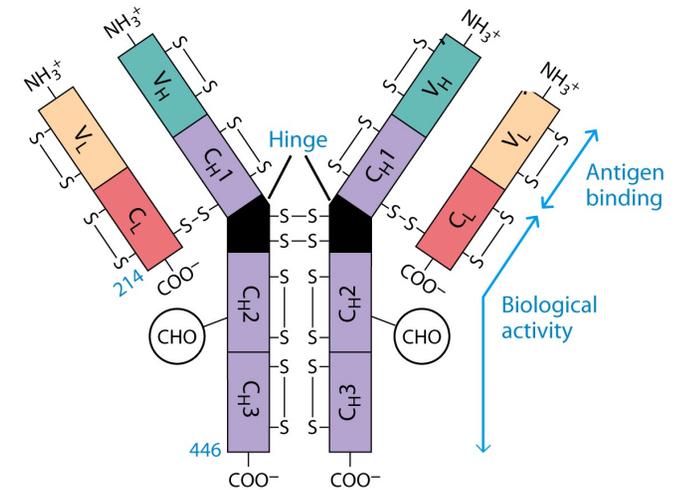


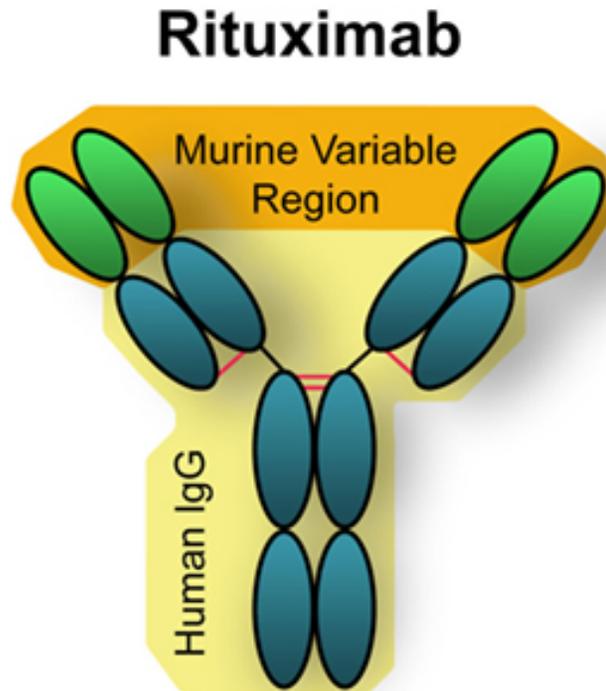
Antibodies as medicine and diagnostics

Engineering antibodies for medicine and diagnostics

- Humanized monoclonal antibodies as drugs
 - Rituximab (1997)
 - LY-CoV555 (2020)
- Antibody fragments (scFvs): bispecific antibodies
 - Blinatumomab (2009)
- Antibody drug conjugates
 - Trodelvy (2020)
- Lateral flow immunochromatographic assay
 - human chorionic gonadotropin, hormone (pregnancy) test (1970s)
 - COVID-19 lateral flow assay (2020)

