### DYNASAN® - VERSATILE EXCIPIENTS FOR SOLID DOSAGE FORMS

The DYNASAN<sup>®</sup> product range is a family of high-purity monoacid triglycerides.

Their chemical nature, raw materials, manufacturing and purification result in products with advantageous properties and impurity profile. 

PRODUCT	CHEMICAL DESCRIPTION	MELTING POINT °C	HYDROXYL VALUE mg KOH/g
DYNASAN® 114	Trimyristin	55 - 58	Max. 10
DYNASAN® 116	Tripalmitin	63 - 68	Max. 10
DYNASAN® 118	Glyceryl Tristearate	69 - 73	Max. 5

\* Additional DYNASAN® Triglycerides from mixed or pure fatty acids are technically feasible; please get in touch with us.

The optimized degree of esterification leads to almost no free OH functionality on the backbone of the excipients and removes formulation and stability challenges in tablets, typically encountered with lubricants based on metallic salts of fatty acids<sup>10</sup>:

- Metal salt impurities interacting with APIs
- API interaction with lubricants due to their ionic characteristics
- Microenvironmental pH shift into alkaline milieu, creating a risk of drug hydrolysis
- Initiation of redox cascades degrading APIs
- Metal ion-mediated degradation of APIs
- lonic excipients and salt counter-ions show reactivity towards amines, which are common functional groups in APIs with metal soaps

Incompatibilities with metallic salts of fatty acids have been reported, e.g. for the following active ingredients: Acetylsalicylic Acid, Captopril, Besylate, Cephalexin, Clopidogrel Besylate, Erythromycin, Glibenclamide, Glimepiride, Ibuprofen, Indomethacin, Ketoprofen, Norfloxacin, Oxacillin, Penicillin G, Temazepam

### DYNASAN° 118

Glyceryl Tristearate / USP-NF

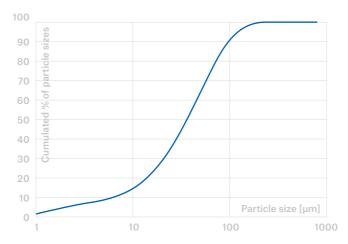
### **Characteristics**

DYNASAN® 118 is a monoacid triglyceride based on C18 fatty acid with a high melting point. It is available as a microfine powder or in flaked form particularly suitable for oral solid dosage forms.

### Uses

- Matrix for lipophilic actives
- Coated formulations for protection of sensitive drug molecules (or APIs) or for taste masking
- Solid Lipid Nanoparticles, Self-Emulsifying Drug **Delivery Systems**
- Processing by hot melt techniques like granulation and extrusion

### Particle Size Distribution of DYNASAN® 118 (microfine powder)



#### Benefits for Tableting<sup>11) 12)</sup>

- Effective lubricant for tablet production in low concentrations (~ 0.5%) without the risks of fatty acid soaps
- No influence on drug release up to a concentration of 1%, model drug Acetylsalicylic Acid
- Higher concentrations from 2 to 5% increase the hydrophobic characteristics of tablet matrix and introduce retardation effects into oral dosage forms for sustained release characteristics
- Enhances the tablet fracture stability due to an increased lubricity factor resulting from particle morphology and particle surface

<sup>10)</sup> Lubricants in Pharmaceutical Solid Dosage Forms, Jinjiang Li and Yongmei Wu, Lubricants 2014, 2, 21-43

<sup>10</sup> Lubricating properties of triacylgycerols related to the release of medicaments: M. Vitkova; M. Chalabala; J. Rak, Faculty of Pharmacy, 832-32, Bratislava, Czecho-Slovakia

12) A. Stamm, Sci. Techn. Pharm. 9 (1), 471 - 478 (1980)



# **LIPIDS FOR SOLID DOSAGE** FORMS

### **PHARMA**

# YOUR FORMULATION CHALLENGES – OUR SOLUTIONS

According to the Biopharmaceutical Classification System (BCS), more than 70% of the APIs used in pharmaceutical preparations are difficult to formulate due to poor solubility and/or permeabilities, and the majority of drug candidates have similar characteristics. Formulating these APIs into stable forms and achieving satisfactory active delivery and dissolution profiles is a key challenge for galenic scientists.

IOI Oleo PHARMA offers a variety of vegetable-derived mono-, di- and triglycerides for a wide range of dosage forms such as tablets, capsules, granules or powders. The lipid nature of our products gives them an inherent compatibility towards these problematic API categories and can help to solve the galenic formulation hurdles. Most of our excipients are multifunctional and suitable for various drug delivery systems. Adopted dosing is a trajectory to accentuate certain characteristics and functionalities depending on the specific formulation task.

Technologies and techniques such as SEDDS, SMEDDS, SNEDDS, SLN, Hot Melt Coating (taste masking), Hot Melt Extrusion or Hot Melt Granulation can be applied to enable delivery or enhance the dissolution profile of poorly soluble drugs.

