

Cover Page

Release Date: March 11, 2022

ClinicalTrials.gov ID: NCT02403505

Unique Protocol ID: IND 28299

Brief Title: Early Phase Clinical Trial About Therapeutic Biological Product Mix for Treating COVID-19 (AD26-BCG)

Official Title: Conducting an Early Phase Clinical Trial to Assess for COVID-19 Antigen Presentation Therapeutic Biological Product Mix Activity That Suggests the Potential for Clinical Benefits of COVID-19 Patients.

Secondary IDs: FWA00015357 [Registry ID: Federal-wide Assurance (FWA) DHHS OHRP]
IORG0007849 [Registry ID: Institution or Organization (IORG) DHHS OHRP]
IRB00009424 [Registry ID: Institutional Review Board (IRB) DHHS OHRP]
IND28299 [Registry ID: Investigational New Drug Application (IND) FDA]
NDA217392 [Registry ID: New Drug Application (NDA) FDA]
BLA761321 [Registry ID: Biologics License Applications (BLA) FDA]

Informed Consent Form

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If you agree to join this study, you will need to sign and date or make your mark below. Before you sign and date or make your mark on this consent form, make sure of the following:15

Informed Consent Form

Study Title: Treat COVID-19 virus infection with Ad26 COVID-19 Spike plus TICE® BCG mix for intradermal injection

ClinicalTrials.gov ID: NCT02403505

Sponsored by: Han Xu, M.D., Ph.D., FAPCR, Sponsor-Investigator

Principal Investigator: Han Xu, M.D., Ph.D., FAPCR, Sponsor-Investigator

Telephone: +1 301-222-7143

Thank you for your interest in our research study. Please read this consent form or ask someone to read it to you. If you decide to join the study, we will ask you to sign and date or make your mark on this form. We will offer you a copy to keep. We will ask you some questions to see if we have explained everything clearly. You can also ask us questions about the study.

The clinical research is the same as treatment. The Ad26 COVID-19 Spike plus TICE® BCG mix for intradermal injection can treat infection of multiple gene mutation COVID-19 virus strains via activate human COVID-19 antigen presentation reaction, so the human antigen presenting cells (APCs) take up and process COVID-19 target antigen into small peptide fragments, and then kill COVID-19 virus directly. The purpose of the research study, i.e., clinical trial NCT02403505 is to answer medical questions.

Key information

- Being in this research study is voluntary. It is your choice.
- BCG should not be given to individuals previously infected with M. tuberculosis. A person given the tuberculin skin test [10 tuberculin units (10 TU) purified protein derivative (PPD)] must return within 48 hours. The positive is skin test reading > 5 mm induration at 48 hours.
- The purpose of this study is to learn more about infection with and recovery from the virus called SARS-CoV-2. Some people know this virus by the name “coronavirus.” It can cause the disease called COVID-19. We will enroll 20 participants who have no symptoms, mild symptoms, and people who need not to be hospitalized without more severe symptoms.
- Your participation in this study will last for about 1 month. Procedures will include intradermal injection and COVID-19 testing by standard RT-PCR assay.
- We will also contact you about 1 month after your last study visit to check on your health.

- Because you have a positive COVID-19 test result you may have been told to be in quarantine or staying at home away from others. Some clinic staff may come to your location to meet with you and have an intradermal injection for this study as needed.
- The risks associated with this study are minimal and may be primarily due to collecting the blood, nasal, saliva, and optional stool samples and the potential for loss of confidentiality.
- There is no direct benefit to you from your participation in the study.

1. About the study

The COVID-19 Telehealth / Telemedicine clinical trial site are doing a study to learn more about SARS-CoV-2 infection and the disease called COVID-19 that it can treat. The Telehealth / Telemedicine clinical trial site includes Medicine Invention Design Incorporation (Health Care Provider - Group or Organization) and online patient referral center.

About 20 participants will take part in this phase 1 clinical study at single study site or multiple study sites across the United States. The sponsor-investigator in charge of this clinical study at this research center is Han Xu, M.D., Ph.D., FAPCR. The Medicine Invention Design Incorporation will be paying for the phase 1 clinical study.

