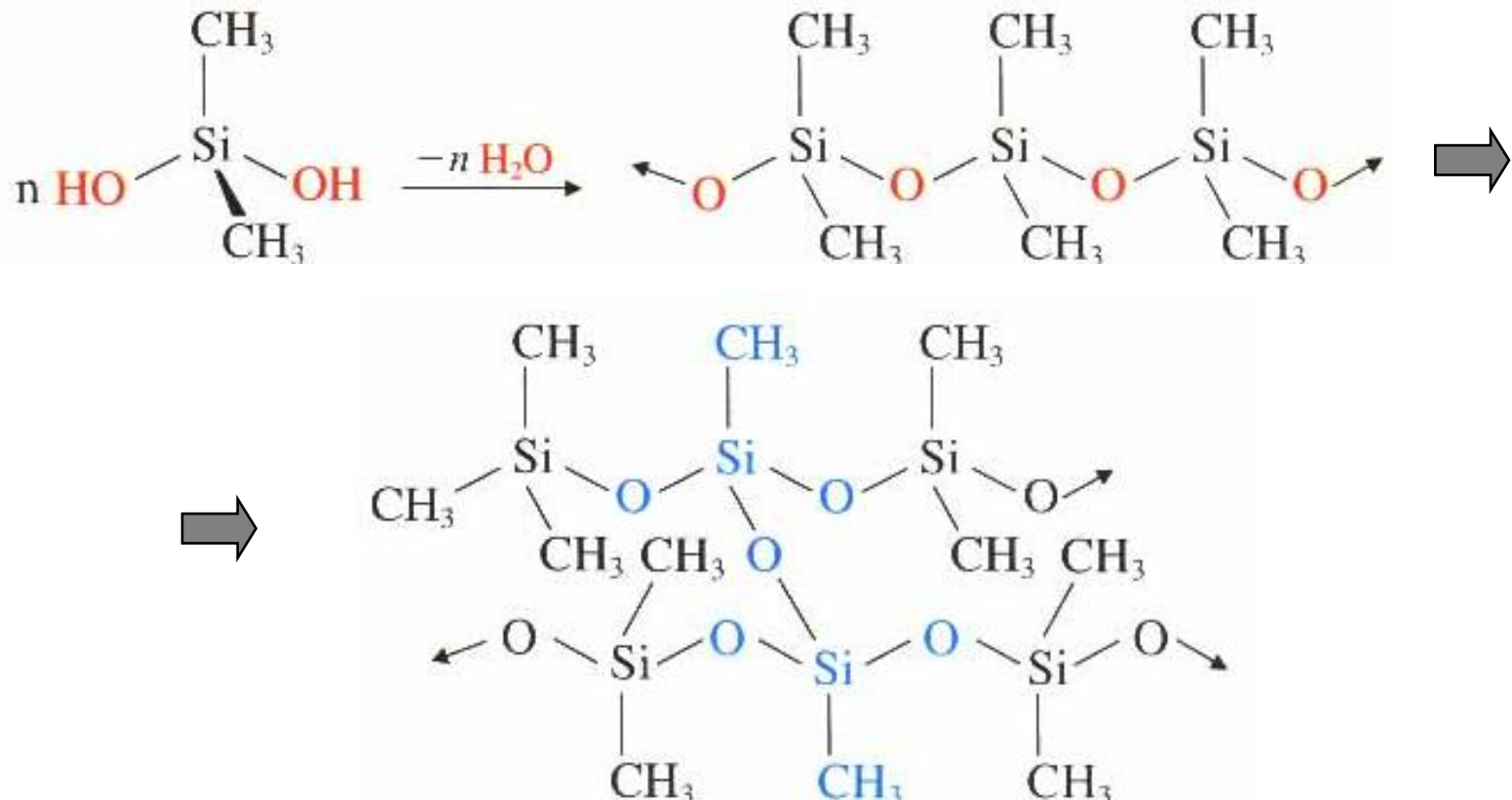


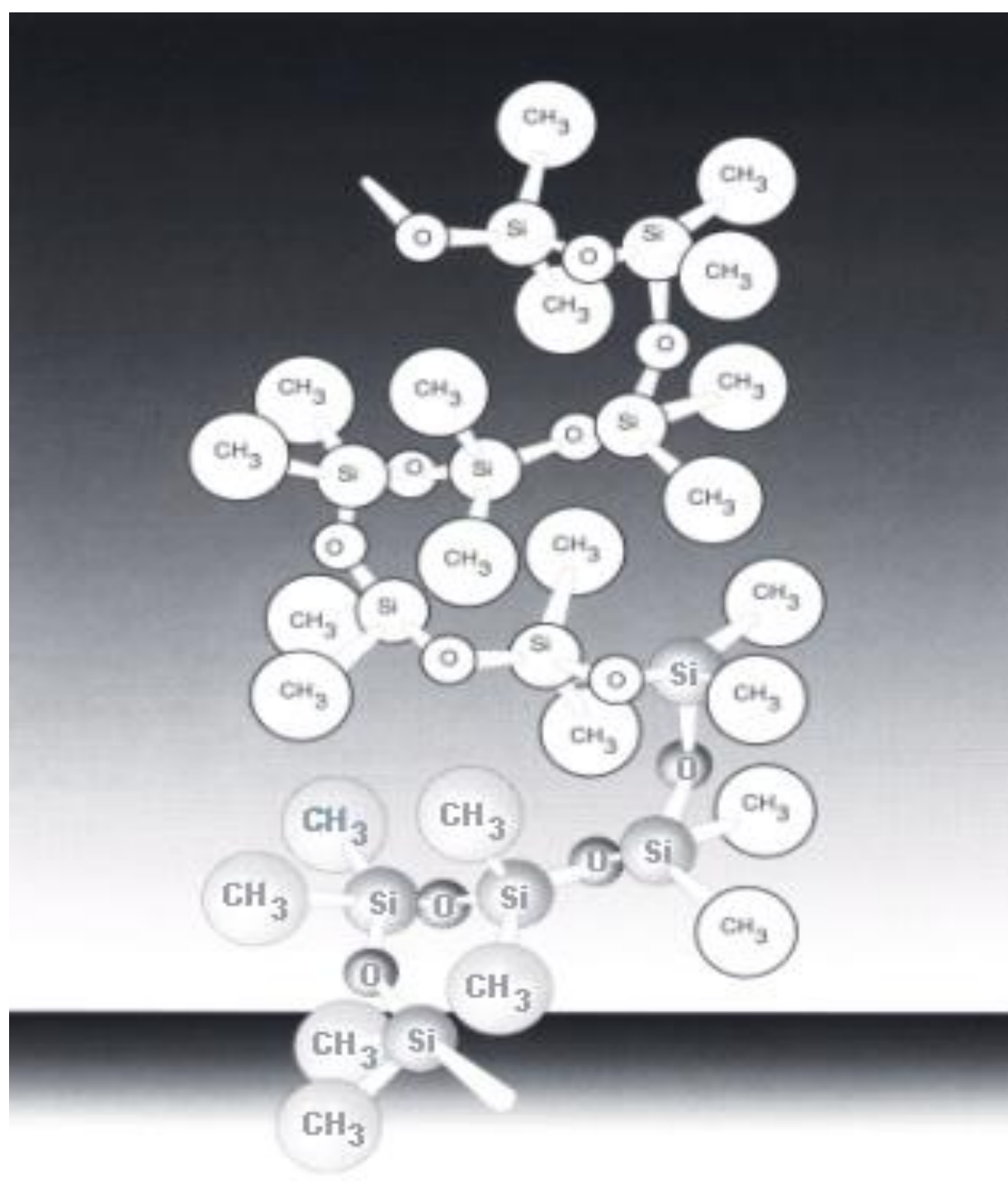
SILICONES



Polímeros de condensación: Siliconas

- Proceden de monómeros del tipo $R_2Si(OH)_2$
- Se utiliza para sellar juntas debido a su carácter hidrofóbico.





SILICATOS Y SILICONAS

El silicio forma una variedad de polímeros naturales inorgánicos, los *silicatos*, que contienen unidades SiO_4

