Supporting Information

Drug-Loaded PLGA Electrospraying Porous Microspheres for the Local Therapy of Primary Lung Cancer via Pulmonary Delivery

Lifei Zhu,^{†,‡} Miao Li,[†] Xiaoyan Liu,[†] and Yiguang Jin^{*,†,‡}

[†]Department of Pharmaceutical Sciences, Beijing Institute of Radiation Medicine, 27 Taiping Road, Beijing 100850, China, e-mail address: jinyg@sina.com

[‡]Anhui Medical University, Hefei 230001, China

L.Z. and M.L. contributed equally to this manuscript.



2Theta (Coupled TwoTheta/Theta) WL=1.54060







2Theta (Coupled TwoTheta/Theta) WL=1.54060



Figure S1. X-ray diffraction (XRD) graphs of PLGA (A), raw oridonin powders (B), the physical mixture of them (C), and the oridonin-loaded PLGA EPMs (D).









Figure S2. Differential scanning calorimetric (DSC) graphs of PLGA (A), raw oridonin powders (B), the physical mixture of them (C), and the oridonin-loaded PLGA EPMs (D).









Figure S3. Infrared (IR) spetra of PLGA (A), raw oridonin powders (B), the physical mixture of them (C), and the oridonin-loaded PLGA EPMs (D).



Figure S4. *In vitro* distribution of non-porous oridonin-loaded solid PLGA microspheres and oridonin-loaded PLGA EPMs using the NGI. The stage numbers indicate the deposition site of lung. Higher numbers indicate the deeper deposition sites. Generally, Stages 2~8 indicate effective lung deposition.