



EN ESTE NÚMERO

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Noticias en la Web

Qdenga - A Promising Dengue Fever Vaccine

Oct 1. Dengue fever is transmitted through mosquito bites, and while mosquito control remains essential, vaccination offers an additional layer of protection. The World Health Organization (WHO) classifies dengue as one of the top 10 global health threats, with cases increasing tenfold from 500,000 in 2000 to over five million in 2019. In 2023 saw an unprecedented surge in dengue cases, with the first-ever global figures exceeding 6.5 million cases and over 6,800 deaths. Cases were reported in over



80 countries across Africa, the Americas, South-East Asia, the Western Pacific, and the Eastern Mediterranean regions. Asia recorded the highest case fatality ratio in 2023. In that year Bangladesh experienced its deadliest dengue outbreak in two decades, with 1,705 deaths out of 321,179 confirmed cases, a fatality rate of 0.53%.

In addition, spatial and temporal shifts in dengue patterns were observed in 2022 and continued in 2023. Nepal and Bangladesh experienced spikes in case numbers earlier than usual. Cases in Nepal shifted from the Kathmandu Valley in 2022 to the southeast Terai region and hill districts in Gandaki province in 2023. India, in 2023, experienced an increase in cases in Kerala and northeastern States bordering Bangladesh compared to the previous year. Vietnam has experienced dengue fever peaks approximately every 10 years between 1980 and 2018. However, the period from 2019 to 2023 saw two epidemic peaks in 2019 and 2022. In 2022, Vietnam reported over 367,000 dengue cases, second only to Brazil globally. WHO estimates about 390 million dengue infections occur worldwide annually, with 96 million symptomatic cases, 500,000 hospitalizations, and 40,000 deaths.

Vulnerable Populations and Treatment

Anyone can contract dengue fever, but children, pregnant women, and individuals with chronic diseases or obesity are at higher risk. Untreated dengue can lead to severe complications, including hypotension, heart failure, kidney failure, hemorrhagic shock, multiple organ failure, cerebral hemorrhage, and coma. Currently, treatment focuses on managing symptoms through blood filtration, plasma exchange, and anti-shock measures. The cost of treating a severe case can cross the economic ability of most of the people.

The approved dengue vaccines are Dengvaxia® and Qdenga® (TAK-003), with varying efficacies and recommended populations. Dengvaxia® has an efficacy of approximately 60–80% for preventing symptomatic dengue, but it is restricted to children aged 9-16 with a confirmed prior dengue infection due to an increased risk of severe disease in seronegative individuals. On the other hand Qdenga® shows higher overall efficacy, with around 80% protection against symptomatic dengue in both seropositive and seronegative individuals aged 4 and older.

Qdenga was invented by Takeda Pharmaceuticals, a Japanese company that developed the live, attenuated dengue vaccine (TAK-003) to protect against all four dengue virus serotypes. Regulatory

bodies that approved Qdenga include the European Medicines Agency (EMA) and the European Commission (EC) in 2022, followed by other countries like Indonesia, Great Britain, Brazil, and Argentina.

As of September 2025, QDenga was authorized in 40 countries and available in 27 countries, including Indonesia (approved Aug 2022, available April 2023), Brazil (approved Mar 2023, available June 2023, Thailand (approved May 2023, available Aug 2023), Argentina (approved Apr 2023, available Oct 2023), (approved February 2024), and is also available in Vietnam.

Efficacy and Impact of Qdenga

The Qdenga vaccine has demonstrated an efficacy of up to 80% in protecting against all four dengue virus types. More importantly, it reduces the risk of hospitalization by up to 90%. This is particularly crucial in different countries like Bangladesh, where a significant portion of the population has been infected with dengue fever at least once. Subsequent infections tend to be more severe, making timely vaccination a vital preventive measure.

A vigorous consultation with all concern stakeholders early engagement with Takeda to secure a substantial supply of the vaccine may emphasis the commitment to reducing the annual burden of dengue fever.

Our people are frequent travelers to dengue-endemic areas and there is already a demand and questions on how and when to use the vaccine. Therefore, the Vaccine Expert group of the Swedish Society for Infectious Diseases Physicians have reviewed the literature to guide and formulate recommendations on the use of Qdenga® as a travel vaccine.

For travelers with previous known self-reported (hospitalized or polyclinic testing) dengue fever, vaccination is recommended before travel to an endemic country.

For dengue naïve travelers vaccination may be considered in individuals aged 4–16 years old irrespective of travel duration.

For travelers aged 17–60 years old are recommended considering vaccination only for longer trips and related to travel destination. They also have suggested a trip for more than six weeks to South-East Asia, a region with among the highest global incidence of dengue fever, to be used as a reference.

Since Qdenga® has not yet been studied in individuals >60 years old, they clearly advise vaccination should be avoided in this group until data are available.

Travelling after only one vaccine dose should be avoided if possible, which is a challenge for most travelers coming with shorter notice for travel medicine advice. It can be considered, especially in individuals with previous dengue fever, given that they receive the second dose after return. As far as we know these recommendations are more cautious than in other non-endemic countries, especially regarding the upper age limit, and should be continuously revised when more information and experience become available. For safely travelling and living, management of aedes mosquitoes along with vaccination is essential.

Fuente: Daily Sun. Disponible en <https://n9.cl/u1cg8>

New research finds HPV vaccine delivers herd immunity, protecting unvaccinated women

Oct 1. A study spanning nearly two decades has found that vaccinating communities against human papillomavirus (HPV) doesn't just protect those who receive the shot, it also helps shield women who haven't been vaccinated.

HPV is the main cause of cervical cancer – a leading cause of death among women, especially in low- and middle-income countries.

The HPV vaccine, if received in early adolescence, is known to be highly effective at preventing cervical cancer in later life.



A girl with a health worker during a HPV vaccination session in Eswatini. Credit: Gavi/2025/Svetlomis Slavchev

Led by researchers at Albert Einstein College of Medicine and published in [JAMA Pediatrics](#), the research analysed data on more than 2,300 adolescent and young adult women aged 13 to 26 years at enrolment, from 2006 to 2023.

The researchers found that HPV vaccination programmes offered significant protection through herd immunity: immunity conferred on unvaccinated people when enough people are vaccinated in a community.

Widespread vaccination would have lowered overall transmission of the virus, benefitting even those not immunised.

Dr Jessica Kahn, professor of paediatrics at the Albert Einstein College said that while randomised controlled studies had clearly shown the value of the HPV vaccine, the team wanted to “evaluate how the vaccine works in real-world settings that included young women who were at relatively high risk for HPV and had different levels of vaccine uptake”.

Dramatic drops in HPV infections

Over the study period, HPV vaccination coverage in Cincinnati, USA, where the study data was taken from, rose from 0% to 82%.

During that time, HPV infections in vaccinated women fell steeply:

- In women covered by the bivalent vaccines against HPV types 16 and 18, infections dropped by 98.4%
- In women covered by the quadrivalent vaccines against HPV types 6, 11, 16 and 18, infections fell by 94.2%
- Infections of types addressed by the newer nonavalent vaccine dropped by 75.7%

Even among women who never received the vaccine, infection rates for HPV types covered by older vaccines declined dramatically.

- Infections with HPV types included in the bivalent vaccine decreased by 71.6%
- Infections with HPV types included in the quadrivalent vaccine dropped by 75.8%

Data for herd immunity from the nonavalent vaccines is still underway.

The decline is attributed to vaccination

“Our analysis of the data indicates that those reductions in infection rates were primarily due to the vaccine’s introduction and not because of changes in sexual behavior or other factors,” said Aislinn DeSieghardt, the

paper's first author and clinical research coordinator at the Cincinnati Children's Hospital Medical Center.

Because the study tracked trends over such a long span and included women at higher risk (for example many had multiple sexual partners or prior STIs) the results offer compelling “real-world” validation of the vaccine's broader population impact.

The study reinforces that strong vaccine uptake in a community can have important collateral benefits, reducing the burden of HPV even among those who are unvaccinated.

Dr Kahn concludes that “By expanding uptake of this highly safe and effective vaccine, and ensuring access to screening and treatment, we can achieve one of the greatest public health victories of our time: the elimination of cervical cancer worldwide.”

Fuente: Gavi. Disponible en <https://n9.cl/v7sag>

New study shows all-in-one coronavirus vaccines could save millions of lives in future pandemics

Oct 2. A new study published in *Nature Communications* reveals that vaccines designed to protect against a wide range of coronaviruses could dramatically reduce deaths and economic disruption and limit the need for lockdowns in the event of another global coronavirus pandemic.

The research, led by Imperial College London and supported by CEPI, modelled the potential impact of broadly protective sarbecovirus vaccines — innovative vaccines that could defend against multiple viruses including a novel sarbecovirus, referred to as SARS-X. Sarbecoviruses are types of coronaviruses that include both SARS-CoV-1, which caused the 2002–2004 SARS outbreak, and SARS-CoV-2, which continues to cause COVID-19.

Broadly protective sarbecovirus vaccines could be manufactured and stockpiled ahead of time and deployed as soon as a new coronavirus is detected, protecting the most vulnerable in the critical early months of a future outbreak while a new vaccine targeting the specific pathogen is being developed.

The modelling shows that, during a SARS-X outbreak, vaccinating high-risk populations aged 60 years and older with a broadly protective sarbecovirus vaccine could reduce mortality by more than half, while also cutting the need for disruptive non-pharmaceutical interventions, such as lockdowns, by more than 60 percent.

“COVID-19 showed us the catastrophic costs of facing a deadly new virus without a vaccine,” said Dr Richard Hatchett, CEO of CEPI. “Broadly protective coronavirus vaccines could change the game. They would buy us valuable time in a future pandemic—saving lives, easing the burden on health systems, and helping societies to stay open—while scientists develop vaccines specifically targeting the new pathogen.”

Vaccines to stay one step ahead of coronaviruses

The study explored multiple scenarios, from SARS-CoV-1-like outbreaks with limited transmission to highly transmissible SARS-CoV-2-like pandemics, and considered a range of assumptions about the size of the vaccine stockpile and the stringency of non-pharmaceutical interventions.

- Modelling shows that broadly protective sarbecovirus vaccines could halve the number of deaths in a future pandemic.
- Stockpiled vaccines would dramatically reduce the need for non-pharmaceutical interventions like lockdowns.
- If they had been available during the COVID-19 pandemic, as many as 65 percent of deaths could have been averted in the first year.



It found that in outbreaks caused by slower-spreading viruses, using broadly protective sarbecovirus vaccines in ring vaccination strategies—where contacts of infected individuals are rapidly immunised—could help to contain transmission and bring an outbreak to an end. For fast-spreading viruses, like SARS-CoV-2, all-in-one shots would not contain outbreaks but could slow the spread of the disease and provide critical protection for those most at risk until a vaccine tailored to the new virus is developed.

In a scenario where it takes 250 days to create such a virus-specific vaccine, broadly protective vaccines could cut deaths in an average upper-middle-income country by more than half.

Even in a scenario in which CEPI's 100 Days Mission is achieved—a goal to develop new vaccines within 100 days of a pandemic pathogen being identified—and a virus-specific vaccine is available in just three months, all-in-one vaccines would provide a crucial safeguard against more transmissible pathogens. Faced with a pathogen with a reproduction rate of $R_0=3.5$ – similar to that of smallpox, for example – broadly protective sarbecovirus vaccines could reduce deaths by 43 percent and the number of days under non-pharmaceutical interventions by almost 70 percent.



Dr Charlie Whittaker, Assistant Professor, Infectious Diseases & Vaccinology, University of California, Berkeley, who led the modelling study, said: “Our work highlights that a stockpiled broadly protective coronavirus vaccine could halve deaths in a future pandemic. Achieving this impact depends on more than just the vaccines alone though - it will require investment in manufacturing and delivery systems to ensure equitable global access, so that everyone, everywhere can benefit when the next pandemic arrives.”

The researchers also looked back at the COVID-19 pandemic to assess the potential impact of broadly protective sarbecoviruses had they been available. They estimate that as many as 65 percent of deaths could have been averted during the pandemic's first year, depending on the size of the stockpile. For example, a moderate stockpile with enough doses to vaccinate 60 percent of older adults, could have saved 64,000 lives in Italy (52 percent of those who died), 56,000 lives in Iran (45 percent) and 89,000 lives in Bangladesh (69 percent) during that period.

CEPI's leadership in broadly protective vaccines

CEPI is a world leader in advancing broadly protective sarbecovirus vaccines, committing over US\$200 million to date in a portfolio of vaccines spanning a diverse range of technologies. From RNA to protein-based candidates, the CEPI-supported vaccines each use different strategies for achieving broad protection.

Dr Hatchett added: “We cannot predict when or where the next coronavirus outbreak will emerge, but we can prepare. By investing in all-in-one vaccines now, CEPI is boosting the world's defences against future coronavirus pandemics.”

The modelling study was conducted by a consortium of researchers from Imperial College London, the University of California, University of New South Wales, Duke-NUS Medical School, the London School of Hygiene & Tropical Medicine, National University of Singapore, National Centre for Infectious Diseases Singapore, Nanyang Technological University Singapore, and CEPI.

The full study is available open access in Nature Communications: Quantifying the impact of a broadly protective sarbecovirus vaccine in a future SARS-X pandemic | Nature Communications

Fuente: CEPI. Disponible en <https://n9.cl/4zd7o>

Alerta mundial: la OMS confirma la nueva cepa de COVID-19 "Stratus" con un síntoma inesperado que la delata

3 oct. La Organización Mundial de la Salud emitió una advertencia sobre una nueva variante del coronavirus que está ganando terreno en los primeros compases del otoño.

