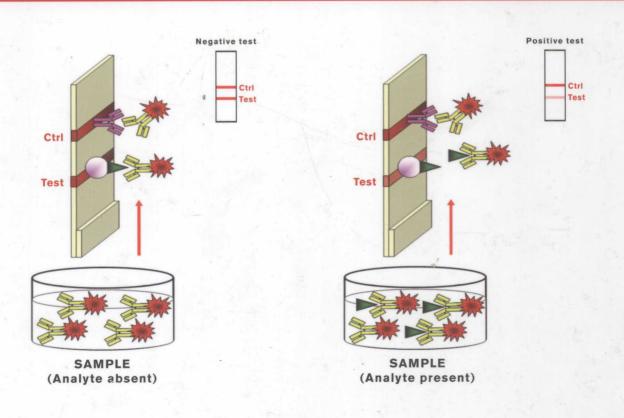
Chemical Analysis of Antibiotic Residues in Food

Edited by
Jian Wang, James D. MacNeil, and Jack F. Kay

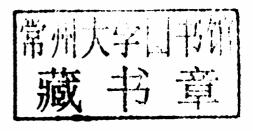




CHEMICAL ANALYSIS OF ANTIBIOTIC RESIDUES IN FOOD

Edited by

JIAN WANG JAMES D. MACNEIL JACK F. KAY





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PREFACE

Food safety is of great importance to consumers. To ensure the safety of the food supply and to facilitate international trade, government agencies and international bodies establish standards, guidelines, and regulations that food producers and trade partners need to meet, respect, and follow. A primary goal of national and international regulatory frameworks for the use of veterinary drugs, including antimicrobials, in food-producing animals is to ensure that authorized products are used in a manner that will not lead to non-compliance residues. However, analytical methods are required to rapidly and accurately detect, quantify, and confirm antibiotic residues in food to verify that regulatory standards have been met and to remove foods that do not comply with these standards from the marketplace.

The current developments in analytical methods for antibiotic residues include the use of portable rapid tests for on-site use or rapid screening methods, and mass spectrometric (MS)-based techniques for laboratory use. This book, Chemical Analysis of Antibiotic Residues in Food, combines disciplines that include regulatory standards setting, pharmacokinetics, advanced MS technologies, regulatory analysis, and laboratory quality management. It includes recent developments in antibiotic residue analysis, together with information to provide readers with a clear understanding of both the regulatory environment and the underlying science for regulations. Other topics include the choice of marker residues and target animal tissues for regulatory analysis, general guidance for method development and method validation, estimation of measurement uncertainty, and laboratory quality assurance and quality control. Furthermore, it also includes information on the developing area of environmental issues related to veterinary use of antimicrobials. For the bench analyst, it provides not only information on sources of methods of analysis but also an understanding of which methods are most suitable for addressing the regulatory requirements and the basis for those requirements.

The main themes in this book include antibiotic chemical properties (Chapter 1), pharmacokinetics, metabolism, and distribution (Chapter 2); food safety regulations (Chapter 3); sample preparation (Chapter 4); screening methods (Chapter 5); chemical analysis focused mainly on LC-MS (Chapters 6 and 7), method development and validation (Chapter 8), measurement uncertainty (Chapter 9), and quality assurance and quality control (Chapter 10).

The editors and authors of this book are internationally recognized experts and leading scientists with extensive firsthand experience in preparing food safety regulations and in the chemical analysis of antibiotic residues in food. This book represents the cutting-edge state of the science in this area. It has been deliberately written and organized with a balance between practical use and theory to provide readers or analytical laboratory staff with a reference book for the analysis of antibiotic residues in food.

JIAN WANG JAMES D. MACNEIL JACK F. KAY

Canadian Food Inspection Agency, Calgary, Canada St. Mary's University, Halifax, Canada University of Strathclyde, Glasgow, Scotland

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The editors are grateful to Dr. Dominic M. Desiderio, the editor of *Mass Spectrometry Reviews*, for the invitation to contribute a book on antibiotic residues analysis; to individual chapter authors, leading scientists in the field,

for their great contributions as the result of their profound knowledge and many years of firsthand experience; and to the editors' dear family members for their unending support and encouragement during this book project.

