



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.

- 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.

Noticias en la Web

Pasteur Institute, Nagasaki University step up disease prevention collaboration

Aug 1. The Pasteur Institute in Ho Chi Minh City and Nagasaki University of Japan, both leading facilities in tropical medicine and infectious diseases in their respective countries, have signed a memorandum of understanding (MoU) to strengthen cooperation in key areas such as epidemiological surveillance capacity building, vaccine development, and response to both existing and emerging diseases.



Associate Professor Dr. Nguyen Vu Trung, Director of the Pasteur Institute, said that under the MoU, the two sides committed to promoting diverse and practical collaborative activities. These include joint conferences and workshops, exchanges of scholars, researchers and students, sharing academic materials, and the development of specific programmes and projects in scientific research, training, and epidemic prevention.

Witnessing the signing ceremony, Nguyen Ngo Quang, Director of the Administration of Science, Technology and Training under the Ministry of Health, suggested that the institute quickly draft a first-year action plan focused on readily implementable items such as short-term expert exchanges, bilateral workshops, and joint research group connections. Additionally, it should proactively seek and mobilise resources from international research support programmes and bilateral or multilateral funding organisations to ensure the financial sustainability of the cooperation.

Masuo Ono, Consul General of Japan in the southern metropolis, emphasised the importance of preventive medicine in preventing outbreaks and the transmission of infectious diseases. He highlighted its role in protecting lives, public health, and ensuring societal stability and prosperity. He expressed his hope that the agreement would strengthen efforts to control infectious diseases in Vietnam and around the world.

Notably, the two institutions have engaged in research and training cooperation since 2004.

Fuente: Vietnam Plus. Disponible en <https://n9.cl/ifsx65>

70% enrolment of participants for Phase III clinical trial of dengue vaccine completed: Centre

Aug 1. India achieves a milestone in dengue prevention. The DengiAll vaccine trial sees 70% participant enrollment. ICMR leads the multi-center study across 20 Indian sites. The budget is approximately Rs 1.3 to 1.5 crore per site. Dengue cases are rising, with over 5 lakh confirmed in 2024. The government monitors outbreaks and provides support to states.

"The trial involves over 10,000 participants, out of which more than 70 per cent enrolment has been completed. The trial has been implemented in 20 sites across India."
Prataprao Jadhav

Seventy percent enrolment of participants for the Phase III clinical trial of the indigenous one-shot dengue vaccine, DengiAll, has been completed, Minister of State for Health, Prataprao Jadhav, told Lok Sabha on Friday.

The Indian Council of Medical Research (ICMR) has undertaken a clinical trial titled phase III, multicentre, randomised, double-blind, placebo controlled study to evaluate the efficacy, immunogenicity and safety of the single-dose Dengue Tetravalent vaccine, live attenuated (Recombinant, Lyophilised) - 'DengiAll', Jadhav said in a written reply.

The sites are JSS Medical College and Hospital, Mysuru (Karnataka), Bangalore Medical College and Research Institute, and All India Institute of Medical Sciences, Bibinagar (Telangana).

The approximate budget for the trial is Rs 1.3 to 1.5 crore per site, Jadhav said.

According to ICMR, there is no specific antiviral treatment or licensed vaccines for dengue and the treatment is supportive in nature, Jadhav said.

A total of 5,73,563 laboratory confirmed dengue cases were reported through the Integrated Health Information Platform (IHIP) in 2024, according to information shared by the National Centre for Disease Control (NCDC).

According to ICMR, all four serotypes of the dengue virus (DENV-1, DENV-2, DENV-3, and DENV -4) are known to circulate and co-circulate in India.

Multiple serotypes can be present in the same geographical region and can even infect the same individual at the same time.

Hence, participants have been enrolled in the ICMR vaccine trial from different zones where these four serotypes circulate in order to test the efficacy of the vaccine against all four serotypes, Jadhav said.

The minister also said that the Union Health Ministry, Director General of Health Services, and National Centre for Vector Borne Disease Control (NCVBDC) regularly monitor the situation of dengue outbreaks across the country for assessing the disease situation, preparedness, technical guidance and to sensitise and forewarn the states.

Under the National Health Mission, adequate budgetary support is provided to states and Union territories for dengue control activities such as epidemic preparedness, monitoring, case management, vector control (provision of domestic breeding checkers, ASHA involvement, insecticide, fogging machines), training, intersectoral convergence, awareness activities, etc.

For surveillance and free of cost diagnosis of dengue, Sentinel Surveillance Hospitals with laboratory facility and Apex Referral laboratories with advance diagnostic facilities have been identified across the country, Jadhav said.

The government of India has developed national guidelines on dengue treatment/management, in consultation with experts, which have been shared with all the states and UTs for implementation.

The guidelines emphasise the preparedness of hospitals for case treatment/management.

In addition, advisories have been issued from time to time to states/UTs for prevention and control of dengue, including case management, the minister said.

Fuente: The Economic Times. Disponible en <https://n9.cl/ldgvbq>



RSV vaccine 82 per cent effective in preventing hospitalisation in older people

Aug 1. A study on the effectiveness of RSV vaccines published yesterday by the UK Health Security Agency (UKHSA) claims they are 82 per cent effective in preventing people aged 75 to 79 from being admitted to hospital with infection.

The study, conducted alongside Nottingham University Hospitals and other NHS trusts, also revealed RSV vaccines are “highly effective” in preventing older people with a chronic respiratory condition and those living with immunosuppression from hospitalisation.



In April, NHS England said it would increase the number of pharmacies taking part in its respiratory syncytial virus vaccines pilot from 50 to as many as 200 in 2025-26.

Twenty-five pharmacies in Mid and South Essex and North-East Essex integrated care boards were commissioned to provide the vaccine in August last year, shortly before two RSV vaccination schemes, one for older adults and a maternal programme, were introduced to the NHS vaccination schedule.

Under the scheme for older adults, people turning 75 can get vaccinated and there is a “one-off catch-up campaign” for adults aged 75 to 79.

The maternal vaccination programme is offered to women from 28 weeks of pregnancy to protect newborns at higher risk of severe illness from RSV.

Another study, *Vaccination in Pregnancy and RSV Hospitalisation in Infants in the UK*, published in *The Lancet Child and Adolescent Health*, found the maternal RSV vaccine was 72 per cent effective in preventing infants needing hospital treatment when their mothers were vaccinated over 14 days before delivery.

According to the UKHSA, the older adults programme reached 62.9 per cent of people as of 30 June 2025, an increase on 60.3 per cent in March, while 20,051 of 36,657 women who gave birth in March this year received an RSV vaccine.

