Turkey Respiratory Complex

Involved Pathogens

Field Situation

Low pathogenic avian influenza

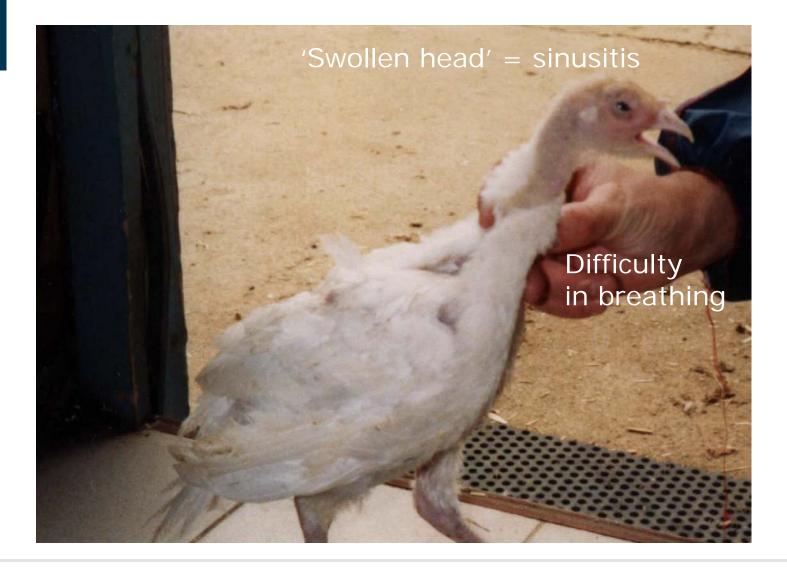
Current Solutions

XXIII. DERZSY NAPOK 2015. június 4-5. Zalakaros

Merial Avian Technical Services – Drs S. LEMIERE (Author & Presenter) & I. DEVAUD (Co-author) June 2015 © 2015 Merial SAS. Lyon, France - All rights reserved - Do not distribute



Turkey respiratory complex



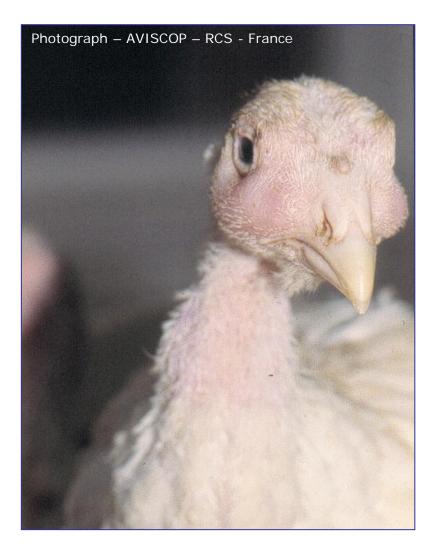
Involved Pathogens

- Primary pathogens = viruses, mainly but not exclusively:
 - Turkey Rhinotracheitis Metapneumovirus
 - Paramyxovirus type 1
 - Etc.



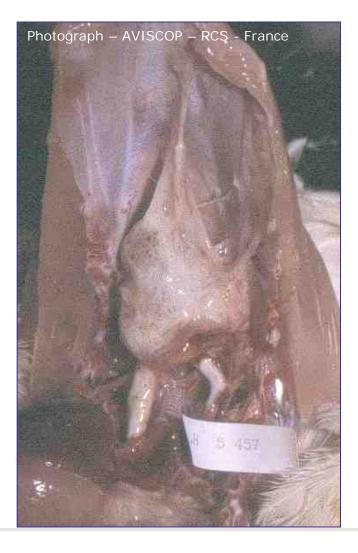


- Secondary infections involving bacteria, mainly but could be primary pathogens:
 - Ornithobacterium rhinotracheale
 - Mycoplasma spp
 - Escherichia coli
 - Etc.