EDITORS

Dr. Jian Wang received his PhD at the University of Alberta in Canada in 2000, and then worked as a Post Doctoral Fellow at the Agriculture and Agri-Food Canada in 2001. He has been working as a leading Research Scientist at the Calgary Laboratory with the Canadian Food Inspection Agency since 2002. His scientific focus is on the method development using liquid chromatography-tandem mass spectrometry (LC-MS/MS) and UPLC/QqTOF for analyses of chemical contaminant residues, including antibiotics, pesticides, melamine, and cyanuric acid in various foods. He also develops statistical approaches to estimating the measurement uncertainty based on method validation and quality control data using the SAS program.

Dr. James D. MacNeil received his PhD from Dalhousie University, Halifax, NS, Canada in 1972 and worked as a government scientist until his retirement in 2007. During 1982–2007 he was Head, Centre for Veterinary Drug Residues, now part of the Canadian Food Inspection Agency. Dr. MacNeil has served as a member of the Joint FAO/WHO Expert Committee on Food Additives (JECFA), cochair of the working group on methods of Analysis and Sampling, Codex Committee on Veterinary Drugs in Foods (CCRVDF), is the former scientific editor for "Drugs, Cosmetics & Forensics" of J.AOAC Int., worked on IUPAC projects, has participated in various consultations on method validation and is the author of

numerous publications on veterinary drug residue analysis. He is a former General Referee for methods for veterinary drug residues for AOAC International and was appointed scientist emeritus by CFIA in 2008. Dr. MacNeil holds an appointment as an adjunct professor in the Department of Chemistry, St. Mary's University.

Dr. Jack F. Kay received his PhD from the University of Strathclyde, Glasgow, Scotland in 1980 and has been involved with veterinary drug residue analyses since 1991. He works for the UK Veterinary Medicines Directorate to provide scientific advice on residue monitoring programmes and manages the research and development (R&D) program. Dr. Kay helped draft Commission Decision 2002/657/EC and is an International Standardization Organization (ISO)-trained assessor for audits to ISO 17025. He served as cochair of the CCRVDF ad hoc Working Group on Methods of Sampling and Analysis and steered Codex Guideline CAC/GL 71-2009 to completion after Dr. MacNeil retired. Dr. Kay now cochairs work to extend this to cover multi-residue method performance criteria. He assisted JECFA in preparing an initial consideration of setting MRLs in honey, and is now developing this further for the CCRVDF. He also holds an Honorary Senior Research Fellowship at the Department of Mathematics and Statistics at the University of Strathclyde.

CONTRIBUTORS

- **Bjorn Berendsen**, Department of Veterinary Drug Research, RIKILT—Institute of Food Safety, Unit Contaminants and Residues, Wageningen, The Netherlands
- **Alistair Boxall**, Environment Department, University of York, Heslington, York, United Kingdom
- Andrew Cannavan, Food and Environmental Protection Laboratory, FAO/IAEA Agriculture & Biotechnology Laboratories, Joint FAO/IAEA Division of Nuclear Techniques in Food and Agriculture, International Atomic Energy Agency, Vienna, Austria
- Martin Danaher, Food Safety Department, Teagasc, Ashtown Food Research Centre, Ashtown, Dublin 15, Ireland
- **Leslie Dickson**, Canadian Food Inspection Agency, Saskatoon Laboratory, Centre for Veterinary Drug Residues, Saskatoon, Saskatchewan, Canada
- **Rick Fedeniuk**, Canadian Food Inspection Agency, Saskatoon Laboratory, Centre for Veterinary Drug Residues, Saskatoon, Saskatchewan, Canada
- **Lynn G. Friedlander**, Residue Chemistry Team, Division of Human Food Safety, FDA/CVM/ONADE/HFV-151, Rockville, Maryland
- **Kevin J. Greenlees**, Office of New Animal Drug Evaluation, HFV-100, USFDA Center for Veterinary Medicine, Rockville, Maryland
- Jack F. Kay, Veterinary Medicines Directorate, New Haw, Surrey, United Kingdom; also Department of Mathematics and Statistics, University of Strathclyde, Glasgow, United Kingdom (honorary position)
- Bruno Le Bizec, Food Safety, LABERCA (Laboratoire d'Etude des Résidus et Contaminants dans les Aliments), ONIRIS—Ecole Nationale Vétérinaire, Agroalimentaire et de l'Alimentation Nantes, Atlantique, Nantes, France