The UKHSA said the highest vaccine maternal coverage was among the Chinese ethnic group at 73.3 per cent while the lowest was among black and black British Caribbean women at 26.4 per cent.

Fuente: Pharmacy Magazine. Disponible en <https://n9.cl/0aita0>

mRNA Technology Shows Early Progress in Developing HIV Vaccines Targeting Diverse Virus Variants

Aug 1. Recent studies have demonstrated early progress in the development of mRNA-based HIV vaccines, addressing longstanding challenges in targeting the virus. Researchers have focused on eliciting neutralizing antibodies capable of binding to and blocking diverse variants of HIV, a critical hurdle that has historically slowed vaccine development efforts.

Neutralizing antibodies are proteins produced by the immune system that play a key role in defending against pathogens such as viruses and bacteria.

The findings suggest that mRNA technology, which has gained prominence in recent years for its role in COVID-19 vaccines, may offer a promising approach to overcoming the complexities associated with HIV's variability. Scientists involved in these studies aim to refine this strategy further to improve its effectiveness across the wide range of HIV strains.

Fuente: GeneOnline. Disponible en <https://n9.cl/f637k>



Pfizer's Phase 3 Study: Evaluating Co-administration of RSV and Shingles Vaccines

Aug 2. Pfizer Inc. is conducting a Phase 3 study titled 'A Study to Evaluate the Safety and Immunogenicity of RSVpreF Coadministered With Herpes Zoster Vaccine in Adults.' This study aims to assess the safety, tolerability, and immune response of a Respiratory Syncytial Virus (RSV) prefusion F subunit vaccine when administered alongside a herpes zoster subunit vaccine in adults aged 50 and above. The significance of this study lies in its potential to enhance vaccine efficacy and convenience for older adults.

The study is testing two biological interventions: the RSV prefusion F subunit vaccine and the herpes zoster subunit (HZ/su) vaccine. Both are administered via intramuscular injection and are intended to prevent RSV and shingles, respectively.

This interventional study follows a randomized, parallel-group design with no masking, focusing primarily on prevention. Participants are divided into two groups: one receiving both vaccines simultaneously and the other receiving them sequentially.

The study began on April 4, 2025, with primary completion expected shortly thereafter. The latest update was submitted on July 30, 2025. These dates are crucial for tracking the study's progress and anticipating results.

Pfizer's ongoing study could influence its stock performance positively by showcasing its commitment to advancing adult vaccination. This could also impact investor sentiment, especially as the pharmaceutical industry remains competitive with ongoing innovations in vaccine development.

The study is currently active but not recruiting, with further details available on the ClinicalTrials portal.

Fuente: THE GLOBE AND MAIL. Disponible en <https://n9.cl/39n19>

Estudio global revela cuántas vidas salvaron las vacunas contra la COVID-19

2 ago. Las vacunas contra COVID-19 evitaron 2,5 millones de muertes, según un nuevo estudio de modelización global.

Entre diciembre de 2020 y finales de 2024 se administraron alrededor de 13.600 millones de dosis de vacunas contra COVID-19 en todo el mundo.

Es decir, por 5.400 dosis de vacunas administradas, se evitó una muerte, precisó en un artículo el doctor John Ioannidis, profesor de medicina y

epidemiología de la Universidad de Stanford, y sus coautores. Se tomaron estimaciones de la tasa de mortalidad por COVID-19 y la eficacia de las vacunas contra la mortalidad y las introdujeron en modelos para tratar de averiguar cuántas muertes se evitaron gracias a la vacunación.

Los datos oficiales sobre el número de personas fallecidas a causa de la COVID-19 ascienden a más de siete millones de personas en el mundo (7.083.769 en informe de la OMS del 5 de enero de 2025), aunque la ONU habla de una cifra varias veces superior: al menos 20 millones.

La Organización Mundial de la Salud (OMS), lideró el extenso estudio que demuestra lo que por años han defendido los expertos: la vacunación es uno de los mayores avances en salud pública. Según el informe, entre 1974 y 2024, las vacunas salvaron al menos 154 millones de vidas en el mundo.

Esto equivale a seis vidas salvadas por minuto durante cinco décadas, una cifra que subraya el poder transformador de la inmunización, especialmente en los países en desarrollo.

Los más beneficiados con vacunas: niños menores de cinco años

Del total de vidas salvadas, la mayoría correspondió a niños menores de cinco años, el grupo más vulnerable a enfermedades prevenibles como el sarampión, la difteria o la tosferina.

El estudio destaca que, sin acceso a vacunas, millones de familias en todo el mundo habrían enfrentado tragedias evitables. La inmunización ha sido especialmente crítica en regiones de África, Asia y América Latina.

El sarampión, la vacuna con mayor impacto

Entre todas las vacunas, la que tuvo el mayor impacto fue la del sarampión, responsable de alrededor del 60% de las vidas salvadas, según el reporte. Esta enfermedad altamente contagiosa ha sido controlada en gran parte del mundo gracias a programas masivos de inmunización infantil.

"Las cifras hablan por sí solas. Vacunar no solo protege al individuo, sino que salva comunidades enteras", indicó un vocero de la OMS.

Riesgos sí existen, pero los beneficios superan con creces

El informe también reconoce que, como cualquier medicamento, las vacunas pueden tener efectos secundarios. Sin embargo, recalca que los beneficios superan ampliamente los riesgos, y que los sistemas de salud deben seguir fomentando la vacunación segura y equitativa.

Los expertos insisten en que los casos graves por reacciones adversas son extremadamente raros, y que evitar las vacunas puede poner en riesgo tanto al individuo como a su entorno.

Un llamado global a mantener y ampliar la cobertura en vacunación: OMS

La OMS hizo un llamado a mantener e incluso ampliar la cobertura de vacunación, especialmente en un

"Estudio internacional revela el verdadero impacto de las vacunas contra la COVID-19 y expone cifras sorprendentes sobre vidas salvadas en todo el mundo."

contexto en el que los movimientos antivacunas y la desinformación amenazan con revertir décadas de progreso.

“Invertir en vacunas es invertir en vida. Es la herramienta más efectiva, equitativa y económica para proteger la salud global”, concluyó el estudio.

La OMS defiende el tratado antipandemias

Entre tanto, el tratado global recientemente alcanzado para luchar contra pandemias y las reformas de las regulaciones sanitarias internacionales acordados en 2024 se aprobaron por consenso de los miembros de la Organización Mundial de la Salud (OMS) y no quitan soberanía a los Estados, defendió el director general de la agencia, Tedros Adhanom Ghebreyesus, en respuesta a recientes críticas desde EE.UU. e Italia.

