

Terapia precoce anti-COVID sul territorio

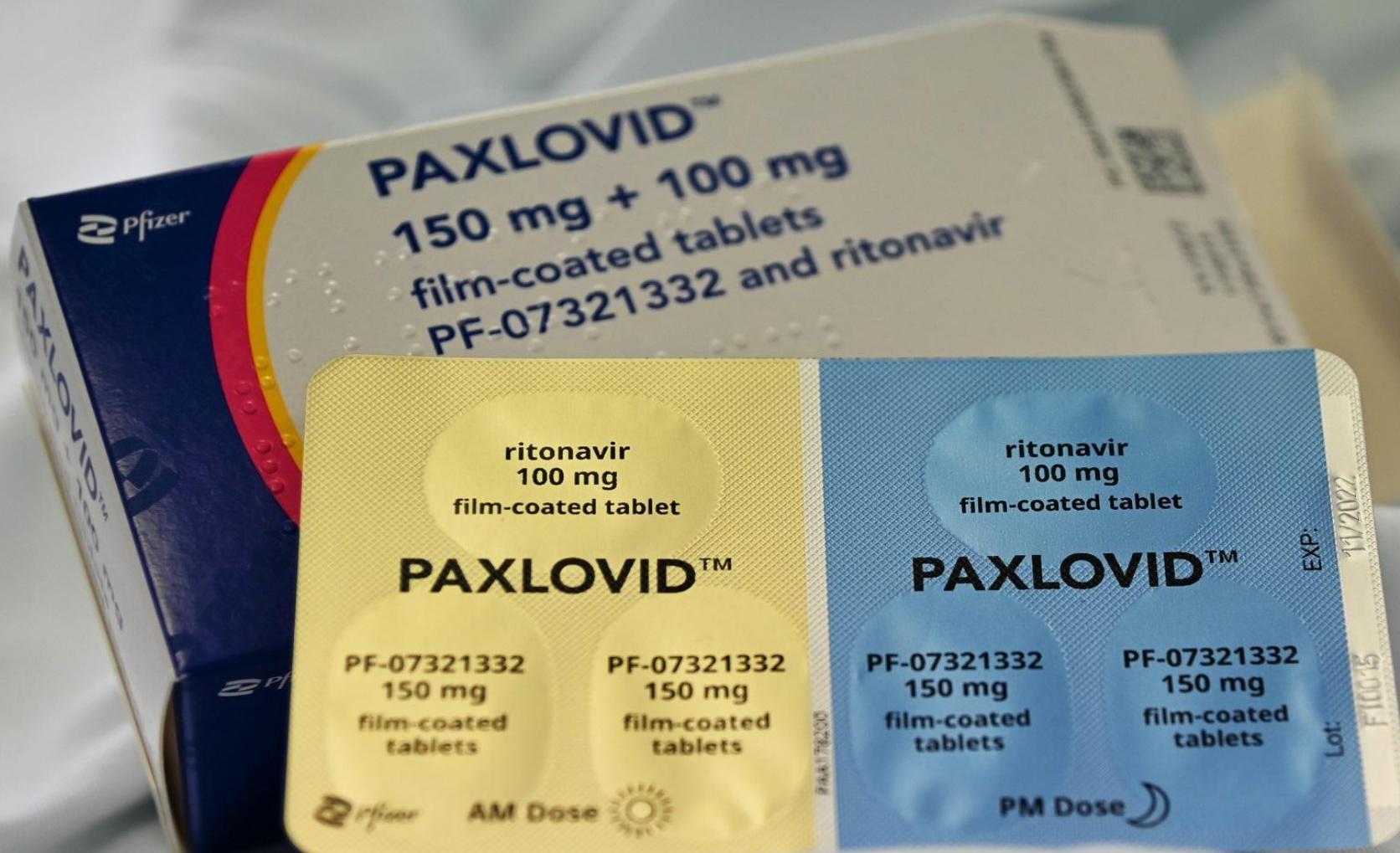
Dr. Simone Vasilij Benatti

Malattie Infettive

ASST- Papa Giovanni XXIII – Bergamo

sbenatti@asst-pg23.it

teleinfe@asst-pg23.it





Nirmatrelvir/ritonavir

Nirmatrelvir è un INIBITORE della PROTEASI di SARS-CoV-2

Ritonavir (RTV) è solo booster

POSOLOGIA:

2 + 1 capsule (intero)

[1 + 1 se EGFR tra 30 e 60]

Con o senza cibo

Ogni 12 h

Per 5 gg

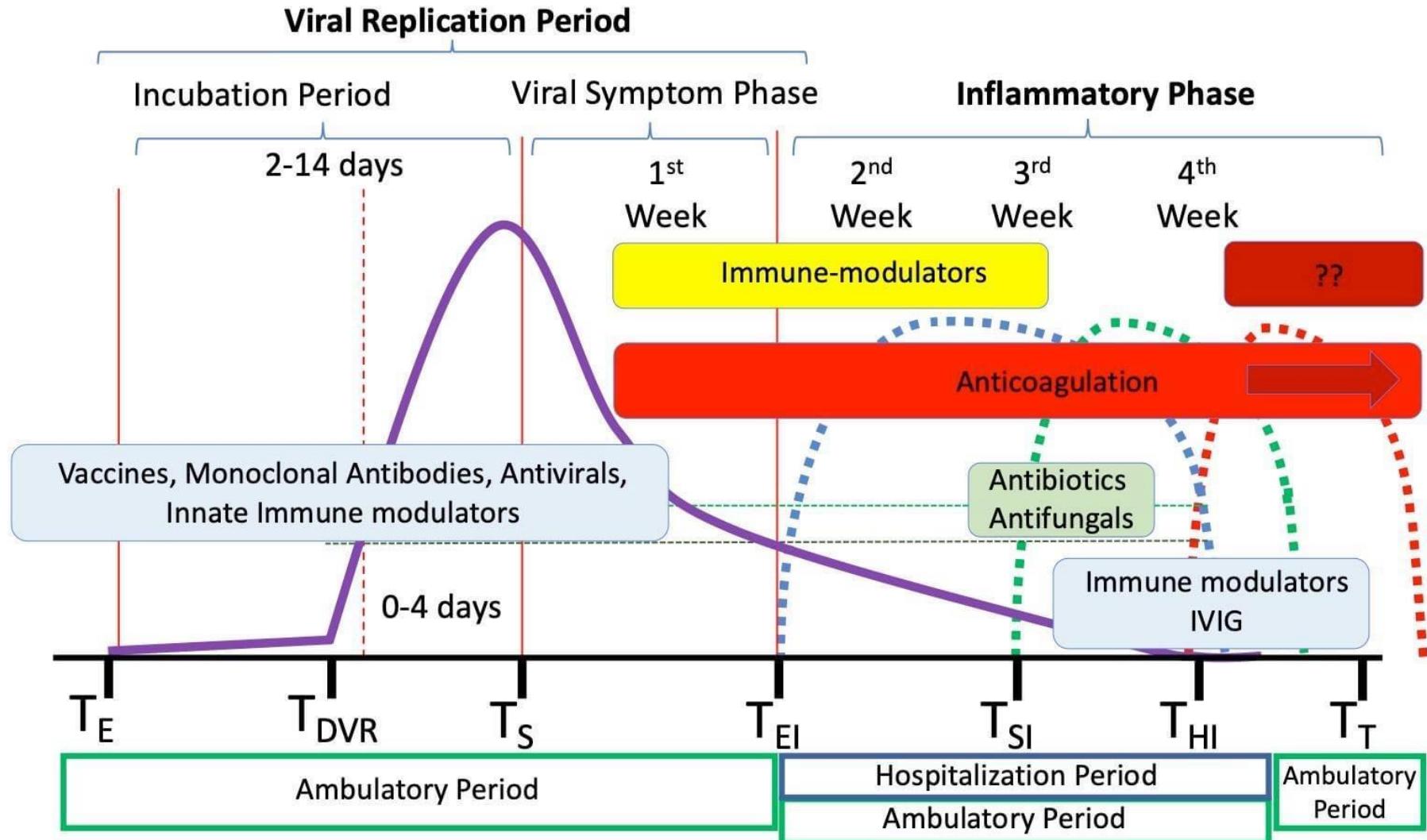
Poi STOP!