Con la llegada de temperaturas más bajas y las primeras lluvias, los virus respiratorios encuentran su entorno ideal para propagarse, y esta nueva cepa denominada Stratus está demostrando una capacidad de transmisión preocupante.

Lo que distingue a esta variante del resto no son solo sus índices de contagio, sino un síntoma particular que está sirviendo como señal de alarma: la ronquera persistente.

Los datos más recientes indican que esta variante ya se detectó en 38 países y representa más del 20% de todos los casos de COVID-19 registrados actualmente a nivel mundial. Aunque las autoridades sanitarias califican el riesgo para la salud pública como bajo, la rapidez de su expansión puso en alerta a los sistemas de vigilancia epidemiológica de todo el planeta.

¿Qué es exactamente la variante Stratus y por qué preocupa a los expertos?

La cepa Stratus, identificada científicamente como XFG, pertenece a lo que los virólogos denominan combinaciones resultantes de la recombinación de dos o más cepas preexistentes. En este caso específico, Stratus es el resultado de la fusión entre las variantes LF.7 y LP.8.1.2, ambas circulantes anteriormente.

Este tipo de recombinaciones genéticas no son inusuales en virus ARN como el SARS-CoV-2, especialmente cuando múltiples variantes coexisten en la misma población. Lo que sí resulta destacable es la velocidad con la que Stratus logró posicionarse como una de las cepas dominantes en un período relativamente corto, superando ya el 20% de prevalencia en el mundo.

A pesar de su rápida propagación, la OMS mantiene una evaluación de riesgo bajo para la salud pública, sugiriendo que, si bien es más transmisible, no parece causar enfermedad más grave que sus predecesoras.

El síntoma revelador: ronquera persistente que marca la diferencia

La característica más llamativa de la variante Stratus es sin duda la ronquera o disfonía que están experimentando la mayoría de los pacientes infectados. Este síntoma, que implica una alteración en la calidad de la voz, va desde una ligera aspereza hasta la pérdida casi completa de la capacidad de hablar con normalidad.

Los especialistas explican que esta afectación vocal sugiere una predilección particular de esta variante por las vías respiratorias superiores, específicamente la laringe y las cuerdas vocales. Mientras que variantes anteriores también podían causar dolor de garganta, la ronquera marcada no había sido un síntoma tan predominante.

Además de la alteración vocal, los pacientes con Stratus continúan presentando el cuadro clínico habitual del COVID-19: congestión nasal, dolor de garganta y de cabeza, fiebre, fatiga extrema y en algunos casos, molestias digestivas.

Fuente: El Cronista. Disponible en <https://n9.cl/swnehj>



AI can strengthen pandemic preparedness

Oct 3. How to identify the next dangerous virus before it spreads among people is the central question in a new Comment in *The Lancet Infectious Diseases*. In it, researchers discuss how AI, combined with the One Health approach, can contribute to improved prediction and surveillance.

"Artificial intelligence cannot by itself prevent pandemics, but the technology can be a powerful supplement to the knowledge and methods we

already use. The better we become at integrating data from humans, animals, and the environment, the better prepared we will be," says Professor Frank Møller Aarestrup from the DTU National Food Institute in Denmark, one of the authors of the Comment.

It was co-authored by Professor Marion Koopmans from the Erasmus Medical Center in the Netherlands. She warns that once a disease starts spreading, it is very hard to bring under control.

"The interventions required are drastic—as we saw during COVID-19. That is why it is crucial to detect new pathogens before they gain a foothold," says Koopmans, noting that once established, new diseases can become persistent challenges, as COVID-19 has also shown.

The team of authors, which also includes experts from Eötvös Loránd University (ELTE) in Hungary, the University of Bologna in Italy, and the UK Animal and Plant Health Agency, speaks from their experience as collaborators over years, focusing on One Health approaches to emerging disease preparedness in the VEO consortium—a European research initiative developing data-driven tools to detect and track emerging infectious diseases.

Pandemics often originate in animals

The outbreaks of diseases such as SARS-CoV-2, avian influenza, and mpox demonstrate the difficulty of controlling new potential epidemics. Many pathogens originate in animals, but when and where they will spill over into humans is unpredictable.

The authors of the Comment highlight how climate change, intensive animal production, and human encroachment into natural habitats increase the risk of so-called spillover events—situations in which pathogens cross from animals to humans and, in the worst case, develop into epidemics.

Spillovers have been likened to sparks: most extinguish, but some ignite fires that spread uncontrollably. Being able to detect such spillovers as early as possible is a challenge that the team has been studying using big data approaches.

AI can reveal patterns in complex datasets

Artificial intelligence can help to analyze such datasets from diverse sources—such as climate, land use, animal production, transport, population movements, and socio-economics. When these datasets are combined, AI can reveal patterns that would otherwise be difficult to discern.

"AI can help us identify where in the world surveillance should be intensified geographically, but also in specific animal species, in wastewater, or in humans. In this way, we can prioritize efforts where the risks are greatest, so-called hotspots," says Aarestrup.



Genetic signals as early warning

Once such hotspots are predicted, metagenomic sequencing can be added as a catch-all approach for detection of pathogens, both known and new ones. Metagenomic sequencing is the analysis of genetic material—in samples from wastewater, air, food, or the environment. It is increasingly used to provide insight into a vast diversity of known and unknown microorganisms. Many of the genetic fragments identified are not yet characterized.

"When we sequence a sample, we may find millions of genetic fragments. Most resemble something familiar and harmless, but we are left with thousands of unknowns. Here, AI can help detect patterns and point to what might be dangerous," explains Aarestrup.

Once it is clear there is a potential pathogen, questions can arise about how dangerous it is. The potential for viruses from animals to infect humans, spread and cause disease in part is embedded in the genetic code. AI-based tools can be used to predict how mutations might alter viral properties.

"We see huge developments in this area. AI-based protein models can provide an indication of what a mutation does to the structure of viruses, and how that then can be translated to risk of spread, or risk of severe disease. While challenging now, we see great potential for the use of AI to speed up risk assessment," says Koopmans.

AI as a co-scientist—opportunities and limitations

The comment also describes early prototypes of so-called AI "co-scientists," capable of conducting an entire research cycle—from hypothesis generation and literature review to data analysis and reporting.

"I envisage AI becoming a recognized competence at the table—on a par with different types of researchers. AI can deliver analyses or suggestions that we as scientists can evaluate. In that way, the technology becomes a supplement that can strengthen our decision-making processes," says Aarestrup.

"That also implies that we need to learn what our future role is as teachers and supervisors. How do we make sure that the novel ways of working provide trustworthy output? Will we be able to recognize mistakes with advancements of AI models? We also need to go back to the classroom. Really exciting," says Koopmans.

The authors conclude that artificial intelligence offers intriguing possibilities for enhancing pandemic preparedness. Still, it must be seen as a complement—not a replacement—to the classical surveillance and research approaches already in use.

Fuente: medicalxpress. Disponible en <https://n9.cl/gvb77>

Cuba vacunará a niñas de nueve años contra el VPH

3 oct. Autoridades sanitarias anunciaron la inclusión de la vacuna contra el Virus del Papiloma Humano (VPH) en el Programa Nacional de Inmunización, con la aplicación a todas las niñas de nueve años en Cuba.

Dagmar García, vicedirectora del Instituto Finlay de Vacunas, explicó que la decisión respondió a la relación comprobada entre la infección persistente por VPH y el cáncer de cuello uterino, una de las principales causas de muerte por cáncer en mujeres a nivel mundial, precisó el diario Granma.

La especialista acotó que se empleará la vacuna CECOLIN, de tipo bivalente, que protege contra los serotipos 16 y 18,



responsables de alrededor del 70 por ciento de los casos de esa enfermedad.

García señaló que la inmunización resulta más efectiva antes del inicio de la vida sexual, por lo cual se determinó aplicarla a niñas de nueve años, en correspondencia con las recomendaciones internacionales.

La vacuna CECOLIN, producida por la empresa china Innovax, cuenta con la precalificación de la Organización Mundial de la Salud y ha demostrado seguridad y eficacia en su uso.

El Programa Nacional de Inmunización, en funcionamiento desde 1962, administra cada año un promedio de 4 800 000 dosis de 13 tipos de vacunas, de las cuales ocho son de producción nacional.

Gracias a ese programa, Cuba eliminó seis enfermedades transmisibles, entre ellas la poliomielitis y el sarampión, y redujo complicaciones graves como el síndrome de rubéola congénita.

Con la incorporación de la vacuna contra el VPH, el país refuerza su estrategia de prevención y se adelanta en más de una década a la aparición de casos de cáncer cervicouterino en las nuevas generaciones.

Fuente: El Artemiseño. Disponible en <https://n9.cl/14ywy>

Europa compra la vacuna de Hipra contra la COVID-19 adaptada a LP.8.1

4 oct. La Autoridad de Preparación y Respuesta ante Emergencias Sanitarias de la Comisión Europea (HERA, por sus siglas en inglés) y la farmacéutica biotecnológica Hipra han firmado un contrato de compra de su vacuna Bimervax, de proteína recombinante monovalente con adyuvante adaptada a la variante LP.8.1 de la COVID-19. El acuerdo, con una duración de hasta dos años, permitirá a los países acceder a hasta 4 millones de dosis de la vacuna proteica de Hipra, sin que exista una obligación mínima de compra. Las entregas se realizarán a tiempo para la campaña de vacunación de este otoño, lo que refuerza la preparación de la Unión Europea frente al virus en los próximos meses.

La firma se ha llevado a cabo en Bruselas el viernes 3 de octubre entre David Nogareda, presidente y CEO de HIPRA, y Laurent Muschel, director de la Health Emergency Preparedness and Response Authority (HERA).

Hadja Lahbib, comisaria europea de Igualdad, Preparación y Gestión de Crisis, señaló que “con el aumento de casos de la variante conocida como ‘Frankenstein’, debemos garantizar que se mantiene la protección frente a esta enfermedad, especialmente para las personas más vulnerables. Con un portafolio diversificado de vacunas, que ahora incluye hasta 4 millones de dosis de esta vacuna proteica, reforzamos nuestra preparación y aseguramos el suministro de soluciones médicas necesarias frente a la amenaza constante de la Covid-19. La vacuna de Hipra sigue un enfoque integral, desde la investigación hasta la producción, íntegramente en Europa, lo que fortalece nuestra autonomía estratégica. Estamos comprometidos con reforzar nuestra seguridad sanitaria para una Europa más segura, saludable y mejor protegida”.

Aunque las vacunas de ARNm ya están disponibles, este contrato de adquisición conjunta diversifica el acceso para los ciudadanos europeos al incluir una vacuna de proteína recombinante. La vacuna de HIPRA, que ha recibido recientemente autorización de comercialización de la Comisión Europea tras demostrar que genera inmunidad frente a la variante LP.8.1 y protección cruzada frente a variantes emergentes como NB.1.8.1 y XFG, está lista para su distribución en viales monodosis y puede almacenarse entre 2 °C y 8 °C durante 12 meses.

Los estudios realizados con la vacuna original y con su adaptación han demostrado que es segura, con menos efectos secundarios que la comparativa de ARNm y genera una respuesta inmunitaria fuerte, duradera y con amplia reactividad cruzada frente a variantes emergentes.

Durante la firma, David Nogareda, presidente y consejero delegado de Hipra, subrayó que “este acuerdo es una muestra de confianza en la capacidad de Hipra para contribuir a la protección de la salud pública. Desde la investigación hasta la fabricación, todo el proceso tiene lugar en Europa, lo que nos permite garantizar calidad, seguridad e independencia estratégica. Con este contrato, reforzamos no solo la respuesta frente a la COVID-19, sino también la preparación de Europa ante futuros retos sanitarios”.

Fuente: DiarioFarma. Disponible en <https://n9.cl/k03dpa>

A next-generation vaccine technology has been developed that has a stronger immune effect than conventional injections

Oct 5. A next-generation vaccine technology has been developed that has a stronger immune effect than conventional injections just by applying it like a lotion without a syringe. It is predicted that it will not only dramatically increase the convenience of vaccination by eliminating pain and rejection, but also increase vaccine efficacy, which can emerge as a new alternative to the vaccine market in the future.

A joint research team from France's National Institute of Health and Medicine (INSERM) and King's College London in England has developed a new platform technology that delivers vaccines in a way that gently increases the skin with special devices, according to the international journal "Cell Reports."

The key to this technique is the gentle tension it puts on the skin. When the skin is relaxed for about 20 minutes

with a device developed by the research team, collagen fibers in the skin are rearranged and hair follicles (fur holes) are temporarily opened. This is the principle that vaccine antigen particles penetrate deeply into the skin effectively through this gap.

The biggest finding of this study is that the act of increasing the skin itself, beyond simple drug injection, acts as a powerful 'immune promoter'. When the skin is relaxed, it has been confirmed that immune cells recognize the fine introduced skin bacteria as external invasion and operate the immediate defense system. In other words, the effectiveness of the vaccine can be maximized without a separate immunostimulator.

In fact, the research team administered the flu vaccine to mice in this way, and as a result, antibodies were formed at a much higher level than the general muscle injection method.

Dr. Stuart Jones, King's College London, UK, who led the study, said, "It has led to a more effective immune response just by increasing the skin than by needle inoculation. This is a practical achievement that can change the existing notion of vaccine delivery."