“La OMS no tiene autoridad para decir a los países lo que deben hacer, no podemos imponerles limitaciones a los viajes, confinamientos, vacunaciones o cualquier otra medida”, aclaró el director general en rueda de prensa el pasado miércoles.

Fuente: Vanguardia. Disponible en <https://n9.cl/5jqtg>



Stanford Medicine: laboratorios con “científicos virtuales” diseñan soluciones biomédicas en días

Aug 2. Un equipo de investigadores de Stanford Medicine creó “científicos virtuales” que colaboran en laboratorios digitales para resolver desafíos biológicos, logrando en pocos días avances como el diseño de candidatos innovadores para vacunas contra variantes recientes de COVID-19. Anticipo del futuro en el que la IA acelera y multiplica los descubrimientos científicos.

La colaboración interdisciplinaria está en el corazón del progreso científico, pero organizar equipos de expertos, coordinar ideas y avanzar a gran escala sigue siendo un enorme reto. Ahora, investigadores de Stanford Medicine liderados por James Zou, PhD, han puesto a prueba una solución disruptiva: laboratorios virtuales impulsados por inteligencia artificial, capaces de simular la dinámica y el ingenio de los mejores equipos humanos, pero trabajando sin descanso ya una velocidad inalcanzable para las personas.

“Las grandes innovaciones ocurren cuando personas de diferentes disciplinas colaboran, pero forman esos equipos y consiguen sinergia es uno de los mayores cuellos de botella de la investigación”, explicó Zou, quien se dirigió al estudio publicado el 29 de julio en *Nature*.

“Hemos visto enormes avances en los agentes de IA capaces de actuar proactivamente, colaborar entre sí y con humanos en lenguaje natural. Eso nos animó a entrenar estos modelos para pensar y debatir como científicos top, buscando soluciones novedosas a grandes problemas.”

¿Cómo funciona el laboratorio virtual?

El laboratorio virtual de Stanford se modela como un auténtico grupo de investigación, con un investigador principal de IA (IA PI), que recibe un reto científico y decide qué otros “agentes científicos” de IA sumarán, asignando roles como inmunología, biología computacional, machine learning y un crítico encargado de identificar fallas o posibles sesgos.

A cada paso, los agentes pueden hacer uso de herramientas especializadas —como AlphaFold para modelado de proteínas— y hasta solicitar acceso a nuevos softwares, que los investigadores humanos incorporan de acuerdo a las necesidades del equipo virtual. Las reuniones, en vez de durar horas, se resuelven en segundos; Además, el grupo realiza decenas de sesiones simultáneas. Sin microgestión, la intervención humana es mínima (alrededor de 1%), oscilando entre límites presupuestarios y validaciones iniciales del desafío. Todo el proceso queda automáticamente registrado, permitiendo a los investigadores monitorear el avance y redireccionar el trabajo cuando sea necesario.



Resultados: IA resuelve en días lo que antes tomaba meses

Para probar la eficacia del laboratorio virtual, el equipo de Stanford encargó el diseño de una estrategia para una vacuna contra las variantes recientes del SARS-CoV-2. Mientras la solución habitual habría sido anticuerpos tradicionales, el equipo de IA se sorprendió proponiendo nanocuerpos: fragmentos más pequeños, simples de modelar y potencialmente más fáciles de producir a escala.

“Desde el inicio, los científicos virtuales decidieron que los nanocuerpos eran más prometedores que los anticuerpos. Argumentaron que, al ser más pequeños, aumentaban la confianza en el modelado y diseño computacional de proteínas”, explicó Zou. El equipo diseñó 92 nuevos nanocuerpos; Tras su síntesis en el laboratorio real del coautor John Pak en Chan Zuckerberg Biohub, comprobaron que más del 90% eran funcionales y solubles. Dos de ellos sobresalieron al unirse con alta afinidad a cepas recientes del virus y también a la variante de Wuhan, clave para una vacuna de amplio espectro.

“Lo que era una idea de ciencia ficción ahora es real: los agentes de IA propusieron una estrategia creativa, razonable y muy rápida. Y los experimentos validaron su efectividad”, resaltó Pak.

Más allá de la COVID-19: ciencia acelerada, nuevas preguntas

La versatilidad del laboratorio virtual es enorme: tras el éxito con el SARS-CoV-2, los investigadores planean explorar otros desafíos biomédicos y crear agentes especializados en análisis de datos complejos, capaces de reevaluar investigaciones previas y formular nuevas hipótesis.

“Muchas veces la IA propone hallazgos más allá de lo publicado por humanos. Esto es tremendamente talentoso para toda la comunidad científica”, afirmó Zou.

Para Pak, lejos de reemplazar a los investigadores, la IA potenciará su productividad: “La IA nos dio más trabajo, porque sugirió más hipótesis para evaluar. Si puede proponer ideas testables, todos ganamos”.

El futuro de la investigación, hoy

El estudio, financiado por becas de la Knight-Hennessy y Stanford Bio-X, ilustra cómo los laboratorios virtuales —combinando IA generativa, modelado estructural y participación humana— pueden multiplicar la capacidad y velocidad de la ciencia moderna. Los responsables de la investigación prevén que esta metodología se expanda en áreas como el desarrollo de fármacos, medicina personalizada y análisis de nuevas enfermedades.

Los datos, los entrecomillados y los resultados experimentales ya hablan por sí mismos: la colaboración entre humanos e “investigadores virtuales” puede marcar un antes y un después en la carrera por los grandes descubrimientos de la biomedicina.

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

Seven more strains covered

Aug 3. Malaysians can now be protected against 20 strains of the bacteria *Streptococcus pneumoniae*, which causes pneumonia.

This follows the recent announcement that the Health Ministry has approved the 20-valent pneumococcal conjugate vaccine (PCV20) by American pharmaceutical company Pfizer.

“As pneumococcal bacterial strains continue to evolve, scientists race to update vaccines for more protection.”

This new vaccine offers protection against an additional seven serotypes of *S. pneumoniae*, also known as pneumococcus, compared to its 13valent predecessor (PCV13).

These include newer strains associated with antibiotic resistance and high death rates.

Professor Dr Mark van der Linden, who heads the national reference centre for Streptococci in Germany, notes that scientists have been racing to keep up with the evolving strains of pneumococcus.

