U.S. Food and Drug Administration (USFDA), 2003. Guidance for Industry Bioavailability and Bioequivalence Studies for Orally Administered Drug Products — General Considerations. [online] available at: http://www.fda.gov/downloads/Drugs/.../Guidances/ucm070124.pdf [accessed 9 September 2012]

U.S. Food and Drug Administration (USFDA), 2012. The FDA's Drug Review Process: Ensuring Drugs Are Safe and Effective. [online] available at: http://www.fda.gov/drugs/resourcesforyou/consumers/ucm143534.htm [accessed 25 April 2013]

Williams, R.T. 1972. Progress report – hepatic metabolism of drugs. Gut, 13: 579 – 585.

Wu, Y., M. Hussain and R. Fassihi. 2005. Development of a simple analytical methodology for determination of glucosamine release from modified release matrix tablets. Journal of Pharmaceutical and Biomedical Analysis, 38: 263–269.