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1. Introduction. Radium-223 is a first-in-class targeted alpha therapy for castration-resistant prostate cancer (CRPC) and symptomatic bone metastases with minimal myelosuppressive effects. The relatively large size of alpha particles, coupled with the high linear energy transfer of emitted particles, results in a short path length and localized area of cell destruction (<100 µm; 2-10 ...

Three-year Safety of Radium-223 Dichloride in Patients ...

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DICTIONARY OF ABBREVIATIONS AND ACRONYMS IN GEOGRAPHIC ...

Background. Current treatment guidelines for type 2 diabetes (T2D) highlight the importance of patient-centered engagement and support with lifestyle interventions and pharmacologic therapies to manage hyperglycemia to individualized glycemic targets. 1 Although there are now 12 classes of antihyperglycemic therapies available for the treatment of T2D, many patients do not achieve adequate ...

Design of DEVOTE (Trial Comparing Cardiovascular Safety of ...

Overview. As per the LawNo9.782, ResolutionNo9, and Ord650/2014, the Brazilian Health Surveillance Agency (Agência Nacional de Vigilância Sanitária) (ANVISA) is the regulatory authority responsible for clinical trial oversight, approval, and inspections in Brazil. ANVISA grants permission for clinical trials to be conducted in Brazil in accordance with the provisions of ResolutionNo466 ...

Brazil | ClinRegs

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Assistance was provided by the Asian Development Bank (ADB) under TA 9047-PAK, for which AF Mercados, Spain and Hagler Bailly Pakistan were engaged as program preparatory technical assistance (PPTA) consultants, to support the design and implementation of the Clean Energy Investment Program complying with a results-based lending modality (RBL).

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Architecture - Atkins

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Federal Register :: Guidelines Establishing Test ...

Mental Health Acronyms. A B C D E F G H I J K L M N O P Q R S T U V W X Y Z. This list of acronyms was compiled from a variety of sources. An acronym is a ...

Mental Health Acronyms

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FY 2018 Report from the Director - fda.gov

Overview. As per the NDPA CTReg and the UC-GCPs, the National Drug Authority (NDA) is the regulatory authority responsible for clinical trial approval and inspections in Uganda. The NDA grants permission for clinical trials to be conducted in Uganda in accordance with the provisions of the NDPA Act.. As stated in the NGHRP, the NDA regulates safety, quality, efficacy, handling and use of drugs ...

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