“There are currently 107 known serotypes.

“We’re not creating new ones; they evolve naturally.

“Pneumococcus reproduces every 30 minutes, so it adapts far faster than humans,” he shares.

The first conjugate vaccine, which was developed specifically for use in young children, targeted the seven serotypes that caused the majority of pneumococcal disease.

This 7-valent pneumococcal conjugate vaccine (PCV7) was introduced in Malaysia over two decades ago.

As vaccination reduced these strains, others emerged – a process called serotype replacement.

“We saw a 40% drop in cases, but new strains emerged.

“That’s why vaccines must continuously evolve.

“We can’t cover all 107, but expanding coverage is essential, especially for high-risk groups,” says Prof van der Linden.

Meanwhile, consultant clinical microbiology and infection physician Dr Teresa Wang Kin Fong from Hong Kong explains that based on data from the Chinese territory, invasive pneumococcal disease is now more common among adults than children.

“We used to give adults the polysaccharide vaccine PPV23, but it had limited effectiveness.

“Pfizer later developed PCV7, and at one point, we [in Hong Kong] implemented a combined adult programme using both PPV23 and PCV7.

“Now with PCV20, we can skip PPV23 entirely.

“PCV20 offers broader protection and simplifies the process to a single dose, making it more convenient for patients and doctors alike.”

In Malaysia, vaccine uptake among adults remains low.

“This is largely due to the outdated perception that vaccines are only for children,” notes Malaysian Society of Infectious Diseases and Chemotherapy past president Prof Dr Zamberi Sekawi.

Adults with chronic conditions like diabetes, high blood pressure and lung disease, are especially vulnerable

to developing severe pneumonia, which can be life-threatening. Vaccinating adults not only protects individuals, but also reduces transmission within households.

“When one family member brings home an infection, the whole household is at risk.

“Vaccinating adults helps break that chain,” Prof Zamberi adds.

This is important as both young children and the elderly are also vulnerable to developing severe pneumonia.

“Children are born with immature immune systems.

“As we age, our immune system weakens – a process called immunosenescence.

“That’s why both groups need the most protection,” says consultant paediatrician Prof Datuk Dr Zulkifli Ismail.

Since the Covid-19 pandemic, parental concerns about vaccination have shifted from cultural or religious questions to fear of side effects.

“I’ve had parents who declined the pneumococcal vaccine, only to return later after their child was hospitalised.

“Sadly, by then it’s too late,” shares Prof Zulkifli.

PCV20 is approved for use in both children and adults starting from the age of six weeks onwards.



Fuente: Press Reader. Disponible en <https://n9.cl/x9seh>

Korea launches free 20-valent pneumococcal vaccine for children in October

Aug 4. The Korea Disease Control and Prevention Agency announced on 4th that it plans to implement free vaccinations for the 20-valent pneumococcal conjugate vaccine (PCV20) starting from Oct. 1.

The target group includes all infants aged 2 months to under 12 years and high-risk adolescents under 18 years; this is part of the national immunization program (NIP).

Pneumococcus is a major bacterial pathogen that causes various diseases such as otitis media, pneumonia, and meningitis in infants. It can cause invasive infections (Invasive Pneumococcal Disease, IPD) that threaten the lives of immunocompromised children, making vaccination very important.

So far, the national immunization program has supported vaccinations for children with the 13-valent conjugate vaccine (PCV13) and the 15-valent conjugate vaccine (PCV15).

The newly introduced PCV20 received approval from the Ministry of Food and Drug Safety last October. It includes an additional 5 serotypes (8, 10A, 11A, 12F, 15B) on top of the 15 serotypes included in the 15-valent vaccine (PCV15), allowing for protection against a total of 20 serotypes.

For healthy children, the vaccination schedule remains the same: a total of 3 vaccinations at 2, 4, and 6 months, followed by an additional vaccination at 12 to 15 months. Children who have already started vaccination with PCV13 can receive cross vaccination with PCV20. However, if vaccination was started with PCV15, it is recommended to complete the vaccination with the same vaccine.

High-risk children and adolescents who are vulnerable to infections due to immunodeficiency, chronic diseases, or cochlear implants are also eligible for PCV20 vaccination. For high-risk children and adolescents,

the vaccination schedule varies according to their age and previous vaccination history, so they should follow a vaccination schedule that fits their situation. The upper age limit for high-risk children eligible for PCV20 support has been raised from the previous 12 years to 18 years.

Fuente: Chosun Biz. Disponible en <https://n9.cl/48c3mv>

Co-Administering Dengue and HPV Vaccines Found Compatible

Aug 4. According to the World Health Organization (WHO), Dengue is a viral infection spread to humans by infected mosquitoes and is a leading cause of febrile illness among international travelers.

Based on the record-setting number of Dengue outbreaks in 2024, the WHO has classified Dengue as a grade 3 emergency, placing about four billion people at risk globally.

"Takeda second generation dengue vaccine Qdenga approved in various countries."

In recent years, the development of effective vaccines to prevent Dengue has been an essential factor in the global effort to reduce outbreaks.

Still, strategies to maximize second-generation vaccine coverage in endemic regions are needed.

According to a new study published in the journal *Vaccines* (Volume 62, August 30, 2025, 127558), dengue-endemic countries, such as the Kingdom of Thailand, have already initiated national Human Papillomavirus (HPV) school-based vaccination programs based on recommendations from the WHO.

These researchers wrote, 'Due to the overlapping age range targeted by many vaccines, including 9vHPV (Merck GARDASIL 9®) and TAK-003 (Takeda's QDENGAR®), integration of Dengue vaccination into these existing programs could be a beneficial approach to increase vaccine coverage and reduce operational costs.'

This phase 3, multicenter clinical trial (NCT04313244) demonstrated the non-inferiority (NI) of the immune response to 9vHPV when co-administered with the live-attenuated dengue vaccine, TAK-003, compared with 9vHPV administered alone, with the primary objective was met.

There was no evidence of interaction caused by the live-attenuated vaccine on the virus-like particle-based 9vHPV immunogenicity and safety.

Similarly, these data suggest that co-administration of TAK-003 with HPV does not compromise TAK-003 immunogenicity or safety.

In the co-administration group, an immune response was elicited against each of the four dengue serotypes, with GMTs of anti-dengue NAb increasing to 494–1691 at M4 post-vaccination.

Participants who had a previous exposure to Dengue (Dengue seropositive at baseline) showed higher post-vaccination GMTs, which can be explained by the presence of immune memory from their prior dengue infection.

As observed in previous studies with TAK-003, almost all participants (99.6%) had tetravalent seropositivity at M4, irrespective of baseline serostatus.

The co-administration of 9vHPV and TAK-003 was well-tolerated, and there was no clinically relevant adverse impact on the safety or reactogenicity, with no AEs leading to vaccine withdrawal or trial discontinuation.

A limitation of this trial is that a TAK-003-only arm was not included, meaning the NI of the TAK-003 immune response could not be directly evaluated after co-administration. One of the reasons behind this design was that the trial was only considered feasible in dengue-endemic regions.

As of August 4, 2025, Thailand's Food and Drug Administration has approved the QDENGAR vaccine and over

30 other countries, but not the United States. Currently, the U.S. is evaluating the first-generation Dengue vaccine (Dengvaxia) in children in Puerto Rico.

Note: This phase 3 clinical trial was sponsored by Takeda Vaccines, Inc., Cambridge, MA, USA, the producer of the QDENG A vaccine.

Fuente: VAX-BEFORE-TRAVEL. Disponible en <https://n9.cl/yI0d3>

China makes significant progress in HIV vaccine development

Aug 4. China's National Center for AIDS/STD Control and Prevention, along with other research teams, has recently completed the Phase I clinical trial of China's first replication-competent Tiantan vaccinia-based AIDS vaccine, marking a major breakthrough in the country's HIV vaccine development efforts, Beijing Daily reported on Monday.

The study innovatively used the Tiantan strain of the vaccinia virus, which was widely used as a smallpox vaccine in China. The trial confirmed the vaccine's safety and its ability to elicit effective immune responses against HIV, according to the report.



AIDS remains one of the major public health challenges facing humanity. Despite progress in antiretroviral therapy and pre-exposure prophylaxis, there are still over a million new HIV infections globally each year.

The research team noted that the high variability of the HIV virus and its ability to attack the immune system pose significant challenges for vaccine development. Globally, over 300 HIV vaccine trials have been conducted, including 11 efficacy trials, most of which have failed to demonstrate protective immune effects.

Compared to replication-defective vectors that provide only transient antigen exposure, replication-competent viral vectors are considered a promising alternative for HIV antigen delivery. Replication-competent vectors, such as adenovirus, vaccinia virus, and cytomegalovirus, have been studied in preclinical and clinical trials.

Against this backdrop, teams from the AIDS Prevention and Control Center under the Chinese CDC, Peking Union Medical College Hospital, and other institutions used the vaccinia virus Tiantan, previously used to eradicate smallpox, as a vector to construct a recombinant vaccine. So far, they have completed a Phase I clinical trial evaluating safety and immunogenicity in 48 healthy participants, Beijing Daily reported.

The study results confirmed the safety of the replication-competent Tiantan vaccinia-based AIDS vaccine in healthy individuals and demonstrated its ability to induce sustained HIV-specific immune responses through this enhanced immunization strategy, laying the foundation for Phase II clinical trials.

Fuente: Global Times. Disponible en <https://n9.cl/kd6ih2>

Departamento de Salud de EEUU elimina \$500 millones para desarrollo de vacunas

5 ago. El Departamento de Salud y Servicios Humanos de Estados Unidos (HHS, por sus siglas en inglés) cancelará contratos y retirará fondos para algunas vacunas en desarrollo contra virus respiratorios como la COVID-19 y la gripe.

El secretario de Salud, Robert F. Kennedy Jr., anunció en un comunicado que se detendrán 22 proyectos, por un valor de \$500 millones de dólares, para desarrollar vacunas con tecnología de ARNm.

Es la última de una serie de decisiones que han cristalizado las dudas de RFK Jr. sobre las vacunas en el departamento de salud del país. Kennedy ha retirado recomendaciones sobre las vacunas contra la COVID-19, despidió al panel que hace recomendaciones de vacunas y se negó a ofrecer un respaldo energético a las vacunas mientras empeoraba un reciente brote de sarampión en el país.

El secretario de Salud criticó las vacunas de ARNm en un video en sus cuentas de redes sociales en el que explica la decisión de cancelar proyectos liderados por las principales compañías farmacéuticas del país, incluidas Pfizer y Moderna, que ofrecen protección contra virus como la gripe, COVID-19 y H5N1.



Health and Human Services Secretary Robert F. Kennedy Jr. speaks during an event in the East Room of the White House, Wednesday, July 30, 2025, in Washington.
AP Photo/Mark Schiefelbein

"Para reemplazar los problemáticos programas de ARNm, estamos priorizando el desarrollo de estrategias de vacunas más seguras y amplias, como las vacunas de virus completo y plataformas novedosas que no colapsan cuando los virus mutan", dice Kennedy en el video.

Expertos en enfermedades infecciosas señalan que la tecnología de ARNm utilizada en las vacunas es segura, y atribuyen la ralentización de la pandemia de coronavirus de 2020 al desarrollo de dichas vacunas durante el primer mandato de Donald Trump. Advirtieron que futuras pandemias serán más difíciles de detener sin la ayuda del ARNm.

"No creo haber visto una decisión de salud pública más peligrosa en mis 50 años en el negocio", dijo Mike Osterholm, experto en enfermedades infecciosas y preparación para pandemias de la Universidad de Minnesota.

Señaló que la tecnología de ARNm ofrece ventajas potenciales como su rápida producción, que es crucial en caso de otra pandemia que requiera una nueva vacuna.

El abandono de los proyectos de ARNm es miope en el contexto de que aún hay preocupaciones sobre una pandemia de gripe aviar, dijo el doctor Paul Offit, experto en vacunas del Hospital Infantil de Filadelfia.

"Ciertamente ha salvado millones de vidas", dijo Offit sobre las vacunas de ARNm existentes.

Los científicos están utilizando el ARNm para más que vacunas contra enfermedades infecciosas, pues investigadores de todo el mundo exploran su uso para inmunoterapias contra el cáncer. Este año en la Casa Blanca, el multimillonario empresario tecnológico Larry Ellison elogió el ARNm por su potencial para tratar el cáncer.

Tradicionalmente, las vacunas han requerido cultivar fragmentos de virus, a menudo en huevos de gallina o grandes cubas de células, y luego purificar ese material. La estrategia de vacunas ARNm comienza con un fragmento de código genético que lleva instrucciones para fabricar proteínas. Los científicos eligen la proteína a atacar, inyectan esa instrucción y el cuerpo produce la suficiente para desencadenar la protección inmunitaria, produciendo su propia dosis de vacuna.

En un comunicado el martes, el HHS dijo que "otros usos de la tecnología de ARNm dentro del departamento no se ven afectados por este anuncio".

La tecnología de ARNm se utiliza en las vacunas aprobadas contra la COVID-19 y el VRS, pero aún no ha

sido aprobada para una vacuna contra la gripe. Moderna, que estaba estudiando una combinación de vacuna de ARNm contra la COVID-19 y la gripe, había dicho que creía que el ARNm podría acelerar la producción de vacunas contra la gripe en comparación con las vacunas tradicionales.

Los proyectos de ARNm abandonados son indicio de un "cambio en las prioridades de desarrollo de vacunas", dijo el departamento de salud en su comunicado, agregando que comenzará a "invertir en mejores soluciones". No se proporcionaron detalles sobre cuáles podrían ser esas otras tecnologías.

"Permítanme ser absolutamente claro, el HHS apoya vacunas seguras y efectivas para cada estadounidense que las desee", dijo Kennedy en el comunicado.

Fuente: Eye Witness News. Disponible en <https://n9.cl/hi9zh>

Identifying high risk "pathogen X" with potential to spark future pandemics

Aug 5. Announcing a new article publication for [Zoonoses](#) journal. In past decades, numerous emerging infectious diseases have markedly affected global health and social-economic development.

Consequently, the concept of "disease X" has been developed and lists of the pathogens with potential to cause future pandemics have been proposed and updated several times by the World Health Organization. This article analyses the major features of pathogens with high probability to cause future COVID-19-like pandemics, on the basis of their transmission routes, animal-human interfaces, herd immunity barriers, and evolutionary and mutational characteristics. High-priority viral infectious diseases with potential to become disease X in the future, are also evaluated on the basis of their transmission pathways, including airborne/respiratory, vector-borne, and direct contact. Overall, viruses had much greater likelihood of becoming "pathogen X" than either bacteria or other microorganisms.

Respiratory viruses, particularly RNA viruses, have notably high potential to become pathogen X. Vector-borne viruses might also become pathogen X, and mosquito-borne viruses would be more likely than tick-borne viruses to do so. Many contagious viral infectious diseases lead to severe clinical outcomes but have relatively low likelihood of causing global pandemics. Coronaviruses in the *Coronaviridae* and influenza viruses in the *Orthomyxoviridae* were identified as top-priority pathogens X with potential to cause the next COVID pandemic.

Fuente: News Medical Life Sciences. Disponible en <https://n9.cl/6sx58z>

Leading science labs in Korea join world's largest vaccine testing network

Aug 5. Two leading scientific organisations based in Korea, the Korea Disease Control and Prevention Agency (KDCA) and International Vaccine Institute (IVI), are joining the world's largest network dedicated to the standardised testing of epidemic and pandemic vaccines.

- ◆ KDCA and IVI join global group of laboratories set up to standardise assessment of new vaccines.
- ◆ Expansion of network to Korea could help the country more quickly respond to future outbreaks.
- ◆ CEPI set up the centralised group of labs in 2020 to respond to the COVID-19 pandemic.
- ◆ KDCA and IVI already working with CEPI on pandemic preparedness and response efforts.

Set up by the Coalition for Epidemic Preparedness Innovations (CEPI), the network aims to accelerate the development of vaccines to respond more quickly to future outbreaks of viral threats in as little as 100 days.



This includes potentially deadly infections like coronaviruses, pandemic influenza, and a novel or as-yet-identified 'Disease X'.

KDCA and IVI are the first institutions headquartered in Korea to join the network now made up of twenty laboratories. They will work together and with scientific groups across Africa, the Americas, Asia, Australasia and Europe as a 'centralised laboratory', using the same tools and protocols to evaluate multiple vaccine candidates created from different developers as though they have been tested 'under one roof'. This can help overcome the current challenge of being unable to easily compare vaccine candidates as different measurements and techniques are used by different laboratories. Minimising variability in the vaccine data could help researchers and regulators based in Korea more rapidly select the best-performing vaccine candidates.

"When multiple vaccine candidates are undergoing testing, differences in how data is collected becomes an issue. As well as potential variations in markers of immunity, there can be distinctions in how and where samples are collected, transported and stored" explains Dr Kent Kester, Executive Director of Vaccine R&D at CEPI. "This impacts the quality and usefulness of the data produced. KDCA and IVI's new membership to our Centralised Laboratory Network will lessen these problems, to more reliably and quickly evaluate potentially life-saving vaccines under development."

Expanding CEPI's Centralised Laboratory Network's footprint to include sites in Korea could also help the region more quickly respond to an emerging local outbreak as time is not wasted shipping samples for testing to laboratories further afield.

"The inclusion of KDCA and IVI in CEPI's Centralised Laboratory Network will facilitate harmonized and expedited evaluation of vaccine candidates during epidemics and pandemics," says Dr. Manki Song, Deputy Director General of Science at IVI. "By joining this global network, KDCA and IVI can further increase their contributions to pandemic preparedness and global health security by enhancing our shared capabilities in clinical sample analysis and vaccine development."

"By joining CEPI's Centralised Laboratory Network together with IVI, KDCA expects to strengthen global partnerships in vaccine evaluation, enhancing preparedness for emerging infectious diseases and potential pandemic preparedness," says Dr. Seungkwan Lim, Commissioner of KDCA. "To strengthen global health security, we will continue to expand our R&D infrastructure for vaccine development, targeting CEPI priority pathogens and Disease X, while deepening global research collaboration."

As part of its broader policy priorities, the Korean government pledged during the June presidential election to expand joint research on infectious diseases, strengthen international cooperation in responding to public health crises, and enhance its infectious disease response systems.

Set up in 2020 in response to the COVID-19 pandemic, CEPI's Centralised Laboratory Network has made significant contributions to the development of several COVID-19 vaccines and vaccines being designed against other emerging infectious diseases. CEPI has provided up to US \$57 million to run the network.

To date, CEPI has allocated approximately US\$350 million allocated to Korean companies and institutions advancing research to prepare for future epidemic and pandemic threats. This includes existing partnerships with KDCA and IVI. Since joining CEPI in 2020, the South Korean government has contributed \$51 million to support CEPI's global efforts.

Fuente: CEPI. Disponible en <https://n9.cl/bb0ns>

La variante “Frankenstein” de la COVID-19 llega a Sudamérica: ¿En qué países hay casos confirmados y cuáles son los síntomas?

