

ANTIFONGIQUES



Enseignant : F. Van Bambeke

FARM2233 – année 2011-2012

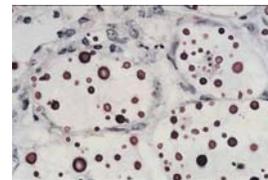
Il y a champignon ...et champignon



onychomycose

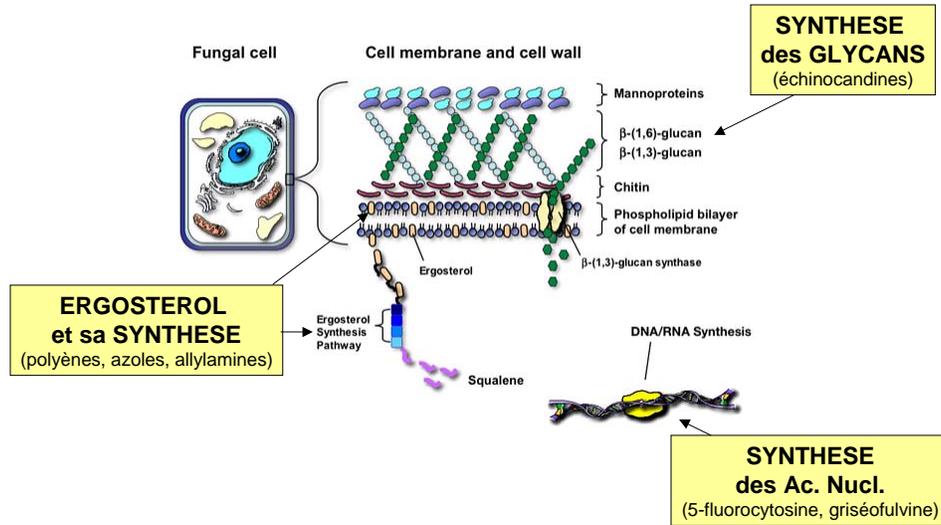


candidose



cryptococose pulmonaire

Cibles des médicaments antifongiques

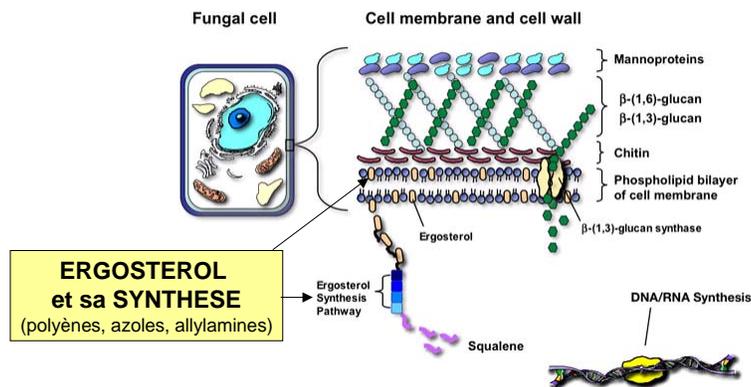


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Cibles des médicaments antifongiques



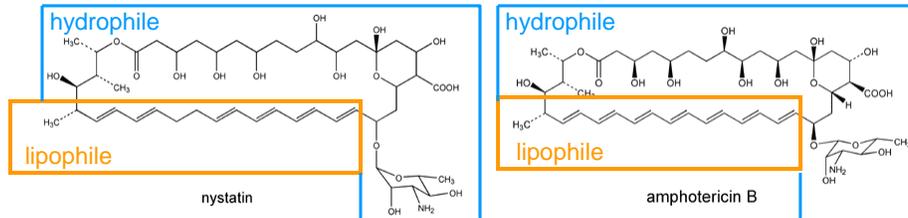
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Polyènes antifongiques

Molécules amphiphiles mais volumineuses
→ mal résorbées

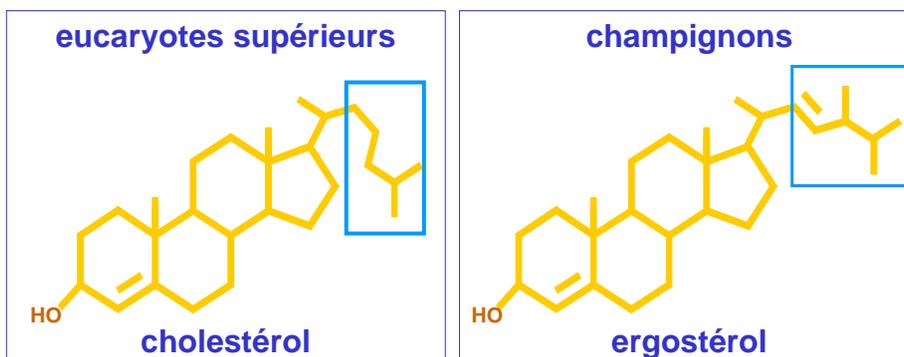


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Cible pharmacologique: les stérols membranaires



