Sinusite maxillaire aiguë

Dr Guillaume Moulis

Cas clinique

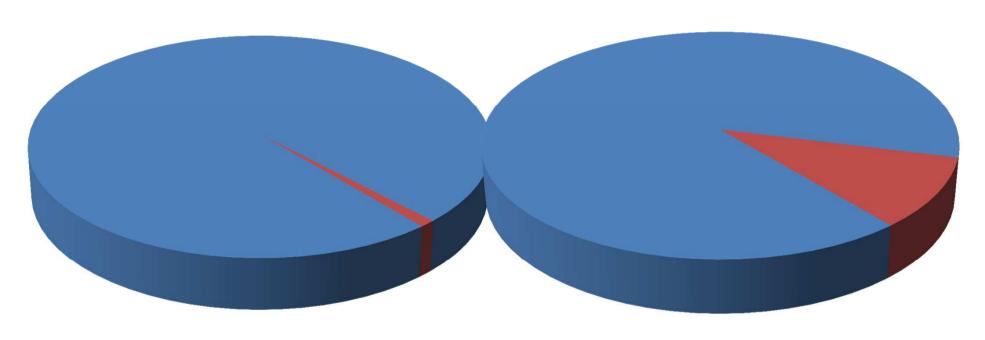
- Lundi
- Homme de 63 ans, informaticien, doit rendre un important rapport pour vendredi
- Depuis 48 heures : rhinite claire, douleur maxillaire droite (EVA : 4), augmentée la tête penchée en avant et par pression
- Terrain
 - HTA équilibrée sous énalapril
 - Hypercholestérolémie sous simvastatine
 - Syndrome anxio-dépressif sous paroxétine
 - Hypertrophie bénigne de la prostate

Antibiotiques?

□Oui

□Non

Fréquence des surinfections bactériennes



ADULTES: 0,5-2% ENFANTS: 5-13%

Sandes & Gwaltney CID 2004

Quelle situation vous ferait prescrire des antibiotiques ?

- **☐** Douleur dentaire
- **□** Douleur rétro-orbitaire
- ☐ Ecoulement nasal purulent
- ☐ Niveau hydro-aérique maxillaire à la radiographie
- ☐ Diabétique de type 2 parfaitement équilibré sous metformine

Quel antalgique?

- **□**Paracétamol
- □lbuprofène
- **□**Tramadol
- **□**Diclofénac
- **□**Prednisone

AINS/paracétamol

EFFICACITÉ ET TOLÉRANCE DE L'ACIDE TIAPROFÉNIQUE (SURGAM ®) DANS LES SINUSITES AIGUES DE L'ADULTE

Résultats d'une étude randomisée contre paracétamol et contre placebo

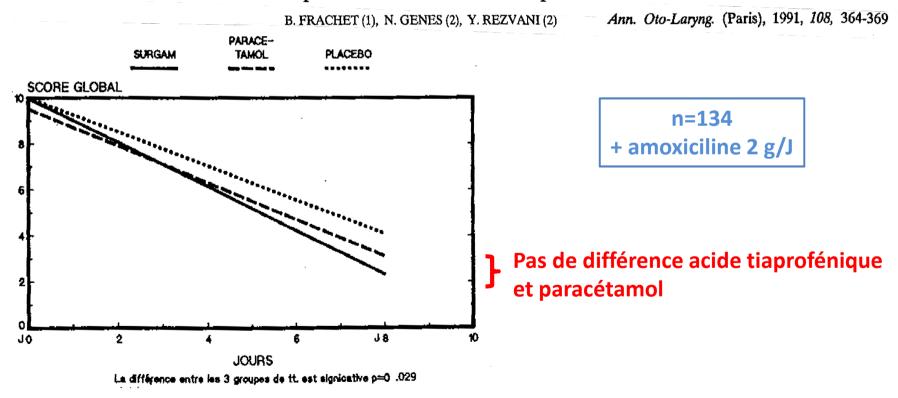


Fig. 1. – Evolution du score global.

Paracétamol en chirurgie de sinus

The Laryngoscope Lippincott Williams & Wilkins, Inc. © 2006 The American Laryngological, Rhinological and Otological Society, Inc.