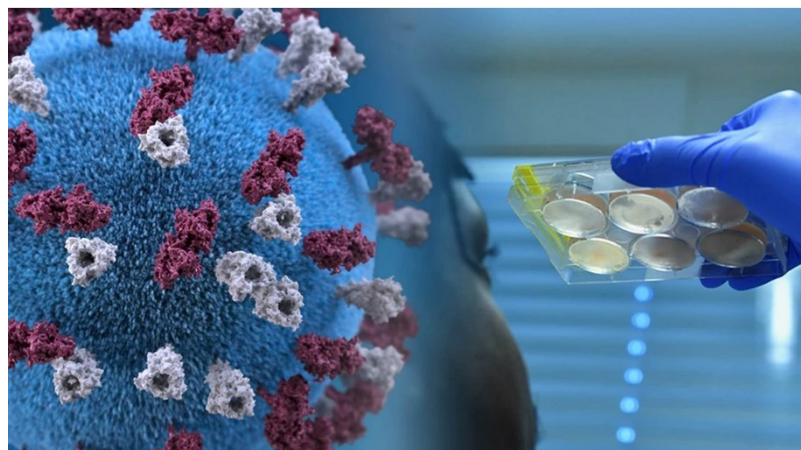
7 ago. La variante “Frankenstein” (XFG) de la COVID-19 ha sido detectada en países de Sudamérica como Argentina y Brasil, generando alerta por su rápida capacidad de mutación.

¿Qué es la variante “Frankenstein” y cuáles son sus síntomas?

La variante “Frankenstein” es una cepa recombinante del SARS-CoV-2. Esto significa que es una combinación de dos variantes preexistentes, la LF.7 y la LP.8.1.2. El nombre proviene de su naturaleza altamente mutada, similar a la descripción de la variante Ómicron en el pasado.

Los síntomas asociados con esta variante son los mismos que los de otras cepas de COVID-19. Estos pueden variar en gravedad y generalmente incluyen:

- ♦ Fiebre.
- ♦ Tos.
- ♦ Dificultad para respirar.



Aunque la Organización Mundial de la Salud (OMS) ha clasificado a la XFG como una “variante bajo monitoreo”, no hay evidencia que sugiera que causa una enfermedad más grave que otras variantes en circulación.

La variante XFG se ha detectado en varios países a nivel mundial. En América Latina, Brasil ha reportado un aumento significativo en la circulación de esta cepa. Específicamente, en Río de Janeiro, se identificaron 46 casos entre el 1 y el 8 de julio de 2025, lo que representó el 62% de los genomas analizados en ese periodo. También se han confirmado casos en otros estados brasileños como São Paulo, Ceará y Santa Catarina.

Por su parte, Argentina ha registrado tres casos de la variante XFG entre la semana epidemiológica 9 y 28 de 2025, según el Boletín Epidemiológico Nacional.

La OMS ha señalado un aumento general de la variante en varias regiones, incluyendo América Latina.

Fuente: TV Azteca Chiapas. Disponible en <https://n9.cl/7b6tl>

Enfermedad X: La OMS se prepara para próxima pandemia letal

7 ago. Mucho se ha escuchado sobre la “enfermedad x”, pero muchos desconocen qué es y cómo la OMS se está preparando para una posible pandemia desde hace años.

El temor ha invadido las redes ante la posibilidad de una próxima pandemia, ya que mucho se ha escuchado sobre la “Enfermedad X”, la cual ha incluido la Organización Mundial de la Salud (OMS) dentro de su lista de patógenos que deben investigar con urgencia, ya que podría ser letal y causar una grave epidemia internacional.

A pesar de su reciente popularización en redes, este término ya había adoptado por la OMS desde el 2017, fecha en que presentaron una lista de patógenos que podían provocar futuros brotes o pandemias, entre las cuales se encontraron “la enfermedad del virus del Ébola, la fiebre de Lassa, el síndrome respiratorio de

Oriente Medio (MERS), el síndrome respiratorio agudo severo (SARS), la fiebre del Valle del Rift, la COVID-19, el Zika y la “enfermedad X”.

La “enfermedad X” no se trata de una enfermedad como tal, sino que es un término que adoptó la Organización Mundial de la Salud para referirse a un patógeno desconocido que podría causar una grave epidemia internacional. De esta forma, se busca realizar investigaciones avanzadas que sirvan para estar preparados para futuras enfermedades no conocidas, intentando así tener una acción rápida en futuras pandemias.

“Es esencial centrarse en los patógenos y familias de virus prioritarios para que puedan ser investigados y desarrollar las contramedidas necesarias a fin de dar una respuesta rápida y eficaz a las epidemias y pandemias”, comentó Michael Ryan, directivo ejecutivo del Programa de Emergencias Sanitarias de la OMS.

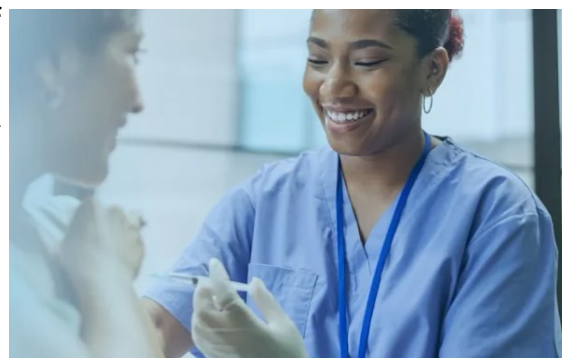
La pandemia por COVID-19, además de los estragos, dejó un serie de aprendizajes en la comunidad científica sobre cómo actuar ante futuras epidemias internacionales. No obstante, debido a que la COVID-19 ya había estado siendo estudiada con anticipación dentro de la lista de patógenos prioritarios, fue posible “desarrollar vacunas seguras y eficaces en tiempo récord”, por lo que este modelo seguirá siendo implementado.

De esta forma, cuando la “enfermedad X” se llegara a presentar, ya se contarán con avances y estudios que permitirán una pronta acción de las autoridades para garantizar la sanidad a los ciudadanos.

Fuente: TV Azteca Chiapas. Disponible en <https://n9.cl/1h06q>

Why long-term follow-up should start early in vaccine trials

Aug 8. In 1974, the World Health Organisation (WHO), at the request of its members, launched the Expanded Program on Immunisation (EPI). The aim was to make life-saving vaccines available to all globally. A study published in Lancet to mark the 50th anniversary of the EPI showed the impact of the initiative on public health. The study's authors found that the program averted 154 million deaths.¹ The EPI program is estimated to have been directly responsible for a 40% reduction in infant mortality during the same period. According to the study, as many as 146 million deaths were averted in children under five years old, including an estimated 101 million in infants under one year old.