Mycoplasma gallisepticum infection:





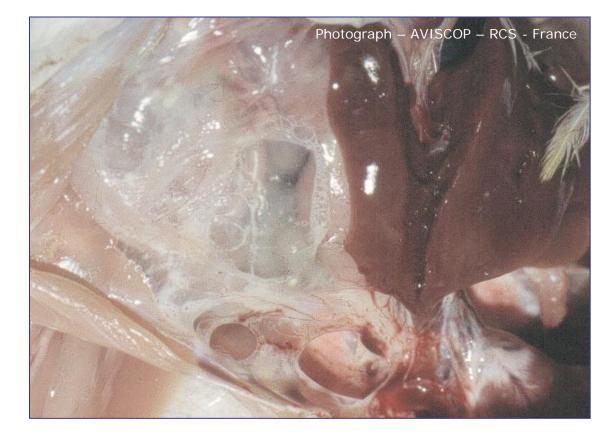
 Mycoplasma gallisepticum infection pericarditis



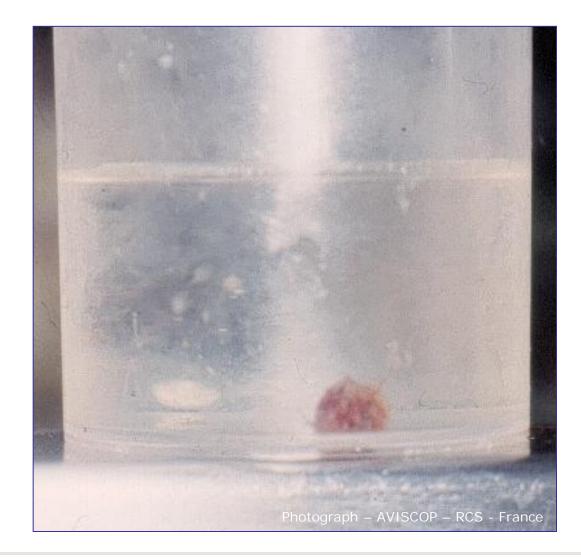
 Mycoplasma gallisepticum infection – pericarditis & perihepatitis

