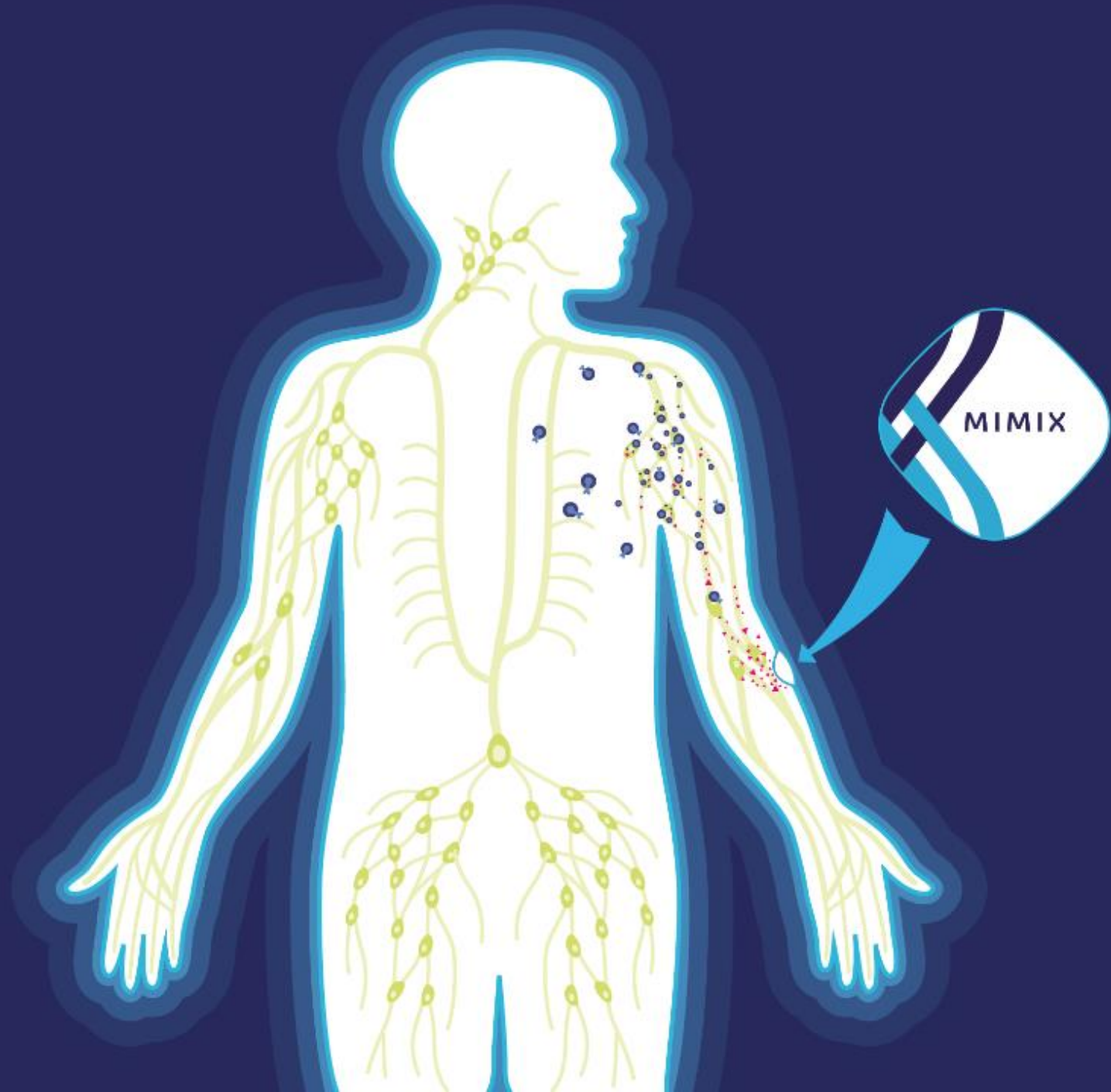


Introducing

 **MIMIX**TM

Radically Effective
Therapies in Oncology
and Infectious Disease



Developed by

 **VAXESS**
technologies

A decorative graphic on the left side of the slide, consisting of several thick, curved, overlapping bands in dark blue, light blue, and white, creating a sense of movement and depth.

Unlocking the Power of Our Immune System with MIMIX™

MIMIX [smart-release therapies](#) are engineered to release at the optimal rate in the skin or other epithelial tissue, the frontline of our body's immune system, to [stimulate powerful antibody and T-cell responses](#).

17 Employees	2 Partners	10 Patent Families	\$15M Non-dilutive funding	\$8.2M VC Funding	 
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breakthrough immune activation

Not only what we deliver, but where and for how long.*



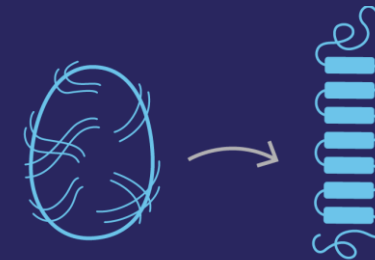
next-gen delivery

MIMIX enables sustained release of biologics into tumors or epithelial tissue



advanced materials

New silk-based material combines stabilization with tunable release to enable the first sustained delivery of biologics

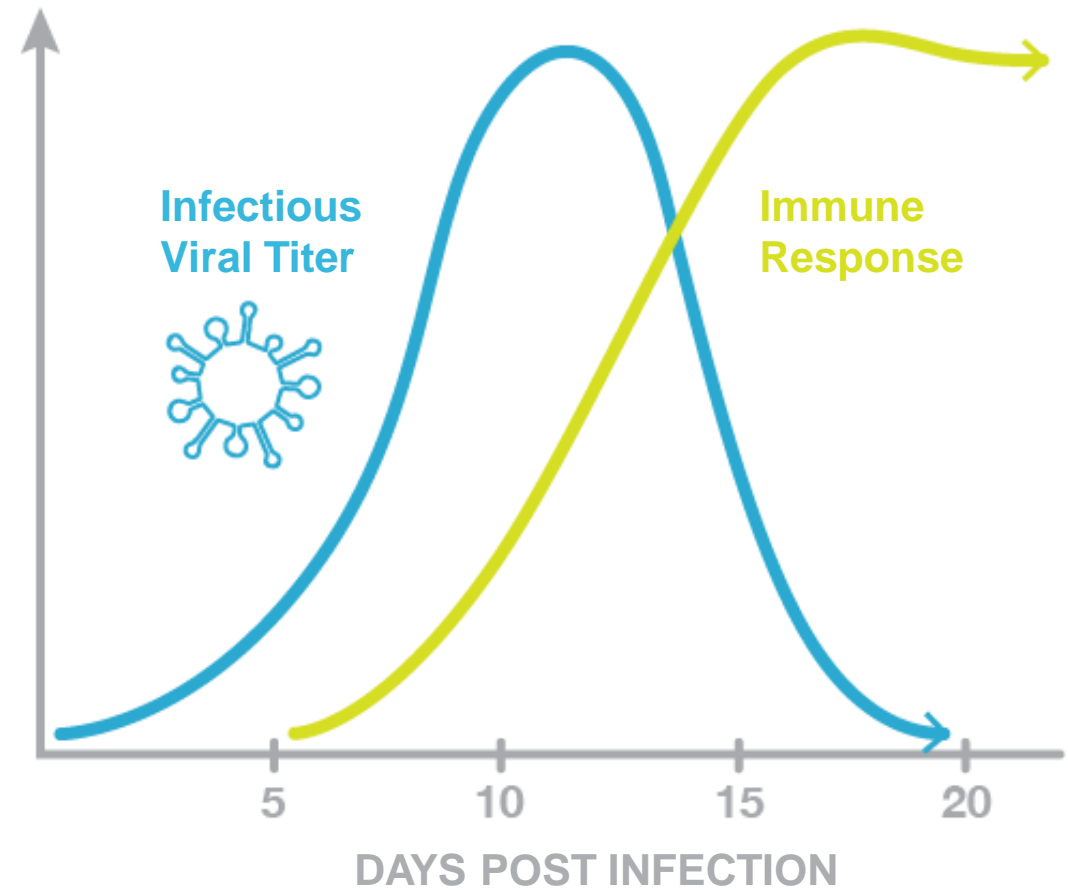


*Wittrup et al. (2019) What, Why, Where, and When: Bringing Timing to Immuno-Oncology. *Trends in Immunology*

THE INSPIRATION

Viruses enter epithelial tissue (ie. skin) and persist in the host over 2+ weeks.

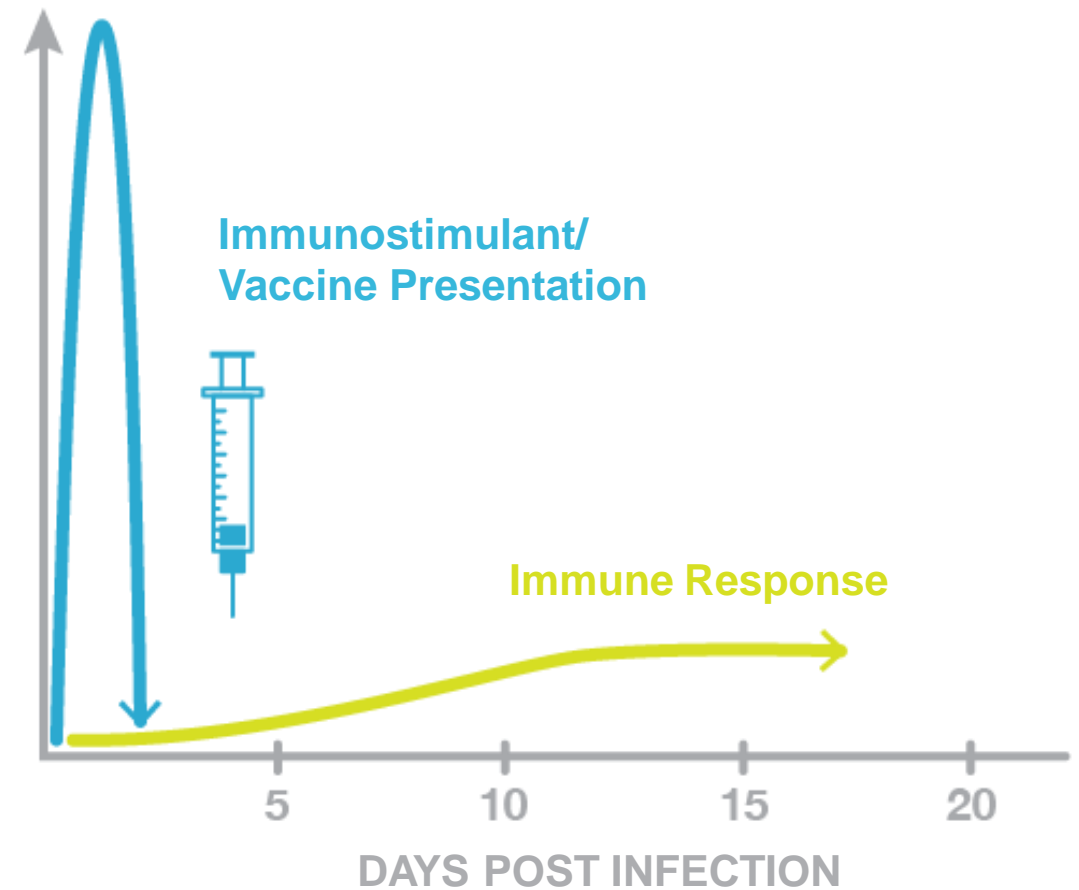
The body generates a strong and durable immune response.



THE DISCONNECT

Vaccines are administered into veins or muscles and cleared from the body in <2 days.

The body doesn't have time to mount a strong response.



It isn't just **WHAT** we deliver,
but **WHERE** and for **HOW**
LONG to enable breakthrough
immune responses.

