List of Publications by Year in descending order

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ΔΜΟΙΤ ΡΛΙΙΠΕΙ

#	Article	IF	CITATIONS
1	Manufacturing of solid dispersions of poorly water soluble drugs by spray drying: Formulation and process considerations. International Journal of Pharmaceutics, 2013, 453, 253-284.	2.6	442
2	Raman spectroscopy in pharmaceutical product design. Advanced Drug Delivery Reviews, 2015, 89, 3-20.	6.6	221
3	Theoretical and Experimental Investigation on the Solid Solubility and Miscibility of Naproxen in Poly(vinylpyrrolidone). Molecular Pharmaceutics, 2010, 7, 1133-1148.	2.3	136
4	Influence of Preparation Methods on Solid State Supersaturation of Amorphous Solid Dispersions: A Case Study with Itraconazole and Eudragit E100. Pharmaceutical Research, 2010, 27, 775-785.	1.7	115
5	Structural and Dynamic Properties of Amorphous Solid Dispersions: The Role of Solid-State Nuclear Magnetic Resonance Spectroscopy and Relaxometry. Journal of Pharmaceutical Sciences, 2014, 103, 2635-2662.	1.6	103
6	Influence of Solvent Composition on the Miscibility and Physical Stability of Naproxen/PVP K 25 Solid Dispersions Prepared by Cosolvent Spray-Drying. Pharmaceutical Research, 2012, 29, 251-270.	1.7	84
7	A Review of PAT Strategies in Secondary Solid Oral Dosage Manufacturing of Small Molecules. Journal of Pharmaceutical Sciences, 2017, 106, 667-712.	1.6	72
8	Orodispersible films: Towards drug delivery in special populations. International Journal of Pharmaceutics, 2017, 523, 327-335.	2.6	70
9	Can compression induce demixing in amorphous solid dispersions? A case study of naproxen–PVP K25. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 81, 207-213.	2.0	62
10	Advances in experimental and mechanistic computational models to understand pulmonary exposure to inhaled drugs. European Journal of Pharmaceutical Sciences, 2018, 113, 41-52.	1.9	57
11	Printing medicines as orodispersible dosage forms: Effect of substrate on the printed micro-structure. International Journal of Pharmaceutics, 2016, 509, 518-527.	2.6	52
12	Progress in spray-drying of protein pharmaceuticals: Literature analysis of trends in formulation and process attributes. Drying Technology, 2021, 39, 1415-1446.	1.7	49
13	Effect of Compression on Non-isothermal Crystallization Behaviour of Amorphous Indomethacin. Pharmaceutical Research, 2012, 29, 2489-2498.	1.7	41
14	Relating Hydrogen-Bonding Interactions with the Phase Behavior of Naproxen/PVP K 25 Solid Dispersions: Evaluation of Solution-Cast and Quench-Cooled Films. Molecular Pharmaceutics, 2012, 9, 3301-3317.	2.3	40
15	Searching for physiologically relevant in vitro dissolution techniques for orally inhaled drugs. International Journal of Pharmaceutics, 2019, 556, 45-56.	2.6	40
16	Characterization of degradation products of amorphous and polymorphic forms of clopidogrel bisulphate under solid state stress conditions. Journal of Pharmaceutical and Biomedical Analysis, 2010, 52, 332-344.	1.4	39
17	Carrier-based dry powder inhalation: Impact of carrier modification on capsule filling processability and in vitro aerodynamic performance. International Journal of Pharmaceutics, 2015, 491, 231-242.	2.6	37
18	An Investigation into the Effect of Spray Drying Temperature and Atomizing Conditions on Miscibility, Physical Stability, and Performance of Naproxen–PVP K 25 Solid Dispersions. Journal of Pharmaceutical Sciences, 2013, 102, 1249-1267.	1.6	36

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19	Key acceptability attributes of orodispersible films. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 125, 131-140.	2.0	33