The research team expected that the technology developed this time could be widely used to deliver not only vaccines but also various biopharmaceuticals such as cell therapy and polymer drugs without pain in the future.

Fuente: MAEIL BUSINESS NEWSPAPER. Disponible en <https://n9.cl/rgppux>



[Picture = Pixabay]

CEPI backs new research into vaccines against multiple deadly filoviruses

Oct 6. CEPI will fund new research conducted by scientists at the Stanford School of Medicine to study whether new vaccines could offer broad protection against multiple deadly pathogens from the filovirus viral family.

Supported by up to \$18 million in CEPI investment, researchers will design and test new vaccine constructs that could provide all-in-one protection against viruses including Zaire Ebolavirus, Marburg and Sudan Ebolavirus, and even filoviruses that we don't yet know about that could spill over from animals into humans in the future.



"A broadly protective filovirus vaccine could transform the world's defences against outbreaks of some of the world's most dangerous pathogens, for the benefit of all," explains CEPI's Executive Director of Research & Development Dr Kent Kester. "If we solve the scientific challenge of developing all-in-one filovirus vaccines now, we can ensure the world is ready to respond at speed to newly emerging filoviruses and potentially take the threat of future filovirus pandemics off the table."

The research will use cutting-edge artificial intelligence to design immunogens – the substance in a vaccine that provokes an immune response – that may be capable of protecting against more than one filovirus. These immunogens will be combined with a ferritin-based protein-nanoparticle vaccine platform to create a range of vaccine candidates that will be tested in preclinical studies to establish proof of concept. The most promising vaccine candidate will be advanced to the point where it is ready to rapidly enter Phase I clinical trials should an unknown filovirus outbreak emerge.

The ferritin nanoparticle vaccine platform is favourable for use in low- and middle-income countries as it does not require complex frozen storage that can impact access to doses in low-resource settings. It has previously been tested in Phase I clinical trials for influenza and COVID-19 vaccines, which generated positive safety data.

Professor Peter S. Kim, the Virginia & D.K. Ludwig Professor of Biochemistry, at the Stanford School of Medicine, and Principal Investigator on the project said: "We are excited to collaborate with CEPI to develop a globally effective broad-spectrum vaccine to protect against future filovirus pandemics."

Preparing for both known and unknown filovirus threats

Filoviruses such as Ebolavirus Zaire, Sudan Ebolavirus and Marburg cause frequent outbreaks in African countries, characterized by high fatality rates and significant societal and economic impacts. A single vaccine capable of combating all of these known threats could offer a cost-effective solution for proactively immunizing those who are most likely to be infected by one or more of the viruses.

Crucially, the project will also equip scientists with the tools they need to respond at pace if faced with an unknown pathogen – a so-called 'Disease X' - from the filovirus family. The scientific knowledge and immunogen designs generated by the research will be added to CEPI's 'Disease X Vaccine Library', a comprehensive collection of vaccine data and information about the most worrying viral families that can be rapidly adapted for use during outbreaks of novel viruses with pandemic potential.

This approach to significantly accelerate vaccine development is central to the 100 Days Mission: a global goal, spearheaded by CEPI and embraced by G7 and G20 nations to develop pandemic-busting vaccines in just 100 days and make the world a safer place for all.

Committing to equitable access

CEPI is committed to enabling equitable access to the outputs of this project, in line with CEPI's Equitable Access Policy. This ultimately includes a commitment to vaccines being available first to populations at risk when and where they are needed, including specific obligations in the agreement related to affordable pricing and supply volumes for public health needs. It also includes sharing of any commercial benefits arising from CEPI's funding. Should the project be successful, CEPI has the option to explore providing further funding to the Stanford School of Medicine to develop its vaccine product and/or its immunogen designs, potentially combining them with other vaccine platforms via CEPI matchmaking for pandemic preparedness and response. Project results, including related data, will be published open access for the benefit of the global scientific community.

Fuente: CEPI. Disponible en <https://n9.cl/7by7t>

Pneumococcal Conjugate Vaccine Booster Maximizes Protection Against IPD

Oct 7. Pneumococcal conjugate vaccines (PCVs) were deemed highly effective in preventing invasive pneumococcal disease (IPD) among children, according to a study published in *Vaccine: X*. The study also revealed that PCV booster doses play a key role in maximizing vaccine type-IPD protection within the same population.

"*Streptococcus pneumoniae* is a bacterial pathogen responsible for a wide range of diseases, from mucosal infections, such as otitis media, sinusitis, and pneumonia, to IPD, which includes bacteremic pneumonia, sepsis, and meningitis," wrote the authors of the study. "High-risk populations include children under 5 years, adults over 65 years, individuals with immunocompromising conditions, those with chronic comorbidities, and individuals with lifestyle risk factors such as smoking, alcohol misuse, and being unhoused."

According to CDC data, IPD incidence falls at 8.3 cases per 100,000 patients. Further stratifying these statistics, the average mortality rate was 1 death per 100,000. Accounting just for children under the age of 18, this age group's incidence rate was 6.9 per 100,000. Finally, these data also showed a trend of greater risk for IPD complications as ages were reported to be lower.

Despite the continued prominence of IPD, especially among older adults and pediatric patients, its common occurrence has been significantly ameliorated through the introduction of PCVs.

Indeed, after the US' implementation of PCV7 in 2000, rates of pediatric IPD have decreased significantly. Because of vaccine protection against IPD, childhood rates of the disease decreased by 72% from 2002 to 2021. Since the introduction of all new PCVs following PCV7, researchers are still trying to determine the proper dosing and administration of PCVs to maximize protection against pediatric IPD.

"While previous research has evaluated overall [vaccine effectiveness (VE)], the impact of different pneumococcal dosing regimens remains unclear," they continued. "Given the importance of optimizing pneumococcal vaccination strategies, we aim to evaluate the effectiveness of PCV7 and PCV13 based on the number of vaccine doses received in this systematic review and meta-analysis."

To go about reassessing PCV7 and PCV13, researchers conducted a systematic review and meta-analysis of studies that explored the VE of each PCV valency among patients 18 years and younger.

More specifically, to determine VE among this population, researchers separated participating patients into 4 groups based on vaccination status at the onset of children's IPD. The 4 groups were a primary plus booster group (1 to 3 primary doses less than 12 months of age plus 1 booster dose 12 months or greater); the 1 primary dose group; 2 primary doses group; and 3 primary doses group (primary doses given <12 months of age and no booster).

With patients telling providers their vaccination status while also reporting IPD, the researchers were better able to determine how effective they were and what the ideal PCV dose should be.

After identifying 1982 studies on their initial search, study experts concluded their review with a total of 25.

"Pooled results showed that children receiving 2 or 3 primary doses without a booster had a higher risk of IPD compared with those completing a schedule with a booster dose (2 + 1 or 3 + 1 schedules)," wrote the authors. "Our findings reinforce the existing evidence supporting a vaccination schedule that includes a booster dose, as we found neither 2 nor 3 primary doses alone consistently achieved a pooled VE of >90% as with a schedule including a booster dose."

Previous evidence has shown that IPD can occur in vaccinated individuals due to PCV7 and PCV13's inability to protect against vaccine serotypes 3 and 19A. However, with the current finding of the booster's significant success over 1, 2, or even 3 doses of the primary vaccine alone, researchers uncovered an ideal PCV regimen for keeping children protected against IPD. As long as primary vaccine doses are accompanied by a subsequent booster, children 18 years and under can become even better protected against IPD.

"Our study reaffirms the high overall effectiveness of PCVs in preventing IPD in children and highlights the essential role of the booster dose in maximizing protection against [vaccine type]-IPD," concluded the authors. "As many countries transition to reduced dosing schedules to improve cost-effectiveness, streamline immunization programs, and address vaccine supply constraints, our findings emphasize the need for a full vaccination series that includes a booster to ensure optimal protection against IPD."

Fuente: Drug Topics. Disponible en <https://goo.su/Hr3Dy7>

New Vaccine Shows Promise Against Typhoid and Invasive Salmonella in First Human Trial

Oct 8. Researchers at the University of Maryland School of Medicine's Center for Vaccine Development and Global Health (CVD) have completed a successful Phase 1 clinical trial of a novel vaccine designed to protect against both typhoid fever and invasive non-typhoidal Salmonella--two major causes of illness and death among children in sub-Saharan Africa.

Results were published today in the journal *Nature Medicine*.

The investigational Trivalent Salmonella Conjugate Vaccine (TSCV) includes sugar molecules taken from the outer coating of the Salmonella typhi bacteria that cause typhoid and the two most common types of invasive Salmonella infections that are not associated with typhoid caused by Salmonella enterica. These sugars are attached to special proteins that help the body recognize and respond to the bacteria more effectively.

In the randomized, placebo-controlled Phase 1 trial, 22 healthy adults in the US received either a low dose (6.25 µg) or high dose (12.5 µg) of TSCV, or a placebo shot. The vaccine was found to be safe and well-tolerated, with only mild, short-lived injection site pain reported.



Importantly, 100 percent of the vaccine recipients developed strong immune responses to all three polysaccharide components, while none of the placebo recipients did.

“These results are highly encouraging,” said study lead investigator Wilbur Chen, MD, MS, Professor of Medicine at UM School of Medicine and Chief of the Adult Clinical Studies section within CVD. “They show that TSCV has the potential to protect children in regions where both typhoid and salmonella are endemic and deadly.”

The vaccine also has the potential to protect Americans against Salmonella infections, one of the leading causes of foodborne illness. Every year, bacteria from raw or undercooked chicken and eggs and contaminated produce cause 1.35 million Salmonella infections in the US and more than 26,000 hospitalizations, according to the Centers for Disease Control and Prevention. The serotypes targeted by the TSCV vaccine are among the most common in US infections.

In fact, some study participants showed pre-existing antibody responses, suggesting prior exposure to the bacteria via foodborne illness. Such priming might have led to stronger and longer-lasting immunity in the adult study volunteers, though researchers remain optimistic about the vaccine’s effectiveness in infants and young children in endemic regions.

The vaccine triggered a strong and balanced immune response, including long-lasting antibodies—even at lower doses. It also activated a specific immune defense involving white blood cells that help clear infections, which hadn’t been seen before with one of the vaccine’s protein components. These findings suggest the vaccine could offer both gut-level and whole-body protection against Salmonella.

“These findings provide a strong foundation for future studies,” said study co-author Myron Levine, MD, DTPH, professor emeritus at UM School of Medicine and CVD founding director. “We plan to explore broader functional assays to identify correlates of protection and evaluate TSCV’s performance in young children—the population most vulnerable to these diseases.”

The vaccine was developed in collaboration with Bharat Biotech International Limited (BBIL), building on their WHO-prequalified Typbar TCV™ platform.

“In 2017, sub-Saharan Africa saw over 420,000 cases of Salmonella disease and 66,000 deaths, primarily among children,” said Mark T. Gladwin, MD, Dean of the UM School of Medicine. “Typhoid fever caused an additional 650,000 cases and nearly 9,000 deaths in the region. A single vaccine that protects against both could be a game-changer for global pediatric health.”

Fuente: School of Medicine, University of Maryland. Disponible en <https://goo.su/A4XiSiL>

Moderna opens Oxfordshire site for vaccine development

Oct 9. The new Moderna Innovation and Technology Centre, which will develop and manufacture mRNA vaccines, has been officially opened by Health and Social Care Secretary Wes Streeting.

Pioneering US biotech firm Moderna will be able to make millions of vaccines for seasonal viruses like COVID-19, create dozens of highly skilled jobs, and help Britain prepare for future pandemics thanks to its cutting-edge research centre in Harwell, Oxfordshire.



The new Moderna Innovation and Technology Centre is the latest addition to the UK's £100 billion life sciences sector to come out of Moderna's 10-year strategic partnership with the government. The partnership also includes an over £1 billion programme of investment in UK R&D being brought forward by the company, tapping into the UK's deep skills base, world-class academic institutions, and the power of the NHS as an engine for research and innovation.

The facility will support around 150 highly-skilled jobs, while enabling groundbreaking work on the use of mRNA vaccine technology to tackle cancer and seasonal infections like flu, RSV and COVID-19, subject to the usual regulatory processes. RNA technology is an emerging field of healthcare with the potential to overhaul how we tackle a wide range of diseases that currently cause ill-health and distress.

It could also be utilised to produce up to 250 million vaccine doses a year in the event of a pandemic, making a critical contribution to national resilience to future health emergencies, by building the UK's capacity to produce cutting-edge therapies at scale, at home, should they be needed.

To support further game-changing investments in life sciences R&D, the government is piloting the Life Sciences Transformational R&D Investment Fund. This new £50 million fund aims to bolster the UK's research and development infrastructure by supporting large-scale, capital-intensive projects in the life sciences sector worth over £100 million.

The fund is designed to attract substantial private investment in UK-based R&D, reinforcing the UK's position as the location of choice for life sciences innovation, and pulling in the investment that is a vital part of economic growth.

Science Minister Lord Vallance said: "The UK's life sciences sector is a fundamental part of our nation's health and economy, and will be critical to our ambitions going forward. The pioneering work Moderna will be doing, here in the UK, on mRNA is a prime example of the opportunity we want to grasp. It's the chance for new treatments that save lives, and new products that drive growth and job creation, all made possible thanks to our world-class institutions, skills, and the NHS. It is this unique mix that makes the UK a life sciences destination that punches well above its weight. The opening of this centre is proof of the brilliant things coming out of the deep and meaningful relationships we're determined to forge with companies like Moderna. This Innovation Centre will strengthen the UK's growth, health, and our resilience."