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#### PRODUCT **CHEMICAL** MONO-MELTING APPLICATIONS DESCRIPTION POINT °C GRAPH 55 - 58 DYNASAN® 114 Trimyristin Lubricants for tablets & capsules Lipophilic matrices for fat-soluble actives DYNASAN® 116 Tripalmitin 63 - 68 \_ Taste masking<sup>1) 2) 3) 4) 5) 6) 7)</sup> Release profile modification in oral solid dosage DYNASAN® 118 Glyceryl Tristearate USP-NF 69 - 73 matrices and implants Body imparting & structure forming in semi-solids Crystallization accelerators and seeding agents in sticks, ovules & suppositories Matrix component for SLN $^{1),\,\,2)\,\,4),\,\,5)}$ IMWITOR® 900 (F) P, Glycerol Monostearate Ph. Eur. 54 - 64 Solid partial glycerides with surface-active properties IMWITOR® 900 K 40-55% Type I & II USP-NF Lipophilic matrices for oral solid dosage forms Granulation, Hot Melt Techniques Tablet lubricant Matrix component for SLN<sup>6)</sup> Emulsion stabilizer Dispersing agent for pigments 42 - 44 WITEPSOL® E 85 Hard Fat Ph. Eur. Lipophilic matrix for use in bio-degradable subcuta-USP-NF neous implants<sup>8)</sup> SOFTISAN® 154 53 - 58 Lipophilic matrix for SLN for controlled drug release<sup>9)</sup> Hydrogenated Palm Oil \_

<sup>1)</sup> Lubricating properties of triacylglycerols related to the release of medicaments: M. Vitkova; M. Chalabala; J. Rak, Faculty of Pharmacy, 832-32, Bratislava, Czecho-Slovakia.

<sup>27</sup> Taste masked lipid pellets with enhanced release of hydrophobic active ingredient: Vaassen, Jonathan; Bartscher, Kathrin; Breitkreutz, Joerg, International Journal of Pharmaceutics (Amsterdam, Netherlands) (2012), 429(1-2), 99-103.

<sup>3</sup> Paclitaxels-containing nanometer liposome for injection and the preparation method thereof: Li, Shubin; Liu, Dan; Song, Jian; Bao, Jie; Ning, Hong; Li, Minghui; Guo, Zhifu; Wang, Hong; Yu, Na; Li, Lei, Faming Zhuanli Shenqing Gongkai Shuomingshu (2009), CN 101366697 A 20090218.

<sup>4)</sup> Active ingredient matrix dosage forms: Francas, Gernot; Przyklenk, Karl-Heinz, Ger. Offen. (2012), DE 102011051304 A1 20121227.

<sup>S)</sup> Optimization of 5-Fluorouracil solid lipid nano-particles: a preliminary study to treat colon cancer: Yassin, Alaa Eldeen B.; Khalid Anwer, Md.; Mowafy, Hammam A.; El-Ba- gory, Ibrahim M.; Bayomi, Mohsen A.; Alsarra, Ibrahim A., International Journal of Medical Sciences (2010), 7(6), 398-408. <sup>6)</sup> New approaches in nanoparticulate drug delivery system - a review: Rakesh Bagul; Vjay Mahajan; Avinash Dhake, International Journal of Current Pharmaceutical Research, Vol 4, Issue 3 (2012).

 $^\eta$  Development of biodegradable implants for extended delivery of proteins, Inaugural Dissertation, Luisa Fernanda Duque Zapata, Freie Universität Berlin

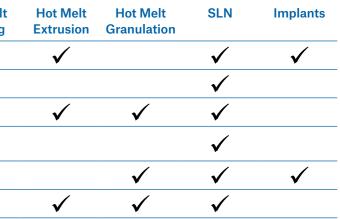
Extruded depot form for sustained release of active ingredient, patent DE102017106216A1, 2017, Alexandra Partenhauser, AMW GmbH

<sup>9)</sup> Process, Optimization, and Characterization of Budesonide Loaded Nanostructured Lipid Carriers for the Treatment of Inflammatory Bowel Disease, Gurpreet Kaur Sinhmar et al., 2005

### **Application Survey of Solid Dosage Forms**

PRODUCT	Lubricant	Taste Masking	Hot Melt Coating
DYNASAN® 114	$\checkmark$		
DYNASAN® 116	$\checkmark$	$\checkmark$	$\checkmark$
DYNASAN® 118	$\checkmark$	$\checkmark$	$\checkmark$
IMWITOR <sup>®</sup> 900 (F) P IMWITOR <sup>®</sup> 900 K	$\checkmark$		
WITEPSOL® E 85		$\checkmark$	$\checkmark$
SOFTISAN® 154		$\checkmark$	$\checkmark$





Identified uses in commercial products and literature

### Approved Ingredients:

- Vitamin preparations
- Expectorants (e.g. ACC/ NAC)
- Antiacidics (e.g. Hydrotalcite)
- Antihelmintics (e.g. Moxidectin)
- Non-steroidal anti-inflammatory drugs (e.g. ASA)
- Non-opioid Analgesics (e.g. Paracetamol)