20	The Influence of Relative Humidity and Storage Conditions on the Physico-chemical Properties of Inhalation Grade Fine Lactose. AAPS PharmSciTech, 2022, 23, 1.	1.5	33
21	Novel polyester-based thermoplastic elastomers for 3D-printed long-acting drug delivery applications. Journal of Controlled Release, 2021, 335, 290-305.	4.8	26
22	Performance indicators for carrier-based DPIs: Carrier surface properties for capsule filling and API properties for in vitro aerosolisation. International Journal of Pharmaceutics, 2018, 536, 326-335.	2.6	24
23	Can 3D printing of oral drugs help fight the current COVID-19 pandemic (and similar crisis in the) Tj ETQq1 1 0.	784314 rg 2.4	BT /Qverlock
24	Formulation performance and processability window for manufacturing a dual-polymer amorphous solid dispersion via hot-melt extrusion and strand pelletization. International Journal of Pharmaceutics, 2018, 553, 408-421.	2.6	22
25	Lyophilized protein powders: A review of analytical tools for root cause analysis of lot-to-lot variability. TrAC - Trends in Analytical Chemistry, 2016, 82, 468-491.	5.8	20
26	Relative Contributions of Solubility and Mobility to the Stability of Amorphous Solid Dispersions of Poorly Soluble Drugs: A Molecular Dynamics Simulation Study. Pharmaceutics, 2018, 10, 101.	2.0	20
27	Formulation and processability screening for the rational design of ethylene-vinyl acetate based intra-vaginal rings. International Journal of Pharmaceutics, 2019, 564, 90-97.	2.6	20
28	Assessment of Dry Powder Inhaler Carrier Targeted Design: A Comparative Case Study of Diverse Anomeric Compositions and Physical Properties of Lactose. Molecular Pharmaceutics, 2018, 15, 2827-2839.	2.3	18
29	Use of PBPK Modeling To Evaluate the Performance of Dissolv <i>lt</i> , a Biorelevant Dissolution Assay for Orally Inhaled Drug Products. Molecular Pharmaceutics, 2019, 16, 1245-1254.	2.3	18
30	Developing HME-Based Drug Products Using Emerging Science: a Fast-Track Roadmap from Concept to Clinical Batch. AAPS PharmSciTech, 2020, 21, 176.	1.5	18
31	Tribo-Charging Behaviour of Inhalable Mannitol Blends with Salbutamol Sulphate. Pharmaceutical Research, 2019, 36, 80.	1.7	17
32	Controlled-Release from High-Loaded Reservoir-Type Systems—A Case Study of Ethylene-Vinyl Acetate and Progesterone. Pharmaceutics, 2020, 12, 103.	2.0	17
33	The effect of material attributes and process parameters on the powder bed uniformity during a low-dose dosator capsule filling process. International Journal of Pharmaceutics, 2017, 516, 9-20.	2.6	16
34	PVP-H2O2 Complex as a New Stressor for the Accelerated Oxidation Study of Pharmaceutical Solids. Pharmaceutics, 2019, 11, 457.	2.0	16
35	Review of sensing technologies for measuring powder density variations during pharmaceutical solid dosage form manufacturing. TrAC - Trends in Analytical Chemistry, 2021, 135, 116147.	5.8	16
36	Impact of simulated lung fluid components on the solubility of inhaled drugs and predicted in vivo performance. International Journal of Pharmaceutics, 2021, 606, 120893.	2.6	16

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37	Drug–Excipient Interactions in the Solid State: The Role of Different Stress Factors. Molecular Pharmaceutics, 2017, 14, 4560-4571.	2.3	15
38	Polyelectrolyte–surfactant–complex nanoparticles as a delivery platform for poorly soluble drugs: A case study of ibuprofen loaded cetylpyridinium-alginate system. International Journal of Pharmaceutics, 2020, 580, 119199.	2.6	15
39	Multi-methodological investigation of the variability of the microstructure of HPMC hard capsules. International Journal of Pharmaceutics, 2016, 511, 840-854.	2.6	14
