

Potential Covid-19 drug to start final trial

Developed by S'pore-based firm, the drug will be tested on 500 Covid patients

✔ LIM MINZHANG

Trials for a potential treatment for Covid-19 are being accelerated, with the third and final phase for the drug due to start in a few weeks' time.

Developed by Singapore-based biotechnology company Tychan, the monoclonal antibody – or immune system protein – known as TY027 will be tested on 500 recently diagnosed Covid-19 patients.

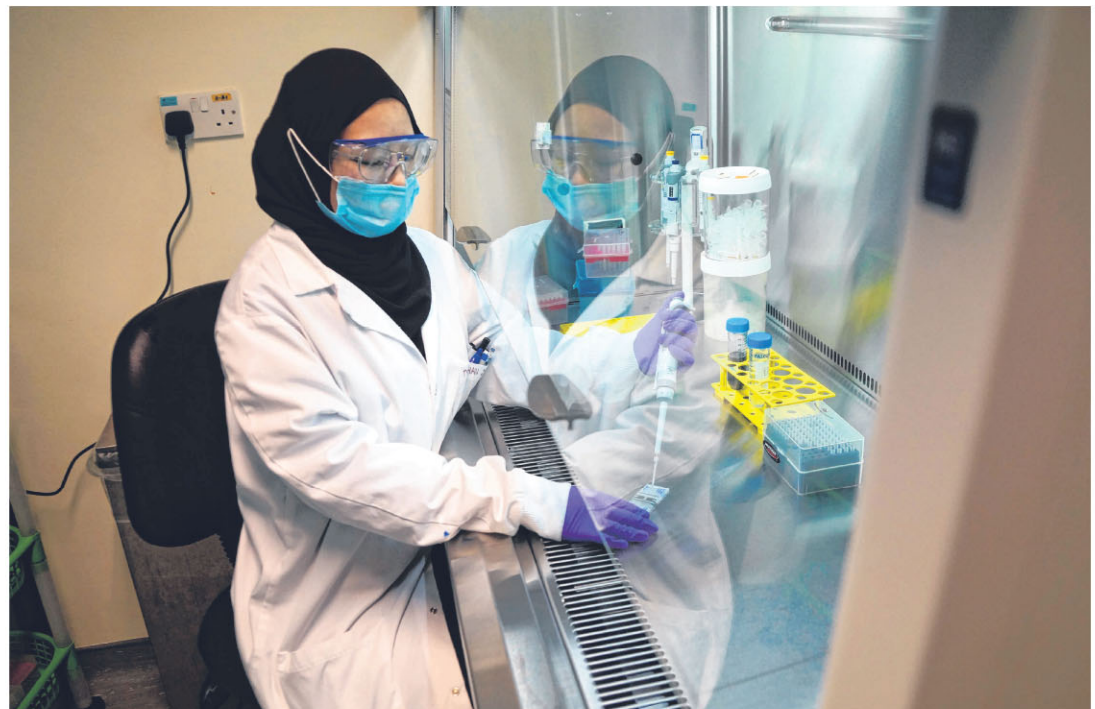
The trial is expected to take a few months, but is the final step before the drug can be approved for treatment here.

Professor Ooi Eng Eong from Duke-NUS Medical School, who is one of the founders of Tychan, told *The Straits Times* yesterday that as a small company, Tychan does not have the resources to run trials with thousands of patients. But if the results are dramatic, there is a chance that fewer than 500 patients would suffice.

“We have to be tighter in how we recruit and the kind of patients that we enrol. The key criterion is for the patients to be in the first seven days of illness,” he said.

“Limiting the kind of patients we can enrol would give ourselves the best chance of showing, in as short a time as possible and in as few patients as possible, that this drug works to prevent severe Covid-19.”

Monoclonal antibodies are immune system proteins that are created in the laboratory and can be specially designed and engineered to target Sars-CoV-2.



Developed by Singapore-based biotechnology company Tychan, TY027 will be tested on 500 recently diagnosed Covid-19 patients. PHOTO: TYCHAN

The advantage is that these antibodies can be developed over several months and produced in large batches. A single injection may last for a few weeks.

Half of the 500 patients will receive a placebo in the double-blind trial so that the drug's effectiveness can be tested, said Prof Ooi. Tychan will work with hospitals under SingHealth to recruit the patients.

PENDING APPROVAL

The recruitment process has not started yet because the final trial is pending approval from the authorities.

In the best-case scenario, TY027 could be approved as a drug treatment for Covid-19 by early next year, or even earlier, said Prof Ooi, who is deputy director of the Emerging Infectious Diseases Programme at Duke-NUS.

Clinical trials are commonly

conducted in four phases, beginning with a small group to test the drug's safety and side effects, then moving on to larger groups to determine its efficacy.

Usually, in phase three, what is being tested is whether the drug works as intended in preventing disease or accelerating recovery.

TY027 is being explored for the treatment of Covid-19 patients in slowing the progression of the disease and speeding up recovery, as well as potentially providing temporary protection against infection.

A six-week phase one trial in June with 23 healthy volunteers yielded good outcomes in terms of safety, said Prof Ooi.

He added: “The safety profile has been excellent, there's minimal side effects... And the results have been very, very encouraging.”

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