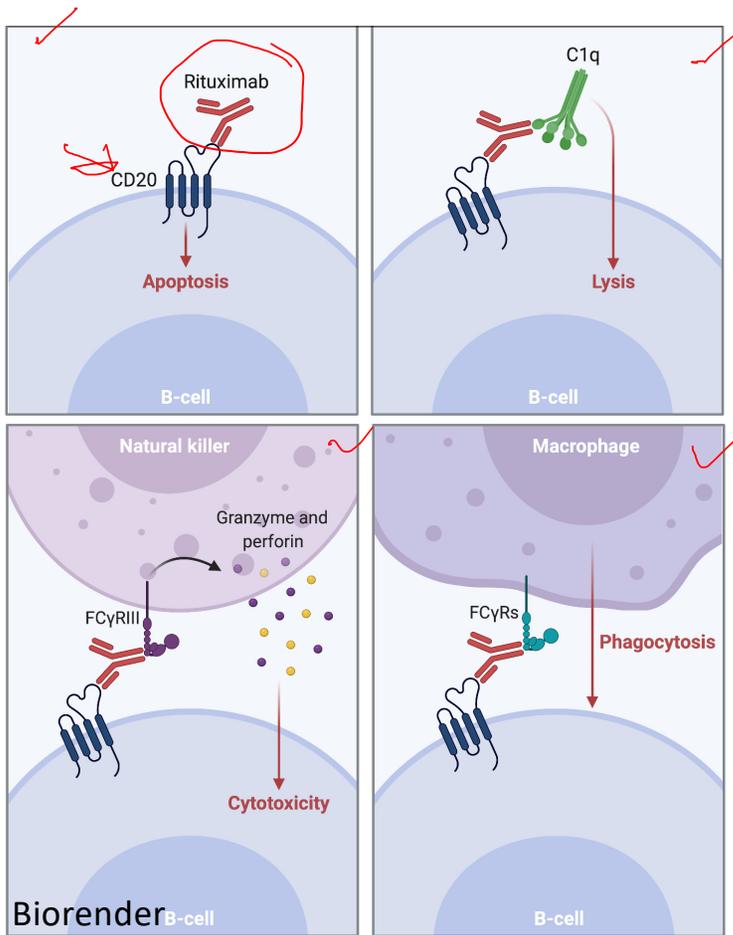


Rituximab is treatment for cancer and autoimmune diseases



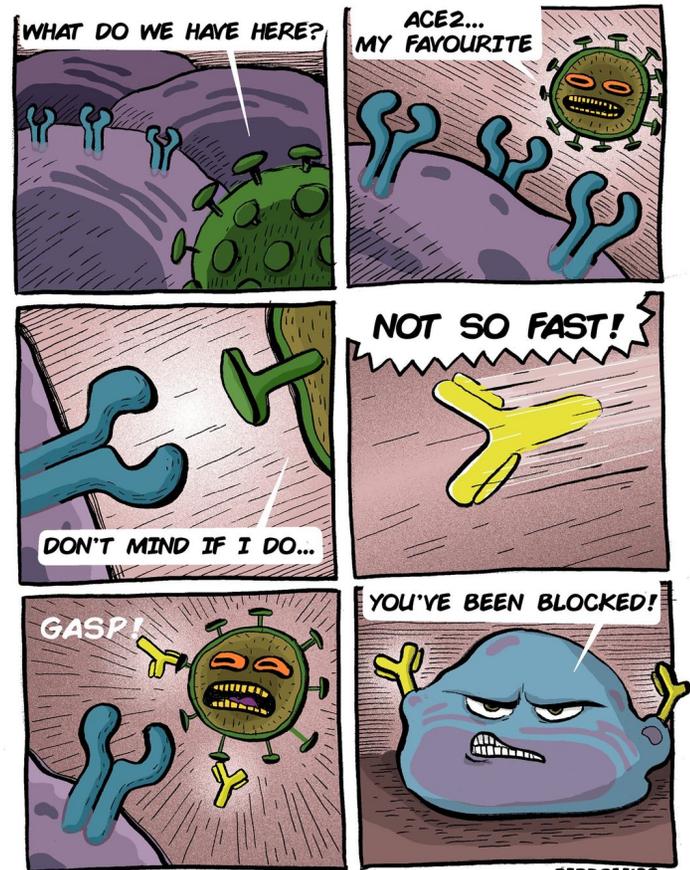
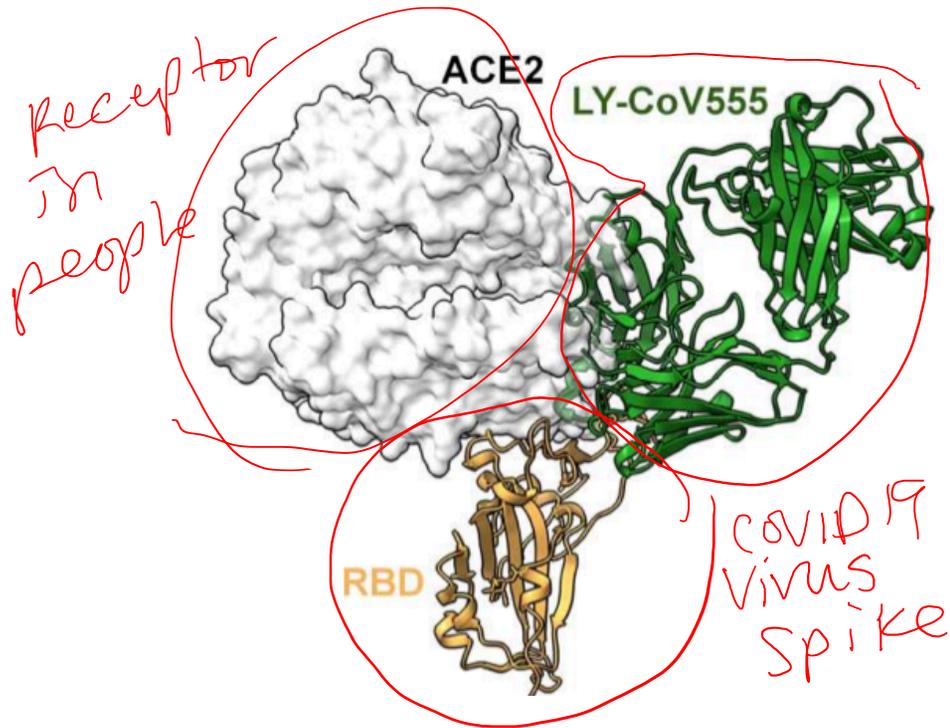
- First antibody FDA approved for cancer treatment in 1997
- Antigen is CD20
- Produced in CHO cells
- Approved for treatment of:
 - non-Hodgkin's lymphoma
 - chronic lymphocytic leukemia
 - rheumatoid arthritis
 - multiple sclerosis

Rituximab targets and kills CD20+ B cells



- Binding of Rituximab to CD20+ cells can result in:
 - CD20 mediated apoptosis
 - Complement-dependent cytotoxicity (direct lysis)
 - antibody-dependent cell-mediated cytotoxicity (Natural Killer cell)
 - antibody-dependent phagocytosis (macrophage phagocytosis)
- Need better mouse models to study effects of immunotherapy to reduce resistance and side effects
 - Anti-human CD20 mouse models don't mimic the human immune system well enough

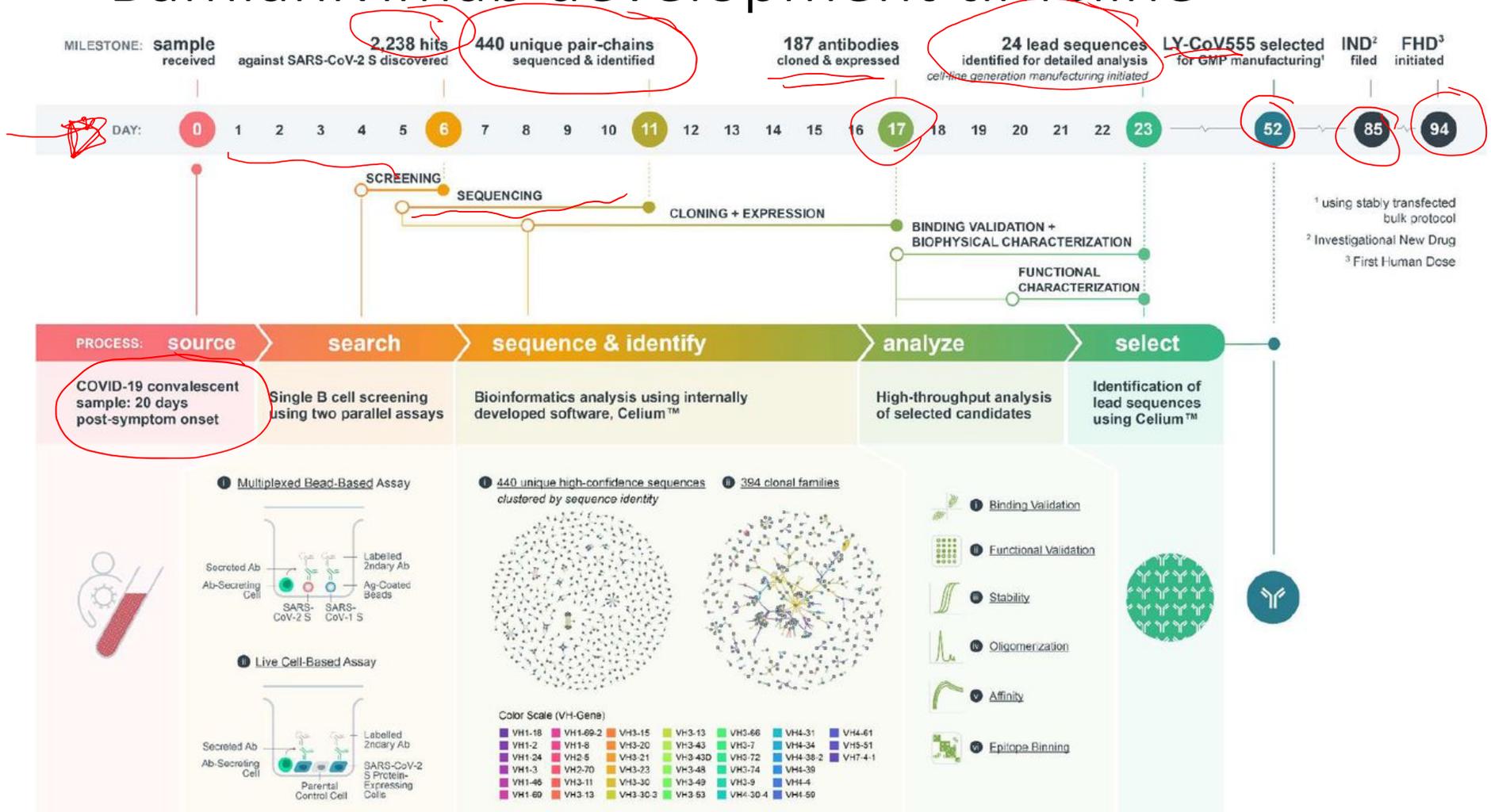
Bamlanivimab (LY-CoV555) first monoclonal antibody therapy approved for treatment of COVID-19



Jones, B. et al. LY-CoV555, a rapidly isolated potent neutralizing antibody, provides protection in a non-human primate model of SARS-CoV-2 infection. N Engl J Med. 2021.