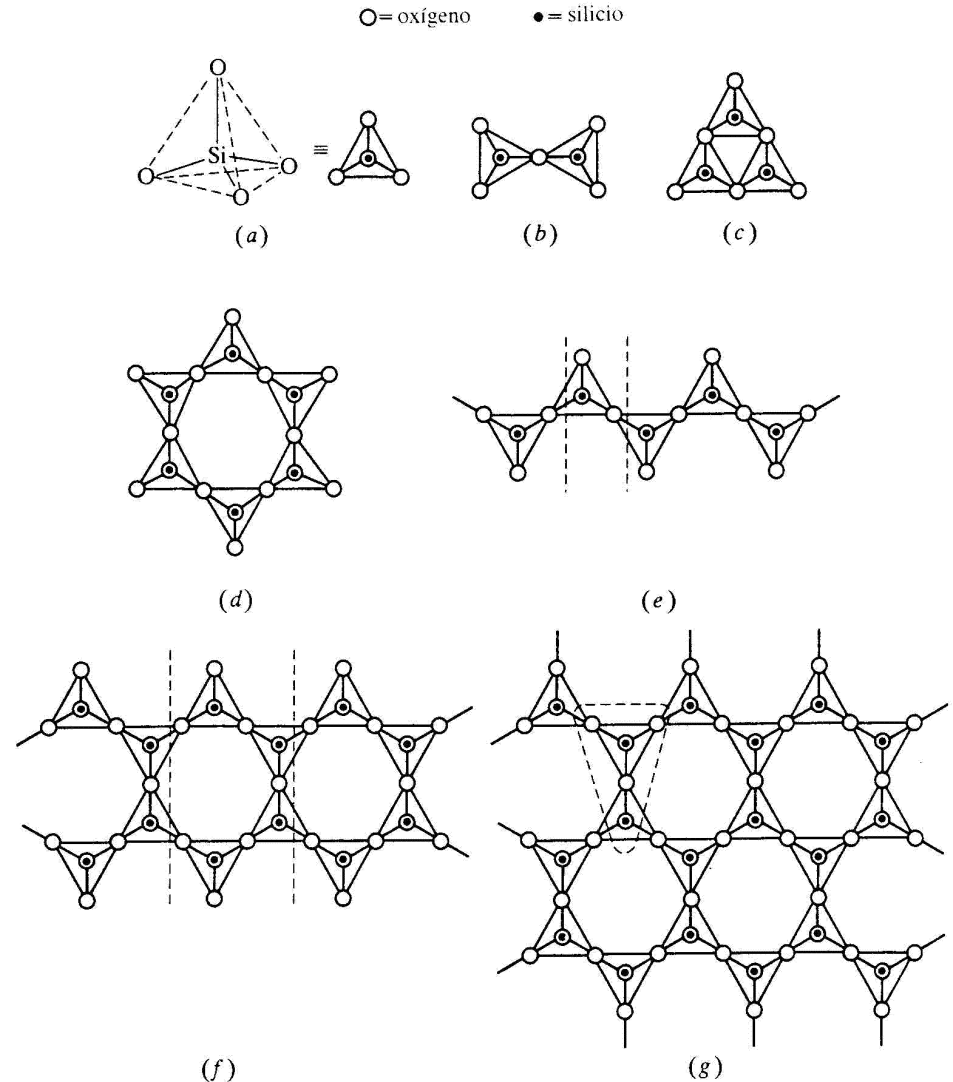