Entro quanto tempo?

- Max 5 gg dall'esordio dei sintomi
- Prima si fa, meglio è
- Solo se appartiene a categoria a rischio
- Il test molecolare non è richiesto

L'importante è che il paziente:

- ✓ Non sia da ricoverare (non deve avere necessità di O₂ – se già in LTOT, non deve essere aumentato...)
- ✓ Non sia una donna in gravidanza (attenzione con contraccettivi orali, evitare gravidanza fino a 7 gg dopo la fine della cura)
- ✓ Abbia EGFR > 30 mL/min



The major phases and suggested time points that are critical in COVID-19. The Pre-exposure period, the Incubation period starting at T_E (Time of exposure), the Detectable Viral Replication Phase T_{VR} (Time of viral replication), the Viral Symptom Phase T_S (Time of symptom onset), the Early Inflammatory Phase T_{EI} (Time of early inflammation), the Secondary Infection Phase starts at T_{SI} (Time of secondary infection) the Hyperinflammatory Phase which starts at T_{HI} (Time of Hyperinflammation onset) and the Tail Phase that starts at T_T (Time of the tail onset)

A chi si può dare?

Tabella-1 - CRITERI ELEGGIBILITA' AIFA (gen/2022)

		Sotrovimab	Remdesivir precoce	Molnupiravir	Nirmatrelvir
Timing	esordio sintomi	≤ 7 gg	≤ 7 gg	≤ 5 gg	≤ 5 gg
Criteri di esclus.	età minima	≥ 12	≥ 18	≥ 18	≥ 12
	peso minimo	> 40 Kg	> 40 Kg	> 40 Kg	> 40 Kg
	gravidanza	No	No	No [®]	No ^{&}
	allattamento	No?	No (stop temporaneamente)		
	EGFR < 30	Sì	No	No	No (se EGFR 30-60: ridurre dose)
	ALT > N	Sì	Solo se < 5 ULN	No	Non se Child C
Efficacia in soggetti non vaccinati		85% ^a	87% ^b	31% ^c	88% ^d
Criteri di inclus.	età > 65	Sì	No		
	BMI >= 30	Sì	Sì		
	Diabete	Non controllato o con compl. croniche	Non controllato		
	Patol. Cardiovasc.	Sì	Solo se grave (SCC, CAD, CMD)		
	Patol. Cerebrovasc.	Sì	No		
	Immunodeficienza	Sì	Sì		
	Patol. Polmonare	Sì	Solo se grave		
	Epatopatia	Sì	No		
	Hb-patie	Sì	No		
	Patol. del neurosviluppo o neurodegener.	Sì	No		
	Neoplasia oncologica/ematol. in fase attiva	Sì	Sì		
	IRC	Sì	Sì (ma EGFR < 30 è criterio di esclusione)		

Effetti collaterali

- Di tipo digestivo
- Diarrea

(Solitamente non rilevanti)

