

Novel gold(I)-based compounds with *in vitro* anticancer activity.

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Abstract:

As a symbol of purity and royalty, gold has accompanied the evolution of civilizations from the beginning of times and has been undoubtedly considered as one of the first metals used by humans. Historically, the importance of this element has been widely known for its applications in jewelry, artistic decoration, electronics or monetary usage. Conditioned by its metallic state, research on gold chemistry has given rise to innovative approaches in medicine and biology; for example, with the gold(I) complex auranofin in treatment of rheumatoid arthritis.¹ However, since the discovery of platinum compounds as anticancer agents in 1960s,² investigations on the efficacy of chemotherapeutic gold-based drugs have gained greater and greater attention. Over the past decades, several organometallic gold complexes have demonstrated potent cytotoxic activity against cancer cells.³ This presentation gives an overview of the design, chemical synthesis and *in vitro* anticancer activity evaluation of two novel classes of gold(I) complexes recently developed by our research group. The first one is made up of a well-established clinical anticancer agent, erlotinib, coordinated to a metallic gold center along with a triphenylphosphine (PPh₃) ligand. On the other hand, the second family of gold(I) complexes involves the chalcone(*E*)-3-(9-anthracenyl)-1-(pyridin-4-yl)prop-2-enone, whose properties are overwhelmingly promising in therapy against cancer, and either one PPh₃ or an N-heterocyclic carbene (NHC) group as co-ligand.

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References

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