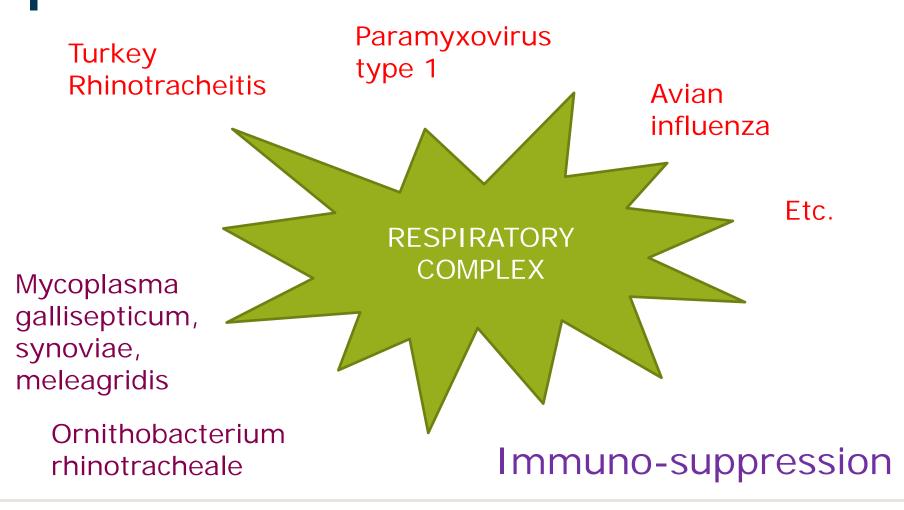


 Mycoplasma gallisepticum infection – airsacculitis

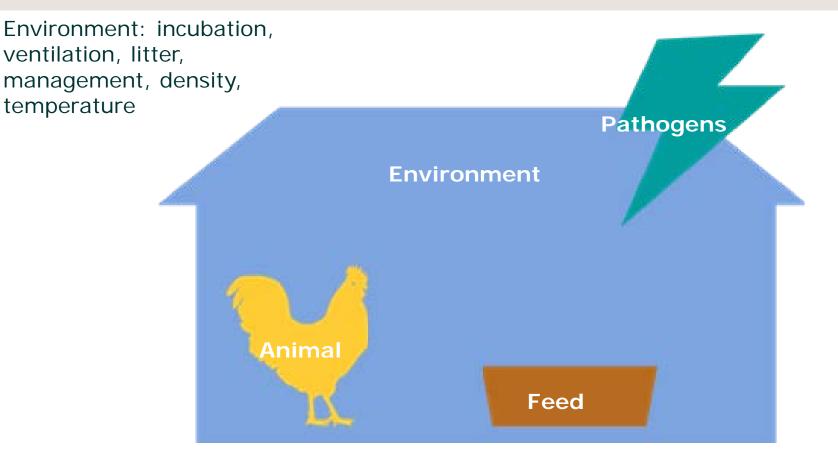


 Mycoplasma gallisepticum infection – pneumonia



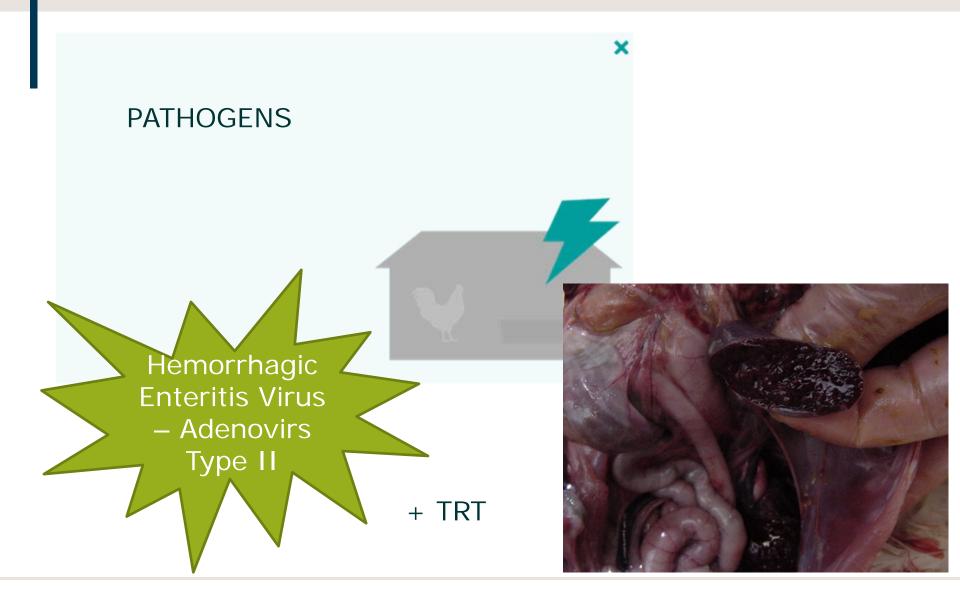


Turkey respiratory complex – Immuno-suppression



Animal: maternal antibodies, genetics, stress, age Feed: unbalanced diet, mycotoxin contamination, environmental toxin contamination

Turkey respiratory complex – Immuno-suppression



Turkey respiratory complex – Mixed infections

- TRT viral infection followed by Ornithobacterium rhinotracheale
- Enhanced clinical score further to mixed infection