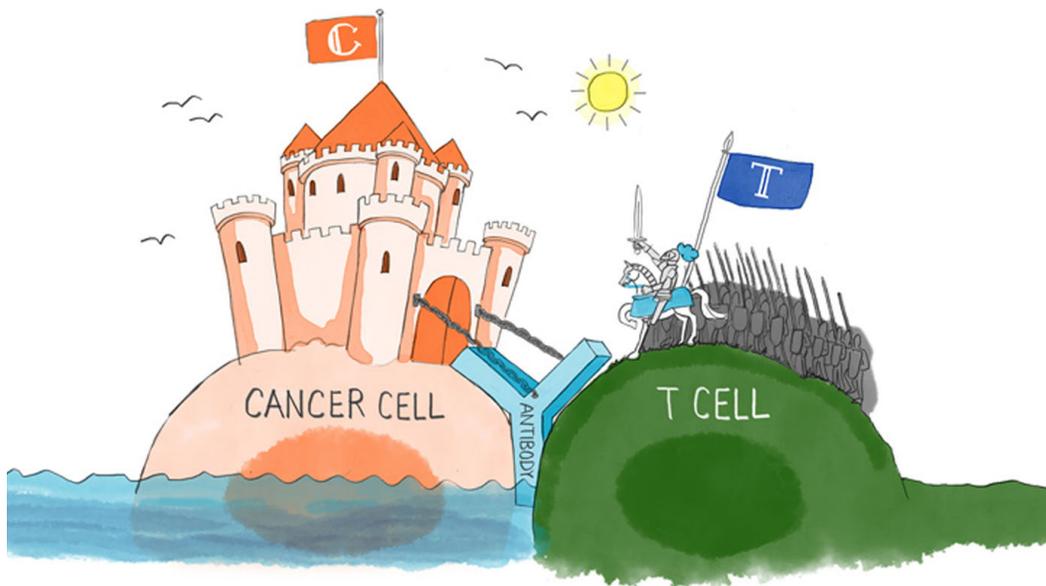
PEDROMICS

Bamlanivimab development timeline



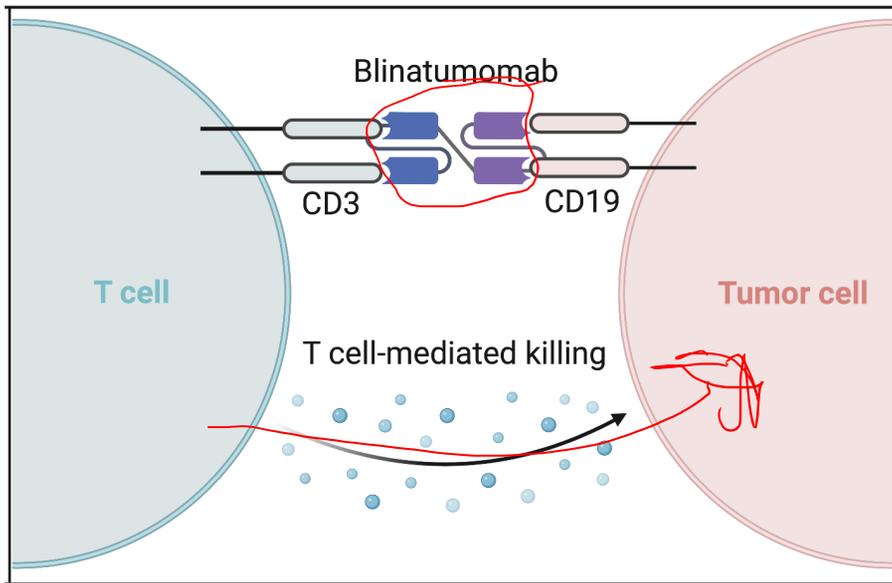
Jones, B. et al. LY-CoV555, a rapidly isolated potent neutralizing antibody, provides protection in a non-human primate model of SARS-CoV-2 infection. N Engl J Med. 2021.

Recombinant antibody production led to development of bispecific antibodies

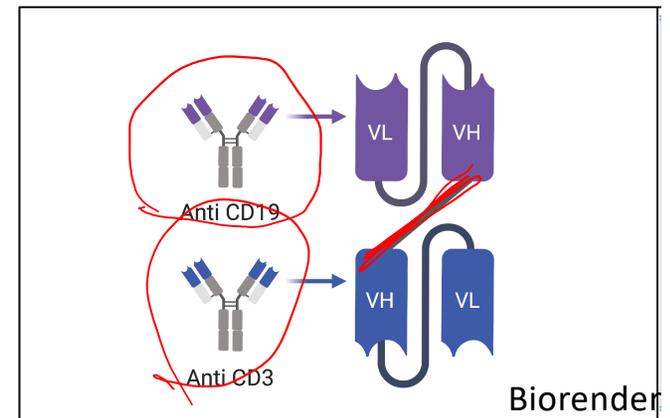


- A bispecific antibody contains two different antigen binding sites (scFvs) in one molecule
- Bispecifics generated in 1980s but not approved for use as drugs until 2009
- Blinatumomab: a bispecific T cell engager (BiTE) antibody against CD19/CD3 for refractory acute lymphoid leukemia

