

Supplementary Materials: A Dry Powder Platform for Nose-To-Brain Delivery of Dexamethasone: Formulation Development and Nasal Deposition Studies

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Table S1. Parameters and their levels included in experimental design.

Formulation Parameters	Low (-1)	Medium (0)	High (+1)
DSP concentration, DSP (%; <i>w/w</i>)	0.02	0.11	0.2
Hypromellose concentration, HPMC (%; <i>w/w</i>)	0.2	0.6	1.0
Process parameters	Low (-1)	Medium (0)	High (+1)
Inlet air temperature, T_{inlet} (°C)	120	140	160
Feed flow rate, FFR (g min ⁻¹)	2.5	3.5	4.5

Table S2. Validation parameters of HPLC method employed for DSP and DB analysis.

Range of linearity						
DSP	1–15 µg/mL					
DB	1–15 µg/mL					
Linearity						
	regression equation*			correlation coefficient		
DSP	y = 35033x – 18226			0.998		
DB	y = 52032x - 12208			0.999		
Precision Data						
	repeatability (RSD %)			intermediate precision (RSD %)		
	low	medium	high	low	medium	high
	2 µg/mL	7.5 µg/mL	15 µg/mL	2 µg/mL	7.5 µg/mL	15 µg/mL
DSP	1.2	1.3	1.2	1.5	0.9	0.8
DB	1.3	1.6	1.0	1.2	1.4	1.1
Accuracy Data						
	accuracy (recovery, mean (%) ± RSD, <i>n</i> = 3)					
	low	medium		high		
	(2 µg/mL)	(7.5 µg/mL)		(15 µg/mL)		
DSP	103.0 ± 0.6	100.0 ± 0.5		99.8 ± 0.2		
DB	99.5 ± 1.3	96.2 ± 1.2		100.4 ± 0.8		
Sensitivity						
	LOD (µg/mL)			LOQ (µg/mL)		
DSP	0.65			1.98		
DB	0.63			1.91		

*x—concentration (µg/mL); y—Area under Curve (µV s); All measurements were performed in triplicate.

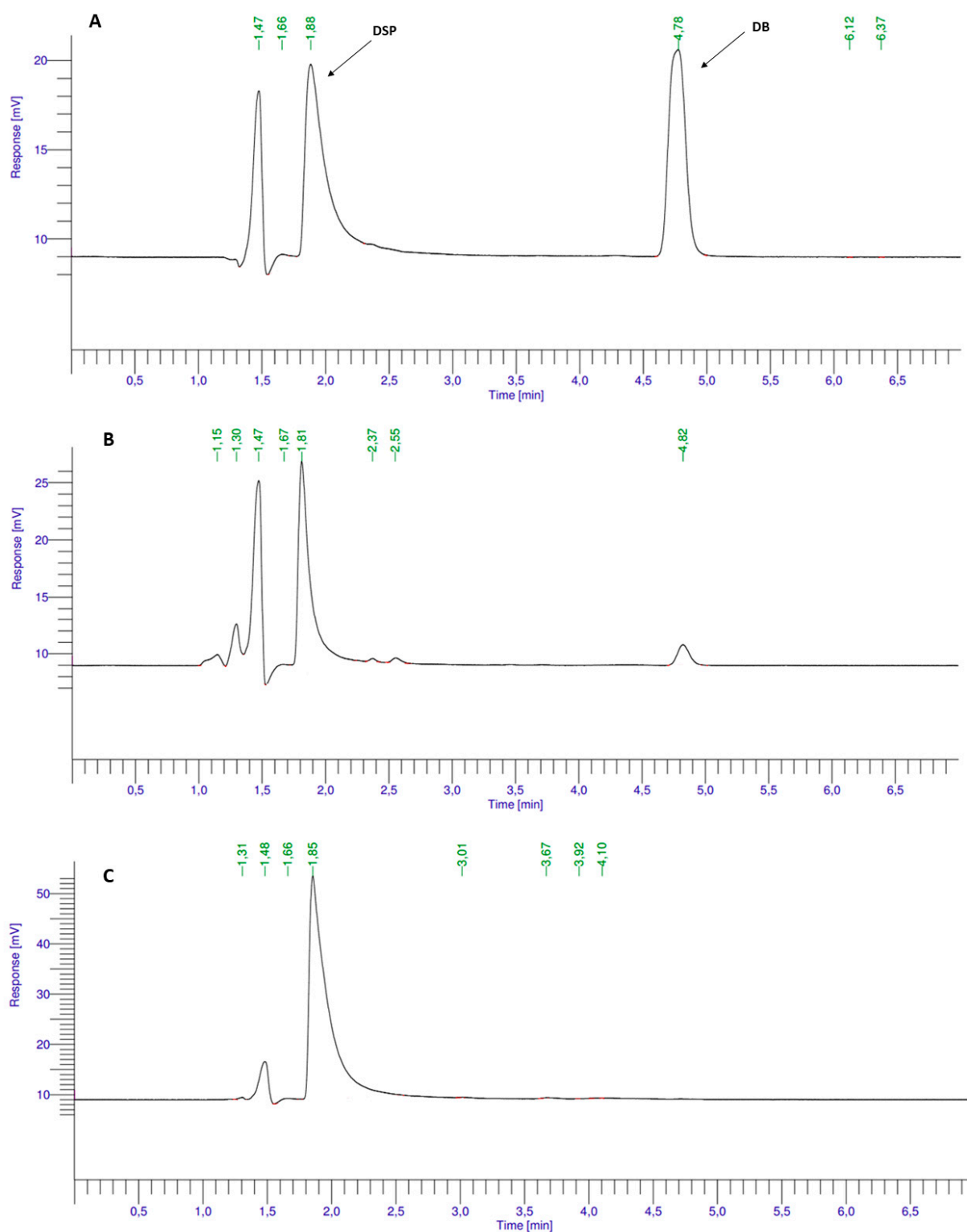


Figure S1. Chromatogram of (A) standard dexamethasone sodium phosphate (DSP) and dexamethasone base (DB) solution (both at $2 \mu\text{g mL}^{-1}$), (B) sample of receptor medium taken during DSP permeability study from DSP powder formulation across Calu-3 cell monolayer, (C) eluate obtained by rinsing of the nasal cavity segment within nasal deposition studies of DSP powder formulation. All chromatograms were recorded at wavelength of 241 nm.