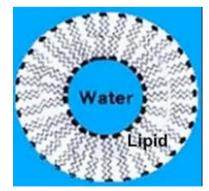


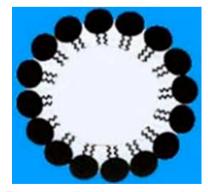
- ➤ They are extremly small vesicles (lower than 300nm)
- ➤ Mainly made of phospholipids
- > The phospholipids are organised in bilayers

Nano LPD's



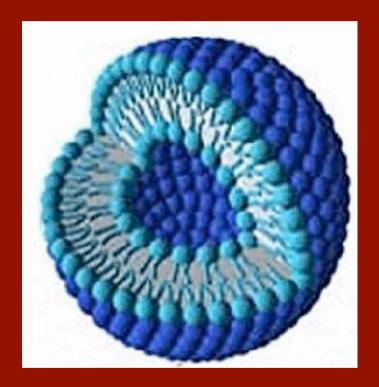
Phospholipids in bilayer

Micela



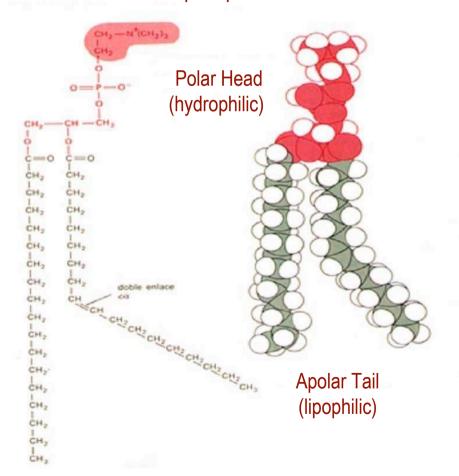
Phospholipids in monolayer

Definition





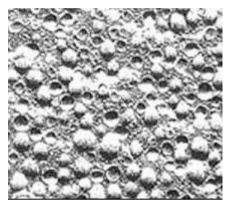
Phospholipid Structure



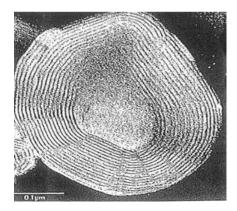
Composition

- ➤ They are mainly made of natural origin phospholipids
- > Their characteristics are :
 - ➤ To have a hydrophilic part (polar head)
 - > and to have a lipophilic part (apolar tail)

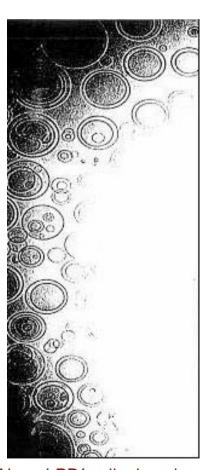




Nano LPD's unilamelars



Nano LPD's multilamelars



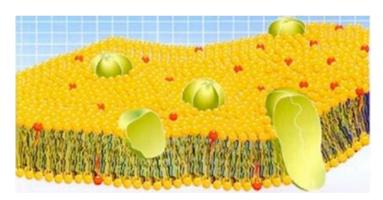
Nano LPD's oligolamelars

Clasification

- ➤ Size:
 - ➤ Small (diameter < 100 nm)
 - ➤ Big (diameter > 100 nm)
- ➤ Number of bilayers:
 - ➤ Unilamelars
 - ➤ Oligolamelars
 - > Multilamelars



Analogy between the cellular membrane structure and Nano LPD's



Cellular Membrane

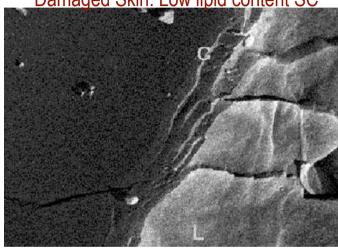


Usages and Advantages

- > They are **natural** delivery systems of active ingredients
- They are controlled and released carrier systems
- They are analogue structures of the cellular membranes (phospholipids)
- They increase the efficacy and decrease the unwanted side effects of the active ingredients (toxicity)