2. Why are we doing this study?

The main purpose of this study is to learn more about SARS CoV-2 infection and how our bodies develop immune responses to it, hopefully helping us to recover from it. Your immune system is what helps to protect you against infections and disease. This information can be used in the future to help develop better treatments for SARS-CoV-2 infection and COVID-19, as well as may help in developing future therapeutic biological products and other prevention strategies.

3. Joining the study

It is completely following up to you whether or not to join the study.

Eligibility

Minimum Age: 24 Years

Maximum Age: 64 Years

Sex: All

Gender Based: No

Accepts Healthy Volunteers: No

Criteria:

- Conducting an initial, small, controlled clinical trial to assess for COVID-19 Antigen Presentation Therapeutic Biologics Mix that suggests the potential for clinical benefit of COVID-19 patients.
- 20 Lighter Than Mild COVID-19 Patients

Inclusion Criteria:

- Lighter Than Mild COVID-19 Patients
- Positive testing COVID-19 by standard RT-PCR assay
- COVID-19 infection without symptoms
- Symptoms of mild illness with COVID-19 that could include fever, cough, sore throat, malaise, headache, muscle pain, nausea, vomiting, diarrhea, and loss of taste or smell, without shortness of breath or dyspnea
- No clinical signs indicative of Moderate, Severe, or Critical Severity
- PPD negative participant is 10 TU skin test reading < 5 mm induration at 48 hours.

Exclusion Criteria:

- PPD positive participant is 10 TU skin test reading > 5 mm induration at 48 hours
- Symptoms of moderate illness with COVID-19, which could include any symptom of mild illness or shortness of breath with exertion
- Clinical signs suggestive of moderate illness with COVID-19
- Symptoms suggestive of severe systemic illness with COVID-19, which could include any symptom of moderate illness or shortness of breath at rest, or respiratory distress
- Clinical signs indicative of severe systemic illness with COVID-19
- Evidence of critical illness
- Respiratory failure
- Shock
- Multi-organ dysfunction / failure
- Allergy

Take your time in deciding. If it helps, talk to people you trust, such as your regular doctor, friends, or family. If you decide not to join this study, or if you leave it after you have joined, your other care and the benefits or rights you would normally have will not be affected.

4. Being in the study

Study Design:

Study Type: Interventional

Primary Purpose: Treatment

Study Phase: Early Phase 1

Interventional Study Model: Single Group Assignment

- Single Usage
- Single Dosage

Number of Arms: 1

Masking: None (Open Label)

Allocation: N/A

Enrollment: 20 [Anticipated]

Study Description

Brief Summary: Conducting an initial, small, controlled clinical trial to assess for COVID-19 Antigen Presentation Therapeutic Biologics Mix activity that suggests the potential for clinical benefit of COVID-19 patients.

1. Treat Infection of Multiple Gene Mutation COVID-19 Virus Strains.
2. Activate human COVID-19 Antigen Presentation Reaction.
3. The human antigen presenting cells (APCs) can treat the COVID-19 virus antigens into small peptide fragments, and then kill COVID-19 virus directly.

Detailed Description:

- Conducting an initial, small, controlled clinical trial to assess for COVID-19 Antigen Presentation Therapeutic Biologics Mix activity that suggests the potential for clinical benefit of COVID-19 patients
- 20 Lower Than Mild COVID-19 Patients
- Positive testing COVID-19 by standard RT-PCR assay
- COVID-19 infection without symptoms
- Symptoms of mild illness with COVID-19 that could include fever, cough, sore throat, malaise, headache, muscle pain, nausea, vomiting, diarrhea, and loss of taste or smell, without shortness of breath or dyspnea
- No clinical signs indicative of Moderate, Severe, or Critical Severity
- PPD negative participant is 10 TU skin test reading < 5 mm induration at 48 hours
- Our trial duration will be 4-week duration.
- Ad26 COVID-19 Spike Organism 1.0 mL plus TICE® BCG Organism 50 MG Mix
- Intradermic Injection, ID
- Our trial duration will be 4-week duration
- Negative testing COVID-19 by standard RT-PCR assay after injection 2 weeks

You will be in the single group, i.e., only one group.