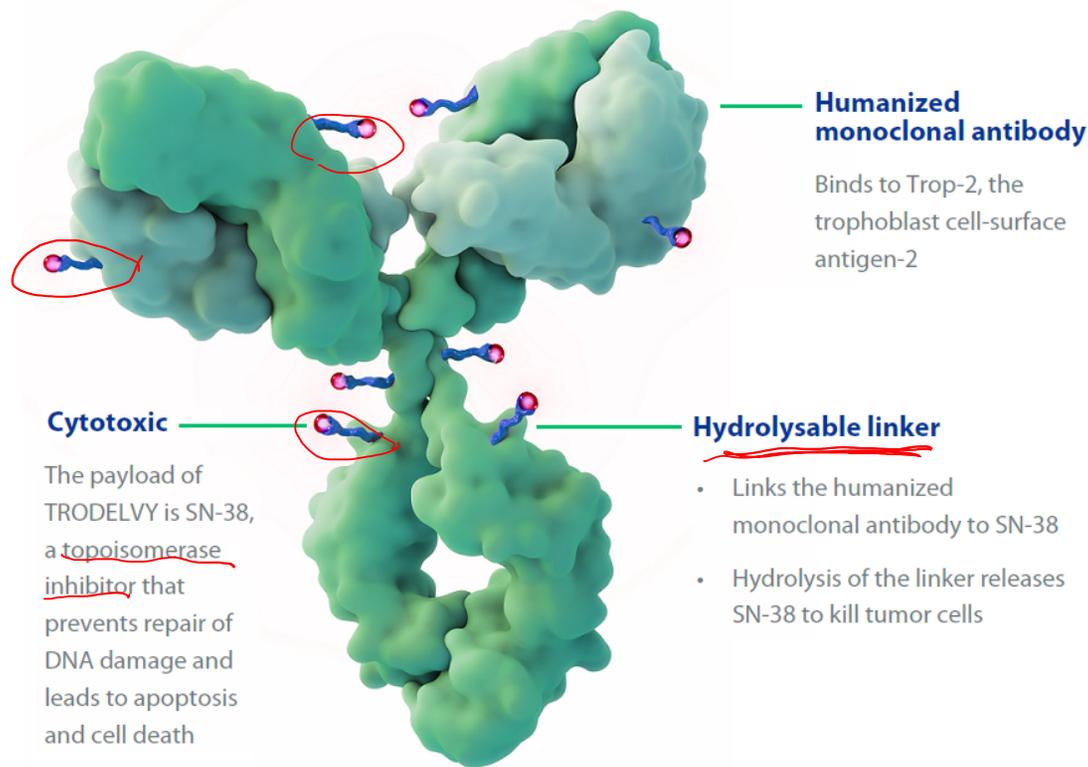
Blinatumomab complexes a T cell with a CD19+ cancer cell resulting in lysis



- ✱ Continuous intravenous administration of the drug is required
- CD19+ targets acute lymphoid leukemia
- CD3+ targets T cells and activates T cell cytotoxicity to kill the leukemia cell

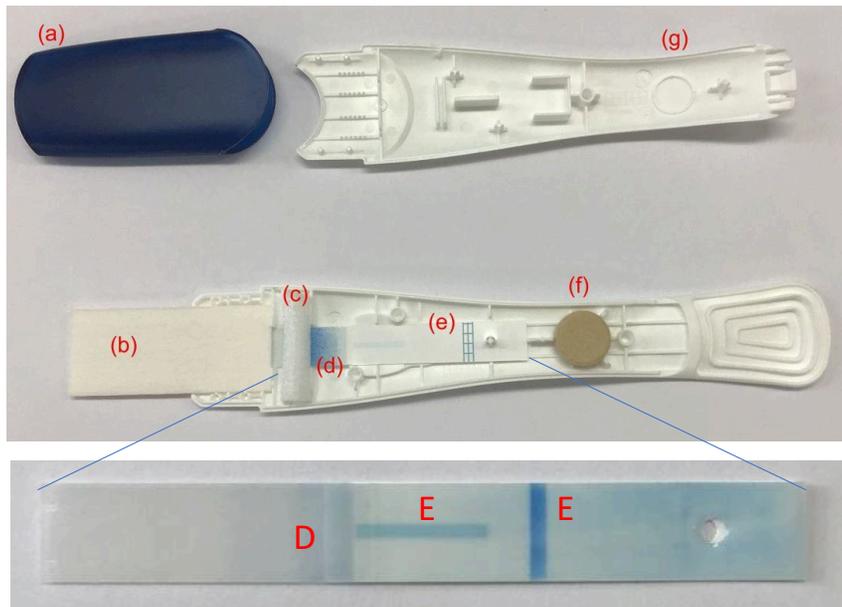


Trodelvy, drug conjugated monoclonal critical for patients with limited treatment options



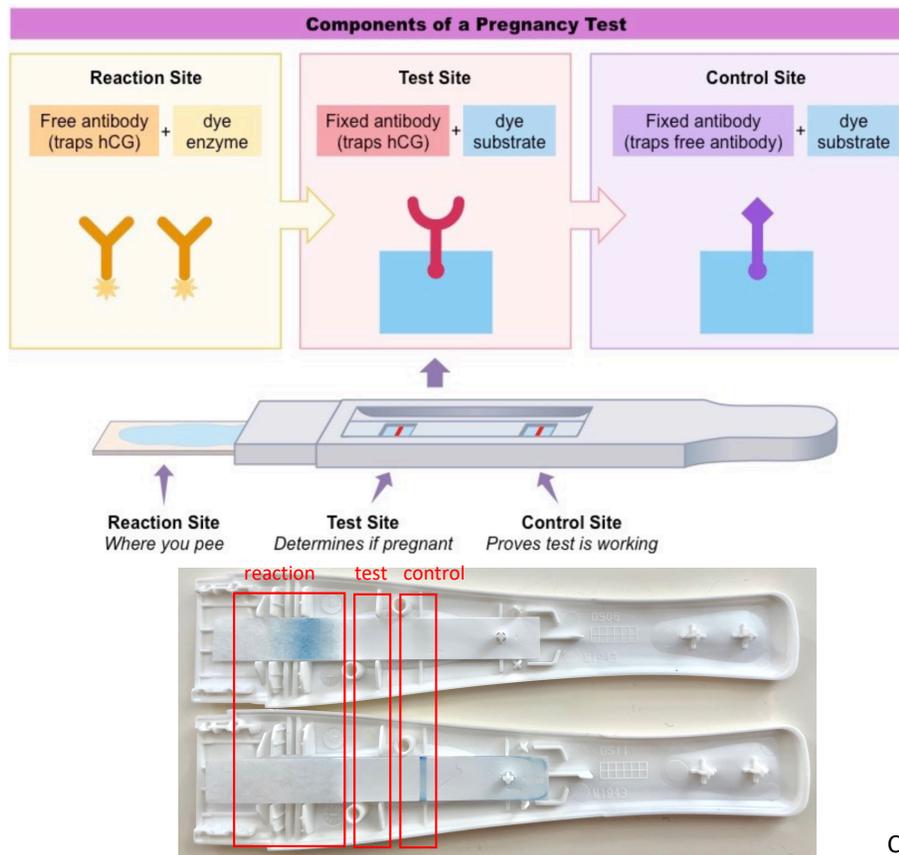
- Triple-negative breast cancer (TNBC) tumor cells limited treatment options are available
- Trop-2 is a surface protein highly expressed by many cancers, including approximately 90% of TNBC tumors
- Once inside the tumor cell, the linker connecting the antibody to the cytotoxic SN-38 is cleaved, releasing the active drug inside

A pregnancy test is a Lateral flow immunochromatographic assay



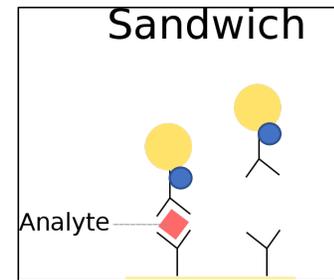
- A pregnancy test is a widely used antibody based diagnostic device
 - anti-hCG (human chorionic gonadotropin, hormone) is a hormone that increases as a pregnancy progresses
- (B) Absorbent pad, a filter helping to remove any proteins or bacteria in the urine that may affect the assay's performance, leaving mostly water and the hCG protein.
- (D) Conjugate pad with free antibody specific to hCG that is conjugated to blue dye
- (E) Nitrocellulose membrane with two antibody test lines

This design is used in many diagnostic devices and only necessitates a specific antibody

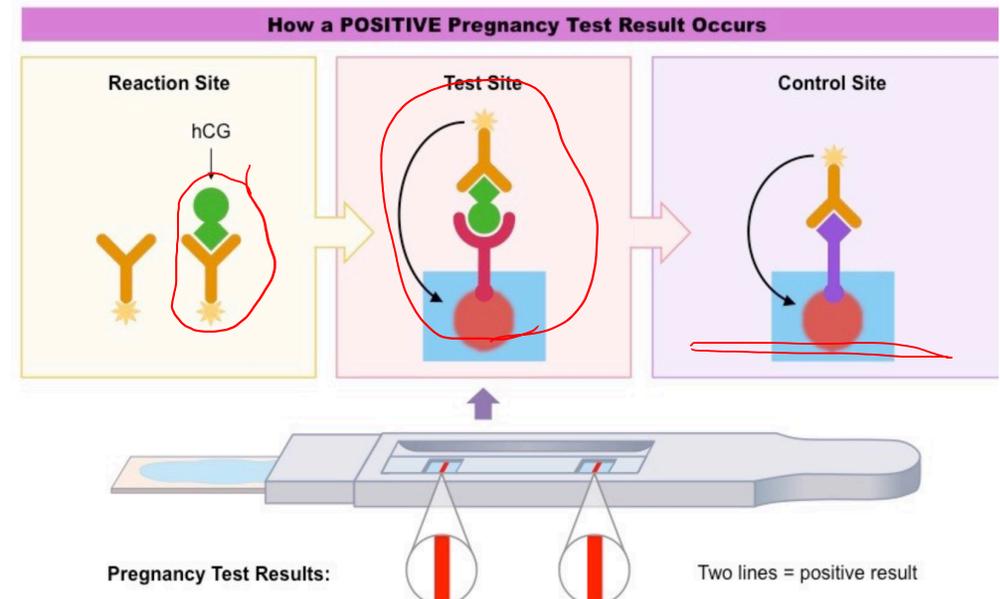
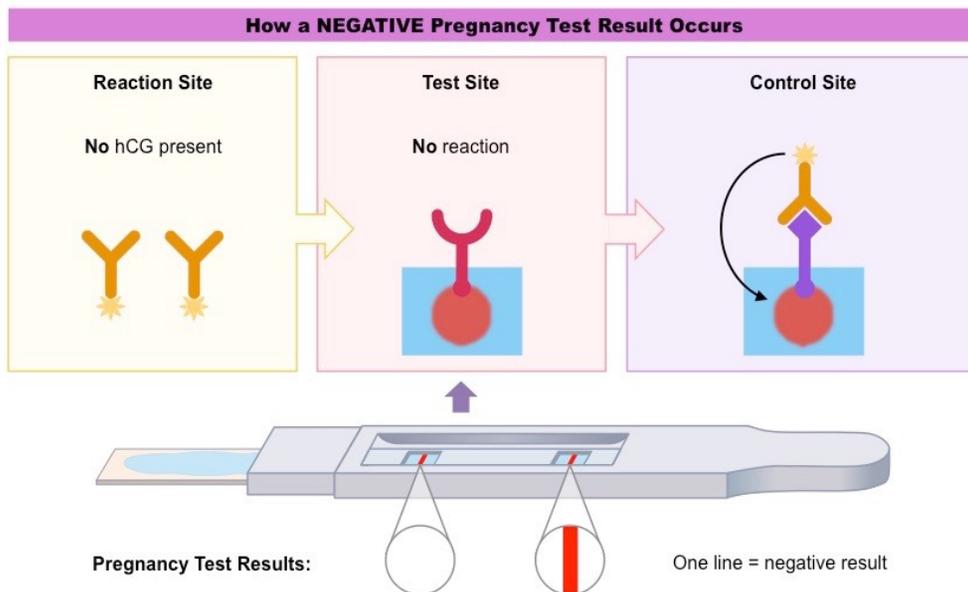


- 3 sites for antibody-antigen interactions:
 - Reaction site: free antibody first encounters hCG antigen
 - Test site: antibody-antigen complex can bind
 - Control site: all antibodies captured here to indicate test complete

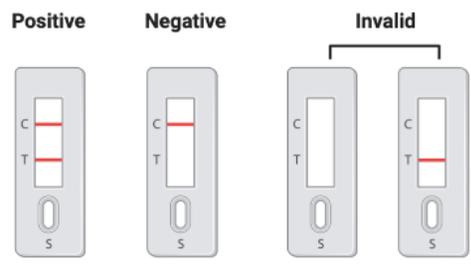
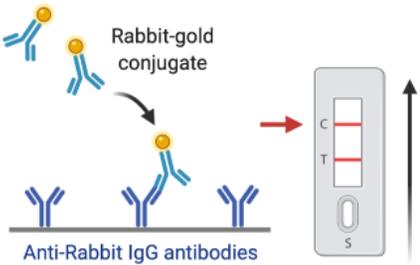
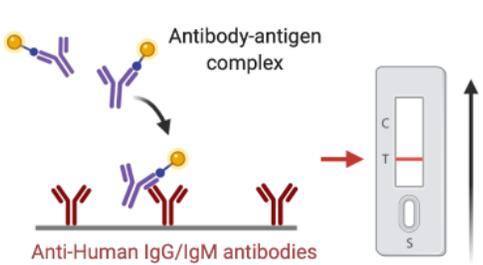
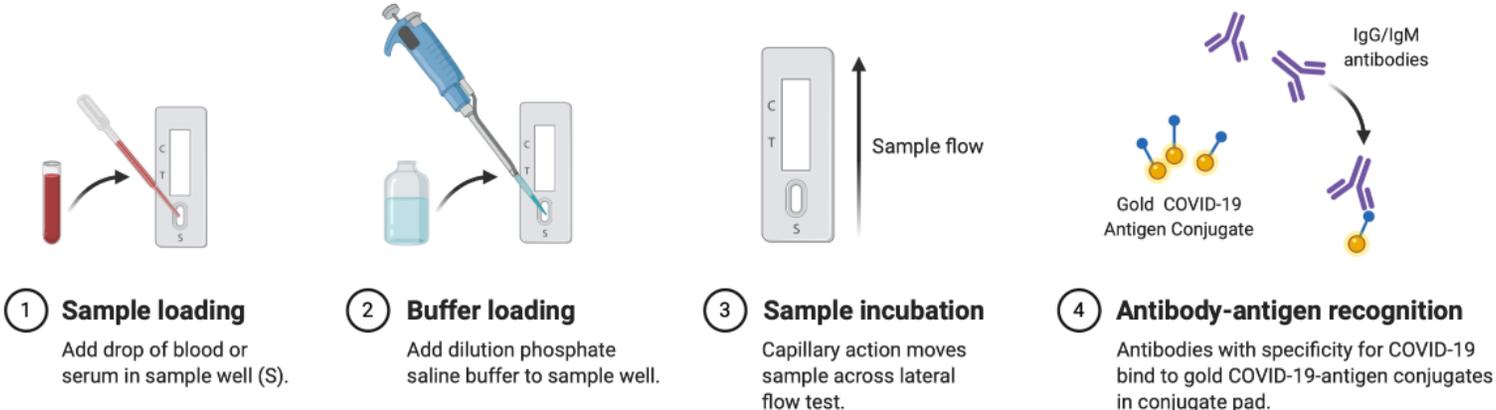
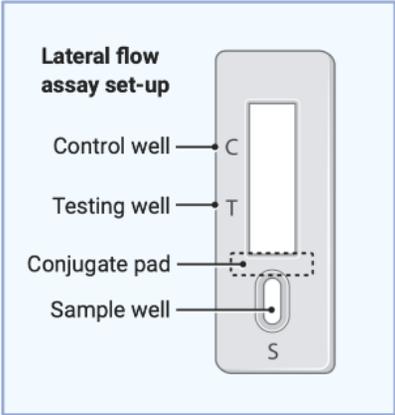
This diagnostic device is modified from a sandwich ELISA



- If test **negative**, free antibodies flow through strip and end at control site (nothing at test site)
- If test is **positive**: hCG binds free antibodies also bind to the antibodies in the test strip, stopping them from flowing through the test strip (some also bind at control).



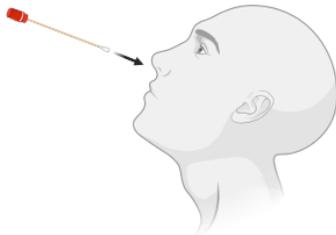
Serologic Diagnostic Test: COVID-19 Detection



COVID-19 Diagnostic Test through RT-PCR

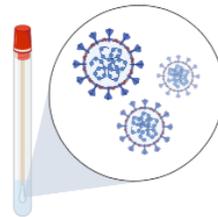
1 Nasopharyngeal swab <15 min

Cotton swab is inserted into nostril to absorb secretions.



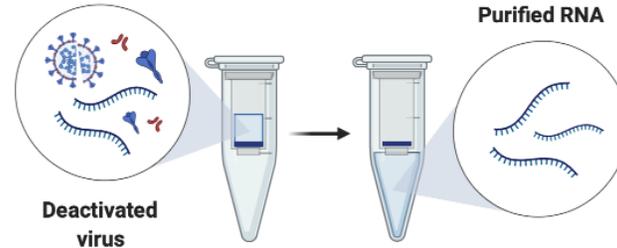
2 Collected specimen 0-72 h

Specimen is stored at 2-8°C for up to 72 hours or proceed to RNA extraction.



3 RNA extraction ~45 min

Purified RNA is extracted from deactivated virus.



4 RT-qPCR ~1 h per primer set

Purified RNA is reverse transcribed to cDNA and amplified by qPCR.



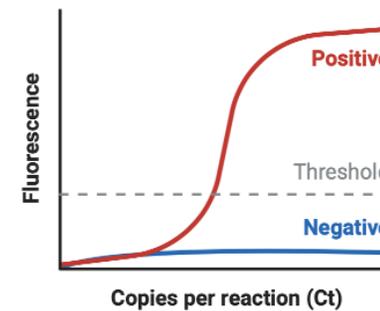
Primers and probes for screening

E_Foward: ACAGGTACGTTAATAGTTAATAGCGT	E gene First-line screening tool
E_Probe1: FAM-ACACTAGCCATCCTTACTGCGCTTCG-BBQ	
E_Reverse: ATATTGCAGCAGTACGCACACA	
RdRp_Foward: GTGARATGGTCATGTGTGGCGG	RdRp gene Confirmatory testing
RdRp_Probe1: FAM-CCAGGTGGWACRTCATCMGGTGATGC-BBQ	
RdRp_Probe2: FAM-CAGGTGGAACCTCATCAGGAGATGC-BBQ	
RdRp_Reverse: CARATGTTAAASACACTATTAGCATA	

* N gene testing is not further used because it is slightly less sensitive.

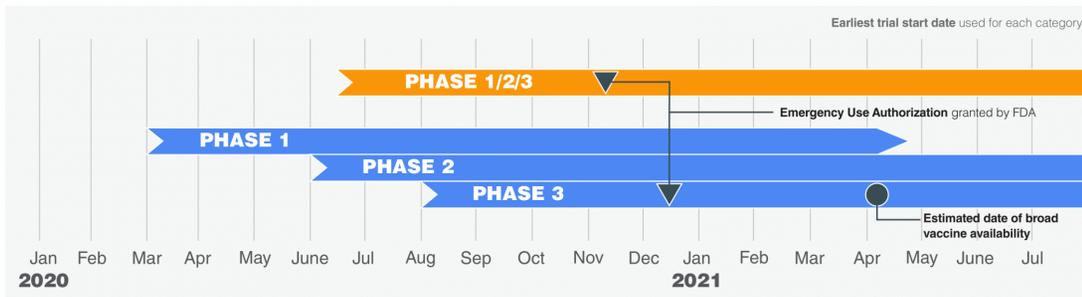
5 Test results real-time

Positive SARS-CoV2 patients cross the threshold line within 40.00 cycles (< 40.00 Ct).



Stunning pace of science in 2020, what's to come?

There are **18 programs** in phase 3 efficacy trials and at least **70 programs** in **Phase 1/2** safety testing.



Published Dec. 8, 2020

ANTIBODIES

Critical to treat early coronavirus infection.

1 MONOCLONAL

- + Simpler to make
- Potentially less effective

2 MONOCLONALS

- Harder to make
- + Potentially more effective

VACCINES

Critical to prevent COVID-19

EMERGING TECHNOLOGY

NUCLEIC ACID / mRNA

Injected gene or mRNA from the virus causes your cells to temporarily produce a foreign protein to kickstart an immune response.

VIRAL VECTOR

Harmless virus shuttles a coronavirus gene into your body to kickstart an immune response.

SUB-UNIT

Purified protein from the coronavirus is injected into the body to kickstart an immune response.

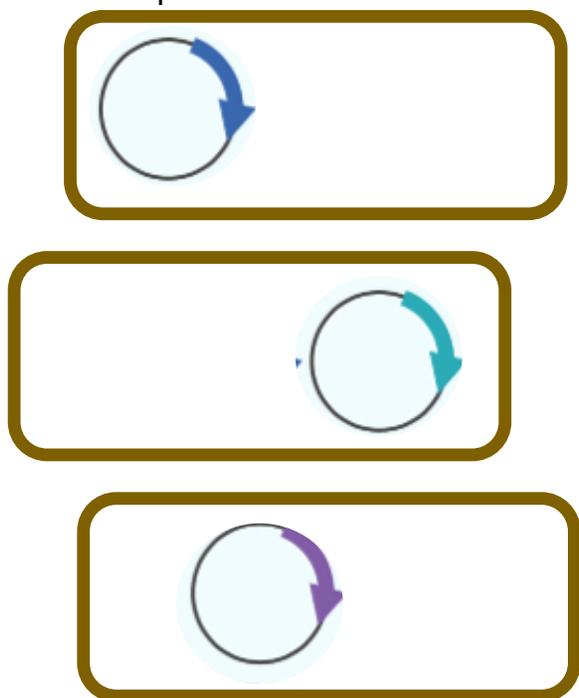
VIRUS

A weakened or killed coronavirus is injected into the body, to kickstart an immune response.

