A Noteworthy Publication from our Faculty **Melanie T. Cushion, PhD** Senior Associate Dean for Research

<u>The celecoxib derivative AR-12 has broad spectrum</u> <u>antifungal activity in vitro and improves the activity of</u> <u>fluconazole in a murine model of cryptococcosis</u>



AR-12 inhibits fungal acetyl CoA synthetase in vitro and is fungicidal at concentrations similar to those achieved in human plasma. AR-12 has a broad spectrum of activity including active against yeasts; molds and dimorphic fungi with minimum inhibitory concentrations of 2-4 µg/mL. AR-12 is active against azoleand echinocandinresistant *Candida* isolates and subinhibitory AR-12 concentrations increase susceptibility of fluconazole-

and echinocandin-resistant*Candida* isolates. AR-12 increases the activity of fluconazole in a murine model of cryptococcosis.

Full Citation: Koselny K, Green J, DiDone L, Halterman JP, Fothergill AW, Wiederhold NP, Patterson TF, **Cushion MT**, Rappelye C, Wellington M, et al. <u>The celecoxib</u> <u>derivative AR-12 has broad spectrum antifungal activity in vitro and improves the</u> <u>activity of fluconazole in a murine model of</u> cryptococcosis. Antimicrob Agents Chemother. 2016 Sep 19. pii: AAC.01061-16.