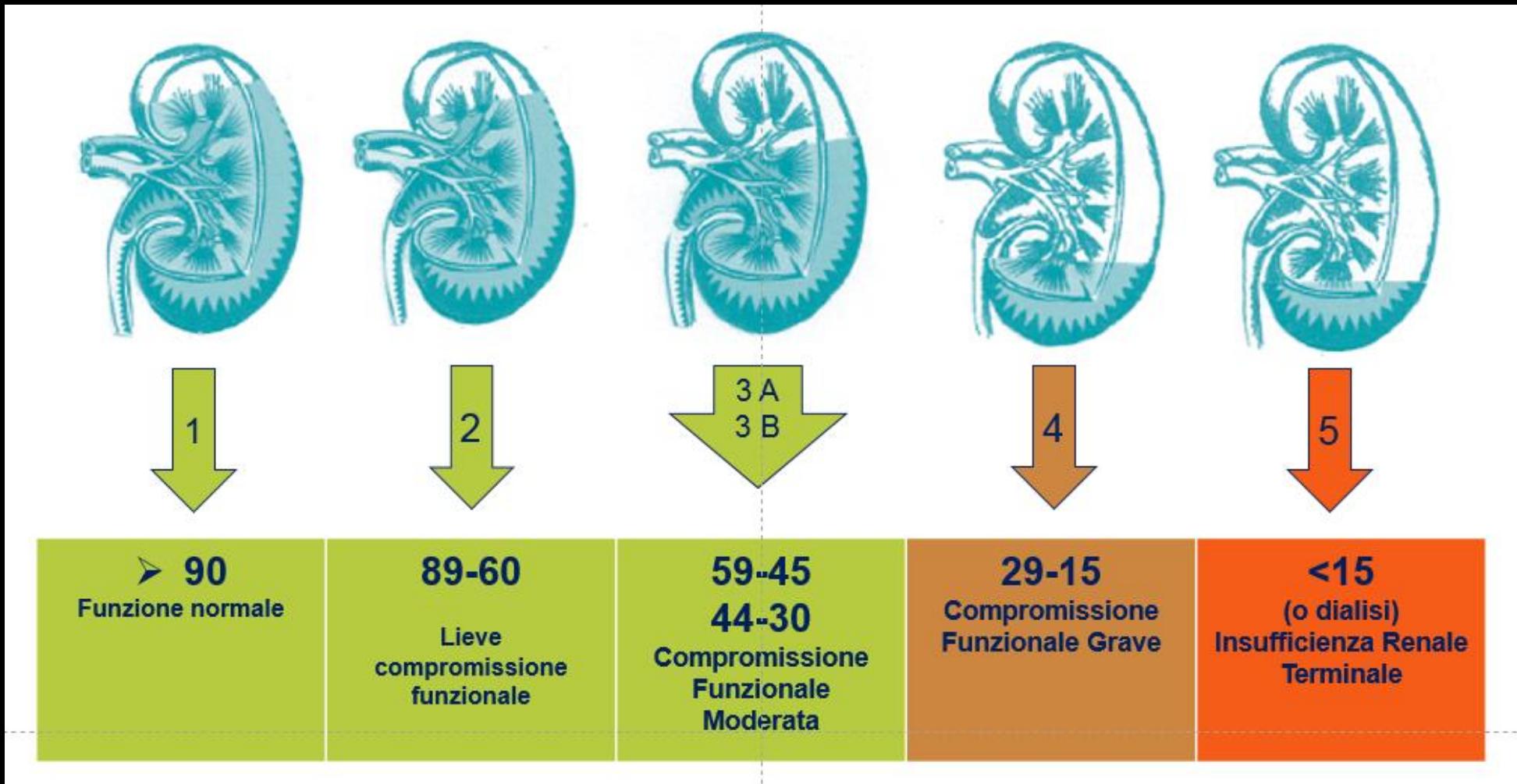
Formula di Cockroft-Gault per EGFR

$$eCrCl = \frac{(140 - \text{Age}) \times \text{Weight (kg)}}{72 \times \text{Creatinine}_{\text{serum}} (\text{mg/dL})} \times 0.85 \text{ if female}$$

Oppure....

Scaricare una App, tipo MedCalc

Stadi insufficienza renale



Interazioni farmacologiche

The screenshot shows the homepage of the COVID-19 Drug Interactions website. At the top left is a white COVID-19 icon. To its right is the text "COVID-19 Drug Interactions". On the right side of the header is the University of Liverpool logo with the text "UNIVERSITY OF LIVERPOOL". Below the header are four navigation links: "About Us", "Interaction Checkers", "Prescribing Resources", and "Contact Us". A green banner across the middle of the page contains the text "New COVID therapy - interactions with molnupiravir now available". The main content area features a large button labeled "Interaction Checker" with the subtext "Access our free, comprehensive and user-friendly drug interaction charts". Below this are two smaller boxes: one for "Prescribing Resources" (printable interaction tables, interaction summary charts and clinical prescribing resources) and one for "Twitter" (@covidDDIs), which encourages users to follow for interaction news and website updates.