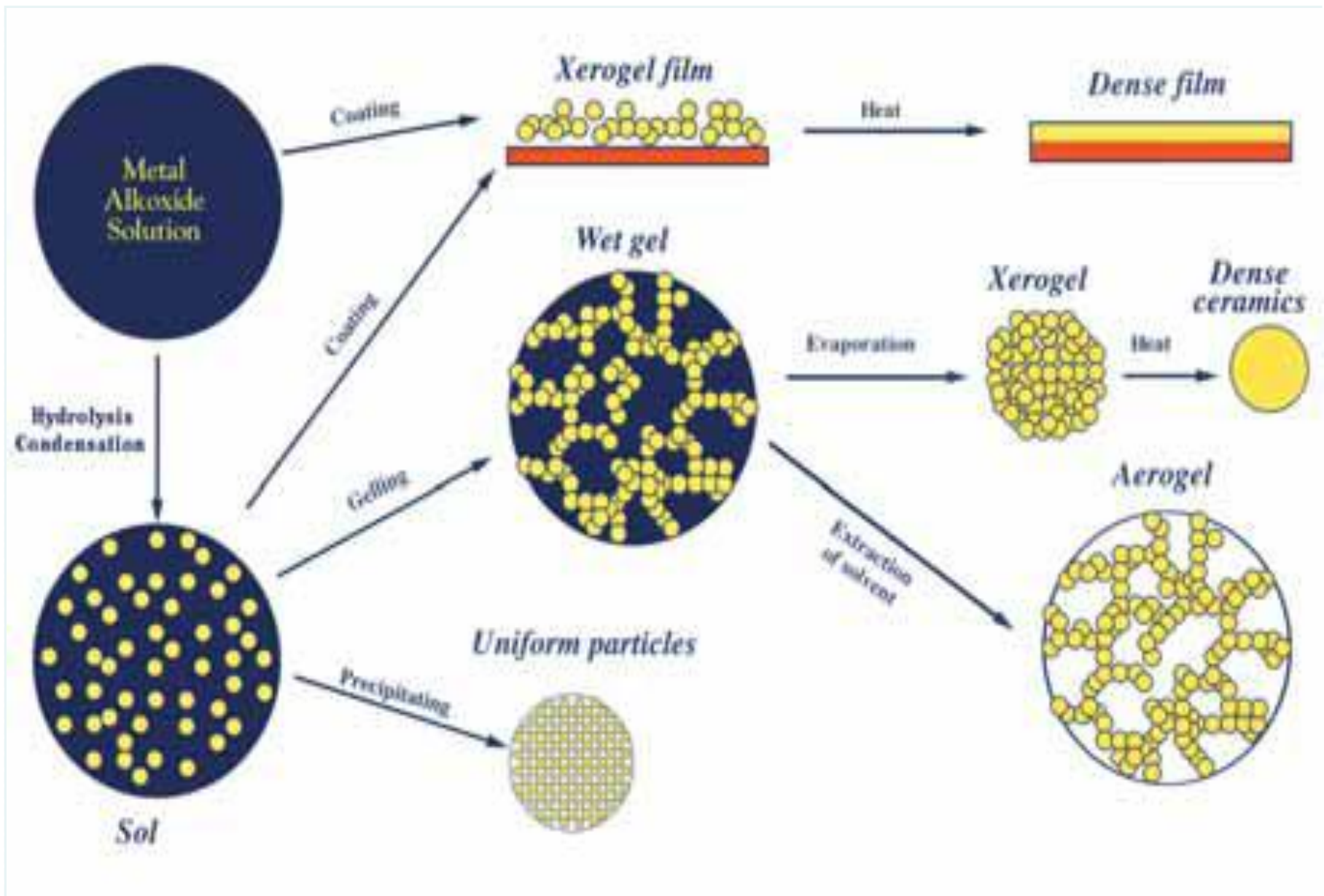