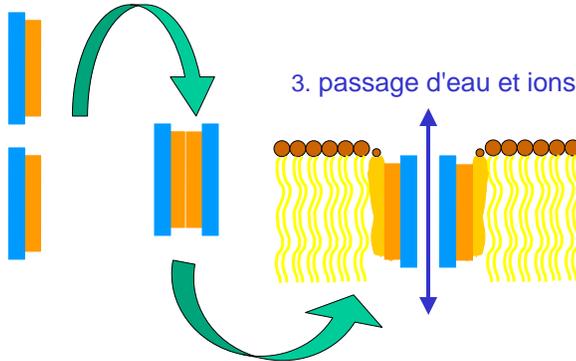
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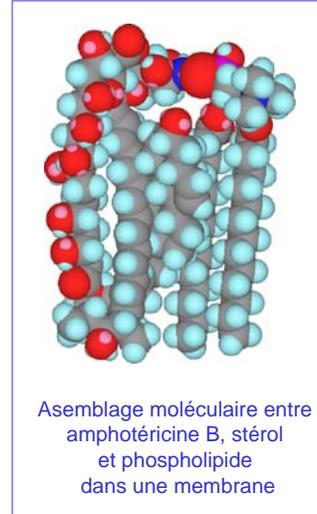
Mécanisme d'action des polyènes antifongiques

1. formation de dimères
peu solubles...



2. Insertion dans la membrane
au niveau de l'ergostérol

3. passage d'eau et ions



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Mécanisme d'action des polyènes antifongiques

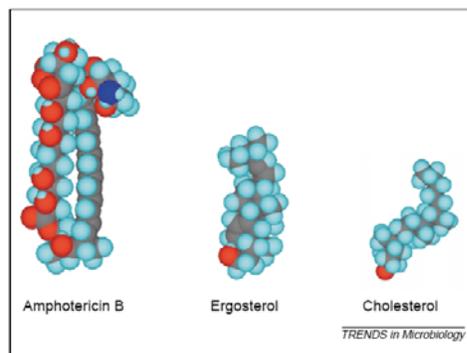
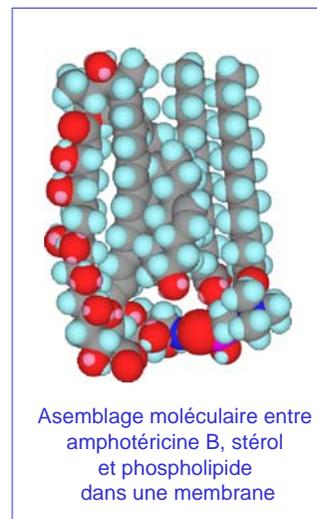


Fig. 3. The polyene antifungal agent, amphotericin B, ergosterol and cholesterol, visualised in three dimensions. Ergosterol, the sterol found in fungal cell membranes, retains a cylindrical shape in all rotations and binds better to the hydrophobic (right-hand) side of the amphotericin B molecule than does cholesterol, with its sigmoid structure. Cholesterol is the membrane sterol found in mammalian cells; the differential binding affinity of amphotericin B for the two sterols is the basis of its selective antifungal action.



Odds, Trends in Microbiology (2003) 11:272-279

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Polyènes antifongiques: propriétés pharmacologiques

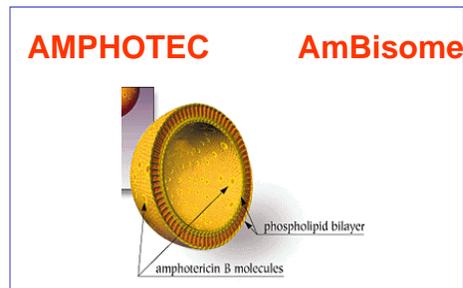
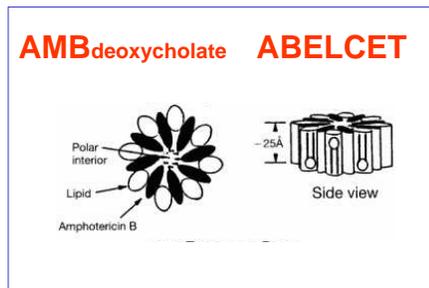
+	-
Fongicides Spectre large, peu de résistance	Résorption mauvaise voire nulle (nystatine) Antagonisme avec les azolés Faible spécificité → toxicité: <ul style="list-style-type: none"> • immédiate: fièvres, frissons, nausées, vomissements, hypotension, arythmies, ... • à court terme: néphrotoxicité • à moyen terme: anémie
nystatine	Amphotéricine B
usage topique <ul style="list-style-type: none"> • oral: candidoses, dermatophyties • décontamination intestinale 	<ul style="list-style-type: none"> • mycoses systémiques • mycoses oropharyngées et digestives graves