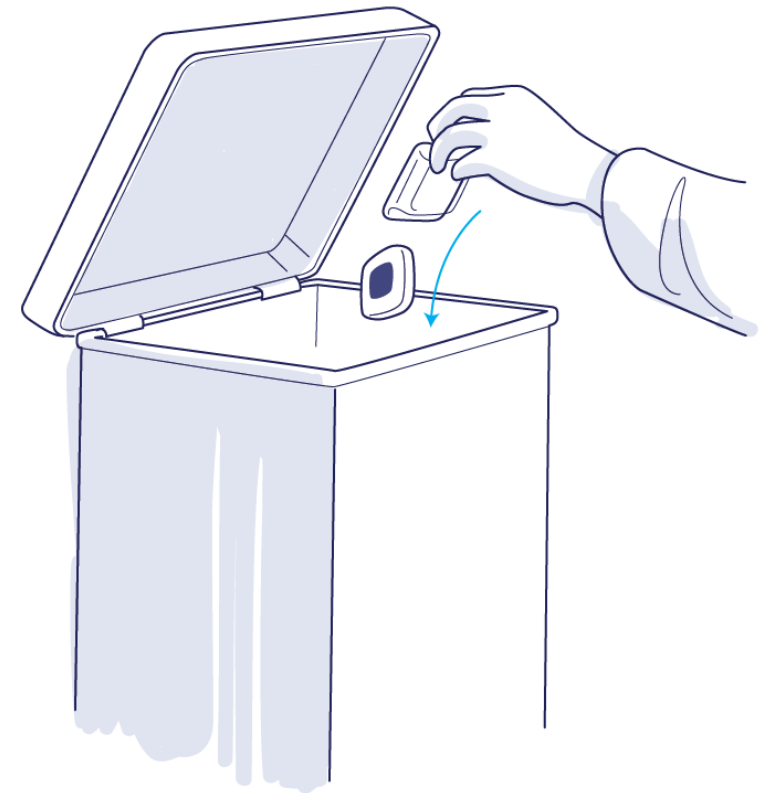
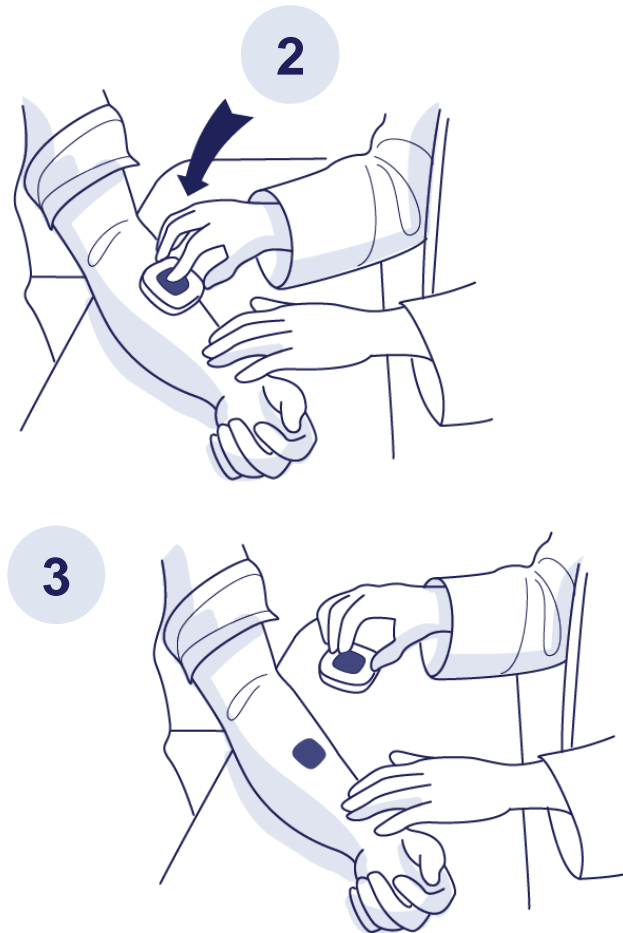
We call it
Infection Mimicry.

Introducing MIMIX smart-release therapies that mimic infections.

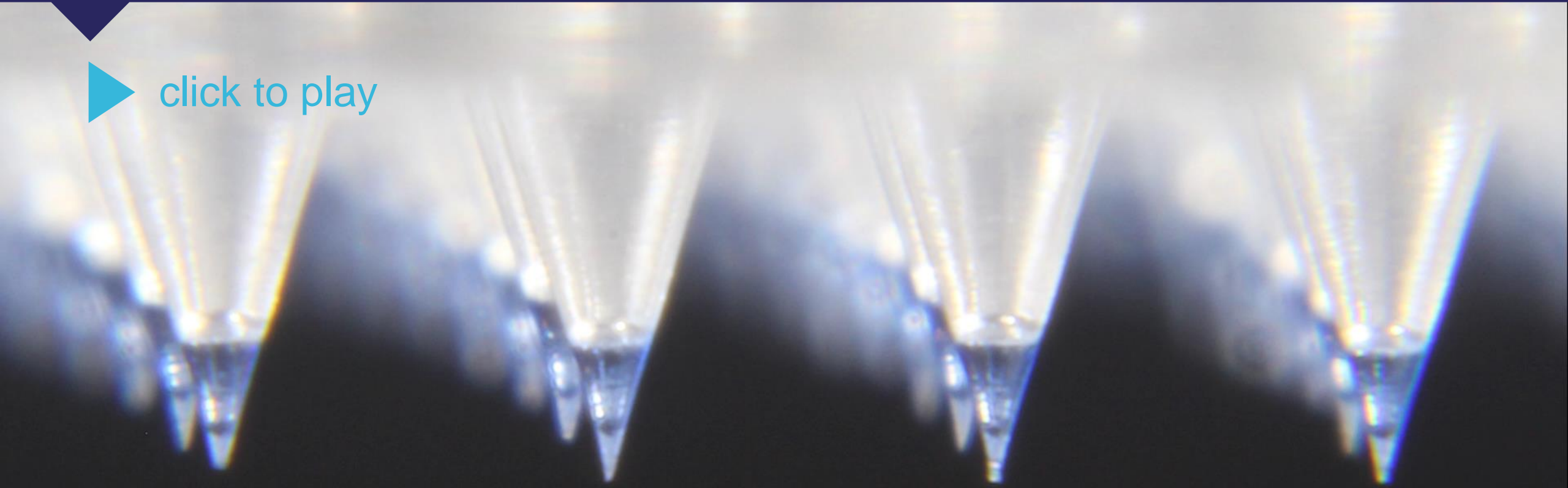
With just minutes of wear-time, MIMIX enables **weeks of sustained release** of treatments into tumors, skin, or other tissue.



How it works

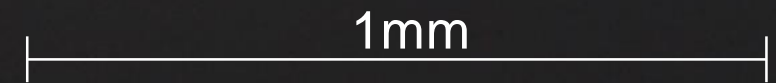


▶ click to play

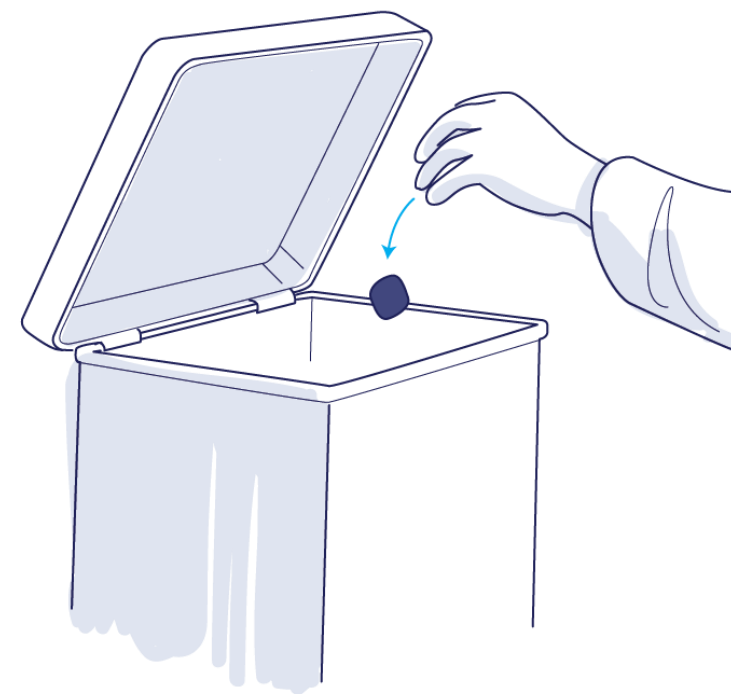
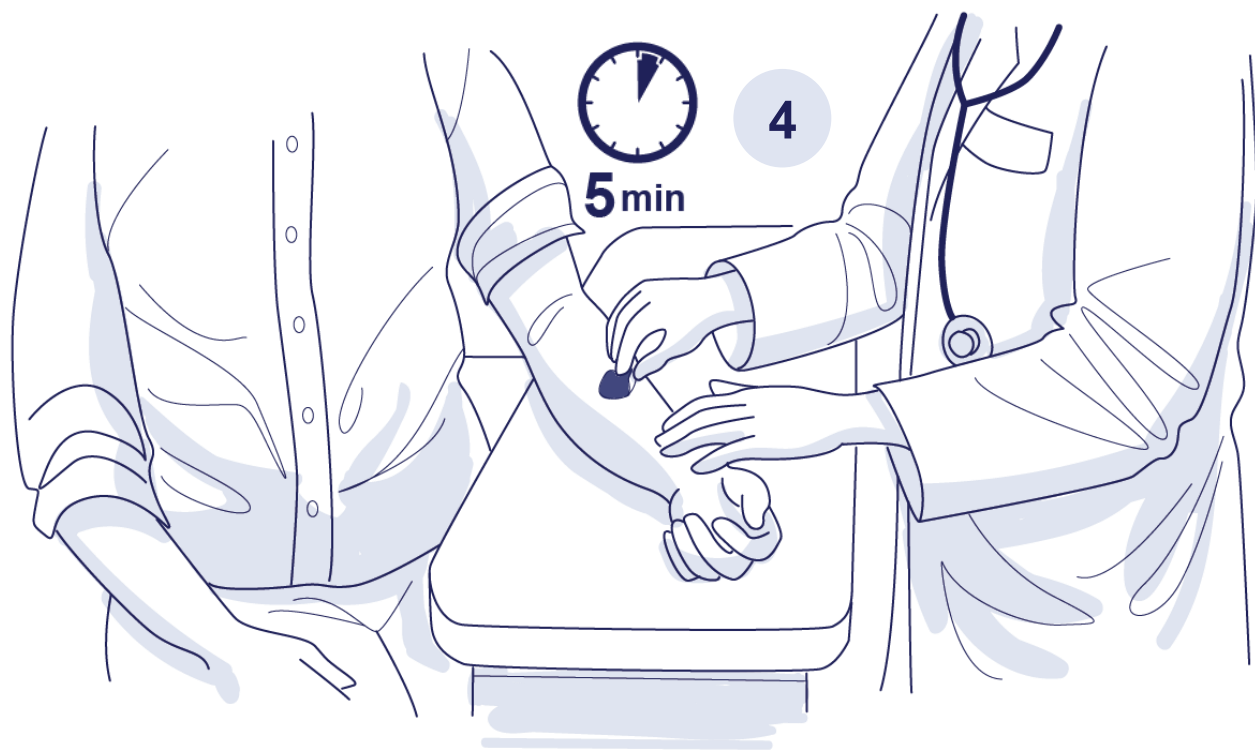


MIMIX™ smart release needles dissolve rapidly upon entering the skin, implanting the slow-release tip and enabling therapeutic release from minutes to months.