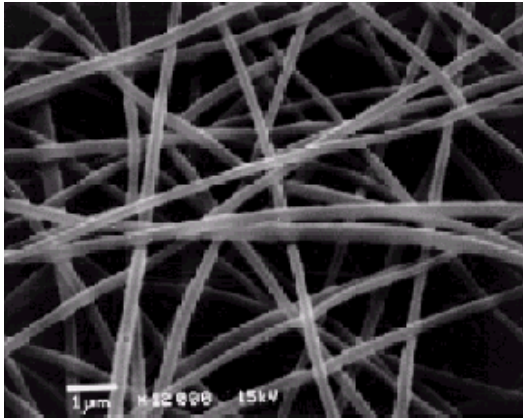
Figura 15.8. Estructuras de los silicatos: (a) orto SiO_4^{4-} , (b) piro $\text{Si}_2\text{O}_7^{6-}$, (c) cíclico $\text{Si}_3\text{O}_9^{6-}$, (d) cíclico $\text{Si}_6\text{O}_{18}^{12-}$, (e) cadena sencilla SiO_3^{2-} , (f) cadena doble $\text{Si}_4\text{O}_{11}^{6-}$ y (g) laminar $\text{Si}_2\text{O}_5^{2-}$.



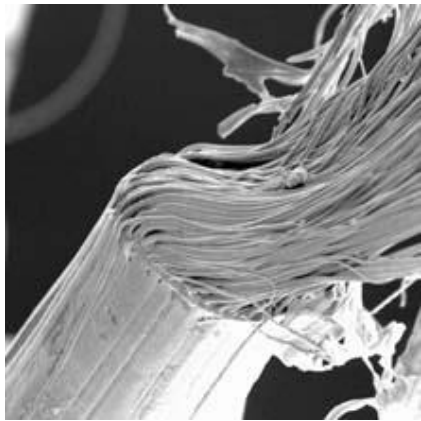
FIBRAS NATURALES



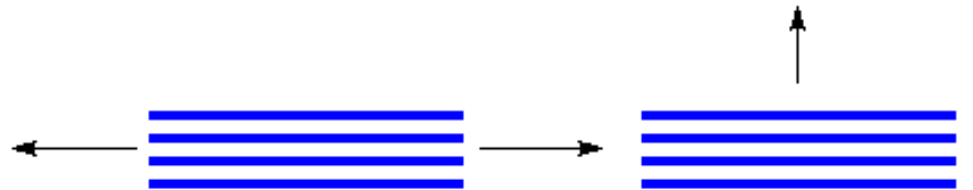
FIBRAS ARTIFICIALES



POLIACRILONITRILO



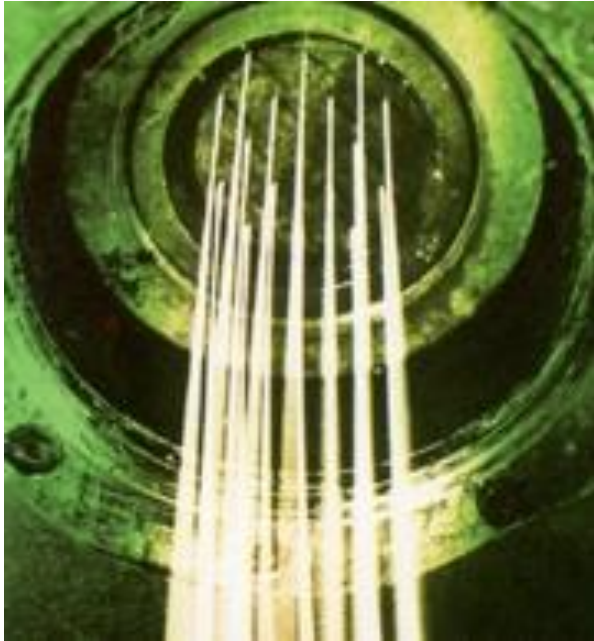
POLIPROPILENO



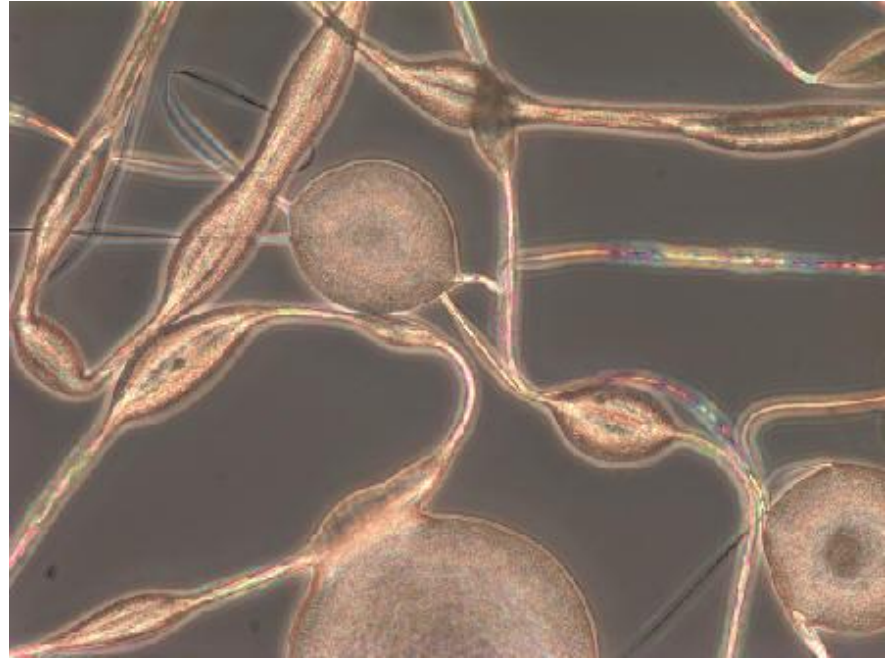
Las fibras orientadas son resistentes cuando usted estira en la dirección de las fibras.

Pero son débiles en ángulos rectos a la dirección de las fibras.

SPINNING (HILADO)



Spinning por fusión o en solución



Electrospinning

Las soluciones de polímero se rotan en un campo eléctrico de alto voltaje. La gota suspendida de polímero se carga y colapsa formando “chorros”.