Damaged Skin. Low lipid content SC

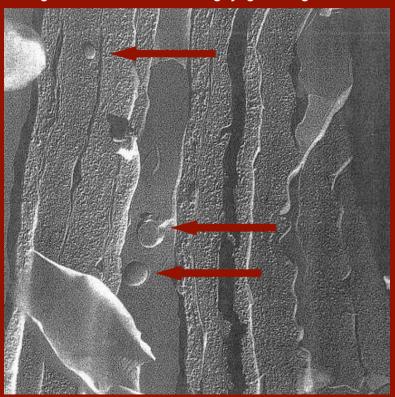


Skin treated with Nano LPD's. Re-epitheliated SC



Usage and Advantages

Image of the Nano LPD's gojng through the SC



Study carried out by Dr. A De La Maza. Departament df Surfactants. CSIC, Barcelona



Interaction Nano LPD's – active ingredient

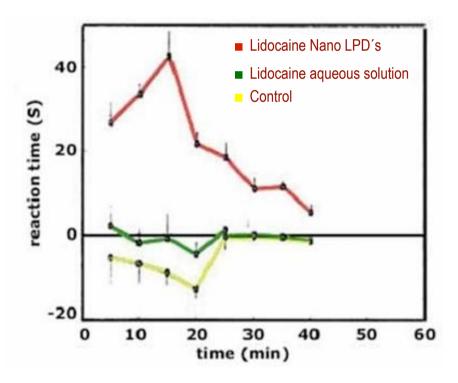


Usage and Advantages

- The active ingredients reallocate at the interface of the Nano-LPD's
- Because of the structure and also the phospholipid bilayer composition, the Nano-LPD's can incorporate: :
 - Hydrophilic Actives (within the vesicle)
 - Lipophilic Actives (between the layers)



Improvement of the analgesic action of the lidocaine (topically applied)



Usages and Advantages

- Prolongation of the bioavailability of the active ingredient
- Better absortion, penetration and diffusion of the active ingredient
- Stabilization of the active ingredient
- Introduction of alternative administration ways of the active ingredient



Menopause

Obesity and

OBESITY:

Metabolic syndrom defined as a pathologic increase of the quantity of adipose tissue.

They are different problems that normally converge

Low hydration

Predisposing

Hormonal Factors Dietetic Factors: (strogens): (bad habits): **Puberty** Fats SALT excess Pregnacy

cellulitis

CELLULITIS:

It is an alteration of the subcuatneous cellular tissue that provokes changes in the microcirculation at the connective tissue giving morphologic, histochemical and biochemical modifications of the tissue

factors

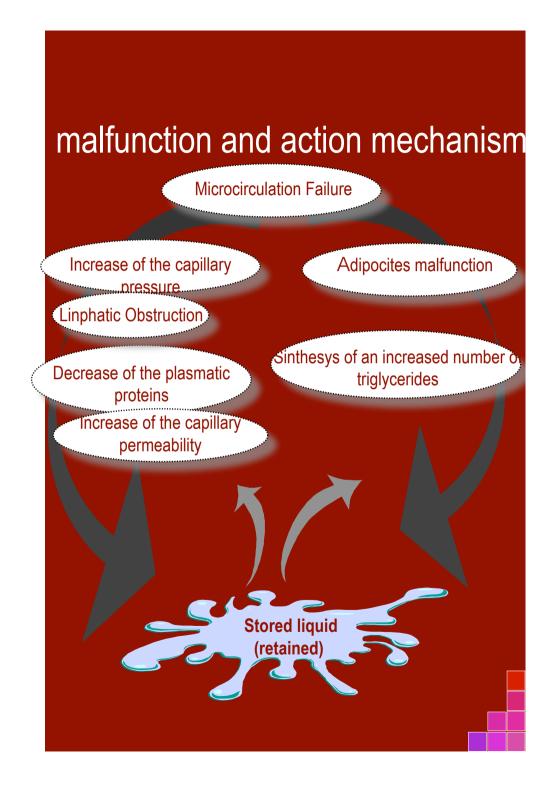
Mechanical factors: (bad habits): Sementary life Very tight cloths Inadequated shoes **Toxics Factors:** (bad habits): Tobacco Alcohol Drugs



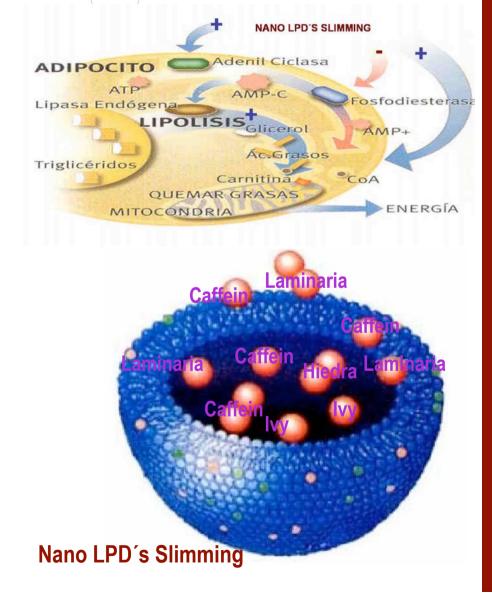
Actions

Required Actions:

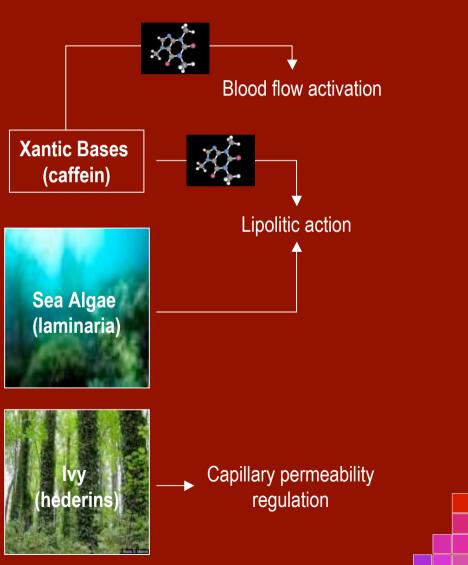
- ➤ Lipolitic action
- > Activation of the blood circulation
- > Regulation of the capillary permeability.







Slimming composition and mechanism of action

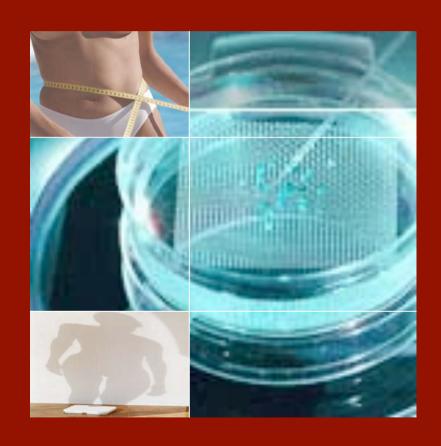




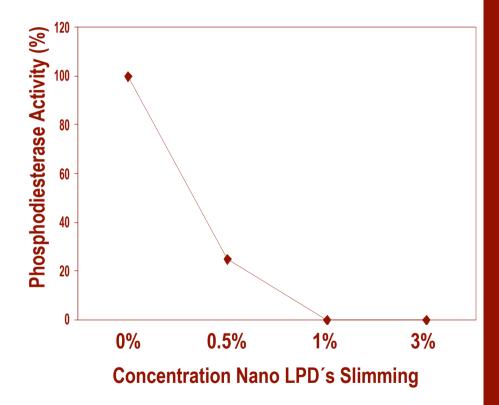
Nano LPD's Slimming Efficacy Test

- > In vitro test:
 - ➤ Phosphodiestarase Inhibition
 - ➤ AMPc Concentration
 - ➤ Free Fatty Acids (FFA)

- > In vivo test:
 - > Assesment of the infiltrated fat decrease at the dermis
 - Perimeter of the thigh area







Slimming Phosphodiesterase Inhibition

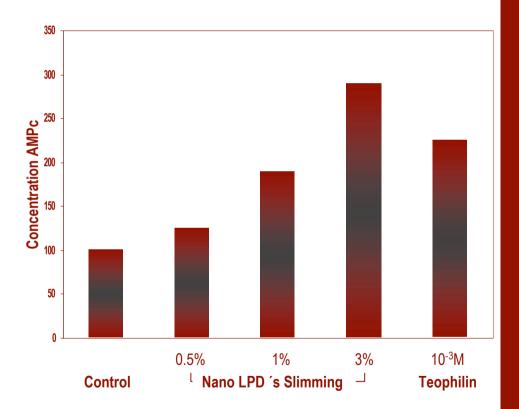
➤ Objective :

Measurament of the phosphodiesterase activity inhibiton (AMPc-PDE), which is responsible of the AMPc (lipase activator)inactivation

> Methodology:

- ➤ Phospodiesterase [3H] cAMP SPA enzyme
- > Enzyme : AMPc-PPE bovine heart
- ➤ Positive Control: teophilin
- ➤ Negative Control :enzyme free culture
- ➤ SPA Technology for the evaluation(Scintillation Proximity Assay)
- ➤ Efficacy assessment of Nano LPD's Slimming at 0.5%, 1% y 3%