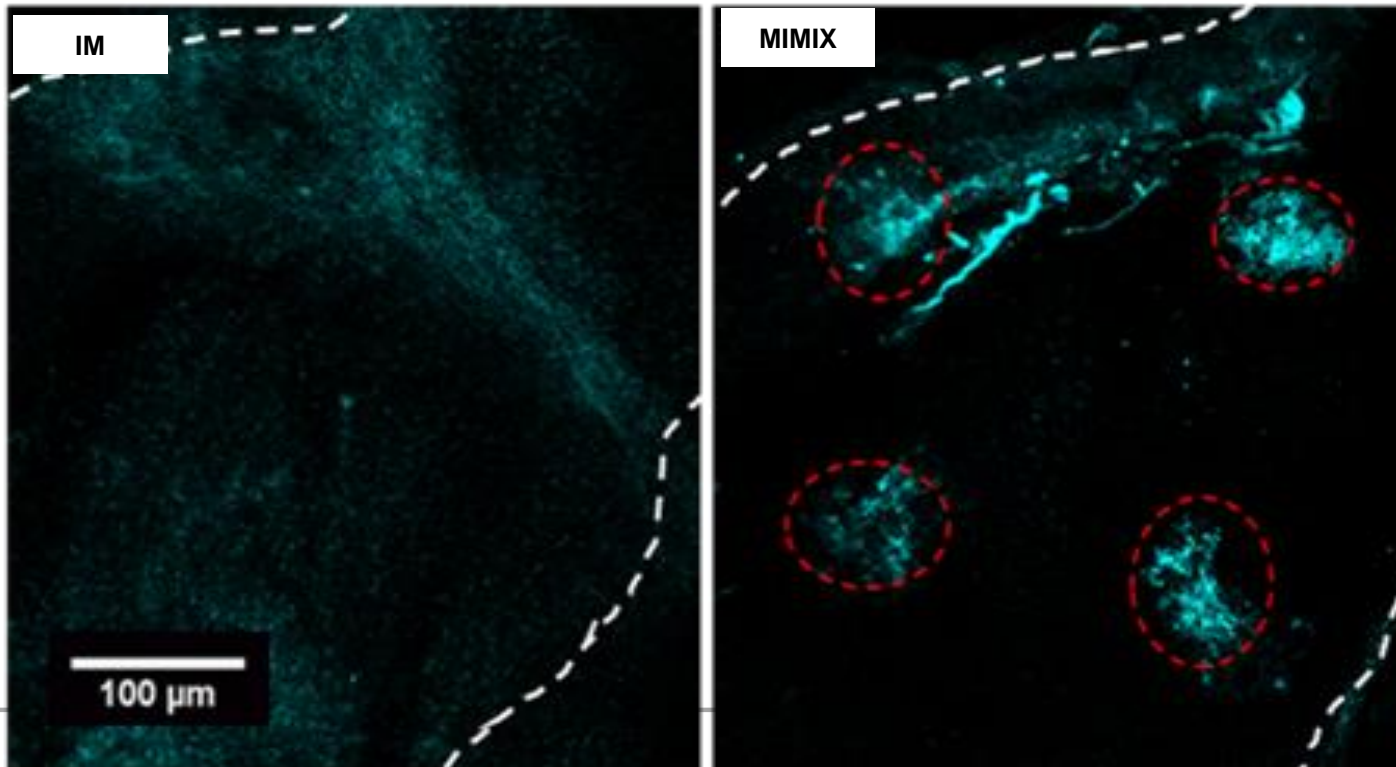
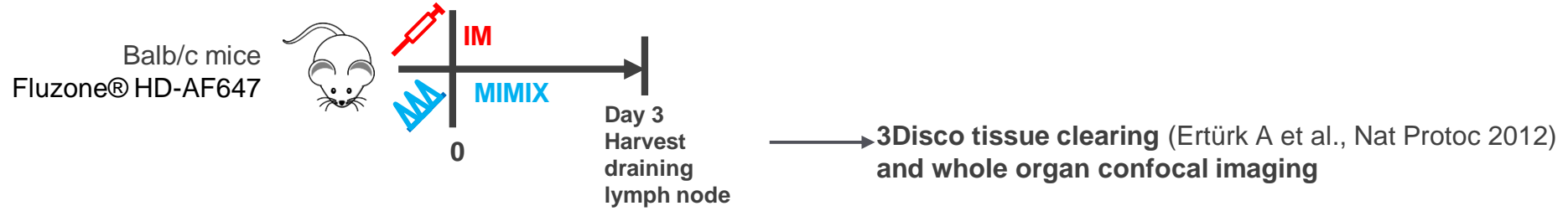
[Click Here for YouTube Video](#)



Dispose of used patches; no sharps.



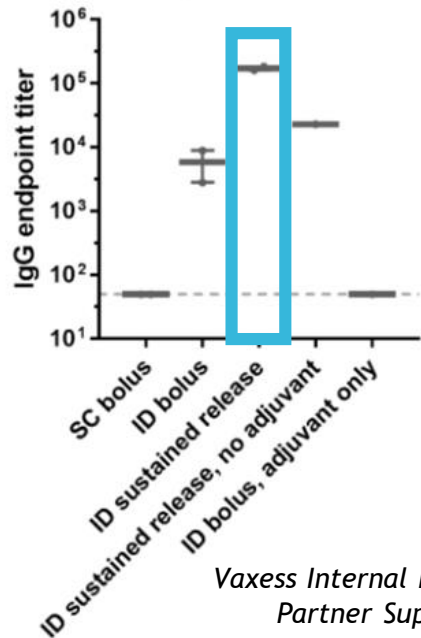
MIMIX enhances antigen retention in draining lymph nodes



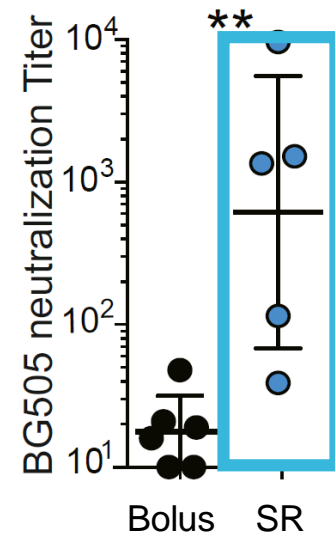
Vaccine -- Cyan
White dotted line – Lymph node (LN) organ outline
Red dotted circles -- Pattern of follicular dendritic cells in LN

Infection mimicry via sustained release drives radical shifts in **quantity** and **quality** of response

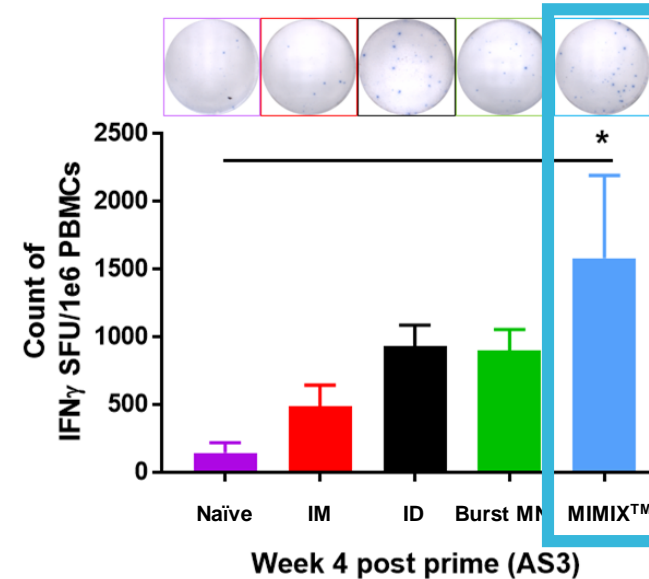
1000x enhancement in antibody response
(HSV Therapeutic Vaccine)



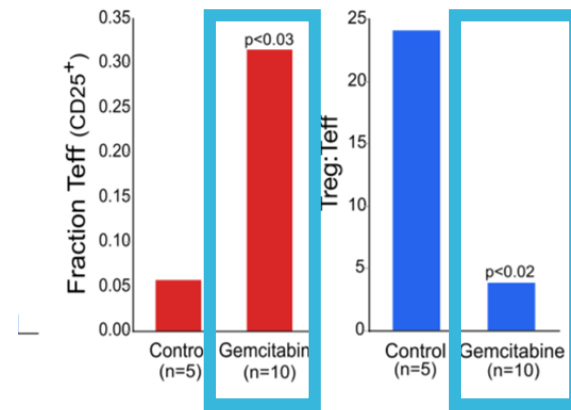
Shift to neutralizing antibodies
(HIV Vaccine (Primates))



4x enhancement in T-cell responses
(Influenza Vaccine)

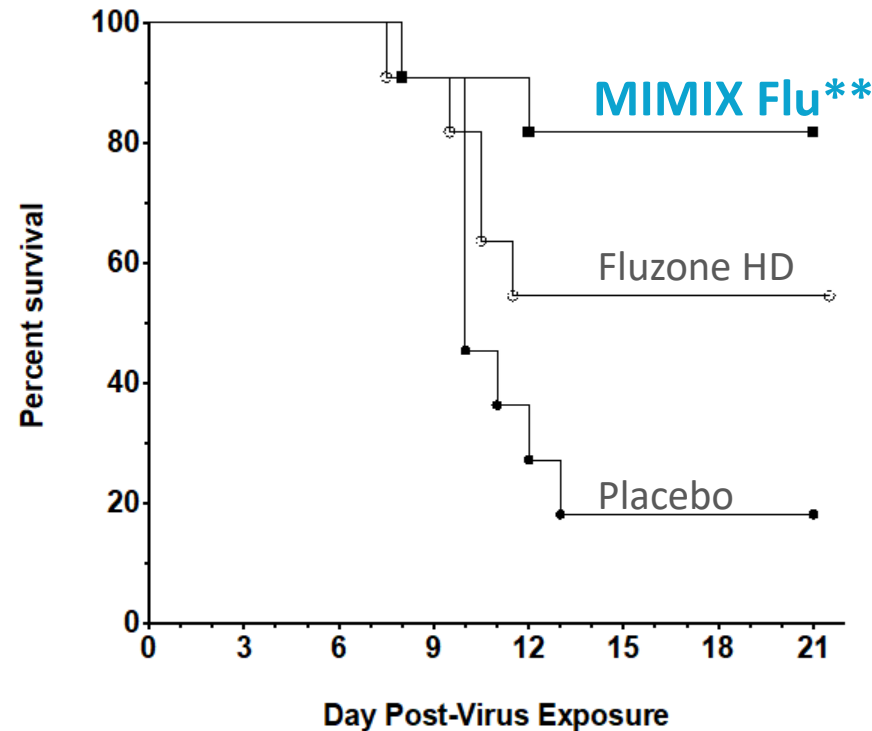
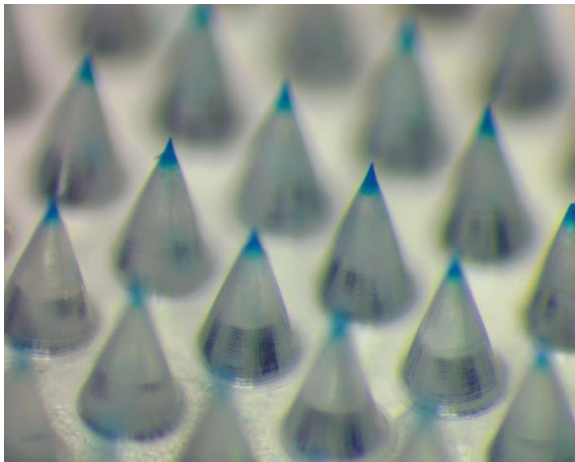
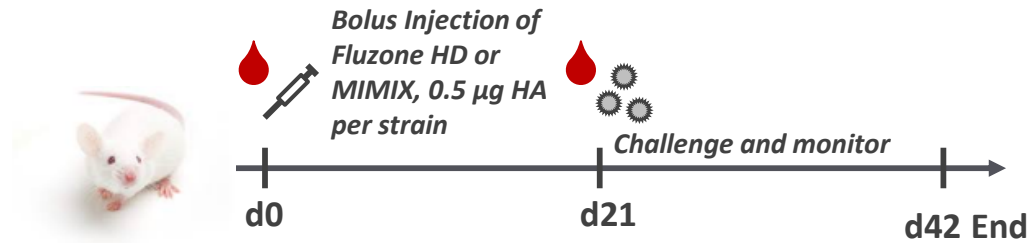


Enhanced T-cell activation, reduced Treg in tumor environment
(Bladder Cancer)



MIMIX-Flu Enhances Protective Efficacy Against Drifted Strains of Influenza

Immunization with 2018-19 seasonal influenza vaccine. Challenge with 2009 H1N1 strain (A/California/04/2009).

