

quality, adverse events, reporting and standards

- [Clinical Practice Guidelines Portal](#)
- [Quality use of Medicines in Australian Hospitals](#)
- [Australian Commission on Safety and Quality Health Care - safe ePrescribing and electronic medication management \(EMM\)](#)
- [Australian National Inpatient Medication Chart \(NIMC\)](#)
- [NEHTA - National E-Health Transition Authority](#)
 - [NEHTA - Clinical Knowledge Manager - development of standardised data for eHealth systems](#)
- [Australian Council on Healthcare Standards Clinical Indicator Program](#)
- [National Guideline Clearing House](#) - initiative of the Agency for Healthcare Research and Quality, and the Department of Health and Human Services in the United States
- [National Institute for Health and Clinical Excellence United Kingdom](#)
- [The Scottish Intercollegiate Guidelines Network \(SIGN\)](#)
- [Coroner's newsletter - Coronial Communique](#)
- [Agency for Healthcare Research & Quality \(United States\)](#)
- [Australian Emergency Medicine Events Register \(EMER\)](#)
- [Aust Fed Govt review of health 2017](#)
- [Vic Gov health incident and clinical risk reporting policy and Incident Severity Rating \(ISR\)](#)
 - ISR rating depends upon:
 - degree of impact
 - level of care
 - treatment required
 - **ISR 1:** the most severe incident, requires notification within 3 days to Govt Sentinel Event Program, and may require a RCA to be concluded and reported within 60 days
 - **ISR 2:** less severe; requires in depth case review by 2 people and reported to hospital's Clinical Governance C'tee
 - **ISR 3:** minimal actual harm
 - **ISR 4:** near-miss
 - ISR 3&4 investigations to be commenced within 2 business days and completed or closed within 7 days
 - ISR 1&2 investigations to be reviewed ASAP and investigation completed within 30 days

do we measure and document too much?

- [NY Times 2016: How Measurement Fails Doctors and Teachers](#)

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