



ANTIVIRaux ACTIFS SUR LES VIRUS RESPIRATOIRES

Enseignant : F. Van Bambeke

FARM2129 – année 2008-2009

02/12/2008

influenza

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Médicaments des infections virales respiratoires

Adenovirus	-
Picornavirus	-
Entero	-
Rhino	-
Orthomyxovirus	
Influenza	inhibiteurs de neuraminidase : zamivir, oseltamivir adamanatanes (influenza A)
Paramyxovirus	-
Parainfluenza	-
Respiratory syncytial virus	ribavirine
SARS virus	-

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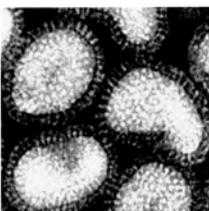
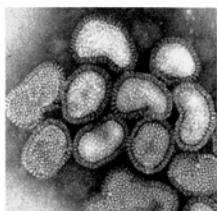
ANTIVIRAUX ACTIFS SUR LE VIRUS INFLUENZA

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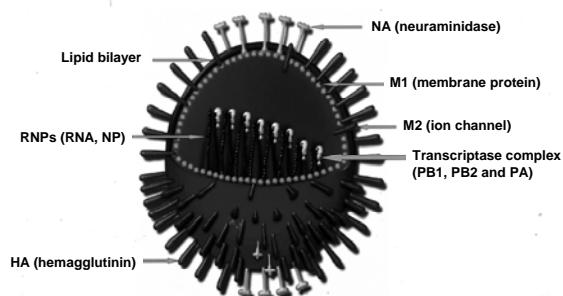
3

Le Virus de l'influenza: constituants



Electron micrographs of purified influenza virions. Hemagglutinin (HA) and neuraminidase (NA) can be seen on the envelope of viral particles. Ribonucleoproteins (RNPs) are located inside the virions.

Diagram of the influenza virus



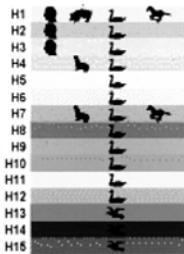
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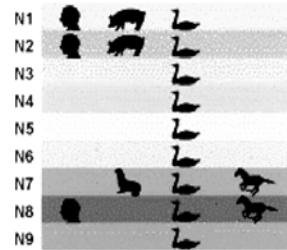
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Virus Influenza: Antigènes de surface

Distribution of Influenza A
Hemagglutinin Subtypes in Nature



Distribution of Influenza A
Neuraminidases in Nature



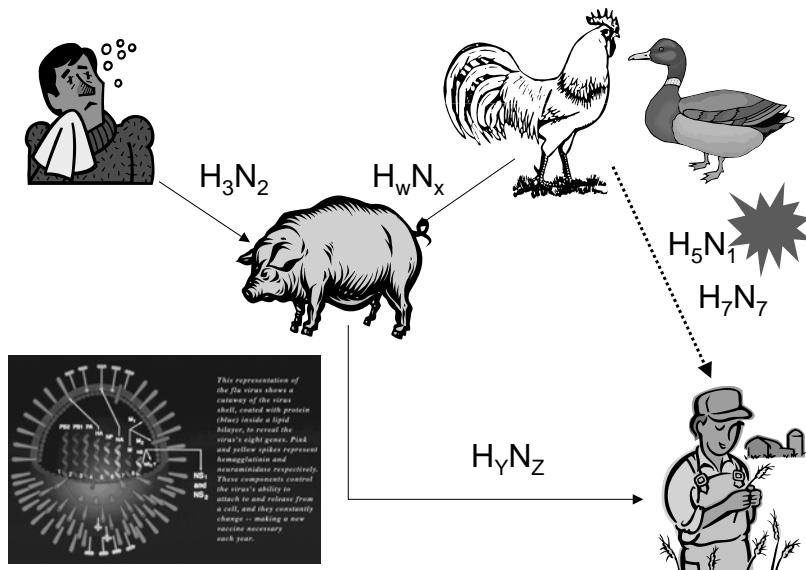
http://www.brown.edu/Courses/Bio_160/Projects1999/flu/mechanism.html

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Influenza A Virus Shift

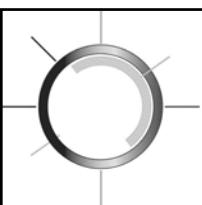


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La Grippe



Commissariat
interministériel Influenza

A distinguer :

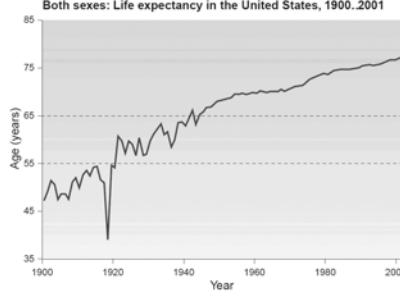
	Grippe saisonnière (Maladie humaine)	→ Attendue chaque hiver
	Cas humains de grippe aviaire (Zoonose)	→ Situation récente en Asie
	Pandémie de grippe (Zoonose évoluant en maladie humaine)	→ Pourrait émerger un jour

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Histoire des pandémies ...

1889-90	A/H2N8	
1900-03	A/H3N8	
1918-19	A/H1N1 <i>Spanish Flu</i>	
1957-58	A/H2N2 <i>Asian Flu</i>	
1968-69	A/H3N2 <i>Hong Kong Flu</i>	
(1977-78)	A/H1N1 <i>Russian Flu</i>	

Both sexes: Life expectancy in the United States, 1900...2001



Life expectancy from 1900 to 2001 showing the impact of the 1918 influenza pandemic. Data are adapted from the National Vital Statistics Reports, Vol. 52, No. 14, February 18, 2004.
(http://www.cdc.gov/nchs/data/dvs/nvsr52_14f12.pdf).

Australian Red Cross 1918



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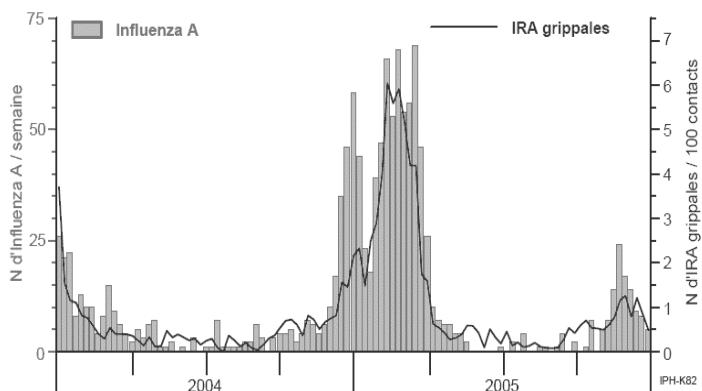
Evolution des "maladies grippales" et de l'isolement d'*Influenza A*

Laboratoires Vigies

Influenza A

- l'évolution du nombre d'*Influenza A* est comparable à celle du nombre d'IRA grippales enregistrées par le réseau de médecins généralistes participant au programme de surveillance des IRA et de la grippe (figure 7).