We will enroll 20 people who have no symptoms, people with mild symptoms, and people who must not be hospitalized because they have no more severe symptoms. It is possible that people who start the clinical study in the single group may be stopped the study if their symptoms change.

You may have about 5 study visits at 1 month, and then we will contact you online at next 2 months to check on your health.

You can enroll online at Telehealth / Telemedicine Clinical Trial Site. Ask you online to take your temperature and report on your COVID-19 symptoms every day for about 30 days. To do this online, we will give you information about an electronic document to use, and we will show you how to use it. But there is no paper form available if you are unable to use the electronic document.

There are 2 visits in the first week (day 1 and day 3), and then visits once in week 3. These visits may take place in the clinic, in your home or wherever you are staying. Some parts of these visits could be completed over the phone or through electronic communication such as text messages and email, in addition to the sample collections done in person. We will also contact you about one month after the final study visit to check on your health.

If you decide to join the study, we will online review your medical history at your enrollment to see if you meet the study requirements. If you are eligible, we will online have you sign and date this consent form and collect more information about your health and medications you take. We online will also ask you about any symptoms you may have related to COVID-19. We may online need to review your medical records for additional information. We will also online ask you to provide the name of a person who can provide information for you if you are unable to do it yourself.

At this first visit, we will do following:

- Take testing COVID-19 by standard RT-PCR assay. BCG should not be given to individuals previously **infected with** *M. tuberculosis*. A person given the tuberculin skin test [10 tuberculin units (10 TU) purified protein derivative (PPD)] must return within 48 hours. The positive is skin test reading > 5 mm induration at 48 hours.
- Collect nasal swab, nasal wash, and saliva samples maybe sometimes. You may be able to do these procedures yourself, or the clinic staff may do them for you.
- Take testing PPD by Intradermal Injection (ID) assay.

The first visit could last from 1 to 2 hours.

For the online remainder of the visits, we want to collect medical information to see how your next steps develop or change over time. At these visits we will:

- Take testing COVID-19 by standard RT-PCR assay at week 1 day 1.
- Take testing PPD by Intradermal Injection (ID) assay at week 1 day 1.
- Review medical information, data, results, and symptoms at week 1 day 3.
- Take Intradermal Injection (ID) of Ad26 COVID-19 Spike plus TICE® BCG mix at week 1 day 3.
- Ask online about any new symptoms you are having since the last time the clinic staff saw you, and after Ad26 COVID-19 Spike plus TICE® BCG mix for intradermal injection you were taking.
- Collect medical information. You may be able to do this procedure yourself, or the research or referral clinic staff can do it.
- Collect medical data. You may be able to do this procedure yourself, or the clinic staff can do it. This may depend on what the clinic staff is allowed to do and may change over time.
- Take testing COVID-19 by standard RT-PCR assay at week 3 day 4. If you are COVID-19 positive, we will refer you to other clinic or hospitals. If you have symptoms change, we will adjust the plan we take for this study and make sure you get the care you need. We will online ask if you will stop to join the clinical trial for SARS-CoV-2. If you do so, we will let you have a referral to other clinic or hospital.
- Take testing COVID-19 by standard RT-PCR assay at week 3 day 4. If you are COVID-19 negative, we will check on your health until to the end of month 3.
- Collect online medical information, data, results, and symptoms every day.
- Remind you online about reporting on your COVID-19 symptoms every day.

These visits could last from a ½ hour to 2 hours.

Table of procedures

Procedure	Online Enrollment	Visit 1 Month 1 Week 1 Day 1	Visit 2 Month 1 Week 1 Day 3	Visit 3 Month 1 Week 3 Day 4	Online Notification	Visit 4 Month 2	Visit 5 Month 3
Medical history *	√	√	√	√	√	√	√
Test COVID-19 (RT-PCR) **		√					
Test PPD ***		√					
Review Results			√	√	√		
Biological Product ID			√				
Online Interview & Online Questionnaire	√	√	√	√	√	√	√

* This includes online reporting on COVID-19 symptoms every day for up to 30 days

** Take testing COVID-19 by standard RT-PCR assay.

*** BCG should not be given to individuals previously infected with M. tuberculosis. A person given the tuberculin skin test [10 tuberculin units (10 TU) purified protein derivative (PPD)] must return within 48 hours. The positive is skin test reading > 5 mm induration at 48 hours.

