

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	An investigation into the use of polymer blends to improve the printability of and regulate drug release from pharmaceutical solid dispersions prepared via fused deposition modeling (FDM) 3D printing. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 108, 111-125.	4.3	208
2	Characterisation of solid dispersions of paracetamol and EUDRAGIT® E prepared by hot-melt extrusion using thermal, microthermal and spectroscopic analysis. International Journal of Pharmaceutics, 2008, 354, 158-167.	5.2	131
3	Development of a Simple Mechanical Screening Method for Predicting the Feedability of a Pharmaceutical FDM 3D Printing Filament. Pharmaceutical Research, 2018, 35, 151.	3.5	111
4	Physicochemical Properties of the Amorphous Drug, Cast Films, and Spray Dried Powders to Predict Formulation Probability of Success for Solid Dispersions: Etravirine. Journal of Pharmaceutical Sciences, 2011, 100, 260-274.	3.3	106
5	Sulforaphane Mediates Glutathione Depletion via Polymeric Nanoparticles to Restore Cisplatin Chemosensitivity. ACS Nano, 2019, 13, 13445-13455.	14.6	106
6	Electrospun Polymer Blend Nanofibers for Tunable Drug Delivery: The Role of Transformative Phase Separation on Controlling the Release Rate. Molecular Pharmaceutics, 2016, 13, 25-39.	4.6	84
7	Insights into the Role of Polymer-Surfactant Complexes in Drug Solubilisation/Stabilisation During Drug Release from Solid Dispersions. Pharmaceutical Research, 2013, 30, 290-302.	3.5	83
8	Characterisation and Prediction of Phase Separation in Hot-Melt Extruded Solid Dispersions: A Thermal, Microscopic and NMR Relaxometry Study. Pharmaceutical Research, 2010, 27, 1869-1883.	3.5	74
9	Microfluidics for pharmaceutical nanoparticle fabrication: The truth and the myth. International Journal of Pharmaceutics, 2020, 584, 119408.	5.2	72
10	An investigation into the effects of thermal history on the crystallisation behaviour of amorphous paracetamol. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 364-371.	4.3	66
11	Impact of Processing Parameters on the Quality of Pharmaceutical Solid Dosage Forms Produced by Fused Deposition Modeling (FDM). Pharmaceutics, 2019, 11, 633.	4.5	63
12	Recent developments in micro- and nanofabrication techniques for the preparation of amorphous pharmaceutical dosage forms. Advanced Drug Delivery Reviews, 2016, 100, 67-84.	13.7	60
13	Controlled Release from Zein Matrices: Interplay of Drug Hydrophobicity and pH. Pharmaceutical Research, 2016, 33, 673-685.	3.5	58
14	Early Stage Phase Separation in Pharmaceutical Solid Dispersion Thin Films under High Humidity: Improved Spatial Understanding Using Probe-Based Thermal and Spectroscopic Nanocharacterization Methods. Molecular Pharmaceutics, 2013, 10, 918-930.	4.6	56
15	An investigation into the crystallisation behaviour of an amorphous cryomilled pharmaceutical material above and below the glass transition temperature. Journal of Pharmaceutical Sciences, 2010, 99, 196-208.	3.3	54
16	Solid lipid microparticles produced by spray congealing: Influence of the atomizer on microparticle characteristics and mathematical modeling of the drug release. Journal of Pharmaceutical Sciences, 2010, 99, 916-931.	3.3	54
17	Compositional Analysis of Low Quantities of Phase Separation in Hot-Melt-Extruded Solid Dispersions: A Combined Atomic Force Microscopy, Photothermal Fourier-Transform Infrared Microspectroscopy, and Localised Thermal Analysis Approach. Pharmaceutical Research, 2011, 28, 2311-2326.	3.5	51
18	Stabilisation of amorphous drugs under high humidity using pharmaceutical thin films. European Journal of Pharmaceutics and Biopharmaceutics, 2013, 84, 555-565.	4.3	44

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19	Microstructure of an Immiscible Polymer Blend and Its Stabilization Effect on Amorphous Solid Dispersions. Molecular Pharmaceutics, 2013, 10, 2767-2780.	4.6	44
20	An investigation into the mechanism of dissolution rate enhancement of poorly water-soluble drugs from spray chilled gelucire 50/13 microspheres. Journal of Pharmaceutical Sciences, 2010, 99, 262-274.	3.3	43
21	Reversing tumor stemness via orally targeted nanoparticles achieves efficient colon cancer treatment. Biomaterials, 2019, 216, 119247.	11.4	43
22	An investigation into the influence of drug–polymer interactions on the miscibility, processability and structure of polyvinylpyrrolidone-based hot melt extrusion formulations. International Journal of Pharmaceutics, 2015, 496, 95-106.	5.2	36
23	Hot melt extruded transdermal films based on amorphous solid dispersions in Eudragit RS PO: The inclusion of hydrophilic additives to develop moisture-activated release systems. International Journal of Pharmaceutics, 2016, 514, 270-281.	5.2	35