Figure 7 : Influenza A : comparaison entre l'évolution par semaine du nombre d'*Influenza A* et celle des IRA grippales (2004-2005)



http://www.ipb.fgov.be/epidemio/epifr/plabfr/plabanfr/05_082f_v.pdf

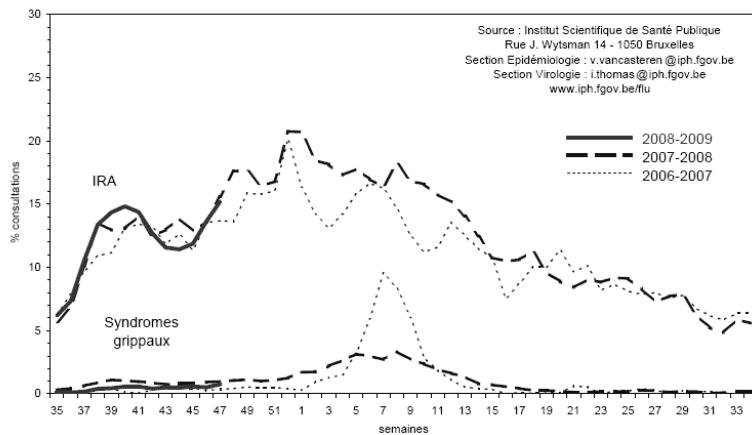
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Epidémiologie annuelle des infections par *Influenza*

Evolution du pourcentage des syndromes grippaux et des infections respiratoires aigües (IRA)
enregistrés par les Médecins Vigies



Source : Institut Scientifique de Santé Publique
Rue J. Wytsman 14 - 1050 Bruxelles
Section Épidémiologie : v.vanasteren@ipb.fgov.be
Section Virologie : i.thomas@ipb.fgov.be
www.ipb.fgov.be/flu

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<http://www.ipb.fgov.be/flu/FR/22FR.htm>

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Voies de transmission

1. aérosolisation



2. Voie aérienne



3. Objets contaminés

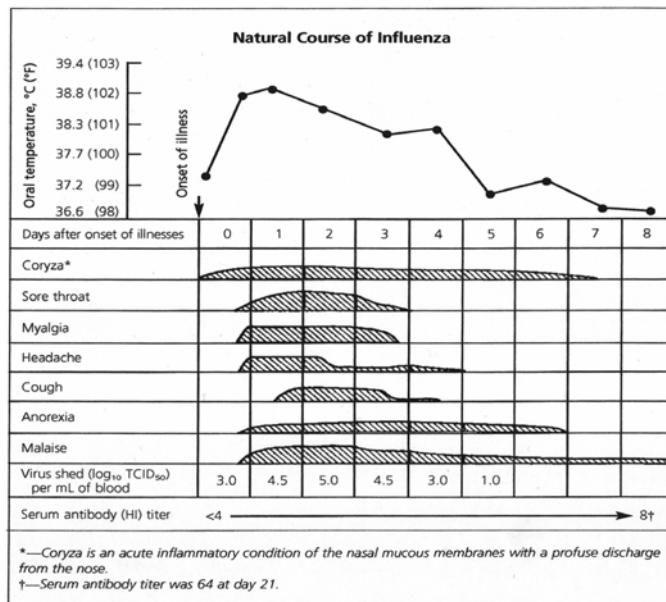
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La grippe: signes cliniques



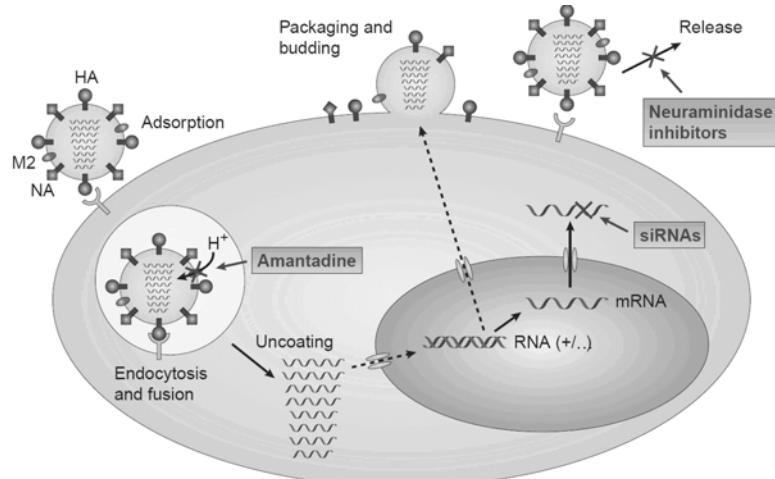
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Cibles des antiviraux