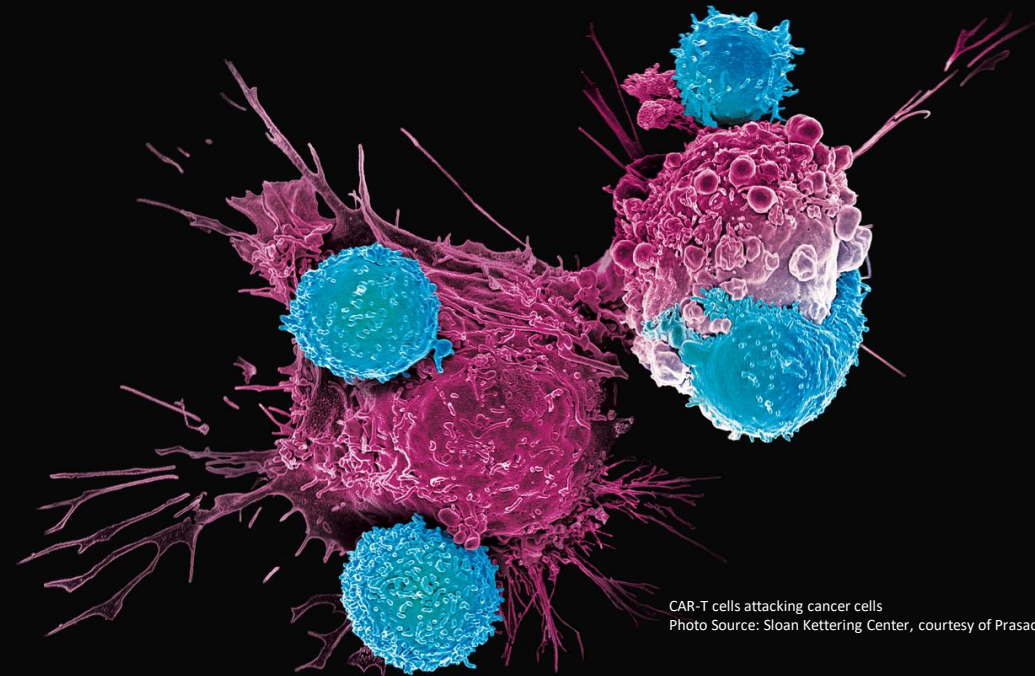


n=21 per group (11 for survival monitoring, 10 for analysis of virus titers and cellular immunity)

(**p < 0.01)

Vaxess is using this approach to **train the immune system at one tumor to attack cancer that has spread throughout the body.**

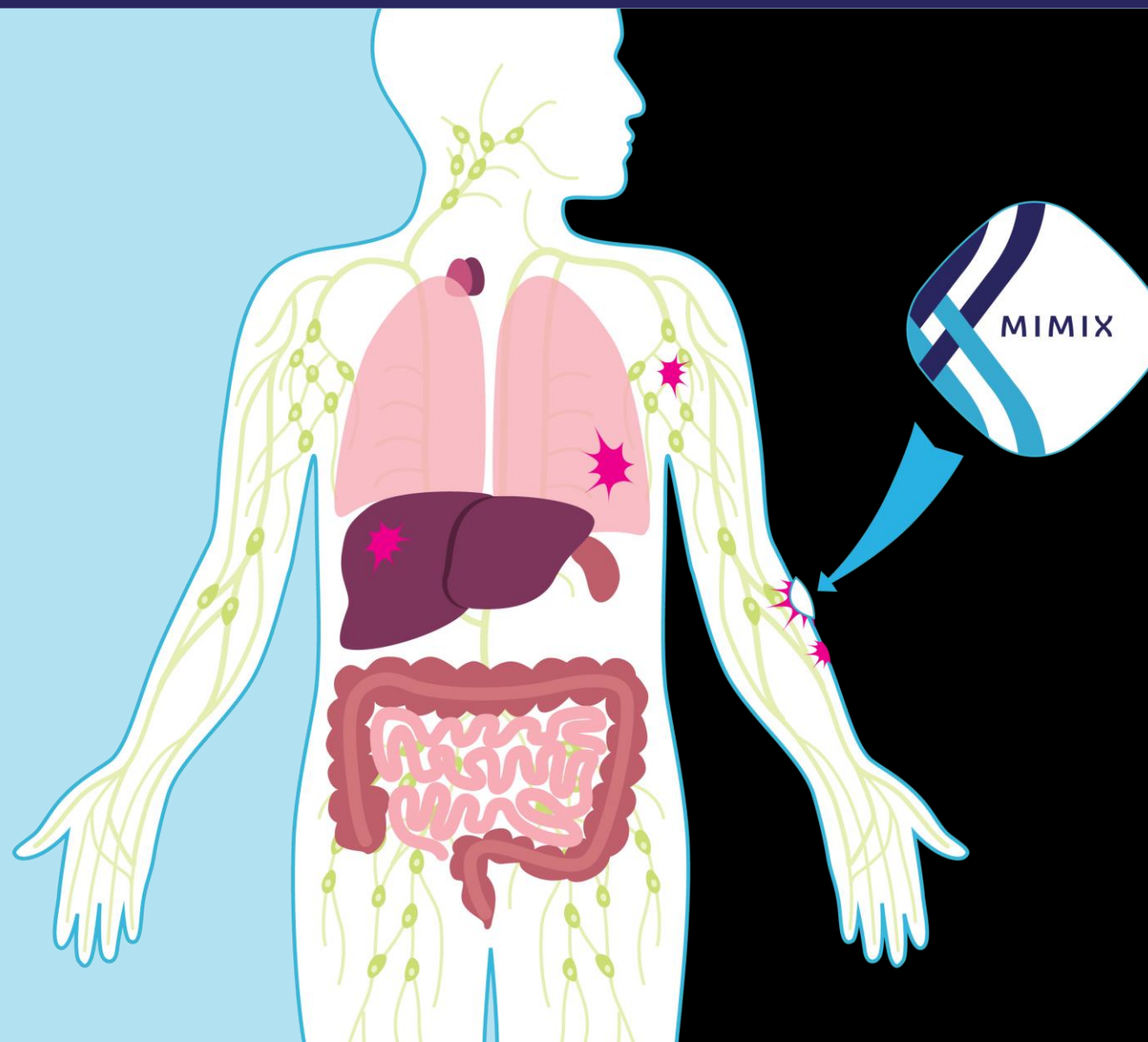
- 1. The right antigen:** The patient's own lysed cancer cells serve as vaccine "antigen"
- 2. In the right place:** T-cell maturation occurs in the lymph nodes, MIMIX drives strong lymph node trafficking
- 3. For the right duration:** T-cell maturation takes time. MIMIX provides 2+ weeks of antigen exposure
- 4. In the right combination:** Immune activation alongside cytokines and checkpoint inhibition
- 5. With less toxicity:** Slow, intratumoral release reduces off-target effects for IL-2 and others



CAR-T cells attacking cancer cells
Photo Source: Sloan Kettering Center, courtesy of Prasad Adusumilli

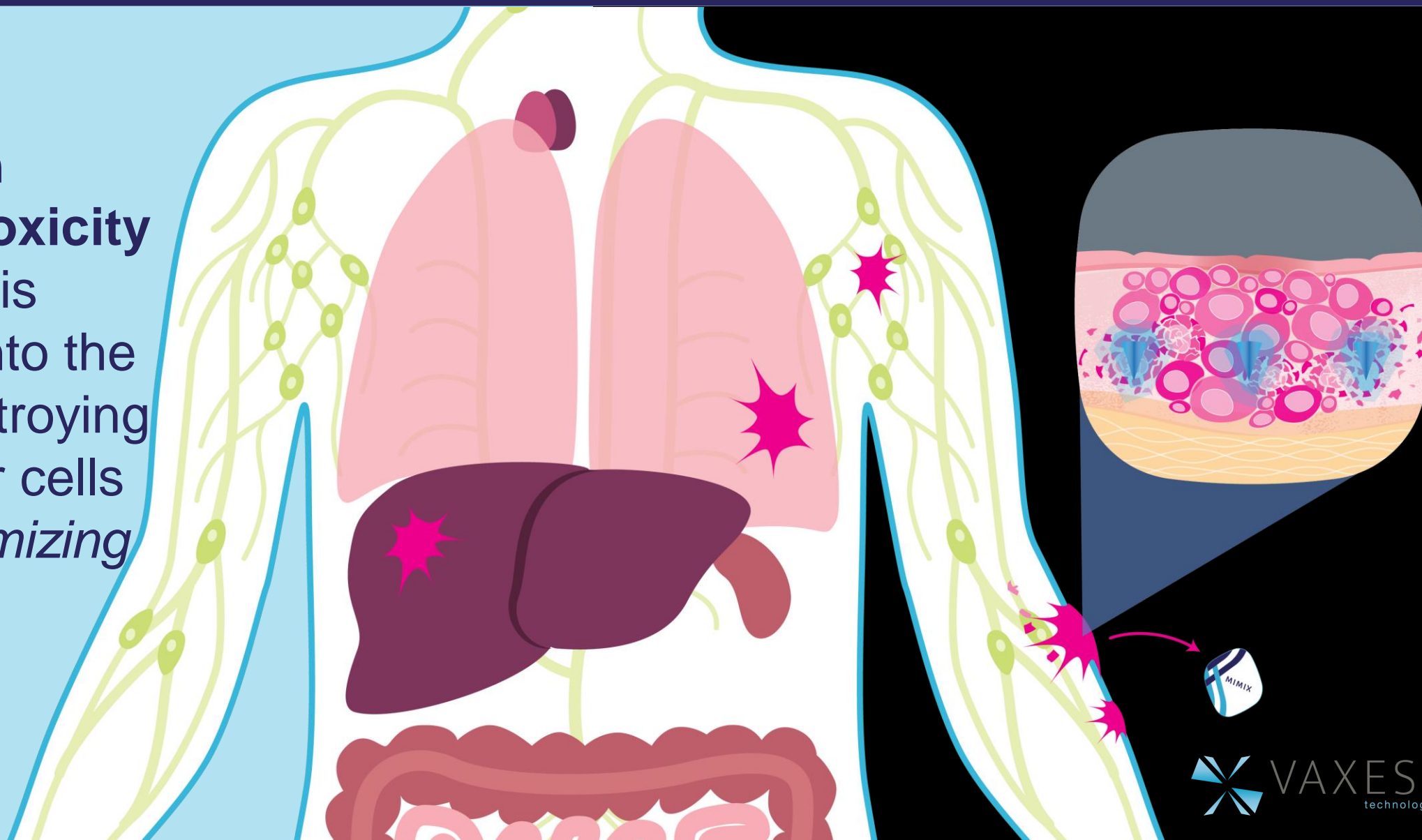
Activating the immune system to fight cancer with MIMIX

The smart release patch is applied to the a skin lesion.



