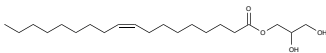
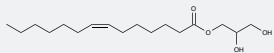
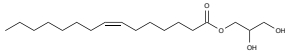
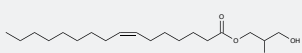
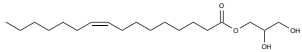
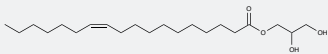
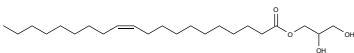


Crystallization in the Lipidic Cubic Phase (LCP) has evolved into an important method for crystallization of membrane proteins, with the lipid Monoolein being the first choice to create a stable LCP. In the past few years however, the short lipids 7.7 MAG<sup>[4,7]</sup>, 7.8 MAG<sup>[6-8]</sup>, 7.9 MAG<sup>[5,7]</sup> have become increasingly popular and the recent progress in applying the highly viscous lipidic cubic phase for serial femtosecond crystallography (LCP-SFX) <sup>[1-3]</sup> further accelerates the success of the LCP method by

- › delivering the crystal-loaded viscous LCP directly into the XFEL beam (thereby reducing sample consumption in comparison to liquid injectors),
- › taking advantage of the inherently small crystals grown in LCP,
- › avoiding tricky crystal mounting from LCP.

LCP Lipid	Lipid structure	Cat-No.	Amount
<b>Monoolein 9.9 MAG</b>		X-LCP-101	1 g
<b>7.7 MAG</b>		X-LCP-105	100 mg
<b>7.8 MAG</b>		X-LCP-106	100 mg
<b>7.9 MAG</b> Stable at low temperatures <sup>[9]</sup>		X-LCP-107	100 mg
<b>Monopalmitolein 9.7 MAG</b>		X-LCP-102	1 g
<b>Monovaccenin 11.7 MAG</b>		X-LCP-103	100 mg
<b>Monoeicosenoin 11.9 MAG</b>		X-LCP-104	1 g

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