40	Establishment of a Molding Procedure to Facilitate Formulation Development for Co-extrudates. AAPS PharmSciTech, 2017, 18, 2971-2976.	1.5	14
41	Pharmaceutical-grade oral films as substrates for printed medicine. International Journal of Pharmaceutics, 2018, 547, 169-180.	2.6	14
42	Feeding of particle-based materials in continuous solid dosage manufacturing: a material science perspective. Drug Discovery Today, 2020, 25, 800-806.	3.2	14
43	How does secondary processing affect the physicochemical properties of inhalable salbutamol sulphate particles? A temporal investigation. International Journal of Pharmaceutics, 2017, 528, 416-428.	2.6	13
44	Analytical and Computational Methods for the Determination of Drug-Polymer Solubility and Miscibility. Molecular Pharmaceutics, 2021, 18, 2835-2866.	2.3	13
45	Understanding Concomitant Physical and Chemical Transformations of Simvastatin During Dry Ball Milling. AAPS PharmSciTech, 2020, 21, 152.	1.5	13
46	Focusing on powder processing in dry powder inhalation product development, manufacturing and performance. International Journal of Pharmaceutics, 2022, 614, 121445.	2.6	12
47	Evolution of the microstructure and the drug release upon annealing the drug loaded lipid-surfactant microspheres. European Journal of Pharmaceutical Sciences, 2020, 147, 105278.	1.9	11
48	Spherical agglomerates of lactose as potential carriers for inhalation. European Journal of Pharmaceutics and Biopharmaceutics, 2021, 159, 11-20.	2.0	11
49	Improving the granule strength of roller-compacted ibuprofen sodium for hot-melt coating processing. International Journal of Pharmaceutics, 2016, 510, 285-295.	2.6	10
50	Density fluctuations in amorphous pharmaceutical solids. Can SAXS help to predict stability?. Colloids and Surfaces B: Biointerfaces, 2018, 168, 76-82.	2.5	10
51	Solid-State Reactivity of Mechano-Activated Simvastatin: Atypical Relation to Powder Crystallinity. Journal of Pharmaceutical Sciences, 2019, 108, 3272-3280.	1.6	10
52	Quantitative Chemical Profiling of Commercial Glyceride Excipients via 1H NMR Spectroscopy. AAPS PharmSciTech, 2021, 22, 11.	1.5	10
53	Near-Infrared Hyperspectral Imaging as a Monitoring Tool for On-Demand Manufacturing of Inkjet-Printed Formulations. AAPS PharmSciTech, 2021, 22, 211.	1.5	10
54	Structural Characterization of Amorphous Solid Dispersions. Advances in Delivery Science and Technology, 2014, , 421-485.	0.4	9

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55	Insights into the processability and performance of adhesive blends of inhalable jet-milled and spray dried salbutamol sulphate at different drug loads. Journal of Drug Delivery Science and Technology, 2018, 48, 466-477.	1.4	9
56	Study of a low-dose capsule filling process by dynamic and static tests for advanced process understanding. International Journal of Pharmaceutics, 2018, 540, 22-30.	2.6	7
57	Insights into DPI sensitivity to humidity: An integrated in-vitro-in-silico risk-assessment. Journal of Drug Delivery Science and Technology, 2019, 52, 803-817.	1.4	7
58	Feasibility of rapidly assessing reactive impurities mediated excipient incompatibility using a new method: A case study of famotidine-PEG system. Journal of Pharmaceutical and Biomedical Analysis, 2020, 178, 112893.	1.4	7
59	Novel Cleaning-in-Place Strategies for Pharmaceutical Hot Melt Extrusion. Pharmaceutics, 2020, 12, 588.	2.0	7
60	Investigation into powder tribo-charging of pharmaceuticals. Part I: Process-induced charge via twin-screw feeding. International Journal of Pharmaceutics, 2020, 591, 120014.	2.6	7