Professor Susan Hopkins, CEO, UK Health Security Agency (UKHSA) said: "The opening of this impressive state of the art facility in Harwell, alongside Moderna's substantial investment in UK research and development, will help ensure the NHS has rapid access to mRNA vaccines if needed in a future pandemic. It will also open doors to new vaccine products, helping support the government's 10 Year Health Plan and the focus on prevention. I'm delighted that UKHSA leads the partnership on behalf of government. It is an exemplar on how we and government can work closely with industry to prepare and respond to emerging health threats. It will support the UK's ambitions towards the 100 Days Mission, helping put the UK at the forefront of pandemic preparedness and new technologies whilst driving economic growth."

Moderna is best-known for developing one of the first COVID jabs put to use during the pandemic. These vaccines were based on messenger RNA technology: an emerging field of healthcare with enormous potential – ranging from cancer therapies currently being trialled on the NHS, to new vaccines for seasonal infections like flu and norovirus.

Moderna are also investing in UK R&D more broadly – ranging from clinical trials, to backing for postgraduate researchers. The company was one of the largest industry sponsors of clinical trials in the UK in 2024, and its investment is helping to get a larger number and broader range of volunteers involved in clinical trials, as part of the UK Vaccine Innovation Pathway. Moderna's 10-year strategic partnership with the government is managed by the UKHSA.

Fuente: PATHOLOGY IN PRACTICE. Disponible en <https://goo.su/E2livU>

Five-Fold Shield: The Latest Advancements in Meningococcal Vaccine Technology

Oct 10. In the realm of public health, the continuous evolution of vaccines plays a pivotal role in safeguarding communities against various infectious diseases. One such groundbreaking development is the introduction of a new meningococcal vaccine that takes immunization to the next level. This vaccine targets not one or two, but an impressive five bacterial strains in a single shot, marking a significant stride in the fight against meningococcal disease. **Understanding Meningococcal Disease:** Meningococcal disease is a serious bacterial infection caused by *Neisseria meningitidis*. It can manifest as meningitis (inflammation of the membranes surrounding the brain and spinal cord) or septicemia (bloodstream infection). This disease can progress rapidly and lead to severe complications, making vaccination a crucial preventive measure. **The Challenge of Multiple Strains:** Traditionally, meningococcal vaccines have focused on specific strains, providing protection against the most prevalent ones. However, the emergence of new strains and the variability in regional

prevalence pose challenges in designing effective vaccines. The new breakthrough addresses this challenge by targeting five bacterial strains in a single vaccine, offering a more comprehensive defense against the diverse nature of meningococcal infections.

Key Features of the New Meningococcal Vaccine

1. **Broad Spectrum Protection:** Unlike previous vaccines that focused on individual strains, this new vaccine covers a spectrum of five bacterial strains, providing a more inclusive defense mechanism.
2. **Reduced Dosage Frequency:** The consolidation of multiple strains into one vaccine means fewer shots are needed for comprehensive protection. This not only simplifies vaccination schedules but also improves adherence to immunization programs.
3. **Enhanced Immune Response:** The formulation of the vaccine is designed to stimulate a robust immune response against each targeted strain, ensuring a higher level of protection compared to previous vaccines.
4. **Global Impact:** Given the vaccine's ability to target multiple strains, it holds promise for global vaccination campaigns, especially in regions where meningococcal disease is prevalent with varying strains.



Potential Challenges and Considerations: While the new meningococcal vaccine represents a significant leap forward, there are considerations to address. These may include:

- regulatory approvals
- distribution logistics
- ongoing surveillance

to monitor the long-term effectiveness and safety of the vaccine. Click [here](#) to see the full scientific article from [Very Well Health](#). The development of a meningococcal vaccine targeting five bacterial strains in one shot is a remarkable achievement in the field of vaccinology. It not only addresses the challenges posed by diverse strains but also offers a more streamlined approach to immunization. As this breakthrough progresses through clinical trials and regulatory processes, it brings hope for a future where meningococcal disease can be more effectively controlled on a global scale, potentially saving countless lives.

Fuente: Asher Longevity Institute ALIVE. Disponible en <https://goo.su/h9wu9b>

Pfizer (PFE) Faces Challenges Amid Drug Pricing Push

Oct 11. Pfizer (PFE) experienced a significant drop last week, with its stock falling 9.41% as the company navigates challenges amid ongoing U.S. efforts to reform drug pricing. The pharmaceutical giant is part of a new direct-to-consumer platform, TrumpRx, which aims to offer sharp discounts on prescription medicines. Meanwhile, President Trump and Commerce Secretary Howard Lutnick's families could see financial gains from these initiatives, according to reports. The recent developments have sparked considerable movement within the healthcare sector.

Pfizer Inc, one of the world's largest pharmaceutical firms, generates annual sales of approximately \$60 billion. Historically, the company sold a variety of healthcare products and chemicals, but now prescription

drugs and vaccines constitute the majority of its sales. Key products include the pneumococcal vaccine Prevnar 13, cancer drug Ibrance, and cardiovascular treatment Eliquis. Pfizer's global reach is significant, with international sales accounting for 40% of total sales, and emerging markets playing a crucial role.

With a market capitalization of \$140.89 billion, Pfizer operates within the healthcare sector, specifically in the drug manufacturers industry. The company's current stock price is \$24.78, reflecting recent market volatility.

Financial Health Analysis

Pfizer's financial metrics present a mixed picture:

- Revenue for the trailing twelve months stands at \$63.83 billion, with a 3-year revenue growth rate of -7.8%, indicating a decline.
- Profitability remains robust with an operating margin of 25.93% and a net margin of 16.84%.
- The gross margin is 72.12%, although it has been in long-term decline at an average rate of -4% per year.
- On the balance sheet, Pfizer maintains a current ratio of 1.16 and a debt-to-equity ratio of 0.69, indicating moderate leverage. The Altman Z-Score of 2.08 places the company in a grey area, suggesting some financial stress.

Valuation & Market Sentiment

Pfizer's valuation metrics are as follows:

- The P/E ratio is 13.18, close to its 2-year low, suggesting potential undervaluation.
- The P/S ratio is 2.21, near a 10-year low, further indicating a favorable valuation.
- Analyst target price is \$29.21, with a recommendation score of 2.5, reflecting a hold position.

Technical indicators show a 14-day RSI of 46.64, suggesting the stock is neither overbought nor oversold. Institutional ownership stands at 65.5%, with insider ownership at 0.28%.

Risk Assessment

Pfizer's financial health is supported by a Piotroski F-Score of 7, indicating a healthy situation. The Beneish M-Score of -2.52 suggests the company is unlikely to be a manipulator.

Sector-specific risks include regulatory changes in drug pricing and competitive pressures in the pharmaceutical industry. The stock's beta of 0.37 indicates lower volatility compared to the market.

Overall, while Pfizer faces challenges in revenue growth and regulatory pressures, its strong profitability and favorable valuation metrics provide a balanced outlook for investors.

- Pfizer (PFE, Financial) stock experienced a significant decline of 9.41% last week amid U.S. drug pricing reform efforts.
- The company is involved in TrumpRx, a new platform aimed at providing discounted prescription medicines.
- Pfizer's financial health shows mixed signals with strong profitability but declining revenue growth.

Fuente: GURU FOCUS. Disponible en <https://n9.cl/20k6f>





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Patentes registradas en Patentscope

Estrategia de búsqueda: (Vaccine) AND DP:([01.10.2025 TO 12.10.2025]) 68 records.

1. WO/2025/199993 CHOLESTEROL-MODIFIED CATIONIC LIPOSOME TUMOR VACCINE, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 02.10.2025

Clasificación Internacional A61K 9/127Nº de solicitud PCT/CN2024/084956 Solicitante SICHUAN UNIVERSITY Inventor/a WEI, Xiawei

The present invention belongs to the technical field of cancer immunotherapy, and particularly relates to a cholesterol-modified cationic liposome tumor vaccine, a preparation method therefor, and use thereof. In order to solve the problems of poor targeting and strong side effects of TLR agonists in anti-tumor treatment, the present invention provides a cationic liposome prepared from a cholesterol-modified 1V209 molecule, a cationic lipid component, cholesterol, and DSPE-PEG2000, and then the cationic liposome and ovalbumin 5 form the tumor vaccine by electrostatic adsorption. Animal experiments show that the vaccine can induce antigen-specific CD8+ T cells, activate lymphocytes, and generate stronger antigen cross-presentation, more memory T cells, antibodies, and cytokines. Prophylactic inoculation with the vaccine can significantly delay the progression of mouse melanoma and lymphoma and prolong the survival of mice. The combination use of the vaccine and a PD-1 checkpoint inhibitor can further enhance the anti-tumor effect. Therefore, the vaccine is a promising cancer vaccine.

2. WO/2025/211492 IMMUNE ADJUVANT COMPRISING IMIQUIMOD AND NOD2 RECEPTOR-STIMULATING COMPONENT OR BACTERIUM

WO - 09.10.2025

Clasificación Internacional A61K 39/39Nº de solicitud PCT/KR2024/006650 Solicitante REPUBLIC OF KOREA(DEFENSE ACQUISITION PROGRAM ADMINISTRATION) Inventor/a KIM, Donghyun

The present invention relates to a vaccine immune adjuvant that utilizes the synergy between imiquimod and a Nod2 receptor-stimulating component or a Nod2 receptor-stimulating bacterium. According to one aspect of the present invention, a vaccine immune adjuvant composition comprising, as active ingredients, imiquimod and a Nod2 receptor-stimulating component or a Nod2 receptor-stimulating bacterium can enhance the efficacy of a vaccine to be greater than that of the administration of imiquimod alone, and thus can be effectively used as a complex immune adjuvant for increasing immune responses of various vaccines such as an SARS-CoV-2 virus vaccine, an influenza A virus vaccine and an anticancer vaccine against melanoma.

3. 20250302932 COMBINATION OF VACCINATION AND OX40 AGONISTS

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18952886 Solicitante CureVac SE Inventor/a Mariola FOTIN-MLECZEK

The present invention relates to a vaccine/agonist combination comprising an RNA vaccine comprising at least one RNA comprising at least one open reading frame (ORF) coding for at least one antigen and a composition comprising at least one OX40 agonist. The present invention furthermore relates to a pharmaceutical composition and a kit of parts comprising the components of such a vaccine/agonist combination. Additionally the present invention relates to medical use of such a vaccine/agonist combination, the pharmaceutical composition and the kit of parts comprising such a vaccine/agonist combination, particularly for the prevention or treatment of tumor or cancer diseases or infectious diseases. Furthermore, the present invention relates to the use of an RNA vaccine in therapy in combination with an OX40 agonist.

4. WO/2025/201659 COMBINATION THERAPY COMPRISING THERAPEUTIC RNA CANCER VACCINES AND ANTI-CTLA4 ANTIBODIES FOR CANCER TREATMENT

WO - 02.10.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2024/058693 Solicitante BIONTECH SE Inventor/a SAHIN, Ugur

The invention provides an RNA cancer vaccine and/or an anti-CTLA4 antibody for use in a method of treating cancer in a subject in need thereof, wherein the subject is administered a combination comprising the RNA cancer vaccine comprising at least one RNA and the anti-CTLA4 antibody. The invention further provides a compositions or kit of parts comprising the RNA cancer vaccine comprising the at least one RNA and the anti-CTLA4 antibody. The invention further provides methods of treating cancer in a subject in need thereof, wherein the method comprises administering to the subject the RNA cancer vaccine comprising the at least one RNA and the anti-CTLA4 antibody. The invention further provides a combination of the RNA cancer vaccine comprising at least one RNA and the anti-CTLA4 antibody.

5. WO/2025/202495 COMBINATION THERAPY COMPRISING THERAPEUTIC RNA CANCER VACCINES AND ANTI-CTLA4 ANTIBODIES FOR CANCER TREATMENT

WO - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/EP2025/058630 Solicitante BioNTech SE Inventor/a SAHIN, Ugur

The invention provides an RNA cancer vaccine and/or an anti-CTLA4 antibody for use in a method of treating cancer in a subject in need thereof, wherein the subject is administered a combination comprising the RNA cancer vaccine comprising at least one RNA and the anti-CTLA4 antibody. The invention further provides a compositions or kit of parts comprising the RNA cancer vaccine comprising the at least one RNA and the anti-CTLA4 antibody. The invention further provides methods of treating cancer in a subject in need thereof, wherein the method comprises administering to the subject the RNA cancer vaccine comprising the at least one RNA and the anti-CTLA4 antibody. The invention further provides a combination of the RNA cancer vaccine comprising at least one RNA and the anti-CTLA4 antibody. The invention further provides a combination of the RNA cancer vaccine comprising at least one RNA and the anti-CTLA4 antibody for use as a medicament.

6. WO/2025/212705 VACCINE TARGETING TO MUCOSAL LYMPHOID TISSUES IN THE GASTROINTESTINAL TRACT

WO - 09.10.2025

Clasificación Internacional A61K 9/10Nº de solicitud PCT/US2025/022620Solicitante MASSACHUSETTS INSTITUTE OF TECHNOLOGYInventor/a IRVINE, Darrell, J.

Viruses, bacteria, and parasites frequently cause infections in the gastrointestinal (GI) tract, but traditional vaccination strategies typically elicit little or no mucosal antibody responses. Disclosed herein is a **vaccine** comprising a nanoemulsion and an immunogen non-covalently conjugated to the surface of the nanoemulsion by an amphiphilic linker, wherein the **vaccine** is suitable for inducing a mucosal antibody response in the gastrointestinal tract. Also disclosed are methods of using the **vaccine** to immunize a subject by intraperitoneal administration of an effective amount of the **vaccine**, alone or with an adjuvant.

