The Volunteer Journey

www.volunteer.gov.bh

Registration requirements and procedures

- Volunteer must be aged 18 years old and above, and in good health
- Volunteer must be a Bahraini resident/or has a residency permit valid during the trial period
- The volunteer will undergo an assessment to evaluate their eligibility to participate in the trials
- Volunteer must register using official documents - ID/passport
- Volunteer must sign the informed consent form before starting the trial

Visit the designated hall at The Bahrain International Exhibition & Convention Centre from 8AM to 6PM to register and start the vaccination process immediately.

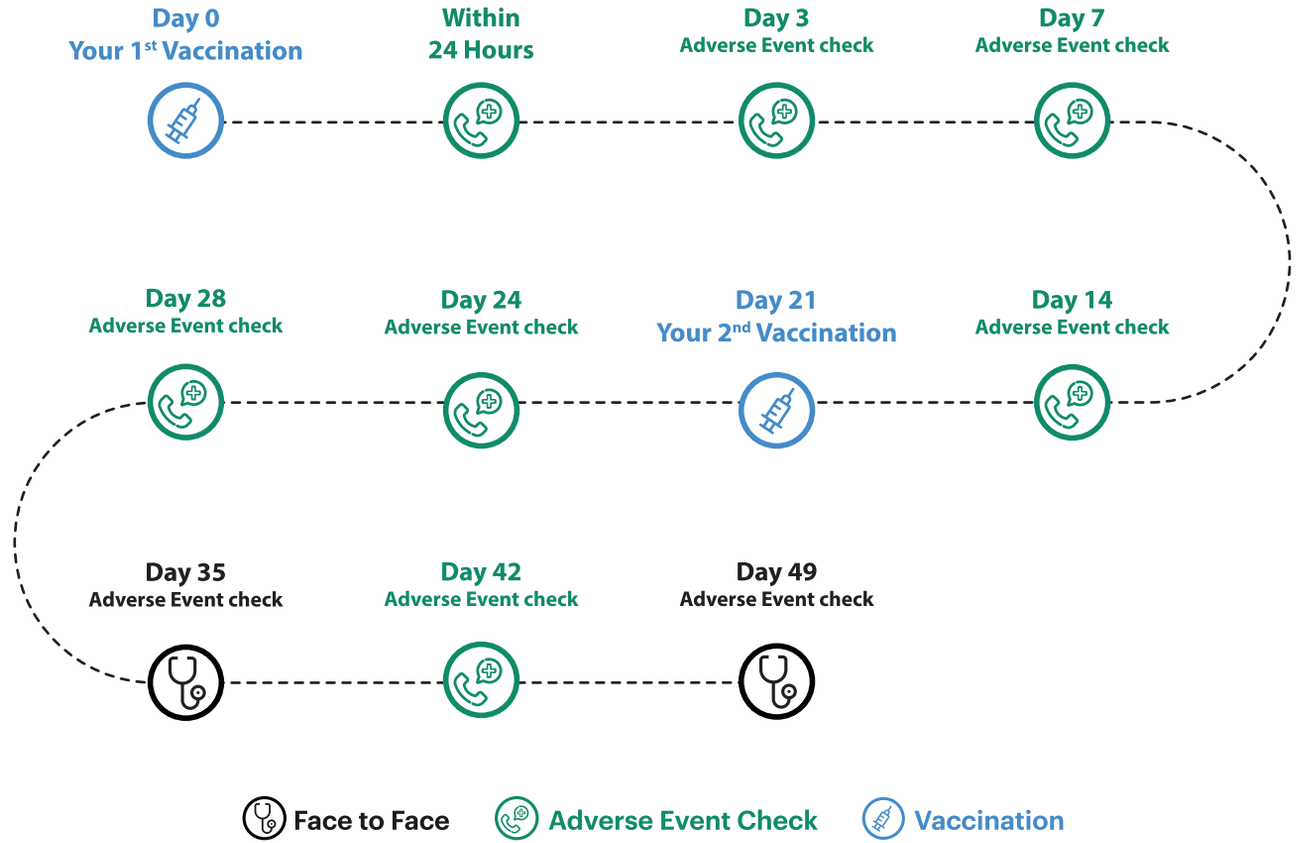
Discover more on www.volunteer.gov.bh

You cannot participate in the trial if:

- Had fever, dry cough, fatigue, nasal obstruction or runny nose within 14 days before vaccination
- Have known allergic reactions to medicines
- Have a personal or family history of convulsion or epilepsy
- Have uncontrollable diabetes, any acute or chronic disease conditions like asthma, severe cardiovascular diseases, liver and kidney diseases, and malignant tumors
- Currently receiving any immunotherapy
- Received blood within 3 months before the trial
- You are pregnant or planning to get pregnant during the study period

Locations for vaccine trials

The allocated hall at the Bahrain International Exhibition & Convention Centre.



After the 49th day you will be regularly contacted by your doctor. A telephone check-up will be conducted once or twice a month, and a physical check-up will be conducted every three months.

#لأجل_الإنسانية
#4Humanity

A clinical cooperation agreement was signed between The Ministry of Health at the Kingdom of Bahrain and the Chinese pharmaceutical Sinopharm (CNBG), to lead the Phase III clinical trial of COVID-19 inactivated vaccine in the Kingdom of Bahrain, engaging multi-national volunteers.

FREQUENTLY ASKED QUESTIONS

Are there related risks?
Some common side effects may occur when receiving this vaccine, including mild pain, redness, induration, and pruritus at the vaccination site.

The volunteer may present mild systemic reactions such as fever, headache, fatigue, nausea, vomiting, diarrhea, cough, allergy, muscle pain, arthralgia, lethargy, etc.

Usually these symptoms are mild and subside on their own. Patients with moderate to severe symptoms will be treated under the supervision of their treating physician and site principal investigator.

Who do I call if I experience any symptoms?
By calling Tel. No. **38817484**.

How long will I be required to participate in the study?
The duration of physical participation is approximately 2 months and the follow-up period might last up to 12 months.

Will my data privacy be respected?
The data of participants will be kept confidential at the Ministry of Health and identification and personal details will not be shared with any entities.

Can I withdraw in the middle of a research study?
The participant is free to opt out at any time and will not bear any responsibility arising from the withdrawal.

This will not affect the participant personally in any shape or form and will not affect any existing medical treatment that he/she is undergoing.

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