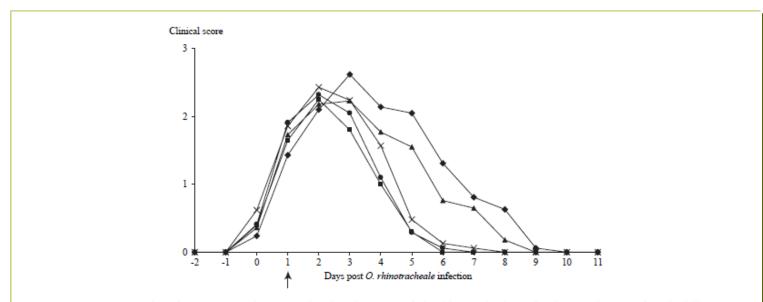


Figure 1. Mean clinical scores in turkeys inoculated with APV and O. rhinotracheale and subsequently treated with different antimicrobial agents: \blacksquare , group E3, 3 days of enrofloxacin (10 mg/kg); \blacklozenge , group E5, 5 days of enrofloxacin (10 mg/kg); \blacktriangle , group A, 5 days of amoxicillin (20 mg/kg); \times , group F, 5 days of florfenicol (20 mg/kg); \blacklozenge , no treatment, control group. Arrow indicates first day of antibiotic treatment.

Turkey respiratory complex – Mixed infections

 AI H9N2 viral infection followed by Ornithobacterium rhinotracheale + other pathogens including viruses

Detection from field samples

Table 4. Laboratory diagnosis of tarkey pallog ms other than AIV in birds from field outbreaks and experimental infection.												
	Testing for the presence of:											
	aMPV	TCoV	ASTRO	ROTA	REO	PARVO	ADENO	M0/MS/MM	в	۸.	OR	T
Outbreak flock	RT/PCR	rRT-PCR	REPOR	RT-PCR	rRT-PCR	PCR	PCR	PERMERTER	PCR	ELISA	PCR	H.ISA
1/A	nog	nog	nog	nog	nog	nog	nog	nog	10	PO8	POS	THE
1/B	nog	nog	neg	nog	neg	nog	nog	nog	100	PO8	neg	100
1/C	nog	nog	neg	nog	neg	nog	nog	nog	10	PO8	POS	THE
1/D	nog	nog	mog	nog	POS	nog	nog	mog	100	POS	neg	10
1/E	nog	nog	nog	nog	neg	nog	nog	nog	100	POS	POS	100
2/A	nog	nog	mog	mog	nog	mog	nog	nog	100	POS	POS	THE
2/8	mag	nog	mog	mog	neg	mog	nog	nog	100	POS	neg	THE
3	mog	nog	POS	POS	mog	POS	nog	nog	100	nog	POS	THE
4	nog	nog	nog	nog	nog	nog	nog	nog	10	POS	POS	THE
Turkeys in fected experimentally	nog	nog	POS	nog	POS	nog	POS	nog	mgTOS*	nt/POS*	PO8/POS*	n#PC8*

"before start of the experiment /14 dpl.

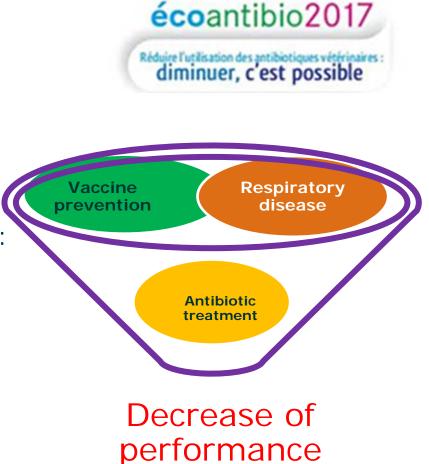
abl? V. avian meltignismovirus; TG/V. taloy coronavirus; ASTRO, atrovirusa; ROTA, neurinus a; REO, novirusa; RORVO, pavovirusa; ADENO, abnovirusa; MG, Mycopisusa gallaspirusa; MS, Mycopisusa gnories; MM, Mycopisusa meltografic; BA, Sordardia artus; ORT, Ornibolastation relacements; p. not trand.

Diagnostic

- Routine testing for monitoring:
 - Serology: mainly ELISA commercial kits for TRT; IHA for PMV1; Etc.
 - Bacteriology: mainly for E. coli, O. rhinotracheale (blood agar), Etc.
 - PCR from tracheal swabs: mainly for M. gallisepticum, M. synoviae, TRT, Etc.