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Formes lipidiques de l'amphotéricine



The various lipid formulations of AMB have demonstrated antifungal efficacy at least equivalent to the conventional product with significantly reduced nephrotoxicity.
 In: THE USE OF LIPID FORMULATIONS OF AMPHOTERICIN B IN CANCER PATIENTS -- Rod Quilitz, PharmD, Department of Pharmacy, H. Lee Moffitt Cancer Center & Research Institute, University of San Francisco, Cal.
<http://www.moffitt.usf.edu/cancjrnl/v5n5/department3.html>

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Formes lipidiques: propriétés pharmacologiques

Parameter	AMB deoxycholate	AMB lipid complex	Liposomal AMB
Dosage* (mg/kg per day)	0.5–1.5	5	3–5 (or higher)
Maximum serum concentration [†]	—	Lower	Higher
Infusion-related toxicity [‡]	High (50%–60%)	Moderate (20%–40%)	Mild (10%–20%)
Decrease in serum potassium	****	**	**
Anaemia	****	+	+
Nephrotoxicity	**** (up to 80%)	+ (15%–25%)	+ (10%–20%)
Prevention of infusion-related toxicity [§]	Required	Required	Generally not required

* Commonly prescribed treatment doses, dose varies with pathogen. High-dose liposomal AMB required for zygomycete infection (≥ 5 mg/kg per day). [†] In comparison with AMB deoxycholate. [‡] Includes fever, chills, headache, joint and muscle pain, and hypotension. Before therapy, a test dose is recommended to identify patients in whom severe infusion-related reactions might occur. [§] Usually comprises "cocktail" of antipyretic, antiemetic and antihistamine drugs. Value of corticosteroids not proven.

Mais cher et administration intraveineuse obligatoire
→ indications limitées

Chen et al. Med J Aust. (2007) 187:404-9.

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Formes lipidiques: indications selon la notice

Abelcept	AmBisome
<p>Infections fongiques généralisées: Abelcept est recommandé dans le traitement des infections fongiques graves, chez les patients n'ayant montré aucune amélioration avec l'amphotéricine B conventionnelle, ou chez les patients ayant développé une insuffisance rénale lors du traitement à l'amphotéricine B, même lorsque ce dernier avait été administré en même temps qu'un litre de solution saline physiologique par jour.</p>	<p>AmBisome est destiné au traitement des formes graves de mycoses systémiques et/ou profondes des patients ne répondant pas à l'amphotéricine B conventionnelle ou des patients représentant une contre-indication à l'administration de celle-ci due à l'existence de lésions rénales. Une réponse positive a été obtenue chez 80 % de ces patients traités pour une candidiase systémique, chez 70 % des patients traités pour une aspergillose et chez 100 % des patients traités pour une cryptococcose....</p>

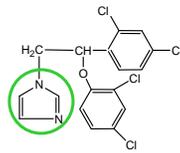
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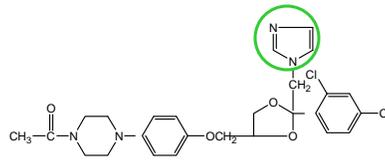
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Dérivés azolés

