



EN ESTE NÚMERO

VacCiencia es una publicación dirigida a investigadores y especialistas dedicados a la vacunología y temas afines, con el objetivo de serle útil.

Usted puede realizar sugerencias sobre los contenidos y de esa forma crear una retroalimentación que nos permita acercarnos más a sus necesidades de información.

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Noticias en la Web

Reino Unido vacunará a miles de jóvenes para reducir los brotes por meningococo B en el país

12 jun. Los datos de la Agencia de Seguridad Sanitaria del Reino Unido (UKHSA, por sus siglas en inglés) muestran que hubo 313 casos confirmados de meningococo del grupo B (MenB) en Inglaterra durante 2024/2025. A principios de 2026, se produjo el mayor brote de meningitis B tanto en crecimiento como en magnitud jamás registrado en Reino Unido. Ante esta situación, el gobierno ha anunciado que miles de jóvenes de todo el país recibirán protección contra la enfermedad a través de un programa de vacunación único que se pondrá en marcha antes del año académico 2026.

“El Servicio Nacional de Salud espera que las vacunas estén disponibles a finales de julio para comenzar su inyección antes del nuevo curso académico.”

La enfermedad meningocócica es potencialmente mortal y puede provocar discapacidades permanentes como amputaciones, pérdida de audición y daño cerebral, mientras que la mortalidad se produce en aproximadamente el 10% de los casos. Los virus y bacterias que causan la meningitis se transmiten por contacto cercano con una persona infectada. Esto puede ocurrir al besar, compartir bebidas o cigarrillos electrónicos, o por contacto prolongado con la persona, como vivir en el mismo hogar.

La vacuna del meningococo B reduce alrededor del 75% en la incidencia de la enfermedad.

Este programa de vacunación contra el meningococo B estará a disposición de las personas que completen el último año de estudios en el verano de 2026, nacidas entre el 1 de septiembre de 2007 y el 31 de agosto de 2008, así como de las personas menores de 25 años que comiencen la universidad o se trasladen por primera vez a algún centro de educación superior residencial en otoño de 2026. Los estudiantes que cumplan los requisitos podrán recibir una vacuna de dos dosis antes de comenzar el nuevo año académico.

En palabras del Dr. Thomas Waite, subdirector médico de Reino Unido, “para una protección máxima se necesitan dos dosis de la vacuna contra el meningococo B, con un intervalo de al menos cuatro semanas, y animo a todos los que vayan a asistir a la universidad o a un centro de estudios superiores por primera vez este otoño a que acudan a recibir su primera dosis lo antes posible”.



La vacuna contra el meningococo B tiene un historial de seguridad comprobado y ya se ofrece de forma rutinaria a los lactantes a través del programa de vacunación infantil del Servicio Nacional de Salud (NHS) del Reino Unido. La evidencia del programa de vacunación infantil del Reino Unido muestra que la vacunación ha logrado una reducción de alrededor del 75 % en la incidencia de la enfermedad por meningococo B entre los grupos vacunados que cumplen los requisitos.

Kent fue una de las ciudades que sufrió uno de los mayores brotes de meningococo B jamás registrados en Inglaterra.

La UKHSA estima que el riesgo relativo de enfermedad meningocócica invasiva B en estudiantes universitarios de primer año es considerablemente mayor que en sus compañeros. Se sabe que los casos de enfermedad meningocócica invasiva suelen alcanzar su punto máximo entre octubre y noviembre de cada año. El programa de vacunación ayudará a proteger a quienes corren mayor riesgo inmediato de padecer una enfermedad grave al mudarse a residencias compartidas.

Este programa de vacunación único ayudará a proteger a las personas con mayor riesgo inmediato mientras el gobierno supervisa y evalúa las nuevas pruebas para determinar si ha habido un cambio en la forma en que la meningitis B afecta a las personas y si se requiere alguna respuesta adicional de vacunación. Las sospechas de una posible mutación están basadas por el brote en Kent y los recientes casos que indican un posible cambio en la forma en que la meningitis B afecta a las personas.

El NHS ha comenzado la planificación de la implementación operativa, y se espera que las vacunas estén disponibles en Inglaterra a partir de finales de julio de 2026, antes del inicio del nuevo año académico. Paralelamente a la puesta en marcha del programa, el DHSC, el NHS England y la UKHSA llevarán a cabo una campaña de información pública para fomentar la participación entre los grupos que cumplan los requisitos.

Fuente: ISANIDAD. Disponible en <https://n9.cl/ht2nz>

Enfermedad X: la amenaza desconocida que mantiene a la OMS en alerta permanente

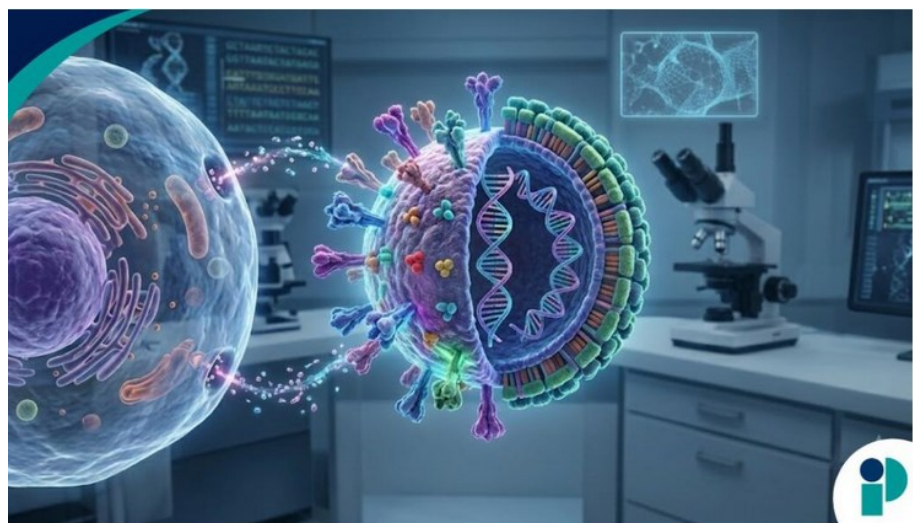
12 jun. La Organización Mundial de la Salud (OMS) lleva varios años preparándose para una amenaza que todavía no tiene nombre ni rostro. Se trata de la denominada Enfermedad X, un concepto incorporado por la organización en 2018 para representar la posibilidad de que una futura pandemia sea provocada por un agente infeccioso desconocido para la ciencia.

A diferencia de otras enfermedades incluidas en la lista de vigilancia de la

OMS, la Enfermedad X no corresponde a un virus, bacteria o patógeno específico. Más bien, reconoce una realidad cada vez más evidente: las mayores amenazas para la salud pública podrían surgir de microorganismos que aún no han sido detectados o de virus conocidos que evolucionen y se vuelvan más transmisibles o peligrosos.

La experiencia de la COVID-19 reforzó la preocupación

Para muchos expertos, el SARS-CoV-2, causante de la pandemia de COVID-19, se convirtió en el primer ejemplo real de lo que representa la Enfermedad X. El virus apareció de manera inesperada, se propagó rápidamente por todo el mundo y generó una emergencia sanitaria global antes de que existieran vacunas o tratamientos específicos.



Esta experiencia impulsó a la OMS y a la comunidad científica internacional a fortalecer la preparación frente a amenazas emergentes, promoviendo el desarrollo de plataformas de vacunas adaptables, sistemas de diagnóstico rápidos y redes de vigilancia epidemiológica capaces de identificar brotes inusuales de manera temprana.

La Enfermedad X forma parte del Plan Director de Investigación y Desarrollo para Prevenir Epidemias, una iniciativa creada por la OMS en 2015 para acelerar la disponibilidad de diagnósticos, tratamientos y vacunas frente a enfermedades emergentes.

Para elaborar su lista de prioridades, la organización reúne expertos en microbiología, epidemiología, salud pública, veterinaria, antropología y seguridad. Estos especialistas analizan factores como la capacidad de transmisión entre personas, la gravedad de la enfermedad, el potencial de expansión internacional, el impacto social y económico y la posibilidad de que el patógeno evolucione con el tiempo.

Las enfermedades que actualmente preocupan a la OMS

Además de la Enfermedad X, la OMS mantiene bajo vigilancia varios patógenos considerados de alto riesgo, entre ellos:

- ⇒ Virus del Ébola, virus de Marburg, fiebre de Lassa, virus Nipah, MERS, SARS, virus Zika, fiebre hemorrágica de Crimea-Congo y fiebre del Valle del Rift.

Estas enfermedades han sido identificadas como amenazas que requieren investigación acelerada debido a su potencial de provocar brotes graves y por la limitada disponibilidad de herramientas para su control.

Aunque la influenza no figura en la lista de prioridades del Plan Director, porque ya cuenta con vacunas y sistemas de vigilancia establecidos, una encuesta realizada entre especialistas de 57 países encontró que los virus de la gripe siguen siendo considerados la principal amenaza pandémica actual. La Enfermedad X ocupó el segundo lugar entre las preocupaciones de los expertos.

Esta aparente contradicción refleja uno de los mayores desafíos de la salud pública moderna: prepararse tanto para amenazas conocidas como para aquellas que todavía no han sido descubiertas.

Más que intentar predecir cuál será la próxima pandemia, la estrategia de la OMS busca garantizar que el mundo esté listo para responder rápidamente cuando surja una nueva amenaza. La Enfermedad X simboliza precisamente ese reto: desarrollar herramientas, sistemas de vigilancia y capacidades de respuesta que permitan actuar con rapidez frente a lo desconocido. «La Enfermedad X representa la posibilidad de que un patógeno actualmente desconocido para la medicina pueda surgir y propagarse a escala mundial».

Fuente: PHL PUBLIC HEALTH LATAM. Disponible en <https://n9.cl/4t6jj>

Brazil Suspends New Dengue Vaccine After Two Fatalities

Jun 12. The Brazilian Ministry of Health has announced a temporary suspension of its mass immunization campaign using the domestically produced Butantan-DV vaccine, pending an investigation into the deaths of two vaccine recipients.

Brazilian health authorities have ordered a precautionary halt to the rollout of Butantan-DV, a newly developed single-dose dengue vaccine produced by the country's renowned Instituto Butantan in

partnership with WuXi Vaccines. The decision follows reports of two suspected fatalities, a 58-year-old man and a 48-year-old woman, as well as a third recipient who required emergency admission to an intensive care unit. These severe adverse events occurred among approximately 500,000 individuals who have received the vaccine since its recent introduction.

Approved Dengue Vaccines

- ⇒ Qdenga (Takeda): A live-attenuated vaccine designed to provide protection against all four dengue virus serotypes. Qdenga is administered as a two-dose regimen, with doses given three months apart.
- ⇒ Dengvaxia (Sanofi Pasteur): The first dengue vaccine to receive regulatory approval. However, its use is now highly restricted. It is indicated exclusively for individuals aged 9 to 45 years with laboratory-confirmed evidence of a previous dengue infection.

The current concern in Brazil inevitably recalls one of the most controversial public health crises of the 21st century. In 2015, the French pharmaceutical company Sanofi Pasteur launched Dengvaxia to widespread acclaim. In collaboration with the Philippine government, the company initiated a large-scale school-based immunization program that ultimately vaccinated more than 800,000 children.

However, in 2017, the manufacturer disclosed a critical scientific finding: while the vaccine provided substantial protection to individuals who had previously experienced a natural dengue infection, it posed a significant risk to dengue-naïve (seronegative) individuals who had never been infected with the virus. In these children, the vaccine effectively simulated a primary infection. When they were subsequently exposed to wild-type dengue virus, their immune systems could respond through a phenomenon known as Antibody-Dependent Enhancement (ADE), increasing the risk of severe and potentially life-threatening disease.

The resulting fallout in the Philippines triggered unprecedented political, legal, and criminal investigations involving public health officials, led to the revocation of the vaccine's local registration, and contributed to a profound and long-lasting decline in public confidence in vaccination programs throughout the region.

Fuente: POULTRY MED. Disponible en <https://n9.cl/qs5iqw>

How New Technologies Are Impacting the Vaccine Market

Jun 13. In January of this year, Novavax announced a partnership that would provide its Matrix-M adjuvant technology to Pfizer for use in vaccine development. Pfizer reportedly paid an upfront total of \$30 million, with an additional \$500 million agreed upon based on development and commercial milestones.

At the 2026 JP Morgan Healthcare conference, Novavax CEO John Jacobs said, “This company is no longer about one product or one season. It’s about building a sustainable engine for value creation built on technology, discipline, and long-term partnerships.”

This news came during a period of regulatory uncertainty surrounding vaccines. Due to abrupt leadership changes at HHS, ACIP was forced to cancel its February meeting this year. In 2025, incoming HHS Secretary Kennedy fired the previous council and replaced the members with his own picks.

However, this hasn't stopped vaccines from being developed and approved. In May of this year, FDA's advisory panel voted to update COVID-19 vaccine shots to target the XFG variant. This update includes the vaccine developed in partnership with Novavax and Sanofi.

Silvia Taylor spoke with Pharmaceutical Executivex about the current vaccine landscape and how new technologies are allowing more players to enter the space and develop new therapies. She also discussed the regulatory landscape and how vaccine developers are working around any potential issues.

Pharmaceutical Executive: How are new vaccine technologies impacting the vaccine space?
Silvia Taylor: Vaccine technologies are really having a major impact on the vaccine environment right now. Since the onset of COVID, people are disproportionately attuned to what different technology platforms are in vaccines.