P. Palese *Nature Medicine* 10 : 2004

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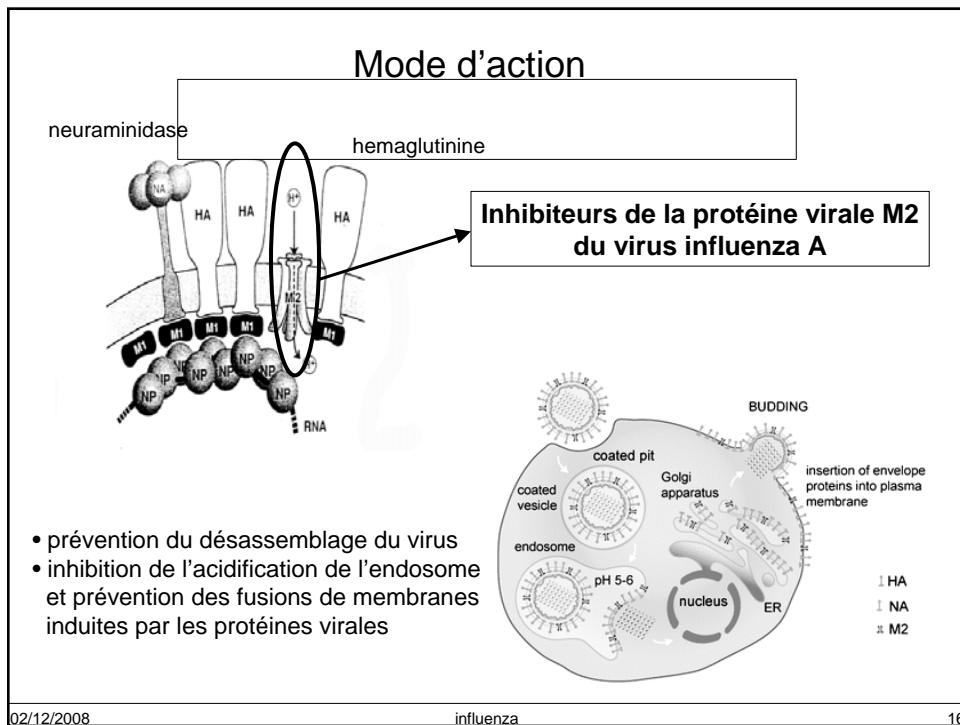
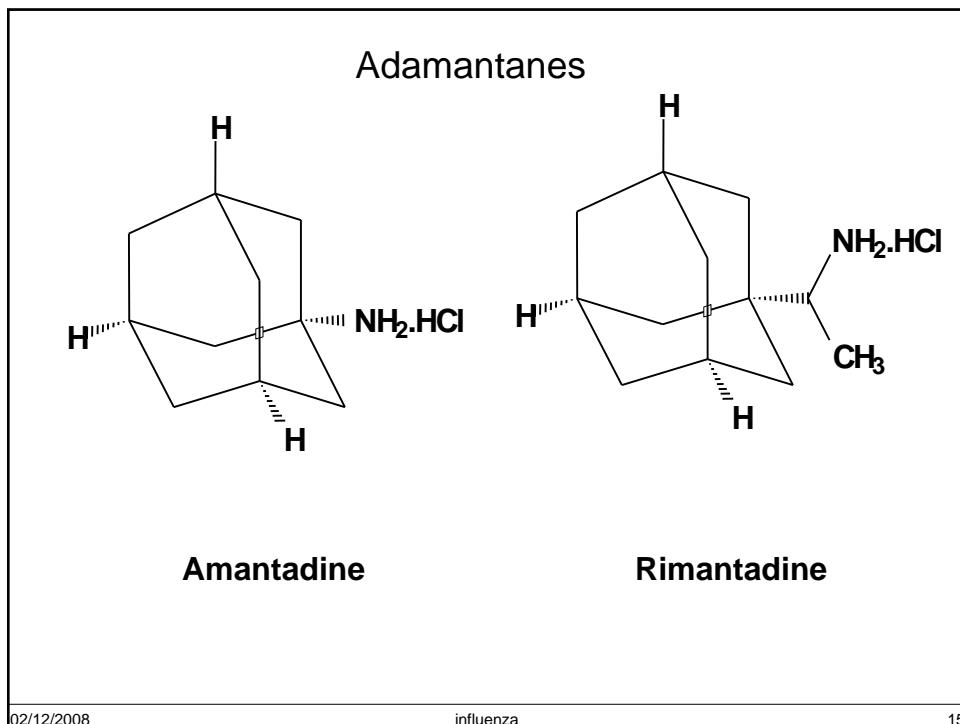
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ADAMANTANES

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INHIBITEURS DE NEURAMINIDASE

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La neuraminidase

Moscona, NEJM (2005) 353:1363-1373

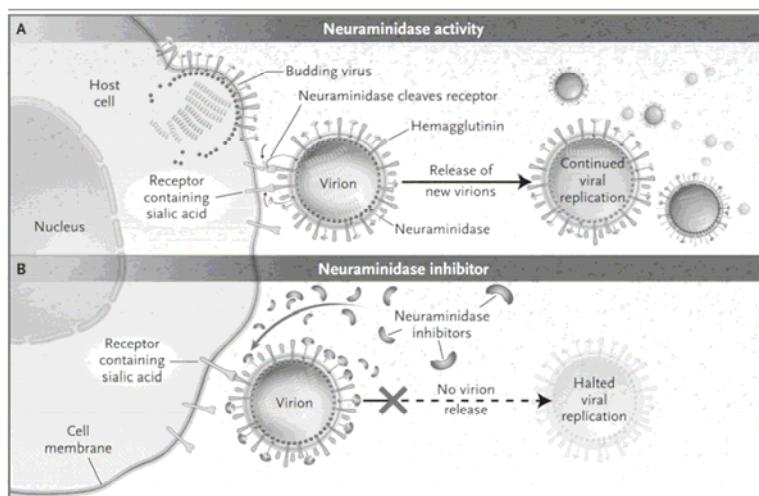


Figure 1. Mechanism of Action of Neuraminidase Inhibitors.

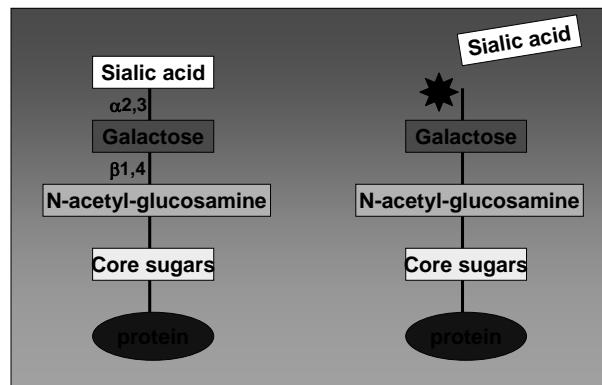
Panel A shows the action of neuraminidase in the continued replication of virions in influenza infection. The replication is blocked by neuraminidase inhibitors (Panel B), which prevent virions from being released from the surface of infected cells.