24	Suppression of Tumor Energy Supply by Liposomal Nanoparticle-Mediated Inhibition of Aerobic Glycolysis. ACS Applied Materials & Interfaces, 2018, 10, 2347-2353.	8.0	35
25	The development of microthermal analysis and photothermal microspectroscopy as novel approaches to drug–excipient compatibility studies. International Journal of Pharmaceutics, 2008, 354, 149-157.	5.2	32
26	A New Low Melting-Point Polymorph of Fenofibrate Prepared via Talc Induced Heterogeneous Nucleation. Crystal Growth and Design, 2015, 15, 5011-5020.	3.0	30
27	Creating Drug Solubilization Compartments via Phase Separation in Multicomponent Buccal Patches Prepared by Direct Hot Melt Extrusion–Injection Molding. Molecular Pharmaceutics, 2015, 12, 4349-4362.	4.6	30
28	Physical stabilization of low-molecular-weight amorphous drugs in the solid state: a material science approach. Therapeutic Delivery, 2014, 5, 817-841.	2.2	29
29	Paracetamol·loaded poly(ε-caprolactone) layered silicate nanocomposites prepared using hot-melt extrusion. Journal of Pharmaceutical Sciences, 2009, 98, 4831-4843.	3.3	26
30	An Investigation into the Mechanisms of Drug Release From Taste-Masking Fatty Acid Microspheres. Journal of Pharmaceutical Sciences, 2008, 97, 3842-3854.	3.3	25
31	An Investigation into the Interaction between Taste Masking Fatty Acid Microspheres and Alkaline Buffer using Thermal and Spectroscopic Analysis. Journal of Pharmaceutical Sciences, 2006, 95, 1022-1028.	3.3	23
32	The effect of processing on the surface physical stability of amorphous solid dispersions. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 88, 897-908.	4.3	22
33	Effects of porosity on drug release kinetics of swellable and erodible porous pharmaceutical solid dosage forms fabricated by hot melt droplet deposition 3D printing. International Journal of Pharmaceutics, 2021, 604, 120626.	5.2	21
34	Characterization of Heterogeneity and Spatial Distribution of Phases in Complex Solid Dispersions by Thermal Analysis by Structural Characterization and X-ray Micro Computed Tomography. Pharmaceutical Research, 2017, 34, 971-989.	3.5	20
35	Probing the molecular interactions between pharmaceutical polymeric carriers and bile salts in simulated gastrointestinal fluids using NMR spectroscopy. Journal of Colloid and Interface Science, 2019, 551, 147-154.	9.4	20
36	Molecular Implications of Drug–Polymer Solubility in Understanding the Destabilization of Solid Dispersions by Milling. Molecular Pharmaceutics, 2014, 11, 2453-2465.	4.6	19

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37	The Development of Direct Extrusion-Injection Moulded Zein Matrices as Novel Oral Controlled Drug Delivery Systems. Pharmaceutical Research, 2015, 32, 2775-86.	3.5	19
38	Preparation and Characterization of Ultrarapidly Dissolving Orodispersible Films for Treating and Preventing lodine Deficiency in the Pediatric Population. Journal of Agricultural and Food Chemistry, 2015, 63, 9831-9838.	5.2	19
39	A SPION-eicosane protective coating for water soluble capsules: Evidence for on-demand drug release triggered by magnetic hyperthermia. Scientific Reports, 2016, 6, 20271.	3.3	19
40	Spatial Characterization of Hot Melt Extruded Dispersion Systems Using Thermal Atomic Force Microscopy Methods: The Effects of Processing Parameters on Phase Separation. Pharmaceutical Research, 2014, 31, 1744-1752.	3.5	18
41	Thermal Analysis by Structural Characterization as a Method for Assessing Heterogeneity in Complex Solid Pharmaceutical Dosage Forms. Analytical Chemistry, 2015, 87, 10848-10855.	6.5	18
42	Characterization of Hydrophilic Polymers as a Syringe Extrusion 3D Printing Material for Orodispersible Film. Polymers, 2021, 13, 3454.	4.5	18
43	Suppression of the coffee-ring effect by tailoring the viscosity of pharmaceutical sessile drops. Colloids and Surfaces A: Physicochemical and Engineering Aspects, 2021, 614, 126144.	4.7	17
44	Evaluation of the Benefits of Microfluidic-Assisted Preparation of Polymeric Nanoparticles for DNA Delivery. Materials Science and Engineering C, 2021, 127, 112243.	7.3	17
45	An investigation into the effects of geometric scaling and pore structure on drug dose and release of 3D printed solid dosage forms. European Journal of Pharmaceutics and Biopharmaceutics, 2022, 177, 113-125.	4.3	17
46	â€~Tablet-in-Syringe': A Novel Dosing Mechanism for Dysphagic Patients Containing Fast-Disintegrating Tablets Fabricated Using Semisolid Extrusion 3D Printing. Pharmaceutics, 2022, 14, 443.	4.5	16
47	On-demand, magnetic hyperthermia-triggered drug delivery: optimisation for the GI tract. Journal of Materials Chemistry B, 2016, 4, 1704-1711.	5.8	15