- **Peter Lees**, Veterinary Basic Sciences, Royal Veterinary College, University of London, Hatfield, Hertfordshire, United Kingdom
- James D. MacNeil, Scientist Emeritus, Canadian Food Inspection Agency, Dartmouth Laboratory, Dartmouth, Nova Scotia, Canada; also Department of Chemistry, St. Mary's University, Halifax, Nova Scotia, Canada
- Ross A Potter, Veterinary Drug Residue Unit Supervisor, Canadian Food Inspection Agency (CFIA), Dartmouth Laboratory, Dartmouth, Nova Scotia, Canada
- Philip Thomas Reeves, Australian Pesticides and Veterinary Medicines Authority, Regulatory Strategy and Compliance, Canberra, ACT (Australian Capital Territory), Australia
- Jacques Stark, DSM Food Specialities, Delft, The Netherlands
- **Sara Stead**, The Food and Environment Research Agency, York, North Yorkshire, United Kingdom
- Alida A. M. (Linda) Stolker, Department of Veterinary Drug Research, RIKILT—Institute of Food Safety Unit Contaminants and Residues, Wageningen, The Netherlands
- **Jonathan A. Tarbin**, The Food and Environment Research Agency, York, North Yorkshire, United Kingdom
- Pierre-Louis Toutain, UMR181 Physiopathologie et Toxicologie Experimentales INRA, ENVT, Ecole Nationale Veterinaire de Toulouse, Toulouse, France
- **Sherri B. Turnipseed**, Animal Drugs Research Center, US Food and Drug Administration, Denver, Colorado
- **Jian Wang**, Canadian Food Inspection Agency, Calgary Laboratory, Calgary, Alberta, Canada

CONTENTS

Pre	Preface					
Ack	nowl	edgmer	nt	xvi		
Edi	Editors					
Con	tribu	itors		XX		
1	Antibiotics: Groups and Properties					
	Phili	p Thoma	as Reeves			
	1.1	1.1 Introduction, 1				
			Identification, 1			
		1.1.2	Chemical Structure, 2			
		1.1.3	Molecular Formula, 2			
		1.1.4	Composition of the Substance, 2			
		1.1.5	pK_a , 2			
		1.1.6	UV Absorbance, 3			
		1.1.7	Solubility, 3			
			Stability, 3			
	1.2	Antibi	otic Groups and Properties, 3			
		1.2.1	Terminology, 3			
		1.2.2	* ·			
			Pharmacokinetics of Antimicrobial Drugs, 4			
		1.2.4				
			1.2.4.1 Spectrum of Activity, 5			
			1.2.4.2 Bactericidal and Bacteriostatic Activity, 6			
			1.2.4.3 Type of Killing Action, 6			
			1.2.4.4 Minimum Inhibitory Concentration and Minimum			
			Bactericidal Concentration, 7			
			1.2.4.5 Mechanisms of Action, 7			
		1.2.5				
		1.2.6	Clinical Toxicities, 7			
		1.2.7				
		1.2.8				
		1.2.9	Environmental Issues, 8			

vi CONTENTS

	1.3	1.3.1 1.3.2 1.3.3 1.3.4 1.3.5 1.3.6 1.3.7 1.3.8 1.3.9 1.3.10 1.3.11 1.3.12 1.3.13 1.3.14 Restric	Groups of Antibiotics, 8 Aminoglycosides, 8 β-Lactams, 10 Quinoxalines, 18 Lincosamides, 20 Macrolides and Pleuromutilins, 21 Nitrofurans, 27 Nitroimidazoles, 28 Phenicols, 30 Polyether Antibiotics (Ionophores), 31 Polypeptides, Glycopeptides, and Streptogramins, 35 Phosphoglycolipids, 36 Quinolones, 36 Sulfonamides, 44 Tetracyclines, 45 ted and Prohibited Uses of Antimicrobial Agents in Animals, 52 asions, 52					
	1.5		•					
		_	ments, 53					
	Refe	rences,	55					
2	Phai	rmacoki	inetics, Distribution, Bioavailability, and Relationship to					
-			Residues					
	Peter	Lees an	d Pierre-Louis Toutain					
	2.1		action, 61					
	2.2 Principles of Pharmacokinetics, 61							
			Pharmacokinetic Parameters, 61					
			Regulatory Guidelines on Dosage Selection for Efficacy, 64					
			Residue Concentrations in Relation to Administered Dose, 64					
		2.2.4						
		225	Clinical Populations, 66					
		2.2.5	Single-Animal versus Herd Treatment and Establishment of					
		2.2.6	Withholding Time (WhT), 66 Influence of Antimicrobial Drug (AMD) Physicochemical					
		2.2.0	Properties on Residues and WhT, 67					
	2.3	Admin	istration, Distribution, and Metabolism of Drug Classes, 67					
	2.5	2.3.1	Aminoglycosides and Aminocyclitols, 67					
		2.3.2	β-Lactams: Penicillins and Cephalosporins, 69					
		2.3.3	Quinoxalines: Carbadox and Olaquindox, 71					
		2.3.4	Lincosamides and Pleuromutilins, 71					
		2.3.5	Macrolides, Triamilides, and Azalides, 72					
		2.3.6	Nitrofurans, 73					
		2.3.7	Nitroimidazoles, 73					
		2.3.8	Phenicols, 73					
		2.3.9	Polyether Antibiotic Ionophores, 74					
		2.3.10	Polypeptides, 75					
		2.3.11	Quinolones, 75					
			Sulfonamides and Diaminopyrimidines, 77					
			Polymyxins, 79					
			Tetracyclines, 79					
	2.4		Guidelines for Residues by Regulatory Authorities, 81					
	2.5		ion, Assessment, Characterization, Management, and					
		Comm	Communication of Risk 82					