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La neuraminidase



La neuraminidase clive l'acide sialique des glycoprotéines de surface cellulaire auquelles sont attachées les nouvelles particules virales

Functions:

- favorise la libération des particules virales
- détruit les récepteurs de surface reconnus par les hémagglutinines
- préviennent l'agrégation des virus à la surface cellulaire
- préviennent l'inactivation par le mucus respiratoire

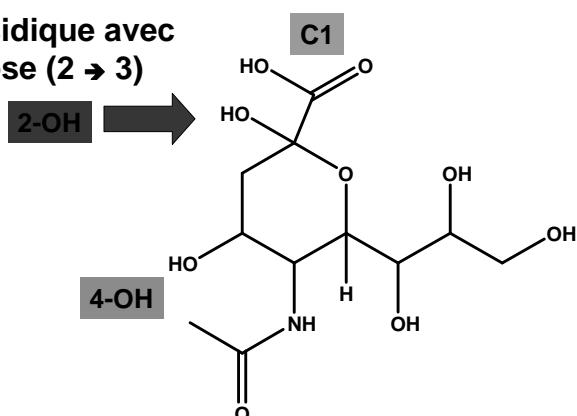
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L'acide sialique

Lien osidique avec galactose (2 → 3)

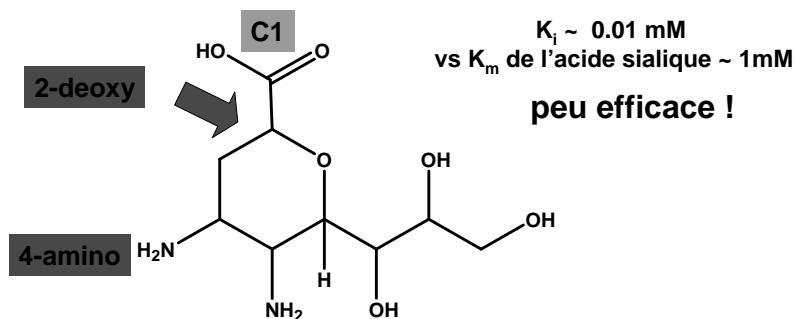


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Découverte du premier inhibiteur ... 1969 !



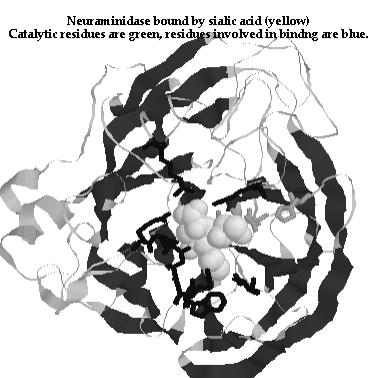
2,4-dideoxy-2,3-didehydro-4-amino-D-N-acetylneuraminic acid
Meindl et al., Hoppe-Seyler's Z. Physiol. Chem., 350:1088-1092, 1969

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De 1969 à 1993...



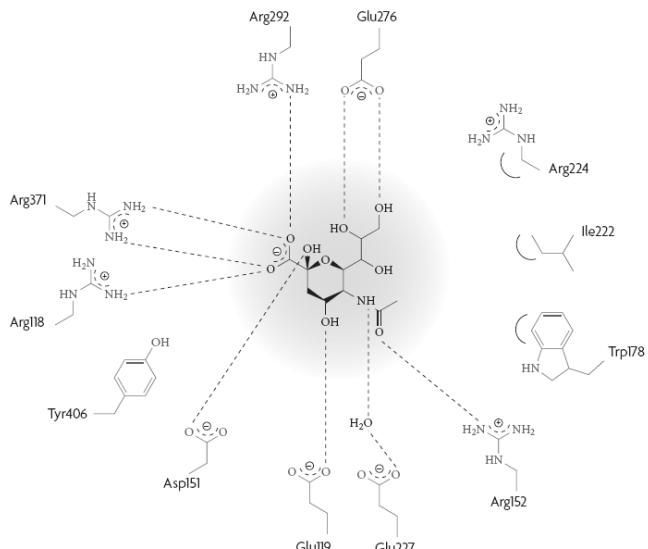
Colman et al., Nature (1983) 303: 41-44

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Site actif de la neuraminidase



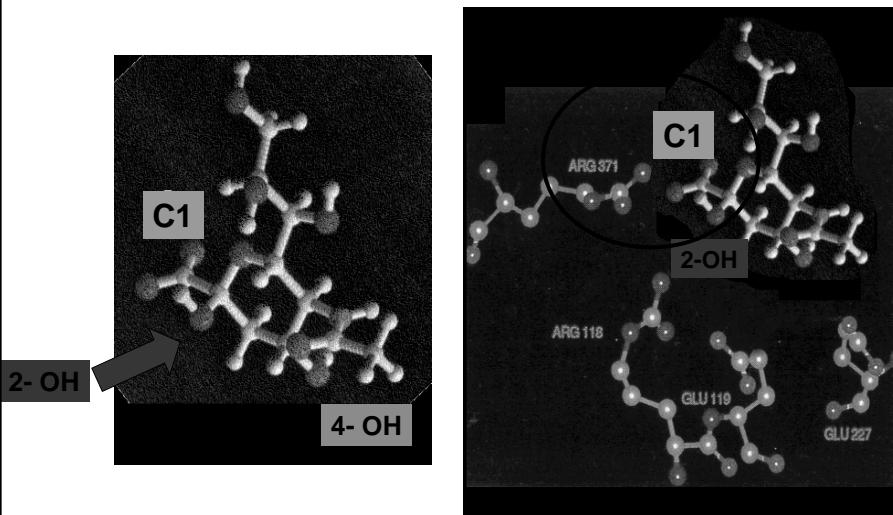
Von Itzstein., Nature Drug Discovery (2007) 6: 967

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De l'acide sialique au zanamivir... (1)



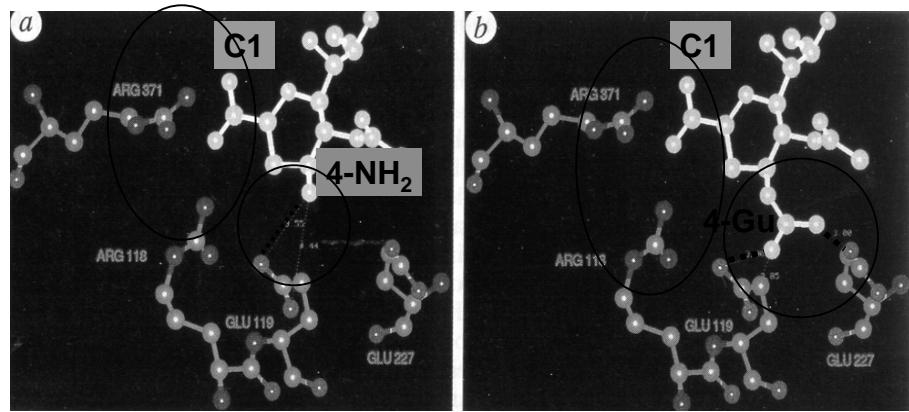
acide sialique ou N-acétyl-neuraminique
interaction COOH avec Arg 371

