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Eliminating Human Error from Your List of Manufacturing Deviations

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To err is human, but is ***human error*** a justified “cause” in GMP investigations? In pharmaceutical manufacturing, upwards of 80% of process deviations can be attributed to human error. While regulatory agencies make clear that all deviations, including those caused by human error, must be fully investigated – can we say that we are effectively doing that as an industry? On today's episode of Voices in Validation we speak with Ginette Collazo as we dig deeper into the identification and reduction of human error in GMP facilities.

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