61

		2.5.1				
		2.5.2	Risk Ass	sessment, 84		
			2.5.2.1	Hazard Assessment, 88		
				Exposure Assessment, 89		
		2.5.3	Risk Cha	aracterization, 90		
		2.5.4	Risk Ma	nagement, 91		
			2.5.4.1	Withholding Times, 91		
			2.5.4.2	Prediction of Withdholding Times from Plasma		
				Pharmacokinetic Data, 93		
			2.5.4.3	International Trade, 93		
		2.5.5		mmunication, 94		
	2.6	Residu	ie Violatio	ons: Their Significance and Prevention, 94		
		2.6.1	Roles of	Regulatory and Non-regulatory Bodies, 94		
		2.6.2	Residue	Detection Programs, 95		
			2.6.2.1	Monitoring Program, 96		
				Enforcement Programs, 96		
				Surveillance Programs, 97		
				Exploratory Programs, 97		
			2.6.2.5	Imported Food Animal Products, 97		
			2.6.2.6	Residue Testing in Milk, 97		
	2.7	Furthe	r Conside	rations, 98		
		2.7.1	Injection	Site Residues and Flip-Flop Pharmacokinetics, 98		
				valence and Residue Depletion Profiles, 100		
		2.7.3		d Usage Data, 101		
				Sales of AMDs in the United Kingdom, 2003–2008, 101		
			2.7.3.2	1 0		
				Medicine in France, 1999–2005, 102		
			2.7.3.3	Global Animal Health Sales and Sales of AMDs for		
				Bovine Respiratory Disease, 103		
	Refe	rences,	104			
3	Anti	biotic 1	Residues	in Food and Drinking Water, and Food Safety		
		Regulations 111				
	Kevin J. Greenlees, Lynn G. Friedlander, and Alistair Boxall					
	3.1					
				d—Where is the Smoking Gun?, 111		
	3.3 How 3.3.1			Residue Concentrations Are Determined, 113		
				gy—Setting Concentrations Allowed in the Human		
		3.3.2	Diet, 11	Residue Concentrations for Substances Not Allowed in		
		3.3.2	Food, 1			
		3.3.3		Residue Concentrations Allowed in Food, 114		
		3.3.3	3.3.3.1	Tolerances, 115		
			3.3.3.2	Maximum Residue Limits, 116		
		3.3.4		onal Harmonization, 117		
	3.4			ner Exposure to Antibiotics in the Natural		
	Э.т		onment, 1			
		3.4.1		t to and Occurrence in Surface Waters and		
			_	vaters, 119		
		3.4.2	Uptake o	of Antibiotics into Crops. 119		
		3.4.2 3.4.3		of Antibiotics into Crops, 119 Antibiotics in the Environment to Human Health, 120		
	3.5	3.4.3		Antibiotics into Crops, 119 Antibiotics in the Environment to Human Health, 120		
		3.4.3	Risks of ary, 120			