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De l'acide sialique au zanamivir... (2)



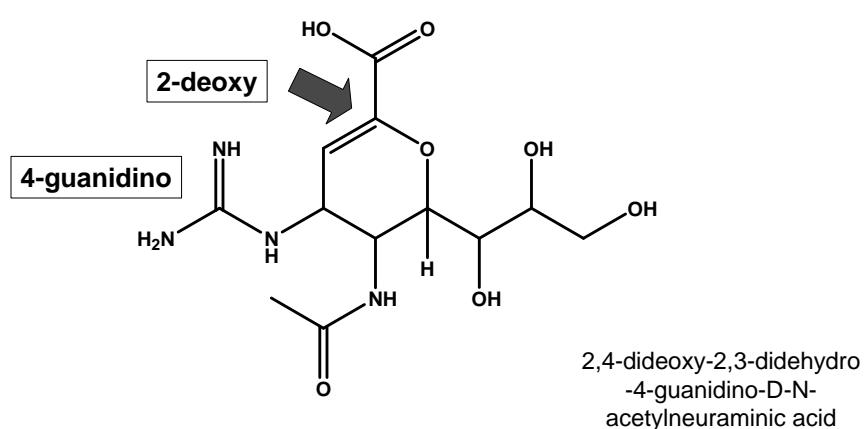
interaction avec 1 puis 2 résidus conservés

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Le zanamivir

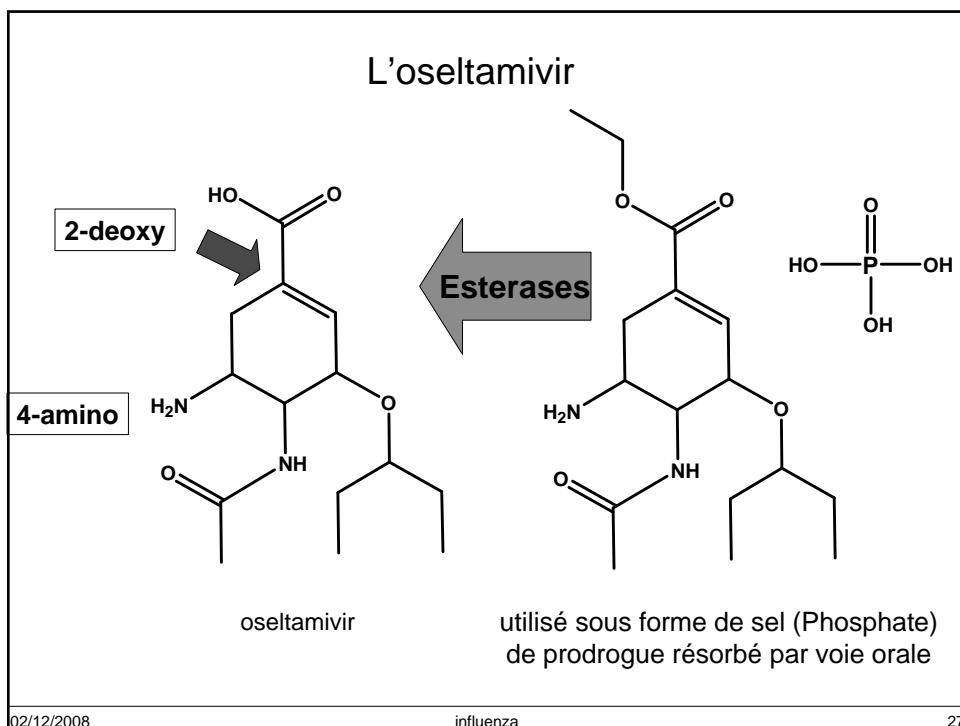


von Itzstein et al., Nature (1993) 363: 418-423, 1993

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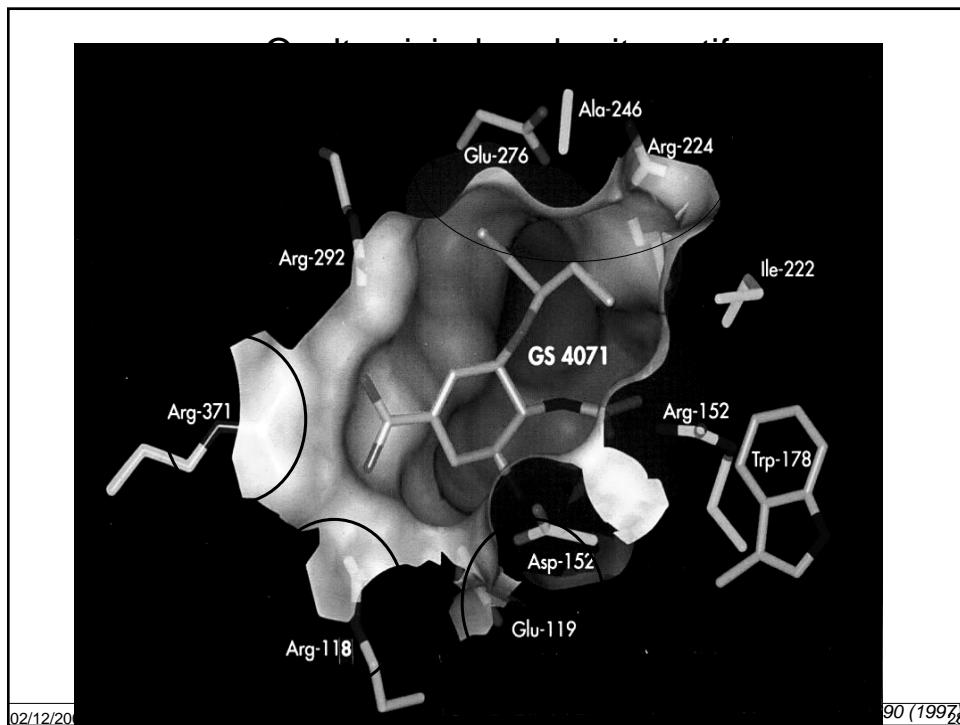
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90 (1997)₂₈

Résistance: mutation de la cible

Neuraminidase

119 Glu → Gly:

- Résistance au zanamivir
(Glu 119 interagit avec guanidinium)

