

## GXP Talk– Question #77: Cleaning Validation in the Lab?



By [Jerry Lanese](#) [Richard Poska](#) Jan 18, 2019 9:54 am PST

Readers are invited to participate and contribute questions, answers, and discussion for this series – please share your successful practices with other readers. This column succeeds when we are able to address current GXP issues submitted by interested readers. Please contact column coordinators Jerry Lanese at [jerry@lanesegroup.com](mailto:jerry@lanesegroup.com), Rich Poska at [richposka@gmail.com](mailto:richposka@gmail.com), or Melissa Carella at [melissa.carella@ivtnetwork.com](mailto:melissa.carella@ivtnetwork.com) with comments, or submissions for publication. We welcome your questions, opinions, or other input. QUESTION #77 Cleaning methods for pharmaceuticals are developed...

**This content is only available to IVT members.**

Get help maintaining your knowledge . [Read More!](#)

If you are already a member and you do not have access to this article, [upgrade your membership](#).

Need help? [Read our FAQs](#).



[Jerry Lanese](#)

John G. (Jerry) Lanese is an independent consultant with a focus on Quality Systems and the components of an effective Quality System. He received a BA and MS from Middlebury College and a Ph.D. in Analytical Chemistry...

[View Author Bio](#)

[Richard Poska](#)

Richard Poska has more than 40 years of experience in pharmaceutical product development and regulatory affairs at major pharmaceutical companies, with the last 13 years focused in the regulatory CMC area. Early in his...

[View Author Bio](#)