4	Sample Preparation: Extraction and Clean-up					
	Alida	lida A. M. (Linda) Stolker and Martin Danaher				
	4.1	1 Introduction 125				
	4.2	Introduction, 125 Sample Selection and Pre-treatment, 126				
	4.3		e Extraction, 127			
	1.5	4.3.1	Target Marker Residue, 127			
		4.3.1 Target Warker Residue, 127 4.3.2 Stability of Biological Samples, 127				
	4.4	Extraction Techniques, 128				
		4.4.1 Liquid–Liquid Extraction, 128				
			Dilute and Shoot, 128			
			Liquid-Liquid Based Extraction Procedures, 129			
			4.4.3.1 QuEChERS, 129			
			4.4.3.2 Bipolarity Extraction, 129			
		4.4.4 Pressurized Liquid Extraction (Including Supercritical Fluid				
			Extraction), 130			
		4.4.5	Solid Phase Extraction (SPE), 131			
			4.4.5.1 Conventional SPE, 131			
			4.4.5.2 Automated SPE, 132			
		4.4.6	Solid Phase Extraction-Based Techniques, 133			
			4.4.6.1 Dispersive SPE, 133			
			4.4.6.2 Matrix Solid Phase Dispersion, 134			
			4.4.6.3 Solid Phase Micro-extraction, 135			
			4.4.6.4 Micro-extraction by Packed Sorbent, 137			
			4.4.6.5 Stir-bar Sorptive Extraction, 137			
			4.4.6.6 Restricted-Access Materials, 138			
		4.4.7	Solid Phase Extraction-Based Selective Approaches, 138			
			4.4.7.1 Immunoaffinity Chromatography, 138			
			4.4.7.2 Molecularly Imprinted Polymers, 139			
			4.4.7.3 Aptamers, 140			
			Turbulent-Flow Chromatography, 140			
		4.4.9	Miscellaneous, 142			
			4.4.9.1 Ultrafiltration, 142			
			4.4.9.2 Microwave-Assisted Extraction, 142			
			4.4.9.3 Ultrasound-Assisted Extraction, 144			
		4.5 Final Remarks and Conclusions, 144				
	References, 146					
_	D!		al Carranina Mathada	152		
5		-	al Screening Methods	153		
	Sara	ara Stead and Jacques Stark				
	5.1	Introduction, 153				
	5.2	Microl	bial Inhibition Assays, 154			
		5.2.1	The History and Basic Principles of Microbial Inhibition			
			Assays, 154			
		5.2.2	The Four-Plate Test and the New Dutch Kidney Test, 156			
		5.2.3	Commercial Microbial Inhibition Assays for Milk, 156			
		5.2.4	Commercial Microbial Inhibition Assays for Meat-, Egg-, and			
			Honey-Based Foods, 159			
		5.2.5	Further Developments of Microbial Inhibition Assays and Future			
			Prospects, 160			
			5.2.5.1 Sensitivity, 160			
			5.2.5.2 Test Duration, 161			
	5.2.5.3 Ease of Use, 161					

		 5.2.5.4 Automation, 161 5.2.5.5 Pre-treatment of Samples, 162 5.2.5.6 Confirmation/Class-Specific Identification, 163
	5.2.6	Conclusions Regarding Microbial Inhibition Assays, 164
5.3	_	Test Kits, 164
	5.3.1	
	5.3.2	Lateral-Flow Immunoassays, 165
		5.3.2.1 Sandwich Format, 166
		5.3.2.2 Competitive Format, 166
	5.3.3	Commercial Lateral-Flow Immunoassays for Milk, Animal
	5 O 4	Tissues, and Honey, 168
	5.3.4	Receptor-Based Radioimmunoassay: Charm II System, 170
	5.3.5	Basic Principles of Enzymatic Tests, 171
		5.3.5.1 The Penzyme Milk Test, 171
	506	5.3.5.2 The Delvo-X-PRESS, 172
~ 1		Conclusions Regarding Rapid Test Kits, 174
5.4		re Plasmon Resonance (SPR) Biosensor Technology, 174
		Basic Principles of SPR Biosensor, 174
	5.4.2	•
	5 1 2	Animal Tissues, Feed, and Honey, 175
	5.4.3	Conclusions Regarding Surface Plasmon Resonance (SPR)
5.5	Enzun	Technology, 176
5.5	5.5.1	ne-Linked Immunosorbent Assay (ELISA), 178 Basic Principles of ELISA, 178
		Automated ELISA Systems, 178
		Alternative Immunoassay Formats, 179
		Commercially Available ELISA Kits for Antibiotic Residues, 179
		Conclusions Regarding ELISA, 180
5.6		al Considerations Concerning the Performance Criteria for
0.0		ning Assays, 181
5.7		ll Conclusions on Bioanalytical Screening Assays, 181
Abbi		ns, 182
	rences,	
Che	mical A	Analysis: Quantitative and Confirmatory Methods 187
Jian	Wang ar	nd Sherri B. Turnipseed
6.1	Introd	uction, 187
6.2		-Class and Multi-class Methods, 187
6.3		natographic Separation, 195
0.0	6.3.1	Chromatographic Parameters, 195
	6.3.2	Mobile Phase, 195
	6.3.3	Conventional Liquid Chromatography, 196
		6.3.3.1 Reversed Phase Chromatography, 196
		6.3.3.2 Ion-Pairing Chromatography, 196
		6.3.3.3 Hydrophilic Interaction Liquid Chromatography, 197
	6.3.4	Ultra-High-Performance or Ultra-High-Pressure Liquid
		Chromatography, 198
6.4	Mass	Spectrometry, 200
	6.4.1	Ionization and Interfaces, 200
	6.4.2	
	6.4.3	Mass Spectrometers, 205
		6.4.3.1 Single Quadrupole, 205
		6.4.3.2 Triple Quadrupole, 206