292 Arg → Lys (R292K):

- Résistance à l'oseltamivir
(Arg 292 interagit avec acide carboxylique mais conformation de la poche enzymatique adhoc pour le zanamivir)

274 His → Tyr (H274Y) et 294 Asn → Ser (N294S)

- Résistance à l'oseltamivir

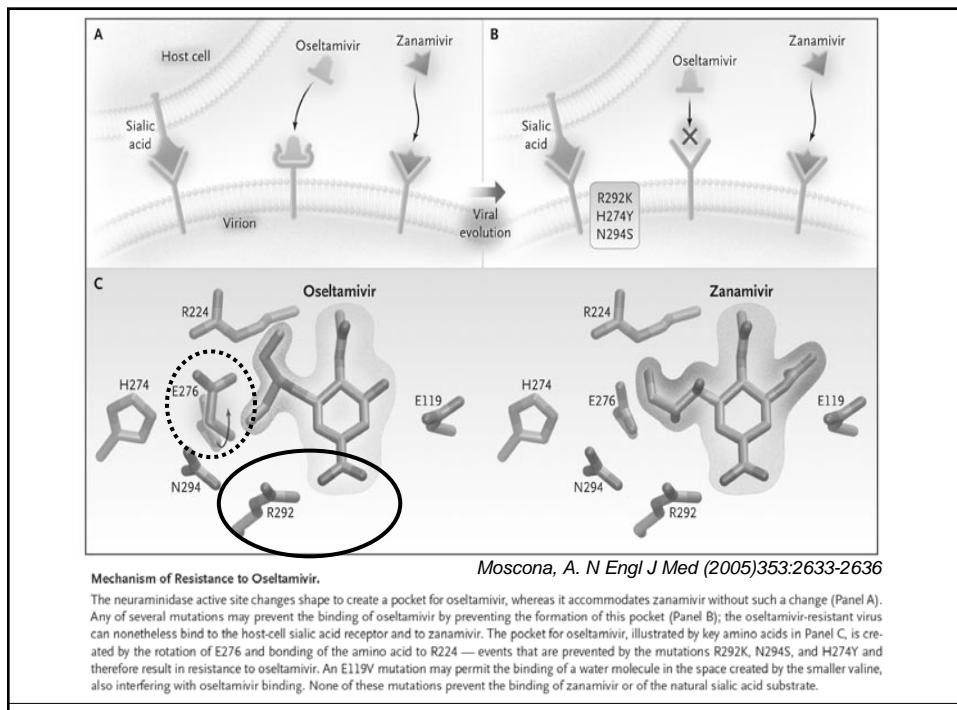
Hemagglutinin

Certaines mutations (198 Thr → Ile) diminuent l'affinité pour le récepteur

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Comparaison des inhibiteurs de neuraminidase

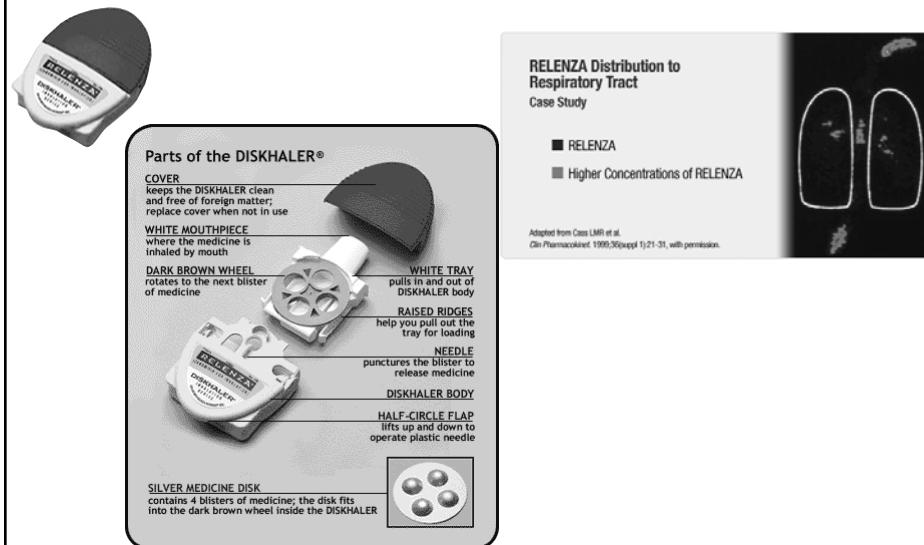
propriété	zanamivir	oseltamivir
spectre	Influenza A et B	
Voie d'administration	Inhalation 10 mg 2X/jour	Voie orale: 75-150 mg 2 X/jour
Traitement (5 jours)	↓ durée des symptômes – grippe saisonnière	
Prophylaxie	4 semaines: ↓ du nb de personnes malades	6 semaines: ↓ du nb de personnes malades
Activité sur H5N1	Peu efficace chez les patients contaminés	
tolérance	Bonne sauf path. respir.	bonne

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Zanamivir: voie d'administration



<http://www.relenza.com/how-to-use-diskhaler.jsp?languages=French>

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Zanamivir: comment l'administrer correctement ?



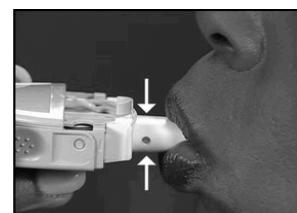
<http://www.relenza.com/how-to-use-diskhaler.jsp?languages=French>

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Zanamivir: comment l'administrer correctement ?



1 dose: = 2 cupules

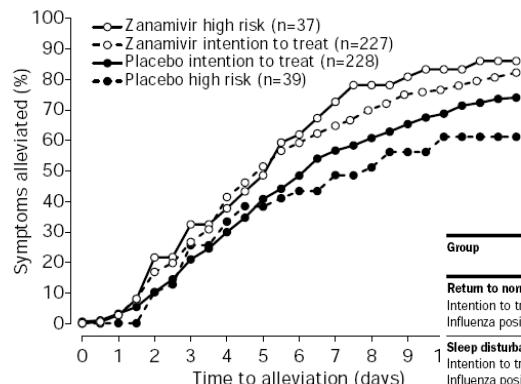
<http://www.relenza.com/how-to-use-diskhaler.jsp?languages=French>

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Zanamivir: efficacité clinique



On gagne 1-2 jours ...