61	Understanding Carrier Performance in Low-Dose Dry Powder Inhalation: An In Vitro–In Silico Approach. Pharmaceutics, 2021, 13, 297.	2.0	7
62	Evaluation of the Physico-mechanical Properties and Electrostatic Charging Behavior of Different Capsule Types for Inhalation Under Distinct Environmental Conditions. AAPS PharmSciTech, 2020, 21, 128.	1.5	7
63	Topologically directed confocal Raman imaging (TD-CRI): Advanced Raman imaging towards compositional and micromeritic profiling of a commercial tablet components. Journal of Pharmaceutical and Biomedical Analysis, 2022, 210, 114581.	1.4	7
64	Forced Solid-State Oxidation Studies of Nifedipine-PVP Amorphous Solid Dispersion. Molecular Pharmaceutics, 2022, 19, 568-583.	2.3	7
65	Continuous low-dose feeding of highly active pharmaceutical ingredients in hot-melt extrusion. Drug Development and Industrial Pharmacy, 2016, 42, 1360-1364.	0.9	6
66	Interplay of Aging and Lot-to-Lot Variability on the Physical and Chemical Properties of Excipients: A Case Study of Mono- and Diglycerides. Molecular Pharmaceutics, 2021, 18, 862-877.	2.3	6
67	Spray-Congealing and Wet-Sieving as Alternative Processes for Engineering of Inhalation Carrier Particles: Comparison of Surface Properties, Blending and In Vitro Performance. Pharmaceutical Research, 2021, 38, 1107-1123.	1.7	6
68	Assessment of Diverse Solidâ^'State Accelerated Autoxidation Methods for Droperidol. Pharmaceutics, 2022, 14, 1114.	2.0	6
69	Can we predict trends in tribo-charging of pharmaceutical materials from first principles?. Powder Technology, 2019, 356, 892-898.	2.1	5
70	Investigation into powder tribo–charging of pharmaceuticals. Part II: Sensitivity to relative humidity. International Journal of Pharmaceutics, 2020, 591, 120015.	2.6	5
71	Towards an Understanding of the Adsorption of Vaporized Hydrogen Peroxide (VHP) Residues on Glass Vials After a VHP Decontamination Process Using a Miniaturized Tool. Journal of Pharmaceutical Sciences, 2020, 109, 2454-2463.	1.6	5
72	Towards predicting the product quality in hot-melt extrusion: Small scale extrusion. International Journal of Pharmaceutics: X, 2020, 2, 100062.	1.2	4

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73	High-Molecular-Weight Hypromellose from Three Different Suppliers: Effects of Compression Speed, Tableting Equipment, and Moisture on the Compaction. AAPS PharmSciTech, 2020, 21, 203.	1.5	3
74	Towards predicting the product quality in hot-melt extrusion: Pilot plant scale extrusion. International Journal of Pharmaceutics: X, 2021, 3, 100084.	1.2	3
75	Phase Behavior of Drug–Lipid–Surfactant Ternary Systems toward Understanding the Annealing-Induced Change. Molecular Pharmaceutics, 2022, 19, 532-546.	2.3	3
76	Polyethylene oxide matrix tablet swelling evolution: The impact of molecular mass and tablet composition. Acta Pharmaceutica, 2021, 71, 215-243.	0.9	2
77	Insights into the Impact of Nanostructural Properties on Powder Tribocharging: The Case of Milled Salbutamol Sulfate. Molecular Pharmaceutics, 2022, 19, 547-557.	2.3	2
78	Development and Validation of a Stability-Indicating UPLC Method for the Determination of Hexoprenaline in Injectable Dosage Form Using AQbD Principles. Molecules, 2021, 26, 6597.	1.7	1
79	Quantitative chemical profiling of cellulose acetate excipient via 13C NMR spectroscopy in controlled release formulations. Journal of Pharmaceutical and Biomedical Analysis, 2022, 217, 114791.	1.4	1
80	A Tribute to Professor Saranjit Singh - A Critical Thinker, Innovator, Mentor, and Educator. Journal of Pharmaceutical Sciences, 2021, , .	1.6	0