6

		_	6.4.3.3 Quadrupole Ion Trap, 208 6.4.3.4 Linear Ion Trap, 209 6.4.3.5 Time-of-Flight, 210 6.4.3.6 Orbitrap, 212 Other Advanced Mass Spectrometric Techniques, 214 6.4.4.1 Ion Mobility Spectrometry, 214 6.4.4.2 Ambient Mass Spectrometry, 214 6.4.4.3 Other Recently Developed Desorption Ionization Techniques, 216 Fragmentation, 216 Mass Spectral Library, 216 gment, 219
	Refe	rences,	
7	_		due Quantitative and Confirmatory Methods 227
	Jona	than A.	Tarbin, Ross A. Potter, Alida A. M. (Linda) Stolker, and Bjorn Berendsen
	7.1	Introd	uction, 227
	7.2	Carba	dox and Olaquindox, 227
		7.2.1	Background, 227
		7.2.2	Analysis, 229
			Conclusions, 230
	7.3		fur and Desfuroylceftiofur, 230
			Background, 230
			Analysis Using Deconjugation, 231
		7.3.3	•
		7.3.4 7.3.5	
	7.4		Conclusions, 233 amphenicol, 233
	7.4	7.4.1	Background, 233
		7.4.2	Analysis by GC-MS and LC-MS, 233
		7.4.3	An Investigation into the Possible Natural Occurrence of CAP, 235
		7.4.4	Analysis of CAP in Herbs and Grass (Feed) Using LC-MS, 236
		7.4.5	Conclusions, 236
	7.5	Nitrof	urans, 236
		7.5.1	Background, 236
		7.5.2	Analysis of Nitrofurans, 236
		7.5.3	Identification of Nitrofuran Metabolites, 237
		7.5.4	Conclusions, 239
	7.6		midazoles and Their Metabolites, 239
		7.6.1	Background, 239
		7.6.2	Analysis, 240
		7.6.3	Conclusions, 241
	7.7		namides and Their N^4 -Acetyl Metabolites, 241
		7.7.1	Background, 241 N^4 -Acetyl Metabolites, 242
		7.7.2 7.7.3	Analysis, 243
		7.7.4	Conclusions, 244
	7.8		yclines and Their 4-Epimers, 244
	, .0	7.8.1	Background, 244
		7.8.2	Analysis, 245
		7.8.3	Conclusions, 246
	7.9		llaneous, 246

