

# Contents

1. Introduction	1
2. General considerations	2
2.1 Modification of the agenda	2
2.2 Initiatives to promote transparency of the MRL-setting process of the Committee	2
2.3 Standards for the generation of data	3
2.4 Deadlines for the submission of data	4
2.5 Initiatives of FAO/WHO Member States and nongovernmental organizations	4
2.6 Residues at the injection site	4
3. Comments on residues of specific veterinary drugs	5
3.1 Anthelmintic agents	6
3.1.1 Moxidectin	6
3.1.2 Tiabendazole (thiabendazole)	8
3.2 Antimicrobial agents	11
3.2.1 Ceftiofur	11
3.2.2 Danofloxacin	15
3.2.3 Dihydrostreptomycin and streptomycin	25
3.2.4 Enrofloxacin	31
3.2.5 Flumequine	35
3.2.6 Gentamicin	43
3.2.7 Spiramycin	44
3.3 Glucocorticosteroid	46
3.3.1 Dexamethasone	46
3.4 Insecticides	46
3.4.1 Cyfluthrin	46
3.4.2 Fluazuron	54
4. Recommendations	60
Acknowledgement	60
References	61
Annex 1	
Reports and other documents resulting from previous meetings of the Joint FAO/WHO Expert Committee on Food Additives	62
Annex 2	
Recommendations on compounds on the agenda, and further toxicological studies and other information required	69