5. New Information

Information about SARS-CoV-2 and COVID-19 is changing every day. We may online ask you to complete additional information. This may be because we would like to obtain your additional information. We will online keep you updated about any new information we learn that may change your decision to be involved in this study.

We may also online contact you after the study ends (for example, to tell you about the study results).

If you agree, we will also online collect medical information, data, results, and symptoms. At the end of this form, we will ask if you allow us to collect your medical information. You can decide not to give these medical data online, but you cannot still be in the study.

We would like online to collect your medical information, data, results, and symptoms. We want to learn if your response to the COVID-19 virus is influenced by the biological product. We also online want to know if SARS-CoV-2 is present in your assay by RT-PCR, and if that changes over time. We will online ask you to do this at most study visits (5 times) as shown in the table above. We will online collect this information. These can be collected online and can be done by yourself, or some clinic staff may be able to do it for you. We will give you online handout that describes these procedures in more detail. We will also ask you questions about your something medical.

6. If you are hospitalized or unable to continue to give consent for study procedures, we will pause your participation.

Some people with COVID-19 develop severe symptoms and must be hospitalized. This can include being put on a ventilator to help you breathe. In the event you are not able to communicate and give consent for the study procedures to continue, we will pause your study participation. At any time that you are awake again and able to communicate, we will ask if you would like to continue participating in the study. We will only resume the study procedures if you give us permission to do so.

It is also possible that you could be admitted to a hospital where the clinic staff are not able to visit you. If this happens, we will pause your study participation. When you are released from the hospital, we will ask if you would like to continue participating in the study.

7. Compensation for Participation

We will give you a receipt document for each study visit you complete.

You do not have to pay anything to be in this study. The study procedures and study visits will be done at no charge to you or your insurance company.

8. The Health Care Provider will use your COVID-19 samples for research related to your response to COVID-19 via biological product Intradermic Injection (ID).

Clinic will send your COVID-19 samples (with your name and other identifying information) to some labs approved by the Health Care Provider for this study, which are located in the United States.

Labs will do RT-PCR testing related to only this study on your COVID-19 samples. Researchers will study only the enrolled cases related to COVID-19, and those biological products that affect how people get treated after infected with the SARS-CoV-2 virus that causes COVID-19.

In enrolled cases, labs will take testing COVID-19 by standard RT-PCR assay, so that help those researchers can contribute to this study.

This RT-PCR test done on your COVID-19 sample is for research purpose, and to check your existing COVID virus. However, the telehealth / telemedicine clinical trial site may online provide you the results of testing by standard RT-PCR assay for SARS-CoV-2 if you choose to know the results. When your COVID-19 samples are no longer needed for this study, the clinical trial site will not continue to store them. You cannot be asked for future studies using your stored samples. You will not be asked to sign a separate consent form for this purpose.

9. Confidentiality

We just only can do limited to protect your private information.

Your study records and samples will be kept in a secure location and kept confidential in clinical trial term. We will label all of your COVID-19 samples and your study records with a code number, not your name or other personal information. However, it is possible to identify you, if necessary. And, if your COVID-19 testing result is positive, your medical information must be open. Meanwhile, we will share your name with the lab that does the RT-PCR tests on your COVID-19 samples, or with anyone else who need to know your name.

Clinic staff will have access to your study records. Your records may also be reviewed by groups who watch over this study to see that we are protecting your rights, keeping you safe, and following the study plan. These groups include:

- The sponsor, Han Xu, M.D., Ph.D., FAPCR, Sponsor-Investigator, Medical Monitor, Safety Officer, IRB Chair, IORG Director -- Health Care Provider - Individual (NPI - 1831468511)
- Any regulatory agency that reviews studies like this US Food and Drug Administration (FDA)
- The Institutional Review Board (IRB), which is Medicine Invention Design Incorporation (MIDI) IRB #1 (IRB00009424)
- The COVID-19 Health Care Provider and people who work for them:
 - ✓ Medicine Invention Design Incorporation (FWA00015357)
 - ✓ Medicine Invention Design Incorporation (IORG0007849)
 - ✓ Health Care Provider - Group/Organization (NPI - 1023387701)
 - ✓ Health Care Provider - Research Clinic/Center (Code - 261QR1100X)
 - ✓ Health Care Provider - Individual (NPI - 1831468511)
Han Xu, M.D., Ph.D., FAPCR, Sponsor-Investigator, Medical Monitor, Safety Officer, IRB Chair, IORG Director
- US Department of Health and Human Services (DHHS):
 - ✓ Health Care Provider - Organization (NPI - 1023387701)
 - ✓ Health Care Provider - Research Clinic/Center (Code - 261QR1100X)
 - ✓ Health Care Provider - Individual (NPI - 1831468511)
- The US Office for Human Research Protections:
 - ✓ FWA00015357
 - ✓ IORG0007849
 - ✓ IRB00009424

All reviewers will take steps to keep your records private.

We cannot guarantee absolute privacy. At this telehealth / telemedicine clinical trial site, we have to report COVID-19 test results to local health authorities, for example, US Centers for Disease Control and Prevention (CDC). The local health authorities may also require that we share your demographic information and information about your COVID-19 test results to support contact tracing.

During your participation in this study, study staff may need to review and where necessary make and keep copies of some of your medical records (for example, records of hospitalizations). Any medical records information requested for this study will be kept confidential in the same manner as your other study records

The results of this study may be published. No publication will use your name or identify you personally.

We may share information from the study with other researchers. We will not share your name or information that can identify you.

A description of this study will be available on <http://www.ClinicalTrials.gov>. This website will not include information that can identify you. At most, the website will include a summary of the results. You can search this website at any time.

10. Risks

The potential risks associated with this study are minimal.

This section describes the risks we know about. There may also be unknown risks. We will tell you if we learn anything new that may affect your willingness to stay in the study.

Risks of taking blood:

In this study, we will take blood. This procedure can cause bruising, pain, feeling faint or dizzy, soreness, redness, swelling, itching, a sore, infection, or bleeding. Taking blood can cause a low blood cell count (anemia), making you feel tired. If you become pregnant, anemia may increase the risk of complications for you or your baby, and so we may reduce or discontinue your blood draws.

Risks of the nasal swab procedure:

The feeling of having a small, soft-tipped swab inserted into your nostril and twirled around may be a little uncomfortable, but it should not be painful. There is a small chance there could be some bleeding, but this is unlikely.

Risks of the nasal wash procedure:

The nasal wash may be uncomfortable, but it should not be painful. There is a small chance that the sterile salt water could go into your throat, causing you to cough or feel like you are choking, but this is unlikely.

Risks of collecting a saliva sample:

There are no known risks of giving a sample of your saliva.

Urine sample collection:

There are no known risks for collecting urine by routine clean catch methods.

Risks of swabbing stool:

There are no known risks of having your stool swabbed to collect a sample.

Risks of collecting stool by a rectal swab:

If you choose to provide a sample by rectal swabbing, you may feel pressure as the swab is inserted into your rectum. It may be uncomfortable, but it is usually not painful. Some people may have a little bit of bleeding.

Risks of disclosure of your personal information:

We will take several steps to protect your personal information. Although the risk is very low, it is possible that your personal information could be given to someone who should not have it. If that happened, you could face discrimination, stress, and embarrassment.

Study records will be kept in secure locations in the study clinic. Study records and specimens will be labeled only with a code. All information entered into a computer is secured and will only be labeled with a code. Your names will be written on some records that are kept in the research clinic. Your study records will only be accessed by study staff given permission to do so. Study records will continue to be kept under confidential conditions according to the sponsor and IRB's guidelines and requirements.

We cannot guarantee absolute privacy. At this telehealth / telemedicine clinical trial site, we have to report COVID-19 test results to local health authorities, for example, US Centers for Disease Control and Prevention (CDC). The local health authorities may also require that we share your demographic information and information about your COVID-19 test results to support contact tracing.

