

Figure S1. An extracted ion chromatogram of free cholesterol (Chl) and cholesteryl esters (CE) of a representative wild type mouse tarsal plate extract. A common analytical ion m/z 369.35 was used to detect the analytes. The following assignments for CE were made based on the m/z values and fragmentation patterns of their (M + H)+, (M + K)+ and (M + Na)+ adducts: C10.0 (1); C12.0- and C14.1-CE (2); C14.0 and C16.1 (3); C15.0 (4); C16.0- and C18.1-CE (5); C17.0-CE (6); C18.0- and C20.1-CE (7); C19.0-CE (8); C20.0- and C22.1-CE (9); C21.0-CE (10); C22.0- and C24.1-CE (11); C23.0-CE (12); C24.0- and C26.1-CE (13); C25.0-CE (14); C26.0- and C28.1-CE (15); C27.0-CE (16); C28.0- and C30.1-CE (17); C29.0 (18); C30.0- and C32.1-CE (19); C34.1-CE (20); cholesteryl esters of (O)-acyl-ω-hydroxy fatty acids (20-22).

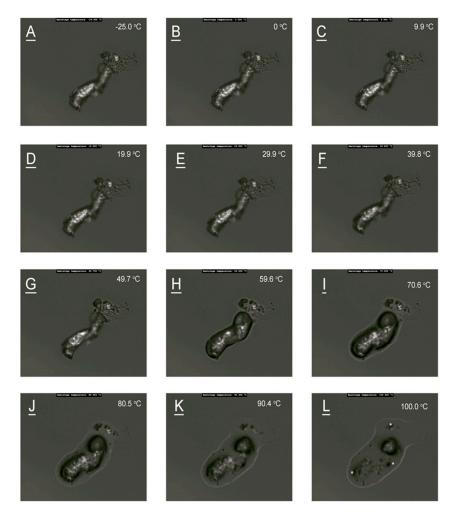


Figure S2. Melting of a *Soat1*-null meibum sample as recorded during a HSPLM experiment. Sample temperatures rose from -25 °C to +100 °C with a rate of 2 °C/min. Bright structures visible within the meibum sample are birefringent lipid aggregates. Note that some aggregates did not melt even at the highest temperature. Dark, non-melting, non-birefringent aggregates of presumably proteinaceous nature are labeled with white asterisks in Panel L.