

Comparison of cytotoxicity evaluation of anticancer drugs between real-time cell analysis and CCK-8 method

Ling Cai^{1#}, Xijiang Qin^{1#}, Zhihui Xu¹, Yiyan Song¹, Huijun Jiang²,
Yuan Wu³, Hongjie Ruan⁴, Jin Chen^{1,5,6*}

¹School of Public Health, Nanjing Medical University, Nanjing 211166, China

²School of Pharmacy, Nanjing Medical University, Nanjing 211166, China

³Department of Medical Oncology, Jiangsu Cancer Hospital, Jiangsu Institute of Cancer Research, The Affiliated Cancer Hospital of Nanjing Medical University, Nanjing 210009, China

⁴Department of gynecology, Obstetrics and Gynecology Hospital Affiliated to Nanjing Medical University, Nanjing, 210004, China

⁵The Key Laboratory of Modern Toxicology, Ministry of Education, School of Public Health, Nanjing Medical University, Nanjing 211166, Jiangsu, China

⁶Center for Global Health, School of Public Health, Nanjing Medical University, Nanjing 211166, Jiangsu, China

*Correspondence and request for materials should be addressed to J.C. (email: okachen30@gmail.com; jchen@njmu.edu.cn)

#Equal contribution

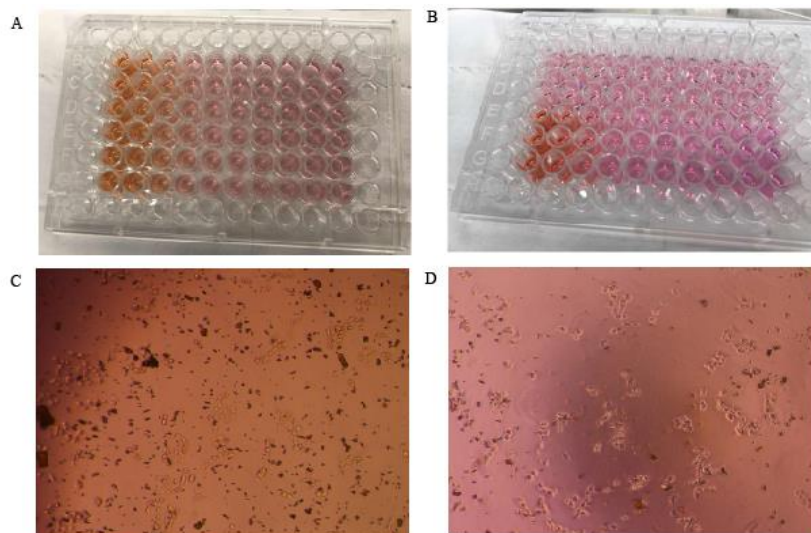


Figure S1 Images of HeLa cultured with curcumin. Photographs of the 96-well plate containing (A) HeLa cells after 24h-treatment of a series concentrations of curcumin and (B) curcumin-treated HeLa cells , of which row B, C and D were rinsed with fresh DMEM while row E, F and G remained unchanged. Images of the cell sample from the same 96-well plate containing 200 $\mu\text{g/mL}$ curcumin before (C) and after (D) replacing the culture medium.

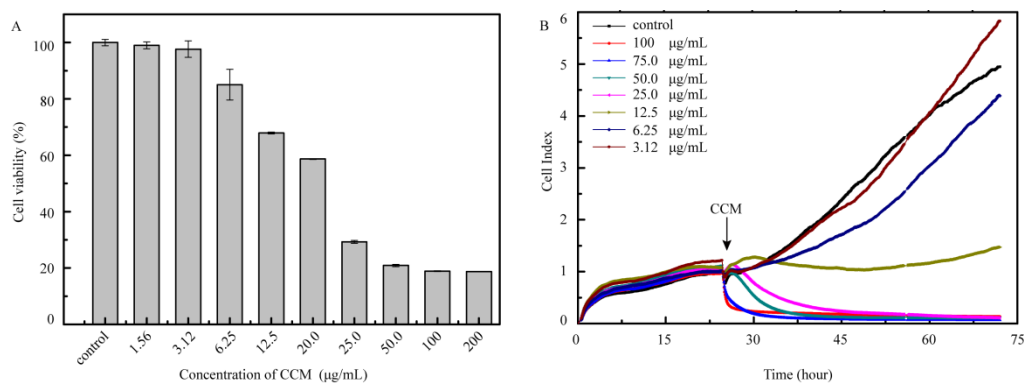


Figure S2 (A) Cell viability of A549 induced by CCM after 24 h in CCK-8 assay; (B) cell growth curve of A549 induced by CCM in RTCA.

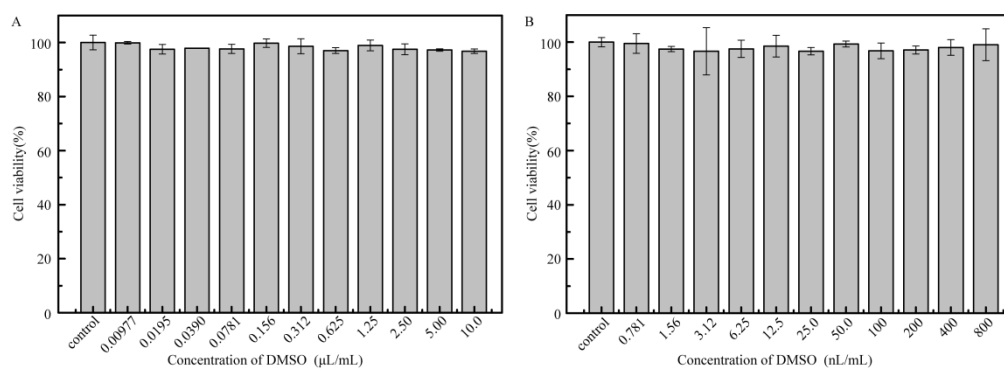


Figure S3 Cell viability of HeLa cells induced by DMSO after 24 h (A) and 48 h (B) in CCK-8 assay.