Acetaminophen is Highly Effective in Pain Treatment After Endoscopic Sinus Surgery

Tatu Kemppainen, MD; Hannu Kokki, MD, PhD; Henri Tuomilehto, MD, PhD; Juhan Seppä, MD, PhD; Juhani Nuutinen, MD, PhD

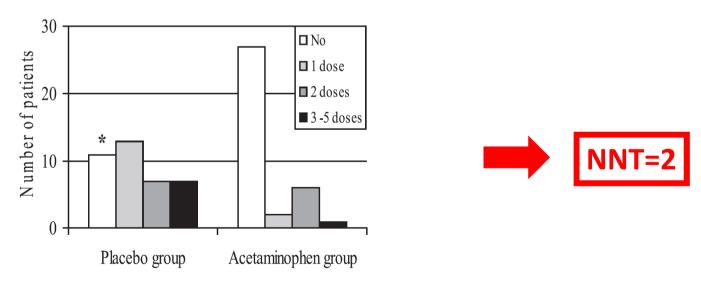


Fig. 1. Number of oxycodone doses in the placebo and acetamin-ophen group. ${}^*P = .001$.

AINS vs. paracétamol après chirurgie

Comparison of Parecoxib and Proparacetamol in Endoscopic Nasal Surgery Patients Yonsei Med J 49(3):383 - 388, 2008

DOI 10.3349/ymj.2008.49.3.383

Yigal Leykin, Andrea Casati, Alessandro Rapotec, Massimiliano Dalsasso, Luigi Barzan, Guido Fanelli, and Tommaso Pellis

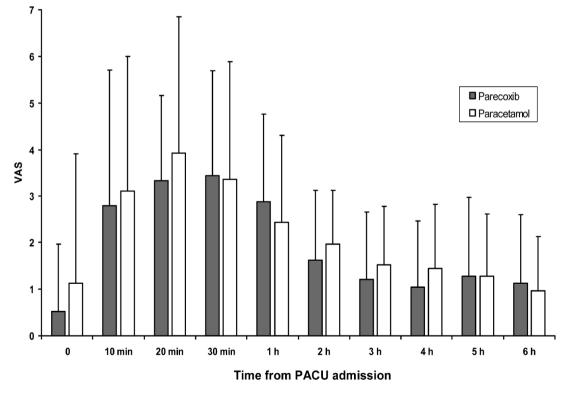


Fig. 1. Mean postoperative VAS measured at PACU admission for the first 6 h.

AINS vs. paracétamol après chirurgie

Comparison of Parecoxib and Proparacetamol in Endoscopic Nasal Surgery Patients Yonsei Med J 49(3):383 - 388, 2008

Yonsei Med J 49(3):383 - 388, 2008 DOI 10.3349/ymj.2008.49.3.383

Yigal Leykin, Andrea Casati, Alessandro Rapotec, Massimiliano Dalsasso, Luigi Barzan, Guido Fanelli, and Tommaso Pellis

Table 3. Incidence of Adverse Events Reported during First 24 h After Surgery in Patients Receiving Proparacetamol or Parecoxib After Endoscopic ENT Surgery

	Group proparacetamol ($n = 25$)	Group parecoxib (n = 25)	
PONV	3 (12%)	2 (8%)	
Gastric pain	0 (0%)	0 (0%)	
Minor bleeding	0 (0%)	2 (8%)	

PONV, post operative nausea and vomit.

Results are presented as number (%).

Corticoïdes en monothérapie

Systemic corticosteroids for acute sinusitis

Roderick P Venekamp¹, Matthew J Thompson², Gail Hayward³, Carl J Heneghan³, Chris B Del Mar⁴, Rafael Perera³, Paul P Glasziou ⁴, Maroeska M Rovers⁵



Cochrane Database of Systematic Review. 2014, ssue 3. Art. No.: CD008115.