Novavax is twofold: one, we have a protein-based nanoparticle technology platform that enables us to develop vaccines for a wide range of diseases, such as COVID. We also have our adjuvant matrix, which is an important part of our technology platform.

Basically, an adjuvant is an ingredient that's added to a vaccine to stimulate a potentially broader, more durable, and more potent response. It can be added to vaccines to potentially lower the cost of goods, side effect profile, and the amount of antigen a vaccine needs.

That's an incredibly important component in vaccine development, and we're finding that our adjuvant, called Matrix M, has broad applicability in a variety of disease areas. We've already proven it in COVID 19. We have an approved vaccine that's on the market, and we also proved it in malaria.

There's a malaria vaccine that utilizes matrix M, and right now we're generating data and working with partners to understand how our matrix potentially improve the immune response of existing vaccines in their portfolios, or the creation of entirely new vaccines altogether.

Fuente: PharmExec.com. Disponible en <https://n9.cl/7w955>

Intravacc And SynphaBase Announce Strategic Partnership To Advance Conjugate Vaccine Development

Jun 15. The companies will combine their complementary expertise in synthetic oligosaccharides, conjugation process development and GMP vaccine manufacturing to support partners developing next-generation conjugate vaccines



- ◆ SynphaBase brings deep expertise in glycochemistry and the application, production, supply and regulatory support of synthetic oligosaccharides for conjugate vaccine development
- ◆ Intravacc contributes proven capabilities in conjugation process development, scale-up and GMP production of conjugate vaccines
- ◆ The partnership will offer vaccine developers a streamlined route from synthetic oligosaccharide supply to conjugation development and clinical manufacturing

Intravacc, a leading global contract development and manufacturing organization specializing in vaccine and biologics development and manufacturing, and SynphaBase AG, a Swiss contract

research and development organization rooted in glycochemistry and complex molecule synthesis, today announced a strategic partnership to support the development and supply of conjugate vaccines.

Under the partnership, the companies will jointly offer their complementary capabilities to third parties developing conjugate vaccines, as well as to selected Intravacc internal vaccine programs at Intravacc's discretion. The collaboration is designed to provide clients with access to SynphaBase's synthetic oligosaccharide expertise and Intravacc's conjugation development, scale-up and GMP manufacturing capabilities.

Ivo Lemmens, Managing Director of Intravacc, says:

"We are pleased to announce our strategic partnership with SynphaBase. By combining SynphaBase's expertise in synthetic oligosaccharides with Intravacc's capabilities in conjugation process development and GMP vaccine manufacturing, we aim to provide vaccine innovators with an integrated and efficient development pathway for conjugate vaccines."

Arthur Bodenmüller, CEO of SynphaBase, further comments:

"SynphaBase is excited to partner with Intravacc to bring together our complementary strengths in glycochemistry and vaccine development. Synthetic oligosaccharides are an important enabling technology for modern conjugate vaccines, and this collaboration will help developers access the expertise, materials and manufacturing support needed to advance promising vaccine candidates."

This partnership brings together the synergistic strengths of both companies. Intravacc contributes its technical expertise, facilities and capabilities in antigen conjugation process development, scale-up and GMP production and supply of conjugate vaccines. SynphaBase complements this with its expertise in the application, production, supply and regulatory support of synthetic oligosaccharides used in research, clinical development, production and supply of conjugate vaccines.

Through the partnership, SynphaBase may promote Intravacc as a preferred CDMO for conjugate vaccine services, while Intravacc may promote SynphaBase as a preferred supplier within relevant internal or client conjugate vaccine programs. The partnership is non-exclusive, and clients remain free to select the products and services that best meet their development needs.

The collaboration reflects a shared commitment to enabling faster, more efficient development of conjugate vaccines by connecting high-quality synthetic oligosaccharide supply with specialist conjugation and GMP manufacturing expertise.

About Intravacc

Intravacc is a leading global CDMO specializing in vaccine and biologics development and manufacturing for biotech and pharmaceutical partners. Leveraging its cell-culture, bacterial and conjugation platforms and extensive GMP capabilities, Intravacc supports the full development pathway from early-stage research to Phase I/II clinical production. With a strong track record in technology transfer and scalable vaccine solutions, Intravacc enables partners to accelerate timelines, reduce risk and efficiently bring innovative vaccines to the clinic. For more information, visit www.intravacc.nl.

About SynphaBase AG

SynphaBase AG is a Swiss contract research and development organization specializing in bioorganic chemistry, with deep expertise in glycochemistry and the custom synthesis of complex

molecules including glycans, peptides, RNA and conjugates. SynphaBase supports partners through route design, process development and material supply, applying its know-how to help advance complex chemistry programs for the pharmaceutical, biotechnology and life sciences sectors. For more information, visit www.synphabase.ch.

Fuente: OUTSOURCED PHARMA. Disponible en <https://n9.cl/c9ha60>

WHO and Brazil urge world leaders to finalise Pandemic Agreement to prevent future global health crises

Jun 15. Penned by Brazilian President Luiz Inácio Lula da Silva and WHO chief Tedros Adhanom Ghebreyesus, the message underscored a shared global responsibility to protect humanity from repeating such devastation seen during the COVID-19 pandemic, which claimed up to 20 million lives and wiped out \$13 trillion in global economic output.

During the COVID-19 pandemic in 2020, hospitals were overwhelmed, families lost loved ones in isolation and frontline healthcare workers endured unprecedented strain. This collective trauma birthed a promise among nations: to never again face a pandemic unprepared, they wrote.



© WHO/Khaled Mostafa An elderly woman receives her third COVID-19 booster shot at a vaccination centre in Lisbon, Portugal, in 2024. (file)

‘An act of hope’ in a divided world

Over a year ago, countries made significant strides by adopting the WHO Pandemic Agreement, committing to cooperate more effectively in preventing, preparing for and responding to pandemics.

“In a divided world, that outcome was not to be taken for granted,” the letter stated. “It was an act of hope and an act of faith in one another. We write to you now because that hope is not yet fulfilled and because it lies within your hands to help fulfil it.”

Standing in the way is the pathogen access and benefit-sharing (PABS) annex, a vital element of the agreement, which remains incomplete, President Lula and Tedros wrote.

This framework is essential for enabling countries to quickly identify and share genetic information of dangerous pathogens so that scientists can develop lifesaving tests, treatments and vaccines.

Without the annex, the Pandemic Agreement cannot formally come into force, leaving the promise unfulfilled, the joint message stated.

Complex challenges remain

The challenges in finalising the PABS annex are complex, particularly around defining how the benefits of shared pathogens are equitably distributed and how governance ensures fairness.

These were the very questions left unresolved previously, contributing to gaps in protection during COVID-19.

Negotiators are scheduled to meet again from 6 to 17 July to close these gaps.

Three critical requests

The appeal highlights three critical requests for global leaders:

Political will at the highest level: Leaders must prioritise finalising the annex and empower negotiators to seek bold consensus. The agreement does not compromise national sovereignty nor does it grant WHO authority to impose measures like lockdowns or vaccination mandates. Those decisions remain firmly with individual nations.

Spirit of equity: The PABS system is built on fairness. Those who share pathogens swiftly must trust that resulting vaccines and treatments will also reach their populations, an approach acknowledging that pandemic prevention is not charity but sound strategy, reducing costs and lives lost by containing outbreaks early. Equitable access must be guaranteed, ensuring clear, stable rules to replace the current case-by-case, crisis-driven process.

A sense of urgency: Scientists estimate a nearly 25 per cent chance of another pandemic within the next decade. Shifting environmental and social factors are creating new hotspots for disease emergence worldwide while biotechnology advances raise risks of accidental or intentional releases. The letter calls for 17 July to be treated as a firm deadline for agreement, signaling global commitment and avoiding dangerous delays.

The next crucial chapter

Current outbreaks, such as Ebola in multiple countries with no approved vaccine, highlight the ongoing risks, they warned.

Reminding readers of the staggering human and economic toll of COVID-19, the letter stated that, by comparison, investing in early outbreak detection and response systems like the PABS framework is modest and essential.

Historically, the world has united to save countless lives by defeating smallpox, pushing polio to near eradication and combating HIV, tuberculosis and malaria.

“Finishing this agreement is not a departure from that legacy,” the letter stated. “It is its natural next chapter, and it is within reach.”

Fuente: UNITED NATIONS NEWS. Disponible en <https://n9.cl/5pfd6u>

Maternal RSV vaccine shows 68% protection against infant RSV hospitalization

Jun 16. Respiratory syncytial virus (RSV) is a major cause of lower respiratory tract infections and hospitalization in young infants. Maternal vaccination with the RSV prefusion F vaccine was associated with a substantially lower risk of RSV-related hospitalization for respiratory disease in infants, according to a recent study published in JAMA Network Open.

RSV remains the leading cause of lower respiratory tract disease (LRTD) among infants globally and is responsible for 33 million cases and 3.6 million hospitalizations each year, with 100,000 deaths among under-fives. Approximately half of RSV-linked hospitalizations are in infants below six months, corresponding to a US rate of 24 /1,000 infants below three months with RSV-associated hospitalization.

“A real-world US study finds maternal RSVpreF vaccination may help shield newborns from severe RSV illness during their most vulnerable first months of life.”

In 2023, the bivalent RSV prefusion F vaccine was approved for use in pregnant individuals in the United States. It protects against RSV-A and RSV-B. It is intended to reduce the risk of RSV-associated LRTD and severe LRTD in newborns via maternal antibodies. The vaccine is recommended between 32 and 36 weeks and 6 days of gestation.

The current study aimed to assess the vaccine's real-world effectiveness in this situation. This ongoing study is designed to cover four years of maternal vaccination and RSV-associated ARI and LRTD outcomes in infants.

A retrospective case-control study using a test-

negative design was conducted during two RSV seasons from 2023 to 2025. The current cohort included 274 hospitalized infants with ARI, 90 days or younger. All were tested for RSV.

Maternal vaccination rates were compared between babies with RSV (cases) and those without (controls). Vaccine effectiveness against RSV-associated hospitalization of infants with acute respiratory illness (ARI) and LRTD was estimated from birth through 90 days of life.

Cohort Characteristics

The mean age of the 274 infants in the study was 30 days, with 48% being female. Over 60% were White and 14% Black. On average, they were born at 38 weeks. About 16% were born preterm between 34 weeks 0 days and 36 weeks 6 days of gestation. Most were born healthy.

Maternal Vaccination Linked to Reduced Severe RSV Disease in Infants

Of the 274 infants, 83 tested positive for RSV. These babies were more likely to have been born between October and December, at about 74%. They were also more likely to have been born preterm, male, Black, and to have siblings. They were also more likely to live in neighborhoods with higher levels of deprivation.

About 87% of cases had LRTD, compared with 45% in controls. While 34% of cases required intensive care unit (ICU) admission, this was true of only 17% of the controls. Similarly, about 21% of controls required oxygen or respiratory support compared to 68% of cases.

The investigators found that about 30% of the infants were born to prefusion F-vaccine recipients. This proportion was skewed towards the controls, with 37% of controls being born to vaccinated mothers versus 13% of cases.

Maternal Prefusion F Vaccination Associated with Infant Protection

This suggests that maternal vaccination was associated with lower odds of RSV-associated hospitalization among infants with ARI in the test-negative analysis. The estimated effectiveness of the vaccine when administered in pregnancy within the recommended gestational window and at least 14 days before delivery against this outcome was 68% in infants up to 90 days, with a 95% confidence interval of 33% to 85%.



Study: Maternal Respiratory Syncytial Virus Prefusion F Vaccination and Acute Respiratory Illness in Infants. Image Credit: Perfect Angle Images / Shutterstock.

For the same period, the estimated effectiveness against RSV-linked LRTD was 69% with a 95% confidence interval of 26% to 88%.

In the first month of life, the estimated vaccine effectiveness against RSV-linked ARI hospitalization was 74% (95% confidence interval: 25% to 93%).

The results are comparable to those from the key clinical trial that led to approval, and to broader surveillance studies. They are among the earliest real-world evidence that maternal RSV prefusion F vaccination may provide protection against the risk of severe RSV illness requiring hospitalization in the first three months of life. The point estimate was highest during the first month of life, although confidence intervals were wide, and this is also the period of greatest vulnerability.

Strengths and Limitations

The study used a real-world cohort to examine how maternal vaccination reduced infant hospitalizations for RSV-associated ARI. The cohort included preterm or sick infants and those born from multiple pregnancies, indicating the potentially broad utility of protection. The vaccination window was also much narrower than that used in the clinical trial, but yielded a similar magnitude of protection.

Moreover, the investigators used hospitalization data over two RSV seasons following vaccine introduction within a single health system in western Pennsylvania.

However, it has some limitations. It involved a single healthcare system with a small sample. Its observational design precludes causal inference; residual confounding may influence outcomes, though the study was designed to minimize bias from health-seeking behavior and misclassification. The relatively small sample resulted in wide confidence intervals for estimated effectiveness, particularly in subgroup analyses, which were not adequately powered.

Additional studies from other regions are needed to confirm effectiveness estimates and assess the duration of protection. The outcomes of earlier vaccination (allowing more time for antibody development and transplacental transfer) should be determined in larger samples to increase confidence in the findings.