Field Situation

- Primary criterion of evaluation:
 - Respiratory virus circulation
 - Serology
 - PCR
- Secondary criterion of evaluation:
 - Immunosuppressive virus circulation
 - Rhinotracheitis virus
 - Paramyxovirus type 1
 - Hemorrhagic enteritis virus



Infectious diseases Complexity of diagnostic

- Complications
- Mortality
- Arthritis
- Breast skin lesions

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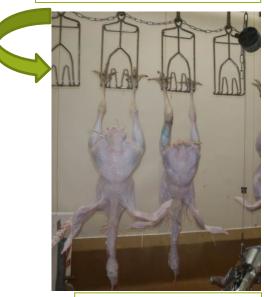
Réduire l'utilisation des antibiotiques vétérinaires : diminuer, c'est possible Slaughter Transformation

- Total condemnation
- Abnormalities originating partial condemnation

HEAL

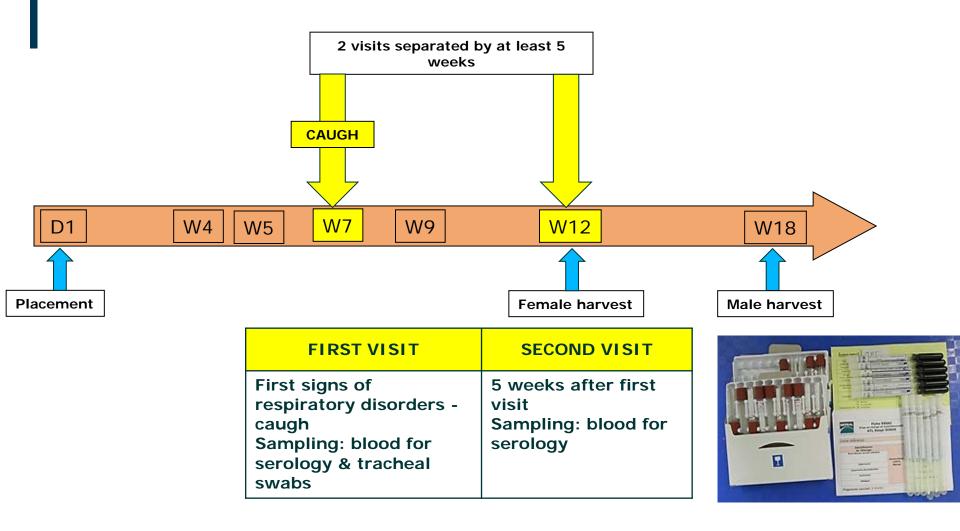
Microbiological counts

Respiratory complex



Partial condemnation

High quality transformed meat

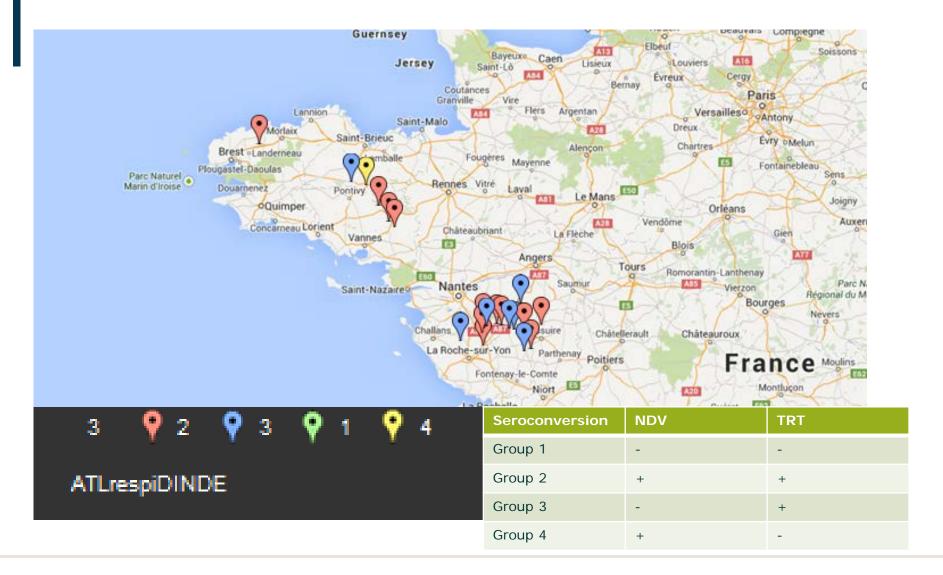




21/30 flocks: evidenced wild virus circulation

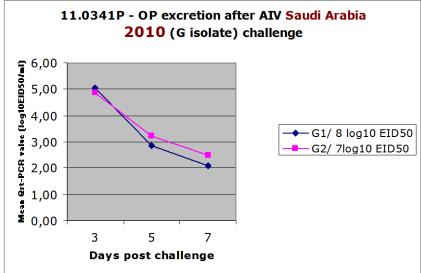
- 70% in Pays de Loire region
- 50% in Brittany

	Sero- conversion	Brittany	Pays de Loire	Negative or vaccine
ND & TRT positives	6	3	3	10
ND positives	5	1	4	/
TRT positives	10	0	10	/
Total positives	21	4	17	/
Total flocks	31	7	24	/