Risks of genetic testing:

If your genetic research data are shared with unauthorized users, you may be at risk of loss of the privacy of your health data. It is unlikely, but the genetic tests done on your samples could show you may be at risk for certain diseases. If others found out, it could lead to discrimination or other problems. However, it is almost impossible for you or others to know your test results from the genetic testing. The results are not part of your study records and are not given to you.

11. Benefits

The study will **not** benefit you directly.

12. Alternatives to Participation

This study is for research purposes only. The only alternative is to not participate in this study. This study may help us learn more about SARS-CoV-2 infection and COVID-19. The study results may be used to guide future research, treatment and prevention that could help others in the future.

13. Your rights and responsibilities

If you join the study, you have rights and responsibilities.

You have many rights that we will respect. You also have responsibilities. We list these in the Bill of Rights and Responsibilities for Research. We will give you a copy of it.

14. Voluntary Participation/Leaving the study

Tell us if you decide to leave the study.

You are free to leave the study at any time and for any reason. Your care at this clinic and your legal rights will not be affected, but it is important for you to let us know. We may take you out of the study at any time.

We may take you out of the study if:

- You do not follow instructions,
- We think that staying in the study might harm you,
- The study is stopped for any reason.

15. Compensation for Injuries

It is unlikely that you will be injured as a result of study participation. If you are injured, the clinic personnel will give you immediate necessary treatment for your injuries and refer you to the local hospital for further care as necessary. You will not have to pay for treatment provided through the public health facilities. If you receive treatment at a private facility, you and/or your medical aid will be responsible for treatment costs. There is no program to pay money or give other forms of compensation for such injuries even if they are study related through Medicine Invention Design, Inc.

By signing and dating this document, you will not lose any of your legal rights or release anyone involved in the research from responsibility for mistakes.

16. Whom to contact about this study

During the study, if you experience any medical problems, suffer a research-related injury, or have questions, concerns or complaints about the study, please contact the study site personnel below:

Han Xu, M.D., Ph.D., FAPCR, Sponsor-Investigator, Medical Monitor, Safety Officer, IRB Chair, IORG Director

Medicine Invention Design Incorporation (MIDI) IRB #1 (IRB00009424)

Health Care Provider - Individual (NPI - 1831468511)

Health Care Provider - Clinical Ethicist (Code - 174V00000X)

Medicine Invention Design Incorporation (FWA00015357)

Medicine Invention Design Incorporation (IORG0007849)

Health Care Provider - Group/Organization (NPI - 1023387701)

Health Care Provider - Research Clinic/Center (Code - 261QR1100X)

Mail: 5545 Burnside Drive, Rockville, MD 20853

Call: 301-222-7143

Fax: 866-458-0099

E-mail: hanxumd@gmail.com

If you have any questions about your rights as a research subject, and/or concerns or complaints regarding this research study, contact:

Sponsor-Investigator, Medical Monitor, Safety Officer, IRB Chair, IORG Director

Tel: 301-222-7143

Fax: 866-458-0099

E-mail: hanxumd@gmail.com

Health Care Provider - Group/Organization

Call: 301-222-7143

Email: midinc@hotmail.com

Enrollment ICF Form
Telehealth / Telemedicine
Clinical Trial

17. Ad26 COVID-19 Spike plus TICE® BCG Mix for Intradermal Injection

The biologics is administered percutaneously utilizing a sterile, single-use multiple puncture device. The multiple puncture device consists of a plastic holder for a thin, wafer-like stainless steel plate from which 36 points protrude (Figure 1). After the biologics is prepared, the skin site is cleansed with an alcohol or acetone sponge and allowed to dry thoroughly.

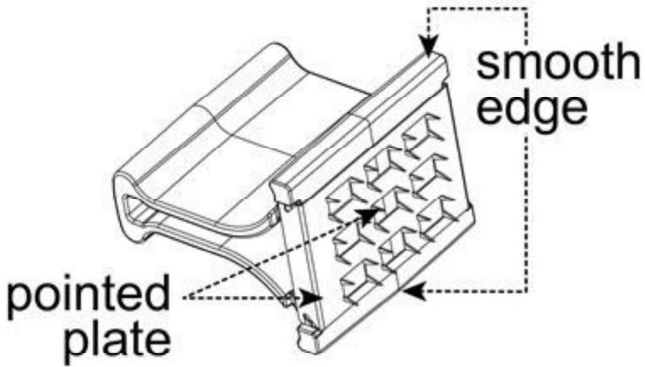


Figure 1

1. Administer the biologics in the deltoid region (Figure 2). Position the arm to maintain a horizontal surface where the biologics is to be placed.