imidazoles

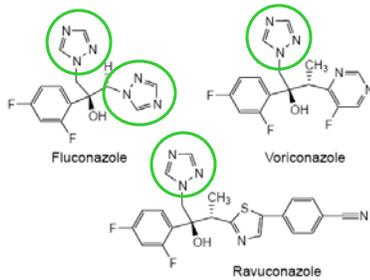


Miconazole



Ketoconazole

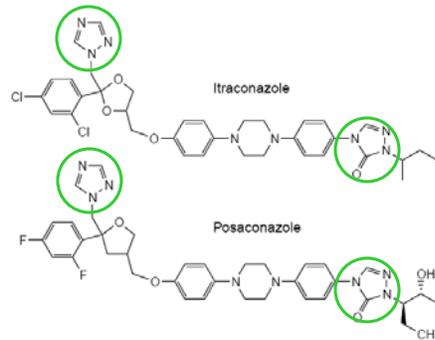
triazoles



Fluconazole

Voriconazole

Ravuconazole



Itraconazole

Posaconazole

Biosynthèse de l'ergosterol

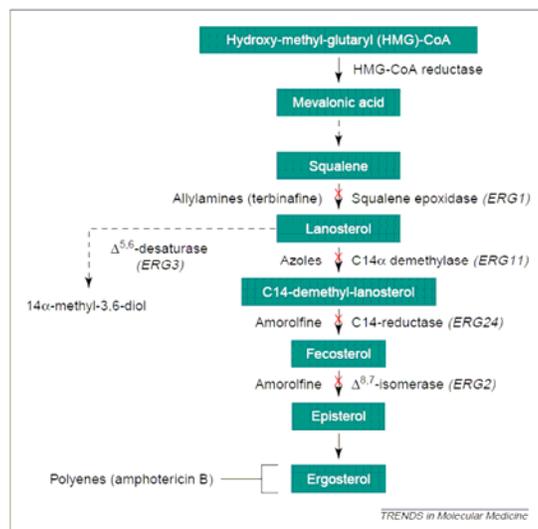
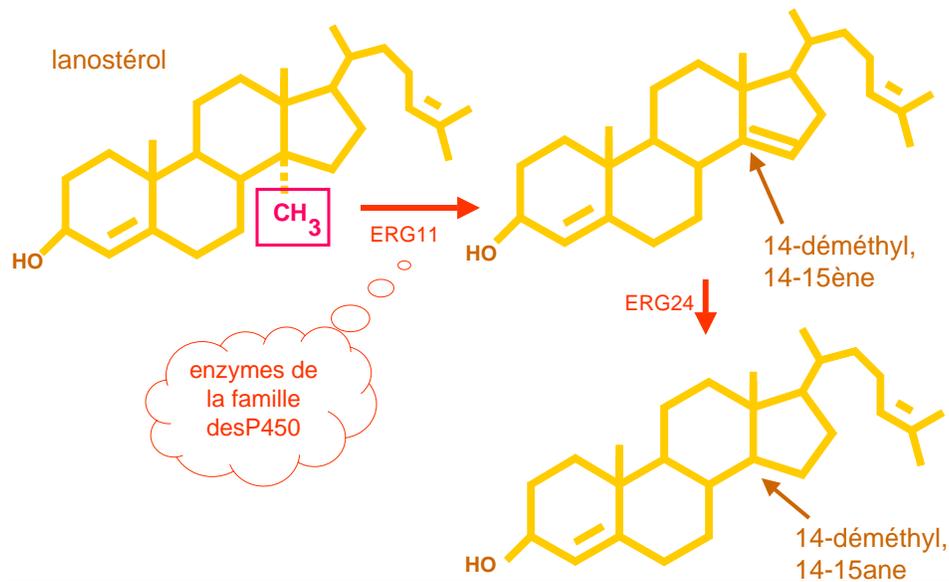


Fig. 2. Mechanism of action of antifungal drugs affecting the ergosterol biosynthetic pathway. The target enzymes are reported on the right with encoding genes in parentheses, whereas the antifungal drugs are reported on the left of the arrows indicating the sequential steps of sterol biosynthesis.

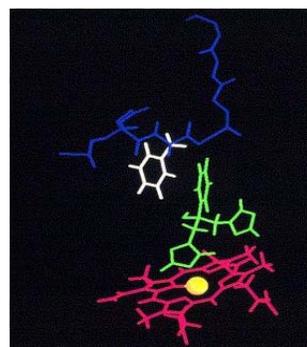
Biosynthèse de l'ergosterol



Dérivés azolés et site actif de Erg11p



Le cytochrome P450 possède un hème essentiel à l'activité oxydo-reductasique



Le cycle imidazole ou triazole se colle à l'hème ...

<http://users.aber.ac.uk/cca/p450/intro.html>

Dérivés azolés et site actif de Erg11p

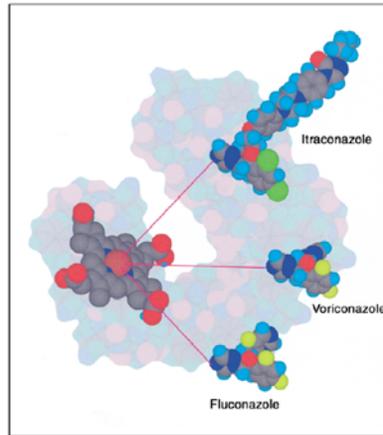


Fig. 4. Cartoon giving an approximate impression of the protoporphyrin moiety located at the active site of Erg11p (CyP51p), the cytochrome P450 enzyme target for imidazole and triazole antifungals. Three triazole antifungals, itraconazole (top), fluconazole (centre) and voriconazole (bottom) are shown in comparable orientations. Arrows link the azole nitrogen atom to the iron atom where the azoles bind to block the active site of the enzyme. The different side chains attached to the common azole pharmacophore in the three examples shown will obviously bind differently to the surrounding regions of the whole P450 protein.

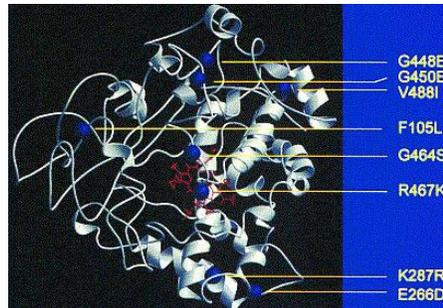
Odds, Trends in Microbiology (2003) 11:272-279

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Mécanismes de résistance aux dérivés azolés



Mechanism	Caused by:	Comments
Alteration in drug target (14 α -demethylase)	Mutations which alter drug binding but not binding of the endogenous substrate	Target is active (i.e., can catalyze demethylation) but has a reduced affinity towards azoles
Alteration in sterol biosynthesis	Lesions in the $\Delta^{5(6)}$ -desaturase	Results in accumulation of 14 α -methyl fecosterol instead of ergosterol
Reduction in the intercellular concentration of target enzyme	Change in membrane lipid and sterols; overexpression of specific drug efflux pumps (CDR1, PDR5, and BEN ¹)	Poor penetration across the fungal membrane; active drug efflux
Overexpression of antifungal drug target	Increased copy number of the target enzyme	Results in increased ergosterol synthesis; contributes to cross-resistance between fluconazole and itraconazole

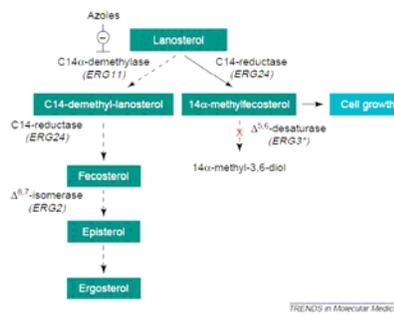
Lupetti et al, Trends Mol Medicine (2002) 8:76-81

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Mécanismes de résistance aux dérivés azolés



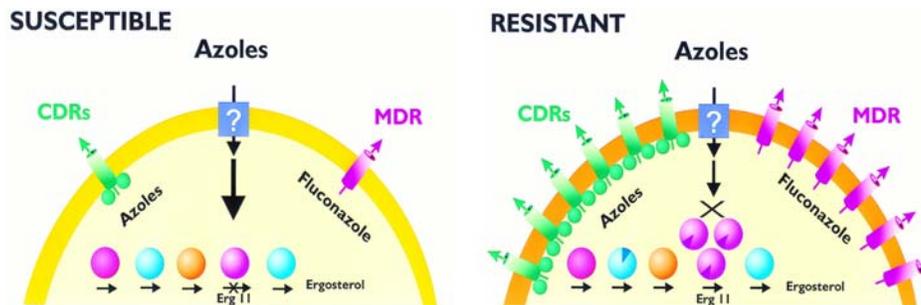
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Mécanismes de résistance aux dérivés azolés



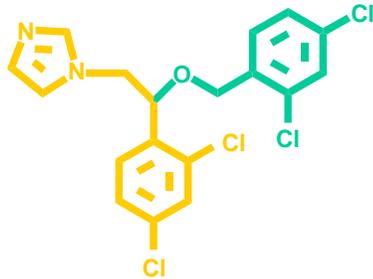
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Miconazole



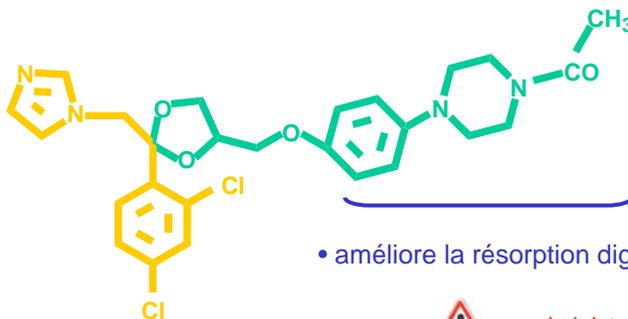
- premier antifongique azolé
- mauvaise résorption digestive
- usage essentiellement topique (candidoses)

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Kétoconazole



- améliore la résorption digestive en milieu acide



co-administration d'anti-acides

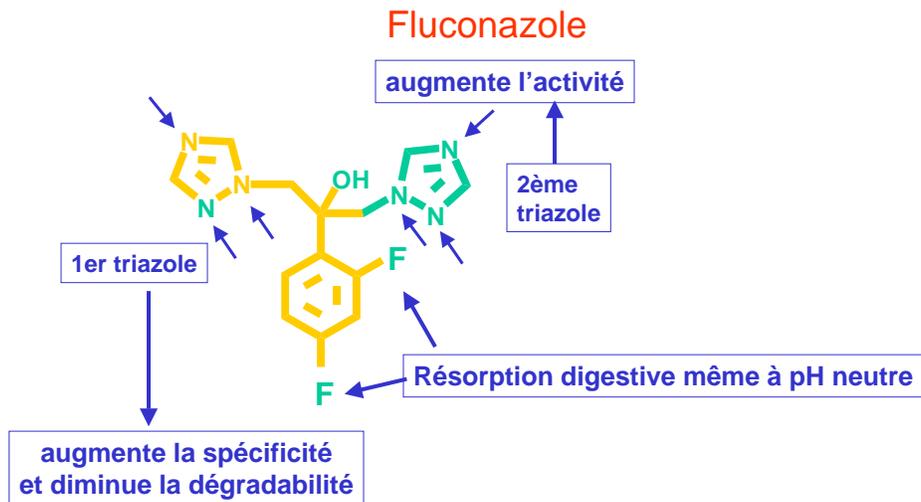
indications (peu utilisé aujourd'hui):

- candidoses (histoplasmoses, coccidioïdomycose)
- prophylaxie des colonisations par levures chez les neutropéniques

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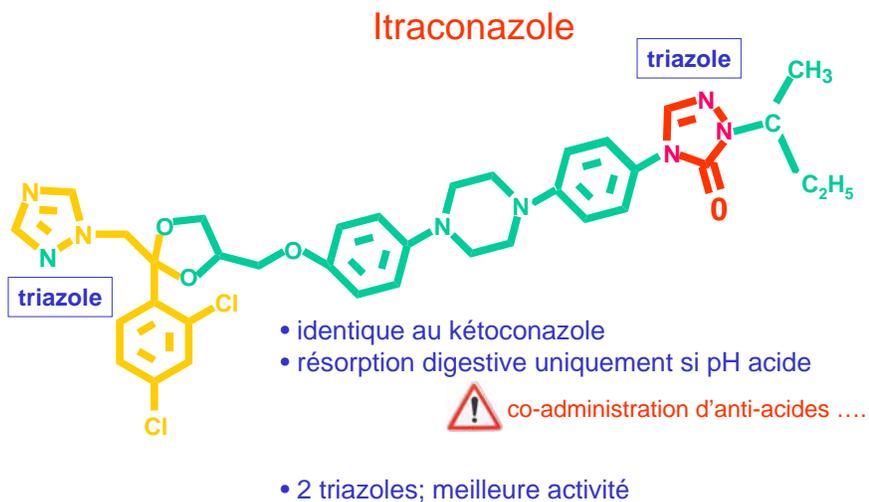
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Indications :

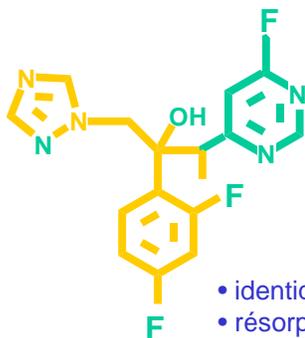
- méningites cryptococciques (pénétration dans LCR)
- candidoses.



Indications :

- candidoses
- aspergilloses

Voriconazole



- spectre large; puissante activité
- pas de résistance croisée avec les autres azolés

- identique au fluconazole
- résorption digestive même à pH neutre

• indications :

- aspergilloses invasives
- infections invasives graves à *Candida* **résistant au fluconazole**.
- Traitement des infections fongiques graves à *Scedosporium* spp. ou *Fusarium* spp.
- en première intention aux patients immunodéprimés, atteints d'infections évolutives, pouvant menacer le pronostic vital.

Fluconazole resistant pseudomembranous candidiasis



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Dérivés azolés et interactions avec CYP (CYP2C19, CYP2C9 et CYP3A4)



	Degree of interaction				Effect	Clinically significant
	FLU	ITC	VOR	POS		
Substrates of CYP3A4 and CYP2C9*	++	+++	+++	++	Increased plasma concentrations of other drug substrates	Yes (some contraindicated)
Inducers of CYP3A4 and CYP2C9†	++	+++	+++	++	Decreased plasma concentrations of triazoles	Yes (some contraindicated)
Warfarin	++	+++	+++	++	Increased prothrombin time	Yes
Phenytoin	+++	+++	+++	+++	Increased phenytoin levels, decreased triazole levels	Yes
Rifampicin	+++	+++	+++	+++	Decreased triazole levels	Yes (contraindicated with ITC, VOR, POS)
Proton-pump inhibitors	++	++	+++	++	Increased proton-pump inhibitor levels, decreased triazole absorption	Yes
Cyclosporine	++	++	+++	++	Toxicity, renal failure	Yes
Tacrolimus	++	++	+++	++	Toxicity, renal failure	Yes
Sirolimus	++	++	+++	++	Toxicity, renal failure	Yes (contraindicated with VOR)
Statins	++	+++	+++	++	Increased statin levels	Yes

FLU = fluconazole; ITC = itraconazole; POS = posaconazole; VOR = voriconazole. ++ = mild, +++ = moderate, ++++ = high, +++++ = very high. * Includes but not restricted to cisapride (contraindicated with FLU, ITC, VOR, POS), terfenadine, astemizole, piroxicam, quinidine, ergot alkaloids (contraindicated with ITC, VOR), sirolimus (contraindicated with VOR), tacrolimus, cyclosporin, statins, warfarin, omeprazole, phenytoin, benzodiazepines, HIV protease inhibitors, non-nucleoside reverse transcriptase inhibitors, and sulfonylurea oral hypoglycaemics. † Includes rifampicin (contraindicated with ITC, VOR, POS), rifabutin (contraindicated with ITC, VOR), long-acting barbiturates (contraindicated with VOR), phenytoin, HIV protease inhibitors, non-nucleoside reverse transcriptase inhibitors.

- **contre-indiqués**: terféfadine, astémizole, cisapride, quinidine, alcaloïdes de l'ergot
- **surveillance étroite** : ciclosporine, tacrolimus, anticoagulants oraux, sulfonylurées
- **adaptation de posologie** : statines, benzodiazépines

Chen et al. Med J Aust. (2007) 187:404-9.

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Dérivés azolés : propriétés pharmacologiques



Pharmacocinétique

- absorption variable; parfois dépendante du pH.
- distribution variable; LCR : fluconazole
 voie orale: itraconazole, fluconazole, voriconazole
- demi vie longue (> 24 heures) sauf kétoconazole (8 h) et voriconazole (6 h)

Effets secondaires

- toxicité hépatique (surtout kétoconazole)
- troubles digestifs (miconazole)
- intolérance cutanée par voie locale

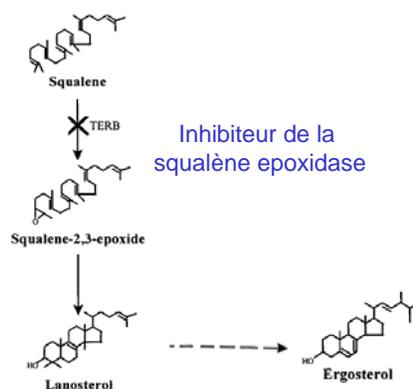
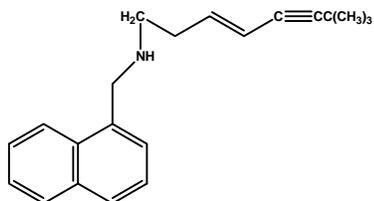
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Allylamines

terbinafine



- actif uniquement sur les dermatophytes
- bien résorbée mais effet de premier passage important
- interaction avec inhibiteurs et inducteurs des cytochromes

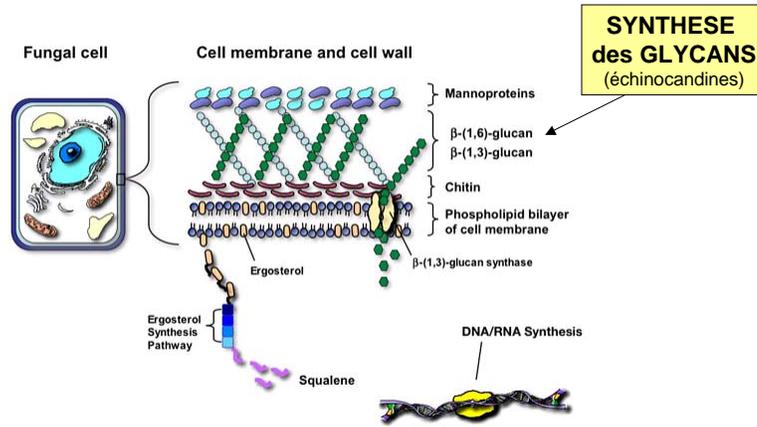


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Cibles des médicaments antifongiques



Echinocandines : caspofungine

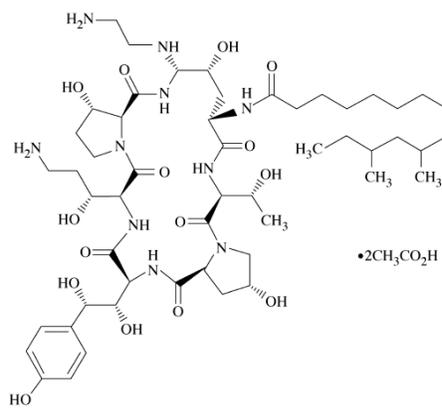


Figure 1. Structure of caspofungin.

Echinocandines

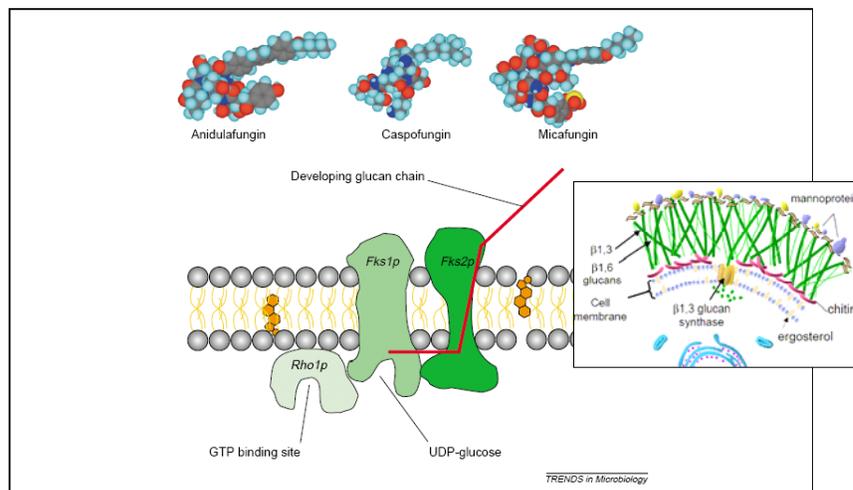


Fig. 5. Cartoon depicting the transmembrane complex of two proteins, Fks1p and Fks2p, involved in synthesis of β -1,3 glucan in the cell walls of *Saccharomyces cerevisiae*. Activity of the complex is regulated by the GTP-binding peptide, Rho1p. Fks1p is the target of the echinocandins, with the structures of the three agents now entering clinical use illustrated above. Evidence for *Candida albicans* suggests that the Fks2p homologue in this species might not be expressed in growing cells. Much remains unknown about the glucan synthase protein complex and the manner of interaction of echinocandins with the complex.

Odds. Trends in Microbiology (2003) 11:272-279

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Caspofungine: propriétés pharmacologiques

Indications:

- candidoses (oropharyngées, oesophagiennes)
- aspergilloses invasives chez les patients qui ne répondent pas ou sont intolérants aux autres traitements

Pharmacocinétique:

- Mauvaise résorption orale
- Accumulation dans l'organisme lors d'un traitement prolongé (25-50 % d'augmentation de l'AUC après 2 à 3 semaines de traitement).
- Métabolisation par le foie; demi-vie, environ 10 h.

Effets secondaires

- phlébite au site d'injection
- fièvre, nausées et vomissements.

Interactions médicamenteuses

- \nearrow AUC de la caspofungine par cyclosporine
- \searrow AUC par inducteurs des cytochromes P450 (rifampicine, inhib. protéases du HIV, phénytoïne, carbamazépine, dexaméthasone).

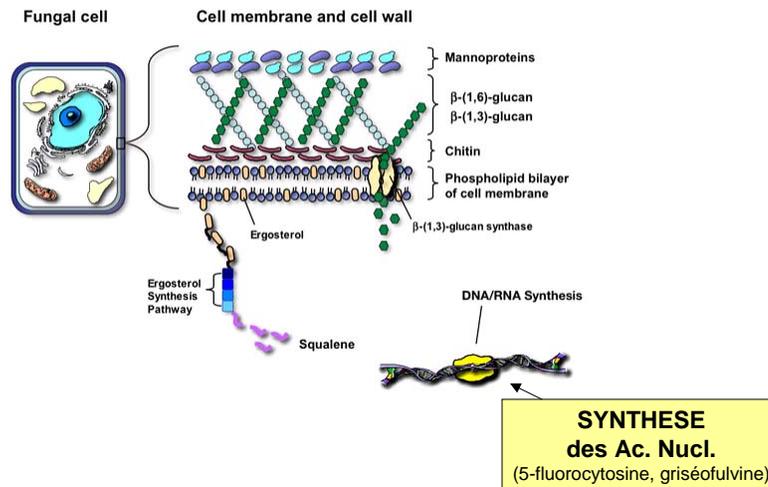
Patients à risque d'infection fongique!

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Cibles des médicaments antifongiques



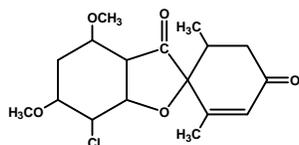
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Inhibiteurs de synthèse des acides nucléiques

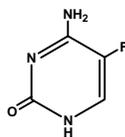
griséofulvine



- infections à dermatophytes (cuir chevelu, peau, ongles)
- adm. orale, résorption \nearrow par aliments riches en graisse
- réactions allergiques, maux de tête, troubles digestifs



5-fluorocytosine



- inhibition de la thymidilate synthase; résistance fréquente
→ utiliser en association avec l'ampho B pour les candidoses et les cryptococcoses
- concentration dans l'urine → infection urinaire
- leuco- et thrombopénie



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