Corticoïdes en monothérapie

Systemic corticosteroids for acute sinusitis

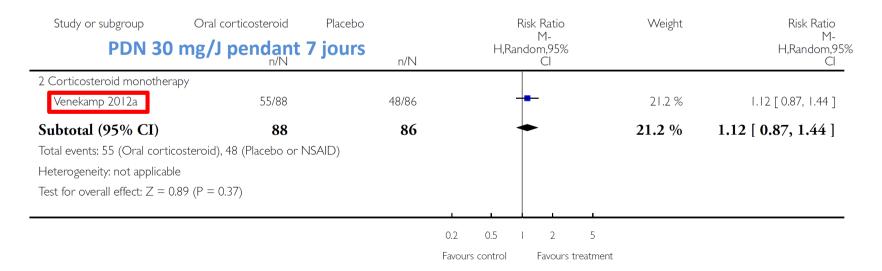
Roderick P Venekamp¹, Matthew J Thompson², Gail Hayward³, Carl J Heneghan³, Chris B Del Mar⁴, Rafael Perera³, Paul P Glasziou ⁴, Maroeska M Rovers⁵

Analysis I.I. Comparison I Oral corticosteroids versus placebo Outcome I Proportion of patients with resolution or improved symptoms at days 3 to 7.

Review: Systemic corticosteroids for acute sinusitis

Comparison: I Oral corticosteroids versus placebo or NSAID

Outcome: I Proportion of patients with resolution or improved symptoms at days 3 to 7



Corticoïdes en monothérapie

CMAJ

RESEARCH

Systemic corticosteroid monotherapy for clinically diagnosed acute rhinosinusitis: a randomized controlled trial

Roderick P. Venekamp MD PhD, Marc J.M. Bonten MD PhD, Maroeska M. Rovers PhD, Theo J.M. Verheij MD PhD, Alfred P.E. Sachs MD PhD

CMAJ, October 2, 2012, 184(14)

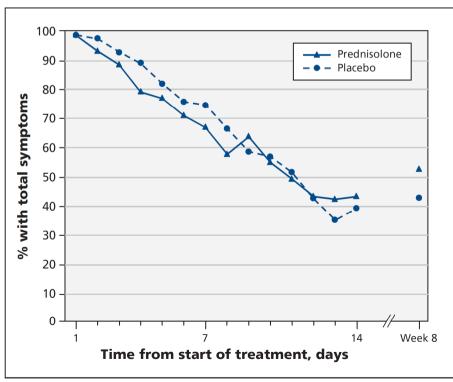


Figure 2: Proportion of patients who had combined symptoms of runny nose, postnasal discharge, nasal congestion, cough and facial pain during follow-up.

Arguments pharmacologiques

- □ Paracétamol
 □ Ibuprofène
 □ Tramadol
 □ Diclofénac
 □ Prednisone
- Lundi
- Homme de 63 ans, informaticien, doit rendre un rapport pour vendredi
- Depuis 48 heures : rhinite claire, douleur maxillaire droite (EVA : 4), augmentée la tête penchée en avant et par pression
- Terrain
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 - Syndrome anxio-dépressif sous paroxétine
 - Hypertrophie bénigne de la prostate

Cas clinique

- Mercredi
- Douleur persistante
- Augmentation de la rhinorrhée purulente
- Examen normal

Antibiotiques?

□Oui

□Non

Antibiotiques?

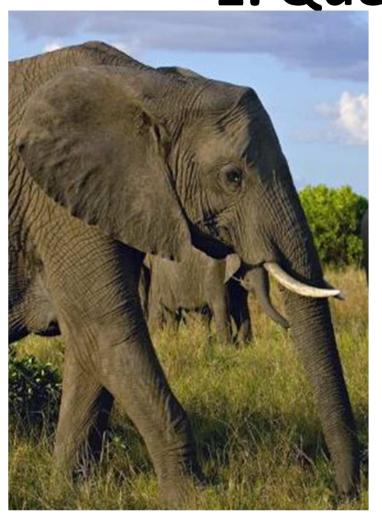
Lequel ?
 □ Amoxicilline + acide clavulanique oral
 □ Azithromycine orale
 □ Cefpodoxime orale
 □ Lévofloxacine orale

• Combien de temps ?