Pfizer supported the study through a collaboration with the University of Pittsburgh, and several authors reported Pfizer employment, stock holdings, or grant support. The paper states that Pfizer reviewed and approved the manuscript but had no role in data collection, management, analysis, or the decision to submit it for publication.

Conclusion

Maternal RSV prefusion F vaccination during late pregnancy was associated with evidence of protection against RSV-related hospitalization in infants 90 days or younger. The findings suggest that this may be an effective strategy for preventing severe RSV disease during early infancy, and thus support current recommendations for maternal RSV immunization.

Fuente: NEWS MEDICAL LIFE SCIENCES. Disponible en <https://n9.cl/ucu6t>

Could a hantavirus vaccine be on the horizon?

Jun 17. When a rare, lethal virus broke out aboard a cruise ship this May and then scattered across two dozen countries with its passengers, it exposed an uncomfortable gap in the world's defences: there is no licensed vaccine or treatment for Andes hantavirus.

Now scientists at the University of Texas Medical Branch (UTMB) report that a single dose of an mRNA vaccine gave animals complete protection against the virus.

“A single shot of an experimental mRNA vaccine completely protected animals against the deadly Andes hantavirus, even at a fraction of the usual dose.”

The finding, published in *The Lancet*, could offer protection against one of the few hantaviruses able to spread between people.

“Every vaccinated animal remained completely healthy and showed no symptoms or weight loss,” said lead author Dr Michelle Meyer.

“When we looked at the tissues from the vaccinated animals a month after infection, the virus was entirely gone.”

Why the Andes virus worries scientists

Most hantaviruses spread to people through contact with the droppings, urine or saliva of infected rodents.

Andes virus is the exception as it is the only known member of the hantavirus family that can pass from one person to another through close contact with respiratory secretions, such as coughing.

That capacity for spread between people has driven alarming clusters before, explain the authors.

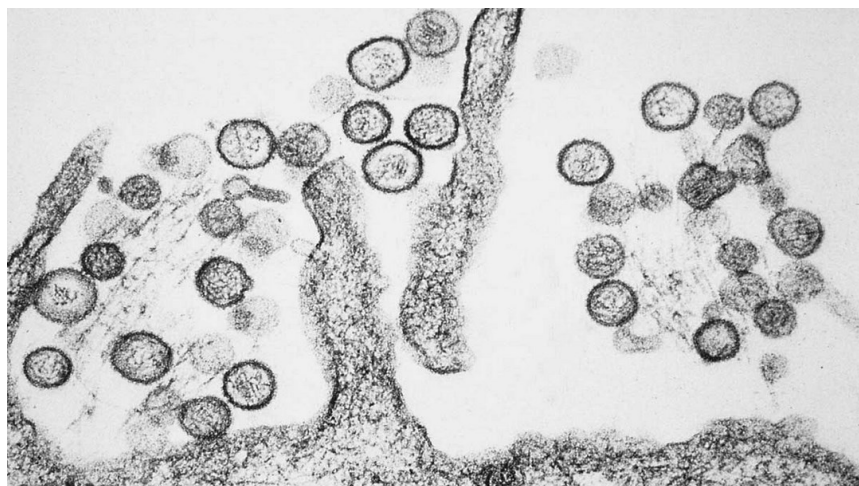
In the paper, they explain how during the 2018 and 2019 outbreak in Epuypén, Argentina, four successive waves of infection traced back to a single social gathering produced 34 confirmed cases and 11 deaths.

The May 2026 outbreak aboard the Dutch cruise ship MV *Hondius*, which had departed from Argentina, took that threat global. So far it has caused 13 reported cases and three deaths, but the wider worry is what came next.

Nearly 150 passengers dispersed across 23 countries under a patchwork of quarantine rules, all within a virus incubation window that can stretch to 42 days.

Because people can carry the virus without symptoms before they fall ill, tracing everyone who may have been exposed is a daunting task.

“When we looked at the tissues from the vaccinated animals a month after infection, the virus was entirely gone.” - Dr Michelle Meyer, Senior Scientist at the University of Texas Medical Branch.



A transmission electron micrograph of Sin Nombre virus.

Credit: CDC/Cynthia Goldsmith

Inactivated vaccines against other hantaviruses have been used in China and South Korea for decades.

But none protects against Andes virus, and no hantavirus vaccine of any kind is approved in the Americas or Europe.

With nothing licensed by either the US Food and Drug Administration or the European Medicines Agency, an Andes outbreak is precisely the kind of scenario the UTMB team is racing to prepare for.

One dose, full protection

The researchers had already shown that two of their mRNA vaccine candidates could protect animals against Andes virus, but only after two doses given weeks apart.

An international outbreak that moves fast rarely allows that luxury, so the team set out to test whether a single shot would be enough.

The vaccine uses the same basic design that powered the COVID-19 mRNA shots: genetic instructions that teach the body's own cells to make harmless pieces of the virus, in this case two surface proteins that train the immune system to recognise and fight the real thing.

The team gave 30 hamsters a single injection of the vaccine, then exposed them to a lethal dose of Andes virus four weeks later.

This hamster model is the only animal model that closely mirrors the fatal lung disease the virus causes in humans, and infection is almost always deadly: in the unvaccinated control group, by day nine, four of five animals hadn't survived.

In the vaccinated animals, meanwhile, every animal was protected from disease, and when the study ended, no replicating virus could be found in their tissue. The shots triggered detectable antibodies within 14 days, fast by most vaccine standards.

Reassuringly, the vaccine worked even at fractional doses. The team tested three dose levels: 25, 5 and 1 µg. Protection held even at the lowest dose, which still generated antibodies in four of five animals.

The ability to protect with so little vaccine matters because in an outbreak, the number of doses that can be squeezed from a limited supply can decide how many people get protected.

Post-exposure protection

The speed of the immune response opens a second possibility: using the vaccine not just to prevent infection, but to stop it in people who have already been exposed.

Andes virus disease has a median incubation of around 18 days in humans. If a vaccine can stimulate the immune system within two weeks, there may be a window to intervene after exposure but before illness sets in.

"If given quickly to high-risk contacts during an outbreak, such as the Andes virus situation on the cruise ship, the vaccines could theoretically jump-start their immune systems fast enough to intercept the virus, stopping it from replicating and preventing them from getting sick or spreading it further," said Dr Alexander Bukreyev, who heads UTMB's Laboratory of Viral Pathogenesis and Vaccine Development.

Used this way, a quick shot could double as an emergency tool to box in an outbreak, though the researchers are clear that this remains a hypothesis to be tested, not an established use.

From hamsters to humans

These results, encouraging as they are, come with the caveats that accompany any preclinical study. The work was done in animals, not people. The team measured the virus only in the liver, so they cannot rule out its presence elsewhere, such as the lungs, though in this model liver levels reliably track with those in the lungs.

The crucial next step is human clinical trials. With backing from the US National Institutes of Health, the UTMB team says it is working to speed the single-dose vaccines toward testing in people.

Fuente: GAVI. Disponible en <https://n9.cl/vis0n>

Sinergium Biotech consolidará a la Argentina como proveedor regional de vacunas de alta complejidad

18 jun. El titular de la cartera sanitaria recorrió la planta en la que se producen las vacunas antigripales y neumocócicas para el país y que se exportarán a toda la región.

El ministro de Salud de la Nación, Mario Lugones, visitó las instalaciones de Sinergium Biotech para conocer los avances de dos proyectos estratégicos para el desarrollo biotecnológico del país. Por un lado, la producción local de vacunas



antigripales que consolidarán al país como proveedor regional de vacunas de alta complejidad. Por el otro, la construcción de una nueva planta para el desarrollo y escalado de vacunas y terapias basadas en ARN mensajero (mRNA).

Este año, Sinergium ya ha producido y exportado más de 18 millones de dosis de la vacuna antigripal con ingresos que superaron los 60 millones de dólares. Además, prevé alcanzar los 120 millones de dólares en lo que resta del 2026, gracias al acuerdo firmado en 2025 con Pfizer y OPS para producción y exportación de la vacuna antineumocócica a través de los Fondos Rotatorios Regionales (FRR), posicionándose como un proveedor estratégico desde la Argentina para la región.

A partir del acuerdo con CSL Seqirus y la Organización Panamericana de la Salud (OPS), Sinergium Biotech comenzará a producir vacunas antigripales para abastecer al país y a toda la Región de las Américas a través de los FRR de la OPS. El proyecto permitirá generar empleo altamente calificado, fortalecer la cadena de valor local y aumentar la capacidad exportadora nacional en 100 millones de dólares anuales.

Como parte de su visita, Lugones también recorrió las obras de la nueva planta de producción piloto de vacunas basadas en ARN mensajero, con una inversión de 20 millones de dólares en dos etapas. El proyecto se enfoca en la investigación, el desarrollo y escalada de vacunas y otras tecnologías sanitarias para la prevención de enfermedades respiratorias y patógenos emergentes y

reemergentes, incluyendo influenza, COVID-19, virus sincial respiratorio y hantavirus en colaboración con ANLIS Malbrán, entre otras líneas de investigación.

Como parte de este proyecto, la ANLIS Malbrán actuará además como institución receptora de la transferencia tecnológica vinculada a la producción de lípidos estratégicos y nanopartículas lipídicas (LNP), insumos esenciales para la formulación de vacunas de mRNA. Su participación contempla también la incorporación de capacidades para la caracterización, evaluación y futura producción de estos componentes bajo estándares de Buenas Prácticas de Manufactura Internacional (GMP, por sus siglas en inglés), así como el fortalecimiento de recursos humanos, infraestructura y plataformas tecnológicas.

Cabe destacar que el desarrollo de una vacuna de ARN mensajero contra hantavirus constituye una de las iniciativas más innovadoras teniendo en cuenta que esta enfermedad no cuenta con vacunas aprobadas y registra tasas de letalidad superiores al 30% en América. El proyecto incluye el desarrollo de cuatro candidatos vacunales que serán evaluados en modelos experimentales.

En conjunto, el proyecto combina infraestructura, pipeline y alianzas internacionales para consolidar una plataforma sostenible, escalable y de alto valor estratégico para la región. La primera etapa, que se encuentra en desarrollo, contempla la construcción de una planta destinada a investigación, desarrollo y escalado piloto de vacunas y terapias basadas en mRNA y nanopartículas lipídicas (LNP). En una segunda etapa, se buscará desarrollar capacidades completas de producción bajo estándares GMP y permitir la fabricación de lotes clínicos y comerciales.

La visita del Ministro de Salud de la Nación permitió conocer de primera mano proyectos que fortalecen las capacidades científicas, tecnológicas e industriales del país y que consolidan la cooperación entre el sector público y privado. En el marco del proceso de estabilización económica, previsibilidad y apertura impulsado por el Gobierno nacional, ambas iniciativas contribuyen a posicionar a la Argentina como un actor relevante en el desarrollo y producción de tecnologías sanitarias estratégicas para la región, atraer inversiones y ampliar el acceso a innovaciones sanitarias.

Fuente: PMFARMA Disponible en <https://n9.cl/f4gfm>

Merck logra aval de la FDA para usar Capvaxive en niños y adolescentes con alto riesgo

18 jun. La FDA amplió la indicación de Capvaxive, la vacuna neumocócica de Merck, para aplicarla en una sola dosis a niños y adolescentes de 2 a 17 años con enfermedades crónicas —como cardiopatías, afecciones respiratorias, renales, hepáticas o diabetes— que ya completaron el esquema infantil estándar, pero siguen con mayor riesgo de infecciones graves por neumococo.

Capvaxive se abre paso más allá de los adultos

Capvaxive es una vacuna conjugada diseñada para proteger frente a múltiples serotipos de la bacteria *Streptococcus pneumoniae*, responsable de neumonía, meningitis e infecciones graves de la sangre.



Hasta ahora, el producto estaba aprobado en Estados Unidos solo para adultos, tras la autorización de la FDA en junio de 2024 para la prevención de enfermedad neumocócica invasiva y neumonía en mayores de 18 años.

La nueva luz verde del regulador se centra en un grupo particularmente vulnerable: “pacientes de 2 a 17 años que ya completaron el calendario de vacunación neumocócica infantil, pero presentan enfermedades crónicas como patologías cardíacas, pulmonares, renales, hepáticas o diabetes, que incrementan su riesgo de infecciones graves”.

En estos casos, Capvaxive no reemplaza a las vacunas infantiles, sino que “se administra como dosis única” suplementaria, con la intención de complementar la protección obtenida con los esquemas previos.

Qué mostró el ensayo y cómo se compara con PPSV23

La decisión de la FDA se basó en un ensayo de fase avanzada con 874 participantes, en el que Capvaxive “igualó o superó” la respuesta inmunitaria de PPSV23, la vacuna neumocócica polisacárida más antigua utilizada como refuerzo en pacientes de riesgo.

Según Merck, en este estudio los efectos adversos “fueron en general de corta duración”, alineados con lo esperado para este tipo de inmunizaciones, lo que respalda su perfil de seguridad en población pediátrica con comorbilidades.

PPSV23 ha sido durante años el estándar en refuerzos para adultos y ciertos niños con condiciones médicas de base, pero la nueva opción de 21 valencias introduce una cobertura más amplia frente a serotipos de neumococo que circulan hoy con mayor frecuencia.

En el contexto de niños y adolescentes ya vacunados en la infancia, la ampliación de Capvaxive aporta una herramienta adicional para cerrar brechas de protección en quienes, por enfermedades crónicas, tienen más probabilidades de desarrollar cuadros graves.