COVID-19 Drug Interactions

UNIVERSITY OF LIVERPOOL

About Us Interaction Checkers Prescribing Resources

Contact Us

New COVID therapy - interactions with molnupiravir now available

Interaction Checker

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Prescribing Resources

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Twitter

@covidDDIs

Follow us on Twitter for interaction news and for the latest additions and changes to the website

Interazioni farmacologiche

If a drug is not listed below it cannot automatically be assumed it is safe to coadminister.

COVID Drugs	Co-medications	Drug Interactions
nirmatre X	amlodip X	<input checked="" type="checkbox"/> Check COVID/COVID drug interactions
<input checked="" type="radio"/> A-Z <input type="radio"/> Class <input type="radio"/> Trade	<input checked="" type="radio"/> A-Z <input type="radio"/> Class	Reset Checker
<input checked="" type="checkbox"/> Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.] (i)	<input checked="" type="checkbox"/> Amlodipine (i)	Switch to table view Results Key
<input checked="" type="checkbox"/> Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.] (i)	<input checked="" type="checkbox"/> Amlodipine (i)	<div style="background-color: #f0e68c; padding: 5px;"><u>Potential Interaction</u></div> <p>Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.]</p> <p>Amlodipine</p>

Interazioni farmacologiche

Potential Interaction 

Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.]

Amlodipine

Quality of Evidence: Very Low 

Summary:

Coadministration has not been studied. Amlodipine is metabolized by CYP3A4. Nirmatrelvir/ritonavir is predicted to increase amlodipine exposure by ~2-fold based on drug-drug interactions studies with amlodipine and indinavir/ritonavir or paritaprevir/ritonavir leading to the recommendation to reduce amlodipine dosage by 50%. However, a dose adjustment can be optional in the case of amlodipine given that patients can be advised to monitor for symptoms of hypotension and to temporarily pause the antihypertensive drug if needed. The inhibitory effect of ritonavir is expected to last up to 3 days after the last administered dose of nirmatrelvir/ritonavir.

Description:

Ritonavir dosed as a pharmacokinetic enhancer or as an antiretroviral agent inhibits CYP3A4 and as a result is expected to increase the plasma concentrations of calcium channel antagonists. Careful monitoring of therapeutic and adverse effects is recommended when this medicine is concomitantly administered with ritonavir.

Paxlovid Summary of Product Characteristics, Pfizer Ltd, February 2022.

Co-administration may increase concentrations of the calcium channel

Attenzione con:

- Anticoagulanti orali (warfarin, DOAC...) → Sospendere? Passare a Clexane?
- Ca++ antagonisti → Sospendere/ridurre?
- Alfa-litici → Sospendere?
- Statine → Sospendere!
- Salmeterolo → Salbutamolo?
- Antirigetto → Valutare alternative

A-Z	Class	Trade	A-Z	Class	Trade	Switch to table view	Results Key
<input checked="" type="checkbox"/> Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.] (i)			<input checked="" type="checkbox"/> Salbutamol (Albuterol) (i)			Switch to table view	Results Key
<input checked="" type="checkbox"/> Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.] (i)			<input checked="" type="checkbox"/> Salmeterol (i)			<div style="background-color: red; color: white; padding: 5px; text-align: center;"><u>Do Not Coadminister</u></div> <div style="background-color: white; border: 1px solid #ccc; padding: 10px; margin-top: 10px;"><p>Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.]</p><p>Salmeterol</p><p>More Info</p></div> <div style="background-color: green; color: white; padding: 5px; text-align: center;"><u>No Interaction Expected</u></div> <div style="background-color: white; border: 1px solid #ccc; padding: 10px; margin-top: 10px;"><p>Nirmatrelvir/ritonavir (5 days) [Please read the interaction details as management of these interactions may be complex.]</p><p>Salbutamol (Albuterol)</p></div>	

Per ogni dubbio o per casi particolari...

- Tel. 339-8756575
- LUN > VEN – h 9.00 > 16.00

Oppure:

- teleinfe@asst-pg23.it

Grazie.