□ Amoxicilline orale

Choix d'un antibiotique

1. Quel microbe?



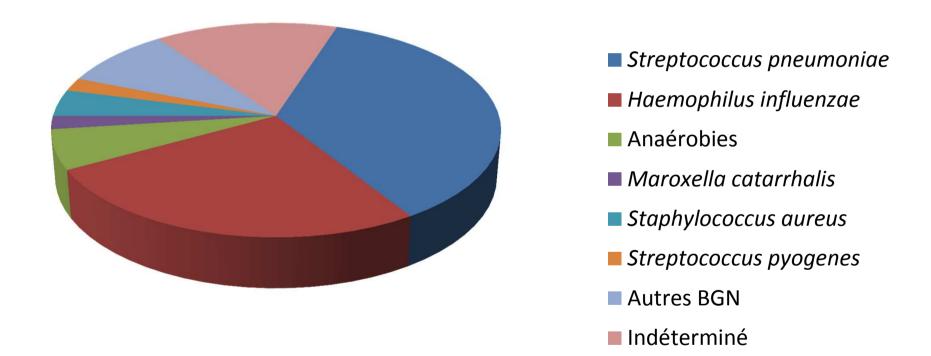


Choix d'un antibiotique

1. Quel microbe?



Spectre bactérien



Sandes & Gwaltney CID 2004

Spectre bactérien

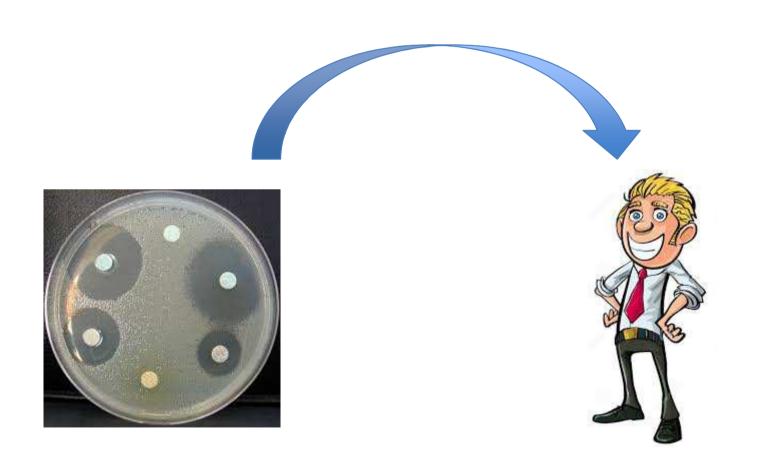
	Pneumocoque	Haemophilus	
Amoxicilline	15-35%	20-35%	
Amoxicilline + AC	15-35%	<10%	
Cefpodoxime	20-60%	+	
Azithromycine	35-70%		
Lévoflofloxacine	+	+	

Choix de l'antibiotique

	1. Microbe
Amoxicilline	
Amoxicilline + AC	
Cefpodoxime	
Azithromycine	
Lévoflofloxacine	

Choix d'un antibiotique

2. Preuves cliniques?



Antibiotics for acute maxillary sinusitis in adults (Review)

Ahovuo-Saloranta A, Rautakorpi UM, Borisenko OV, Liira H, Williams Jr JW, Mäkelä M



Cochrane Database of Systematic Review 2014, Issue 2. Art. No.: CD000243.

Antibiotics compared to placebo for uncomplicated acute maxillary sinusitis Patient or population: patients with uncomplicated acute maxillary sinusitis Settings: primary care Intervention: antibiotics: penicillin V, amoxicillin and azithromycin Comparison: placebo Illustrative comparative risks* (95% CI) No of participants **Quality of the evidence** Comments **Outcomes** Relative effect (95% CI) (studies) (GRADE) Corresponding risk **Assumed risk Placebo Antibiotics** Failure defined as a lack Failure defined as a lack RR 0.66 Adverse effects⁸: Clinical failure rate 1084 $\oplus \oplus \oplus \bigcirc$ of full recovery or im- of full recovery or im- (0.47 to 0.94) patients randomised and **moderate**² antibiotics 8% to 59%; Follow-up: 7 to 15 days provement 1058 evaluated placebo 6% to 38% provement (5 studies¹)



90 per 1000 (64 to 128)