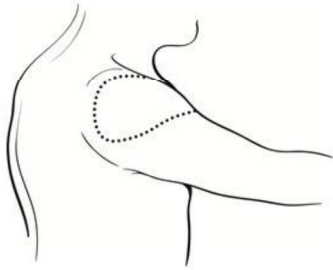


Figure 2

2. Drop the dose of 1 mL of the biologics from the syringe and needle onto the cleansed surface of the skin (Figure 3) and spread over a 1" by 2" area using the smooth edge of the multiple puncture device (Figure 4).

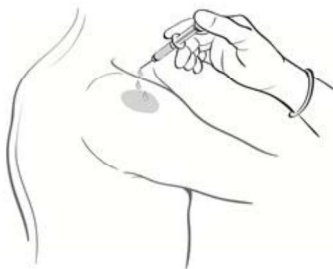


Figure 3

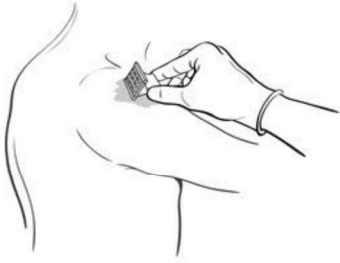


Figure 4

3. Grasp the arm firmly from underneath, tensing the skin. Center the multiple puncture device over the biologics and apply firm downward pressure such that the device points are well buried in the skin (Figure 5).

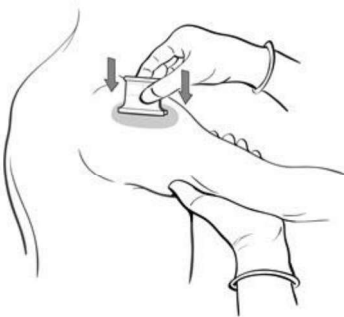


Figure 5

4. Maintain pressure for 5 seconds. Do not "rock" the device. Release the pressure underneath the arm and remove the device. In a successful procedure the points puncture the skin. If the points do not puncture the skin, the puncture procedure must be repeated.
 5. After successful puncture, spread the biologics as evenly as possible over the puncture area with the smooth edge of the device (Figure 4). An additional 1–2 drops of the biologics may be added to ensure a very wet vaccination site.
 6. Use the multiple puncture device once and discard in a standard biohazardous sharps container.
 7. Loosely cover the site and keep dry for 24 hours.
 8. Advise the patient that the biologics contains live organisms. Although the biologics will not survive in a dry state for long, infection of others is possible.
- BCG should not be given to individuals previously infected with M. tuberculosis.
 - **The biologics can use Needle-Free Jet Injector or Needle-Free Syringe**

18. Your permissions, signature and date

In the section titled “New Information” of this form, we told you about collecting your medical information. Please make your mark in the box next to the option you choose.

I agree to provide my medical information.

I do not agree to provide my medical information.

If you agree to join this study, you will need to sign and date or make your mark below. Before you sign and date or make your mark on this consent form, make sure of the following:

- You feel that you understand what the study is about and what will happen to you if you join. You understand what the possible risks and benefits are.
- You agree to have the clinic staff visit the place where you are staying or your hospital room to do the study visits and sample collections.
- You have had your questions answered and know that you can ask more.
- You agree to join this study.
- You will not be giving up any of your rights by signing and dating this consent form.

Participant’s name (print)	Participant’s signature or mark	Date	Time
Study staff conducting consent discussion (print)	Study staff signature	Date	Time

For participants who are unable to read or write, a witness should complete the signature and date block below:

Witness’s name (print)	Witness’s signature	Date	Time
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*Witness is impartial and was present for the entire discussion of this consent form.