7.20250314636METHOD OF PREPARING AND EXPANDING A POPULATION OF IMMUNE CELLS FOR CANCER THERAPY, POTENCY ASSAY FOR TUMOR RECOGNITION, BIOLOGICAL **VACCINE** PREPARATION AND EPITOPETARGET FOR ANTIBODIES

US - 09.10.2025

Clasificación Internacional G01N 33/50Nº de solicitud 18864625Solicitante FUNDAÇÃO D. ANNA DE SOMMER CHAMPALIMAUD E DR. CARLOS MONTEZ CHAMPALIMAUD - CEInventor/a Carolina MENDONÇA GORGULHO

The present invention relates to a method of preparing and expanding a population of immune cells, a potency assay for tumor recognition, a biological **vaccine** preparation to provide anti-tumor response or an antiviral response for cancer therapy and epitopes targets for antibodies which are useful for the construction of chimeric antigen receptors.

The present invention is based on the fact that private or commonly shared tumor-associated antigens or private target antigens can be recognized by clinically relevant immune cells. Such target antigens could be used to prepare a biological **vaccine** preparation to provide anti-tumor response or an antiviral response by expanding a certain set of T-cells or B-cells and boosting the immune response in cancer therapy.

The present invention guides the selection of viable target antigens in designing an anti-tumor **vaccine** to remove potentially harmful autoimmune responses or pro-tumorigenic immune responses and aids to select the biologically and clinically most relevant set of immune cells specifically directed against cancer cells harvested from tumor infiltrating lymphocytes or from different anatomical sites for the active cellular therapy of patients with cancer.

8.20250312315CANCER **VACCINE** COMPOSITIONS AND METHODS OF USE THEREOF

US - 09.10.2025

Clasificación Internacional A61K 31/4178Nº de solicitud 19084504Solicitante Colorado State University Research FoundationInventor/a Terry OPGENORTH

The disclosure provides a cancer **vaccine** composition comprising inactivated cancer cells and an adjuvant, wherein the inactivated cancer cells are incapable of replication. Also provided is a method for producing a cancer **vaccine** composition, the method comprising contacting cancer cells with light (e.g., UV light) in the presence of a photosensitizer (e.g., riboflavin).

9. 20250302939 **VACCINE** FOR HUMAN T-LYMPHOTROPIC VIRUS-1

US - 02.10.2025

Clasificación Internacional A61K 39/21Nº de solicitud 18864213Solicitante The United States of America, as represented by the Secretary, Department of Health and HumanInventor/a Genoveffa Franchini

Provided herein is a nucleic acid-based **vaccine** for human T-cell leukemia virus type 1 (HTLV-1). In some aspects, the **vaccine** includes a combination of nucleic acid molecules encoding HTLV-1 gag protein and one or both of Type A HTLV-1 Envelope (Env) and Type C HTLV-1 Env. In some aspects, the **vaccine** includes a combination of nucleic acid molecules encoding HIV-1 gag protein and one or both of Type A HTLV-1 Envelope (Env) and Type C HTLV-1 Env. When administered to a subject, the Env and Gag proteins are expressed in the host and form HTLV-1 virus-like particles (VLPs) that are secreted from cells within the host and elicit an immune response that inhibits HTLV-1 infection.

10. 4628166 IMPFSTOFFADJUVANS UND HERSTELLUNGSVERFAHREN DAFÜR UND VERWENDUNG DAVON

EP - 08.10.2025

Clasificación Internacional A61P 35/00Nº de solicitud 25195745Solicitante CHENGDU MAXVAX BIOTECHNOLOGY LLCInventor/a CHEN DEXIANG

A **vaccine** adjuvant, and a preparation method therefor and a use thereof. The **vaccine** adjuvant is a MA105 immunologic adjuvant, and comprises (1) QS-21: 50 µg/ml to 300 µg/ml; (2) Poly I:C: 400 µg/mL to 3000 µg/mL; and (3) lipid molecules constituting a vector, the vector being a mixture of a cationic liposome and a neutral liposome.

11. WO/2025/199604 RECOMBINANT **VACCINE** PROTEINS FOR THE PREVENTION OF AVIAN NECROTIC ENTERITIS

WO - 02.10.2025

Clasificación Internacional A61K 39/08Nº de solicitud PCT/CA2024/050383Solicitante UNIVERSITÉ DE MONTRÉALInventor/a GAUCHER, Marie-lou

Avian necrotic enteritis (NE) is a complex and multifactorial enteric disease of commercial chickens and turkeys that has major economic consequences on the poultry industry worldwide. There is thus a need for improved and efficient vaccines for the prevention of diseases associated to pathogenic *Clostridium perfringens* in the poultry industry. The present application discloses immunogenic polypeptides unique to pathogenic strains of *Clostridium perfringens* which causes avian NE, as well as **vaccine** compositions comprising one or more of the immunogenic polypeptides, immunogenic fragments thereof, or nucleic acids encoding such immunogenic polypeptides and fragments. Methods and uses of the **vaccine** compositions for inducing or eliciting an immune response against *Clostridium perfringens*, for preventing or treating infection by *Clostridium perfringens*, and/or for preventing or treating avian NE in an animal, such as a poultry, are also disclosed.

12. WO/2025/202803A A PROCESS FOR PREPARING LACTOBACILLUS-BASED RECOMBINANT **VACCINE** CANDIDATE AGAINST MULTIPLE SALMONELLA SEROVAR IN POULTRY

WO - 02.10.2025

Clasificación Internacional C12N 15/74Nº de solicitud PCT/IB2025/052370Solicitante AHMAD, Syed MudasiInventor/a AHMAD, Syed Mudasi

The process comprises providing a recombinant **vaccine** construct, wherein said construct comprises genetically modified *Lactobacillus plantarum* NC8 as a live vector; modifying the genetic structure of said *Lactobacillus plantarum* NC8 to express conserved *Salmonella* antigens, including PagN, SopE2, and FliC, anchored by a ptrk 892 backbone with a constitutive promoter, phosphoglycerate mutase (PGM); incorporating Signal Lp_2145 and cAM12 Anchor sequences into said genetic construct to enhance surface expression of recombinant proteins on *Lactobacillus plantarum* NC8; administering said recombinant **vaccine** orally to poultry, leveraging the probiotic properties of *Lactobacillus plantarum* NC8 for effective colonization of the poultry gastrointestinal tract; inducing a prolonged and intensified immune response by ensuring sustained high-level expression of target antigens through the utilization of the robust constitutive promoter, phosphoglycerate mutase (PGM); and optimizing immunogenicity through the surface expression of recombinant proteins on *Lactobacillus plantarum* NC8, fostering a robust and precisely targeted immune response.

13. 20250312433PROCESS FOR PRODUCING OF **VACCINE** FORMULATIONS WITH PRESERVATIVES
US - 09.10.2025

Clasificación Internacional A61K 39/09Nº de solicitud 18863176Solicitante PFIZER INC.Inventor/a Michael David BRUCHSALER

The present invention relates to a process for the production of a conjugate **vaccine** comprising a preservative. The invention relates in particular to a process for the production of a conjugate **vaccine** where the preservative is hydrophobic and viscous (such as 2-phenoxyethanol (2-PE)).

14. 4628095FUSIONSPROTEIN UND GRANULIERTES ANTIGEN DAMIT
EP - 08.10.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23887963Solicitante UNIV XIAMENInventor/a LI SHAOWEI

Provided are a fusion protein and a composition, a kit, a granulated antigen, a **vaccine**, and a pharmaceutical composition comprising same. The present invention also relates to use of the fusion protein and the composition, the kit, and the granulated antigen comprising same in the preparation of a pharmaceutical composition or a **vaccine**. The granulated antigen is particularly suitable for the production and vaccination of a **vaccine** and has advantages in the prevention and/or treatment of viral infections.

15. WO/2025/206323CELL THERAPY
WO - 02.10.2025

Clasificación Internacional A61K 48/00Nº de solicitud PCT/JP2025/012812Solicitante OSAKA UNIVERSITYInventor/a ISHII, Hideshi

The present disclosure provides a novel modality for a new target disease (e.g. pancreatic cancer). The present disclosure applies a mRNA CAR-T vaccine to CAR-T for a new target disease (e.g., pancreatic cancer). In the present disclosure, it has been revealed, by in vitro and in vivo experiments, that a mRNA CAR-T vaccine that targets FAP as an antigen can achieve a potent anti-tumor effect in in vivo editing. As for a mechanism for inhibiting the effect of a mRNA CAR-T vaccine, the pathways involving regulatory T cells and the like have also been revealed. As for a synergistic effect with an immune checkpoint agonist, a "condition" for achieving a potent anti-tumor effect in vivo has also been revealed through much trial and error.

16. 4626574 HERSTELLUNG UND HERSTELLUNG EINES FLAVIVIRUSIMPFSTOFFS IM GROSSMASSSTAB

EP - 08.10.2025

Clasificación Internacional B01D 15/36Nº de solicitud 23829266 Solicitante TAKEDA VACCINES INC Inventor/a SANTANGELO JOSEPH DAVID

The present invention provides methods for large-scale flaviviral vaccine production and manufacture. The methods provided herein are specifically contemplated for large-scale production and manufacture of live, attenuated flaviviral vaccines such as live, attenuated, dengue virus vaccines. Further, the methods provided herein pertain to formulation of live, attenuated, monovalent, divalent, trivalent, or tetravalent viral vaccine products.

17. WO/2025/210229 AN IMMUNOGENIC OR VACCINE COMPOSITION COMPRISING A LIVE ATTENUATED AVIAN METAPNEUMOVIRUS (AMPV) FOR USE IN THE PREVENTION AND/OR TREATMENT OF AN AMPV INFECTION

WO - 09.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/EP2025/059289 Solicitante HIPRA SCIENTIFIC, S.L.U. Inventor/a GUIRADO CACERES, Evelina

An immunogenic or vaccine composition comprising a live attenuated avian metapneumovirus (aMPV) for use in the prevention and/or treatment of an aMPV infection The present invention refers to an immunogenic or vaccine composition comprising a live attenuated avian metapneumovirus (aMPV) for use in the prevention and/or treatment of an aMPV infection, wherein said use comprises administering the composition to laying birds, or for use in active immunization of laying birds.

18. 20250302896 VESICULAR STOMATITIS VIRUS MARBURG VIRUS VACCINE

US - 02.10.2025

Clasificación Internacional A61K 35/76Nº de solicitud 19063559 Solicitante International AIDS Vaccine Initiative, Inc. Inventor/a Christopher L. PARKS

The present invention relates to a vesicular stomatitis virus vaccine vector encoding a MARV glycoprotein (rVSVΔG-MARV-GP). Vaccination with as little as 200 plaque-forming units was 100% efficacious against MARV lethality and prevented development of viremia. rVSVΔG-MARV-GP vaccination induced MARV GP-specific serum IgG, and virus-neutralizing activity in serum was detectable in animals vaccinated with the highest doses.

19. 20250312428 HETEROLOGOUS PRIME BOOST VACCINE

US - 09.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 19237140 Solicitante BOEHRINGER INGELHEIM INTERNATIONAL GMBH Inventor/a Guido WOLLMANN

The present invention pertains to the provision of a vaccine comprising a first component (K) and a second component (V), wherein the first component (K) comprises a complex in which a cell penetrating peptide, an antigenic domain and a TLR agonist are functionally linked and the second component (V) comprises an oncolytic recombinant vesicular stomatitis virus expressing an antigenic domain. The invention further pertains to the use of the inventive vaccine in the treatment of cancer. The invention also provides a recombinant vesicular stomatitis virus expressing an antigenic domain and its use in cancer vaccines.

20. 4627580 VERFAHREN ZUR BEWERTUNG DER AUFFÄLLIGKEIT EINES EPITOPS GEGENÜBER DEM REPERTOIRE VON T-ZELL-REZEPTOREN

EP - 08.10.2025

Clasificación Internacional G16B 20/00Nº de solicitud 23817365 Solicitante IMMUNEWATCH BV Inventor/a MEYSMAN PIETER

The current invention relates to a method of predicting T-cell response to a query epitope with a known epitope-TCR binding comprising calculating conspicuousness score of said epitope based on the number of known TCR sequences or TCR clusters responsive to said epitopes or by a centrality metric for each said epitopes in the epitope-TCR graph. The invention further relates to a second method to predict T-Cell response of any arbitrary epitopes using a machine learning algorithm. The T-cell response of epitopes can be used for the selection of molecules for vaccine and/or non-immunogenic compositions. The invention further relates to a third method for producing vaccine, to a vaccine, a biological and a data base.

21. 4626405 PNEUMOKOKKENKONJUGAT-IMPFFSTOFFFORMULIERUNGEN

EP - 08.10.2025

Clasificación Internacional A61K 9/08Nº de solicitud 23817825 Solicitante PFIZER Inventor/a DIOP AWA

The present invention relates to new vaccine formulations comprising conjugated *Streptococcus pneumoniae* capsular saccharide antigens (glycoconjugates) and uses thereof. Vaccine formulations of the present invention will typically comprise at least one glycoconjugate from a *S. pneumoniae* serotype in a formulation designed to facilitate resuspension.