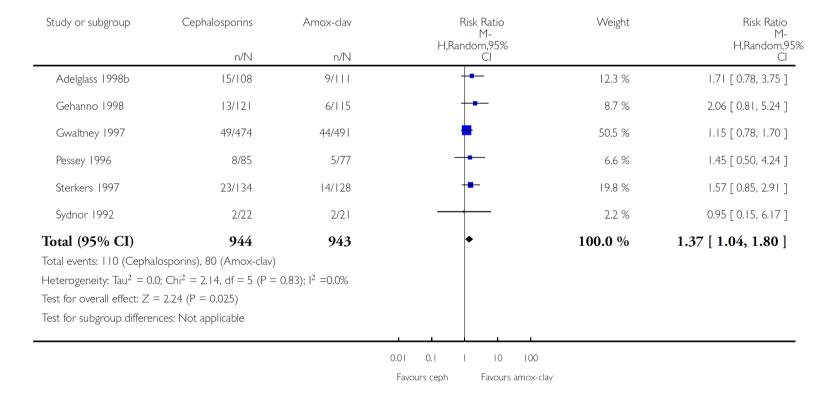
136 per 1000

Analysis 2.1. Comparison 2 Cephalosporin/macrolide versus amoxicillin-clavulanate, Outcome 1 Ceph versus amox-clav; clinical failure defined as a lack of full recovery or improvement at 7 to 15 days of follow-up.

Review: Antibiotics for acute maxillary sinusitis in adults

Comparison: 2 Cephalosporin/macrolide versus amoxicillin-clavulanate

Outcome: I Ceph versus amox-clav; clinical failure defined as a lack of full recovery or improvement at 7 to 15 days of follow-up

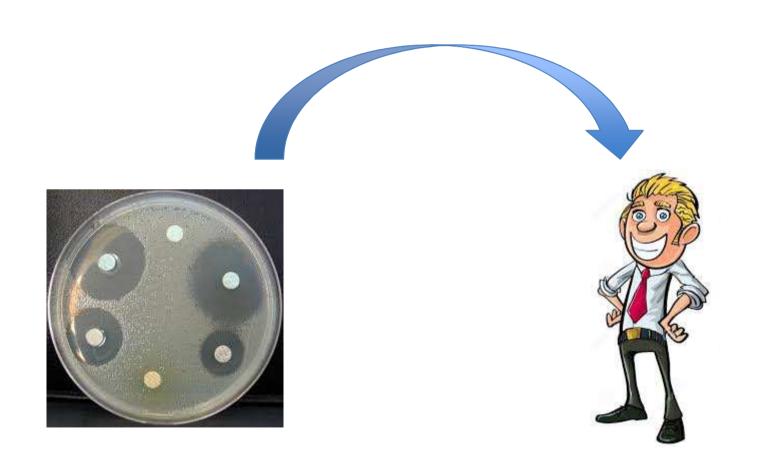


Choix de l'antibiotique

	2. Preuves cliniques
Amoxicilline	
Amoxicilline + AC	
Cefpodoxime	
Azithromycine	
Lévoflofloxacine	

Choix d'un antibiotique

3. Effets indésirables?

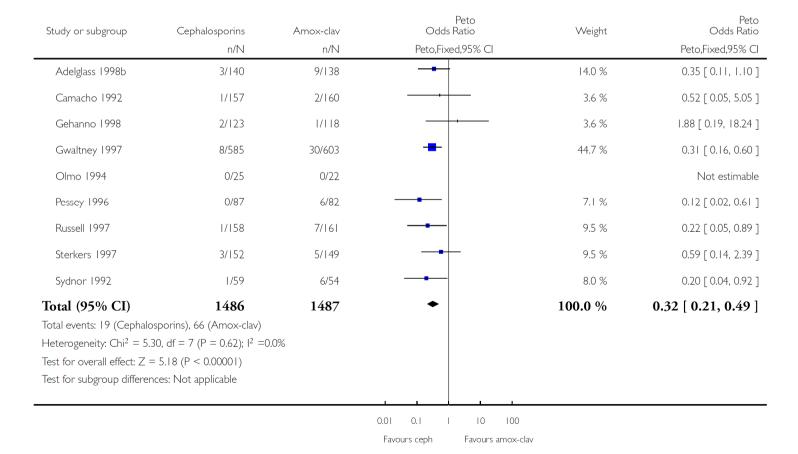


Analysis 2.3. Comparison 2 Cephalosporin/macrolide versus amoxicillin-clavulanate, Outcome 3 Drop-outs due to adverse effects (cephalosporins).