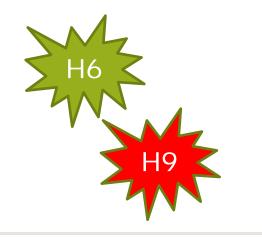
- Virus with low pathogenicity in laboratory conditions:
 - Virus shedding
 - Poor clinical signs in chickens, unless model of co-infection
 - Turkey species more susceptible





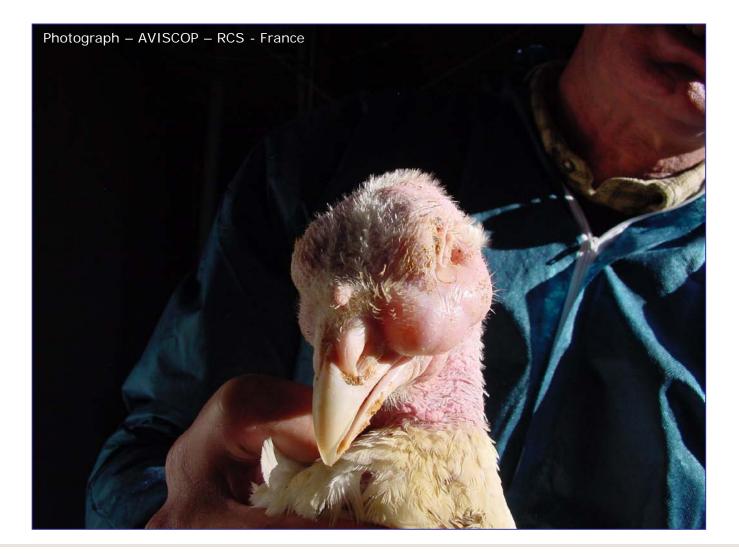








Photograph – AVISCOP – RCS - France AX Make 211







Photograph – AVISCOP – RCS - France





+ pancreatitis associated with AI H9N2

Current Solutions

Current solutions – Optimized vaccination programs

- Respiratory diseases:
 - TRT live vaccine 3 time application from day old during the rearing period (for instance D1-D17-D56)
 - PMV1 live vaccine 2 time application during the rearing period (for instance D28-D56)
- Immuno-suppressive diseases:
 - Adenovirus type II live vaccine application around D28 of age further to maternally-derived antibody waning

Current solutions – Rational use of antibiotics

- Antibiograms + strategy of use:
 - More & more antibiotic resistance
 - Selection of the efficient antibiotic
- Macrolides:
 - Mostly efficient against O. rhinotraeale newer generation
 - Efficient against Mycoplasmas

Current solutions – Biosecurity



Thank you for your attention!