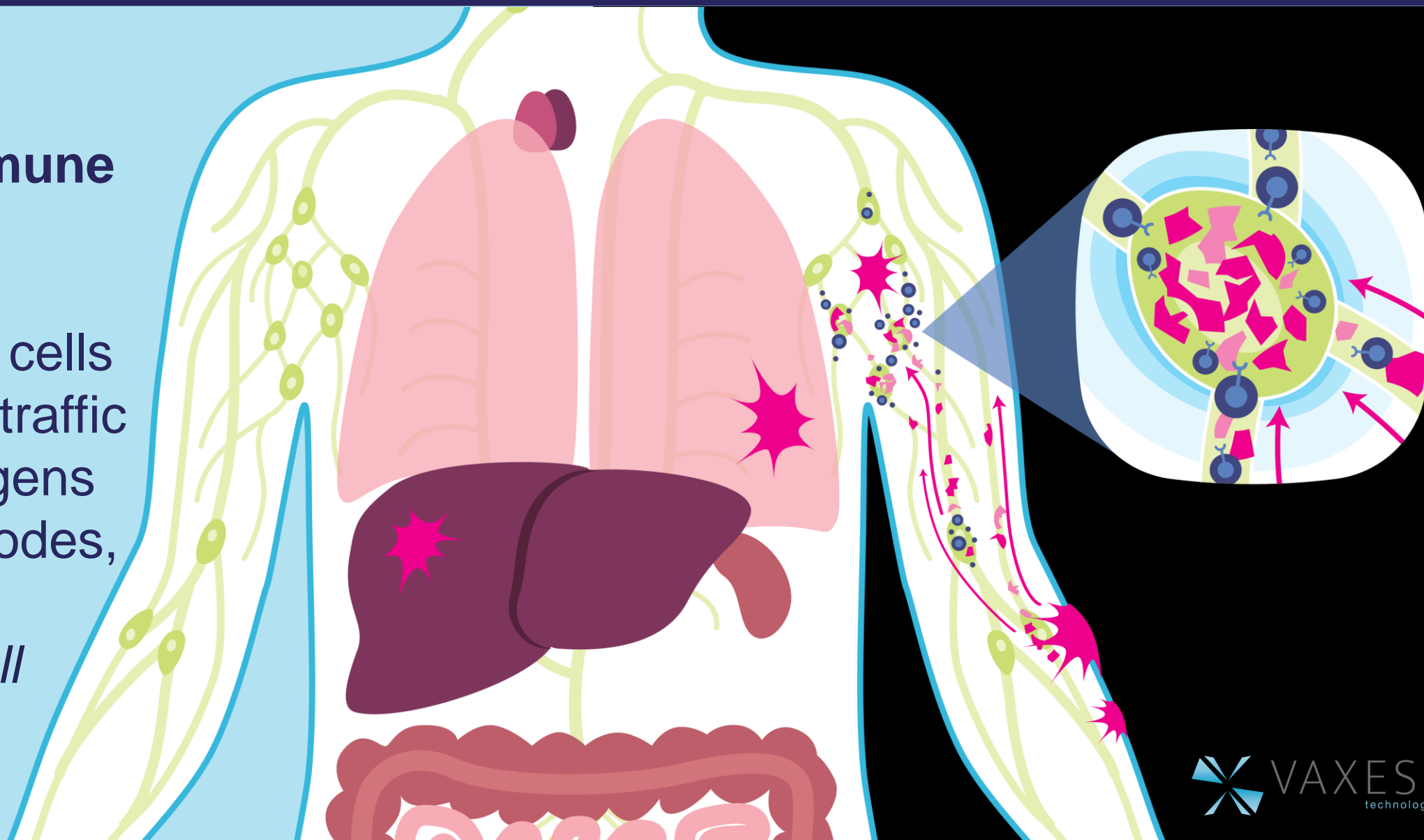
Cell Death Without Toxicity

The agent is released into the tumor, destroying local tumor cells *while minimizing systemic toxicity*



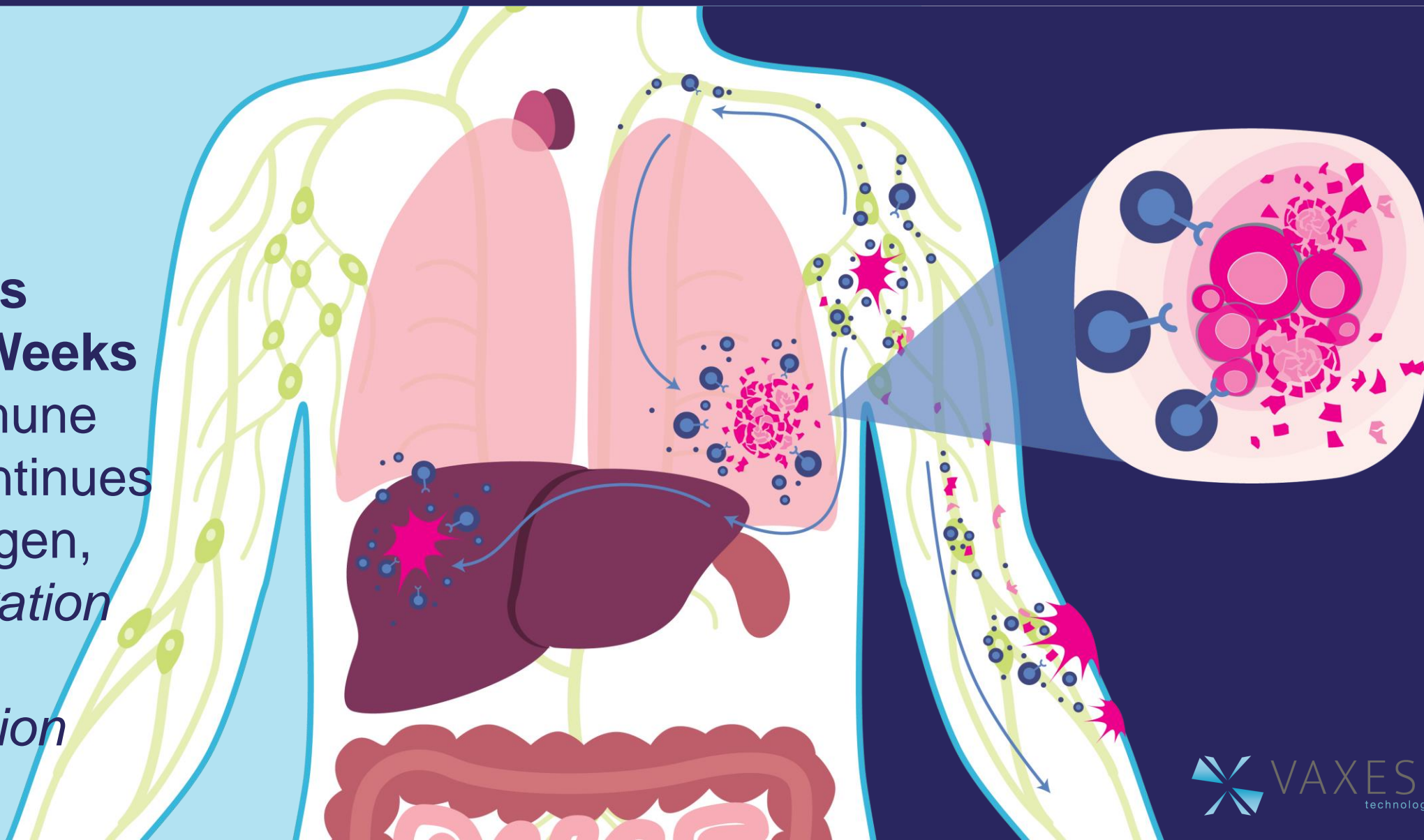
MIMIX Immune Activation

Antigen-presenting cells in the skin traffic tumor antigens to lymph nodes, *triggering potent t-cell responses*



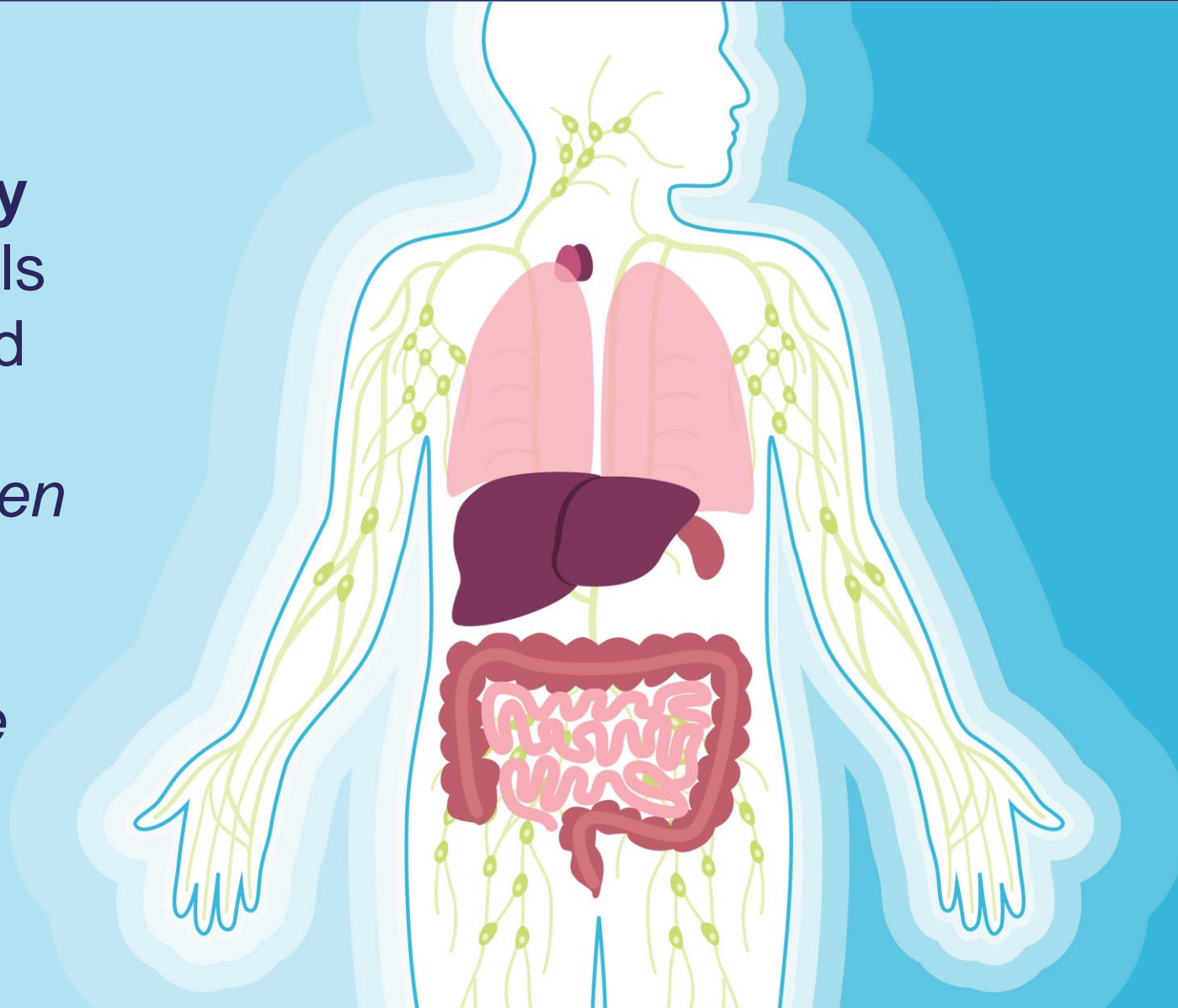
**Building
Powerful
Responses
Over 2-3 Weeks**

As the immune system continues to see antigen, *T-cell activation and differentiation increases*

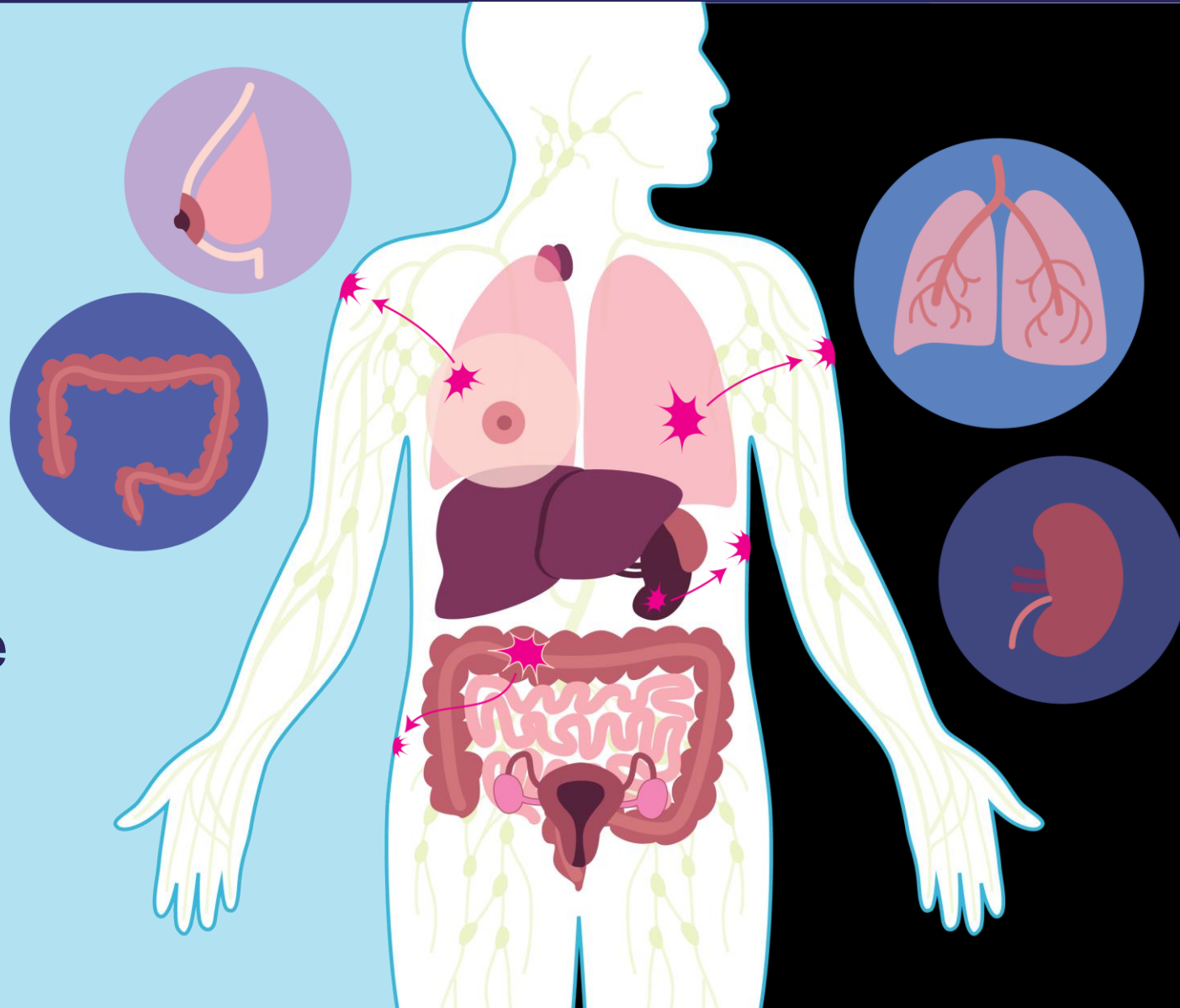


MIMIX Memory

The cancer cells are cleared and *the immune system has been trained, potentially enabling future activation if cancer returns*