Review: Antibiotics for acute maxillary sinusitis in adults

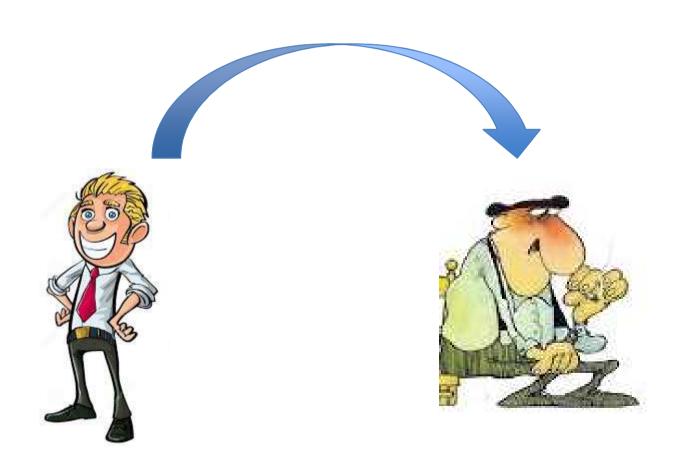
Comparison: 2 Cephalosporin/macrolide versus amoxicillin-clavulanate

Outcome: 3 Drop-outs due to adverse effects (cephalosporins)



Choix d'un antibiotique

3. Effets indésirables?



Arguments pharmacologiques

- Lequel?
 - ☐ Amoxicilline + acide clavulanique oral
 - **□** Azithromycine orale
 - ☐ Cefpodoxime orale
 - ☐ Lévofloxacine orale
 - **□** Amoxicilline orale

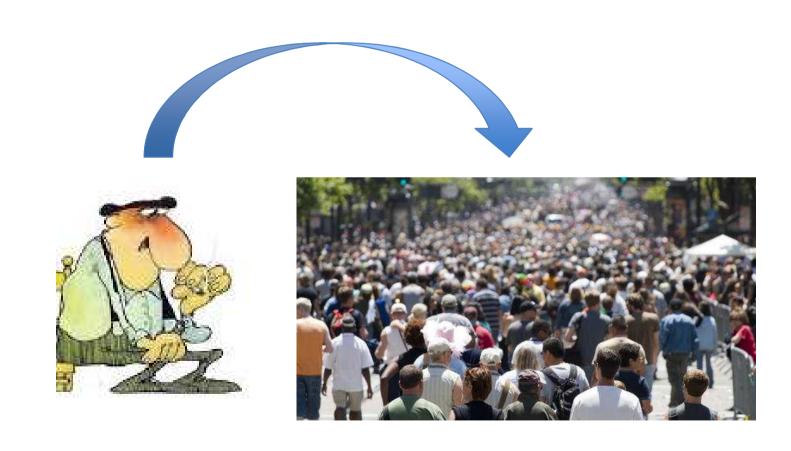
- Terrain
 - 63 ans
 - HTA équilibrée sous énalapril
 - Hypercholestérolémie sous simvastatine
 - Syndrome anxio-dépressif sous paroxétine
 - Hypertrophie bénigne de la prostate

Choix de l'antibiotique

	3. Effets indésirables
Amoxicilline	
Amoxicilline + AC	
Cefpodoxime	
Azithromycine	
Lévoflofloxacine	

Choix d'un antibiotique

4. Dégâts collatéraux ?



Choix d'un antibiotique

4. Dégâts collatéraux ?





Choix de l'antibiotique

	4. Ecologie
Amoxicilline	
Amoxicilline + AC	
Cefpodoxime	
Azithromycine	
Lévoflofloxacine	

Choix de l'antibiotique

	1. Microbe	2. Preuves cliniques	3. Effets indésirables	4. Ecologie
Amoxicilline				
Amoxicilline + AC				
Cefpodoxime				
Azithromycine				
Lévoflofloxacine				

Cas clinique

- La date limite pour rendre son rapport approche.
- « Je veux des corticoïdes »



Systemic corticosteroids for acute sinusitis

Roderick P Venekamp¹, Matthew J Thompson², Gail Hayward³, Carl J Heneghan³, Chris B Del Mar⁴, Rafael Perera³, Paul P Glasziou ⁴, Maroeska M Rovers⁵



Cochrane Database of Systematic Review. 2014, ssue 3. Art. No.: CD008115.