22. WO/2025/207938 NUCLEIC ACID VACCINES AGAINST COCCIDIOIDOMYCOSIS

WO - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/US2025/021841 Solicitante UNIVERSITY OF WASHINGTON Inventor/a FULLER, James Thomas

A trivalent vaccine induces robust antibody and mucosal and systemic IFN- γ and Th17 T cell responses against *Coccidioides* and affords complete protection from fungal dissemination and disease. The combination of the three antigens of this trivalent vaccine can exert synergistic protection against lethality,

disease and fungal dissemination. These robust responses can be further enhanced by use of one or more additional antigens described herein.

23. 20250312434 LIPOPOLYSACCHARIDE (LPS) DEFICIENT ACINETOBACTER BAUMANNII MULTIVALENT **VACCINE**

US - 09.10.2025

Clasificación Internacional A61K 39/104Nº de solicitud 18702767 Solicitante VAXDYN S.L. Inventor/a Juan José INFANTE VIÑOLO

The invention refers to a composition comprising inactivated cells deficient in LPS from the genus *Acinetobacter* and/or outer membrane vesicles from the same and their use for the manufacture of a medicament, preferably a **vaccine**, for the prevention of diseases caused by *K. pneumoniae*, *P. aeruginosa*, *E. coli*, and/or *A. pleuropneumoniae*, and optionally *A. baumannii*.

24. 20250302941 APPLICATION OF AN IMMUNOGENIC OR **VACCINE** COMPOSITION

US - 02.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18283067 Solicitante EYE VACC Inventor/a Joël Blomet

The present invention relates to an immunogenic or **vaccine** composition comprising a marker, for use thereof in prevention and/or treatment of an infection caused by at least one pathogen, characterized in that it is administered on the ocular mucosa and/or the urogenital mucosa.

25. WO/2025/200576 RECOMBINANT RSV F PROTEIN AND USE THEREOF

WO - 02.10.2025

Clasificación Internacional C07K 19/00Nº de solicitud PCT/CN2024/137843 Solicitante XIAMEN UNIVERSITY Inventor/a ZHENG, Zizheng

Provided are a recombinant respiratory syncytial virus (RSV) fusion (F) protein, and a fusion protein, **vaccine**, immunogenic composition, kit and pharmaceutical composition comprising same. Further provided is a use of the recombinant RSV F protein, fusion protein, **vaccine**, immunogenic composition, kit and pharmaceutical composition in preventing and/or treating RSV infection or diseases and/or symptoms caused by RSV infection.

26. 4622666 ABGESCHWÄCHTES AFRIKANISCHES SCHWEINEPESTVIRUS UND VERWENDUNG DAVON IN IMPFSTOFFZUSAMMENSETZUNGEN

EP - 01.10.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23809585 Solicitante INTERVET INT BV Inventor/a VAN DEN BORN ERWIN

The present invention relates to African Swine Fever (ASF) attenuated viruses wherein the genes EP402R and EP153R have been inactivated, which can be used as a **vaccine**. The attenuated viruses protect pigs against subsequent challenge with virulent virus. The present invention also relates to the use of such attenuated viruses to treat and/or prevent ASF.

27. WO/2025/213131 RNA COMPOSITIONS FOR DELIVERY OF ORTHOPOX ANTIGENS AND RELATED METHODS

WO - 09.10.2025

Clasificación Internacional A61K 39/275Nº de solicitud PCT/US2025/023311 Solicitante BIONTECH SE Inventor/a ZUIANI, Adam

The present disclosure provides pharmaceutical compositions for delivery of orthopoxvirus antigens (e.g., mpox, variola, borealpox, ectromelia, cowpox, volepox, modified vaccinia virus Ankara, or vaccinia) and related technologies (e.g., components thereof and/or methods relating thereto). For example, the present disclosure provides polyribonucleotides encoding one or more mpox antigens, fragments, or fusions thereof.

28. WO/2025/202543A **VACCINE** COMPOSITION COMPRISING AN ADENOVIRAL VECTOR AND AN ADENOVIRAL VECTOR DELIVERY ENHANCING AGENT

WO - 02.10.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/FI2025/050148 Solicitante ROKOTE LABORATORIES FINLAND OY Inventor/a YLÄ-HERTTUALA, Seppo

The present invention is directed to a **vaccine** composition comprising i) an adenoviral vector delivery enhancing agent selected from the group consisting of SYN3, and SYN3 analogs, and ii) an adenoviral vector comprising, at an insertion site in the viral genome, a nucleic acid sequence encoding an immunogenic protein generating adaptive immune response upon exposure to a host organism. The present invention is also directed to a method for delivering an adenoviral vector to a mammalian subject, said method comprising a step of administering onto the nasal and/or oral mucosa of the subject a dose of a delivery enhancing agent selected from the group consisting of SYN3 and SYN3 analogs and a dose of an adenoviral vector, wherein the first dose of said delivery enhancing agent and the first dose of said adenoviral vector are administered simultaneously or sequentially.

29. 20250302931 COLORECTAL CANCER TUMOR CELL VACCINES

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18947958 Solicitante NEUVOGEN, INC. Inventor/a Bernadette Ferraro

The present disclosure provides an allogeneic whole cell cancer **vaccine** platform that includes compositions and methods for treating and preventing colorectal cancer. Provided herein are compositions containing a therapeutically effective amount of cells from one or more cancer cell lines, some or all of which are modified to (i) inhibit or reduce expression of one or more immunosuppressive factors by the cells, and/or (ii) express or increase expression of one or more immunostimulatory factors by the cells, and/or (iii) express or increase expression of one or more tumor-associated antigens (TAAs), including TAAs that have been mutated, and which comprise cancer cell lines that natively express a heterogeneity of tumor associated antigens and/or neoantigens, and/or (iv) express one or more tumor fitness advantage mutations, including but not limited to driver mutations. Also provided herein are methods of making and preparing the colorectal cancer **vaccine** compositions and methods of use thereof.

30. WO/2025/208217 A RECOMBINANT BACTERIAL STRAIN AS TUBERCULOSIS **VACCINE** CANDIDATE, AND METHODS OF PRODUCING THEREOF

WO - 09.10.2025

Clasificación Internacional C12N 15/74Nº de solicitud PCT/CA2025/050473 Solicitante UNIVERSITY OF SASKATCHEWAN Inventor/a CHEN, Jeffrey Matthew

The invention describes a recombinant Mycobacterium strain such as *M. smegmatis* that is recombinantly converted to express the *M. tb* complex ESX protein expression system, e.g. *M. tb* complex ESX-1 protein expression system. The recombinant strain exports virulence effector proteins (such as EspA, EspB, EspC, EsxA or EsxB) similar to *M. tb*, when administered to a subject, and is therefore useful as a potential tuberculosis **vaccine** candidate. The recombinant strain is non-pathogenic and the vaccinated subject does not give a false positive result when tested with any standard tuberculosis diagnostic test e.g. CCT, as the recombinant strain does not sensitize the vaccinated host to *M. tb* antigens or protein derivate of *M. bovis*.

31. 4628096 REKOMBINANTER EXPRESSIONSVEKTOR ZUR HERSTELLUNG VON VIRUSÄHNLICHEN PARTIKELN ODER NANOPARTIKELN DER FUSS- UND MUNDERKRANKUNG UND IMPFSTOFFZUSAMMENSETZUNG DAMIT

EP - 08.10.2025

Clasificación Internacional A61K 39/135Nº de solicitud 23895042 Solicitante OPTIPHARM CO LTD Inventor/a LEE JI HOON

The present invention relates to a recombinant expression vector for preparing foot-and-mouth disease virus-like particles or nanoparticles, and a **vaccine** composition using same. The present invention provides virus-like particles or nanoparticles, and a preparation method therefor, the particles or nanoparticles being produced by expressing, simultaneously, VP4 with VPI, VP2 and VP3 or expressing VP1, VP2 and VP3 while excluding VP4, from among FMDV structural proteins. The virus-like particles or nanoparticles, prepared according to the method of the present invention, can be effectively used in the prevention of diseases caused by foot-and-mouth disease viral infection.

32. WO/2025/202630 HEMORRHAGIC FEVER VACCINES

WO - 02.10.2025

Clasificación Internacional C07K 14/005Nº de solicitud PCT/GB2025/050635 Solicitante CAMBRIDGE ENTERPRISE LIMITED Inventor/a BILLMEIER, Martina

Designed polypeptide sequences are described, and their use as vaccines against pathogens which cause hemorrhagic fever, such as viruses of the Ebolavirus, Lassa Fever virus and Marburg virus families. The designed sequences include those which comprise the designed Tri-LEMvac **vaccine**, and the bivalent Lassa virus **vaccine**. Nucleic acid molecules encoding the polypeptides, as well as vectors, fusion proteins, pseudotyped virus particles, pharmaceutical compositions, combined preparations, cells, and their use as vaccines against viruses of the Ebolavirus, Lassa Fever virus and Marburg virus families are also described.

33. WO/2025/200475 SYSTEM AND METHOD FOR PREPARING OUTER MEMBRANE VESICLES WITH GUARANTEED IMMUNE EFFECTS

WO - 02.10.2025

Clasificación Internacional C12P 1/04Nº de solicitud PCT/CN2024/131238 Solicitante HUBEI UNIVERSITY OF MEDICINE Inventor/a HAO, Mingyue

Provided are a system and method for preparing outer membrane vesicles with guaranteed immune effects. The system comprises a strain acquisition module, a strain screening module, an outer membrane vesicle yield promotion module, an outer membrane vesicle extraction module, an outer membrane vesicle purification module, an immune efficacy evaluation module, and a yield evaluation module, wherein the strain acquisition module is connected to the strain screening module, the outer membrane vesicle yield promotion module, and the outer membrane vesicle extraction module, respectively. By means of a genetic modification operation, the *alr* gene of *Brucella* is knocked out, which in turn improves the yield and immune efficacy of the outer membrane vesicles. Moreover, by means of strictly controlling the *Brucella* culture conditions, an inducing environment is established to induce *Brucella* to secrete outer membrane vesicles, thereby further improving the yield of outer membrane vesicles, and in turn obtaining outer membrane vesicles with a high yield and good immune effects, which meets the requirements of large-scale vaccine production, and provides a high-quality vaccine adjuvant for the research into, development of and application of *Brucella* vaccines.

34. 4628510 CHIMÄRER MULTIANTIGEN-POXVIRUS-IMPfstoff UND VERWENDUNG DAVON

EP - 08.10.2025

Clasificación Internacional C07K 19/00Nº de solicitud 23896693 Solicitante INST MICROBIOLOGY CAS Inventor/a GAO FU

The present application relates to a multi-immunogen chimeric or mixed antigen for a poxvirus, particularly monkeypox virus, related products, and preparation method and use thereof. The chimeric or mixed antigen of the present application comprises three immunogens: (1) a monkeypox virus A35R protein or an antigenic fragment thereof (or a peptide fragment derived therefrom), (2) a monkeypox virus M1R protein or an antigenic fragment thereof (or a peptide fragment derived therefrom), and (3) a monkeypox virus B6R protein or an antigenic fragment thereof (or a peptide fragment derived therefrom), and can elicit an immune response against two types of infectious virus particles, namely intracellular mature virus particles (IMV) and extracellular envelope virus particles (EEV), thereby efficiently inducing specific immunoprotective effects against the monkeypox virus. Moreover, the poxvirus vaccine of the present application features good safety, rapid response, and scalable production capacity, thus showing excellent clinical application prospects.

35. 20250302942 VACCINE COMPOSITION AGAINST CORONAVIRUS

US - 02.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18553042 Solicitante LG CHEM, LTD. Inventor/a Yoon Ae NOH

The present invention relates to: a conjugate in which a receptor-binding domain (RBD) of coronavirus and a hepatitis B surface antigen (HBsAg) are linked; virus-like particles including the conjugate; and a coronavirus **vaccine** composition containing the conjugate and/or virus-like particles.

36. 4626470 IMPFSTOFFE MIT NEUEN NANOPARTIKELGERÜSTEN

EP - 08.10.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23898663 Solicitante SCRIPPS RESEARCH
INST Inventor/a HE LINLING

The present invention provides novel engineered nanoparticle scaffold sequences that are derived from the 13-01 protein. Relative to the known 13-01 protein or variants thereof, the novel 13-01 derived scaffold sequences of the invention contain an extended N-terminal helix. Also provided in the invention are **vaccine** constructs that contain various immunogenic proteins displayed on the novel nanoparticle scaffold sequences described herein. The **vaccine** constructs of the invention include, e.g., nanoparticles displaying tandem repeats of influenza M2e proteins or HCV E2 core proteins.

37. 20250302944 COMBINATION OF EPITOPES AND USE THEREOF, **VACCINE** CONSTRUCT, METHOD OF INDUCING AN IMMUNE RESPONSE, METHOD FOR THE IDENTIFICATION OF EPITOPES

US - 02.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18850231 Solicitante FUNDAÇÃO ZERBINI Inventor/a Jorge Elias KALIL FILHO

COMBINATION OF EPITOPES AND USE THEREOF, **VACCINE** CONSTRUCT, METHOD OF INDUCING AN IMMUNE RESPONSE, METHOD FOR THE IDENTIFICATION OF EPITOPES The present invention refers to a combination of epitopes comprising at least eight T cell epitopes from the SARS-CoV-2, as well as the use of said combination ("set of epitopes"). Said epitopes are widely recognized by CD4+ T-lymphocytes of the overwhelming majority of COVID-19 convalescent individuals.