**Beyond
Melanoma**
Many other
tumors
metastasize to
the skin,
providing suitable
“antigen” for
MIMIX immune
activation



MIMIX-MR: Translating to impact global health

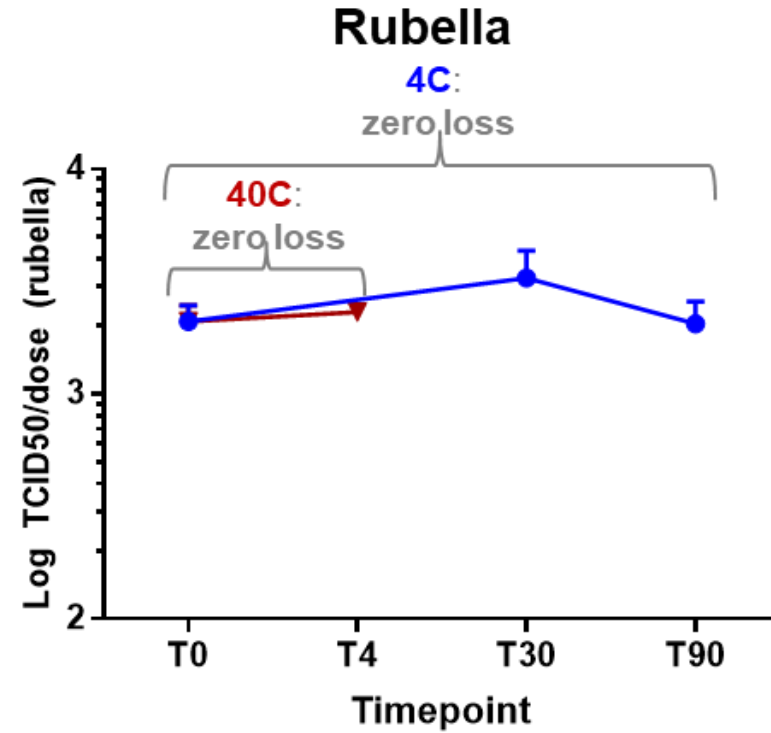
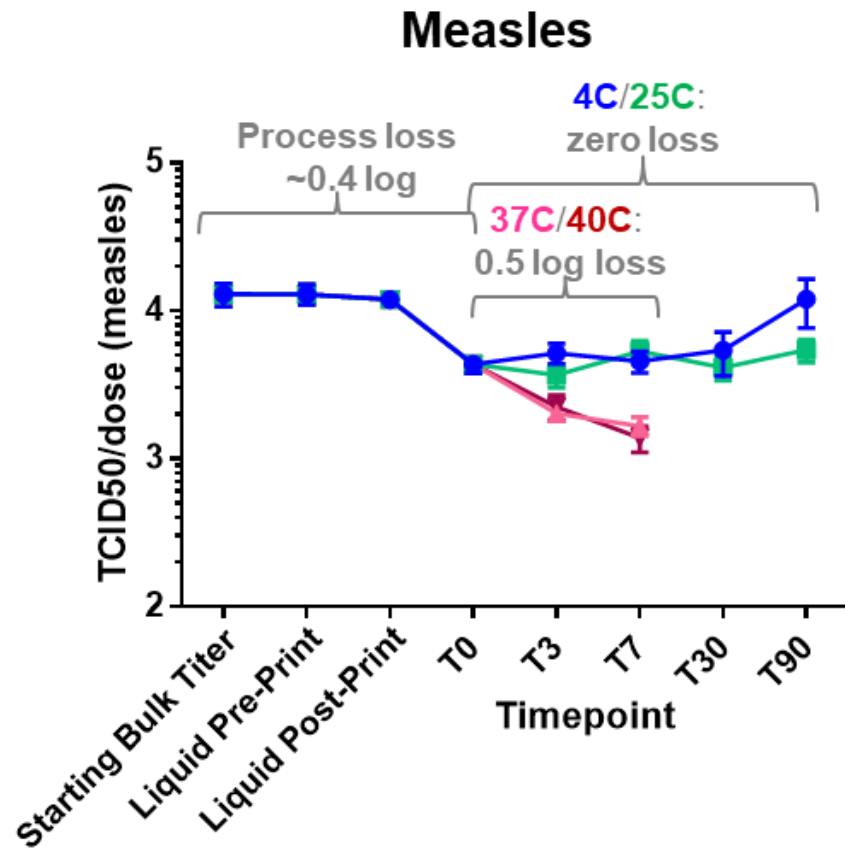
BILL & MELINDA
GATES *foundation*



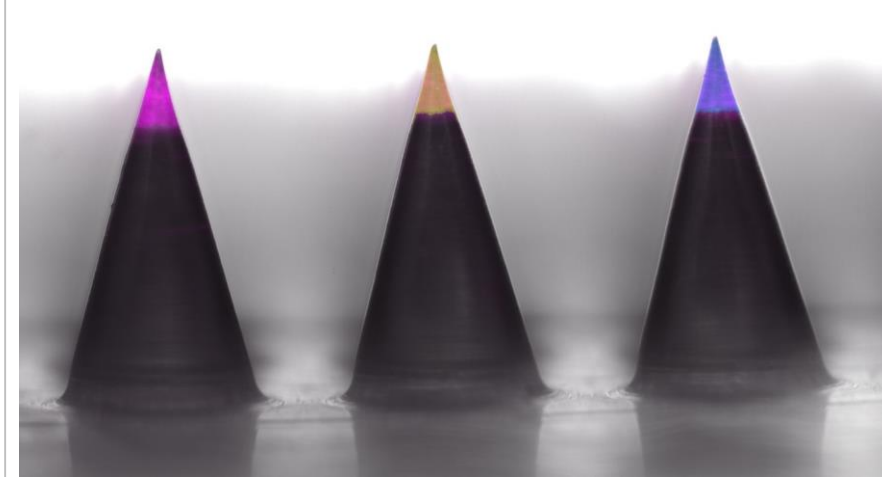
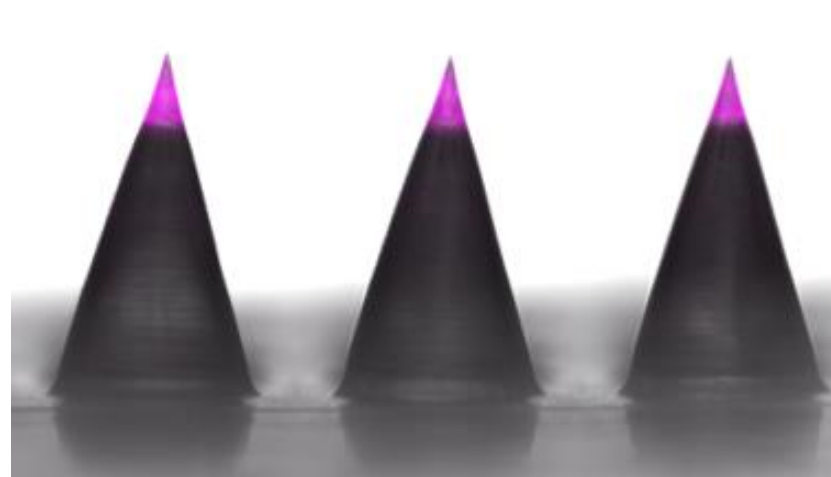
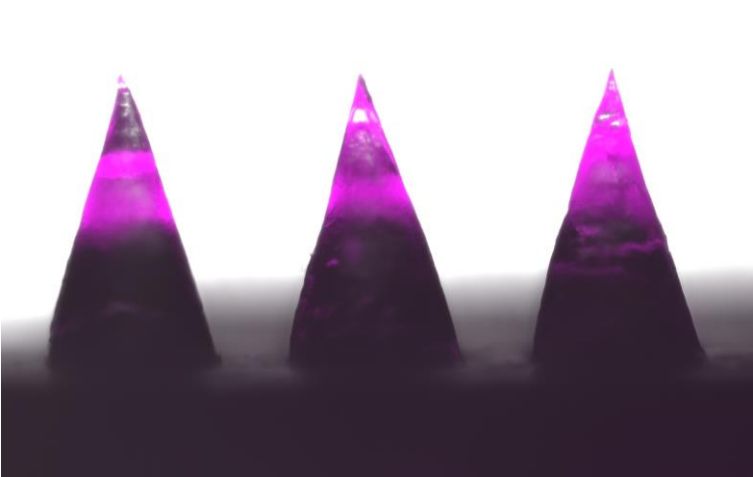
Measles-rubella (MR) microneedle abbreviated Target Product Profile

Variable	Minimum Target
Target Population	Routine: Infants 9-18 months of age, Campaign: >9 mo age, children, adolescents, and adults at risk
Device safety	<ul style="list-style-type: none"> • May tolerate increase in local reactogenicity • Waste that is considered infectious, sharps waste that is disposed of according to instructions for use, WHO level and relevant local requirements
Target Population	Routine: Infants 9-18 months of age, Campaign: >9 mo age, children, adolescents, and adults at risk
Vaccine Volume (cm ³ /dose)	One dose per patch; primary container <1 cm ³ /dose; secondary packaging <5 cm ³ /dose
Stability / Shelf Life	Shelf life: 24 months at 2- 8° C Stability: Not freeze sensitive
Target Procurement Price	≤US\$1.40
Manufacturing Capacities	Up to 50 million patches per year

