



PRODUCT DATA SHEET

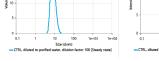
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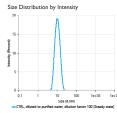
Lipodisq[™] Control Sterile Solution

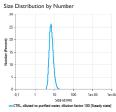
Nano-formulated aqueous solution: Ready-to-use

Cat. No.: |AX-700-100 Lot. No.:

Synonyms	Detergent-free nano-formulation made of styrene-maleic acid lipid particles (SMALP)		
Empirical Formula	N/A		
Concentration	N/A		
Size	Iml		
MW	N/A		
CAS	N/A		
Purity	N/A		
Solution pH	7.00 - 7.50		
Solubility	Soluble in water, PBS, Tris and other physiological solutions as formulated in a proprietary, thermostable, aqueous lipid nanoparticulate formulation (Lipodisq [™] , Malvern Cosmeceutics Ltd., Malvern UK). Avoid the use of buffers with divalent ions such as Ca or Mg or pH <6.5 or >8.0, which can cause particle instability.		
Formulation	Lipodisq [™] are nanosized lipid-based discoidal particles that can be manufactured to incorporate hydrophobic, poorly water-soluble compounds, such as lipids, lipoproteins and glycolipids.		
Appearance	Colourless clear aqueous solution		
Handling	Keep sterile. Avoid skin and eye contact.		
Activity	Cell culture tested (human macrophage cell line) (MTT). Recommended starting dilution: 1:200 or higher. Optimal working concentrations depend on the applications and need to be determined. Published procedures using Lipodisq TM formulations (Curcumin and IAXO TLR4 antagonists) in vivo rodent models at 3-10mg/kg. Recommended route of administration is subcutaneous (s.c.) with or or nasal application as a possible alternative, which needs to be optimised.		
Shipping	Ambient		
Storage	2-8°C		
Stability	12 months after receipt (unopened and as supplied)		
MSDS	Available on request		







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- A nanoparticle (11-40nm) drug delivery system comprising a discoidal phospholipid bilayer membrane stabilised by a chaperone molecule annulus.
- Internal properties of the phospholipid membrane support the disposition and stabilisation of drug molecule candidates and preserve the native conformation of membrane molecules.
- The resulting encapsulated actives are rendered water-soluble and specialised for intra-cellular penetration/delivery via endosomal uptake mechanisms.
- Lipodisq[™] solutions show a good safety profile and are suitable for in vitro and in vivo investigations.
- For a customizable biodegradable Lipodisq[™] version with a higher concentration of actives or an alternative lipid option, contact Innaxon.

Component	Concentration	CAS#	EC#
Water (sterile)	QS	7732-18-5	231-791-2
Poly(styrene maleic acid)	25mg/ml	26762-29-8	607-996-I
Lecithin	I 0mg/ml	92128-87-5	295-786-7

Lipodisq[™] References

Lipodisq[™] Technology

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DISCLAIMER: THIS PRODUCT IS NOT INTENDED OR APPROVED FOR HUMAN, DIAGNOSTICS, OR VETERINARY USE. THIS PRODUCT IS FOR RESEARCH USE ONLY (RUO).

MATERIAL SAFETY DATA: This material should be considered hazardous until information to the contrary becomes available. Do not ingest, swallow, inhale or allow to enter the bloodstream. Avoid contact with the eyes, or the skin, or clothing. Wash thoroughly after handling. Access to this material must be restricted to personnel, who are appropriately experienced, qualified, competent, and properly trained to use it.





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Cat. No.: IAX-700-100 **Lot. No.:**

Lipodisq[™] References

- [9] Nano-size uni-lamellar lipodisq improved in situ auto-phosphorylation analysis of E. coli tyrosine kinase using (19)F nuclear magnetic resonance. Li D, et al. Protein Cell (2015); 6:229
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