

## Molecular Partners' Covid-19 therapy proves inactiv



[Elizabeth Cairns](#)

Ensovibep has become the latest Covid-19 treatment candidate to crash out of the [NIH-sponsored Activ-3 basket trial](#), wiping 34% off Molecular Partners' share price. A planned futility analysis of the cohort evaluating ensovibep, a darpin Molecular Partners licensed to Novartis, concluded that enrolment should be stopped. Activ-3 is looking at potential therapies for hospitalised patients, but no agent in the trial has yet succeeded. The study's [clinicaltrials.gov](#) entry was also edited this month to state that Astrazeneca's MAb combo AZD7442 was no longer being administered either, but no explanation has been offered for this. As a result, Pfizer's IV antiviral PF-07304814 (lufotrelvir) is the only project still standing in Activ-3, at least until any new ones are added. There is some comfort for Molecular Partners in that Activ-3 backed ensovibep's safety profile, which was consistent with the standard of care, Gilead's Veklury, the company said. Ensovibep is in a separate phase 2/3 trial, [Empathy](#), in Covid-19 outpatients, with interim data on the first 400 expected in early 2022. But in this niche [Pfizer's Paxlovid, an oral drug related to lufotrelvir, has set a standard](#) that will be very hard to beat – as Molecular Partner's shareholders clearly understand.

### Activ-3: NIH-funded phase 3 trial in hospitalised patients

Company	Project	Description	Outcome
Lilly/Abcellera	Bamlanivimab	Infused MAb	326 pt arm closed for futility <a href="#">Oct 2020</a>
Vir Biotechnology	Sotrovimab	Intramuscular or infused MAb	344 pt arm closed for futility <a href="#">Mar 2021</a>
Brii Biosciences	BR11-196 and BR11-198	Infused MAb combination	343 pt arm closed for futility <a href="#">Mar 2021</a>
Molecular Partners	Ensovibep	Infused antiviral	470 pt arm closed for futility <a href="#">Nov 2021</a>
Astrazeneca	AZD7442 (tixagevimab + cilgavimab)	Infused MAb combination	Arm closed Nov 2021, reason unclear
Pfizer	Lufotrelvir	Infused antiviral	Arm ongoing

*All patients received a backbone of Gilead's Veklury as standard of care. Source: [clinicaltrials.gov](#).*

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