

Confidential Documents reveal Moderna sent mRNA Coronavirus Vaccine Candidate to University Researchers weeks before emergence of Covid-19

BY THE DAILY EXPOSE ON JUNE 18, 2021 • (8 COMMENTS)



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U.S. Gov't Sent "mRNA Coronavirus Vaccine Candidates" to University Researchers WEEKS BEFORE "COVID" Outbreak in China!

A confidentiality agreement shows potential coronavirus vaccine candidates were transferred from Moderna to the University of North Carolina in 2019, nineteen days prior to the emergence of the alleged Covid-19 causing virus in Wuhan, China.

The confidentially agreement which can be viewed [here](#) states that providers 'Moderna' alongside the 'National Institute of Allergy and Infectious Diseases' (NIAID) agreed to tranfer 'mRNA coronavirus vaccine candidates' developed and jointly-owned by NIAID and Moderna to recipients 'The Universisty of North Carolina at Chapel Hill' on the 12th December 2019.

PUBLIC HEALTH SERVICE

MATERIAL TRANSFER AGREEMENT

This Material Transfer Agreement ("MTA") has been adopted for use by the National Institutes of Health, the Food and Drug Administration and the Centers for Disease Control and Prevention, collectively referred to herein as the Public Health Service ("PHS") in all transfers of research material (Research Material) whether PHS is identified below as its Provider or Recipient.

Providers: *National Institute of Allergy and Infectious Diseases, National Institutes of Health ("NIAID")*
ModernaTX, Inc ("Moderna")

Recipient: The University of North Carolina at Chapel Hill

1. Provider agrees to transfer to Recipient's Investigator the following Research Material:

mRNA coronavirus vaccine candidates developed and jointly-owned by NIAID and Moderna.

Found on page 105 of the agreement

The material transfer agreement was signed the December 12th 2019 by Ralph Baric, PhD, at the University of North Carolina at Chapel Hill, and then signed by Jacqueline Quay, Director of Licensing and Innovation Support at the

**MATERIAL TRANSFER AGREEMENT
SIGNATURE PAGE**

FOR RECIPIENT:

Recipient's Investigator



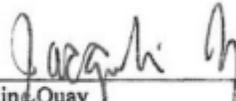
Ralph Baric, PhD
Professor

Date: 12/12/2019

Mailing Address for Materials:

Attention: Dr. Rachel Graham, Department of
Epidemiology, University of North Carolina at
Chapel Hill, 135 Dauer Drive, 2101 McGavran-
Greenberg Hall, CB #7435, Chapel Hill, NC 27599-
7435

Duly Authorized



Jacquelin Quay
Director, Licensing & Innovation Support, OTC

Date: 12/16/19

Mailing Address for Notices:

The University of North Carolina at Chapel Hill
Office of Technology Commercialization
109 Church Street, Chapel Hill, NC 27516

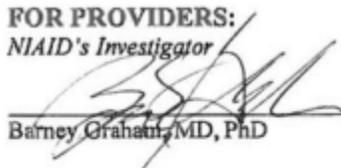
Tel: 919-966-3929 Fax: 919-962-0646

Recipient signatories found on page 107

The agreement was also signed by two representatives of the NIAID, one of whom was Amy F. Petrik PhD, a technology transfer specialist who signed the agreement on December 12th 2019 at 8:05 am. The other signatory was Barney Graham MD PhD, an investigator for the NIAID, however this signature was not dated.

Tel: 919-966-3895 Fax: _____

FOR PROVIDERS:
NIAID's Investigator



Barney Graham, MD, PhD

Date: _____

Duly Authorized

Amy F.

Amy Petrik, PhD
Technology Transfer Specialist, TTIPO, NIAID

Digitally signed by Amy
F. Petrik -S
Date: 2019.12.12
08:05:22 -05'00'

Date: _____

Mailing Address for Notices:

Technology Transfer and Intellectual Property Office
National Institute of Allergy and Infectious Diseases
Department of Health and Human Services
Suite 6D, MSC 9804
5601 Fishers Lane
Rockville, MD 20852
Tel: 301/496-2644 Fax: 240-627-3117

NIAID signatories found on page 107

The final signatories on the agreement were Sunny Himansu, Moderna's Investigator, and Shaun Ryan, Moderna's Deputy General Counsel. Both signatories were made on December 17th 2019.

Moderna's Investigator



Sunny Himansu, PhD

Date: 12/17/2019

Duly Authorized



Shaun Ryan
Deputy General Counsel

Date: 12/17/19

Mailing Address for Notices:

ModernaTX, Inc.
200 Technology Square
Cambridge, MA 20139
Attn: General Counsel

Moderna signatories found on page 108

All of these signatures were made prior to any knowledge of the alleged emergence of the novel coronavirus. It wasn't until December 31st 2019 that the World Health Organisation (WHO) became aware of an alleged cluster of viral pneumonia cases in Wuhan, China. But even at this point they had not determined that an alleged new coronavirus was to blame, instead stating the pneumonia was of "unknown cause".

INFORMATION

31 Dec 2019

WHO's Country Office in the People's Republic of China picked up a media statement by the Wuhan Municipal Health Commission from their website on cases of 'viral pneumonia' in Wuhan, People's Republic of China.

The Country Office notified the International Health Regulations (IHR) focal point in the WHO Western Pacific Regional Office about the Wuhan Municipal Health Commission media statement of the cases and provided a translation of it.

WHO's Epidemic Intelligence from Open Sources (EIOS) platform also picked up a media report on ProMED (a programme of the International Society for Infectious Diseases) about the same cluster of cases of "pneumonia of unknown cause", in Wuhan.

Several health authorities from around the world contacted WHO seeking additional information.

It was not until January 9th 2020 that the [WHO reported](#) Chinese authorities had determined the outbreak was due to a novel coronavirus which later became known as SARS-CoV-2 with the alleged resultant disease dubbed COVID-19. So why was an mRNA coronavirus vaccine candidate developed by Moderna being transferred to the University of North Carolina on December 12th 2019?

The same Moderna that have had an mRNA coronavirus vaccine authorised for emergency use only in both the United Kingdom and United States to allegedly combat Covid-19.

What did Moderna know that we didn't? In 2019 there was not any singular coronavirus posing a threat to humanity which would warrant a vaccine, and evidence suggests there hasn't been a singular coronavirus posing a threat to humanity throughout 2020 and 2021 either.

Considering the fact a faulty PCR test has been used at a high cycle rate, hospitals have been empty in comparison to previous years, statistics show just 0.2% of those allegedly infected have died within 28 days of an alleged positive test result, the majority of those deaths by a mile have been people over the age of 85, and a mass of those deaths were caused by a drug called midazolam, which causes respiratory depression, and respiratory arrest.

Perhaps Moderna and the National Institute of Allergy and Infectious Diseases would like to explain themselves in a court of law?