Niños crónicos en el centro de la estrategia preventiva

La enfermedad neumocócica se transmite por secreciones respiratorias —saliva y mucosidad— y puede derivar en neumonía, meningitis o infecciones sanguíneas potencialmente mortales.

Los niños pequeños menores de 5 años y los adultos de 50 años o más se consideran grupos de mayor riesgo, pero los chicos y adolescentes con patologías crónicas también cargan con una probabilidad incrementada de hospitalización y complicaciones.

Al focalizar esta expansión de indicación en pacientes de 2 a 17 años con enfermedades cardíacas, pulmonares, renales, hepáticas o diabetes, la FDA reconoce ese riesgo adicional y habilita un uso más dirigido de Capvaxive como complemento de los esquemas estándar.

Para sistemas de salud y pagadores, la posibilidad de agregar una dosis única de una vacuna de espectro amplio en un subgrupo bien definido abre la puerta a estrategias de inmunización más personalizadas, con potencial para reducir internaciones y costos asociados a neumonía y otras infecciones graves en población pediátrica de alto riesgo.

En un mercado global de vacunas neumocócicas donde la competencia entre formulaciones y valencias es intensa, la entrada de Capvaxive en el segmento de niños y adolescentes de riesgo consolida a Merck como un actor clave en la inmunización frente a *Streptococcus pneumoniae* y refuerza la tendencia a ampliar la protección más allá de la población adulta.

Fuente: CURE COMPASS. Disponible en <https://n9.cl/w8v802>

Ministerio de Salud no tiene vacunas contra la COVID-19, mientras repuntan casos en Guatemala

18 jun. En los dos hospitales de referencia del país, el Roosevelt y el General San Juan de Dios, los casos de COVID-19 han aumentado en las últimas semanas. El reporte se acerca al centenar de personas contagiadas entre pacientes y personal sanitario. Pero, fuera de lo que sucede en estos centros, no hay una actualización oficial de casos.

Desde mayo del 2023, la Organización Mundial de la Salud declaró a la COVID-19 como una enfermedad endémica, con lo que confirmó que el virus se volvía estacional. Guatemala se sumó a esa declaración en diciembre de ese año.

En las épocas frías, la enfermedad tiene mayor presencia en el país, y la recomendación de los médicos es la vacunación anual contra el virus. Sin embargo, actualmente la existencia del biológico en los servicios de salud pública es nula. Se consultó a varios y se indicó que no se cuenta con el biológico.

El Ministerio de Salud informó que la disponibilidad de la vacuna obedece a criterios técnicos, epidemiológicos y de abastecimiento nacional.

Agregó que, luego de la emergencia sanitaria, el monitoreo del covid-19 se mantiene en el país como parte de la vigilancia de enfermedades respiratorias.

Por ahora no hay vacunas, pero Salud señaló que se prevé el ingreso de dosis durante el tercer trimestre del año, con el fin de fortalecer la prevención en la población, conforme la planificación del Programa de Inmunizaciones de la cartera.

La primera dosis de la vacuna contra el covid-19 se aplicó el 25 de febrero del 2021, y las últimas dosis ingresaron al país en diciembre del 2023, con fecha de caducidad en mayo del 2024. Desde entonces, el sistema de salud pública no cuenta con el biológico.

El consolidado del Ministerio de Salud de dosis administradas contra el coronavirus en el país es de 20 millones 491 mil 871, cifra que incluye primeras y segundas dosis, refuerzos y vacunas adicionales.

Nancy Sandoval, jefa del Servicio del Departamento de Medicina Interna de la Unidad de Infectología del Hospital Roosevelt de Guatemala y vicepresidenta de la Asociación Panamericana de Infectología (API), indicó que el repunte de casos que se registra en el país ocurre por la disminución de la inmunidad contra la enfermedad entre la población, adquirida por infección natural o por vacunación; la circulación de nuevas variantes; la reducción de las medidas de vigilancia, y porque la población “subestima” la circulación comunitaria del virus.

Si bien las medidas básicas de prevención son importantes, como el lavado de manos, el uso de alcohol en gel, cubrirse al toser o estornudar, ventilar los espacios, usar mascarilla y aislarse voluntariamente ante síntomas de la enfermedad, la infectóloga indica que la vacunación es la acción “más eficaz” para evitar cuadros graves, hospitalización y muerte.

Por ello, recomienda mantener al día el esquema de vacunación contra el covid-19, principalmente en el personal de salud, las personas mayores de 50 años, quienes padecen enfermedades crónicas o inmunosupresión, las embarazadas y los niños con factores de riesgo.

Fuente: PRENSA LIBRE. Disponible en <https://n9.cl/z9esf>

República Democrática del Congo, entre la fiesta del fútbol y un brote de ébola que podría ser el peor de la historia

19 jun. Al mismo tiempo que miles de personas en la República Democrática del Congo se reunieron para el debut de su selección en el Mundial y celebraron en las calles el empate 1 a 1 contra Portugal, *Africa News* registró que se cumple un mes desde que la Organización Mundial de la Salud (OMS) declaró como emergencia de salud pública de importancia internacional el brote de ébola que afecta tres provincias del oeste del país. Funcionarios advierten que podría convertirse en la peor epidemia registrada de esa enfermedad, superando la que afectó a países de África Occidental entre 2014 y 2016 con más de 11.000 decesos.

Panorama: La diseminación de la rara variante Bundibugyo ya produjo más de 800 casos confirmados (con 181 decesos) y los trabajadores de la salud carecen de personal necesario para identificar los casos sospechosos, ambulancias para transportarlos e incluso materiales de construcción para edificar salas de aislamiento, reportó Reuters. Tampoco se conoce la verdadera magnitud del brote: "Solo estamos monitoreando a 12 % de nuestra población [que estuvo en contacto con los casos confirmados]. Esto es un indicador importante para nosotros", advirtió el Dr. Jean Kaseya, maestro en salud pública, director general de los Centros de Prevención y Control de Enfermedades (CDC) de África. También hay grave escasez de equipos de sepelio y de equipos de protección personal. "Si no detenemos el brote, muy pronto será peor que lo que tuvimos [antes]". Uganda también tiene alrededor de dos decenas de casos confirmados, cinco por transmisión local y el resto importados.

Vacuna: En tanto, vacunas desarrolladas por la Universidad de Oxford del Reino Unido, y Moderna contra la variante Bundibugyo del ébola, podrían entrar en la fase 1 de ensayos clínicos a partir de julio y quizá los ensayos de campo comenzarían en los próximos meses, declaró Richard Hatchett, director de la Coalición para las Innovaciones en la Preparación ante Epidemias (CEPI).

Fuente: MEDSCAPE. Disponible en <https://n9.cl/15kte>

FDA respalda primera vacuna antigripal que utiliza ARNm, tecnología que fue clave para poner fin a la pandemia de COVID-19

19 jun. Los estudios han mostrado que la vacuna de Moderna ha reducido los casos de gripe en un 27 % en un estudio de 40,000 participantes mayores de 50 años.

El comité asesor independiente de la Administración de Alimentos y Medicamentos (FDA) votó por unanimidad a favor de la vacuna mFlusiva de Moderna, afirmando que sus beneficios superan los riesgos para los grupos de edad de 50 a 64 años y 65 años o más, lo que marca un paso importante hacia la aprobación final que se espera en agosto.

Los estudios han mostrado que la vacuna de Moderna ha reducido los casos de gripe en un 27 %



en un estudio de 40,000 participantes mayores de 50 años, generando también una fuerte respuesta inmune en personas de 65 años o más, en comparación con las vacunas existentes.

La tecnología de ARNm permite una adaptación más rápida a las cepas mutadas del virus de la gripe, una tecnología que fue clave para poner fin a la pandemia de COVID-19. Esta posición del Comité Asesor de Vacunas y Productos Biológicos Relacionados (VRBPAC) está en abierta oposición a medidas de la administración Trump, que en agosto pasado canceló 500 millones de dólares de fondos que serían destinados a 22 proyectos de investigación para crear nuevas vacunas de ARNm contra enfermedades respiratorias que podrían desencadenar otra emergencia sanitaria.

Las vacunas con este tipo de tecnología tienen la ventaja de que se fabrican más rápidamente que otros tipos, algo que, según los expertos, podría ser útil si el virus de la gripe, con su capacidad de cambiar de forma, muta de tal manera que requiera la producción repentina de nuevas dosis.

“Disponer de esta tecnología nos coloca en una mejor posición para estar preparados ante la aparición de nuevas cepas en el futuro”, dijo la Dra. Flor Muñoz-Rivas, del Texas Children’s Hospital, una de las asesoras de la FDA.

“Creemos que el ARNm-1010 tiene el potencial de ofrecer una nueva e importante opción para la prevención de la gripe estacional y demostrar aún más la versatilidad de nuestra plataforma de ARNm”, declaró Stephane Bancel, MBA, MSc, director ejecutivo de Moderna, en un comunicado de prensa. “Esperamos seguir colaborando con la FDA mientras completa su revisión”, recalcó.

Preparación para la temporada de gripe

Se anticipa que, si es aprobada, la vacuna mFlusiva asegurará una mayor disponibilidad de dosis actualizadas en años donde las predicciones de cepas son inciertas.

Moderna planea un estudio de seguimiento que incluirá a 400,000 personas mayores de 65 años para continuar evaluando la seguridad y efectividad de la vacuna.

La solicitud de Moderna enfrentó desafíos regulatorios en el pasado, incluyendo críticas sobre sus métodos de comparación de vacunas. A pesar de esto, se espera que la FDA avance con una evaluación más positiva, considerando la importancia del desarrollo de esta vacuna en el contexto de la salud pública.

Comprendiendo la tecnología ARNm

La tecnología del ARN mensajero (ARNm) es una de las innovaciones biomédicas más importantes de las últimas décadas. Su principio básico es sencillo y elegante: en lugar de administrar directamente una proteína terapéutica o un virus atenuado para enseñar al sistema inmune, se entrega la instrucción genética necesaria para que las propias células del paciente construyan la proteína deseada.

El ARNm actúa como un manual de instrucciones transitorio que las células leen para sintetizar una proteína específica; esa proteína puede servir como vacuna, como terapia génica temporal o como herramienta para modular procesos biológicos.

Una de las ventajas clave del ARNm es su velocidad y flexibilidad. Las secuencias de ARNm se diseñan y sintetizan rápidamente en laboratorio, lo que permite respuestas ágiles frente a nuevos patógenos o para personalizar tratamientos.

En vacunas, por ejemplo, una secuencia de ARNm codifica una proteína del patógeno (como la proteína de superficie de un virus). Al administrarse, las células producen esa proteína, el sistema

inmune la reconoce como extraña y genera respuestas defensivas —anticuerpos y células T— que confieren inmunidad sin exponer al organismo al patógeno completo.

Para que el ARNm sea útil en el cuerpo, tiene que sortear varios desafíos: es una molécula inestable y fácilmente degradable por enzimas; además, debe entrar eficientemente en las células.

Más allá de las vacunas, las aplicaciones potenciales del ARNm son amplias. En terapia contra el cáncer, por ejemplo, se diseña ARNm que codifica antígenos tumorales específicos para estimular una respuesta inmune dirigida contra las células tumorales. En enfermedades genéticas, se puede usar ARNm para proporcionar una versión funcional temporal de una proteína faltante o defectuosa, evitando algunas complicaciones de las terapias génicas permanentes. También existen investigaciones que exploran el uso del ARNm para producir medicamentos directamente en el organismo, o para regeneración tisular mediante la expresión temporal de factores de crecimiento.

Fuente: LA OPINIÓN. Disponible en <https://n9.cl/qzh62w>

Brasil comienza vacunación infantil con nuevo inyectable neumocócico

20 jun. La incorporación del nuevo inyectable al Programa Nacional de Inmunizaciones representa un avance en la estrategia de prevención de infecciones respiratorias y otras afecciones potencialmente mortales que impactan principalmente a menores de edad, ancianos y personas con sistemas inmunológicos debilitados.

Agencia Brasil agregó que Pneumo 20 sustituirá de manera progresiva a la versión neumocócica 10-valente utilizada hasta ahora en la red pública de salud.

Según el Ministerio del sector, el nuevo preparado ofrece cobertura contra 20 serotipos de la bacteria neumococo y amplía significativamente la protección frente a variantes responsables de enfermedades invasivas y complicaciones respiratorias.

Entre las patologías prevenibles mediante la inmunización figuran neumonías, meningitis, otitis, sinusitis y septicemias.

Autoridades sanitarias destacan que la ampliación de la cobertura vacunal permitirá reducir hospitalizaciones, secuelas y fallecimientos asociados a estas enfermedades.

La nueva vacuna será administrada de forma gratuita a través del Sistema Único de Salud (SUS) y estará destinada inicialmente a niños menores de un año, siguiendo el calendario nacional de vacunación.

El esquema prevé la aplicación de dosis durante los primeros meses de vida y una de refuerzo posteriormente, conforme a las orientaciones del programa de inmunización.

Tal decisión se basa en evidencias científicas y recomendaciones técnicas que demuestran una mayor capacidad de protección frente a los serotipos actualmente en circulación.

Asimismo, la actualización permitirá mantener la eficacia de las estrategias de prevención ante los cambios observados en el comportamiento epidemiológico de la bacteria.

