**Supporting Information**

**Enhancement of the anticancer effect of atorvastatin-loaded nanoemulsions by improving oral absorption via multivalent intestinal transporter-targeting lipids**

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**Running head:** Enhanced oral absorption of atorvastatin for chemotherapy

**Table S1.** Inhibitors and concentrations used in the transport study, listed with their functions.

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| Inhibitor | Concentration | Function |
| Act D | 3.2 μM | Inhibitor of ASBT-mediated transport |
| CFZ | 10 μM | Inhibitor of OSTα/β blocking the transport of bile acid across the basolateral membrane |
| PA | 0.2 mM | Inhibition of SMVT-mediated transport |
| Chlorpromazine | 32 μM | Inhibitor of clathrin-mediated endocytosis |
| MBCD | 10 mM | Inhibitor of caveola/lipid raft-mediated endocytosis (cholesterol depletion) |
| Genistein | 0.1 mM | Inhibitor of caveola/lipid raft-mediated endocytosis (broad inhibitor of protein tyrosine kinase) |
| Amiloride | 0.1 mM | Inhibitor of macropinocytosis |
| Brefeldin A | 90 μM | Inhibitor of ER/Golgi pathway |
| Cys A | 10 µM | Inhibitor of P-gp-mediated efflux |

Act D, actinomycin D; ASBT, apical sodium-dependent bile acid transporter; CFZ, clofazimine; OSTα/β, organic solute transporter α and β; PA, pantothenic acid; SMVT, sodium-dependent multivitamin transporter; MBCD, methyl-β-cyclodextrin; ER, endoplasmic reticulum; Cys A, cyclosporine A; P-gp, P-glycoprotein.