# PREVENTIVE THERAPEUTIC



This infographic will provide a visual overview of the key differences between preventive and therapeutic vaccines.



#### Viral vector vaccines

Use a modified version of an alternative virus as a vector, such as measles, influenza and adenovirus. **Example: Oxford–AstraZeneca COVID-19 vaccine.** 

#### Live-attenuated vaccines

Use a weakened (attenuated) form of the pathogen. Example: Measles, mumps and rubella (MMR combined vaccine).

#### Messenger RNA (mRNA) vaccines

mRNA encoding a protein (antigen) from the pathogen is delivered into the body. mRNA is transcribed and the resulting protein triggers an immune response. **Example: Pfizer-BioNTech COVID-19 vaccine – first mRNA vaccine to be authorized for human use.** 

#### Subunit, recombinant, polysaccharide and conjugate vaccines

Utilize a specific component of a pathogen – like a sugar, capsid or protein – to trigger an immune response. **Example: HPV (Human papillomavirus) vaccine.** 

#### Inactivated virus

A "dead" form of the pathogen is used to trigger an immune response. Several doses may be required, compared to a live-attenuated vaccine, to increase duration of immunity. **Example: Hepatitis A vaccine.** 

### **THERAPEUTIC VACCINES**

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Instead of immunizing to prevent a disease, therapeutic vaccines utilize a patient's own immune system to fight an existing disease, by stimulating it to target specific diseased





LET'S TAKE A CLOSER LOOK AT SOME EXAMPLES...

This is an example of how a peptide vaccine could be generated for a cancer patient, using sequencing methods to understand the molecular biology of their tumor and generate synthetic peptides.

#### **PEPTIDE VACCINES**

Peptides, typically 20—30 amino acids in length, are synthesized to create an immunogenic molecule that represents a specific epitope of an antigen.

Peptide vaccines are dependent on adjuvants and specific delivery systems to be effective as they can be poorly immunogenic.



Here is an example of how *in vivo* DC-targeted vaccines are produced.

#### DENDRITIC CELL VACCINES

Dendritic cells (DC) are the most potent antigenpresenting cells in the immune system.

DC-based vaccines can be broadly grouped into: *In vivo* DC-targeted vaccines and *Ex vivo* antigen-loaded vaccines



Currently, three therapeutic vaccines are approved by the US Food and Drug Administration for cancer treatment:



#### THERAPEUTIC HIV VACCINES

A therapeutic HIV vaccine would be utilized in patients diagnosed with HIV to:



Reduce viral load such that it is undetectable without the need for regular antiretroviral therapy (ART)

Slow down or precent the progression of HIV to acquired

Help to prevent transmission of HIV

immundeficiency syndrome (AIDS)

To date, a therapeutic HIV vaccine has not been approved for human use. However, a number of ongoing clinical trials are testing different modalities: DNA, protein, peptide, viral vectors and DC-based vaccines – the latter offering the most promising results so far.

# **FUTURE HORIZONS FOR VACCINE R&D**

The COVID-19 pandemic demonstrated the importance of vaccines for protecting and improving human health, injecting a renewed sense of energy for vaccine R&D. What does the future look like?

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#### Accelerated timelines

A preventive vaccine was developed against SARS-CoV-2 in 326 days. Global health authorities have agreed that the next pandemic vaccine will be delivered in 100 days. Several vaccine developers are exploring strategies for achieving this aim.

## Increased funding for R&D and vaccine access

Coalition for Epidemic Preparedness Innovations (CEPI) recently announced its \$3.5 billion pandemic preparedness strategy, where funds will be allocated to late-stage clinical trials, the continuation of vaccine development for priority pathogens and other R&D endevors. Over the next five years, The Global Fund for Children's Vaccines will give > \$150 million worth of vaccines to improve immunization programs in Africa, Asia and Latin America.

**Novel modalities** 

Following several decades of research, the first mRNA vaccine was authorized for human use in 2020. In the coming years we will likely see an increase in the application of mRNA technology for both preventive and therapeutic vaccines across different disease areas.