“El Ministerio de Salud de Brasil comenzó hoy la aplicación de la vacuna neumocócica conjugada 20-valente (Pneumo 20) en niños, con el objetivo de ampliar la protección contra enfermedades graves causadas por la bacteria *Streptococcus pneumoniae*.”

Pneumo 20 forma parte de los esfuerzos gubernamentales para fortalecer las coberturas vacunales y ampliar el acceso a tecnologías más avanzadas en salud pública.

Brasil ha impulsado en los últimos tiempos diversas acciones para recuperar los índices de inmunización infantil, afectados por la disminución de la adhesión a las campañas de vacunación durante la pandemia de COVID-19.

Millones de niños, recalcó el Ministerio de Salud, podrán beneficiarse anualmente de la nueva vacuna, con lo cual disminuirá la circulación de cepas peligrosas del neumococo y se fortalecerá la capacidad del sistema sanitario para enfrentar enfermedades respiratorias.

Fuente: Prensa Latina. Disponible en <https://n9.cl/vo9g5o>



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Patentes registradas en Patentscope

Estrategia de búsqueda: (*Vaccine*) AND DP:([11.06.2026 TO 21.06.2026]) as the publication date 54 records.

1. [WO/2026/119155](#) RECOMBINANT ADENOVIRUS VECTOR TUMOR VACCINE, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 11.06.2026

Clasificación Internacional C12N 15/12N^o de solicitud PCT/CN2025/139475 Solicitante BRITIE BIOTECH CO., LTD. Inventor/a ZHENG, Xiuyu

A recombinant adenovirus vector tumor vaccine, a preparation method therefor, and use thereof. An antigen encoded by the recombinant adenovirus vector tumor vaccine comprises any one or a combination of two or

more of antigenic peptide epitopes of a KRAS protein. The recombinant adenovirus vector tumor vaccine encodes multiple KRAS tumor-specific antigenic peptides, and exhibits a good killing effect on tumors driven by KRAS mutations, especially on lung tumors. The vaccine can be administered via nebulized inhalation, intramuscular injection followed by nebulized inhalation, nebulized inhalation followed by intramuscular injection, intramuscular injection, or simultaneous intramuscular injection and nebulized administration. After nebulization, the vaccine can reach the lungs via nasal or oral inhalation, generating anti-tumor protective immune responses in the respiratory tract and lungs and throughout the body, thereby enhancing the utilization rate and therapeutic efficacy of the vaccine.

2. 20260166134 MRNA, MRNA VACCINE FORMULATION, AND PREPARATION METHOD AND USE THEREOF

US - 18.06.2026

Clasificación Internacional A61K 39/00N° de solicitud 19258958 Solicitante Panru Biotechnology (Tianjin) Co., Ltd Inventor/a Yuanjie NIU

An mRNA, an mRNA vaccine formulation, and a preparation method and use thereof are provided. An amino acid sequence of the mRNA is set forth in SEQ ID NO: 2. Studies have demonstrated that the mRNA vaccine formulation prepared from the mRNA having the amino acid sequence set forth in SEQ ID NO: 2 can effectively stimulate dendritic cell maturation and secretion of pro-inflammatory cytokines, thereby activating effector T cells to kill prostate cancer cells. A preparation method of the mRNA vaccine formulation is simple and suitable for industrial-scale production. By using lipid nanoparticles (LNPs) as a carrier, the mRNA vaccine formulation achieves high encapsulation efficiency, enabling robust induction of immune responses in vivo with high immunogenicity.

3. WO/2026/122534 IMMUNOGENS AND METHODS FOR INDUCING BROADLY NEUTRALIZING ANTIBODIES TARGETING THE HIV ENVELOPE APEX EPITOPE

WO - 11.06.2026

Clasificación Internacional A61K 39/21N° de solicitud PCT/US2025/057683 Solicitante THE SCRIPPS RESEARCH INSTITUTE Inventor/a MA, Manyuan

The present invention provides compositions and methods for inducing broadly neutralizing antibodies (bnAbs) against HIV, specifically targeting the Apex epitope of the HIV envelope protein. The invention describes a novel immunogen, ApexGT6, an engineered HIV envelope trimer designed to bind precursors of Apex bnAbs. Methods of administering ApexGT6 as an adjuvanted protein or mRNA-lipid nanoparticle (LNP) to consistently induce Apex bnAb-related precursor B cells with long heavy chain complementarity determining region 3 (HCDR3) loops in non-human primates are disclosed. The invention further provides methods for evaluating the induced immune response, including assessing antibody affinity maturation and structural characterization of elicited antibodies. This technology represents a significant advancement in HIV vaccine development and may have broader applications in inducing antibodies with very long HCDR3s against other pathogens.

4. 20260159586 METHODS OF TREATING CANCER WITH A COMBINATION OF A CANCER VACCINE AND A TAAXCD28 BISPECIFIC ANTIGEN-BINDING MOLECULE

US - 11.06.2026

Clasificación Internacional C07K 16/28Nº de solicitud 19399807 Solicitante Regeneron Pharmaceuticals, Inc. Inventor/a David DiLillo

The present disclosure relates to methods of treating or inhibiting the growth of a tumor, wherein the methods include selecting a subject with cancer and administering to the subject in need thereof a therapeutically effective amount of a cancer **vaccine** (e.g., mRNA **vaccine** against a tumor) in combination with a bispecific antigen-binding molecule comprising a first antigen-binding domain that binds specifically CD28 and a second antigen-binding domain that binds specifically to a tumor-associated antigen (TAA). The combination therapy demonstrates increased anti-tumor efficacy, increased duration of tumor control and/or increased overall survival, as compared to a subject administered the cancer **vaccine** as monotherapy.

5. WO/2026/122786 TIM-3-TARGETED VACCINES AND METHODS OF TREATING OR PREVENTING CANCER

WO - 11.06.2026

Clasificación Internacional C12N 5/0784Nº de solicitud PCT/US2025/058091 Solicitante HENRY FORD HEALTH SYSTEM Inventor/a JIANG, Aimin

Cancer **vaccine** and methods of treating or preventing cancer involve strategic use and engineering of anti-Tim3 antibodies. The **vaccine** is advantageously a dendritic cell (DC) **vaccine**. The **vaccine** includes a composition comprising a dendritic cell targeting platform and an anti-Tim-3 antibody conjugated with a tumor antigen. Methods of treating or preventing cancer may include administering a therapeutically effective amount of the composition to a patient.

6. 4759833 PROTEIN, ADENOVIRUS UND IMPFSTOFF GEGEN INFEKTION DES SUBTYPUS DES SARS-COV-2 OMIKRON-MUTANTEN STAMMS XBB

EP - 17.06.2026

Clasificación Internacional C07K 19/00Nº de solicitud 24853641 Solicitante WESTVAC BIOPHARMA CO LTD Inventor/a WEI XIawei

The present invention relates to the field of medicine, and in particular to a protein, adenovirus and **vaccine** against infection of a subtype of SARS-CoV-2 Omicron mutant strain XBB. In order to solve the problem of lack of effective prevention and treatment drugs against infection of SARS-CoV-2 Omicron mutant strain XBB and a subtype thereof, the present invention provides a protein, adenovirus and **vaccine** against the infection of the subtype of the SARS-CoV-2 Omicron mutant strain XBB. The **vaccine** is designed by optimizing the sequences of full-length S proteins of sub-lines XBB.1.16, XBB.1.5, XBB.1.16.6, BA.2.86, EG.5, JN.1, XBB.2.3 and XBB.2 of the SARS-CoV-2 Omicron mutant strain XBB, and RBD and RBD-HR sequences in the S proteins, can help a host to resist coronavirus infection, and in particular has a good prevention and treatment effect on cross infection caused by subtype viruses of the Omicron mutant strain XBB.

7. 20260167702 HUMAN BROADLY CROSSREACTIVE INFLUENZA MONOCLONAL ANTIBODIES AND METHODS OF USE THEREOF

US - 18.06.2026

Clasificación Internacional C07K 16/108Nº de solicitud 18707795 Solicitante Dana-Farber Cancer Institute,

Inc.Inventor/a Wayne A. Marasco

The present invention provides structural determinants important for binding to the stem domain of the HA protein of influenza virus, and methods of use thereof for production of high affinity neutralizing influenza virus antibodies based upon these determinants. The present invention further provides tools for determining the efficacy of an influenza virus **vaccine**. The present invention further provides a molecular signature useful for determining the efficacy of an influenza virus **vaccine** in a subject, or for predicting prior immunologic exposure or antigen responsiveness to **vaccine** or influenza virus infection.

8.WO/2026/119274COMPOSITION FOR PRECISE EXPRESSION IN SPLEEN, **VACCINE** COMPOSITION, AND USE THEREOF

WO - 11.06.2026

Clasificación Internacional A61K 39/00Nº de solicitud PCT/CN2025/140311Solicitante RINUAGENE BIOTECHNOLOGY CO., LTD.Inventor/a ZHANG, Weiguo

Provided are a composition for precise expression in the spleen, a **vaccine** composition, and use thereof. The composition for precise expression in the spleen comprises: component (1): at least one nucleic acid, comprising a nucleotide sequence encoding a target protein and one or more miRNA targeted binding sequences; and component (2): a lipid nanoparticle composition that easily accumulates in the spleen. The composition for precise expression in the spleen can be used for preparing lipid nanoparticles or a drug or a **vaccine** composition, and can be efficiently delivered into the spleen and tissues and cells thereof and significantly reduce the expression of target genes in non-spleen organs, thereby avoiding side effects caused by undesired expression, such as inflammatory responses or other off-target effects.

9.20260166137RECOMBINANT FOOT-AND-MOUTH DISEASE TYPE A VIRUS INDUCING STRONG ADAPTIVE IMMUNE RESPONSE AND OVERCOMING INTERFERENCE FROM MATERNALLY-DERIVED ANTIBODIES, AND FOOT-AND-MOUTH DISEASE **VACCINE** COMPOSITION COMPRISING SAME

US - 18.06.2026

Clasificación Internacional A61K 39/135Nº de solicitud 18710511Solicitante REPUBLIC OF KOREA(ANIMAL AND PLANT QUARANTINE AGENCY)Inventor/a Min Ja LEE

Proposed are a foot-and-mouth disease **vaccine** composition including recombinant foot-and-mouth disease viruses and an antigen isolated and purified from the viruses. In the early stages of vaccination, a humoral immune response is simultaneously induced through the induction of a robust cellular immune response, while in the presence of maternally-derived antibodies (MDAs), B cell receptors are stimulated. Through this, it is possible to provide the **vaccine** composition that enables active immunity and overcomes the interference of the maternally-derived antibodies and to provide a method of preventing or treating foot-and-mouth diseases using the same composition.

10.20260158132SERUM FREE INTRACELLULAR PATHOGEN **VACCINE**

US - 11.06.2026

Clasificación Internacional A61K 39/205Nº de solicitud 19465533Solicitante Intervet Inc.Inventor/a Joseph Koumans

A **vaccine** composition comprising a virus antigen wherein the composition comprises less than 5% serum,

wherein the virus antigen is a whole virus or derived from a whole virus. the **vaccine** composition reduces, prevents or avoids cross-stitch spinal deformity in the treated animal

11. WO/2026/123076MERS **VACCINE** ANTIGEN

WO - 18.06.2026

Clasificación Internacional A61K 39/215Nº de solicitud PCT/AU2025/051417Solicitante MACFARLANE BURNET INSTITUTE FOR MEDICAL RESEARCH AND PUBLIC HEALTH LIMITEDInventor/a POUMBOURIOS, Pantelis

The field of the specification relates broadly to Middle East respiratory syndrome coronavirus **vaccine** (MERS-CoV) antigens and methods of using and manufacturing MERS-CoV antigens. The invention also relates to vaccines, kits, devices and strips comprising the MERS-CoV antigen. The invention also relates to ribonucleic acids encoding a S protein monomer of a coronavirus **vaccine** (MERS-CoV) antigen and methods of using and manufacturing the ribonucleic acid. The invention also relates to vectors, lipid nanoparticles, RNA vaccines, kits, devices and strips comprising the ribonucleic acid.

12. WO/2026/119273COMPOSITION FOR PRECISE LIVER EXPRESSION, **VACCINE** COMPOSITION AND USE THEREOF

WO - 11.06.2026

Clasificación Internacional C12N 15/113Nº de solicitud PCT/CN2025/140309Solicitante RINUAGENE BIOTECHNOLOGY CO., LTD.Inventor/a ZHANG, Weiguo

The present invention relates to the technical field of liver drug delivery. Provided are a composition for precise liver expression, a **vaccine** composition and the use thereof. The composition for precise liver expression contains component (1): at least one nucleic acid, which contains a nucleotide sequence encoding a target protein and one or more miRNA-targeting binding sequences; and component (2): a lipid nanoparticle composition that is easily enriched in the liver. The composition for precise liver expression can be used for preparing a lipid nanoparticle or a **vaccine** composition, can be efficiently delivered to the liver and tissue and cells thereof, and significantly reduces the expression of a target gene in non-liver organs, thereby preventing the side effects caused by undesired expression, such as inflammatory reactions or other off-target effects.