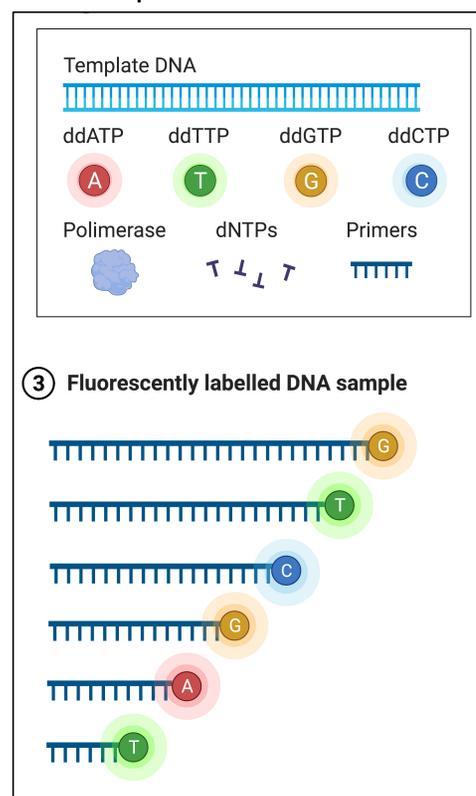
ESTABLISHED TECHNOLOGY

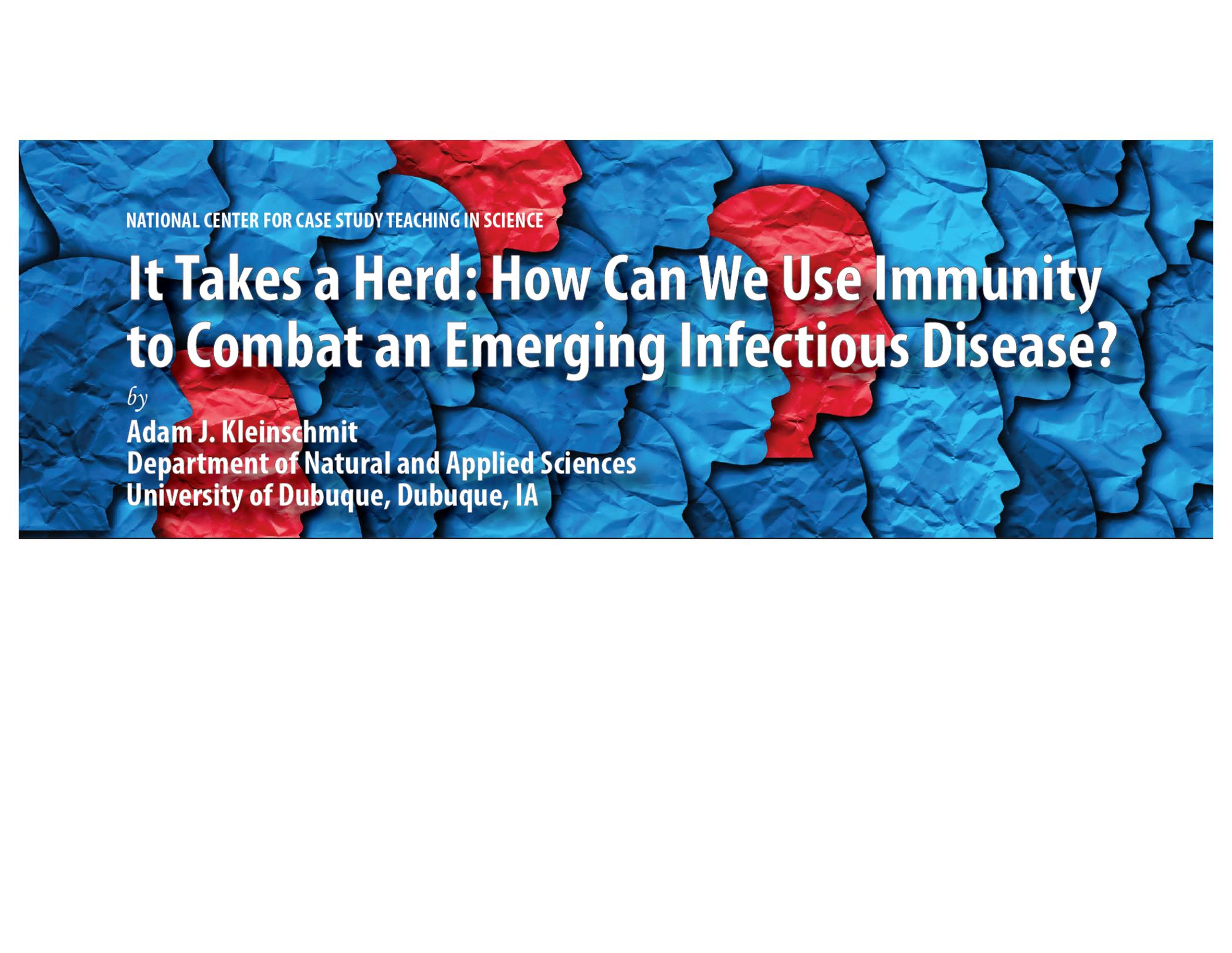
Today in lab, M1D4: Purify and sequence clone

Harvest scFv mutant plasmids from *E. coli*



Sequence each clone





NATIONAL CENTER FOR CASE STUDY TEACHING IN SCIENCE

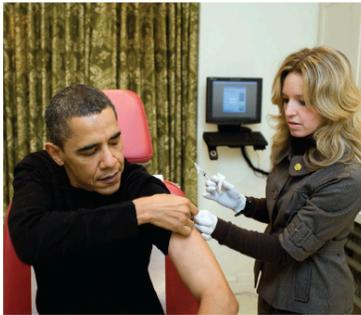
It Takes a Herd: How Can We Use Immunity to Combat an Emerging Infectious Disease?

by

Adam J. Kleinschmit
Department of Natural and Applied Sciences
University of Dubuque, Dubuque, IA

Immunity, ability to resist disease through preventing development of pathogen or by counteracting effects of the pathogen