263

	7.10	7.9.1 7.9.2 7.9.3				
	Abbi	reviation	ns, 253			
8	References, 254 Method Development and Method Validation					
			and James D. MacNeil			
	8.1	Introdu	uction, 263			
	8.2	Source	es of Guidance on Method Validation, 263			
		8.2.1	Organizations that Are Sources of Guidance on Method Validation, 264			
			8.2.1.1 International Union of Pure and Applied Chemistry (IUPAC), 264			
			8.2.1.2 AOAC International, 264			
			8.2.1.3 International Standards Organization (ISO), 264			
			8.2.1.4 Eurachem, 265			
			8.2.1.5 VICH, 265			
			8.2.1.6 Codex Alimentarius Commission (CAC), 265			
			8.2.1.7 Joint FAO/WHO Expert Committee on Food Additives (JECFA), 265			
			8.2.1.8 European Commission, 266			
			8.2.1.9 US Food and Drug Administration (USFDA), 266			
	8.3 The Evolution of Approaches to Method Validation for Veterinary Drug					
			ues in Foods, 266			
		8.3.1	Evolution of "Single-Laboratory Validation" and the "Criteria Approach," 266			
		8.3.2				
		8.3.3				
	0.4	8.3.4	Codex Alimentarius Commission Guidelines, 267			
8.4 Method Performance Characteristics, 2						
	8.5	8.5.1	onents of Method Development, 268 Identification of "Fitness for Purpose" of an Analytical			
		0.5.1	Method, 269			
		8.5.2	Screening versus Confirmation, 270			
		8.5.3	Purity of Analytical Standards, 270			
		8.5.4	Analyte Stability in Solution, 271			
		8.5.5	Planning the Method Development, 271			
		8.5.6	Analyte Stability during Sample Processing (Analysis), 272			
		8.5.7	Analyte Stability during Sample Storage, 272			
		8.5.8	Ruggedness Testing (Robustness), 273			
	0 6	8.5.9	Critical Control Points, 274			
	8.6	8.6.1	onents of Method Validation, 274 Understanding the Requirements, 274			
		8.6.2	Management of the Method Validation Process, 274			
		8.6.3	Experimental Design, 275			

9

10

8.7	Performance Characteristics Assessed during Method Development and			
		med during Method Validation for Quantitative Methods, 275		
	8.7.1	Calibration Curve and Analytical Range, 275		
	8.7.2	Sensitivity, 277		
	8.7.3	Selectivity, 277		
		8.7.3.1 Definitions, 277		
		8.7.3.2 Suggested Selectivity Experiments, 278		
		8.7.3.3 Additional Selectivity Considerations for Mass		
	0.7.4	Spectral Detection, 279		
	8.7.4	Accuracy, 281		
	8.7.5	Recovery, 282		
	8.7.6	Precision, 283		
	8.7.7	Experimental Determination of Recovery and Precision, 283		
		8.7.7.1 Choice of Experimental Design, 283		
	070	8.7.7.2 Matrix Issues in Calibration, 286		
	8.7.8 8.7.9	Measurement Uncertainty (MU), 287 Limits of Detection and Limits of Quantification, 287		
0 0		Decision Limit ($CC\alpha$) and Detection Capability ($CC\beta$), 289 icant Figures, 289		
8.8	_			
8.9	rences,	Thoughts, 289		
Kere	rences,	209		
Moo	curomo	ent Uncertainty	295	
			293	
Jian	wang, A	ndrew Cannavan, Leslie Dickson, and Rick Fedeniuk		
9.1	Introd	uction, 295		
9.2	Genera	al Principles and Approaches, 295		
9.3	Worke	ed Examples, 297		
	9.3.1	EURACHEM/CITAC Approach, 297		
	9.3.2	Measurement Uncertainty Based on the Barwick-Ellison		
		Approach Using In-House Validation Data, 302		
	9.3.3	Measurement Uncertainty Based on Nested Experimental Design		
		Using In-House Validation Data, 305		
		9.3.3.1 Recovery (R) and Its Uncertainty $[u(R)]$, 306		
		9.3.3.2 Precision and Its Uncertainty $[u(P)]$, 312		
		9.3.3.3 Combined Standard Uncertainty and Expanded		
		Uncertainty, 312		
	9.3.4	Measurement Uncertainty Based on Inter-laboratory Study Data, 312		
	9.3.5	Measurement Uncertainty Based on Proficiency Test Data, 317		
	9.3.6	Measurement Uncertainty Based on Quality Control Data and		
		Certified Reference Materials, 319		
		9.3.6.1 Scenario A: Use of Certified Reference Material for		
		Estimation of Uncertainty, 320		
		9.3.6.2 Scenario B. Use of Incurred Residue Samples and		
		Fortified Blank Samples for Estimation of		
		Uncertainty, 324		
Refe	erences,	325		
Qua	lity As	surance and Quality Control	327	
Andr	ew Canr	navan, Jack F. Kay, and Bruno Le Bizec		
10.1		uction, 327		
	10.1.1 Quality—What Is It?, 327			