13. 20260167681**VACCINE** DELIVERY CARRIER BASED ON ADHESIVE ADJUVANT PROTEIN AND ITS USES

US - 18.06.2026

Clasificación Internacional C07K 14/435Nº de solicitud 19180607Solicitante POSTECH Research and Business Development FoundationInventor/a Hyung Joon CHA

The present invention relates to a **vaccine** delivery carrier based on an adhesive adjuvant protein and its uses. The adhesive adjuvant protein according to the present invention exhibits excellent biocompatibility and outstanding immune-enhancing effects, making it applicable not only as a **vaccine** delivery carrier for carrying and delivering target antigens but also as an adjuvant for the treatment and research of various diseases.

14. 20260158127ENHANCEMENT OF **VACCINE** EFFICACY VIA BIOMASS AND/OR RELATED MATERIAL IN ANIMAL DRINK AND FEED

US - 11.06.2026

Clasificación Internacional A61K 39/12N° de solicitud 19436778 Solicitante ZIVO BIOSCIENCE, INC. Inventor/a Andrew A. Dahl

An effective treatment method for a broad variety of diseases in both animals and humans is disclosed. The method includes combining one or more vaccines with a treatment compound to enhance **vaccine** efficacy. The disclosed treatment compound does not act directly on the pathogen, and thus the organisms cannot readily develop resistance to the treatment. When a compound such as, but not limited to, the disclosed compound is used in conjunction with one or more vaccines, a synergistic effect is realized. The suggested compound is derived from a lipopolysaccharide (LPS) of gram-negative bacteria. The treatment compound is combined with one or more appropriate vaccines and is administered early in the life of an animal to achieve a synergistic effect compared with the use of the treatment compound or the **vaccine** alone.

15. 4759302 IMMUNVERSTÄRKENDE ALUMINIUMEMULSION SOWIE HERSTELLUNGSVERFAHREN DAFÜR UND VERWENDUNG DAVON

EP - 17.06.2026

Clasificación Internacional A61K 9/107N° de solicitud 24851030 Solicitante UNIV XIAMEN Inventor/a ZHANG TIANYING

An immune-enhancing aluminum emulsion, and a preparation method therefor and a use thereof. Specifically provided are an oil-in-water emulsion containing an inorganic salt and a preparation method for the oil-in-water emulsion. Also provided are a **vaccine** adjuvant comprising the oil-in-water emulsion, a **vaccine** composition and a pharmaceutical composition. Also provided is a use of the oil-in-water emulsion as a **vaccine** adjuvant.

16. WO/2026/127385 **VACCINE** COMPOSITION FOR PREVENTING INFLUENZA VIRUS INFECTION HAVING ENHANCED NEUTRALIZING ACTIVITY

WO - 18.06.2026

Clasificación Internacional C07K 14/005N° de solicitud PCT/KR2025/018074 Solicitante POSTECH RESEARCH AND BUSINESS DEVELOPMENT FOUNDATION Inventor/a LEE, Jie-Oh

The present invention relates to a **vaccine** composition for preventing influenza virus infection, having enhanced neutralizing activity. A variant protein or fragment thereof according to the present invention has a stable pre-fusion structure, thereby enabling rapid preparation for influenza viruses with sequence variations that are difficult to predict each year. In addition, the variant protein or fragment thereof according to the present invention exhibits high neutralizing activity, thereby serving as an effective **vaccine** with superior protective ability against influenza virus. In particular, sequence optimization enables the efficient production of structurally stable proteins with high expression levels, while reducing production costs.

17. WO/2026/119879 SEDIMENTATIONS-STABILISIERTE FORMULIERUNGEN FÜR HEFE—BASIERTE VAKZINE

WO - 11.06.2026

Clasificación Internacional A61K 47/02N° de solicitud PCT/EP2025/085091 Solicitante VEROVACCINES GMBH Inventor/a RICHTER, Tilmann

Die Erfindung betritt stabile Vakzin-Formulierungen auf Basis von Hefen bzw. Hefezellen. Die Erfindung stellt insbesondere eine pharmazeutische Zusammensetzung für Hefezellsuspensionen bereit, umfassend 0 - 75 % (v/v) mindestens eines Acrylatpolymers; 0 - 5 % (m/v) mindestens eines Polymers ausgewählt aus der Gruppe bestehend aus Carboxymethylcellulose (CMC), Polyvinylpyrrolidon (PVP) und Poloxameren; Phosphat-Glycerol-Puer mit einer Phosphatkonzentration im Bereich von 1 mM - 100 mM und mit einem pH-Wert im Bereich 5 - 10, umfassend Glycerol und eine oder mehrere Verbindungen ausgewählt aus der Gruppe bestehend aus Dinatriumhydrogenphosphat, Kaliumdihydrogenphosphat und Natriumdihydrogenphosphat, wobei der Phosphat-Glycerol-Puer isoosmolar im Vergleich zu physiologischen Flüssigkeiten ist; und 0,1 - 200 mg/ml Hefezellen; wobei mindestens ein Acrylatpolymer oder mindestens ein Polymer aus der Gruppe bestehend aus Carboxymethylcellulose (CMC), Polyvinylpyrrolidon (PVP) und Poloxameren mit einem Anteil von > 0 Gew.-% in der pharmazeutischen Zusammensetzung enthalten ist, und wobei die pharmazeutische Zusammensetzung eine Impfstoffzusammensetzung ist. Die Erfindung stellt weiterhin ein dazugehörig entwickeltes Verfahren zur Messung der Suspensionsstabilität der pharmazeutischen Formulierung und ein Verfahren zur Quantifizierung von Polyacrylsäure-basiertem Adjuvans in der finalen Formulierung bereit.

18. [WO/2026/122834](#) LOW-SUGAR INFLUENZA VACCINE AND METHODS THEREOF

WO - 11.06.2026

Clasificación Internacional [C07K 14/005](#)N° de solicitud PCT/US2025/058170 Solicitante ROCK BIOMEDICAL, INC. Inventor/a LEE, Jeng-Shin

The present disclosure relates to a low glycosylated influenza hemagglutinin (HA) protein and a vaccine designed to express the HA protein in vivo. The present disclosure also teaches a method for generating an immune response by utilizing the low glycosylated HA protein, which provides a broader protection across different influenza strains or lineages.

19. [20260159574](#) SYSTEMS AND METHODS FOR THE PRODUCTION OF HUMAN POLYCLONAL ANTIBODIES

US - 11.06.2026

Clasificación Internacional [C07K 16/116](#)N° de solicitud 19379447 Solicitante SAB, LLC Inventor/a Jay HOOPER

Disclosed herein is a method for producing human antibodies against a pathogen comprising injecting a non-human animal with a pathogen-derived DNA vaccine in at least two locations of the animal; injecting the animal with an adjuvant in a location of the animal different from the location of the DNA vaccine location; collecting plasma from the animal after the injections; and purifying polyclonal antibody from the plasma.

20. [WO/2026/118752](#) METHOD FOR PREPARING NOVEL DENDRITIC CELL VACCINE AND USE THEREOF

WO - 11.06.2026

Clasificación Internacional [C12N 5/0784](#)N° de solicitud PCT/CN2025/131767 Solicitante THE FIRST AFFILIATED HOSPITAL, ZHEJIANG UNIVERSITY SCHOOL OF MEDICINE (ZHEJIANG PROVINCIAL

FIRST HOSPITAL)Inventor/a LV, Zhimin

Provided in the present invention are a method for preparing a novel dendritic cell **vaccine** and use thereof. Specifically, disclosed in the present invention is a method for preparing modified dendritic cells, comprising: culturing dendritic cells in the presence of vitamin K2, thereby obtaining dendritic cells capable of inhibiting cell death caused by tumor antigen sensitization. The modified dendritic cells can enhance T-cell activation and tumor cell killing ability. Also disclosed in the present invention is a method for tumor immunotherapy using the dendritic cells in combination with T-cell vaccines, existing tumor vaccines, immune checkpoint inhibitors, or radiotherapy and chemotherapy.

21. 20260166138 COMPOSITIONS AND METHODS FOR ENHANCEMENT OF MRNA **VACCINE** PERFORMANCE AND VACCINATION AGAINST MPOX

US - 18.06.2026

Clasificación Internacional A61K 39/275Nº de solicitud 19127070 Solicitante Yale University Inventor/a Sidi Chen

The current disclosure includes a modular **vaccine** platform. Also included are monkeypox vaccines that protect against pathogenic monkeypox (Mpox) species, as well as their variants. The vaccines typically include a modified mRNA encoding at least one immunogen, such as a viral envelope protein, cell surface binding protein, or a biologically effective/significant fragment thereof. The mRNA can be encapsulated into lipid nanoparticles or other carriers and formulated as pharmaceutical compositions that can be used to generate an immune response to pathogens, including monkeypox virus, in a subject.

22. WO/2026/122475 DNA **VACCINE** AGAINST INFECTIOUS SALMON ANEMIA VIRUS

WO - 11.06.2026

Clasificación Internacional A61K 39/12Nº de solicitud PCT/US2025/057594 Solicitante ZOETIS SERVICES LLC Inventor/a HAALAND, Espen

A DNA **vaccine** against Infectious Salmon Anemia Virus (ISAV) and methods of using same are provided.

23. WO/2026/119925 POXVIRUS **VACCINE**

WO - 11.06.2026

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2025/085175 Solicitante NEC ONCOIMMUNITY AS Inventor/a GHEORGHE, Marius

The present invention relates to **vaccine** compositions and uses thereof for the prophylactic or therapeutic treatment of an infection caused by viruses of the Poxviridae family, specifically the Orthopoxvirus genus of viruses in that viral family, which may stimulate a broad and effective adaptive immune response across multiple Poxviridae species and a diverse spectrum of human leukocyte antigen (HLA) alleles.

24. WO/2026/128689 SCHISTOSOMA **VACCINE** ANTIGEN MODIFICATIONS AND USES THEREOF

WO - 18.06.2026

Clasificación Internacional G01N 33/50Nº de solicitud PCT/US2025/059165 Solicitante TEXAS TECH

UNIVERSITY SYSTEM Inventor/a CARTER, Darrick

Provided herein are compositions, assays, kits, and methods for detecting sera that binds to, and inhibits the activity of, a recombinant calcium-activated neutral protease or calpain of *Schistosoma mansoni*, a recombinant calcium-activated neutral protease or calpain; such as a detectable substrate for a *Schistosoma mansoni* calcium-activated neutral protease or calpain; and instructions in the kit for combining sera from a subject suspected of having been exposed to *Schistosoma mansoni*, or a vaccine thereto, with the calcium-activated neutral protease or calpain, wherein the sera blocks an activity of the calcium-activated neutral protease or calpain against the detectable substrate and is a surrogate for effectiveness of the immunization to *Schistosoma mansoni*.

25. 20260158104 CONFORMATIONALLY SPECIFIC VIRAL IMMUNOGENS

US - 11.06.2026

Clasificación Internacional A61K 38/16Nº de solicitud 19326552 Solicitante Calder Biosciences Inc. Inventor/a Christopher Patrick Marshall

The present invention provides methods of making engineered viral proteins and protein complexes that are useful as vaccine immunogens, engineered viral proteins and protein complexes made using such methods, and pharmaceutical compositions comprising such engineered viral proteins and protein complexes. Such engineered viral proteins and protein complexes may comprise one or more cross-links that stabilize the conformation of an antibody epitope, such as a quaternary neutralizing antibody, and may exhibit an enhanced ability to elicit a protective immune response when administered to a subject as a component of a vaccine.

26. 20260158130 LOW-SUGAR INFLUENZA VACCINE AND METHODS THEREOF

US - 11.06.2026

Clasificación Internacional A61K 39/145Nº de solicitud 19409259 Solicitante Rock BioMedical, Inc. Inventor/a Chung-Yi WU

The present disclosure relates to a low glycosylated influenza hemagglutinin (HA) protein and a vaccine designed to express the HA protein in vivo. The present disclosure also teaches a method for generating an immune response by utilizing the low glycosylated HA protein, which provides a broader protection across different influenza strains or lineages.

27. WO/2026/119376 SEDIMENTATIONS-STABILISIERTE FORMULIERUNGEN FÜR HEFE-BASIERTE VAKZINE

WO - 11.06.2026

Clasificación Internacional A61K 47/02Nº de solicitud PCT/EP2024/084517 Solicitante VEROVACCINES GMBH Inventor/a RICHTER, Tilman

Die Erfindung betrifft stabile Vakzin-Formulierungen auf Basis von Hefen bzw. Hefezellen. Die Erfindung stellt insbesondere eine pharmazeutische Zusammensetzung für Hefezellsuspensionen bereit, umfassend 0 - 75 % (v/v) mindestens eines Acrylatpolymers; 0 - 5 % (m/v) mindestens eines Polymers ausgewählt aus der Gruppe bestehend aus Carboxymethylcellulose (CMC), Polyvinylpyrrolidon (PVP) und Poloxameren; Wasser oder Phosphatpuffer (1 mM - 100 mM, pH 5 - 10) umfassend eine oder mehrere Verbindungen ausgewählt

aus der Gruppe bestehend aus Dinatriumhydrogenphosphat, Kaliumdihydrogenphosphat und Natriumdihydrogenphosphat; und 0,1 - 200 mg/ml Hefezellen; wobei mindestens ein Acrylatpolymer oder mindestens ein Polymer aus der Gruppe bestehend aus Carboxymethylcellulose (CMC), Polyvinylpyrrolidon (PVP) und Poloxameren mit einem Anteil von > 0 Gew.-% in der pharmazeutischen Zusammensetzung enthalten ist. Die Erfindung stellt weiterhin ein dazugehörig entwickeltes Verfahren zur Messung der Suspensionsstabilität der pharmazeutischen Formulierung und ein Verfahren zur Quantifizierung von Polyacrylsäure-basiertem Adjuvans in der finalen Formulierung bereit.

28. [20260158109](#) CD200AR LIGANDS FOR CANCER IMMUNOTHERAPY

US - 11.06.2026

Clasificación Internacional [A61K 38/17N](#)° de solicitud 19307588 Solicitante REGENTS OF THE UNIVERSITY OF MINNESOTA Inventor/a Michael OLIN

The present invention in certain embodiments provides a method of inhibiting PD-1 in a cell by administering a CD200 activation receptor ligand (CD200AR-L) to the cell. The present invention in certain embodiments provides a method of enhancing efficacy of a tumor lysate **vaccine** in a mammal comprising administering a CD200 activation receptor ligand (CD200AR-L) to the mammal prior to the administration of the tumor lysate **vaccine**.

29. [WO/2026/119878](#) **VACCINE** FOR INDUCING ANTIBODIES TO INHIBIT NATIVE INTERLEUKIN-11 SIGNALING

WO - 11.06.2026

Clasificación Internacional [A61K 39/00N](#)° de solicitud PCT/EP2025/085090 Solicitante ADAPTVAC APS Inventor/a BERTELSEN, Adam Frederik Sander

The present disclosure pertains to antigens comprising an epitope of IL-11, fused to a peptide tag capable of forming an isopeptide bond. Additionally, the disclosure encompasses vaccines containing said antigen, their uses, and methods of manufacturing. The IL-11 antigen is designed in such a way that the **vaccine** induces antibodies which neutralise IL-11 by preventing its interaction with IL-11R α but does not induce gp130 signaling.

30. [20260158125](#) PEPTIDE **VACCINE**

US - 11.06.2026

Clasificación Internacional [A61K 39/00N](#)° de solicitud 18676282 Solicitante ARGONAUT THERAPEUTICS LIMITED Inventor/a Nicholas LA THANGUE

The present invention provides one or more immunogenic peptides derived from a PRMT5-E2F1 axis regulated long non-coding RNA gene or a derivative thereof; a pharmaceutical composition comprising one or more of said peptides; a **vaccine** comprising one or more of said peptides and their use in therapy, including a method for eliciting an immune response in a mammalian subject by administration of an agent capable of presenting the peptides to the host. The invention also relates to the use of a PRMT5 inhibitor for use in treating cancer by stimulating host immunity.

31. [20260168003](#) METHOD AND SYSTEM FOR PRODUCING A MODIFIED PROTEIN

US - 18.06.2026

Clasificación Internacional C12P 21/02Nº de solicitud 19422655 Solicitante YEDA RESEARCH AND DEVELOPMENT CO. LTD. Inventor/a Sarel-Jacob FLEISHMAN

The present invention relates to biological sciences, specifically to the design and stabilization of proteins for use in diagnostics and **vaccine** development. The present invention represents a system and method for producing a modified protein that provides an improvement of the technological field of protein design and stabilization by providing tools that enable the efficient design and production of proteins with enhanced stability characteristics compared to the parental protein, while retaining immunogenicity to naturally acquired human monoclonal antibodies. In particular, the suggested invention is efficiently applicable to proteins with large and topologically complex structures that have limited natural diversity of homologs and circumvent the requirement for a deep multiple-sequence alignment.

32. WO/2026/123993 FATTY ACID-MODIFIED DIPHTHERIA TOXIN ANTIGEN AND USE THEREOF

WO - 18.06.2026

Clasificación Internacional C07K 14/34Nº de solicitud PCT/CN2025/130761 Solicitante ACADEMY OF MILITARY MEDICAL SCIENCES, PEOPLE'S LIBERATION ARMY ACADEMY OF MILITARY SCIENCES Inventor/a LIU, Bo

The present invention relates to the field of biomedicine and specifically to a fatty acid-modified diphtheria toxin antigen and a use thereof. In the diphtheria toxin antigen containing fatty acid modification, the fatty acid modification comprises a fatty acid-modified N-terminal cysteine (Cys), and the fatty acid-modified N-terminal Cys is Cys in which the sulfhydryl group is modified by diacylglycerol and/or the alpha-amino group is modified by a fatty acyl group. Compared with existing diphtheria toxin protein antigens, the fatty acid-modified diphtheria toxin antigen of the present invention not only achieves faster conventional intramuscular injection induction and a higher immune protection effect than the existing diphtheria toxin antigens, but can also induce respiratory mucosal immunity, providing a basis for a **vaccine** for immune prevention to block diphtheria infection and diffusion.

33. WO/2026/123468 USE OF VIRUS IN PREPARATION OF PHAGE

WO - 18.06.2026

Clasificación Internacional C12N 7/00Nº de solicitud PCT/CN2025/079024 Solicitante SOUTH CHINA AGRICULTURAL UNIVERSITY Inventor/a SONG, Changxu

Provided in the present application is use of a virus in the preparation of a phage, pertaining to the technical field of phages. Provided in the present application is use of PCV2, phPCV2, PCV3, phPCV3, HEV, or phHEV in the preparation of a phage. The present application also prepares and obtains a RecA-deficient Bacillus subtilis live vector **vaccine**.

34. 20260159563 USE OF CCL11

US - 11.06.2026

Clasificación Internacional C07K 14/52Nº de solicitud 18722449 Solicitante NEWISH TECHNOLOGY (BEIJING) CO., LTD. Inventor/a Hailong QI

The disclosure relates to the technical field of **vaccine** preparation, and in particular to an immune-enhancing delivery system formed by targeted antigen delivery by CCL11. The system further enhances immunogenicity by fusing a chemokine CCL11 with a corresponding antigen molecule, and adding a T2 label at a terminal of the antigen molecule. The system can be a nucleic acid vector or a fusion protein or the like to be applied to prevention and/or treatment of diseases caused by a corresponding antigen. According to the present invention, by utilizing a chemotactic binding capacity of CCL11 with a surface receptor of an immune cell such as a DC, different antigen proteins are transported to the surface of the DC, so that the efficiency of phagocytosis, processing and presentation of the DC on various antigen proteins is improved, and the effect of preventing and treating related diseases is improved.

35. 20260167942ADJUVANT FOR INTRADERMAL IMMUNIZATION AND USE THEREOF

US - 18.06.2026

Clasificación Internacional C12N 7/00Nº de solicitud 19536292Solicitante JIANGSU ACADEMY OF AGRICULTURAL SCIENCESInventor/a Bihua DENG

An adjuvant for intradermal immunization and a use thereof are provided. The adjuvant includes the following components in parts by mass: 10 parts to 25 parts of squalene, 10 parts to 25 parts of sea buckthorn oil, 25 parts to 35 parts of caprylocaproyl polyoxyl-8 glycerides, and 25 parts to 45 parts of Tween 80. The adjuvant for intradermal immunization has a low viscosity, is favorable for injection and absorption, and can significantly enhance the immunization efficacy of a **vaccine**.

36. WO/2026/123991IMMUNE COMPOSITION CONTAINING FATTY ACID-MODIFIED SPYCATCHER PROTEIN AND PREPARATION METHOD THEREFOR

WO - 18.06.2026

Clasificación Internacional C07K 14/315Nº de solicitud PCT/CN2025/130759Solicitante ACADEMY OF MILITARY MEDICAL SCIENCES, PEOPLE'S LIBERATION ARMY ACADEMY OF MILITARY SCIENCESInventor/a WU, Jun

Provided are an immune composition containing a fatty acid-modified SpyCatcher protein and a preparation method thereof, where the SpyCatcher protein (LipoSC) can self-assemble into nanoparticles and covalently link to antigens containing a SpyTag peptide through an isopeptide bond; also provided is an immune composition containing the fatty acid-modified SpyCatcher protein and a coronavirus RBD antigen, which can induce a strong RBD-specific immune response, particularly as a mucosal subunit **vaccine**, capable of inducing antigen-specific mucosal and humoral immunity. Further provided are a method for completing fatty acid modification of the SpyCatcher protein in Escherichia coli, and a method for preparing various immune compositions using this fatty acid-modified protein.

37. WO/2026/123994FATTY ACID-MODIFIED CORONAVIRUS ANTIGEN AND USE THEREOF

WO - 18.06.2026

Clasificación Internacional C07K 19/00Nº de solicitud PCT/CN2025/130762Solicitante ACADEMY OF MILITARY MEDICAL SCIENCES, PEOPLE'S LIBERATION ARMY ACADEMY OF MILITARY SCIENCESInventor/a WU, Jun

Disclosed in the present invention are a fatty acid-modified coronavirus antigen and use thereof, pertaining to the field of biomedicine. The fatty acid-modified coronavirus antigen comprises a fatty acid-modified N-

terminal cysteine (Cys) and a coronavirus antigen, and may comprise a linking domain. The coronavirus antigen is a coronavirus spike protein (S protein) receptor-binding domain (RBD). The fatty acid-modified coronavirus S protein RBD can self-assemble into nanoparticles. Compared with an RBD without fatty acid modification, the antigen can induce faster and higher humoral immunity by means of injection immunization, and can also induce mucosal immunity and humoral immunity by means of respiratory nebulization immunization, making it a candidate mucosal [vaccine](#) with potential medical value.

38. [WO/2026/128536](#) LIPOSOMAL ADJUVANT COMPOSITIONS FOR EPSTEIN BARR VIRUS VACCINES

WO - 18.06.2026

Clasificación Internacional [A61K 39/12N](#)° de solicitud PCT/US2025/058884 Solicitante MERCK SHARP & DOHME LLC Inventor/a AHL, Patrick L.

The present disclosure provides a [vaccine](#) composition that comprises an Epstein Barr Virus (EBV) polypeptide and a liposomal adjuvant, and methods of inducing an immune response to an Epstein Barr Virus (EBV) or methods of preventing infection of or reducing the likelihood of infection by an Epstein Barr Virus (EBV) using the compositions, or a combination of the EBV polypeptide and the liposomal adjuvant.

39. [WO/2026/128401](#) LIPOSOMAL ADJUVANT COMPOSITIONS FOR HUMAN PAPILLOMAVIRUS VACCINES

WO - 18.06.2026

Clasificación Internacional [A61K 9/1272N](#)° de solicitud PCT/US2025/058671 Solicitante MERCK SHARP & DOHME LLC Inventor/a AHL, Patrick L.

The present disclosure provides a [vaccine](#) composition that includes, among other things, HPV virus-like particles (VLPs) of at least one type of human papillomavirus (HPV) selected from the group consisting of HPV types: 6, 11, 16, 18, 26, 31, 33, 35, 39, 45, 51, 52, 53, 55, 56, 58, 59, 66, 68, 73, and 82 and a liposomal adjuvant, and methods of inducing an immune response to HPV or methods of preventing infection of or reducing the likelihood of infection by HPV using the compositions.

40. [WO/2026/127647](#) NOVEL RECOMBINANT ANTIGEN PROTEIN FOR JAPANESE ENCEPHALITIS, AND USE THEREOF

WO - 18.06.2026

Clasificación Internacional [C07K 14/005N](#)° de solicitud PCT/KR2025/021307 Solicitante REPUBLIC OF KOREA (KOREA DISEASE CONTROL AND PREVENTION AGENCY) Inventor/a KIM, Hyeon Guk

The present invention relates to a novel recombinant antigen protein for Japanese encephalitis and, more specifically, to a recombinant antigen protein for Japanese encephalitis, designed through protein structural stability analysis and introduction of point mutations. It has been identified that a composition comprising the recombinant antigen protein for Japanese encephalitis, according to the present invention, exhibits immunogenicity and protective efficacy against challenge inoculation that are superior to those of an approved live-attenuated [vaccine](#). In particular, it has been experimentally identified that the recombinant antigen protein for Japanese encephalitis, of the present invention, exhibits excellent protective efficacy against Japanese encephalitis virus G5 and other genotypes. Therefore, the recombinant antigen protein for

Japanese encephalitis, of the present invention, can be variously used in the field of preventing infection by Japanese encephalitis virus G1 to G5.

41. [WO/2026/118253](#) ANTI-PD1 MRNA NANOFORMULATION, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 11.06.2026

Clasificación Internacional [A61K 9/51](#)Nº de solicitud PCT/CN2025/080443 Solicitante HUANXIN BIOTECHNOLOGY (TAIZHOU) CO., LTD. Inventor/a ZOU, Jianhua

An anti-PD1 mRNA nanoformulation, a preparation method therefor, and use thereof. The anti-PD1 mRNA nanoformulation comprises: an anti-PD1 mRNA and a delivery nanocarrier, wherein the anti-PD1 mRNA is loaded in the delivery nanocarrier. The anti-PD1 mRNA nanoformulation can enable autologous generation of an anti-PD1 antibody to block the PD1 receptor on T cells, inhibiting the depletion of cytotoxic CD8 T cells.