Measles Rubella MN Stability – 1 week at 40°C and 3 Months at 4°C

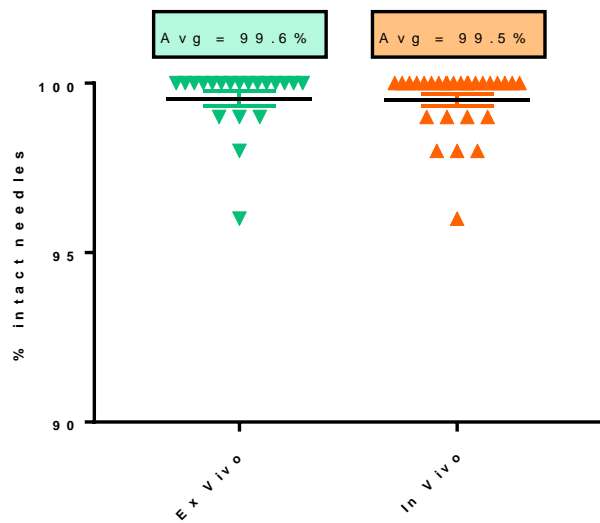


Optimizing for Consistency, Delivery Efficiency, and Flexibility

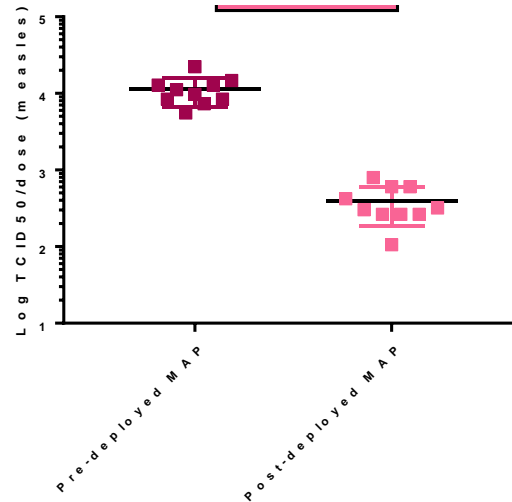


Manufacturing Improvements Drove High Quality Patches with Consistent Delivery

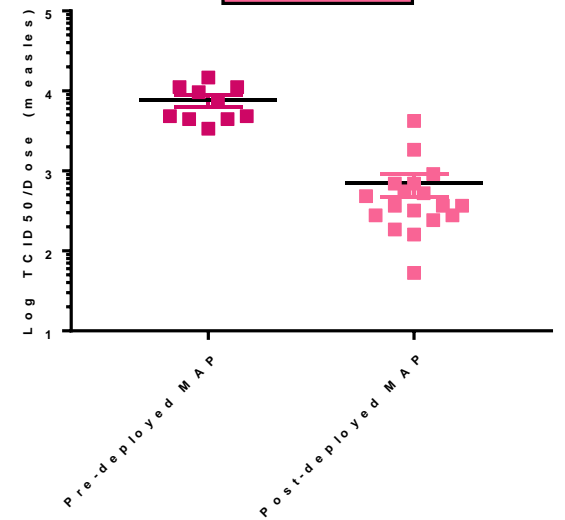
99% needle demolding Efficiency*



97% dose delivered ex vivo



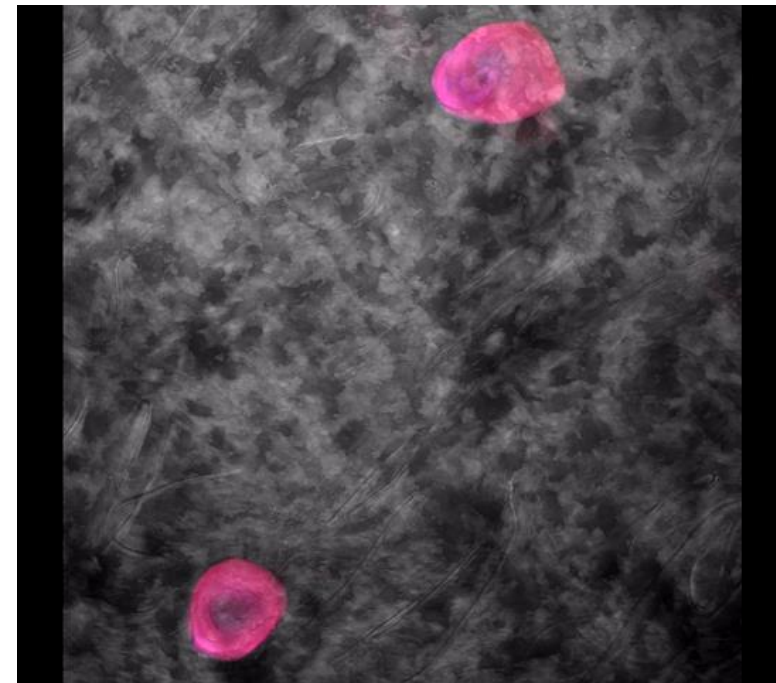
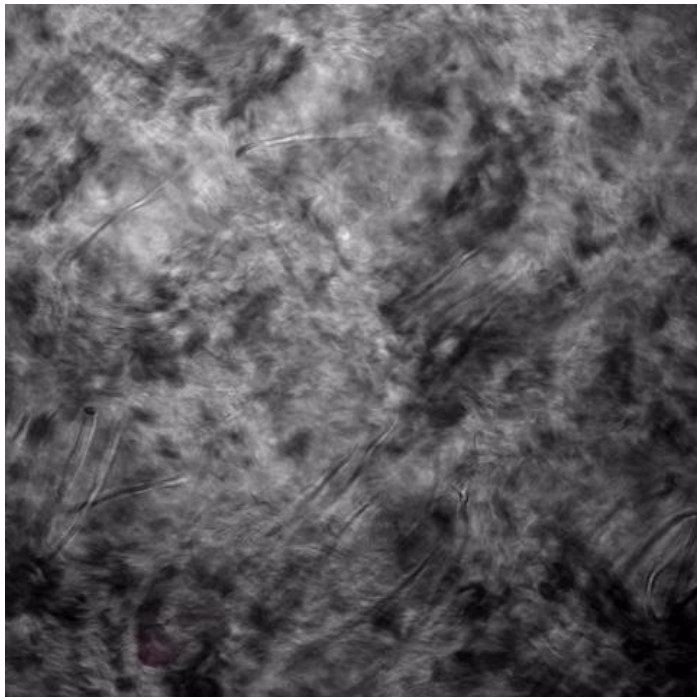
91% dose delivered in vivo



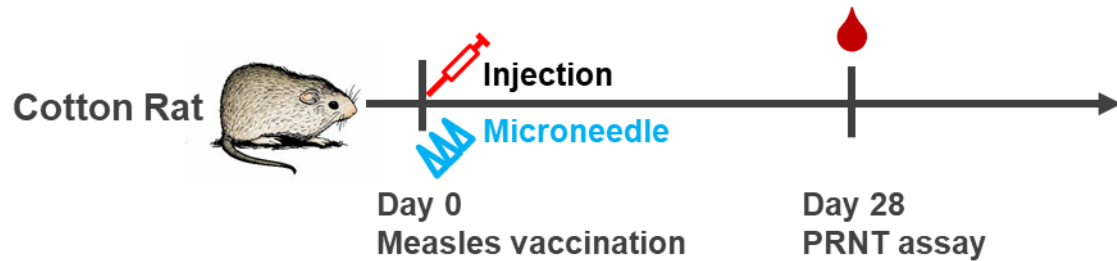
*Demolding efficiency (% intact needles)

Validating Deployment Depth Using Confocal Microscopy

Tips were imaged in the skin using confocal microscopy to validate implantation, tip morphology and deployment depth



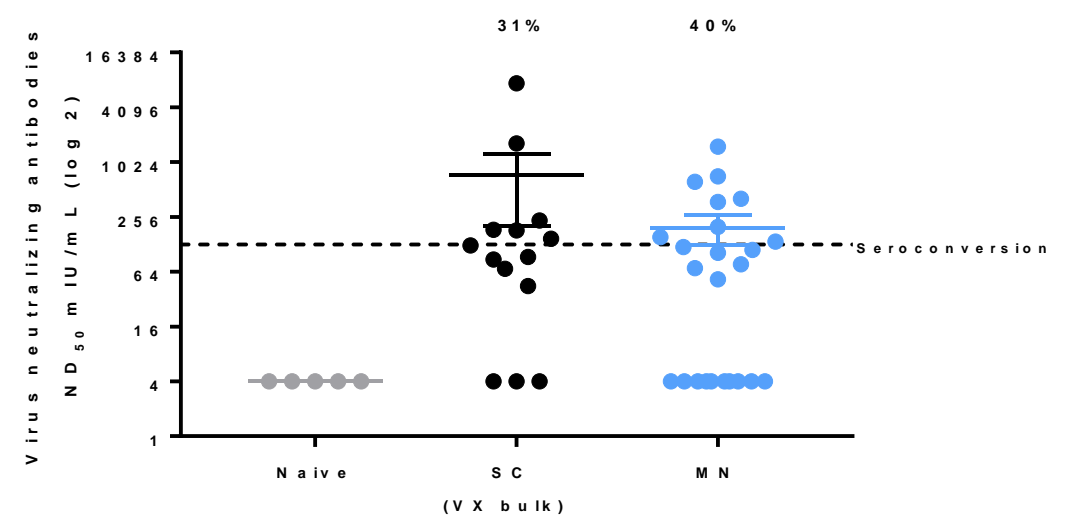
Measles Microarray Patch Vaccine In Vivo Study



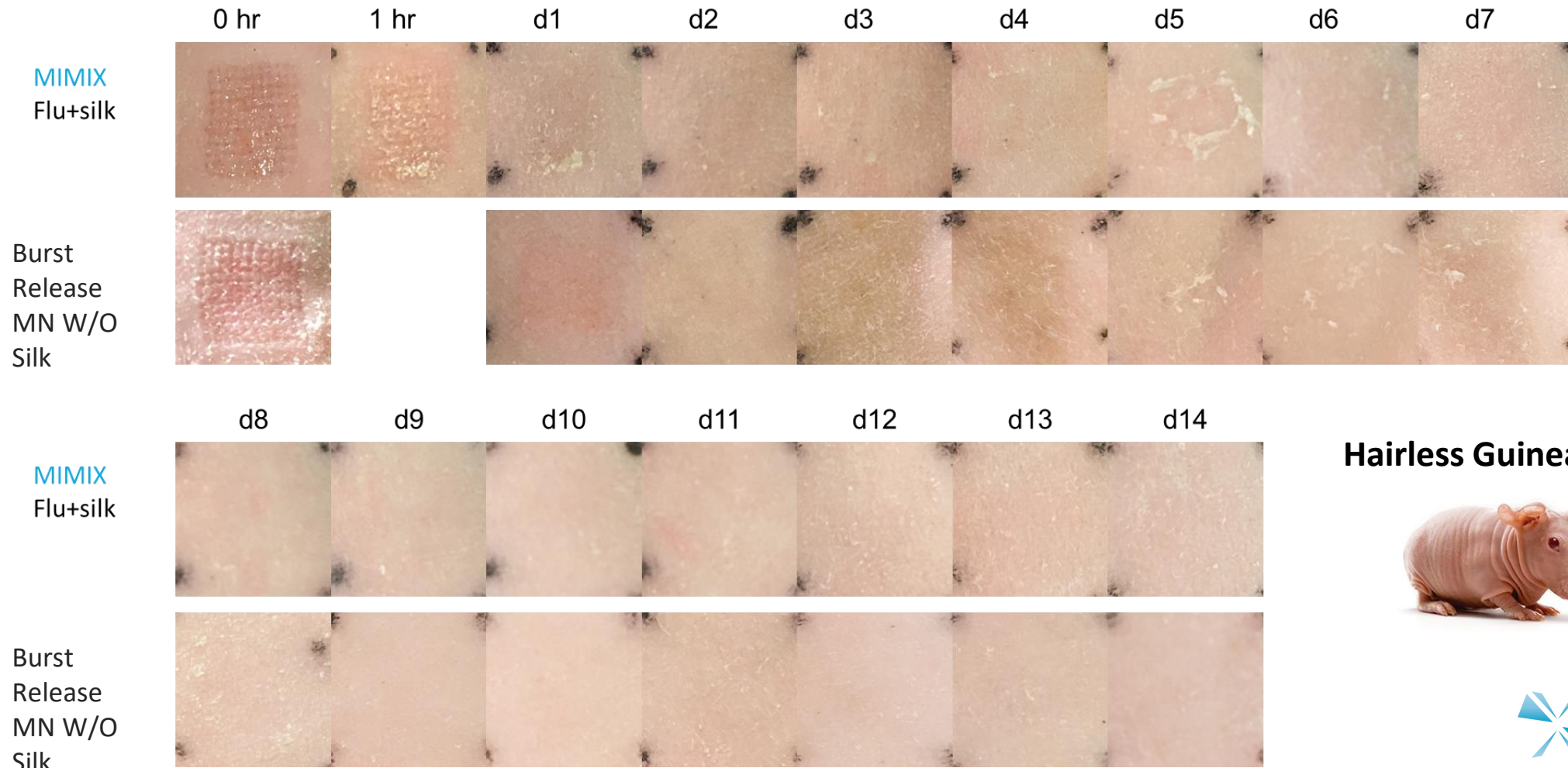
Results: MNs had comparable seroconversion to traditional SC injection