48	Preparation and Characterization of an Electrospun PLA-Cyclodextrins Composite for Simultaneous High-Efficiency PM and VOC Removal. Journal of Composites Science, 2020, 4, 79.	3.0	15
49	Lipid based intramuscular long-acting injectables: Current state of the art. European Journal of Pharmaceutical Sciences, 2022, 178, 106253.	4.0	15
50	Improvement of the in vitro safety profile and cytoprotective efficacy of amifostine against chemotherapy by PEGylation strategy. Biochemical Pharmacology, 2016, 108, 11-21.	4.4	14
51	An investigation into the use of low quantities of functional additives to control drug release from hot melt extruded solid dispersions for poorly soluble drug delivery. International Journal of Pharmaceutics, 2020, 579, 119172.	5.2	14
52	An investigation into the formations of the internal microstructures of solid dispersions prepared by hot melt extrusion. European Journal of Pharmaceutics and Biopharmaceutics, 2020, 155, 147-161.	4.3	13
53	The Development of Modulated, Quasi-Isothermal and Ultraslow Thermal Methods as a Means of Characterizing the α to γ Indomethacin Polymorphic Transformation. Molecular Pharmaceutics, 2012, 9, 1087-1099.	4.6	12
54	Novel Thermal Imaging Method for Rapid Screening of Drug–Polymer Miscibility for Solid Dispersion Based Formulation Development. Molecular Pharmaceutics, 2018, 15, 5625-5636.	4.6	12

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55	Molecular Indicators of Surface and Bulk Instability of Hot Melt Extruded Amorphous Solid Dispersions. Pharmaceutical Research, 2015, 32, 1210-1228.	3.5	10
56	Hot melt extruded zein for controlled delivery of diclofenac sodium: Effect of drug loading and medium composition. International Journal of Pharmaceutics, 2020, 585, 119503.	5.2	10
57	Drop-on-demand printing of personalised orodispersible films fabricated by precision micro-dispensing. International Journal of Pharmaceutics, 2021, 610, 121279.	5.2	10
58	Towards controlling the crystallisation behaviour of fenofibrate melt: triggers of crystallisation and polymorphic transformation. RSC Advances, 2018, 8, 13513-13525.	3.6	8
59	The Use of Quasi-Isothermal Modulated Temperature Differential Scanning Calorimetry for the Characterization of Slow Crystallization Processes in Lipid-Based Solid Self-Emulsifying Systems. Pharmaceutical Research, 2015, 32, 1316-1324.	3.5	7
60	The use of polymer blends to improve stability and performance of electrospun solid dispersions: The role of miscibility and phase separation. International Journal of Pharmaceutics, 2021, 602, 120637.	5.2	7
61	Direct Granule Feeding of Thermal Droplet Deposition 3D Printing of Porous Pharmaceutical Solid Dosage Forms Free of Plasticisers. Pharmaceutical Research, 2022, 39, 599-610.	3.5	7
62	The Effects of Solid Particle Containing Inks on the Printing Quality of Porous Pharmaceutical Structures Fabricated by 3D Semi-Solid Extrusion Printing. Pharmaceutical Research, 2022, 39, 1267-1279.	3.5	7
63	A Laser Imaging and Neutron Reflection Investigation Into the Monolayer Behaviour of Fatty Acids Used for Taste Masking Microspheres. Journal of Pharmaceutical Sciences, 2008, 97, 1864-1877.	3.3	6
64	A multi-technique characterization of the stability of surfactant containing solid dispersion based buccal patches prepared by hot melt injection moulding. International Journal of Pharmaceutics, 2017, 528, 547-562.	5.2	6
65	New insights into the effects of porosity, pore length, pore shape and pore alignment on drug release from extrusionbased additive manufactured pharmaceuticals. Additive Manufacturing, 2021, 46, 102196.	3.0	6
66	Disc-shaped polyoxyethylene glycol glycerides gel nanoparticles as novel protein delivery vehicles. International Journal of Pharmaceutics, 2015, 496, 1015-1025.	5.2	5
67	Nanostructural Analysis of Water Distribution in Hydrated Multicomponent Gels Using Thermal Analysis and NMR Relaxometry. Molecular Pharmaceutics, 2015, 12, 2068-2079.	4.6	4
68	Thermal Analysis of Pharmaceuticals. Advances in Delivery Science and Technology, 2016, , 363-387.	0.4	3
69	Automation Potential of a New, Rapid, Microscopy-Based Method for Screening Drug–Polymer Solubility. ACS Omega, 2020, 5, 11402-11410.	3.5	3
70	Emerging molecular mechanisms and genetic targets for developing novel therapeutic strategies for treating bladder diseases. European Journal of Pharmaceutical Sciences, 2022, 173, 106167.	4.0	3
71	Moisture Uptake of Polyoxyethylene Glycol Glycerides Used as Matrices for Drug Delivery: Kinetic Modelling and Practical Implications. Pharmaceutical Research, 2013, 30, 1123-1136.	3.5	2
72	Precision coating of ocular devices/contact lenses by nanoelectrospray additive printing. Materials and Design, 2022, 219, 110782.	7.0	1