Study Design Objective

Diacerein will be administered to 6 subjects and the pharmacokinetic parameters will be assessed. This is followed by the oral administration of glucosamine to the same 6 subjects on a daily basis. Diacerein is then administered and the pharmacokinetic parameters assessed again to compare them with the initial pharmacokinetic findings (TRB Artrodar monograph, 1999). The objective of the present study conducted is to assess the significance of the change in the pharmacokinetic parameters in the 6 subjects post glucosamine administration.