Types of Nanomaterials				Advantages and Disadvantages
Organic Nanoparticles				
Lipid and surfactant- derived nanoparticles	Vesicular	Liposome	Vesicles composed of a bilayer of phospholipids	<b>Advantages:</b> Biodegradable, biocompatible, amphiphilic (ability to compartmentalize and s penetration
				Disadvantages: Lower medication stacking; lower reproducibility and lower chemical sta
		Ethosome	Liposome formulation with inclusion of a very high concentration of ethanol	Advantages: Higher efficiency and penetration of cosmetic delivery into the skin
				Disadvantages: Poor yield problems, stability and possibility of coalescence
		Transferosome	Liposome formulation with inclusion of edge activators (i.e. softening surfactants)	Advantages: Superior penetration through skin barrier
		Niosome	Vesicles composed of non-ionic surfactants	Advantages: Cheaper raw materials; higher efficiency, penetration, bioavailability and prophospholipid-based vesicles)
				<b>Disadvantages:</b> Higher cost of production; physical and chemical instability; leakage of the
	Non- vesicular	Solid lipid nanoparticles	Inner lipid core (solid at body temperature) formed only by solid lipids prepared by microemulsion and high-pressure homogenization	Advantages: Potential carriers of chemically labile molecules; occlusive properties that pro- duration of action, ease of large-scale production; higher bioavailability and biodegradab
				Disadvantages: Lower shelf life and decreased drug encapsulation
		Nanostructured lipid carrier	red Inner lipid core (solid at body temperature) formed by solid and liquid lipids prepared by microemulsion and high-pressure homogenization	Advantages: Potential carriers of chemically labile molecules; occlusive properties that pro of oils in lipid matrix decreases crystallinity, improving loading capacity, long-term encap large-scale production
				Disadvantages: Lower duration of action
		Nanoemulsion (o/w)	Biphasic systems composed of oil-phase and liquid-phase and one or more emulsifying agents	Advantages: If non-ionic surfactants are used, results in more stability against agglomera droplets makes the material transfer faster, enhancing delivery performance, fluid and transfer faster.
				Disadvantages: Less thermodynamic stability when compared to microemulsion; more di
Polymeric	Nanosphere	Polymeric matrix in which numerous active ingredients can be entrapped or into which they can be adsorbed		Advantages: Improved performance of chemically labile, poorly water-soluble or volatile n their beneficial effects
Nanoparticles	Nanocapsule	Liquid core surrounded by polymeric shell – the inner core holds the bioactive agent while the polymeric shell controls its release		Advantage: Improved performance of chemically labile, poorly water-soluble or volatile mo sustained release of the loaded active molecules; resolution of incompatibility issues bet
				Disadvantages: An additional purification step is required after nanocapsule formation
Nanocrystals	Clusters made up of thousands of molecules joined together in a fixed pattern to form a group with sizes ranging from 10 to 400 nm stabilized by surfactant/polymeric coating			Advantages: Possible solution to poorly water-soluble agents; large surface area and poor adhesivenes, and dissolution rate; 100% drug loading ability
				Disadvantages: Possibility of aggregation; not appropriate for aqueous APIs; only stable to
Dendrimers	Three-dimensional nanostructured macromolecules that are extensively branched			<b>Advantages:</b> Higher solubility of lipophilic drugs; controlled-release drug formulation; mai shelf life of formulation
				Disadvantages: Not good materials for hydrophilic drugs; cellular toxicity; higher manufact
Inorganic Nanoparticles				
Titanium dioxide and zinc oxide	Inorganic UV rad	liation filters		Advantages: TiO2 has a higher sun protection factor at the nanoscale, which makes it mo transparency, in contrast with its original color due to the large surface-area-to-volume r
	ZnO : more effect	tive for UVA		Disadvantages: Inhalation of a large amount of these NPs is harmful
	TiO2: more effec	tive for UVB		
Gold and silver	Gold NPs play a	substantial role in fixing	g skin damage and improving skin surface, grace and flexibility.	Advantages: Stability, biocompatibility, antifungal, antibacterial and anti-aging benefits
	Silver nanoparticles can be used as successful inhibitors of various microorganisms.			Disadvantages: The safety of colloidal silver in nanostructures concerning its use in oral a
Silica	Mainly composed of amorphous silica nanodispersions with a size range of 5 nm to 100 nm; can deliver both hydrophilic and lipophilic entities to their respective targets by encapsulation			Advantages: Hydrophilic surfaces favor extended distribution and low manufacturing cost
				Disadvantages: Concerns about their safety; size and surface changes are factors must b
Carbon Black	Colorant - CI77266			Advantages: Micron-sized NPs have a lower propensity of causing cytotoxicity, aggravation compared to NPs
				Disadvantages: Can be used in cosmetic items when there is no danger of being breathe
tris-Biphenyl triazine	Novel, powerful and photostable filter			Advantages: Broad-spectrum UV protectant
				<b>Disadvantages:</b> Not dangerous if applied to solid, unbroken skin; concerns related to posselected tissues
Bucky Balls (Buckminsterfullerene/ C60)	Carbon fullerene is a three-dimensional spherical compound that comprises a carbon ring with an odd number of atoms			Advantages: Antioxidative properties (potent scavenging ability of free radical oxygen spec
				<b>Disadvantages:</b> Fullerenes alone have limited applications due to their hydrophobic natur improved their aqueous solubility

solubilize both hydrophilic and lipophilic materials) and higher skin

bility; may trigger an immune response

longed stability of drugs (especially when compared to

he drug; time-consuming production

romote skin hydration and penetration of bioactive agents; higher bility

omote skin hydration and penetration of bioactive agents; the inclusion osulation and stability of the system leading to higher shelf life, ease of

ation and precipitation and the large interfacial area displayed by their ransparent appearance and amphiphilic

ifficult to prepare; acid sensitive; lower duration of action

nolecules; sustained release of the loaded active molecules prolonging

olecules; protection of ingredients; masking of undesirable odors; tween formulation components; sustained release formulation

r crystallinity lead to higher drug solubility, particle distribution,

o a certain extent

ntenance of the stability of the drug in cosmetic formulations; higher

cturing cost

re effective and results in a superior restorative effect due to its ratio

and dermal cosmetic items is ambiguous

is; improve the adequacy, surface and shelf life of cosmetics

be considered while surveying its toxicity

on and changes in phagocytosis in human monocytes when

ed in

sible harmful impacts with the potential to bioaccumulate in

cies)

re but the use of surface-active agents in a suitable concentration has