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April 18th, 2013- 1:00 PM EST In
October 2011, CLSI published EP 23
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In 2011, the Clinical and Laboratory Standards Institute published EP23-A, Laboratory Quality Control Based on Risk Management, providing an

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Parvin is now a consultant. “ EP23 defines patient risk as the combination of the probability of occurrence of patient harm and the severity of that harm, ” Parvin said. “ The higher the expected severity of harm to the patient, the lower the

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Probability of occurrence has to be in order for the risk to be acceptable. ” .

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Review key aspects of risk management. Describe the various types of control processes. Identify

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- EP23 describes good laboratory practice for developing a quality control plan based on manufacturer ' s information, applicable regulatory and accreditation requirements, and the individual healthcare and laboratory

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healthcare professions strongly
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management and provides guidance
for laboratories to map their
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identify control processes that can
manage risk. This guideline can help
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Looking Ahead to Patient Risk
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ISO. Clinical laboratory medicine –In vitro diagnostic medical devices –Validation of user quality control procedures by the manufacturer. ISO 15198. Geneva, Switzerland: International Organization for Standardization; 2004. James H. Nichols, CLSI EP23™—Laboratory

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Management, 2012

Evolving Clinical Laboratory
Management Through ...
NABL 112:2016 clause 5.6 mentions
that the laboratory shall use two
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EP23 (2009) strongly recommends a laboratory to define its quality requirements in form individualised quality control plan for laboratories as a part of risk management.

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