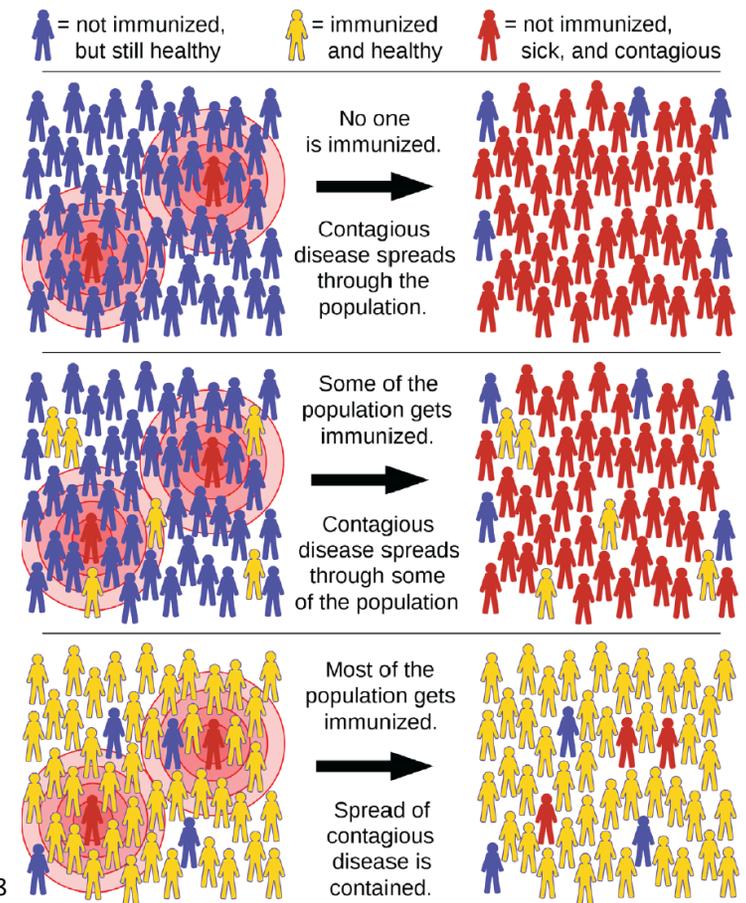
- Active immunity: host immune's system directly produces neutralizing antibodies
- Passive immunity: antibodies produced in a donor organism and transferred to recipient

	Natural	Artificial
Passive	<p>(A) Antibodies passed in breast milk or through placenta.</p> 	<p>(B) Antibodies harvested and transferred from another person, animal, or genetically engineered microbe.</p> 
Active	<p>(C) Illness and recovery.</p> 	<p>(D) Vaccination.</p> 

Herd immunity, indirect protection from infectious disease when a sufficient percentage of a population has become immune

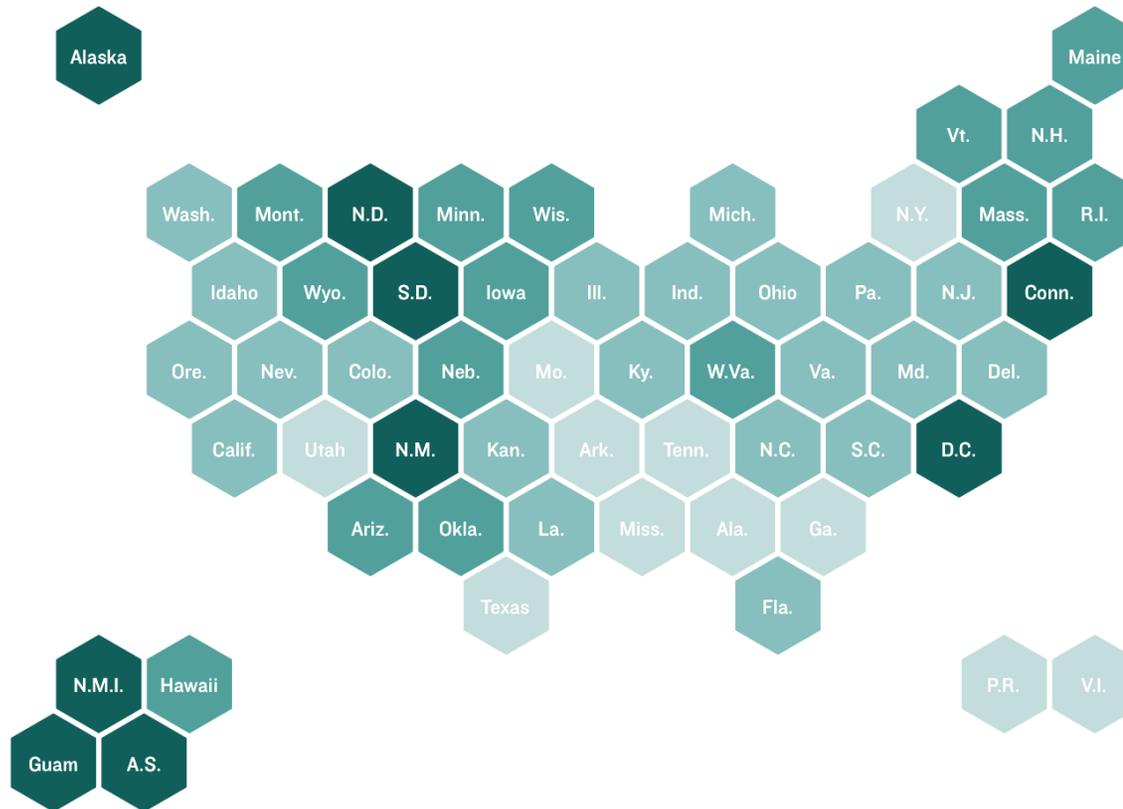
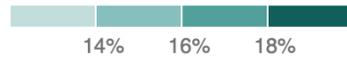
- In most cases, a diverse population can't all develop active immunity due to medical or physiological reasons
- The number of people that need immunity for herd protection depends on how transmittable the pathogen is
- Estimates for COVID-19 suggest ~70%* of the US population of 330million would need immunity to provide the benefit of herd immunity
- Massachusetts at 17.9% with one dose

*<https://www.mayoclinic.org/diseases-conditions/coronavirus/in-depth/herd-immunity-and-coronavirus/art-20486808>



Which States Have Vaccinated More Of Their Population?

Percentage of state's population that has received at least one dose of COVID-19 vaccine as of Feb. 28



Structure of our Discussion Today!

- Part 1, Individual Research (press releases/news articles):
 - (1) Reaching herd immunity,
 - (2) acquired active immunity through vaccination,
 - (3) naturally acquired immunity by infection
 - (4) acquired passive immunity through the creation of therapeutic monoclonal antibody cocktails.
- Part 2, Debrief Groups
 - Share expertise across groups
- Part 3, Whole Group Discussion
 - Discuss recent Massachusetts State policies that are designed to quickly reach herd immunity while providing vaccine rollout in an ethical way

Ethics discussion guidelines

- **We are accountable for our words and their impact.**
- Listen actively and with an ear to understanding others' views. Without interrupting.
 - Don't just think about what you are going to say while someone else is talking.
- Criticize ideas, not individuals.
 - We all can learn something from each other, even if your views don't necessarily align.
- Avoid blame, speculation, and inflammatory language.
- Avoid assumptions about any member of the class or generalizations about social groups.
 - Do not ask individuals to speak for their (perceived) social group.
- **Personal information that comes up in the conversation should be kept confidential.**
 - Take the message out of the classroom, not the messenger.