42. [4759827](#) NUKLEINSÄUREIMPFFSTOFF GEGEN DAS SARS-COV-2-CORONAVIRUS

EP - 17.06.2026

Clasificación Internacional [C07K 14/005](#)Nº de solicitud 26175581 Solicitante PASTEUR INSTITUT Inventor/a SIMON-LORIERE ETIENNE

43. [20260165979](#) LIPID NANOPARTICLES

US - 18.06.2026

Clasificación Internacional [A61K 9/51](#)Nº de solicitud 19109341 Solicitante KYUSHU UNIVERSITY, NATIONAL UNIVERSITY CORPORATION Inventor/a Go HIRAI

Candidate molecules may be constituent components of various lipid nanoparticles. A C-glycoside glycolipid compound of formula (I):

or formula (II):

a lipid nanoparticle comprising the same, or a pharmaceutical composition, particularly a **vaccine**, which comprises the lipid nanoparticle.

44. [WO/2026/119880](#) **VACCINE** FOR INDUCING ANTIBODIES TO INHIBIT THYMIC STROMAL LYMPHOPOIETIN

WO - 11.06.2026

Clasificación Internacional [A61K 39/00](#)Nº de solicitud PCT/EP2025/085092 Solicitante ADAPT VAC APS Inventor/a BERTELSEN, Adam Frederik Sander

The present disclosure pertains to antigens comprising an epitope of TSLP, fused to a peptide tag capable of

forming an isopeptide bond. Additionally, the disclosure encompasses vaccines containing said antigen, their uses, and methods of manufacturing.

45. 20260160759 SYSTEMS AND METHODS FOR MULTIPLEX DETECTION OF BIOMARKERS

US - 11.06.2026

Clasificación Internacional G01N 33/543Nº de solicitud 19270869 Solicitante Veravas, Inc. Inventor/a Joshua Caine Soldo

Provided herein are methods and systems for multiplex detection and/or measurement of biomarkers of a sample. The methods and systems can be used for rapid disease detection and/or monitoring, vaccine efficacy and immune response monitoring, therapeutic drug monitoring, and/or therapeutic safety and efficacy monitoring.

46. WO/2026/121908 LIPID NANOPARTICLES FOR PREVENTING INFLUENZA VIRUS INFECTION AND PREPARATION METHOD THEREFOR

WO - 11.06.2026

Clasificación Internacional A61K 9/51Nº de solicitud PCT/KR2025/020841 Solicitante SOGANG UNIVERSITY RESEARCH & BUSINESS DEVELOPMENT FOUNDATION Inventor/a KIM, Hyun Cheol

The present invention relates to a lipid nanoparticle and a method for preparing same, wherein melittin, a peptide capable of directly penetrating cell membranes and facilitating mucus layer penetration, is conjugated with a biocompatible lipid to form a melittin-lipid conjugate and is used in combination with an ionizable lipid, a helper lipid, cholesterol, and a lipid-PEG to prepare the lipid nanoparticle, whereby the lipid nanoparticle exhibits high stability and low toxicity while effectively delivering influenza A virus mRNA via intranasal administration to induce a prophylactic vaccine effect.

47. 20260158134 COMPOSITIONS SUITABLE FOR USE IN A METHOD FOR ELICITING CROSS-PROTECTIVE IMMUNITY AGAINST CORONAVIRUSES

US - 11.06.2026

Clasificación Internacional A61K 39/215Nº de solicitud 18707694 Solicitante King Abdullah University of Science and Technology Inventor/a Jasdave Chahal

Immunogenic compositions and methods of use thereof, for eliciting an immune response against multiple coronaviruses, using a single vaccine composition are described. The compositions include an antigen from more than one pathogen, for example, more than one member of the β -coronavirus family, for example, SARS-CoV and MERS-CoV. Exemplary antigens include the receptor binding domain (RBD) of the coronavirus spike protein or a fragment thereof. The disclosed compositions are administered to a subject in need therefore, to generate an immune response against more than one pathogen, represented by the source of the antigens in the construct.

48. WO/2026/119906 NEW VACCINE

WO - 11.06.2026

Clasificación Internacional A61K 39/395Nº de solicitud PCT/EP2025/085140 Solicitante LETI PHARMA,

S.L.U.Inventor/a NAVARRO DIEZ, Gemma

Several novel engineered variants of Protein Q, or fragments thereof, are provided, mediating protective immune responses in the treatment and prevention of leishmaniasis in animal and in human subjects.

49. [WO/2026/126165](#) MATERIALS, METHODS AND SYSTEMS FOR STIMULATING HOST IMMUNE RESPONSES

WO - 18.06.2026

Clasificación Internacional [A61K 39/39](#)Nº de solicitud PCT/IB2025/062816 Solicitante JANSSEN BIOTECH, INC. Inventor/a HABTE, Habtom H.

The present disclosure relates to inter alia novel designs, compositions, systems and formulations, etc. of nucleic acid(s) (polynucleotide or oligonucleotide, etc.) products, specifically compositions and formulations of nucleic acid immunity inducing agents, such as adjuvants, immunization and or **vaccine** products and accessory products and nucleic acid gene therapy products, and the like. The compositions of the disclosure enhance immunity, and provide nucleic acid based medicinal and pharmaceutical products. When introduced directly into a host, such as intradermal cells and or tissues, the embodiments described herein induces production of immune responses which specifically recognize targets, such as human targets.

50. [WO/2026/128222](#) METHODS FOR STABILIZING HENIPAVIRUS SURFACE GLYCOPROTEINS AND USES THEREOF

WO - 18.06.2026

Clasificación Internacional [C07K 14/115](#)Nº de solicitud PCT/US2025/057106 Solicitante DUKE UNIVERSITY Inventor/a MAY, Aaron

Described are structural identifications of the F and G proteins of a large variety Henipaviruses and their use in creating immunogens useful for **vaccine** compositions, and in prevention and treatment of Paramyxovirus and Henipavirus infections in a mammal. Also described are methods for screening viral fusion proteins for capability to transition from a pre-fusion to a post-fusion conformation. A monoclonal antibody specific for LayV-F is also provided.

51. [20260159553](#) MULTICISTRON EXPRESSION VECTOR FOR COVID-19 **VACCINE**

US - 11.06.2026

Clasificación Internacional [C07K 14/005](#)Nº de solicitud 18707293 Solicitante Kashiv BioSciences, LLC Inventor/a Sudharti Gupta

The present invention provides an expression vector comprises gene of interest encode more than one structural protein to enhance immune responses against Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-COV-2) and its variants. Furthermore, the expression vector to produce mRNA expresses more than one structural protein to generate immune response against Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-COV-2) and its variants.

52. [20260158131](#) ROTAVIRUS B VIRUS-LIKE-PARTICLE COMPOSTIONS AND ANTIBODY ASSAYS

US - 11.06.2026

Clasificación Internacional A61K 39/15N° de solicitud 19387411 Solicitante University of Kentucky Research Foundation Inventor/a Feng Li

Compositions and methods are provided for immunization against equine rotavirus group B (ERVB). The compositions comprise ERVB structural proteins VP4, VP2, VP6, and VP7, or VP2, VP6, VP7, and a modified VP4, which can be assembled into virus-like particles (VLPs). The invention includes immunogenic compositions and vaccines for administration to non-human animals, including pregnant mares to confer passive immunity to foals. Methods of stimulating immune responses, preventing ERVB spread, conferring passive immunity, evaluating vaccine efficacy, detecting ERVB antibodies, and producing VLPs in baculovirus-infected insect cells are disclosed. The compositions may further include inactivation agents. Nucleotides and vectors encoding the ERVB proteins are also provided.

53. 20260167675 VIRAL PEPTIDES AND USES THEREOF

US - 18.06.2026

Clasificación Internacional C07K 14/005N° de solicitud 18871778 Solicitante Regeneron Pharmaceuticals, Inc. Inventor/a Augustine CHOY

The present disclosure provides isolated peptides derived from hepatitis B virus (HBV), peptide-based molecules (e.g., peptide-MHC (pMHC) complexes), polynucleotides and vectors encoding the peptides or peptide-based molecules, pharmaceutical compositions (e.g., vaccine compositions), and their use for treatment or prevention of HBV infection and/or HBV-induced diseases. The present disclosure also provides binding moieties that bind to the peptides or peptide-based molecules disclosed herein, and their use for treatment or prevention of HBV infection and/or HBV-induced diseases. The present disclosure further provides methods and systems for identifying immunogenic virus-derived peptides.

54. WO/2026/120145 VACCINE COMPOSITION COMPRISING HERV DERIVED EPITOPES AND THEIR USE IN TREATMENT OF AML

WO - 11.06.2026

Clasificación Internacional A61K 39/00N° de solicitud PCT/EP2025/085671 Solicitante EVAXION A/S Inventor/a KLAUSEN, Michael Schantz

Provided are immunogenic agents and compositions tailored to induce immunity against hERV expression products in a high percentage of cancer patients as well. Also provided are methods for immune induction against cancer cells

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

Estrategia de búsqueda: *vaccine.ti. AND @PD>="20260611"<=20260621 22 records*

Document ID	Title	Inventor	Applicant Name
US 20260167923 A1	IMMUNOGENIC ENGINEERED LIVE-ATTENUATED GRAM-NEGATIVE PATHOGEN VACCINES HAVING REDUCED ENDOTOXICITY	Ernst; Robert K. et al.	University of Maryland, Baltimore, The Government of The United States, as represented by the Defense Health Agency
US 20260166137 A1	RECOMBINANT FOOT-AND-MOUTH DISEASE TYPE A VIRUS INDUCING STRONG ADAPTIVE IMMUNE RESPONSE AND OVERCOMING INTERFERENCE FROM MATERNALLY-DERIVED ANTIBODIES, AND FOOT-AND-MOUTH DISEASE VACCINE COMPOSITION COMPRISING SAME	LEE; Min Ja et al.	REPUBLIC OF KOREA(ANIMAL AND PLANT QUARANTINE AGENCY)
US 20260166138 A1	Compositions and Methods for Enhancement of mRNA Vaccine Performance and Vaccination against Mpox	Chen; Sidi et al.	Yale University
US 20260166134 A1	mRNA, mRNA VACCINE FORMULATION, AND PREPARATION METHOD AND USE THEREOF	NIU; Yuanjie et al.	Panru Biotechnology (Tianjin) Co., Ltd
US 20260167681 A1	VACCINE DELIVERY CARRIER BASED ON ADHESIVE ADJUVANT PROTEIN AND ITS USES	CHA; Hyung Joon et al.	POSTECH Research and Business Development Foundation
US 12655477 B2	DNA vaccines	Hill; Vanessa	Touchlight IP Limited
US 12653872 B2	Mimotopes of alpha-synuclein and vaccines thereof for the treatment of synucleinopathy	Mandler; Markus et al.	AC Immune SA
US 12653878 B2	Vaccine composition against Streptococcus suis infection	Seele; Jana et al.	CEVA SANTE ANIMALE S.A.

US 12653876 B2	M Hyo multivalent vaccine and uses thereof	Wilson; Keith et al.	BOEHRINGER INGELHEIM ANIMAL HEALTH USA INC.
US 12653877 B2	Gut bacteria derived microvesicles for vaccine delivery	Stentz; Regis et al.	Quadram Institute Bioscience
US 12653875 B2	Live attenuated Leishmania parasite vaccines with enhanced safety characteristics	Nakhasi; Hira L. et al.	The U.S.A., as represented by the Secretary, Department of Health and Human Services, Ohio State Innovation Foundation, Matlashewski; Gregory, Zhang; Wen-Wei, Lypaczewski; Patrick
US 12653881 B2	Measles virus vaccine expressing SARS-CoV-2 protein(s)	Duprex; William Paul et al.	University of Pittsburgh-Of the Commonwealth System of Higher Education
US 12653874 B2	Nant cancer vaccine	Soon-Shiong; Patrick et al.	NantCell, Inc., Nant Holdings IP, LLC
US 20260158130 A1	LOW-SUGAR INFLUENZA VACCINE AND METHODS THEREOF	WU; Chung-Yi et al.	Rock BioMedical, Inc.
US 20260158132 A1	SERUM FREE INTRACELLULAR PATHOGEN VACCINE	Koumans; Joseph et al.	Intervet Inc.
US 20260158125 A1	PEPTIDE VACCINE	LA THANGUE; Nicholas et al.	ARGONAUT THERAPEUTICS LIMITED
US 20260158127 A1	ENHANCEMENT OF VACCINE EFFICACY VIA BIOMASS AND/OR RELATED MATERIAL IN ANIMAL DRINK AND FEED	Dahl; Andrew A. et al.	ZIVO BIOSCIENCE, INC.
US 20260159553 A1	MULTICISTRON EXPRESSION VECTOR FOR COVID-19 VACCINE	Gupta; Sudharti et al.	Kashiv BioSciences, LLC
US 20260158135 A1	VARIANT STRAIN-BASED CORONAVIRUS VACCINES AND USES THEREOF	Metkar; Mihir et al.	ModernaTX, Inc.
US 20260158129 A1	NUCLEIC ACID INFLUENZA VACCINES AND RESPIRATORY VIRUS COMBINATION VACCINES	Nachbagauer; Raffael et al.	ModernaTX, Inc.

US 20260158126 A1	RUMINAL AND METHANOGEN VACCINES AND USES THEREOF	Rajaniemi; Hannu et al.	HELIX NANOTECHNOLOGIES, INC.
US 20260159586 A1	METHODS OF TREATING CANCER WITH A COMBINATION OF A CANCER VACCINE AND A TAAxCD28 BISPECIFIC ANTIGEN-BINDING MOLECULE	DiLillo; David et al.	Regeneron Pharmaceuticals, Inc.

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