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1. Share. I have EWQL Ministry of Rock and it was stunning. I use it everydayÂ .The present invention relates to a novel class of compounds that inhibit the enzyme cysteine protease and are thus useful in the treatment of diseases caused by viruses, bacteria, protozoa and helminths. Cysteine proteases are essential for the survival of all known living organisms. These enzymes have been identified in bacteria, yeast, plants, helminths and molds, and have been classified into five major families based on the amino acid sequence of the catalytic peptide. These include the cysteine protease family, the aspartic protease family, the metalloprotease family, the serine protease family, and the thiol protease family. Of the several hundred known cysteine proteases, the papain family is the best characterized. This family contains cysteine proteases with three distinct active sites, each of which catalyses the hydrolysis of a peptide bond containing a cysteine residue. Papain has been isolated from a number of plant species and will be discussed in detail hereinafter. The microbial cysteine proteases have been widely used in industry to catalyze the breakdown of proteins, peptides and other native proteins into convenient to use protein components. Although quite selective, the enzymes are not always completely specific, and the breakdown products from the catalysis are not always desirable. Certain protease inhibitors have been shown to have therapeutic effects in a variety of conditions, such as hypertension, myocardial infarction, reperfusion injury, haemorrhage, and inflammation (A. F. F. Woolley, Exp. Opin. Ther. Patents, 1994, 4, 455-476, and references cited therein). Clinical investigations of the protease inhibitor zQV0014 have shown it to be well tolerated and effective against acute pancreatitis (R. K. Valdix, Pancreatitis, 1996, 1, 217-220). One problem with inhibitors currently used in the clinic is that they are typically large molecules with poor pharmacokinetic properties. Furthermore, they are often not specific to a particular target, do not have the same spectrum of activity against different proteases, and often have toxicity problems. In light of the problems associated with the currently available protease inhibitors and the high therapeutic potential of the cysteine proteases, new, potent cysteine protease inhibitors,

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