Oral corticosteroids versus placebo or NSAID for acute sinusitis

Patient or population: patients with acute sinusitis

Settings: primary and secondary care

Intervention: oral corticosteroids versus placebo or NSAID

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
	Assumed risk Control	Corresponding risk Oral corticosteroids versus placebo or NSAID			
Proportion of patients with resolution or improved symptoms at days 3 to 7 - Co-treatment with antibiotics	541 per 1000	758 per 1000 (585 to 980)	RR 1.40 (1.08 to 1.81)	869 (4 studies)	⊕⊕⊖⊝ low ^{2,3}



Figure 3. Forest plot of comparison: I Oral corticosteroids versus placebo or NSAID, outcome: I.I Proportion of patients with resolution or improved symptoms at days 3 to 7.

1.1.1 Co-treatment with a		Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI
1.1.1 Co-treatment with ar Cannoni 1990							
Cannoni 1990	70						
	79	103	40	100	20.4%	1.92 [1.47, 2.49]	
Gehanno 2000	158	208	136	209	28.1%	1.17 [1.03, 1.32]	-
Klossek 2004	60	103	49	104	20.5%	1.24 [0.95, 1.61]	+-
Ratau 2004 Subtotal (95% CI)	15	21 435	10	21 434	9.9% 78.8 %	1.50 [0.89, 2.53] 1.40 [1.08, 1.81]	•
Total events	312		235				

Oral corticosteroids versus placebo or NSAID for acute sinusitis

Patient or population: patients with acute sinusitis

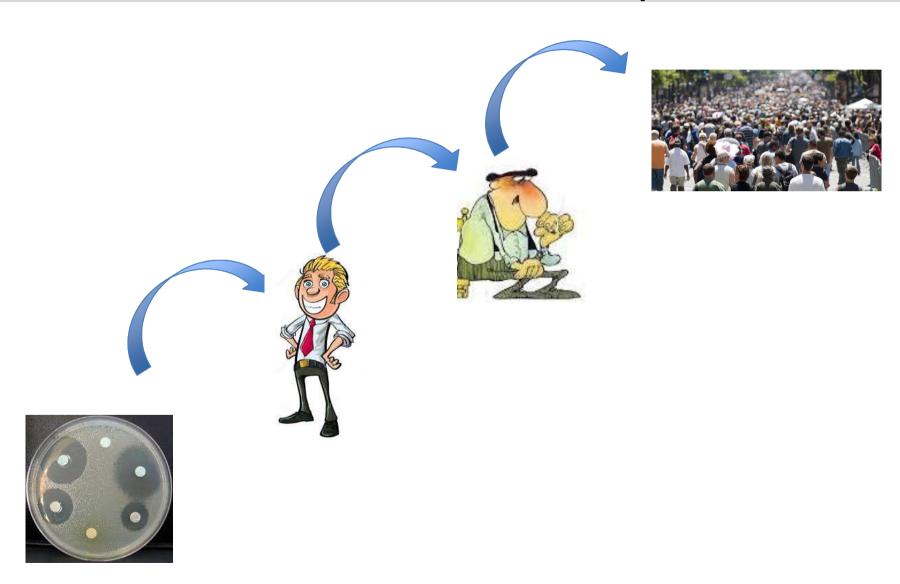
Settings: primary and secondary care

Intervention: oral corticosteroids versus placebo or NSAID

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
	Assumed risk	Corresponding risk			
	Control	Oral corticosteroids versus placebo or NSAID			
Proportion of patients	Study population		RR 1.40	869 (4 studies)	⊕⊕○○ !au23
with resolution or im- proved symptoms at days 3 to 7 - Co-treat- ment with antibiotics	541 per 1000	758 per 1000 (585 to 980)	(1.08 to 1.81)	(4 studies)	low ^{2,3}



Take-homme messages : choix d'un antibiotique



Take-homme messages : sinusite maxillaire aiguë non-compliquée

- Viral dans l'immense majorité des cas
 - Antalgique => paracétamol
 - Si échec : antibiotique
 - amoxicilline
 - puis [amoxicilline + AC] ou lévofloxacine
 - Intérêt discutable des corticoïdes en plus



- Si foyer dentaire, immunodépression ou non maxillaire
 - Antibiotique d'emblée : amoxicilline + acide clavulanique

