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Or Research Institution Belongs ("Cluster") Or The Private Hospital In Which The Trial Is To Be Conducted ("Ethics Committee") • The Medical Council Of Hong Kong And The Chinese Medicine Council Of Hong Kong, Which Investigate Noncompliance By Medical Practitioners And Chinese Medicine Practitioners With The Code Of Professional Jan 5th, 2024

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• Integrating Research And Care Is An Important Public Health Goal: Both Care Givers And Investigators Should Know What Happens To Their Patients • Integrated Systems Can Avoid Duplication Of Data Entry (e.g. Demographics, Concomitant Meds, Comorbidities) • EHRs Are An Important Resource For Identifying And Recruiting Patients For Studies Feb 10th, 2024

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Narratives, And/or Completed Case Report Forms. The Team Also Warned That The EMA Had Been Releasing Documents In Multiple Tranches And Unsuitable Formats, Making Independent Analysis Of Clinical Trial Outcomes Exceedingly Difficult.13 In Contrast, In 2011, The EMA Reportedly Granted A Full Request For CSRs Of Trials For The Controversial May 2th, 2024

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Precautions To Ensure Patient Safety Require That Clinical Investigators Must Report To The Sponsors All Serious Adverse Events On An Expedited Basis, Regardless Of Whether They Are Considered Drug-related Or Not. Events Which Cannot Be Analysed As Single Cases Need To Be Assessed On An Aggregate Basis And Reported If There Is A Difference In The Reporting Rates Between The Drug And The ... Apr 9th, 2024

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