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June 30th, 2016 - Technical report no 26 revised 2008 Article in PDA journal of pharmaceutical science and technology PDA 62 5 Suppl TR26 2 60 · February 2008 with 3 102 Reads Source PubMed

July 18th, 2018 - 1 PDA J Pharm Sci Technol 2005 Sep Oct 59 3 Suppl TR39 1 12 Technical report 39 cold chain guidance for medicinal products maintaining the quality of temperature sensitive medicinal products through the transportation environment'

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October 7th, 2018 - The PDA Technical Report on Low Endotoxin Recovery provides a science based and data driven strategy in dealing with the LER phenomenon The author of this article who acted as co lead of the TR authoring team provides first hand information that allows companies to develop product specific solutions to the LER problem'

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October 10th, 2018 - PDA Technical Report 26 is the most detailed comprehensive and descriptive document in respect to liquid filter validation It has not been meant as an industrial standard but is often enough used as such Filter users have to be aware about it because it is utilized by regulatory authorities Others will follow e g ISO 13408 2 Conclusion M W Jornitz May

2002 Conclusion cont Validation'

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October 1st, 2018 - The technical report concludes with a comprehensive reference section The revised PDA TR33 is a culmination of industry best practices that have been successfully used by multinational firms and accepted by global regulatory agencies when validating and implementing alternative and rapid microbiological methods'

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'GMP Compliance PDA Technical Report No 49 Points To

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October 11th, 2018 - PDA Technical Report 3 Revised 2013 TR 3 Validation of Dry Heat Processes Used for Depyrogenation and Sterilization PDA has revised TR 3 originally issued in 1981 The revision offers a modern scientific approach to dry heat depyrogenation and sterilization processes and includes recommendations for use by industry and regulators'

'PDA Technical Report 7 TR 7 Depyrogenation Putra

October 3rd, 2018 - *The Depyrogenation report consists of 14 chapters each written by an authority in the field Each chapter describes the different methods of depyrogenating solutions and devices*"**PDA Technical Report 72 TR 72 TP3 Global**

October 11th, 2018 - The Parenteral Drug Association's PDA latest technical report 72 TR 72 Passive Thermal Protection Systems for Global Distribution Qualification and Operational Guidance was released in October and is the first of its kind to outline selection and user guidelines for passive thermal protection systems"*PDA Technical Report 22 TR 22 Revised 2011 PROCESS*

October 9th, 2018 - *Technical Report No 22 Revised 2011 Process Simulation for Aseptically Filled Products originally published in 1996 The Task Force charged with updating the document ensured that the new version reflects the con?"***Product Catalog Parenteral Drug Association**
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October 8th, 2018 - News from the PDA PDA Technical Report No 61 TR 61 Steam In Place has been released The primary objective of the task force responsible for this technical report was to develop a scientific technical report on Steam In Place SIP processes that

provides recommendations for use by industry and regulators" PDA's New Technical Report for Biotech Cleaning Validation

October 13th, 2018 - The authors encourage biotech manufacturers to consult PDA Technical Report No 49 for a detailed perspective on current practices and issues in biotech cleaning validation'

'COLD CHAIN COMPLIANCE Qualifying Cold Chains Writing

October 10th, 2018 - Technical Report 39 revised 27 The PDA's TR39 was created in 2005 and revised in 2007 to harmonise it with EU regulatory expectations with the objective of providing ??guidance to industry on the essential principles and practices of transporting temperature sensitive medicinal'

'PDA publishes Technical Report on Cleaning Validation

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current scientific understanding of the causes of and control strategies for bioburden in pharmaceutical production systems with a special emphasis on biofilms in fluid handling systems'

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Performance Qualification Process Validation ? Responsibilities ? Summary" **PDA Technical
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