38. 20250302935 DETECTION OF A GENETIC FUSION OR DELETION THAT RESULTS IN EXPRESSION OF A NEOANTIGEN

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 19067594 Solicitante Flagship Pioneering Innovations VI, LLC Inventor/a Anthony P. Shuber

The invention provides methods of detecting a sequence modification (e.g., a genetic fusion or deletion) associated with cancer development that results in expression of a neoantigen. The neoepitope serves as the basis for manufacture of a **vaccine**, which is administered to a subject to induce an immune response against those cells producing the neoantigen.

39. 20250302938 RECOMBINANT SUBUNIT BASED UNIVERSAL INFLUENZA AND RESPIRATORY VIRUS VACCINES

US - 02.10.2025

Clasificación Internacional A61K 39/145Nº de solicitud 18863015 Solicitante GEORGIA STATE UNIVERSITY RESEARCH FOUNDATION, INC. Inventor/a Sang-Moo KANG

In accordance with the purpose(s) of the present disclosure, as embodied and broadly described herein, the disclosure relates to universal influenza vaccines and methods of making the same. Also disclosed is method for vaccinating a subject for influenza A that involves administering a cross-protective influenza vaccine disclosed herein to a subject in need thereof by intranasal, intramuscular, subcutaneous, transdermal, or sublingual administration.

40. 20250312427 MRNA VACCINES COMPRISING IL-4 AND/OR IL-13 RNA AND USES THEREOF

US - 09.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18685004Solicitante NEOVACSIInventor/a Beatrice DROUET

An mRNA vaccine including at least one RNA molecule that encodes at least one cytokine (preferably IL-4, IL-13 or fragments thereof) and at least one T cell epitope, for treating or preventing disorders associated with aberrant IL-4 and/or IL-13 expression or activity, in particular asthma, atopic dermatitis and allergic disorders.

41. 20250312439 VACCINE FOR PREVENTING OR TREATING CORONAVIRUS INFECTION

US - 09.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18866792Solicitante PHION THERAPEUTICS LTDInventor/a Helen MCCARTHY

The present specification relates to vaccines comprising an mRNA polynucleotide encoding a coronavirus antigen; and an amphipathic cell penetrating RALA peptide.

42. WO/2025/207952 MATERIALS AND METHODS FOR TREATMENT OF MELANOMAS AND OTHER CANCERS

WO - 02.10.2025

Clasificación Internacional A61K 39/09Nº de solicitud PCT/US2025/021863Solicitante TUHURA BIOSCIENCES, INC.Inventor/a LAWMAN, Michael J.P.

The present disclosure generally relates to DNA vaccine compositions and methods of use thereof to enhance an immune response against cancer, and in particular to methods of enhancing an immune response to immune checkpoint inhibitor resistant cancers and rendering the cancer susceptible to immune checkpoint inhibitor therapies.

43. 20250312425 COMPOSITIONS AND METHODS COMPRISING ADA-1 FOR WOUND HEALING AND VACCINE ENHANCEMENT

US - 09.10.2025

Clasificación Internacional A61K 38/50Nº de solicitud 19172548Solicitante Drexel UniversityInventor/a Kara Lorraine Spiller

Compositions and methods incorporating Adenosine deaminase-1 (ADA-1) as an adjuvant are provided. Polynucleotides encoding novel coding sequences, polypeptides, vectors containing same, and methods of use are provided. Compositions and methods incorporating Adenosine deaminase-1 (ADA-1) in a hydrogel are also provided. Compositions and methods comprising ADA-1 for treating wounds are also provided.

44. WO/2025/211732 LIPID NANOPARTICLES COMPRISING GINSENOSE OR DERIVATIVE THEREOF, AND USE THEREOF

WO - 09.10.2025

Clasificación Internacional A61K 9/51Nº de solicitud PCT/KR2025/004311 Solicitante GENOMICTREE, INC. Inventor/a AN, Sungwhan

The present invention relates to lipid nanoparticles comprising ginsenoside or a derivative thereof, and use thereof, and, specifically, to: lipid nanoparticles containing vegetable ginsenoside or a derivative thereof included in ginseng or red ginseng, instead of animal cholesterol for the delivery of therapeutic or vaccine nucleic acids; gene-inorganic nanoparticles in which the corresponding lipid nanoparticles are encapsulated, and a preparation method therefor; and a nucleic acid delivery composition comprising same.

45. 2639923 SYSTEM AND METHOD FOR FORECASTING EFFECT OF PATHOGEN ON POPULATION

GB - 08.10.2025

Clasificación Internacional G16H 50/80Nº de solicitud 202404459 Solicitante AIRFINITY LTD Inventor/a MATTHEW DAVID LINLEY

A mathematical simulation model to the input data to generate output data providing a forecast indicative of an effect of the pathogen on the population, wherein the mathematical simulation model is configured to use at least a Susceptible-Infected-Recovered model (SIR model). The SIR model is applied to a historical part of the input data to generate an initial iteration of the output data, the SIR model trained on a recently-acquired part of the input data to improve an accuracy of the SIR model when generating a subsequent iteration of the output data. Vaccine forecast for mitigating pathogen impacts, Fourier computation and variations in infection rates over time may be used to predict epidemiological burden, hospital admissions and deaths. Input data may be acquired in part using sensor arrangements.

46. WO/2025/212898 PRE-FUSION STABILIZED CCHFV GP38-GN-GC GLYCOPROTEIN COMPLEX

WO - 09.10.2025

Clasificación Internacional C07K 14/005Nº de solicitud PCT/US2025/022968 Solicitante BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM Inventor/a MCLELLAN, Jason

Provided herein are engineered proteins comprising engineered CCHFV glycoprotein ectodomain. Such engineered CCHFV glycoprotein ectodomains may exhibit improved conformational homogeneity and/or biophysical stability. The ectodomains may be incorporated into nanoparticles or virus-like particles. Methods are also provided for use of the engineered CCHFV glycoprotein ectodomains as diagnostics, in screening platforms, and/or in vaccine compositions.

47. 20250302934 SET DOMAIN-CONTAINING 2 (SETD2) VACCINE

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18864271Solicitante INSTITUTE FOR CANCER RESEARCH D/B/A THE RESEARCH INSTITUTE OF FOX CHASE CANCER CENTERInventor/a Philip ABBOSH

Pharmaceutical compositions comprising one or more peptides derived from aberrantly translated retained introns (ATaRIs), methods for treating cancer with the same, and methods of immunizing a subject against cancer or eliciting an immune response to one or more peptides derived from ATaRIs are provided herein.

48.20250312436STABILIZATION OF VIRUS-BASED THERAPEUTIC AGENT

US - 09.10.2025

Clasificación Internacional A61K 39/205Nº de solicitud 19170865Solicitante ELAREX INC.Inventor/a Jeremy Andrew IWASHKIW

A virus-based active agent is mixed with trehalose and water and dried. The mixture may also contain one or more of carbohydrate, such as pullulan and albumin. The mixture may be dried to a moisture content of 0.1-10%. The drying may be under vacuum but optionally without producing a foam. Some or all of the drying may be at a temperature in the range of 15-400C, or at a temperature in the range of 1-15° C., or both. The active agent may be based on a vesicular stomatitis virus (VSV) or an adenovirus (AdV). The dried mixture may be stored at a temperature in the range of 1-55° C. A composition includes a virus, which may be a derived or modified form of a virus such as VSV or AdV. The composition may be used for a virus-based **vaccine**.

49.WO/2025/212893COMPOSITIONS AND METHODS FOR TREATING AND/OR PREVENTING LYME DISEASE

WO - 09.10.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/US2025/022962Solicitante THE MEDICAL COLLEGE OF WISCONSIN, INC.Inventor/a LOCHHEAD, Robert Bruce

Disclosed herein are compositions and methods for treating, and for the preventative treatment of Lyme disease. The compositions can include a conjugate **vaccine** polypeptide.

50.4628897VERFAHREN, ZUSAMMENSETZUNGEN UND SYSTEME ZUM NACHWEIS VON CORONAVIRUSNEUTRALISIERENDEN ANTIKÖRPERN

EP - 08.10.2025

Clasificación Internacional G01N 33/68Nº de solicitud 25181055Solicitante LABORATORY CORP AMERICA HOLDINGSInventor/a PETROPOULOS CHRISTOS J

The present disclosure relates to methods, compositions, and systems for detecting whether a subject exposed to a coronavirus has developed a neutralizing antibody response. Also disclosed are methods for determining whether a patient infected by a coronavirus is likely to respond to treatment with an antibody preparation. Also disclosed are methods for detecting the level of neutralizing antibody response in a sample of serum from a subject exposed to a coronavirus or to a coronavirus **vaccine**.

51.4626469COXIELLA BURNETTI-IMPFFSTOFF

EP - 08.10.2025

Clasificación Internacional A61K 39/02Nº de solicitud 23821335Solicitante MOREDUN RES INSTITUTEInventor/a MCNEILLY TOM

The disclosure provides a phase II Coxiella burnetii variant that can be used to raise protective immune responses. The disclosure further provides vaccines and compositions comprising the same as well as antigens and LPS derived therefrom, both of which may also be used to raise protective immune responses. The disclosure further relates to the treatment of disease and/or conditions caused or contributed to by Coxiella burnetii.

52.20250302943RE-FOCUSING PROTEIN BOOSTER IMMUNIZATION COMPOSITIONS AND METHODS OF USE THEREOF

US - 02.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18707713Solicitante King Abdullah University of Science and TechnologyInventor/a Magnus Albert Rueping

Booster immunization compositions and methods of use thereof, are provided. The disclosed compositions use adjuvanted proteins as a booster vaccine targeting functionally relevant pathogen B and T cell epitopes to re-focus a patient's adaptive antibody and cytotoxic T-cell response. The pharmaceutically active ingredient of the re-focusing boost vaccines include of one or several recombinant protein molecules in a full-length or truncated yet functional (meaning the overall antigenic tertiary structure is retained) version of disease-related protein or protein domain, a smaller subunit or specific or modified epitope or combination of shorter epitopes derived from that initial prime antigen sequence. Methods for re-focusing an immune response in a subject, to augment an existing (not sufficiently protective immune response) and effectively neutralize a pathogen of interest in a mammalian host organism include administering to a subject whose immune system has been primed by a previous infection/vaccination against the pathogen.

53.20250302837INHIBITORS OF CHYMASE FOR USE IN THE SELECTIVE RESOLUTION OF THROMBI IN THROMBOTIC OR THROMBOEMBOLIC DISORDERS

US - 02.10.2025

Clasificación Internacional A61K 31/513Nº de solicitud 18854816Solicitante SOCPRA SCIENCES SANTÉ ET HUMAINES S.E.C.Inventor/a Pedro D'ORLÉANS-JUSTE

The present invention covers the use of chymase inhibitors in general and more in particular substituted bicyclically substituted uracils of general formula (I) as described and defined herein, and 3-methylbenzo-[b]thiophene)-2-sulfonamido derivatives of general formula (II) for manufacturing pharmaceutical compositions for the treatment or prophylaxis of stroke, pulmonary embolism, deep or superficial vein thrombosis, thrombotic microangiopathy, thrombotic microangiopathy in hypercoagulable states after infection, inflammation, transplantation, disseminated intravascular coagulation, vaccine-induced immune thrombotic thrombocytopenia, vascular access site thrombosis or occlusion.

54.3615005COMPOSITIONS AND METHODS FOR VACCINE DELIVERY

PL - 06.10.2025

Clasificación Internacional A61K 9/127Nº de solicitud 18791678SolicitanteInventor/a SHEETIJ DUTTA

55. 20250312458 FORMULATIONS FOR NEOPLASIA VACCINES

US - 09.10.2025

Clasificación Internacional A61K 47/12Nº de solicitud 19074044 Solicitante The Broad Institute, Inc. Inventor/a EDWARD F. FRITSCH

The present invention relates to neoplasia vaccine or immunogenic composition formulation for the treatment or prevention of neoplasia in a subject.

56. WO/2025/210141 DELIVERY SYSTEM USEFUL FOR TOPICAL ADMINISTRATION OF RNA

WO - 09.10.2025

Clasificación Internacional A61K 9/107Nº de solicitud PCT/EP2025/059116 Solicitante BOEHRINGER INGELHEIM VETMEDICA GMBH Inventor/a SIGOILLOT-CLAUDE, Cecile

The present invention relates to a composition useful for the topical administration of RNA, wherein the composition comprises (i) an aqueous phase containing RNA and (ii) an oil phase, which in one embodiment are in mixture with two specific surfactants. Particularly, the present invention pertains to the formulation of mRNA encoding for an antigenic protein of interest. More particular, a microemulsion to cross the skin barrier is provided, which allows for alternative routes for mRNA vaccine applications. The microemulsion may be applied topically and can go through the skin via the transfollicular route. Thereby a delivery system is enabled which allows to (1) avoid the use of needles for vaccination and (2) target more specifically the mucosal component of immunity.

57. 20250302929 TUMOR NEOANTIGENIC PEPTIDES

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18549993 Solicitante INSTITUT CURIE Inventor/a Sebastian AMIGORENA

The present disclosure provides tumor neoantigenic peptide sequences and nucleotide sequences encoding such peptide sequences; a vaccine or immunogenic composition capable of raising a specific T-cell response comprising one or more of the neoantigenic peptides, or comprising nucleic acid encoding one or more of the neoantigenic peptides; an antibody, or an antigen-binding fragment thereof, a T cell receptor (TCR), or a chimeric antigen receptor (CAR) that specifically binds such neoantigenic peptides; methods of producing such antibodies, TCRs or CARs; polynucleotides encoding such neoantigenic peptides, antibodies, CARs or TCRs, optionally linked to a heterologous regulatory control sequence; immune cells that specifically bind to such neoantigenic peptides; and dendritic cells or antigen presenting cells that have been pulsed with one or more of the neoantigenic peptides; and methods of using such products in particular therapeutic uses of these products.