Nano LPD's Slimming AMPc Concentration

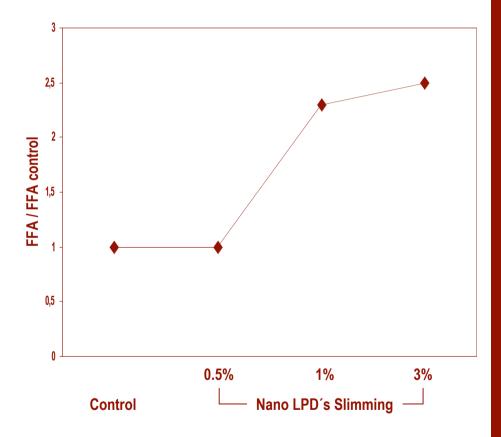
➤ Objetive :

To asses the increment of AMPc in human adipocytes after having applied with Nano LPD's Slimming

➤ Methodology:

- > Enzymatic Inmunotest (Biotrack kit, Amersham, RPN 225)
- ➤ Culture : human adipocytes
- ➤ Positive Control: teophlin
- ➤ Negative Control: culture
- ➤ Efficacy evaluation of Nano LPD's Slimming at 0.5%, 1% y 3%





Slimming Free Fatty Acids (FFA)

➤ Objetive :

To asses the liberation of FFA in human adipocytes after having applied with Nano LPD's Slimming

➤ Methodology:

- > Enzymatic colorimetric measurement
- Waco Chemicals 994-75409 (Kit Nefa-C)
- > Culture : human adipocytes
- ➤ Negative Control: culture
- ➤ Efficacy evaluation of Nano LPD's Slimming at 0.5%, 1% y 3%



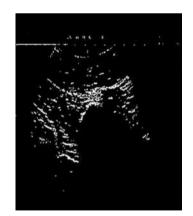
Nano LPD's Slimming in vivo test



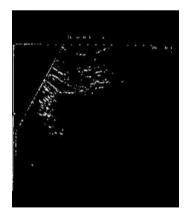
Left leg



Right leg



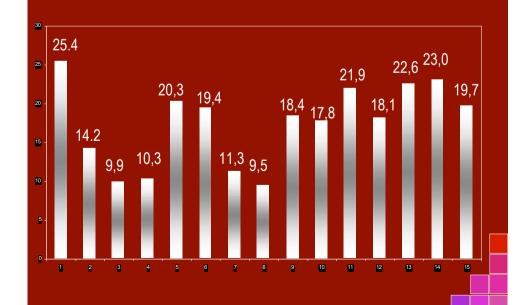
Volunteer 10



➤ Objetive :

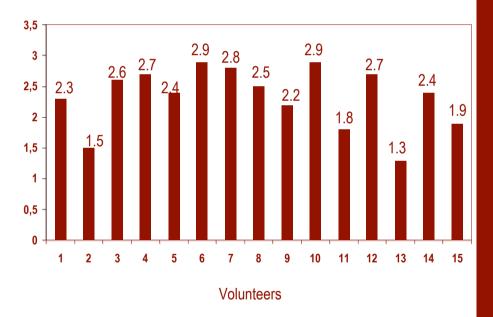
To asses the decrease of infiltrated fat at the volunteers dermis with cellulitic symptomatology

- ➤ Methodology:
 - > Echographic measurement at the thigh areas
 - > Application of a cream twice a day containing 5% of Nano LPD's Slimming vs. Placebo cream
 - > Echography at the beginning and at the of the treatment (28 days)
- > Results: 17.5% average reduction of the infiltrated fat





Perimeter variations (cm.)



Slimming thigh perimeter

➤ Objetive :

To asses the decrease of thigh area perimeter through application of a cream containing 5% of Nano LPd's Slimming compared to a placebo

- ➤ Methodology:
 - ➤ Centimetric measurement of the thigh perimeter
 - ➤ Measurement at the beginning and at the end of the treatment (28 days)
 - ➤ Echography first day of the treatment and at the end of the treatment (28 days)
- > Results: 2.3 cm. of perimeter average diminution



- ➤ New technology :
 - ➤ Nanosystems
 - > Natural
 - > Controlled and sustained release

- ➤ Natural active ingredients of proven efficacy
 - > Laminaria
 - ➤ Caffein
 - ≽lvy