Group	No of patients	Median	No of patients	Median	Difference (95% CI)	p
Return to normal activities (days)						
Intention to treat	228	9.0	227	<7.0	2.0 (0 to 4.0)	<0.001
Influenza positive	160	9.0	161	<7.0	2.0 (0.25 to 4.0)	<0.001
Sleep disturbance (days of 13)						
Intention to treat	228	3	223	3.0	0 (-1.0 to 1.0)	0.088
Influenza positive	160	3.0	159	2.0	1.0 (0 to 1.5)	0.047
Number of paracetamol tablets (days 1-4)						
Intention to treat	228	12	224	14.0	-2 (-6 to 0)	0.291
Influenza positive	160	13	159	14.0	-1 (-5 to 2)	0.854
Number of cough mixture spoonfuls (days 1-14)						
Intention to treat	228	9	224	7.0	2 (-3 to 5)	0.738
Influenza positive	160	12	159	7.0	5 (-1 to 9)	0.045

Table 3: Sleep disturbance, return to normal activities, and use of relief medications

The Lancet (1998) 352: 1877-1881

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Zanamivir: efficacité en prophylaxie

Table 2. Efficacy of Zanamivir in Prevention of Influenza Infection and Disease*

Outcome	Frequencies in Study Groups, No. (%)		Odds Ratio (95% CI)	Estimated Risk Ratio (95% CI)	Efficacy, 1 - Risk Ratio (95% CI), %
	Placebo (n = 554)	Zanamivir (n = 553)			
Laboratory-confirmed clinical influenza	34 (6)	11 (2)	0.31 (0.14-0.64)†	0.33 (0.17-0.61)	67 (39-83)
Laboratory-confirmed influenza with fever	19 (3)	3 (<1)	0.15 (0.03-0.53)†	0.16 (0.06-0.45)	84 (55-94)
All febrile illnesses	58 (10)	33 (6)	0.54 (0.34-0.86)‡	0.57 (0.38-0.86)	43 (14-62)
Influenza infection with or without illness	77 (14)	53 (10)	0.66 (0.44-0.97)§	0.69 (0.50-0.96)	31 (4-50)

*CI indicates confidence interval.

†P≤.001.

‡P = .009.

§P = .03.

JAMA (1999) 282: 31-35

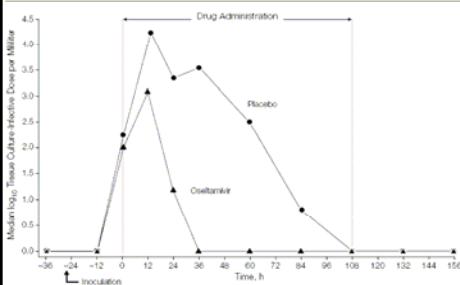
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Et l'oseltamivir ?

Figure 3. Effect of Oral Oseltamivir Treatment on Vital Titers in Nasal Lavages Following Experimental Influenza A/Texas/36/91(H1N1) Infection



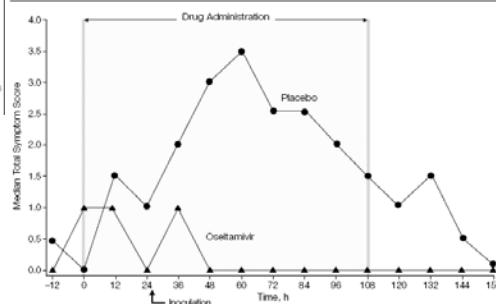
Traitements :
On gagne 1-2 jours ...

JAMA (1999) 282:31-36

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Prophylaxie : efficace

Figure 2. Effect of Oral Oseltamivir Prophylaxis on Illness Following Experimental Influenza A/Texas/36/91(H1N1) Inoculation



The total symptom score area under the curve value was lower in the combined oseltamivir groups ($n = 21$) compared with placebo ($n = 12$); $P = .02$. Fourteen symptoms related to influenza were included in the score.

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Zanamivir: effets secondaires et contre-indications

Important Safety Information

RELENZA IS NOT RECOMMENDED FOR TREATMENT OR PROPHYLAXIS OF INFLUENZA IN INDIVIDUALS WITH UNDERLYING AIRWAY DISEASE (SUCH AS ASTHMA OR CHRONIC OBSTRUCTIVE PULMONARY DISEASE).

- ◆ Serious cases of bronchospasm, including fatalities, have been reported during treatment with RELENZA in patients with and without underlying airway disease. Many of these cases were reported during postmarketing, and causality was difficult to assess
- ◆ RELENZA SHOULD BE DISCONTINUED IN ANY PATIENT WHO DEVELOPS BRONCHOSPASM OR DECLINE IN RESPIRATORY FUNCTION; immediate treatment and hospitalization may be required
- ◆ RELENZA has not been proven effective for treatment of influenza in individuals with underlying airways disease.
- ◆ If treatment with RELENZA is considered for a patient with underlying airway disease, the potential risks and benefits should be carefully weighed. If a decision is made to prescribe RELENZA for such a patient, this should be done only under conditions of careful monitoring of respiratory function, close observation, and appropriate supportive care including availability of fast-acting bronchodilators
- ◆ Common adverse events in treatment and prophylaxis studies with RELENZA were nausea, diarrhea, sinusitis, viral respiratory infections, headaches, nasal signs and symptoms. The incidence of these adverse events was similar in both groups for RELENZA and placebo-treated groups

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Bénéfice potentiel des inhibiteurs de neuraminidase

Thérapeutique:

- ↘ durée des symptômes de 1-2 jours
- ↘ risque de transmission du virus
- ↘ risque de complications (sinusite, bronchite)
- ↘ usage d'antibiotiques

Prophylaxie:

- Prévention saisonnière de l'infection

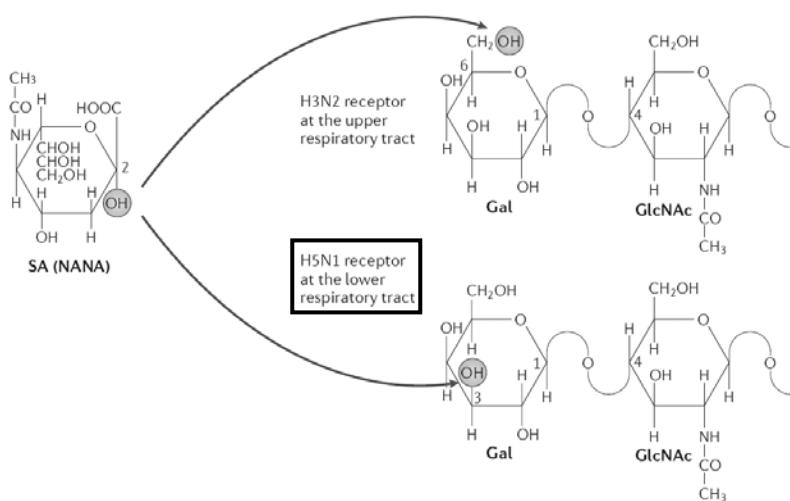
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Panique à bord: A quand la grippe aviaire ?

Pourquoi est-elle si redoutable ?



De Clercq, *Nature Drug Discovery* (2006) 5:1015-25

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Panique à bord: A quand la grippe aviaire ?

Stockpiling of Antivirals



Objectif :
30% de la population belge devrait
avoir accès au traitement dès 2008



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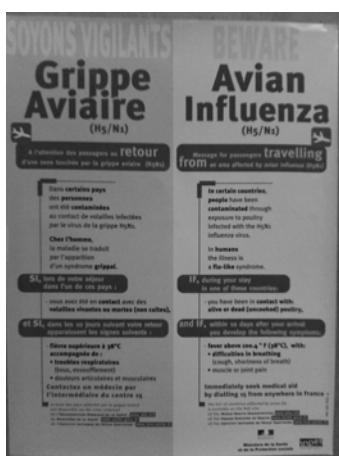
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Panique à bord: A quand la grippe aviaire ?

Conseils aux voyageurs

- Vérifier la liste des pays concernés
- Éviter le contact avec la volaille
- Hygiène des mains en toutes circonstances
- Hygiène culinaire, dont la cuisson
- Pas d'antiviraux en préventif
- Consulter un médecin si fièvre < 7 jours après le retour



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Pharmacothérapie de la grippe

- Traitement symptomatique
 - antipyrétiques
 - éviter aspirine chez les enfants (syndrôme de Reye)
 - selon les symptômes: décongestionnants, antitussifs
- Traitement antiviral
 - intérêt assez limité en traitement ...
 - commencer < 36 heures après le début des symptômes
 - prophylaxie de l'environnement familial ?
- Suivre la survenue de complications (personnes à risque!)
- Vaccination !

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Recommandations de vaccination



- Personnes à risque de complication:
 - > 65 ans
 - personnes vivant en institutions
 - co-morbidités
- Personnes susceptibles de transmettre la maladie à des personnes à risque (personnel médical, ...)

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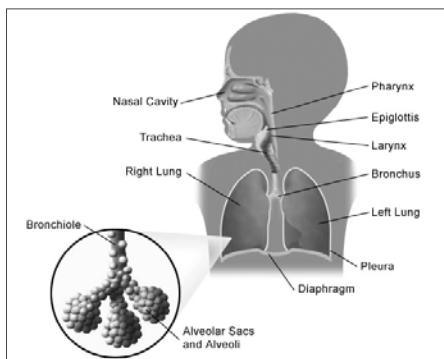
ANTIVIRAUX ACTIFS SUR RESPIRATORY SYNCYTIAL VIRUS (RSV)

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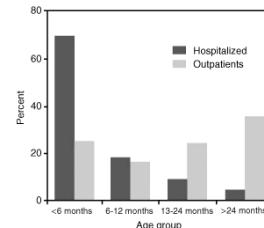
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Infections respiratoires à RSV



Syndrome	Percentage Caused by RSV
Bronchiolitis	43-90
Pneumonia	5-40
Tracheobronchitis	10-30
Croup	3-10
Asymptomatic	0.3



Difference in age distribution of inpatients and outpatients with respiratory syncytial virus infection in Rochester, NY.

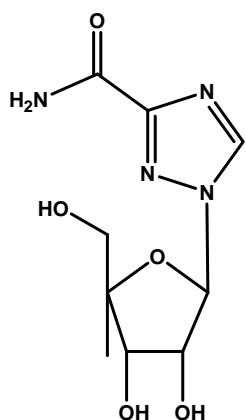
Of infants requiring hospitalization, 70% were younger than 6 months. In comparison, 25% of the children treated as outpatients were younger than 6 months and 38% were older than 2 years.

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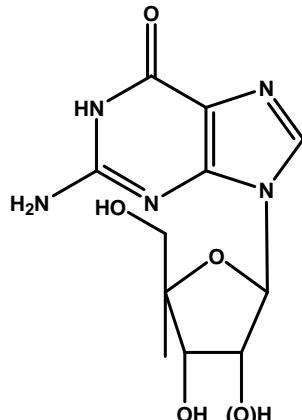
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la ribavirine, un analogue de la guanosine



RIBAVIRINE



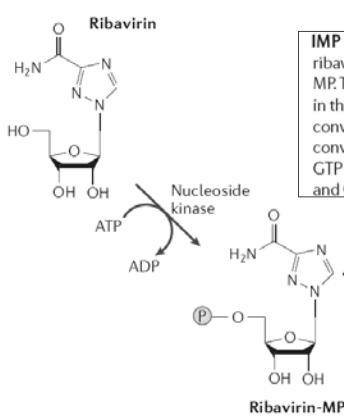
(deoxy)GUANOSINE

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Mode d'action de la ribavirine



IMP dehydrogenase inhibition.

ribavirin, which is converted intracellularly to its 5'-monophosphate derivative, ribavirin-MP. The latter inhibits inosine 5'-monophosphate (IMP) dehydrogenase, a crucial enzyme in the biosynthesis of RNA, including viral RNA. IMP dehydrogenase is responsible for the conversion of IMP into xanthosine 5'-monophosphate (XMP) which, in turn, is further converted to GMP (guanosine 5'-monophosphate), GDP (guanosine 5'-diphosphate) and GTP (guanosine 5'-triphosphate). The latter serves as substrate, together with ATP, UTP and CTP, in the synthesis of RNA.

De Clercq, Nature Drug Discovery, 2006

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Indications et propriétés pharmacologiques

Aérosol : traitement des infections à RSV
mais peu utilisé ... affection bénigne

Voie générale: adjuvant dans le traitement de l'hépatite C

Effets secondaires:

- Détérioration de la fonction respiratoire
- Anémie hémolytique + hypoplasie médullaire
(Concentration dans les globules rouges)

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Traitement des infections à RSV

Traitement symptomatique

- antipyrétique
- apport de fluides
- oxygène si nécessaire
- bronchodilatateurs
- corticoïdes



Traitement antiviral

(ribavirine)



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