58. WO/2025/206402 AFRICAN SWINE FEVER VACCINE-PRODUCING RECOMBINANT ERI SILKWORM AND USE THEREOF

WO - 02.10.2025

Clasificación Internacional A01K 67/60Nº de solicitud PCT/JP2025/013137 Solicitante OKINAWA UKAMI SERICULTURE CO., LTD. Inventor/a MAEGAWA, Kenichi

The purpose of the present invention is to provide a method for producing an ASF antigen protein, said method having low endotoxin contamination and reduced workload in comparison with conventional methods. This purpose is achieved by a method using a recombinant eri silkworm that expresses an African swine fever virus antigen protein, in particular a recombinant eri silkworm that expresses an African swine fever virus antigen protein such as EP402R, i177L, P30, P54, pp62 and P72.

59. 20250304623 PEPTIDES FOR INTRACELLULAR DELIVERY

US - 02.10.2025

Clasificación Internacional C07K 7/08Nº de solicitud 18876364 Solicitante UNIVERSIDADE DE SANTIAGO DE COMPOSTELA Inventor/a Javier MONTENEGRO GARCÍA

The present invention relates to short peptides or salts thereof which can be modified with appropriate hydrophobic tails to generate amphiphilic molecules useful for the intracellular delivery of molecules of biological interest. Thus, the invention also relates to the amphiphilic molecules and to the complexes between said amphiphilic molecules and the molecules of biological interest. In addition, the invention refers to the use of said peptides, said amphiphilic molecules and said complexes for the delivery of molecules of biological interest. The invention relates to said peptides or salts thereof, said amphiphilic molecules and said complexes for use in medicine, particularly for use as a **vaccine**. Methods for the preparation of the amphiphilic molecules and the complexes of the invention are also contemplated. Finally, the invention relates to novel amino acids useful in the synthesis of the peptides and molecules of the invention.

60. 20250304995 MODIFIED PARAPOXVIRUS HAVING INCREASED IMMUNOGENICITY

US - 02.10.2025

Clasificación Internacional C12N 15/86Nº de solicitud 18258502 Solicitante PRIME VECTOR TECHNOLOGIES GMBH Inventor/a Ralf AMANN

The present invention relates to a modified *Parapoxvirus*, preferably a *Parapoxvirus* vector, having an increased immunogenicity, a biological cell containing said modified *Parapoxvirus*, a pharmaceutical composition, preferably a **vaccine**, containing said modified *Parapoxvirus* vector and/or said cell, and a new use of said modified *Parapoxvirus*.

61. WO/2025/201437 STING PROTEIN MUTANT, NUCLEIC ACID ENCODING STING PROTEIN MUTANT, AND USE OF STING PROTEIN MUTANT

WO - 02.10.2025

Clasificación Internacional C07K 14/47Nº de solicitud PCT/CN2025/085227 Solicitante RINUAGENE BIOTECHNOLOGY CO., LTD. Inventor/a ZHANG, Weiguo

Provided are a STING mutant protein, a nucleic acid encoding the protein, a vector, a composition or **vaccine** comprising the protein or nucleic acid or vector, and a use of the protein in stimulating an immune response or treating cancer.

62. 20250302933 ATTENUATED YET IMMUNOGENICALLY POTENTIATED TUMOR EXTRACELLULAR VESICLE COMPOSITIONS AND USES THEREOF

US - 02.10.2025

Clasificación Internacional A61K 39/00Nº de solicitud 19097584Solicitante KOREA INSTITUTE OF SCIENCE AND TECHNOLOGYInventor/a In-San KIM

The present disclosure relates to attenuated yet immunogenically potentiated tumor-derived extracellular vesicle compositions and uses thereof. The present disclosure was completed by confirming that the extracellular vesicles secreted by tumor cells are attenuated yet immunogenically potentiated through a process of treating tumor cells with verteporfin, and the extracellular vesicles of the present disclosure can be used as a **vaccine** composition, etc.

63.20250302940PATHOGEN-LIKE ANTIGEN-BASED **VACCINE** AND PREPARATION METHOD THEREFOR

US - 02.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18267379Solicitante RONGSEN BIOTECHNOLOGY (BEIJING) CO., LTDInventor/a Baidong HOU

The present application relates to a pathogen-like antigen (PLA) complex, and a preparation method therefor and an application thereof. The PLA complex consists of structurally-modified *Escherichia coli* bacteriophage virus-like particles (VLPs) and antigens displayed thereon, and nucleic acids are encapsulated inside the VLPs.

64.WO/2025/209940EBV **VACCINE** AND ANTIBODIES

WO - 09.10.2025

Clasificación Internacional A61P 31/20Nº de solicitud PCT/EP2025/058562Solicitante HELMHOLTZ ZENTRUM MUENCHEN DEUTSCHES FORSCHUNGSZENTRUM FÜR GESUNDHEIT UND UMWELT (GMBH)Inventor/a HAMMERSCHMIDT, Wolfgang

The present invention relates, inter alia, to a composition comprising the BILF2 glycoprotein of Epstein-Barr-Virus (EBV), a nucleic acid encoding the BILF2 glycoprotein of EBV, or an extracellular vesicle (EV) or EB-virus-like particle (VLP) displaying at its surface the BILF2 glycoprotein of EBV, for use in a method of eliciting antibodies against said BILF2 glycoprotein which reduce infectivity of EBV for human epithelial cells. Preferably, said composition elicits antibodies that prevent infection of human epithelial cells by EBV.

65.20250316334NEOANTIGEN IDENTIFICATION, MANUFACTURE, AND USE

US - 09.10.2025

Clasificación Internacional G16B 20/30Nº de solicitud 18431595Solicitante GRITSTONE BIO, INC.Inventor/a Thomas Boucher

Disclosed herein is a system and methods for determining the alleles, neoantigens, and **vaccine** composition as determined on the basis of an individual's tumor mutations. Also disclosed are systems and methods for obtaining high quality sequencing data from a tumor. Further, described herein are systems and methods for identifying somatic changes in polymorphic genome data. Finally, described herein are unique cancer vaccines.

66.WO/2025/208014HAPTEN CONJUGATE **VACCINE** AGAINST XYLAZINE

WO - 02.10.2025

Clasificación Internacional A61K 39/385Nº de solicitud PCT/US2025/021994Solicitante THE SCRIPPS RESEARCH INSTITUTEInventor/a JANDA, Kim, D.

The application discloses compositions comprising a xylazine hapten having the Formula (I). The application further discloses xylazine hapten compositions and vaccines against xylazine. The application further discloses methods of preventing xylazine overdose. The application further discloses methods of preventing use disorder of xylazine. The application further discloses methods of generating antibodies against xylazine. The application further discloses methods of treating consumption of a substance laced with xylazine.

67.20250312440CHEMICAL SCREEN OF MODULATORS FOR **VACCINE** ADJUVANTS

US - 09.10.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18869655Solicitante THE UNIVERSITY OF CHICAGOInventor/a Aaron ESSER-KAHN

The disclosure provides for methods and compositions that can be used for modulating an immune response. Described are methods comprising administering an effective amount of (a) an adjuvant; and (b) one or more of bafetinib (INNO-406), LY3009120, MK-8353 (SCH900353), amodiaquine, zanubrutinib (BOB-3111), Ku55933, Tucidinostat, PD318088, WNK463, and TRx0237 (LMTX) mesylate to a subject. The methods may be for the vaccination of subjects, or for the treatment or prevention of cancer, graft rejection, graft versus host disease, a bacterial infection, or a viral infection in a subject. The method may be for modulating an immune response in vivo, in vitro, or ex vivo. The methods also include immune activation of a population of immune cells in vitro or ex vivo. Also described is a method for identifying the efficacy of an adjuvant.

68.WO/2025/202674LANGERHANS CELLS TARGETING HIV-1 VACCINES

WO - 02.10.2025

Clasificación Internacional C07K 16/28Nº de solicitud PCT/IB2024/000163Solicitante INSTITUT NATIONAL DE LA SANTÉ ET DE LA RECHERCHE MÉDICALEInventor/a CARDINAUD, Sylvain

Developing an effective HIV-1 **vaccine** is contingent on generating protective antibodies (Abs). Novel antigen delivery methods are needed to enhance immune responses. One promising approach involves directing antigens to dendritic cells (DC) through fused monoclonal antibodies (mAbs) to amplify both cellular and humoral responses. Here, the inventors aimed to refine Langerhans cells (EC) targeting by designing three Env monochains instead of 2 (EC³. Env³) mimicking natural Env conformation. The inventors demonstrated that EC³. Env³ construct (i) elicited a rapid and potent Env-IgG response, (ii) enhanced the avidity of anti-Env IgG that was accompanied by a marked expansion of Tfh and GC B cells and (iii) swiftly induced the formation of structured germinal centers in dLN, indicative of a robust immune response. Finally, significant Tier-1 NeutAb induction was observed in rabbits immunized with the construct. In conclusion, HIV Env antigen can be adaptively targeted to LC as a timer, intensifying both the magnitude and quality of humoral responses. The present invention thus relates to the use of such a construct as LC targeting HIV-1 vaccines.

Patentes registradas en United States Patent and Trademark Office (USPTO)

Estrategia de búsqueda: *vaccine.ti. AND @PD>="20251001"<=20251012* 33 records

Document ID	Title	Inventor	Applicant Name
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US 20250312315 A1	CANCER VACCINE COMPOSITIONS AND METHODS OF USE THEREOF	OPGENORTH; Terry et al.	Colorado State University Research Foundation
US 20250312442 A1	EPSTEIN-BARR VIRUS VACCINES	Ciamarella; Giuseppe et al.	ModernaTX, Inc.
US 20250312428 A1	Heterologous Prime Boost Vaccine	WOLLMANN; Guido et al.	BOEHRINGER INGELHEIM INTERNATIONAL GMBH
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US 20250313858 A1	SYNTHETIC MODIFIED VACCINIA ANKARA (SMVA) BASED CORONAVIRUS VACCINES	DIAMOND; Don J. et al.	CITY OF HOPE
US 20250312429 A1	ANTI-GUCY2C VACCINES AND VACCINATION	Snook; Adam E et al.	THOMAS JEFFERSON UNIVERSITY
US 20250312439 A1	VACCINE FOR PREVENTING OR TREATING CORONAVIRUS INFECTION	MCCARTHY; Helen et al.	PHION THERAPEUTICS LTD
US 20250312427 A1	mRNA VACCINES COMPRISING IL-4 AND/OR IL-13 RNA AND USES THEREOF	DROUET; Beatrice et al.	NEOVACS
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US 20250314636 A1	METHOD OF PREPARING AND EXPANDING A POPULATION OF IMMUNE CELLS FOR CANCER THERAPY, POTENCY ASSAY FOR TUMOR RECOGNITION, BIOLOGICAL VACCINE PREPARATION AND EPITOPETARGET FOR ANTIBODIES	MENDONÇA GORGULHO; Carolina et al.	FUNDAÇÃO D. ANNA DE SOMMER CHAMPALIMAUD E DR. CARLOS MONTEZ CHAMPALIMAUD - CE, Human Cell Co.
US 20250312431 A1	VACCINES AGAINST MORITELLA VISCOSA	FUREVIK; Anette et al.	Zoetis Services LLC
US 20250312438 A1	HUMANIZED CONSTRUCTS, VACCINES, AND METHODS	Julien; Jean-Philippe et al.	THE HOSPITAL FOR SICK CHILDREN
US 20250312440 A1	CHEMICAL SCREEN OF MODULATORS FOR VACCINE ADJUVANTS	ESSER-KAHN; Aaron et al.	THE UNIVERSITY OF CHICAGO
US 20250312425 A1	COMPOSITIONS AND METHODS COMPRISING ADA-1 FOR WOUND HEALING AND VACCINE ENHANCEMENT	Spiller; Kara Lorraine et al.	Drexel University
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US 20250302941 A1	Application of an Immunogenic or Vaccine Composition	Blomet; Joël et al.	EYE VACC
US 20250302940 A1	PATHOGEN-LIKE ANTIGEN-BASED VACCINE AND PREPARATION METHOD THEREFOR	HOU; Baidong et al.	RONGSEN BIOTECHNOLOGY (BEIJING) CO., LTD, INSTITUTE OF BIOPHYSICS, CHINESE ACADEMY OF SCIENCES
US 20250302942 A1	VACCINE COMPOSITION AGAINST CORONAVIRUS	NOH; Yoon Ae et al.	LG CHEM, LTD.
US 20250302936 A1	MULTIVALENT PNEUMOCOCCAL VACCINES	Besin; Gilles R. et al.	Affinivax, Inc.
US 20250304636 A1	DAP12 CONSTRUCTS AND THEIR USE TO ENHANCE DC VACCINES AND IMMUNOTHERAPIES	Eksioglu; Erika A.	H. LEE MOFFITT CANCER CENTER AND RESEARCH INSTITUTE, INC.
US 20250302939 A1	VACCINE FOR HUMAN T-LYMPHOTROPIC VIRUS-1	Franchini; Genoveffa et al.	The United States of America, as represented by the Secretary, Department of Health and Human
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