Neutralizing Ab Responses to Measles Vaccine (Day 28)



Evaluating Skin Responses in Hairless Guinea Pigs



Hairless Guinea Pig

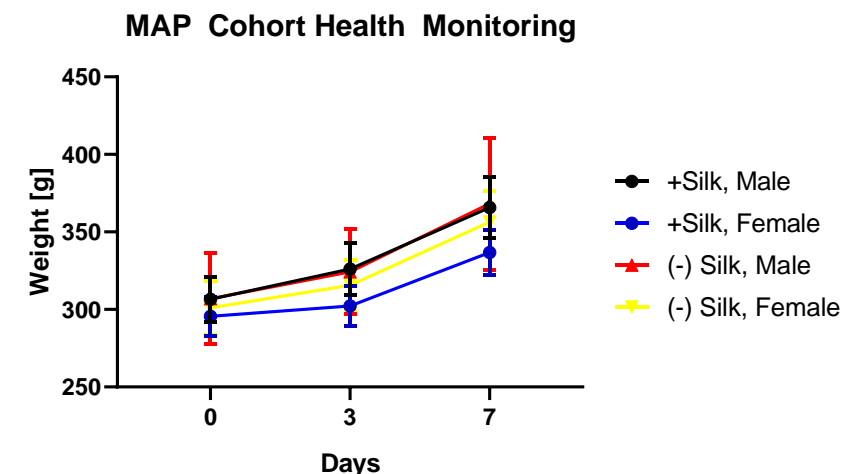
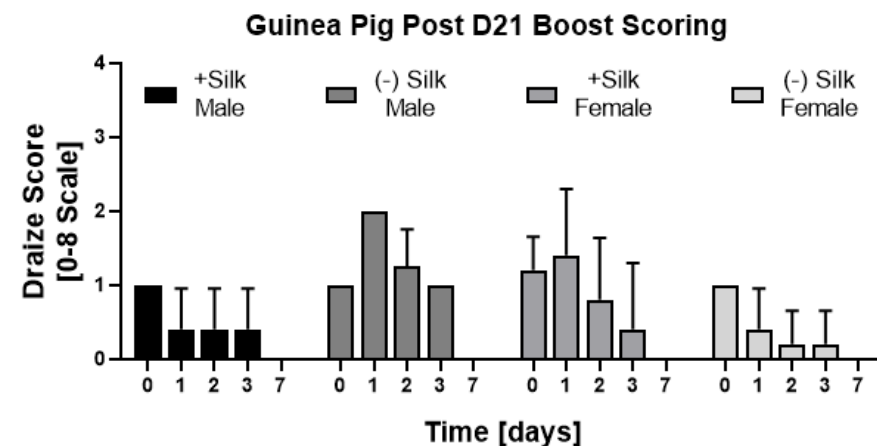
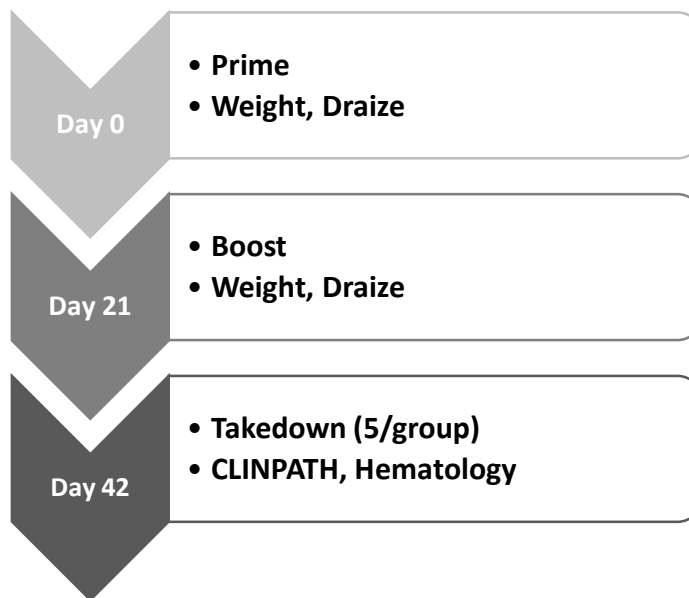


Validating Safety/Tox/Reactogenicity

Hairless Guinea Pig

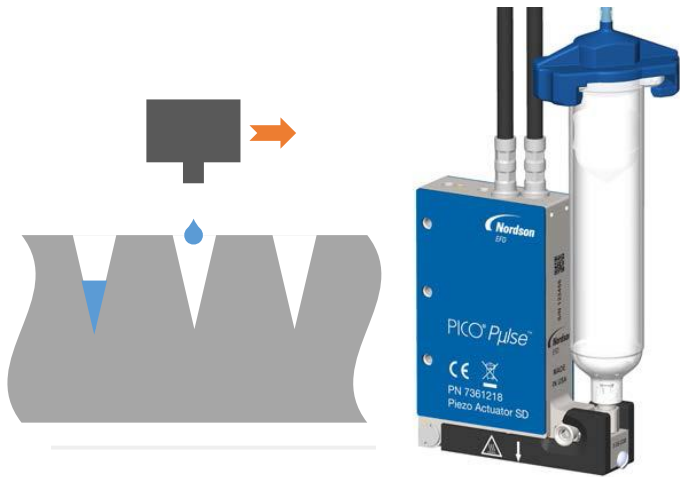
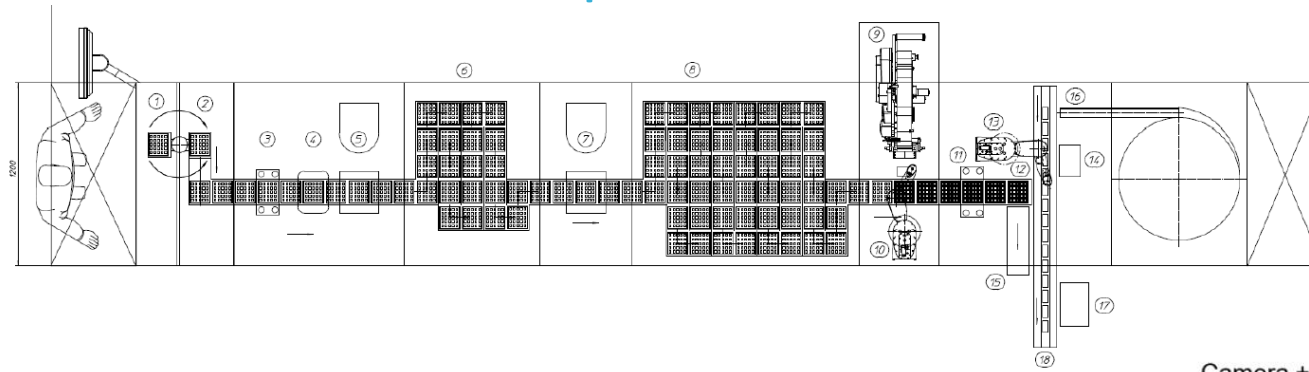


Group #	Type	Sex	Nair?	N
1	MAP	Male	No	5
2	MAP	Female	No	5
3	MAP (-silk)	Male	No	5
4	MAP (-silk)	Female	No	5

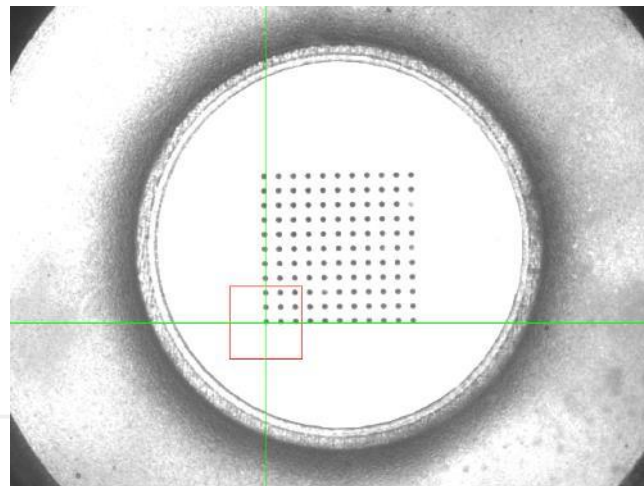


All patches were well tolerated. No safety concerns or evidence of toxicity for silk through MAP, ID or SQ

20M Patch/Year Scalable Manufacturing Line Plan Developed and Unit Ops Validated

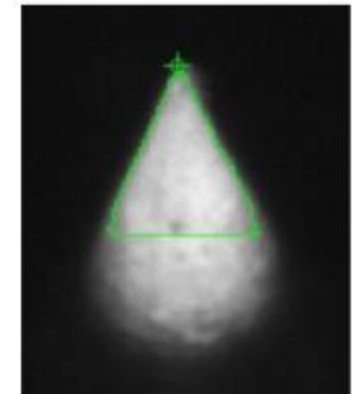
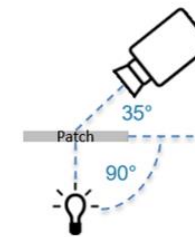


Individual needle filling



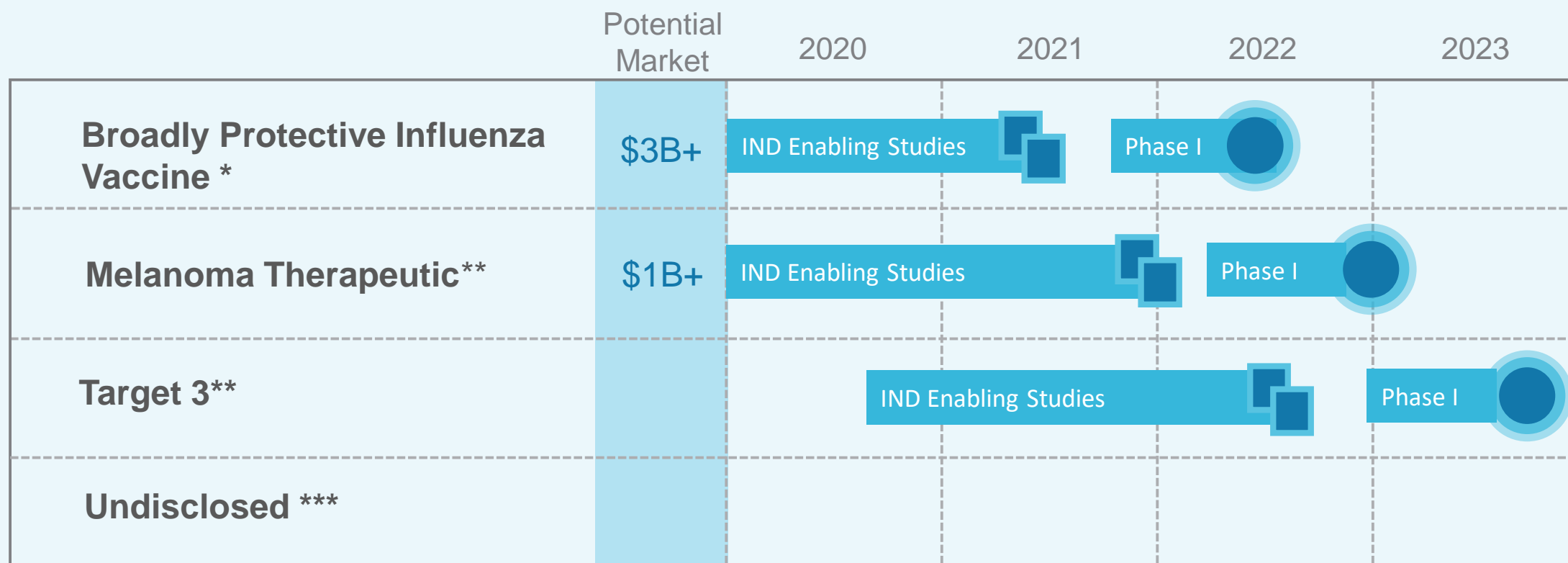
Vision-guided printing screenshot

Camera + light setup:



Machine vision inspection of demolded needle

Vaxess is moving multiple programs into the clinic in 2021-2022



*In-house Development with API from GC Pharma

** In-house Development with off-patent molecule

***Partner Joint Development

MIMIX™



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Proudly funded by:

