



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.

- Un nuevo enfoque de vacunas basadas en proteínas podría permitir respuestas más rápidas a la Enfermedad X.
- Noticias más recientes en la Web sobre vacunas.
- Artículos científicos más recientes de Medline sobre vacunas.
- Patentes más recientes en Patentscope sobre vacunas.
- Patentes más recientes en USPTO sobre vacunas

Un nuevo enfoque de vacuna basadas en proteínas podría permitir respuestas más rápidas a la Enfermedad X

Una nueva investigación está destinada a investigar un enfoque pionero que utiliza tecnología de marcado de proteínas en nanopartículas, lo que podría acelerar la producción de vacunas contra amenazas epidémicas y pandémicas, incluyendo una futura "Enfermedad X". El diseño de las nanopartículas también podría facilitar una administración de vacunas más potente y específica, y ayudar a reforzar la respuesta inmunitaria.

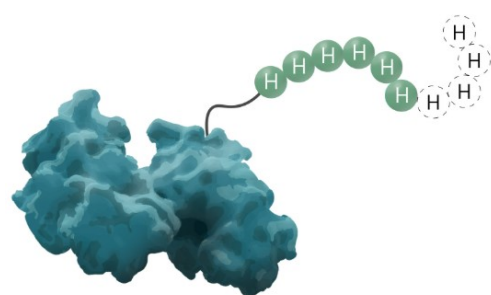
El equipo de expertos, con sede en la empresa biotecnológica privada POP Biotechnologies, Inc. (POP BIO), recibirá hasta 1,5 millones de dólares estadounidenses de CEPI para avanzar en la investigación de la plataforma de vacunas basadas en proteínas SNAP™ (Particulación Espontánea de Antígenos en Nanoliposomas) de POP BIO.



Esta tecnología ha sido diseñada para desarrollar rápidamente los candidatos vacunales potentes en nanopartículas, a la vez que purifica antígenos. Se añaden a la vacuna fragmentos de proteínas inertes de un patógeno causante de enfermedad, conocidos como antígenos, para enseñar al sistema inmunitario a reconocerlo y defenderse.

“Los antígenos añadidos a las formulaciones de vacunas deben purificarse para garantizar su seguridad, eficacia y consistencia. Sin embargo, los métodos de purificación convencionales pueden ser costosos e implicar múltiples pasos, que suelen durar varios días, lo que ralentiza el proceso de desarrollo de la vacuna”, afirmó el Dr. Kent Kester, Director Ejecutivo de I+D de Vacunas de la Coalición para las Innovaciones en Preparación para Epidemias (CEPI).

Mediante un enfoque innovador, la tecnología patentada de SNAP utiliza una pequeña etiqueta proteica, denominada polihistidina o His6, que se adhiere a los antígenos de la vacuna (para una técnica de purificación comúnmente utilizada en la investigación básica de proteínas) y también se utiliza para integrarlos en pequeños liposomas esféricos combinados con cobalto metálico para favorecer la respuesta inmunitaria. Gracias a su excelente relación costo-beneficio, se le conoce como una de las etiquetas de afinidad proteica más utilizadas hasta la fecha.



Por su pequeño tamaño, que varía entre cuatro y diez aminoácidos, rara vez interfiere con la función de la proteína diana. Además, su baja inmunogenicidad y versatilidad en condiciones nativas y desnaturizantes la distinguen de otras. El principio de purificación subyacente de la etiqueta His es la interacción entre los iones metálicos y el anillo imidazol de la histidina.

En comparación con los métodos de purificación tradicionales, esta técnica mejorada podría eliminar rápidamente los posibles contaminantes de los antígenos de la vacuna en tan solo 30 minutos.

“Como cada día cuenta durante un brote incipiente, un desarrollo más rápido de vacunas purificadas podría ayudar a contener con mayor rapidez una amenaza viral nueva o reemergente de rápida propagación antes de que alcance proporciones pandémicas, de acuerdo con la Misión de los 100 Días”, afirma el Dr. Kester.

El versátil diseño “plug and play” de SNAP también podría ser beneficioso en un brote, ya que el antígeno procedente del patógeno causante de la enfermedad podría integrarse fácil y rápidamente en la plataforma de la vacuna para un desarrollo y despliegue más rápidos de las vacunas. Con menos etapas y menos

complejas involucradas en su proceso de purificación, la tecnología SNAP también podría ser más eficiente que los métodos tradicionales de purificación de vacunas basadas en proteínas, produciendo mayores rendimientos de antígenos que ayudan a ampliar los suministros de vacunas disponibles y permiten inmunizar a más personas durante un brote.

“El uso dual de la etiqueta histidina para facilitar la purificación de proteínas y luego convertirlas de forma fácil, rápida y estable en potentes candidatos vacunales en nanopartículas ha sido fundamental para el impulso que POP BIO ha alcanzado”, afirma Jonathan Lovell, cofundador de dicha compañía biotecnológica. “Es importante destacar que, cuando se utiliza la etiqueta his para anclar proteínas en la superficie de las nanopartículas SNAP, no se produce una respuesta inmunitaria significativa contra la etiqueta en sí, como se observó en pruebas clínicas en humanos”.

La tecnología SNAP ya se ha probado en un ensayo a gran escala de fase III de etapa avanzada de la vacuna EuCorVac-19 contra la COVID-19, donde se obtuvieron resultados positivos de seguridad y eficacia. A principios de este año, la revista Cell Biomaterials publicó resultados positivos para una vacuna contra la gripe aviar H5N1 desarrollada con la plataforma SNAP de POP BIO en modelos preclínicos.

El proyecto, explorará primero el potencial de la tecnología para proteger contra el síndrome de fiebre grave con trombocitopenia (SFTS), una enfermedad viral transmitida por garrapatas que afecta a Asia Oriental, en ensayos preclínicos. Liderado por POP BIO, el proyecto será llevado a cabo por un equipo internacional de investigadores de la Universidad de Buffalo y la Universidad Médica SUNY Upstate en EE. UU., así como del Instituto Nacional de Enfermedades Infecciosas, dentro del Instituto Japonés para la Seguridad Sanitaria. CEPI está explorando opciones para seleccionar antígenos para las pruebas de sus socios actuales, incluyendo aquellos que utilizan IA para el diseño de antígenos.

Si el proyecto tiene éxito, la plataforma SNAP tiene el potencial de adaptarse rápidamente para desarrollar candidatos vacunales contra otros patógenos, incluyendo la Enfermedad X.

CEPI y POP BIO se comprometen a facilitar el acceso equitativo a los resultados de su colaboración, de acuerdo con la Política de Acceso Equitativo de CEPI. Los resultados del proyecto, incluyendo los datos relacionados, se publicarán en acceso abierto para beneficio de la comunidad científica mundial.

Acerca de POP Biotechnologies

POP Biotechnologies, Inc. es una empresa biotecnológica privada que desarrolla nuevas terapias y vacunas basadas en su tecnología patentada de liposomas de porfirina-fosfolípido (PoP). La plataforma PoP, con licencia exclusiva de la Fundación de Investigación de la Universidad Estatal de Nueva York (SUNY-RF), fue inventada por el cofundador, el Dr. Jonathan Lovell, en SUNY Buffalo. POP Biotechnologies opera desde la incubadora de SUNY Buffalo en el Parque de Investigación Baird.

Acerca de la tecnología SNAP™ de POP BIO

La tecnología SNAP™ (Particulación Espontánea de Antígenos Nanoliposomales) de POP BIO permite el desarrollo rápido de potentes vacunas basadas en partículas mediante el uso de un sistema de liposomas PoP modificado con cobalto (CoPoP). Esta plataforma permite la presentación estable y espontánea de antígenos proteicos y peptídicos en liposomas, lo que mejora sustancialmente la inmunogenicidad.

Fuentes:

CEPI. Disponible en <https://n9.cl/6mvvg>

Cube Biotech. The His-Tag: Fundamentals And Principles. Disponible en <https://n9.cl/lrlfk>

POP BIO. Introducing SNAP: Spontaneous Nanoliposome Antigen Particles. Disponible en <https://n9.cl/9xy71>

Noticias en la Web

Study Estimates COVID-19 Vaccinations Prevented More Than 2.5 Million Deaths in 2020 to 2024

Aug 18. COVID-19 vaccination during 2020 to 2024 was beneficial, but estimates are more conservative than previous calculations that focused on the first year of vaccinations, according to a study published online July 25 in JAMA Health Forum.

John P.A. Ioannidis, M.D., D.Sc., from the Stanford University School of Medicine in California, and colleagues calculated lives and life-years saved by COVID-19 vaccination worldwide from the onset of vaccination campaigns through Oct. 1, 2024, in a comparative effectiveness study.

The researchers found that more than 2.5 million deaths were averted in the main analysis (one death averted per 5,400 vaccine doses administered); 82 percent were among people vaccinated before any infection. A suggested 1.4 to 4.0 million lives were saved in sensitivity analyses. A preponderance of benefit was seen during the pre-omicron period in some sensitivity analyses. An estimated 14.8 million life-years were saved, with one life-year saved per 900 vaccine doses; the sensitivity range was 7.4 to 23.6 million life-years. Overall, 76 percent of life-years saved were among people aged 60 years or older; only 2 percent of the total was contributed by long-term care residents. Very small contributions to the total benefit were seen for children and adolescents and young adults aged 20 to 29 years (0.01 and 0.07 percent of lives saved, respectively; 0.1 and 0.3 percent of life-years saved, respectively).

“These estimates are substantially more conservative than previous calculations that focused mainly on the first year of vaccination, but clearly demonstrate an important overall benefit from COVID-19 vaccination over the period 2020-2024,” coauthor Stefania Boccia, D.Sc., Ph.D., from the Università Cattolica del Sacro Cuore in Rome, said in a statement.



Most life-years saved were among people aged 60 years or older; estimates are more conservative than those that focused on the first year of vaccination.

Fuente: Hematology Advisor. Disponible en <https://n9.cl/ftbr7v>

Vaccine line project delays impact SKAN's bottom line

Aug 21. SKAN Group has come out with its financial results for the first half of 2025.

The Switzerland-based company is a provider of isolators, cleanroom devices, and decontamination processes, primarily focused on the aseptic production of biopharmaceutical substances.

The company stated that H1 was marked by “two contrasting developments”

The two developments concern a very strong order intake, contrasted by weak net sales.

The strong order intake was attributed to filling lines for ADCs (antibody drug conjugates), which SKAN said



were in particularly high demand.

These modern oncology drugs require complex, large-scale aseptic-toxic systems with the highest quality and safety requirements.

SKAN's success rate for tenders for these products was over 40%.

This success delivered order intake at 20.2% above 2024, with the order backlog reaching a record level of CHF 386.4m.

So orders were coming in, but why were sales so low in comparison to 2024.

In terms of incoming money, SKAN attributed the weak performance to project delays. This means that final payments on these can not be taken, so do not contribute to H1. These payments can often be about 20% of total project revenue.

Specifically, these project delays centre around vaccine lines. "This development is primarily attributable to the global shift in vaccine-related demand and strategic priorities by some pharmaceutical companies," SKAN stated. "Several lines ordered during the COVID phase are only slowly being put into operation."

SKAN does believe that it will be able to compensate for these temporary shortfalls in the second half of the year due to its full order books and pipeline.

Other influencing factors in the company's operations this year, were acquisitions and tariffs.

July saw two important majority share acquisitions for SKAN; Metronik and ABC Transfer. These acquisitions focus on digitalisation and aseptic transfer, respectively. Both of these will add significant value to the company's offering in the pharmaceutical industry.

Current developments in US customs policy also represent a challenge to businesses with operations in the US. "[The tariffs] confirm the strategic relevance of the plan to expand the SKAN site in the US and establish local production."

SKAN added that suitable options are currently being evaluated for this US expansion.

Fuente: CLEANROOM TECHNOLOGY. Disponible en <https://n9.cl/8fef9>

Japan's Takeda weighs India for global trials to speed up drug launches

Aug 22. Japanese drugmaker Takeda Pharmaceutical is exploring conducting global clinical trials in India to speed up the launch of its innovative therapies in the country, its India head told Reuters. The move comes as India's clinical trials market expands, driven by diverse patient pools, cost efficiency, and a rapidly growing hospital network. Grand View Research projects the market to exceed Rs. 17,456 crore (US\$ 2 billion) by 2030. General manager of Takeda India, Ms. Annapurna Das, said the company is evaluating opportunities to leverage India's clinical trial ecosystem and is open to partnerships with local academia, healthcare providers, and technology firms to advance innovation. Takeda's goal is to integrate India's research and development ecosystem into its global pipeline while expanding patient access to therapies in oncology, neuroscience, gastrointestinal health, and inflammation.



Takeda plans to launch several key therapies in India over the next two to three years, including a lung cancer

drug this year and a dengue vaccine in partnership with local manufacturer Biological E., awaiting approval from India's drug regulator. The company recently established an innovation centre in Bengaluru, expanding from just over 500 to 750 employees, specialising in artificial intelligence (AI), data science, engineering, and design to support its global digital transformation. This strategic focus on India underscores Takeda's long-term commitment to the country as a growth market. It reflects the increasing role of India in global pharmaceutical research and innovation.

Fuente: India Brand Equity Foundation IBEF. Disponible en <https://n9.cl/zx0la>

Sanofi's New Vaccine Trial: A Potential Game-Changer for Infant Health

Aug 22. Sanofi is currently conducting a Phase 3 clinical trial to evaluate the safety and effectiveness of a new 21-valent pneumococcal conjugate vaccine (PCV21) in healthy infants and toddlers. The study, officially titled A Phase 3, Randomized, Modified Double-blind, Active-controlled, Parallel-group, 2-arm



Study to Investigate the Safety and Immunogenicity of a 21-valent Pneumococcal Conjugate Vaccine in Healthy Infants and Toddlers, aims to determine if PCV21 can safely produce antibodies when administered alongside routine pediatric vaccines.

The trial involves two groups: one receiving the experimental PCV21 vaccine and the other receiving the active comparator, the 15-valent pneumococcal vaccine (Vaxneuvance). Both vaccines are administered via intramuscular injection at specified intervals, with additional routine vaccines given concurrently.

This interventional study follows a randomized, parallel assignment model with quadruple masking to ensure unbiased results. The primary goal is prevention, focusing on the vaccine's ability to generate an immune response.

Key dates for the study include its start on May 22, 2025, with the most recent update submitted on August 19, 2025. These dates are crucial as they mark the progression and current status of the trial.

The outcome of this study could significantly impact Sanofi's market position, potentially boosting investor confidence and stock performance if results are favorable. It also positions Sanofi competitively within the vaccine industry, particularly against other pneumococcal vaccine producers.

The study is ongoing, with further details available on the ClinicalTrials portal.

Fuente: TIP RANKS. Disponible en <https://n9.cl/y8lau>

Equipo científico chileno está entre los más productivos en avances contra el virus sincitial

22 ago. El virus respiratorio sincitial (VRS) es la principal causa de infecciones agudas respiratorias en niños menores de 2 años en el mundo. Una reciente publicación científica de *Frontiers* hizo una revisión sistemática y un análisis bibliométrico de los últimos cinco años, para determinar las características, puntos críticos y fronteras de la producción científica mundial.

El profesor Alexis Kalergis, director del Instituto Milenio en Inmunología e Inmunoterapia (IMI), -equipo que actualmente está desarrollando la primera vacuna chilena contra el VRS- aparece en los top 10 de autores más productivos en este campo de investigación del mundo. El grupo de científicos que está desarrollando la primera vacuna chilena contra el virus respiratorio sincitial, que lidera el académico, contribuyó un número

importante de publicaciones de alto impacto en los últimos 5 años, que han fortalecido conocimiento fundamental sobre el VRS.

Este reporte denominado “Puntos críticos de investigación y tendencias mundiales del virus respiratorio sincitial en los últimos años”, de los autores Xiaoli Tao, Zhuping Ma, Hongxia Yuan, Wei Zhao, Jingyu Liu, Jing Tian, del Centro de Innovación Colaborativa para la Prevención y el Control de Zoonosis, y de la División de Enfermedades Infecciosas del Primer Hospital Afiliado de la Universidad Médica de Jinzhou, China, destacaron los avances donde se recuperó la versión del Science Citation Index Expanded (SCI-Expanded) de la Web of Science Core Collection (WoSCC) para publicaciones e información de registros publicados entre 2020 y 2024.

“Recibimos con mucho agradecimiento y humildad este reconocimiento, el cual es compartido junto a un extenso grupo de colaboradora(e)s y científica(o)s jóvenes que, como equipo, aspiramos a que desde la ciencia que desarrollamos en la universidad podamos impactar positivamente a la sociedad”, manifestó el académico UC. En el reporte se utilizó un paquete de software bibliométrico para analizar los indicadores bibliométricos, y las tendencias de investigación y los puntos críticos del VRS se visualizaron mediante VOSviewer y Citespace. Se evaluó la influencia de los artículos con el Global Citation Score (GCS). Se buscaron un total de 7.238 artículos y comentarios.

“Este equipo ha demostrado que, a través de la investigación científica básica, fundamental, aplicada y clínica, y su difusión, podemos aportar al desarrollo de políticas públicas en la búsqueda de una mejor calidad de vida para nuestra población, a nivel nacional e internacional. La importancia que reviste el reconocimiento que se nos hace como grupo es destacar el aporte científico que realizamos desde nuestro país hacia el mundo y que se basa en la calidad de las publicaciones y el impacto de nuestra investigación científica en varias disciplinas, lo que además reconoce el gran potencial científico que tiene nuestro país”, agregó el experto.



Estados Unidos es el país más productivo en el campo de la investigación sobre el VRS y también el país con la cooperación más estrecha con otros países e instituciones. La revista más influyente en este campo es “VIRUSES BASEL”, con 246 publicaciones.

El análisis de cocitación de referencias mostró que los temas relacionados con el VRS con mayor enfoque son: "pandemia de COVID-19"; "prefusión del virus respiratorio sincitial"; "academia estadounidense" y "vacuna proteica". De 2020 a 2024, los análisis de grupos y ráfagas de palabras clave mostraron que "Virus Respiratorio Sincitial", "Infección" y "Niños"; "Coinfección viral", "antivirus" y "vacunas" son actualmente focos de investigación. El área de investigación en este campo se distribuye principalmente entre "Inmunología", "Pediatria", "Farmacia", "Farmacéutica" y "Bioquímica Biología Molecular".

Fuente: Pontificia Universidad Católica de Chile. Disponible en <https://n9.cl/8s685y>

Evolving Perspectives on Pneumococcal Vaccination Across Adult

Aug 25. Community-acquired pneumonia is one of the leading infectious causes of hospital admission; further, it is associated with high rates of morbidity and mortality. *Streptococcus pneumoniae* remains the most common bacterial cause of pneumonia, accounting for approximately 227,000 hospitalizations annually and an estimated 25% mortality rate within 1 year of hospitalization. Additional complications of pneumococcal disease can range from non-invasive syndromes (acute otitis media, sinusitis) to invasive pneumococcal disease (IPD), including meningitis and bacteremia.



Prevention of pneumococcal disease through routine vaccination has led to substantial declines in IPD, particularly in serotypes covered in the pneumococcal conjugate vaccines (PCV). Since 2010, the Centers for Disease Control and Prevention (CDC) Advisory Committee on Immunization Practices (ACIP) has recommended the routine use of the 13-valent pneumococcal vaccine (PCV-13) for all children aged 14 to 59 months. This recommendation has led to direct declines in PCV-13 serotype pneumococcal disease in children and an indirect decline in adults, highlighting the importance of vaccination efforts to protect at-risk patients. Adults at highest risk of IPD include those older than 50 years, those aged 19 to 49 years who smoke or have certain comorbidities (eg, heart disease, diabetes, chronic obstructive pulmonary disease), and immuno-compromised individuals. Additionally, racial disparities in pneumococcal disease continue to disproportionately affect certain groups, with IPD rates higher in non-Hispanic Black and non-Hispanic American Indian or Alaska Native adults compared to the general population.

To mitigate IPD caused by non-PCV-13 serotypes, several new pneumococcal vaccines have been approved for the prevention of IPD, including a 15-valent (PCV-15), 20-valent (PCV-20), and 21-valent (PCV-21) pneumococcal vaccine. Based on the CDC's Active Bacterial Core Surveillance, approximately 54% to 62% and 77% to 85% of IPD cases between 2018-2022 were due to pneumococcal vaccine serotypes contained in PCV-20 and PCV-21, respectively. Unfortunately, overall risk-based pneumococcal vaccination coverage among adults remains low, with only 22% of adults aged 19 to 64 years who have risk factors receiving at least 1 dose of pneumococcal vaccine. Rates of vaccination additionally differ by race and health care accessibility, with vaccination rates generally lower among Black or Hispanic adults and those without a usual place to receive health care.

Expanding Protection: ACIP's Latest Adult Pneumococcal Vaccine Recommendation

As of October 23, 2024, the ACIP expanded the recommended use of a PCV from the previous recommendation of adults 65 years or older to include all PCV-naïve adults 50 years and older. The recommendation for PCV in adults aged 19 to 49 years with risk factors remains unchanged.^{12,15} Available vaccine options include PCV-15 (Vaxneuvance; Merck), PCV-20 (Prenar 20; Pfizer), and PCV-21 (Capvaxive; Merck). The 3 PCVs offer protection against different pneumococcal serotypes. If PCV-15 is selected, administration of pneumococcal polysaccharide vaccine (PPSV-23) is recommended at least 1 year later (or at least 8 weeks later if the patient is immunocompromised) to confer protection against 8 additional serotypes not covered in PCV-15.^{15,17} If PCV-20 or PCV-21 is used, a follow-up PPSV-23 dose is not needed, as these 2 vaccines confer broader protection of prevalent serotypes compared to PCV-15.

Of note, PCV-20 confers protection against the same 15 serotypes as PCV-15 plus 5 additional serotypes. Although PCV-21 offers more broad protection against 11 serotypes not prevalent in PCV-15 or PCV-20, it does not include serotype 4; this is an important consideration in regions such as the western United States, where this serotype remains highly prevalent among adults younger than 65 years who smoke, are homeless, use injectable drugs, or have underlying conditions (eg, alcoholism, chronic lung disease). Vaccine selection should take into account local serotype distribution and individual patient risk factors. Individuals who previously started the pneumococcal vaccine series with PCV-13 or PPSV-23 should receive an updated PCV vaccine; timing and product selection will vary depending on the initial vaccine received.

The expansion of the recommendation for PCV administration to adults at least 50 years of age aims to reduce disease burden in patients aged 50 to 64 years, increase access to preventative care in patients at high risk for IPD, and reduce racial disparities seen in IPD. The shift in recommendation for the administration of expanded serotype vaccines (eg, PCV-20, PCV-21, or PCV-15 plus PPSV-23 instead of PCV-13) is designed to combat the substantial burden of non-PCV-13 serotypes that account for 42% and 75% of IPD in children and adults, respectively.¹⁸ The expanded age recommendation attempts to ensure that patients at moderate to high risk for IPD are readily identified and able to access vaccines. Previous data suggested that over 80% of patients aged 50 to 64 years who were hospitalized with pneumococcal pneumonia had at least 1 risk factor (eg, heart disease, lung disease, liver disease, diabetes) that put them at higher risk of IPD. According to 2022 Behavioral Risk Factor Surveillance System Data, among the general population, an estimated 32% to 54% of adults aged 50 to 64 years qualify for a risk-based pneumococcal vaccine, although only an estimated 37% of eligible adults aged 50 to 64 years were vaccinated. Rates of vaccination were higher in patients older than 65 years who were identified based on an age cutoff instead of a risk-based system, suggesting that a streamlined age-based approach may be easier to follow.

In addition to capturing patients at high risk based on comorbidities, previous data suggested that peak rates of IPD occur at a younger age (55-59 years) in Black males, highlighting a population that previously would not have been identified as candidates for pneumococcal vaccination using comorbidity-based risk assessment. By simplifying the PCV recommendation to include adults older than 50 years, patients who would benefit most from pneumococcal vaccination may be more readily identified, which may help reduce pneumococcal-associated morbidity and mortality and improve public health.

Clinical Considerations for Pharmacists

Pharmacists can play a key role in preventing pneumococcal infection by administering pneumococcal vaccines in the community and health care settings. Identifying patients eligible for pneumococcal vaccination should involve stepwise decision-making. Key factors to consider include age, underlying medical conditions, and vaccine history.

For adults at least 50 years of age or aged 19 to 49 years with risk factors (alcoholism, chronic heart disease, chronic liver disease, chronic lung disease, cigarette smoking, or diabetes) who have not previously received a pneumococcal vaccine, PCV-15, PCV-20, or PCV-21 should be administered. If PCV-15 is administered, an additional dose of PPSV-23 is required in at least 1 year, making this approach less preferable for patients with limited health care access. For special populations (eg, patients with an immunocompromising condition, cochlear implants, or cerebrospinal fluid leaks) who have not previously received a pneumococcal vaccine, PCV-15, PCV-20, or PCV-21 may be administered, although PCV-15 should be followed by PPSV-23 in 8 weeks or more, compared to at least 1 year in non-immunocompromised patients.

Additional considerations for selection between pneumococcal vaccines, particularly PCV-20 and PCV-21, include the patient's geographic location, as serotype protection differs between vaccines. For areas where more than 30% of IPD is caused by serotype 4, PCV-20 or PCV-15 followed by PPSV-23 at least 1 year later is preferred, as PCV-21 lacks serotype 4 protection. For all other areas, both vaccines are viable options, although PCV-21 offers broader coverage of more serotypes compared to PCV-20. Streamlined vaccine recommendations may make vaccinations more readily offered and available during routine pharmacist interactions.

For patients who previously started the pneumococcal vaccine series, the timing and selection of vaccine will differ depending on the prior vaccine schedule. For adults who previously were given PPSV-23 only, a single dose of PCV-15, PCV-20, or PCV-21 should be administered at least 1 year after the last PPSV-23 dose. For adults who previously were given PCV-13 only, a single dose of PCV-20 or PCV-21 is recommended at least 1 year after PCV-13, and PPSV-23 is no longer needed. For adults who were given PCV-13 and PPSV-23 (1 or 2 doses), a single dose of PCV-20 or PCV-21 may be recommended 5 years or longer after the last pneumococcal vaccine depending on age, risk factors, and shared clinical decision-making.

Fuente: Drug Topics. Disponible en <https://n9.cl/e099zk>

Vaccine Integrity Project Fills Information Gap

Aug 26. This fall, guidance for protection against the flu, COVID, and other respiratory infections remains consistent with previous years, with no new safety signals, based on evidence reviewed by the Vaccine Integrity Project and presented in an online webinar.

The Vaccine Integrity Project (VIP) was created by the University of Minnesota's Center for Infectious Disease Research and

Policy (CIDRAP) to provide evidence on vaccine safety and effectiveness as a resource for clinicians and for medical societies.

Overall, the experts found that vaccination against COVID, respiratory syncytial virus (RSV), and influenza yielded similar or better outcomes compared to no vaccination. Additionally, the VIP experts found no sufficient evidence of new safety signals to justify the CDC's rescinding of the previous recommendation to vaccinate pregnant women and healthy children against COVID this season, said CIDRAP director Michael Osterholm, PhD, MPH, in a press release.

The VIP is a response to the gaps left by federal changes that have altered the dissemination of evidence-based scientific information, according to CIDRAP. In advance of the upcoming respiratory virus season, the VIP presented its first live-streamed webinar on August 19 with data on influenza, COVID, and RSV vaccines, focusing on indications for pregnant women, children, and immunocompromised individuals.

For the presentation, a team of 24 infectious disease experts and epidemiologists reviewed data from 1406 full-text articles published within the last few years and identified 590 for detailed examination. Jake Scott, MD, of Stanford University, established the VIP's credentials by reviewing their strict research criteria and outlined the specific safety events analyzed for three vaccines of interest: COVID vaccines, RSV vaccines and antibodies, and influenza vaccines.



Pregnancy Protection

Some infections, such as influenza, are more severe during pregnancy and in fact may increase the risk of adverse outcomes, including preterm birth and stillbirth, said Caitlin Dugdale, MD, an infectious diseases physician at Massachusetts General Hospital and assistant professor of medicine at Harvard Medical School, Boston. Dugdale presented the VIP's evidence base for vaccines in pregnancy.

The researchers focused on safety and effectiveness of COVID, RSV, and influenza, and found no studies with new epidemiologic trends related to RSV or influenza; the one study involving epidemiology of COVID showed a 7% incidence of long COVID among vaccinated pregnant patients compared to a 12% incidence among unvaccinated pregnant patients. As for COVID vaccine safety, no concerns were identified for outcomes of miscarriage, stillbirth, or congenital anomalies with either the Pfizer or Moderna vaccines, she said.

Considerations for Children

Harleen Marwah, MD, a pediatrician at Mass General Brigham for Children, presented the evidence review for the flu, COVID, and RSV vaccines in children.

Children are especially vulnerable to respiratory infections because they are in critical stages of development, Marwah said. The VIP experts reviewed data on myocarditis and Guillain-Barré syndrome associated with COVID vaccination. The studies of myocarditis were consistent with previous research, and new data showed that risk decreased with subsequent booster doses, she said. Similarly, pediatric studies of co-administered COVID mRNA vaccines and flu vaccines showed no increased risk of serious adverse events.

Immunocompromised Individuals

Michael Abers, MD, a physician scientist at Albert Einstein College of Medicine, New York City, presented the evidence base for vaccination of immunocompromised individuals against COVID, RSV, and influenza.

Vaccines work by training the immune system to more rapidly recognize pathogens, but someone with a compromised immune system may be less responsive to vaccination, Abers said in his presentation.

However, the most recent data showed that vaccines for COVID were similarly safe and effective among immunocompromised individuals and the general population. The RSV vaccine was especially effective among patients with solid organ transplants, he noted. The flu vaccine, while slightly less effective among immunocompromised individuals compared to immunocompetent individuals, was still effective, he said.

Support for Society Guidelines

The goal of the VIP webinar was to assemble information that could be shared with professional societies to make recommendations, Abers said in a discussion period.

Based on the data reviewed by the VIP, the American Academy of Pediatrics (AAP) released its own recommended childhood vaccination schedule, independent of the CDC. Other societies, including the Infectious Diseases Society of America, the American College of Obstetricians and Gynecologists, and the American Academy of Family Physicians, are expected to release independent vaccine guidelines based on the VIP evidence review.

Evidence Clinicians Need

"The work of the Vaccine Integrity Project is urgently needed because of the myriad questions [about vaccines] that have now been raised," David J. Cennimo, MD, associate professor of medicine & pediatrics at Rutgers New Jersey Medical School in Newark, told Medscape Medical News.

“Unfortunately, the past work of ACIP and CDC which has been grounded in vaccine science and public health has been destabilized, often with unfounded speculations and misinterpretation of data,” said Cennimo, who is not involved in the VIP.

“I commend CIDRAP and Dr Osterholm for their steadfast leadership and commitment to protecting the nation’s health,” Cennimo added. “They have worked to recreate the unbiased, evidence-based research presentations that characterized the ACIP’s gold standard of open debate.” The medical community will need these data not only to make informed decisions, but also to support educated and informed discussions for shared decision-making with parents and families, he said.

“For instance, the safety and efficacy of COVID vaccines in children are actually supported by a plethora of data, contrary to recent reports,” he said.

“As clinicians, we must become familiar with these debates as well as the incorrect messages permeating our media,” Cennimo told Medscape Medical News. He expressed confidence that the AAP and other medical organizations will strengthen the vaccine message, but also expressed concerns about vaccine funding, which is tied to CDC recommendations.

Fuente: Medscape. Disponible en <https://n9.cl/z6gh5>

ImmunoPrecise Advances AI Discovered and Developed Universal Dengue Vaccine Initiative: Moves into Preclinical Antibody Generation

Aug 27. ImmunoPrecise Antibodies Ltd. (NASDAQ: IPA) (“IPA” or the “Company”), a biotherapeutics company delivering advanced solutions in biologics and drug discovery today announced the next milestone in its universal dengue vaccine program. Building on its June announcements confirming the discovery and validation of a uniquely conserved dengue epitope using its LENSai™ platform powered by patented HYFT® technology, the Company has updated that they are now advancing to pre-clinical manufacturing for *in vivo* (in animal) testing and virus neutralization analysis.



The manufactured product will be used in proprietary immunization protocols at IPA’s Canadian facility. These studies, conducted in a rabbit model, are designed to evaluate whether the vaccine candidate elicits robust monoclonal antibody (mAb) responses—a crucial step toward translational development.

“This is the natural next step in our journey from AI discovery to real-world validation,” said Dr. Jennifer Bath, CEO of ImmunoPrecise. “We’ve shown that HYFT technology can identify vaccine targets that others miss — unique targets that are conserved, specific, and immunologically relevant. Now we’re now moving into manufacturing and live immunization studies, where the question becomes: can this target elicit a powerful and protective antibody response? This is where discovery turns into translational reality, with dengue as the first example of what our platform can achieve across infectious diseases.”

The science behind the step:

- ♦ AI Discovery → Manufacturing: HYFT-guided analysis revealed a discontinuous dengue epitope conserved across all four virus serotypes.

- ♦ **Validated Safety & Immunogenicity:** In silico screens confirmed immune activation potential (B- and T-cell responses), structural stability, and comprehensive safety against host proteomes.
- ♦ **Now Entering Immunization Studies:** The product is being synthesized and will be used in rabbit models to assess whether it elicits specific monoclonal antibodies — an essential proof point for vaccine development.

A platform approach, not just a vaccine:

This progression highlights IPA's end-to-end advantage: combining AI-native discovery with in-house wet-lab validation. With regulators like the U.S. FDA encouraging predictive, human-relevant approaches that reduce reliance on animal testing, ImmunoPrecise's in silico-to-in vivo workflow represents a forward-looking model for next-generation vaccine R&D.

By applying the same methodology across pathogens, IPA is positioning its LENSai/HYFT platform as a versatile discovery engine — not only for dengue, but also for other infectious diseases and immuno-oncology targets.

About ImmunoPrecise Antibodies Ltd.

ImmunoPrecise Antibodies Ltd. (NASDAQ: IPA) is a biotherapeutics company focused on the discovery and development of next-generation biologics. The Company combines scientific expertise with proprietary technologies—such as its LENSai™ platform—to accelerate drug discovery and improve decision-making across complex biological systems. IPA supports global partners in advancing novel therapeutics, diagnostics, and translational research. For more information, visit www.ipatherapeutics.com

Fuente: MORNING STAR. Disponible en <https://n9.cl/wjqggt>

Boosting mRNA Vaccines with Cutting-Edge Technology

Aug 27. Messenger RNA (mRNA) vaccines revolutionized public health during the COVID-19 pandemic, representing a groundbreaking shift away from traditional vaccine design. Unlike conventional vaccines that introduce weakened or inactivated viruses to train the immune system, mRNA vaccines operate on a genetic blueprint level. They deliver encoded instructions directly into human cells, enabling these cells to produce viral proteins that then trigger an immune response. This innovative method allowed Pfizer-BioNTech and Moderna to rapidly develop highly effective COVID-19 vaccines, setting the stage for a new era in vaccinology.

Building on this success, researchers at Yale University have now unveiled a novel technological advance that enhances the immunogenic power and effectiveness of mRNA vaccine platforms.

This breakthrough, recently detailed in the prestigious journal *Nature Biomedical Engineering*, promises to significantly broaden the potential applications of mRNA vaccines beyond infectious diseases like COVID-19, extending to challenging conditions such as cancer and autoimmune disorders. The team, led by Sidi Chen, associate professor of genetics and neurosurgery, sought to understand why mRNA vaccines, despite their triumphant pandemic debut, often underperformed in clinical trials for other diseases.

A critical bottleneck identified by the researchers lies in the behavior of antigens — the molecular flags



presented by infected or abnormal cells that alert the immune system. For an antigen to be effectively recognized and to induce a robust immune response, it must be displayed on the surface of cells. However, Chen and his team discovered that many antigens generated through mRNA vaccines remain trapped inside the cell's interior, inaccessible to immune surveillance. This intracellular sequestration critically limits the vaccine's capacity to provoke a protective immune response, hindering its efficacy against a range of diseases.

To overcome this challenge, Yale scientists engineered a sophisticated molecular vaccine platform (MVP) that effectively upgrades the delivery and presentation of vaccine-derived antigens. Their approach involves fusing what they refer to as a "cell-GPS" module to the proteins produced by mRNA instructions. This GPS-like component comprises natural membrane-associated elements such as signal peptides and transmembrane anchors, which are essential in normal biological processes for directing proteins to their correct cellular locations — namely, the cell membrane.

Signal peptides are short amino acid sequences that act like postal codes, guiding the nascent proteins through cellular trafficking pathways to ensure they reach the surface. Transmembrane anchors secure these proteins to the extracellular membrane, stabilizing their position where immune cells can detect them. Incorporation of these elements into the vaccine design guarantees that the antigens will be displayed robustly on the cell surface, vastly improving immune visibility and subsequent activation of both antibody- and T cell-mediated responses.

In rigorous laboratory experiments, this MVP framework was tested across multiple disease models including mpox virus (formerly monkeypox), human papillomavirus (HPV), and the varicella-zoster virus responsible for shingles. Remarkably, the enhanced antigen expression translated into significantly amplified immune responses: elevated levels of neutralizing antibodies, greater activation of cytotoxic T lymphocytes, and improved overall immunogenicity. These results underscore the platform's versatility and its potential to redefine mRNA vaccine effectiveness against a spectrum of viral infections and potentially malignant conditions.

The implications of this research extend far beyond virology. By ensuring precise antigen localization, the MVP technology addresses one of the principal limitations that have constrained the broader adoption of mRNA vaccines in oncology and immunology. Diseases such as cancer and autoimmune disorders, which require a finely tuned immune activation profile, may profoundly benefit from this targeted approach. Chen emphasizes that this innovation represents a foundational step toward expanding the versatility of mRNA-based immunotherapies.

Moreover, the platform's modular nature allows for rapid adaptation to different antigens and disease targets, a crucial advantage in the battle against emerging pathogens and evolving health threats. This flexibility is particularly vital given the increasing incidences of viral mutations and the complexity of tumor-associated antigens. By integrating natural cellular machinery into vaccine design, the researchers have crafted a robust system that harmonizes synthetic biology with immunological precision.

The study also benefits from a collaborative environment at Yale, involving interdisciplinary expertise from immunobiology, molecular biophysics, and therapeutic radiology. Co-senior authors Carolina Lucas and Daniel DiMaio contribute insights from their respective fields, enriching the study's multi-faceted approach to solving complex biological challenges. Their combined efforts highlight the importance of integrating diverse scientific perspectives to overcome hurdles in next-generation vaccine development.

As the scientific community intensifies the push for more effective immunization strategies, the MVP platform constitutes a critical advance in mRNA vaccine science. It not only revitalizes interest in mRNA technology for diseases that have eluded effective vaccination but also inspires confidence in the adaptability of this platform for future biomedical applications. The fusion of natural protein trafficking elements with lipid nanoparticle mRNA delivery opens new avenues for precision immunotherapy, with the promise to uplift global health outcomes.

Looking ahead, clinical translation of this innovative vaccine platform will require extensive validation in human trials to confirm safety, immunogenicity, and efficacy. Nonetheless, the promising preclinical results afford optimism. With improved antigen presentation capability, vaccines developed through MVP technology could lead to a new generation of preventive and therapeutic measures against a broad array of infectious agents and immune-related diseases.

This breakthrough exemplifies how fundamental insights into cellular biology can be harnessed to refine and expand the capabilities of revolutionary technologies like mRNA vaccinology. The research team plans to further explore applications of their platform, aiming to tackle formidable health challenges including HIV and autoimmune conditions. Their work not only furthers scientific understanding but also paves the way for tangible innovations in medicine poised to transform disease prevention and treatment worldwide.

Fuente: Bioengineer.org. Disponible en <https://n9.cl/wzkjt>

La OPS llama a reforzar vacunación y vigilancia ante la propagación de resistencia de la tosferina a los antibióticos en las Américas

27 ago. Frente al resurgimiento de la tosferina en varios países de la región y la aparición y propagación de cepas resistentes a los antibióticos, la Organización Panamericana de la Salud (OPS) reiteró la importancia de fortalecer la vacunación y los sistemas de vigilancia. La preocupación fue compartida durante una reunión reciente con la Red Latinoamericana y del Caribe de Vigilancia de la Resistencia a los Antimicrobianos (ReLAVRA+), en la que se discutieron los hallazgos más recientes y las estrategias de respuesta frente a esta enfermedad altamente contagiosa.



“La vacunación, la vigilancia y el uso responsable de los antibióticos son fundamentales para evitar que la tosferina vuelva a convertirse en una amenaza grave de salud pública”, afirmó Pilar Ramón-Pardo, jefa del Programa Especial de Resistencia Antimicrobiana de la OPS. “Aún estamos a tiempo de contener este problema, pero necesitamos actuar: aumentar la cobertura vacunal, fortalecer la detección temprana y mejorar nuestra capacidad de respuesta ante brotes”, añadió.

Uno de los principales desafíos es que el tratamiento habitual se basa en antibióticos del grupo de los macrólidos, como la azitromicina, claritromicina y eritromicina. Sin embargo, mutaciones genéticas en la bacteria, particularmente en el gen 23S rRNA, están reduciendo la eficacia de estos medicamentos, dificultando el tratamiento de pacientes y la prevención en contactos cercanos.

Durante la pandemia de COVID-19, el uso extensivo e inadecuado de antibióticos como la azitromicina pudo haber contribuido a la aparición de cepas resistentes de *Bordetella pertussis*. Desde 2024, se han reportado casos en Brasil, Estados Unidos, México y Perú. Su detección fue posible gracias al fortalecimiento de los sistemas de vigilancia y diagnóstico, que incluyen el uso de pruebas moleculares y métodos estandarizados en laboratorios de referencia. Este avance, apoyado por alianzas internacionales, ha sido clave para identificar y contener estos casos.

Dado el uso generalizado de estos antibióticos y la alta movilidad internacional, existe el riesgo de una propagación más amplia si no se refuerza la respuesta sanitaria.

La resistencia a los antimicrobianos (RAM) ocurre cuando las bacterias desarrollan mecanismos para evadir el efecto de los medicamentos, volviéndolos ineficaces. En el caso de la tosferina, esto puede limitar las opciones de tratamiento, dificultar el control de brotes, y aumentar el riesgo de complicaciones graves, especialmente en contextos con baja cobertura vacunal.

Coberturas de vacunación en descenso

La tosferina, también conocida como coqueluche o tos convulsiva, puede prevenirse con tres dosis de la vacuna DPT en niños menores de un año, y refuerzos durante la infancia y la adolescencia. Aun así, en personas infectadas —especialmente en lactantes— puede causar cuadros graves, con riesgo de complicaciones o incluso la muerte.

La enfermedad ha resurgido con fuerza en las Américas. Mientras en 2023 se notificaron 4.139 casos, en 2024 la cifra ascendió a 43.751. En los primeros siete meses de 2025, nueve países han reportado más de 18.595 casos y 128 defunciones. Este resurgimiento está relacionado con la disminución de las tasas de vacunación y resalta la necesidad de reforzar la vigilancia activa y estandarizada.

Durante la pandemia, las coberturas regionales de la primera dosis (DTP1) y la tercera (DTP3) cayeron a mínimos históricos: 87% y 81% en 2021, respectivamente. En 2023 se observó una recuperación parcial (90% y 88%), aunque aún por debajo del 95% recomendado por la OPS y con disparidades significativas dentro de los países. Además de la vacunación infantil, la OPS también recomienda vacunar a las embarazadas —especialmente en contextos de brote— y al personal de salud que está en contacto con recién nacidos.

Laboratorios fortalecen la vigilancia

Fortalecer los laboratorios nacionales de referencia con métodos de diagnóstico confiables y estandarizados es esencial para detectar la resistencia de manera oportuna y guiar la respuesta de las autoridades de salud. Iniciativas colaborativas como la red ReLAVRA+ y la Red de Vigilancia Genómica Regional (PAHOgen) de la OPS, junto con otras alianzas estratégicas, han permitido estandarizar metodologías y ampliar la capacidad de vigilancia en varios países de la región.

Como parte de su cooperación técnica, la OPS organizó el 19 de agosto un seminario virtual de la red ReLAVRA+, que reunió a laboratorios nacionales de referencia y especialistas en microbiología de la región. Durante el encuentro, expertos de Colombia, México, Brasil y Perú compartieron hallazgos recientes, metodologías para detectar cepas resistentes y estrategias de vigilancia y respuesta.

Ante este panorama, la OPS insta a los países a fortalecer el diagnóstico, reforzar la formación técnica del personal de salud, mantener o incrementar las coberturas de vacunación, y establecer sistemas de vigilancia activos y estandarizados que permitan una respuesta ágil y eficaz frente a la resistencia a antimicrobianos.

Fuente: Organización Panamericana de la Salud OPS. Disponible en <https://n9.cl/k5em4>

Raina Biosciences unveils breakthrough generative AI platform for mRNA therapeutics featured in Science

Aug 28. Raina Biosciences Inc., ("Raina"), an mRNA technology and therapeutics company, today announced the publication of data from its generative AI platform in *Science*. The data supports Raina's pioneering approach to mRNA design using its GEMORNA platform to generate novel sequences with superior drug properties over existing mRNA discovery methods. Founded by a team with deep RNA therapeutics and AI expertise, the Company's mission is to transform the mRNA-based therapeutics landscape by accelerating drug discovery timelines and opening new therapeutic areas for mRNA with best-in-class AI-designed therapeutics.

The *Science* article, titled, "Deep generative models design mRNA sequences with enhanced translational capacity and stability," marks a significant landmark for optimizing novel mRNAs with enhanced expression and durability to facilitate a wide range of therapeutic mRNA applications.

The article, which was published online today by the journal *Science*, is summarized as follows:

- ⇒ Raina's GEMORNA platform designs superior linear and circular mRNA drug molecules with optimized expression levels and durability, addressing two major issues that have limited the effectiveness of mRNA-based therapeutics
- ⇒ GEMORNA is highly differentiated from prior RNA language models developed for predictive tasks through its direct design of novel sequences from a near infinite design space
- ⇒ GEMORNA-generated mRNAs:
 - Elicited significantly higher immune response in mice compared to a leading commercially available mRNA vaccine sequence, potentially supporting mRNA therapeutic vaccine applications for diseases such as cancer
 - Achieved up to a 150-fold increase in human erythropoietin (hEPO) expression compared to an optimized benchmark, potentially supporting mRNA medicines for gene therapy
 - Demonstrated a 5-fold increase in CD19 CAR expression and a 2-fold improvement in durability compared to a patented benchmark, resulting in nearly 100% anti-tumor efficacy in primary human T cells, potentially supporting mRNA therapies for in vivo CAR-T

"Raina's GEMORNA platform is built upon a decade of the team's foundational work in synthetic biology and artificial intelligence," said Jicong Cao, Ph.D., Chief Executive Officer and co-founder of Raina Biosciences, and corresponding author of the *Science* paper. "We are excited to work with industry-leading pharma and biotech companies to expand the usage of mRNA-based therapeutics while we prepare to build an internal pipeline."

"Raina has the potential to transform mRNA therapeutics by rapidly and reliably generating novel sequences with greater performance, precision and efficacy," said Timothy Lu, M.D., Ph.D., Raina's Chairman of the Board and former MIT faculty member. "The GEMORNA platform could be a sea change for biopharma companies pursuing mRNA-based medicines beyond traditional infectious disease vaccines ranging from neoantigen cancer vaccines, in vivo CARs, to gene editing or gene therapy applications."

Based in Boston, Raina Biosciences is a spinout company from the Massachusetts Institute of Technology

(MIT), founded by leaders in mRNA and AI with a strong track record in the biotech industry. Jicong Cao, Ph.D. (co-founder, Chief Executive Officer) is a former mRNA researcher at MIT and co-founder of Bota Biosciences. He Zhang, Ph.D. (co-founder, Chief Technology Officer and first author of the Science paper) was previously a Senior Staff Scientist at Baidu Research. Timothy Lu, M.D., Ph.D., Raina's co-founder and Chairman of the Board, is a serial biotech entrepreneur having co-founded a number of biotechnology and biopharmaceutical companies, and served on the MIT faculty from 2010–2022. Joel Edwards, Chief Corporate Development Officer, brings over 25 years of leadership in corporate strategy and deal-making across biotech, including his tenure as Vice President of Corporate Strategy at Ionis Pharmaceuticals. Raina is also supported by its key scientific advisor Jeff Collier, Ph.D., Bloomberg Distinguished Professor and Inaugural Director of the RNA Innovation Center at Johns Hopkins University. Raina closed an angel round of \$5.7 million upon company formation.

About Raina Biosciences

Raina is pioneering the world's first and leading generative AI platform for mRNA-based therapeutics. Using its proprietary platform, GEMORNA, Raina navigates new frontiers in mRNA engineering to design sequences with optimal profiles for expression, durability, and stability. Raina is partnering with leading pharma and biotech companies to create next generation and best-in-class mRNA-based therapeutics.

Fuente: EurekAlert. Disponible en <https://n9.cl/9yooj>

Bi TECNOLÓGICA mixta Cuba-Vietnam elevará producción de fármacos

29 ago. La producción conjunta de medicamentos esenciales para Cuba y Vietnam, así como exportar a naciones de Asia y el mundo figura entre los objetivos esenciales de la empresa mixta Genfarma, destacó este viernes BioCubafarma en un comunicado.

Establecida en Vietnam, fue creada tras la firma del acuerdo constitutivo el pasado 19 de mayo de 2025 entre BCF S.A. (de BioCubaFarma) y la empresa vietnamita Genfarma Holdings, destacó la institución cubana en un comunicado.

En ocasión de la visita de la presidenta de BioCubaFarma Mayda Mauri a esa nación asiática la empresa resaltó que Genfarma representa un modelo de cooperación en el sector biofarmacéutico que presenta, además, entre sus objetivos, la transferencia tecnológica para llevar tecnología médica cubana de vanguardia a Vietnam, con una clara vocación internacional.

Este aspecto es fundamental para la sostenibilidad económica del proyecto y para cumplir con su objetivo de generar ingresos que serían utilizados en el desarrollo de la industria y la producción de medicamentos para la población cubana, subraya el comunicado.

La visita de la presidenta Mayda Mauri Pérez a la empresa mixta en Vietnam representa un paso significativo en la materialización de este proyecto estratégico para ambos países.



Esta iniciativa no sólo fortalece los lazos históricos entre Cuba y Vietnam, sino que establece un modelo innovador de cooperación sur-sur en el sector de la biotecnología farmacéutica, con potencial para impactar positivamente en la salud de ambos pueblos y de la región asiática.

La concreción exitosa de esta empresa mixta demostrará cómo naciones hermanadas pueden desarrollar proyectos mutuamente beneficiosos que aprovechen las capacidades científicas y tecnológicas para el bienestar de sus poblaciones, destacó el texto.

Mega-empresa de casi 12 años de fundación, BioCubaFarma se consolida como entidad de ciclo cerrado que investiga, desarrolla, produce, comercializa, o sea, desde la gestación del producto, hasta que llega al paciente.

En su gestión sustentable y sostenible garantiza todo lo necesario para el cuidado de la salud de los cubanos, exporta medicamentos, medios y sistemas diagnósticos, equipos médicos, tecnologías de avanzada en la producción de medicamentos y alimentos, así como servicios de alta tecnología.

Fuente: Cuba MinRex. Disponible en <https://n9.cl/frka6>

Ente africano lanzó proyecto sobre respuesta ante pandemias

29 ago. La Autoridad Intergubernamental para el Desarrollo (IGAD) lanzó en Uganda el Proyecto de Preparación y Respuesta ante Pandemias, una iniciativa que fortalecerá la vigilancia, alerta temprana y sistemas de laboratorios para mejorar la resiliencia regional.

En la estrategia de tres años participan junto al ente africano el Fondo Pandémico, la Federación Internacional de Sociedades de la Cruz Roja y de la Media Luna Roja y la Organización Mundial de la Salud colaboran en la propuesta.

La jefa de la Misión de la IGAD en Kampala, Joselyn Bigirwa, el proyecto es oportuno y esencial, dada la creciente frecuencia y complejidad de las emergencias de salud pública en el Cuerno de África.

Bigirwa precisó que está centrado en mejorar las funciones transfronterizas críticas, esenciales para salvaguardar la seguridad sanitaria regional.



Este hito subraya el compromiso de la IGAD de salvar vidas, proteger los medios de vida y promover un Cuerno de África más seguro y saludable mediante la acción colectiva, subrayó un comunicado de prensa.

La región del ente africano, integrada por Djibouti, Etiopía, Kenya, Somalia, Sudán del Sur, Sudán del Sur y Uganda, se enfrenta a riesgos persistentes de brotes de enfermedades infecciosas con importantes consecuencias para la salud pública, socioeconómica y mundial.

El texto reconoció que los sistemas sanitarios frágiles se ven continuamente tensados por las epidemias recurrentes, exacerbadas por los altos niveles de circulación transfronteriza de personas, ganado y bienes, lo que aumenta el riesgo de transmisión de enfermedades.

Al margen del lanzamiento, el secretario ejecutivo de la IGAD, Workneh Gebeyehu, entregará suministros y equipos críticos de preparación para pandemias al Ministerio de Salud de Uganda.

Fuente: Granma. Disponible en <https://n9.cl/q7xq65>

México y Brasil firman acuerdo para producir medicamentos y vacunas

29 ago. Las autoridades sanitarias de México y Brasil firmaron este jueves dos memorándums de entendimiento para impulsar la producción de vacunas y principios activos para desarrollar medicamentos, así como para reforzar la vigilancia epidemiológica y el combate de enfermedades como el dengue.

En un comunicado, la Secretaría de Salud destacó que el acuerdo de colaboración permitirá reforzar la producción nacional de medicamentos esenciales “para garantizar la soberanía sanitaria regional”.

La firma de los acuerdos de colaboración se selló entre autoridades sanitarias de ambos países y fue encabezada por el secretario de Salud de México, David Kershenobich, y el vicepresidente de Brasil, Geraldo Alckmin.

El primero de estos acuerdos fue entre la Comisión Federal para la Protección contra Riesgos Sanitarios (Cofepris) y la Agencia Nacional de Vigilancia Sanitaria de Brasil (ANVISA).

Con ello, se busca el reconocimiento recíproco de capacidades regulatorias, lo que permitirá agilizar la homologación de registros sanitarios, facilitar ensayos clínicos y coordinar la certificación de plantas farmacéuticas, incluso en terceros países.

“El reconocimiento mutuo entre ANVISA y Cofepris permitirá acelerar la llegada de nuevas moléculas y medicamentos a la población”, dijo el titular de Salud.

El segundo memorándum se celebró entre Laboratorios de Biológicos y Reactivos de México (Birmex) y la Fundación Fiocruz.

Lo anterior permitirá la cooperación en la producción de vacunas y principios activos para desarrollar “medicamentos estratégicos”, incluyendo factores de coagulación, albúminas e inmunoglobulinas, vitales para los sistemas de salud de México y Brasil.

La Secretaría de Salud destacó que bajo este acuerdo también se buscará combatir el dengue y otras arbovirosis, mediante el uso de la vacuna desarrollada por el Instituto Butantan, el intercambio de métodos de control biológico y la vigilancia epidemiológica.

Crearán Comité Bilateral de Salud

Para dar seguimiento puntual a estos compromisos y con el fin de que se traduzcan en resultados concretos, se creará un Comité Bilateral de Salud México–Brasil que se reunirán cada mes.



“Celebro este encuentro entre Brasil y México, no sólo de palabra, sino porque están aquí sus delegaciones, la nuestra, y hemos logrado implementar lo que discutimos, asegurando que trascienda en verdaderos proyectos de colaboración en nuestros países. Creo que en la región latinoamericana tenemos una gran responsabilidad”, manifestó David Kershenobich.

El secretario de Salud también resaltó los programas sociales como “Salud Casa por Casa”, que prioriza la atención a personas mayores y con discapacidad, como ejemplos de políticas que pueden inspirar una mayor cooperación sanitaria regional.

“México y Brasil comparten una relación bilateral larga y profunda que está a la altura de nuestras responsabilidades como las dos economías más grandes de América Latina, bajo el liderazgo de la presidenta Claudia Sheinbaum y del presidente Luiz Inácio Lula da Silva, con una visión humanista que prioriza la salud como derecho, el desarrollo científico y tecnológico, y el impulso a la producción local para robustecer la soberanía sanitaria en un escenario geopolítico”, aseguró David Kershenobich.

Por su parte, la titular de Cofepris, Armida Zúñiga, subrayó que estos acuerdos se alinean con el Plan México, impulsado por el gobierno de la presidenta Claudia Sheinbaum.

Fuente: EL ECONOMISTA. Disponible en <https://n9.cl/bi1619>

FDA approves updated Covid-19 vaccines for limited groups as US cases continue to rise

Aug 31. The U.S. Food and Drug Administration approved updated COVID-19 vaccines but for a more limited group: adults age 65 and older and younger people who are at higher risk from COVID-19.

"FDA has now issued marketing authorization for those at higher risk: Moderna (6+ months), Pfizer (5+), and Novavax (12+). These vaccines are available for all patients who choose them after consulting with their doctors," US Health and Human Services Secretary Robert F. Kennedy Jr. said in a post on X.

Kennedy said in this post that he had promised "to keep vaccines available to people who want them, especially the vulnerable," but the move may limit access to COVID-19 shots for people who were routinely able to get them in the past.

It may be significantly more difficult for infants and young children to be vaccinated, although they are especially vulnerable to COVID-19. Last respiratory virus season, there were 48 COVID-19 hospitalizations for every 100,000 children under 5 – nearly seven times the rate for older children and more than twice the rate for adults under 50.

Emergency use authorizations for COVID-19 vaccines are rescinded, Kennedy said on Wednesday, which means Pfizer's COVID-19 vaccine is no longer available for children younger than 5. Moderna's vaccine is approved for children as young as 6 months, but only if they have an underlying condition that puts them at higher risk. The FDA had already approved Novavax's COVID-19 vaccine, but only for people 65 and older and those 12 and up who have at least one underlying condition that puts them at higher risk of severe illness.

"Our health care system is now solidly anti-children and anti-science. The data are clear: young children—especially infants—remain highly vulnerable to severe illness and hospitalization from COVID-19. By restricting access to safe, evidence-based vaccines, federal leaders are choosing ideology over science," Fatima Khan, co-founder of the nonprofit grassroots group Protect Their Future, which advocates for vaccine access for kids, told CNN. "Denying children a critical tool to prevent avoidable tragedies will be a lasting stain on every policymaker who allowed it to happen."

In May, Kennedy announced that the U.S. Centers for Disease Control and Prevention would no longer recommend COVID-19 vaccines for healthy children or pregnant women. The CDC's immunization schedule was updated to reflect that children would be able to get the vaccines after consulting with a health care provider — what's known as "shared decision-making." For pregnant women, there is no recommendation.

Last week, the American Academy of Pediatrics broke from the CDC to recommend COVID-19 shots for infants and young children. The American College of Obstetricians and Gynecologists also reaffirmed support for COVID-19 vaccination during pregnancy,

FDA Commissioner Dr. Marty Makary and Dr. Vinay Prasad, director of FDA's Center for Biologics Evaluation and Research, also said in May there's not enough evidence that healthy children and adults get clinically meaningful benefit from regular COVID-19 shots, and they want to see more placebo-controlled trials, particularly in adults 50 to 64, before recommending the shots for other groups.

COVID-19 levels have been on the rise for months in the U.S., with transmission increasing in most states, according to the U.S. Centers for Disease Control and Prevention. Surveillance data from WastewaterSCAN suggests that virus levels are about half of what they were during the peak of last summer's surge, but they match the peak from the summer of 2023 – and epidemiologist Caitlin Rivers noted in her newsletter that there are no signs of slowing yet, so activity will likely continue to increase in the weeks ahead. There were about 1.7 Covid hospitalizations for every 100,000 people during the first week of August, CDC data shows, with the highest rates among seniors and children under 5.

Fuente: WCVB. Disponible en <https://n9.cl/75bfg>



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Estrategia de búsqueda: (Vaccine) AND DP:([18.08.2025 TO 31.08.2025]) as the publication date 67 records.

1. WO/2025/176145 NINE-VALENT HUMAN PAPILLOMAVIRUS VACCINE AND USE THEREOF

WO - 28.08.2025

Clasificación Internacional A61K 39/295 N° de solicitud PCT/CN2025/078034 Solicitante BEIJING HEALTH GUARD BIOTECHNOLOGY INC. Inventor/a LIU, Yongjiang

Provided are a nine-valent human papillomavirus vaccine and use thereof. The nine-valent human papillomavirus vaccine comprises HPV6, HPV11, HPV16, HPV18, HPV31, HPV33, HPV45, HPV52, and HPV58 type truncated L1-VLP antigens. All antigen components in the provided nine-valent HPV vaccine are derived from L1, a major capsid protein of the HPV, and truncated. The yield of the L1 protein is improved by means of a truncated L1 protein expressed by an Escherichia coli system, thereby reducing the production

cost of the vaccine. Optimizing the particle size of each type of VLP antigens and the particle size of the aluminum hydroxide adjuvant, combined with control over the ratio of each type of VLP antigens to the adjuvant and the buffer system and the like, enables the obtained nine-valent HPV vaccine to induce a high-level immune response, especially to demonstrate a significant clinical effect.

2. 20250269009 VACCINE COMPOSITION FOR NOVEL CORONAVIRUS INFECTION

US - 28.08.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18267355 Solicitante RONGSEN BIOTECHNOLOGY (BEIJING) CO., LTD Inventor/a Baidong HOU

The present application relates to a novel coronavirus pathogen-like antigen (PLA) vaccine, a preparation method therefor and an application thereof. The PLA vaccine consists of structurally-modified *Escherichia coli* virus-like particles (VLPs) and novel coronavirus antigens displayed thereon, and nucleic acid is encapsulated inside of the VLPs. The novel coronavirus PLA vaccine of the present invention formed by passing through modifications effectively prevents the aggregation or precipitation of particles, facilitating the production of the vaccine and ensuring the stability of vaccine efficacy; in addition, relative to conventional vaccines which require additionally adding an additional adjuvant, the PLA-SARS-CoV2 vaccine of the present invention is capable of inducing a significantly higher level of specific antibodies and neutralizing antibodies, having a significantly higher efficacy in challenge tests relative to conventional vaccines.

3. 824117 PRODUCTION OF POXVIRUSES FROM QUAIL CELL CULTURES

NZ - 29.08.2025

Clasificación Internacional C12N 7/00Nº de solicitud 824117 Solicitante BAVARIAN NORDIC A/S Inventor/a HAUSMANN, Jürgen

The present invention relates to methods of producing poxvirus viral vector-based vaccine products from avian cell lines. In some embodiments, the avian cells are suspension quail cell lines. Pharmaceutical compositions such as vaccines produced by the methods of the invention are also provided. In some embodiments, the poxvirus viral vector is Modified Vaccinia Virus Ankara ("MVA") or recombinant MVA. In some embodiments, the recombinant MVA encodes heterologous antigens and can be used to produce a vaccine against the antigens. In some embodiments, the recombinant MVA encodes antigens of Respiratory Syncytial Virus (RSV) and the avian cells are used to produce a vaccine against RSV comprising the recombinant MVA and/or the encoded antigens.

4. 20250269006 METHODS FOR PREDICTING EFFICACY OF A MODIFIED LIVE PORCINE REPRODUCTIVE AND RESPIRATORY SYNDROME VIRUS (PRRSV) VACCINE

US - 28.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 18866887 Solicitante ELANCO US INC. Inventor/a Mark Hammer

Methods are provided for eliciting heterologous immunogenicity against heterologous porcine reproductive and respiratory syndrome virus (PRRSV) strains to allow assessment of innate immunity and adaptive immunity. In other aspects are provided methods for determining the efficacy of a vaccine against PRRSV. In

still other aspects are provided methods for predicting the efficacy of a vaccine against PRRSV in pigs suspected of having an infection with PRRSV.

5. WO/2025/179050 VACCINE COMPOSITIONS AND METHODS OF USE

WO - 28.08.2025

Clasificación Internacional A61K 40/42Nº de solicitud PCT/US2025/016645 Solicitante BIONTECH US INC. Inventor/a FOLEY, Kendra

The present disclosure provides a therapy with vaccine compositions including multiepitopic polypeptides, recombinant nucleic acids encoding multiepitopic polypeptides and methods of use thereof. Also provided herein are T-cell receptors (TCRs) that can recognize one or more epitopes of the multiepitopic polypeptides. The therapy provided herein can comprise (i) a TCR, a recombinant nucleic acid encoding the TCR, or cells comprising the TCR or the recombinant nucleic acid encoding the TCR, and (ii) a vaccine composition described herein. The methods and compositions provided herein can be used for the treatment of cancer.

6. 824298 VACCINE COMPOSITION COMPRISING A SYSTEM FOR DELIVERING AN INACTIVATED WHOLE BACTERIUM VIA CATIONIC POLYSACCHARIDE NANOPARTICLES WITHOUT ANY ADJUVANT

NZ - 29.08.2025

Clasificación Internacional A61K 9/51Nº de solicitud 824298 Solicitante VAXINANO Inventor/a BETBEDER, Didier

The invention relates to the field of vaccine compositions. The invention more particularly relates to a prophylactic vaccine composition that is intended for mammals and birds and comprises a killed whole bacterium, said bacterium being covered with a cationic agent, in particular cationic nanoparticles.

7. 202430707 CYCLIN D1 BASED CANCER VACCINE

DK - 19.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud PA 2024 30707 Solicitante Anders Kaare Nørgaard Inventor/a Anders Kaare Nørgaard

The present disclosure provides a vaccine against cancers that express both high levels of Cyclin D1 and MHC-II. That includes hormone positive breast cancers and some prostate cancers and melanoma. Cyclin D1 is not presented well on MHC-I, and it expresses very little on non-cancerous tissues, so that it is possible, based on a genomic test of the individual, to design a polytope based on epitopes from Cyclin D1 that in most cases display on MHC-II and not on MHC-I, such that the cancer after vaccination with the polytope will reveal itself to the immune system and get eradicated while there are little side effects in that the vaccine will not make the immune system harm the healthy tissue. Since the invention utilizes the fact that the cancer expresses MHC-II, it may work with other antigens than Cyclin D1 as long as the cancer expresses MHC-II.

8. WO/2025/171792 ANTI-TUMOR COMBINATION VACCINE

WO - 21.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/CN2025/077276Solicitante SHAOXING YUERAN BIOMEDICAL TECHNOLOGY CO., LTD.Inventor/a XU, Honghui

Provided is use of a combination of three or more of the following microorganisms in the preparation of an anti-tumor combination **vaccine**: Bordetella pertussis, Salmonella typhi, Salmonella paratyphi A, Salmonella paratyphi B, Staphylococcus aureus, Listeria monocytogenes, Escherichia coli, Proteus bacillus vulgaris, Lactic acid bacteria, Bifidobacterium longum, Bordetella pertussis, Corynebacterium diphtheriae and Clostridium tetani, Clostridium acetobutylicum, Salmonella typhimurium, and Streptococcus pyogenes. The anti-tumor combination **vaccine** is broad-spectrum, safe and non-toxic, and thus has the prospect of clinical applications.

9.3826600ISFORET VACCINEKØLESKAB

DK - 25.08.2025

Clasificación Internacional A61J 1/16Nº de solicitud 19742747Solicitante B Medical Systems S.à.r.l.Inventor/a RIES, Gilles

An ice-lined **vaccine** refrigerator (10) comprises: a **vaccine** storage compartment (15); an electrically powered cooling circuit (16), the electrically powered cooling circuit being configured to generate an ice-lining and to cool the **vaccine** storage compartment; an AC power inlet (17) adapted for connection to an external supply of AC power; and a refrigerant compressor (21) forming part of the electrically powered cooling circuit and adapted to be powered by the external supply of AC power through the AC power inlet. Reliability is improved by using a DC powered compressor and an AC/DC convertor (24) to convert AC power received at the AC power inlet to DC power to power the compressor.

10.20250269005METHODS FOR TREATING, AMELIORATING OR PREVENTING INFECTIONS USING DRUG AND VACCINATION COMBINATION TREATMENT

US - 28.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 18290146Solicitante TOPELIA AUST LIMITED (ACN 652 771 670)Inventor/a Thomas Julius BORODY

In alternative embodiments, provided are methods for treating, ameliorating, decreasing the chances of having any adverse effects from, decreasing the severity of adverse effects from, or preventing an infection, or boosting or enhancing natural immunity acquired by an infected individual, by administration of: an inactivated infectious causative agent of the infection, or an antibiotic and/or an anti-viral drugs and a **vaccine** directed to a causative agent of the infection and/or an attenuated and/or a live, viable or infectious causative agent of the infection. In alternative embodiments, the infection is bacterial or viral. In alternative embodiments, the viral infection is a coronavirus infection such a Covid-19 infection. In alternative embodiments, methods as provide herein prevent or decrease the prevalence or severity of "**vaccine** breakthrough infections" after vaccination, where external mutants of COVID-19 infect patients in spite of the fact that they have undergone immunization, for example, to prevent a mutant or variant COVID-19 infection. In alternative embodiments, an antiviral combination administered in coordination with a **vaccine** comprises PF-07321332 or PAXLOVID™ and/or ritonavir, or ivermectin, doxycycline and a zinc or a zinc salt. In alternative embodiments, methods as provided herein are used to prevent in vivo mutations of such mutant infectious agent to enhance the efficacy of an administered vaccination; in other words, methods as provided herein are used to prevent in vivo replication of an acquired viral mutant or variant infectious

agent, and thus also prevents ongoing mutations of the viral infectious agent because using the combination antiviral co-therapy where there is no replication of infectious agent and so there is no possible further mutation of the infectious agent.

11. 4604997 REKOMBINANTE PROTEINIMPFSTOFFE MIT ENANTIOSPEZIFISCHEM KATIONISCHEM LIPID R-DOTAP UND VERFAHREN ZUR VERWENDUNG DAVON

EP - 27.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud 23880615 Solicitante PDS BIOTECHNOLOGY CORP Inventor/a CONN GREGORY

Provided herein are **vaccine** compositions including recombinant protein antigens derived from computationally optimized broadly reactive influenza antigen (COBRA) proteins and an immunomodulator, and methods of use thereof. The **vaccine** compositions include one or more COBRA proteins, and the immunomodulator is a cationic lipid. The cationic lipid includes R-DOTAP. The methods of use of the **vaccine** compositions includes methods of inducing a humoral immune response against influenza viruses, methods of inducing polyfunctional CD8+ and CD4+ effector T cells against influenza viruses, methods of inducing memory T cells against influenza viruses, methods of enhancing immunity against influenza viruses, and methods of inducing balanced Th1/Th2 immune response against influenza viruses in a subject.

12. 20250262288 CELLULAR ADJUVANTS FOR VIRAL INFECTION

US - 21.08.2025

Clasificación Internacional A61K 39/108Nº de solicitud 19203533 Solicitante NantBio, Inc. Inventor/a Kayvan Niazi

Two-component **vaccine** formulations and methods are contemplated where the **vaccine** has an adjuvant component and a therapeutic component. The therapeutic component comprises preferably a recombinant therapeutic virus encoding a therapeutic antigen while the adjuvant component comprises a non-host cell or immune stimulating portion thereof. Notably, use of the adjuvant component will result in significant uptake of the therapeutic component into immune competent cells, even in the absence of receptors for entry of the therapeutic component. In addition, such adjuvant also stimulates expression of the therapeutic antigen.

13. 0002845571 **VACCINE** BASED ON AN ADENO-ASSOCIATED TYPE 9 VECTOR ENCODING SPIKE PROTEIN FOR THE PREVENTION OF SARS-COV-2 CORONAVIRUS INFECTION

RU - 21.08.2025

Clasificación Internacional C12N 15/50Nº de solicitud 2024131226 Solicitante Inventor/a Фролова Мария Евгеньевна (RU)

FIELD: medicine. SUBSTANCE: invention relates to molecular biology, biotechnology, and medicine. A genetic construct is described for the expression in human or animal cells of a polynucleotide encoding a full-length spike protein of the SARS-CoV-2 virus containing mutations: partial deletion of the RRAR furine site with the removal of all arginines (R) included in the site, as well as mutations K986P, V987P and D614G. The construct is represented by the adeno-associated AAV9 virus, into which the specified polynucleotide is cloned. A recombinant adeno-associated viral vector AAV9 is described, which provides expression in human or animal cells of a polynucleotide encoding a full-length spike protein of the SARS-CoV-2 virus with a genetic

construct. The application of a genetic construct or recombinant adeno-associated viral vector AAV9 for the induction of an immune response against the SARS-CoV-2 virus is presented. EFFECT: the developed genetic construct and a vaccine based on it expand the arsenal of means for inducing specific immunity to the SARS-CoV-2 virus and can be used to effectively stimulate the humoral and cellular response in order to prevent coronavirus infection caused by SARS-CoV-2. 9 cl, 16 dwg, 6 tbl, 5 ex

14. 4601686 VERWENDUNG EINES DENGUE-IMPfstOFFS BEI PERSONEN MIT SCHWANGEREN UND/ODER STILLEN

EP - 20.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23805387 Solicitante TAKEDA VACCINES INC Inventor/a KLAS SHERI DENÉT

A dengue vaccine for use in a method of protecting against dengue disease in a pregnant and/or breastfeeding human subject.

15. 4223311 INDIVIDUALISIEREDE VACCINER MOD CANCER

DK - 25.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 23169466 Solicitante BioNTech SE Inventor/a SAHIN, Ugur

The present invention relates to the provision of vaccines which are specific for a patient's tumor and are potentially useful for immunotherapy of the primary tumor as well as tumor metastases. In one aspect, the present invention relates to a method for providing an individualized cancer vaccine comprising the steps: (a) identifying cancer specific somatic mutations in a tumor specimen of a cancer patient to provide a cancer mutation signature of the patient; and (b) providing a vaccine featuring the cancer mutation signature obtained in step (a). In a further aspect, the present invention relates to vaccines which are obtainable by said method.

16. 3609535 HBV-VACCINE

DK - 18.08.2025

Clasificación Internacional A61K 39/29Nº de solicitud 18719248 Solicitante Oxford University Innovation Ltd. Inventor/a BARNES, Eleanor

The invention relates to a multi-HBV immunogen viral vector vaccine comprising: a viral vector comprising an immunogen expression cassette, wherein the expression of a protein encoded by the expression cassette is arranged to be driven by a promoter, wherein the immunogen expression cassette encodes: a) HBV Core; b) a modified HBV polymerase (P_{mut}), wherein the modification is a mutation to wild-type HBV polymerase to substantially remove polymerase function; c) HBV surface antigen (HbsAg); and d) an intergenic sequence that is arranged to cause expression of at least the HBV surface antigen (HbsAg) as a separate protein from the HBV core and the modified HBV polymerase (P_{mut}), wherein the intergenic sequence is downstream (3') of the sequences encoding the HBV core and the modified HBV polymerase (P_{mut}) and upstream (5') of the sequence encoding the HBV surface antigen (HbsAg); and related compositions, vaccination methods and methods of treatment or prophylaxis of HBV infection.

17. WO/2025/174962 NOVEL SAFE POTENT SINGLE VECTOR PLATFORM VACCINES AGAINST MELIOIDOSIS AND GLANDERS

WO - 21.08.2025

Clasificación Internacional A61K 39/104Nº de solicitud PCT/US2025/015720 Solicitante THE REGENTS OF THE UNIVERSITY OF CALIFORNIA Inventor/a HORWITZ, Marcus A.

Embodiments of the invention include single vector platform vaccine compositions for preventing the disease Melioidosis caused by *Burkholderia pseudomallei* in humans and animals. Embodiments of the invention include single vector platform vaccine compositions for preventing the disease glanders caused by *Burkholderia mallei* in humans and animals. Embodiments of the invention include methods of immunizing a susceptible host against *Burkholderia pseudomallei* and/or *Burkholderia mallei* using the disclosed compositions.

18. 20250263448 VETERINARY VACCINES AND METHODS FOR THE TREATMENT OF PASTEURELLA MULTOCIDA INFECTIONS IN FOOD PRODUCTION ANIMALS

US - 21.08.2025

Clasificación Internacional C07K 14/285Nº de solicitud 18858473 Solicitante ENGINEERED ANTIGENS INC. Inventor/a Trevor MORAES

Disclosed are novel veterinary vaccine compositions comprising a *P. multocida* PmSLP protein or an immunogenically equivalent portion thereof. The vaccine compositions may be used to ameliorate, treat or prevent pathogenic infections of food production animals, such as bovine and porcine animals, caused by *Pasteurella multocida*. Related methods and uses are also disclosed.

19. 20250262292 VETERINARY VACCINES AND METHODS FOR THE TREATMENT OF PASTEURELLA MULTOCIDA INFECTIONS IN FOOD PRODUCTION ANIMALS

US - 21.08.2025

Clasificación Internacional A61K 39/102Nº de solicitud 18858510 Solicitante ENGINEERED ANTIGENS INC. Inventor/a Trevor MORAES

Disclosed are novel veterinary vaccine compositions comprising a *P. multocida* PmSLP protein or an immunogenically equivalent portion thereof. The vaccine compositions may be used to ameliorate, treat or prevent pathogenic infections of food production animals, such as bovine and porcine animals, caused by *Pasteurella multocida*. Related methods and uses are also disclosed.

20. WO/2025/172363 VACCINE AGAINST BOVINE VIRAL DIARRHOEA VIRUS AND BOVINE RESPIRATORY DISEASE

WO - 21.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2025/053726 Solicitante HIPRA SCIENTIFIC, S.L.U. Inventor/a GIBERT LLEIXA, Marta

The present invention relates to an immunogenic or vaccine composition for use in the treatment and/or prevention of Bovine Viral Diarrhoea Virus type 1 (BVDV-1) and/or Bovine Viral Diarrhoea Virus type 2 (BVDV-2) infection in cattle and associated Bovine Respiratory Disease.

21. 20250270296 POTENT NEUTRALIZING ANTIBODIES FOR PREVENTION AND TREATMENT OF COVID-19

US - 28.08.2025

Clasificación Internacional C07K 16/10Nº de solicitud 18028687Solicitante Academia SinicaInventor/a Han-Chung WU

Potent neutralizing antibodies for prevention and treatment of covid-19. Human chimeric antibodies (RBD-chAbs) specifically against SARS-COV-2 Spike(S) receptor-binding domain (RBD) are disclosed. Antibody cocktails or vaccine compositions comprising the RBD-chAbs are also disclosed. The RBD-chAbs, the antibody cocktails, and the vaccine compositions are effective for protection and/or treatment of COVID-19 and are potent against COVID-19 variants including United Kingdom variant B.1.1.7 (Alpha), South African variant B.1.351 (Beta), Brazil variant PI (Gamma), California variant B.1.429 (Epsilon), New York variant B.1.526 (Iota), Indian variants B.1.617.1 (Kappa) and B.1.617.2 (Delta).

22. 20250269010 NOVEL NUCLEIC ACID MOLECULE

US - 28.08.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18276086Solicitante ST PHARM CO., LTD.Inventor/a Kyungjin KIM

The present invention relates to a nucleic acid molecule for preventing or treating viral infection or cancer, comprising a nucleic acid encoding a signal peptide and a nucleic acid encoding an antigen. Furthermore, it relates to a vaccine composition for preventing or treating viral infection or cancer, comprising the nucleic acid molecule.

The nucleic acid molecule according to the present invention has excellent intracellular protein expression rate and protein secretion ability to the outside of the cell. In addition, when administered in vivo, it allows the subject to acquire humoral immunity, such as inducing antigen-specific neutralizing antibodies, and cellular immunity, such as increasing the amount of immune cells directly involved in killing the virus. Thus, it can be useful as a vaccine for the prevention and treatment of viral infection or cancer.

23. 4604994 DENGUE-IMPfstoffFORMULIERUNG

EP - 27.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23804875Solicitante TAKEDA VACCINES INCInventor/a KOMMAREDDY SUSHMA

The present invention relates to a dengue vaccine formulation comprising a tetravalent dengue virus composition comprising a live attenuated dengue virus serotype 1, a live attenuated dengue virus serotype 2, a live attenuated dengue virus serotype 3, and a live attenuated dengue virus serotype 4, at least one non-reducing disaccharide, at least one poloxamer, urea, at least one amino acid having a positively charged side chain at neutral pH, tromethamine, and human serum albumin.

24. 4606887 KULTURVERFAHREN FÜR FISCHZELLEN UND DAFÜR VERWENDETES ZELLKULTURMEDIUM SOWIE ORALER IMPfstoff FÜR FISCHE

EP - 27.08.2025

Clasificación Internacional C12N 5/071Nº de solicitud 23879808Solicitante BIOSIENSE CO
LTDInventor/a OKUTANI ASUKA

The present disclosure provides a cell culture medium for use in culturing of fish cells, a method for culturing fish cells, and an oral vaccine for fish. In the present disclosure, the cell culture medium for use in culturing of fish cells preferably comprises serum obtained from adult fish.

25.4604990NEUARTIGE T-ZELLEN-AKTIVIERENDE IMMUNTHERAPEUTIKUM ZUR BEHANDLUNG
VON MUCIN-1-PROTEIN-EXPRIMIERENDEM MENSCHLICHEM KREBS

EP - 27.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 23880573Solicitante PDS BIOTECHNOLOGY
CORPInventor/a BEDU-ADDO FRANK

Provided herein are multiepitope peptides including at least one mucin 1 (MUC1) peptide, the multiepitope peptides have MHC affinity for at least one of HLA serotype and are recognized by a CD4+ T cell receptor and/or by a CD8+ T cell receptor. Also provided herein are compositions comprising the multiepitope peptides and a cationic lipid, including vaccine compositions. In various aspects, the cationic lipid is R-DOTAP. The invention also provides methods of use of the multiepitope peptides and of the compositions and vaccine compositions. The methods of use include methods of treating cancer and method of inducing a MUC-specific polyfunctional cytolytic T cell response in a subject.

26.4604995KOMBINATIONSIMPFSTOFFE GEGEN CORONAVIRUS-INFEKTION, INFLUENZA-
INFEKTION UND/ODER RSV-INFEKTION

EP - 27.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23806135Solicitante BIONTECH SEInventor/a SAHIN
UGUR

This disclosure relates to the field of RNA to prevent or treat multiple infectious agents. In particular, the present disclosure relates to methods and agents for vaccination against coronavirus infection, influenza infection, and/or RSV infection and inducing effective coronavirus, influenza virus, and/or RSV antigen-specific immune responses such as antibody and/or T cell responses. Specifically, in one embodiment, the present disclosure relates to methods comprising administering to a subject (i) a bivalent RNA vaccine encoding peptides or proteins comprising epitopes of SARS-CoV-2 spike proteins (S proteins) and (ii) a tetravalent RNA vaccine encoding peptides or proteins comprising epitopes of hemagglutinin (HA), for inducing an immune response against coronavirus S proteins, in particular S proteins of SARS-CoV-2, and influenza proteins, in particular HA proteins of type A and type B influenza viruses, in the subject.

27.4601681SAPONIN-DMLT-ADJUVANTEN UND ZUGEHÖRIGE VERWENDUNGEN

EP - 20.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 23878219Solicitante Q VANT BIOSCIENCES
INCInventor/a NORTON ELIZABETH

A composition, preferably an immune adjuvant system, for vaccines containing a saponin component and a dmLT component. The combination of the saponin component and the dmLT component shows a synergistic effect in the treatment of various conditions, illnesses and diseases. The methods of use for prophylactic use or therapeutic treatment are disclosed. Exemplary adjuvant compositions include a double-mutant heat-labile

toxin adjuvant derived from an *Escherichia coli* enterotoxin and a saponin, optionally with an additional vaccine component (e.g., an antigen), particularly when used in a vaccine.

28. WO/2025/175807 METHOD FOR PREPARING STREPTOCOCCUS PNEUMONIAE CAPSULAR POLYSACCHARIDE

WO - 28.08.2025

Clasificación Internacional C08B 37/00Nº de solicitud PCT/CN2024/126421 Solicitante BEIJING MINHAI BIOTECHNOLOGY CO., LTD. Inventor/a LI, Yuelong

The present invention relates to the technical field of biologics, and in particular to a method for preparing a *Streptococcus pneumoniae* capsular polysaccharide. The present invention provides a method for preparing a *Streptococcus pneumoniae* capsular polysaccharide. The method comprises: using β -propiolactone to treat a *Streptococcus pneumoniae* fermentation culture, separating and collecting a supernatant, purifying the supernatant, and collecting a capsular polysaccharide. In the present invention, β -propiolactone is used to treat the *Streptococcus pneumoniae* fermentation culture, thereby reducing residual impurities such as proteins, and achieving relatively high polysaccharide yield; in addition, the β -propiolactone is susceptible to hydrolysis, and the hydrolysate is non-toxic and harmless and has higher safety. The *Streptococcus pneumoniae* capsular polysaccharide prepared by the method of the present invention have quality control indexes meeting requirements, and can be used for preparing a *Streptococcus pneumoniae* capsular polysaccharide vaccine and a polysaccharide conjugate vaccine.

29. WO/2025/172775 MACHINE LEARNING APPROACH TO PREDICT VIRUS OR CANCER MUTATIONS FOR VACCINE PRODUCTION OR DRUG DESIGN

WO - 21.08.2025

Clasificación Internacional G16B 20/50Nº de solicitud PCT/IB2025/050405 Solicitante NEC LABORATORIES EUROPE GMBH Inventor/a SIARHEYEU, Raman

A computer-implemented, machine learning method for predicting a top-k most likely mutated molecular sequences includes encoding a molecular sequence at a first time into a first vector in a latent space. A second vector is generated, by a time-varying mutation models, in the latent space using as input the first vector. The second vector indicates a time-varying influence of the molecular sequence on a mutated version of the molecular sequence at a subsequent time. The second vector is decoded to generate a prediction of the top-k most likely mutated molecular sequences for the molecular sequence at the subsequent time. The method has applications including, but not limited to, use cases in computational biology and medical AI and healthcare for optimizing vaccine design or supporting decision making in diagnosis and treatment of patients.

30. 20250269007 NUCLEIC ACID MOLECULES AND USES THEREOF

US - 28.08.2025

Clasificación Internacional A61K 39/125Nº de solicitud 19204075 Solicitante CureVac SE Inventor/a Susanne RAUCH

The present invention is directed to an artificial nucleic acid and to polypeptides suitable for use in treatment or prophylaxis of an infection with Norovirus or a disorder related to such an infection. In particular, the present

invention concerns a Norovirus **vaccine**. The present invention is directed to an artificial nucleic acid, polypeptides, compositions and vaccines comprising the artificial nucleic acid or the polypeptides. The invention further concerns a method of treating or preventing a disorder or a disease, first and second medical uses of the artificial nucleic acid, polypeptides, compositions and vaccines. Further, the invention is directed to a kit, particularly to a kit of parts, comprising the artificial nucleic acid, polypeptides, compositions and vaccines.

31. 20250263442 RECOMBINANT CEDAR VIRUS CHIMERAS

US - 21.08.2025

Clasificación Internacional C07K 14/005Nº de solicitud 18557247Solicitante THE HENRY M. JACKSON FOUNDATION FOR THE ADVANCEMENT OF MILITARY MEDICINE, INC.Inventor/a Christopher C. Broder

Described herein are replication-competent recombinant Cedar virus chimeras are described that are engineered to express antigenic surface or soluble proteins/polypeptides of a non-CedV henipavirus, such as of a pathogenic henipavirus, such as Nipah virus or Hendra virus. **Vaccine** compositions containing the recombinant Cedar virus chimeras are also described, as are therapeutic methods and uses for protecting against pathogenic henipavirus infection.

32. 20250269008 VACCINATION OF IMMUNOCOMPROMISED SUBJECTS

US - 28.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud 19182727Solicitante Seqirus UK LimitedInventor/a Giuseppe DEL GIUDICE

Disclosed herein are methods for enhancing immune responses to a **vaccine** in immunocompromised individuals, including those receiving a slatin therapy. Related products are also provided.

33. 4604998 MATERIALIEN UND VERFAHREN ZUR BEHANDLUNG VON EPSTEIN-BARR-VIRUS (EBV) UND EBV-INDUZIERTEN ERKRANKUNGEN

EP - 27.08.2025

Clasificación Internacional A61K 39/245Nº de solicitud 23794305Solicitante UNIV WIEN MEDInventor/a VIETZEN HANNES

The present invention relates to means and methods to prevent and/or treat Epstein-Barr virus (EBV) and EBV-induced diseases, such as EBV infection, infectious mononucleosis (IM), malignant or non-malignant post-transplant lymphoproliferative disorder (PTLD) and other EBV-associated diseases. In particular, the invention provides a SQAPLPCVL peptide that can be used in a treatment or a method of treatment to induce an EBV-specific immune response in a subject. The SQAPLPCVL can be used in a treatment or method of treatment as a **vaccine** against EBV and EBV-induced diseases. It is preferred herein that Epstein-Barr virus (EBV) and/or EBV-induced diseases are prevented.

34. WO/2025/178495 METHOD OF PREPARING A MICRONEEDLE ARRAY

WO - 28.08.2025

Clasificación Internacional A61M 37/00Nº de solicitud PCT/NL2025/050088Solicitante CERAVX B.V.Inventor/a DE OLIVEIRA, Daniel Campling

The invention relates to a method of preparing a microneedle array, comprising the steps of dissolving a water-soluble copolymer, comprising maleic anhydride and isobutylene monomers, in an aqueous solvent to obtain an aqueous solution, adding a ceramic material to said aqueous solution to obtain an aqueous ceramic slurry for gel casting, adding at least a portion of said aqueous ceramic slurry to a mould to obtain a layer of aqueous ceramic slurry in said mould, degassing said layer of aqueous ceramic slurry to obtain a degassed layer of aqueous ceramic slurry in said mould, gelling said degassed layer of aqueous ceramic slurry to remove said aqueous solvent from said aqueous ceramic slurry and removing said mould to obtain a gelled tape, drying said gelled tape to obtain a green tape, sintering said green tape to obtain said microneedle array. Further, the invention provides for a green tape, comprising a base plate and a set of microneedles integrated with said base plate, a microneedle array, comprising a base plate and a set of microneedles integrated with said base plate, a system for enabling transport of a substance through a material barrier, and a use of a microneedle array for intradermal drug or vaccine delivery, diagnostics, cosmeceuticals,, sensing of biomarkers found in the skin and monitoring of physiological conditions of the body.

35. 20250269003 NANO-ENHANCED VACCINE

US - 28.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18859144Solicitante University of Virginia Patent FoundationInventor/a Mark Kester

Provided are composition that include stable TLR4 agonist (e.g., KDO2) containing nanoliposomes. In some embodiments, the TLR4 agonist (e.g., KDO2) containing nanoliposome include a lipid component comprising, consisting essentially of, or consisting of DSPC, DOPE, PEG(2000)-PE, one or more TLR4 agonists (e.g., KDO2), Cholesterol, Rhodamine or DiD, and optionally DOTAP and/or DHP. In some embodiments, the TLR4 agonist (e.g., KDO2) containing nanoliposomes are cationic, anionic, or neutral liposomes. In some embodiments, the TLR4 agonist (e.g., KDO2) containing nanoliposome encapsulate one or more immunogenic peptides, which can be peptides associated with malignant melanoma, which optionally can be subsequences of tyrosinase, gp100, MAGE-1,2,3,6, Melan-A/MART-1, and/or MAGE-3. Also provided are methods for treating and/or preventing malignant melanoma and for inducing anti-melanoma immune responses in subjects using the presently disclosed compositions.

36. WO/2025/172435 HBV ANTIGEN FORMULATION FOR TREATING HEPATITIS B

WO - 21.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2025/053853Solicitante HELMHOLTZ ZENTRUM MÜNCHEN - DEUTSCHES FORSCHUNGSZENTRUM FÜR GESUNDHEIT UND UMWELT (GMBH)Inventor/a PROTZER, Ulrike

The invention relates to methods of vaccination, comprising administering to a human subject a first dose of an HBcAg particle, an HBsAg, and c-di-AMP; a second dose of the HBcAg particle, the HBsAg, and c-di-AMP; and a dose of a vaccine vector. The invention also relates pharmaceutical compositions, combinations of HBcAg particles, HBsAg and c-di-AMP, and kits for use in said methods.

37. 20250262291 MICROPARTICLES FROM STREPTOCOCCUS PNEUMONIAE AS VACCINE ANTIGENS

US - 21.08.2025

Clasificación Internacional A61K 39/09Nº de solicitud 19200040Solicitante ZalVac ABInventor/a Birgitta Henriques Normark

An isolated *Streptococcus pneumoniae* membrane vesicle microparticle (MP), wherein said MP comprises: the protein Ply at the level of ≥ 0.070 $\mu\text{g}/\text{mg}$ total protein in the MP; and/or the protein LytA at the level of ≥ 0.070 $\mu\text{g}/\text{mg}$ total protein in the MP; and/or the protein PspC at the level of ≥ 0.130 $\mu\text{g}/\text{mg}$ total protein in the MP; and/or the protein RrgB at the level of ≥ 0.020 $\mu\text{g}/\text{mg}$ total protein in the MP. Compositions comprising such MPs. Uses thereof in particular in immunization, as well as methods of manufacture thereof.

38.20250273291SELECTION OF DIVERSE CANDIDATE PEPTIDES FOR PEPTIDE THERAPEUTICS

US - 28.08.2025

Clasificación Internacional G16B 15/30Nº de solicitud 19207999Solicitante Genentech, Inc.Inventor/a William John THRIFT

A method for developing a therapeutic such as, for example, a peptide vaccine. A machine learning model is trained using a metric learning algorithm, training peptide sequence data, and training allele presentation data corresponding to the training peptide sequence data. Peptide sequence data identifying peptide sequences that correspond to peptides is received. A peptide sequence vector is generated, via a machine learning model, for each peptide sequence using the peptide sequence data to form a plurality of peptide sequence vectors. An output is generated using the plurality of peptide sequence vectors. The output provides an indication of similarity between peptide sequences of the plurality of peptide sequences. A group of candidate peptides is selected from the plurality of peptides for development of the therapeutic based on the output such that the group of candidate peptides includes at least two dissimilar candidate peptides.

39.4604926NEUARTIGER NICHT-HLA-BESCHRÄNKTER T-ZELL-IMPfstoff FÜR KREBS MIT EXPRESSION VON TCR-GAMMA-ALTERNATIVEM LESERASTERPROTEIN (TARP)-PROTEIN

EP - 27.08.2025

Clasificación Internacional A61K 9/127Nº de solicitud 23880501Solicitante PDS BIOTECHNOLOGY CORPInventor/a BEDU-ADDO FRANK

Novel compositions of TCR Gamma Alternate Reading Frame Protein (TARP) peptides combined with cationic lipids such as the DOTAP and specifically R-DOTAP, induce high levels of TARP-specific polyfunctional cytolytic T-cells. Compositions and methods of use are provided. The compositions comprise N-terminal and C-terminal overlapping peptide sequence pairs duplicating the critical central antigenic region of TARP and encompassing the entire protein selected and designed to be effectively processed by antigen-presenting cells to prime cytotoxic T cells specific for TARP-derived T cell peptide antigens when delivered in combination with immunostimulatory nanoparticles composed of R-DOTAP cationic lipids.

40.WO/2025/176928NANOESTRUCTURAS VÍRICAS SINTÉTICAS Y SUS USOS

WO - 28.08.2025

Clasificación Internacional A61K 9/50Nº de solicitud PCT/ES2025/070090Solicitante FUNDACIÓN INSTITUTO DE INVESTIGACIÓN MARQUÉS DE VALDECILLAInventor/a GARCÍA HEVIA, Lorena

Nanoestructuras víricas sintéticas y sus usos La presente invención se refiere a una nanoestructura que comprende un núcleo magnético recubierto por uno o más componentes celulares, donde dicho núcleo magnético comprende nanopartículas magnéticas, al método de obtener dicha nanoestructura, y al empleo

de dicha nanoestructura como medicamento, como vacuna, o para el análisis in vitro de compuestos capaces de inhibir virus, para la validación in vitro de anticuerpos o como sistema in vitro de biopsia intracelular.

41. 824463 NUCLEOCAPSID ANTIGEN IMMUNOTHERAPY FOR COVID-19 FUSION PROTEINS AND METHODS OF USE

NZ - 29.08.2025

Clasificación Internacional A61K 39/215Nº de solicitud 824463 Solicitante VAKSTON, INC. Inventor/a LANCASTER, Thomas M.

The present disclosure provides recombinantly manufactured fusion proteins comprising a SARS-CoV-2 nucleocapsid protein (N-protein) fragment or an analog thereof linked to a human Fc fragment for use in relation to the 2019 Novel Coronavirus (COVID-19). Embodiments include the administration of the fusion proteins to patients that have recovered from COVID-19 as a booster vaccination, to antibody naive patients to produce antibodies to the SARS-CoV-2 virus to enable the patients to become convalescent plasma donors, to patients who have been infected by the SARS-CoV-2 virus and have contracted COVID-19 in order to limit the scope of the infection and ameliorate the disease, and as a prophylactic COVID-19 vaccine. Exemplary Fc fusion proteins and pharmaceutical formulations of exemplary Fc fusion proteins are provided, in addition to methods of use and preparation.

42. WO/2025/179103 PENTAVALENT AND OCTAVALENT INFLUENZA VIRUS IMMUNOGENS, COMPOSITIONS AND METHODS OF USE THEREOF

WO - 28.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud PCT/US2025/016726 Solicitante UNIVERSITY OF GEORGIA RESEARCH FOUNDATION, INC. Inventor/a ROSS, Ted M.

Provided are an immunogenic multivalent (e.g., pentavalent or octavalent) composition, vaccine, and formulations thereof, composed of five or eight, isolated, non-naturally occurring, immunogenic antigens derived from influenza viruses that are capable of eliciting a broadly reactive immune response, e.g., a broadly reactive neutralizing antibody response directed against influenza virus antigens, following introduction into a subject. Also provided are immunogens and virus-like particles (VLPs) comprising the multivalent composition. Methods of generating an immune response in a human or non-human subject by administering the multivalent composition are provided. In particular, the multivalent immunogen comprises isolated, broadly reactive hemagglutinin (HA) protein antigens of influenza virus strains, such as H1, H3, and IBV, and/or viral neuraminidase (NA) N1 and N2 protein antigens, which may be recombinant and/or recombinantly produced.

43. WO/2025/179238 CORONAVIRUS VACCINE

WO - 28.08.2025

Clasificación Internacional A61P 31/14Nº de solicitud PCT/US2025/016933 Solicitante BIONTECH SE Inventor/a SWANSON, Kena Anne

This disclosure relates to the field of RNA to prevent or treat coronavirus infection. In particular, the present disclosure relates to methods and agents for vaccination against coronavirus infection and inducing effective

coronavirus antigen-specific immune responses such as antibody and/or T cell responses against various SARS-CoV-2 variants of concern.

44. WO/2025/172592 ALPHA SYNUCLEIN THERAPEUTIC VACCINE

WO - 21.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/EP2025/054124 Solicitante AC IMMUNE SA Inventor/a STAFFLER, Günther

The present invention relates to immunogenic compositions that can be employed for the prevention, alleviation or treatment of a condition associated with diseases, disorders and abnormalities associated with alpha synuclein (alpha synuclein, α -synuclein, A-synuclein, aSynuclein, A-syn, α -syn, aSyn, a-syn) aggregates including, but not limited to, Lewy bodies and/or Lewy neurites, such as Parkinson's disease, Multiple System Atrophy, Lewy Body dementia (LBD; dementia with Lewy bodies (DLB) ("pure" Lewy body dementia), Parkinson's disease dementia (PDD)), or Diffuse Lewy Body Disease.

45. 20250273288 ANTIGEN PREDICTIONS FOR INFECTIOUS DISEASE-DERIVED EPITOPES

US - 28.08.2025

Clasificación Internacional G16B 15/30Nº de solicitud 18908082 Solicitante Gritstone Bio, Inc. Inventor/a Joshua Klein

Disclosed herein is a system and methods for determining the alleles, antigens, and infectious disease-based vaccine composition as determined on the basis of a patient's expressed HLA alleles. Additionally described herein are unique infectious disease-derived vaccines.

46. 823892 VACCINE FOR STREPTOCOCCUS EQUI SUBSP. ZOOEPIDEMICUS

NZ - 29.08.2025

Clasificación Internacional A61K 39/09Nº de solicitud 823892 Solicitante University of Saskatchewan Inventor/a COSTA, Matheus

Provided herein is a composition comprising a live strain of *S. zooepidemicus* and a pharmaceutically acceptable carrier, wherein the live strain of *S. zooepidemicus* contains a mutated M protein trans-acting positive regulator (MGA) gene that results in impaired DNA binding. Also provided are methods and uses to eliciting an immune response against an infection by *S. zooepidemicus* in a subject, comprising administering to the subject an effective amount of the composition described herein. Also provided is a method of generating strains of *S. zooepidemicus* with reduced virulence.

47. 20250263704 NUCLEIC ACID-POLYPEPTIDE NANO-PHARMACEUTICAL COMPOSTION FOR TREATING AND PREVENTING HUMAN PAPILLOMA VIRUS INFECTION

US - 21.08.2025

Clasificación Internacional C12N 15/113Nº de solicitud 18561596 Solicitante SIRNAOMICS, INC. Inventor/a Alan Lu

Disclosed is a nucleic acid-polypeptide nano-pharmaceutical composition for treating and preventing human papilloma virus infection. A small interfering nucleic acid siRNA molecule used for inhibiting and treating various diseases caused by a HPV infection can block the virus replication life cycle by means of targeted inhibition of the expression of the HP16/18 key gene, reduce a viral infection and finally remove viruses. A pharmaceutical composition based on the siRNA molecule comprises a siRNA molecule and another molecule, including a siRNA molecule for inhibiting PD-1/PD-L1, a small molecule compound against a HPV infection, a therapeutic mRNA/neoantigen vaccine, and the like. The siRNA molecule and other anti-HPV drugs are coupled by means of a specific chemical bond to form a new coupled molecule, and the composition further includes a pharmaceutically acceptable polypeptide polymer nano-introduction carrier. In some embodiments, the carrier is a histidine-lysine polypeptide polymer nanocarrier.

48. 20250263740 NON-REPLICATING BOVINE INFECTIOUS LYMPHOMA VIRUS (BLV) AND CELLS FOR PRODUCING SAME

US - 21.08.2025

Clasificación Internacional C12N 15/86Nº de solicitud 18857565 Solicitante THE UNIVERSITY OF TOKYO Inventor/a Yoko AIDA

An object of the present invention is to provide a novel non-replicating bovine leukemia virus (BLV) and a producing cell thereof. According to the present invention, there is provided a bovine leukemia virus (BLV) in which at least a part of the function of a pol gene is deficient. Also, according to the present invention, there is provided a non-replicating BLV-producing cell comprising a gene of a BLV in which at least a part of the function of a pol gene is deficient. The present invention is advantageous in that it can provide a BLV vaccine which is highly immunogenic, and is highly safe without replicating in an infected subject.

49. 20250262294 VACCINE

US - 21.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud 18032367 Solicitante THE PIRBRIGHT INSTITUTE Inventor/a Munir Iqbal

The present invention relates to a genetically engineered protein comprising: at least one binding domain which is capable of binding to a cell surface protein on an avian antigen presenting cell; and a) at least one antigenic polypeptide or b) at least one binding domain which is capable of binding to at least one antigenic polypeptide. The present invention also relates to avian vaccines comprising at least one binding domain which is capable of binding to a cell surface protein on an avian antigen presenting cell; and a) at least one antigenic polypeptide or b) at least one binding domain which is capable of binding to at least one antigenic polypeptide and to the use of such vaccines to treat and/or prevent disease in avian subjects.

50. WO/2025/179112 IMMUNOGENIC COMPOSITIONS FOR THE PREVENTION OF HERPES ZOSTER

WO - 28.08.2025

Clasificación Internacional A61K 39/25Nº de solicitud PCT/US2025/016744 Solicitante CUREVO, INC. Inventor/a DE LA ROSA, Guy, R.

The present disclosure provides immunogenic and vaccine compositions comprising a Varicella Zoster Virus (VZV) glycoprotein E (gE) antigen comprising SEQ ID NO: 2 and an SLA adjuvant for use in the prevention of herpes zoster.

51. 20250268997A SUBUNIT CRYPTOCOCCUS VACCINE

US - 28.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18705007Solicitante University of MassachusettsInventor/a Stuart Levitz

Provided herein are subunit vaccines against *Cryptococcus*.

52. 20250270656METHODS AND SYSTEMS FOR DEVELOPING PERSONALIZED VACCINE BY IDENTIFICATION AND PRIORITIZATION OF MUTATION-DERIVED NEOANTIGENS

US - 28.08.2025

Clasificación Internacional C12Q 1/6886Nº de solicitud 19205975Solicitante Personal Genome Diagnostics Inc.Inventor/a Victor Velculescu

Cancer immunology provides promising new avenues for cancer treatment but validation of potential neoantigens to target is costly and expensive. Analysis of MHC binding affinity, antigen processing, similarity to known antigens, predicted expression levels (as mRNA or proteins), self-similarity, and mutant allele frequency, provides screening method to identify and prioritize candidate neoantigens using sequencing data. Methods of the invention thereby save time and money by identifying the priority candidate neoantigens for further experimental validation.

53. 4605412VERFAHREN ZUR HERSTELLUNG REKOMBINANTER TRANSFERRINBINDENDER PROTEINE UND IMPFSTOFFZUSAMMENSETZUNGEN DAMIT

EP - 27.08.2025

Clasificación Internacional C07K 14/22Nº de solicitud 23805182Solicitante SERUM INST OF INDIA PVT LTDInventor/a KARALE ABHIJEET JAGANNATH

The present disclosure relates to manufacturing transferrin binding proteins. Specifically, the present disclosure relates to a simple, scalable, commercially viable fermentation and purification process for obtaining recombinant transferrin binding protein (rTbp-B) along with high recovery, low impurity/ aggregate content, and at the same time retains the integrity of the protein. The method uses a single chromatographic step and does not require tagging of proteins as compared to multiple chromatographic steps used previously and still manages to provide r-Tbp-B with at least 95 % purity.

54. 824465PREFUSION-STABILIZED HMPV F PROTEINS

NZ - 29.08.2025

Clasificación Internacional A61K 39/155Nº de solicitud 824465Solicitante BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEMInventor/a MCLELLAN, Jason

Provided herein are engineered hMPV F proteins. In some aspects, the engineered F proteins exhibit enhanced conformational stability and/or antigenicity. Methods are also provided for use of the engineered F proteins as diagnostics, in screening platforms, and/or in vaccine compositions.

55. 20250262287ARTHROSPIRA PLATENSIS ORAL VACCINE DELIVERY PLATFORM

US - 21.08.2025

Clasificación Internacional A61K 39/015Nº de solicitud 19202734Solicitante Lumen Bioscience, Inc.Inventor/a James ROBERTS

The present disclosure provides oral antigenic compositions comprising a recombinant *Spirulina* comprising at least one exogenous antigenic epitope. Oral antigenic compositions of the present disclosure can be used as vaccines. Oral antigenic compositions of the present disclosure can be used to induce a protective immune response to infectious microorganism, tumor antigens, or self-antigens.

56.20250270298MULTIVALENT ANTI-CAMPYLOBACTER ANTIBODIES AND VACCINE

US - 28.08.2025

Clasificación Internacional C07K 16/12Nº de solicitud 18692967Solicitante BiomEdit, LLCInventor/a Arvind Kumar

One or more antibodies, particularly nanobodies or VHH single domain antibodies, directed against one or more *Campylobacter* bacteria targets with roles in *Campylobacter* colonization, particularly *Campylobacter jejuni*, in animals, particularly in poultry are provided. The nanobodies are useful in reducing or blocking *Campylobacter* colonization or infection. The invention provides methods for reducing or blocking *Campylobacter* colonization or infection, for improving food safety, and for reducing bacterial gastroenteritis in animals, including humans.

57.4604992IMPFSTOFF

EP - 27.08.2025

Clasificación Internacional A61K 39/09Nº de solicitud 23793721Solicitante GLAXOSMITHKLINE BIOLOGICALS SAInventor/a GOMES MORIEL DANILO

The present invention relates to compositions comprising at least one Group A Streptococcus antigen, an aluminium salt and (a) a TLR7 agonist, or (b) a benzonaphthyridine compound. The present invention further relates to methods of making an immunogenic composition comprising at least one antigen, an aluminium salt and (a) a TLR7 agonist, and/or (b) a benzonaphthyridine compound. The present invention also relates to vaccines comprising the compositions, and methods of using and uses of the compositions.

58.0002845642VACCINE FORMULATIONS AGAINST CORONAVIRUS

RU - 25.08.2025

Clasificación Internacional C07K 16/10Nº de solicitud 2022122879SolicitanteInventor/a СМИТ, Гейл (US)

FIELD: biotechnology. SUBSTANCE: described is, different from naturally occurring spike (S) glycoprotein of coronavirus 2 associated with acute respiratory syndrome of transient development (SARS-CoV-2), where different from the naturally occurring S-glycoprotein SARS-CoV-2 contains: (i) an inactivated furin cleavage site; and (ii) proline at amino acid positions 973 and 974; where the amino acids are numbered in accordance with SEQ ID NO: 2. Described is nucleic acid which codes glycoprotein. Presented is a host cell containing vectors with nucleic acid for glycoprotein expression. Disclosed is an immunogenic composition containing: (a) glycoprotein; (b) an adjuvant; and (c) a pharmaceutically acceptable buffer; immunogenic composition containing: (a) a nanoparticle comprising glycoprotein and a core comprising a non-ionic detergent; (b) an adjuvant; and (c) a pharmaceutically acceptable buffer. Disclosed is a method for stimulating the immune response against SARS-CoV-2 in a human, comprising administering a glycoprotein or an immunogenic

composition to the human. EFFECT: invention widens the range of agents for stimulating the immune response against SARS-CoV-2 in humans. 36 cl, 46 dwg, 5 tbl, 8 ex

59. WO/2025/172883 PROCESS FOR THE PURIFICATION OF OMVS FROM GRAM-NEGATIVE BACTERIA, AND RELATIVE USES

WO - 21.08.2025

Clasificación Internacional C07K 14/22Nº de solicitud PCT/IB2025/051536 Solicitante DI CIOCCIO, Vito Inventor/a DI CIOCCIO, Vito

The present invention relates to a process for the purification of vesicles released from the outer membrane (OMVs) of Gram-negative bacteria, the OMVs thus obtained and their relative uses as antigen carriers in vaccine formulations and as a system for conveying the administration of molecules for therapeutic applications.

60. 4601683 GEGEN GRUPPE-B-STREPTOCOCCUS (GBS) GERICHTETER MAP-IMPFFSTOFF

EP - 20.08.2025

Clasificación Internacional A61K 39/02Nº de solicitud 23878304 Solicitante CHILDRENS MEDICAL CT CORP Inventor/a THOMPSON CLAUDETTE

Technologies for the prevention and/or treatment of GBS infections. The technology relates to compositions, including vaccines compositions and methods comprising an immunogenic complex that is a GBS multiple antigen presenting system (MAPS-GBS), where two or more biotinylated GBS polysaccharide antigens are joined together by non-covalent associations with one or more bifunctional fusion proteins comprising, in any order, (i) a sialic acid binding protein (SBD), a GBS polypeptide antigen and (iii) a biotin-binding moiety (BBD), thereby facilitating the linking of multiple GBS polysaccharide antigens together in the complex to form a MAPS-GBS immunogenic complex. The polysaccharide antigens that are linked can be on the same polysaccharide macromolecule or on distinct polysaccharide macromolecules.

61. 4606817 HERSTELLUNG UND ANWENDUNG VON FLAGELLINIMMUNPOTENZIERENDEN DERIVATEN MIT TLR5-AKTIVITÄT UNTER VERWENDUNG EUKARYOTISCHER ZELLEXPRESSIONSSYSTEME

EP - 27.08.2025

Clasificación Internacional C07K 14/28Nº de solicitud 23879958 Solicitante UNIV NAT CHONNAM IND FOUND Inventor/a RHEE JOON HAENG

The present invention relates to the manufacture and application of flagellin immunopotentiating derivatives with TLR5 activity using an eukaryotic cell expression system. The flagellin derivative according to the present invention has a high immunostimulatory effect and can be used in various vaccine and immunotherapy compositions.

62. 823890 RSV F VACCINE FORMULATIONS

NZ - 29.08.2025

Clasificación Internacional A61K 39/155Nº de solicitud 823890Solicitante Novavax, Inc.Inventor/a TIAN, Jing-Hui

Disclosed herein are RSV F glycoproteins and nanoparticles comprising the same, which are suitable for use in vaccines. The nanoparticles present antigens from pathogens surrounded to and associated with a detergent core resulting in enhanced stability and good immunogenicity. Dosages, formulations, and methods for preparing the vaccines and nanoparticles are also disclosed.

63.20250269002CTA **VACCINE** CASSETTES

US - 28.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 19044124Solicitante Gritstone Bio, Inc.Inventor/a Karin Jooss

Disclosed herein are compositions that include antigen-encoding nucleic acid sequences having multiple iterations of CTA epitope-encoding sequences or Cancer Testis Antigen (CTA)-encoding nucleic acid sequences and KRAS-encoding nucleic acid sequences. Also disclosed are nucleotides, cells, and methods associated with the compositions including their use as vaccines.

64.20250269004ENTEROCOCCUS FAECALIS **VACCINE** AND USES THEREOF

US - 28.08.2025

Clasificación Internacional A61K 39/09Nº de solicitud 18857949Solicitante Vaxcyte, Inc.Inventor/a Jeffery C. Fairman

The present disclosure provides immunogenic compositions comprising at least one recombinant polypeptide antigen derived from an *Enterococcus* bacterium (e.g., *E. faecalis*, *E. faecium*, *E. durans*). The disclosure further provides methods, and uses of the immunogenic compositions, for protecting or treating a subject from infection by an *Enterococcus* bacterium. Such infections may cause, or worsen, conditions such as root canal failure, endocarditis, bacteremia, urinary tract infections, prostatitis, intraabdominal infection, cellulitis, dysbiotic gastrointestinal tract, prosthetic joint infection, or wound infections.

65.824300NUCLEIC ACID MOLECULES

NZ - 29.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 824300Solicitante ASTRAZENECA ABInventor/a LOO, Yueh-Ming

The present disclosure relates to a nucleic acid molecule comprising 5'-UTR and/or 3'-UTR sequences that yield high translation levels. Aspects of the disclosure further relate to nucleic acid molecules suitable for use as a **vaccine** in the treatment and prevention of infectious diseases, including those caused by a coronavirus, compositions comprising said nucleic acid molecules and methods of treating or preventing infectious diseases.

66.20250264467NOVEL HCV NUCLEIC ACIDS AND ENCODED GLYCOPROTEIN ANTIGENS AND METHODS

US - 21.08.2025

Clasificación Internacional G01N 33/569Nº de solicitud 19053963Solicitante Purdue Research FoundationInventor/a Devika Sirohi

The present disclosure provides for novel nucleic acid constructs and methods for expressing HCV E2 protein on a cell surface as an active antigen. The present disclosure provides for a method for evaluation of protein expression. Vaccine formulations, constructs and expression vectors are all within the scope of this disclosure.

67.20250269035INHALABLE LIPID NANOPARTICLE, LIPID-CHARGED MOLECULE CONJUGATE, AND LIPID-ZWITTERIONIC MOLECULE CONJUGATE

US - 28.08.2025

Clasificación Internacional A61K 47/54Nº de solicitud 19205101Solicitante INSTITUTE OF CHEMISTRY, CHINESE ACADEMY OF SCIENCESInventor/a Xueguang LU

Disclosed herein are a lipid-charged molecule conjugate, an inhalable lipid nanoparticle, a preparation method therefor, and use thereof. The lipid-charged molecule conjugate of the present invention comprises a lipid unit and a charged unit. The charged unit is selected from a negative charge unit and/or a positive charge unit. The lipid nanoparticle of the present invention comprises an ionizable lipid, an auxiliary phospholipid, cholesterol, a pegylated lipid, and the lipid-charged molecule conjugate. After the lipid nanoparticle of the present invention is administered by means of atomization inhalation, the anti-atomization stability of the lipid nanoparticle and the nucleic acid delivery efficiency are improved. When the lipid nanoparticle was used to encapsulate mRNA vaccine, inhaled mRNA-LNP can activate humoral, cellular, and mucosal immune responses.

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

Estrategia de búsqueda: vaccine.ti. AND @PD>="20250817"<=20250831 18 records

Document ID	Title	Inventor	Applicant Name
US 20250270656 A1	METHODS AND SYSTEMS FOR DEVELOPING PERSONALIZED VACCINE BY IDENTIFICATION AND PRIORITIZATION OF MUTATION-DERIVED NEOANTIGENS	Victor et al	Personal Genome Diagnostics Inc.
US 20250269009 A1	VACCINE COMPOSITION FOR NOVEL CORONAVIRUS INFECTION,HOU	Baidong	"RONGSEN BIOTECHNOLOGY (BEIJING) CO., LTD,INSTITUTE OF BIOPHYSICS, CHINESE ACADEMY OF SCIENCES"

US 20250257101 A1	Filovirus Consensus Antigens, Nucleic Acid Constructs and Vaccines Made Therefrom, and Methods of Using Same	Weiner; David B. et al.	The Trustees of the University of Pennsylvania, THE WISTAR INSTITUTE OF ANATOMY AND BIOLOGY, Inovio Pharmaceuticals, Inc.
US 20250259721 A1	DIGITAL VACCINE SYSTEM, METHOD AND DEVICE	Kaput; James et al.	VYDIANT, INC
US 20250255949 A1	STREPTOCOCCUS SUI VACCINE COMPOSITION COMPRISING IMMUNOGENIC FUSION POLYPEPTIDES	Frosth; Sara et al.	Intervacc AB
US 20250255946 A1	LOW-DOSE NEOANTIGEN VACCINE THERAPY	Jooss; Karin et al.	Gritstone bio, Inc.
US 20250255954 A1	STABILIZED SPIKE PROTEIN AND METHOD OF USE THEREOF AS A CORONAVIRUS DISEASE 2019 (COVID-19) VACCINE	Wu; Yuanhan et al.	Wu; Yuanhan, Walker; Susanne, Kulp; Daniel, The Wistar Institute of Anatomy and Biology
US 20250255944 A1	IMMUNOACTIVATOR, VACCINE ADJUVANT, AND METHOD FOR INDUCING IMMUNITY	SUZUKI; Ryo et al.	TEIKYO UNIVERSITY
US 20250255948 A1	PNEUMOCOCCAL CONJUGATE VACCINES AND METHODS OF USE THEREOF	Buchwald; Ulrike K. et al.	Merck Sharp & Dohme LLC
US 20250255952 A1	SELF-AMPLIFYING mRNA VACCINE BASED ON Z7 GENOME	Bai; Fengwei et al.	Bai; Fengwei, Huang; Faging
US 12383612 B2	Vaccine to protect a pig against Actinobacillus pleuropneumoniae	Witvliet; Maarten Hendrik et al.	Intervet Inc.

US 12383609 B2	Plasmodium sporozoite NPDP peptides as vaccine and target novel malaria vaccines and antibodies binding to	Lanzavecchia; Antonio et al.	INSTITUTE FOR RESEARCH IN BIOMEDICINE, SEATTLE CHILDREN'S HOSPITAL, SCHWEIZERISCHES TROPEN—UND PUBLIC HEALTH-INSTITUT
US 12383610 B2	Vaccine for protection against Streptococcus suis serotype 9, sequence type 16	Jacobs; Antonius Arnoldus Christiaan	Intervet Inc.
US 12383616 B2	Recombinant herpes zoster vaccine composition and application thereof	Shi; Li et al.	IMMUNE-PATH BIOTECHNOLOGY (SUZHOU) CO., LTD.
US 20250249090 A1	NEW FELINE HERPES VIRUS VACCINE	Vissek; Callie Ann et al.	Boehringer Ingelheim Vetmedica GmbH
US 20250249088 A1	THERAPEUTIC AND VACCINE CANDIDATES AGAINST SARS-CoV-2	MUTHUMANI; Kar	GENEONE LIFE SCIENCE, INC.
US 20250249087 A1	CONSTRUCTION OF NOVEL HUMAN INFLUENZA VIRUS VACCINE AND APPLICATION	TAN; Yeping et al.	Wuhan JinYiTai Biological Co., Ltd.
US 12377140 B2	Multivalent pneumococcal vaccines	Besin; Gilles R. et al.	Affinivax, Inc.

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