Corticoïdes en monothérapie

CMAJ

RESEARCH

Systemic corticosteroid monotherapy for clinically diagnosed acute rhinosinusitis: a randomized controlled trial

Roderick P. Venekamp MD PhD, Marc J.M. Bonten MD PhD, Maroeska M. Rovers PhD, Theo J.M. Verheij MD PhD, Alfred P.E. Sachs MD PhD

CMAJ, October 2, 2012, 184(14)

Table 4: Adverse events reported during the study*					
	Group; no. c				
Adverse event	Prednisolone n = 88	Placebo n = 86	<i>p</i> value†		
First week					
Gastric complaint	11/86	8/84	0.5		
Diarrhea	14/86	10/85	0.4		
Increased appetite	14/86	7/84	0.1		
Mood disturbance	13/83	15/85	0.7		
Sleep disturbance	24/85	28/86	0.5		
Second week					
Gastric complaint	7/88	5/85	0.6		
Diarrhea	12/87	8/86	0.4		
Increased appetite	8/87	3/85	0.1		
Mood disturbance	9/88	11/84	0.6		
Sleep disturbance	12/87	15/85	0.5		

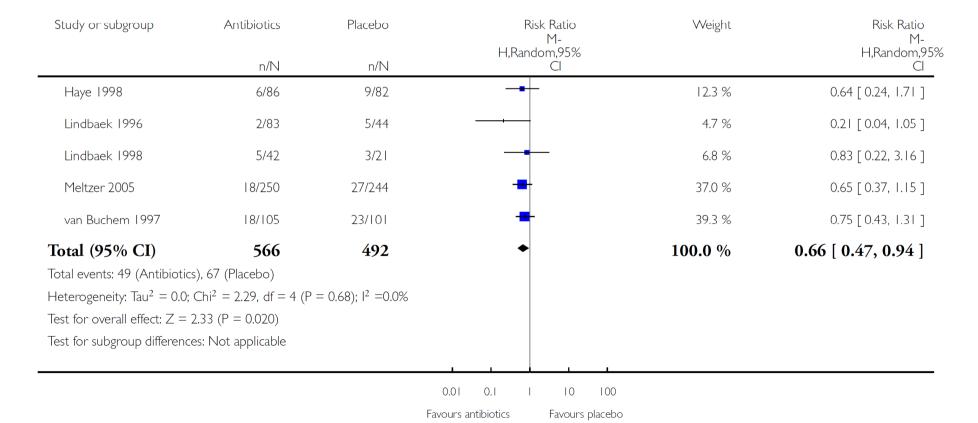
^{*}Two serious adverse events not related to drug use were reported: 1 hospital admission for anemia on day 49 of the study in the placebo group and 1 hospital admission for wasp sting–induced anaphylaxis on day 28 of the study in the prednisolone group. $t\chi^2$ test.

Analysis I.I. Comparison I Antibiotics versus placebo, Outcome I Clinical failure defined as a lack of full recovery or improvement at 7 to 15 days of follow-up.

Review: Antibiotics for acute maxillary sinusitis in adults

Comparison: I Antibiotics versus placebo

Outcome: I Clinical failure defined as a lack of full recovery or improvement at 7 to 15 days of follow-up



Antibiotics compared to other antibiotics for uncomplicated acute maxillary sinusitis

Patient or population: patients with uncomplicated acute maxillary sinusitis

Settings: primary care

Comparisons: non-penicillin antibiotic versus beta-lactamase sensitive penicillins (n = 8), non-tetracycline versus tetracycline (n = 5, of which one study was also included in the comparison of non-penicillin antibiotic versus beta-lactamase sensitive penicillins), macrolides versus amoxicillin-clavulanate (n = 11), cephalosporins versus amoxicillin-clavulanate (n = 10) and miscellaneous comparisons (n = 21)

Outcomes	Impact	No of participants (studies)	Quality of the evidence (GRADE)	Comments
Clinical failure rate (failure defined as a lack of full recovery or improvement or a lack of full recovery) Follow-ups: 7 to 15 days and 16 to 60 days		54 studies representing 10 different comparisons		Adverse effects ³

Figure 2. Funnel plot of comparison: I Oral corticosteroids versus placebo or NSAID, outcome: I.I Proportion of patients with resolution or improved symptoms at days 3 to 7.