Recent outbreaks of familiar vaccine-preventable illnesses such as measles (in the US and abroad³) and pertussis (in the EU⁴ and especially the Czech Republic⁵) have reignited debates about vaccines. Vaccination programs can be cost-effective and life-saving public health tools. A crucial factor in any vaccine program is the ability to demonstrate the vaccine's efficacy and safety. Demonstrating safety and efficacy ensures that the vaccine is effective for its intended use, and that the benefits outweigh the potential risks, safeguarding patient well-being. Vaccine programs only succeed when the public feel confident in the safety of the vaccines in question. Similarly, vaccine trials only succeed when participants are willing to join them. Trial participants put their health in the hands of vaccine developers for the greater good of humanity. In doing so, they need to be confident that potential trial-related risks are mitigated and managed by the vaccine trial team. While long-term follow up (LTFU) was always a critical element of vaccine trials, there has been a recent push from regulators to follow up for longer periods. This means sponsors need to be prepared to gather data that demonstrates vaccine safety and efficacy over a more extended term.

This data can be used to increase trust with physicians and patients and to demonstrate vaccines' effectiveness and safety.

How COVID-19 changed vaccine trials

Before the COVID-19 pandemic, vaccine trials were a relatively niche area of clinical research. Typically, trials took between 5 and 10 years to conduct efficacy and safety studies. The urgency of the COVID-19 pandemic led to an unprecedented acceleration of vaccine development timelines. At the same time, regulators fast-tracked the regulatory review and approval process for vaccines. ICON provided clinical trial services to support one of the largest and most expeditious randomised clinical trials ever conducted. Our team helped accelerate the sponsor's mission to develop the world's first safe and effective investigational vaccine for COVID-19 vaccine to be approved. This partnership redefined industry expectations in terms of trial management and speed, while never compromising safety, quality and integrity. However, the speed with which vaccines became available may have unintentionally contributed to public fears about vaccine safety. Addressing these fears requires both effective communication and long-term data. While communication is largely in the hands of physicians and public health officials, sponsors have a significant role to play in gathering the necessary long-term data.

Demonstrating long-term safety starts in the early phases

When it comes to demonstrating long-term safety, vaccine developers should start with the end in mind. Planning for long-term follow up (LTFU) when designing study protocols will ensure the right data is captured both during and after a vaccine trial. Today sponsors can draw from a wide array of real world data (RWD) to understand the long-term impact of their vaccine or treatment. Clinical trial tokenisation (CTT) enables sponsors to follow patients for years after the trial ends by substituting Personal Identifiable Information (PII) with a unique token or deidentifier. This token is then used as a reference point so that real world healthcare data can be analysed to understand trends in disease, indication and post-exposure safety/effectiveness patterns without compromising patient privacy. Patient data can be captured discreetly allowing sponsors to track large volumes of real world patient data for cohorts of study participants. This provides real world evidence (RWE) of vaccine efficacy and interactions with other illnesses or drugs. However, to leverage CTT, sponsors must ensure that participants consent to being followed for a specified period of time once the trial is over.

When designing a study sponsors should consider what the minimum follow up period will be. While this may be dictated by the regulatory requirements, it can be worth considering a longer follow up period as well as the possibility of extension studies. Sponsors must also decide whether to deploy passive or active surveillance of participants and what data points to capture. Special attention may be necessary to account for population-specific groups such as older people, pregnant women or immunocompromised individuals as well as those with comorbidities. Sponsors will need to demonstrate to regulators that they have received informed consent and that the data is secured to protect patient privacy.

The future of vaccine trials

Having begun this article with a look back at the half-century of vaccination through the WHO EPI, it seems apt to close with a look to the future. As sponsors and supporters of vaccine trials we have a responsibility to conduct long-term follow up that demonstrate the continued safety of vaccines for years to come. By incorporating long-term follow up, RWD, and RWE strategies in clinical trials from as early as possible,

sponsors will help ensure the success of their vaccine programs. In doing so they can support individuals to confidently make informed decisions about vaccination.

This National Immunisation Awareness Month we thank both past and future vaccine trial participants for contributing to the development of safe and effective vaccines which save lives.

Fuente: ICON. Disponible en <https://n9.cl/tqyzn>

GSK's Phase 3b Study on Meningococcal Vaccine: A Potential Game Changer?

Aug 8. GlaxoSmithKline (GSK) is conducting a Phase 3b study titled A Phase 3b, Open-Label, Multi-Center Study to Assess the Immune Response And Safety of The Meningococcal Group B Vaccine Rmenb+Omv Nz When Administered to Healthy Participants Aged 10 To 20 Years Old, Who Were Primed During the First 2 Years Of Life. The study aims to evaluate the immune response and safety of a booster dose of the meningococcal group B vaccine, rMenB+OMV NZ, in adolescents and young adults who were initially vaccinated as infants. This research is significant as it seeks to confirm whether a booster dose enhances immunity in previously vaccinated individuals compared to those who have never received the vaccine.



The intervention being tested is the rMenB+OMV NZ vaccine, also known as Bexsero. It is a biological intervention designed to boost immunity against meningococcal group B infections in participants who were vaccinated as infants.

This interventional study is randomized with a parallel assignment model and is open-label, meaning no masking is involved. The primary purpose is prevention, focusing on comparing immune responses between previously vaccinated (primed) participants and those who are vaccine-naïve.

The study began on July 7, 2025, with the latest update on August 6, 2025. These dates are crucial as they mark the study's progress and timeline for completion, which is yet to be determined.

The outcome of this study could significantly impact GSK's stock performance and investor sentiment, as successful results may enhance the marketability of the Bexsero vaccine. In the competitive landscape of vaccine development, positive findings could position GSK favorably against other pharmaceutical companies working on similar vaccines.

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

Fuente: THE GLOBE AND MAIL. Disponible en <https://n9.cl/8cszp>

Vacuna contra el VPH protegerá frente al 90 % de los casos de cáncer de cuello uterino

10 ago. El Perú da un paso decisivo en la lucha contra el cáncer de cuello uterino. El Ministerio de Salud (Minsa) incorporó la vacuna nonavalente contra el Virus del Papiloma Humano (VPH) al Esquema Nacional de Vacunación, una dosis capaz de prevenir 9 variantes del virus responsable del 90 % de los casos de cáncer de cuello uterino y otros tumores asociados.

"El Minsa incorporó dicha dosis al Esquema Nacional de Vacunación. Será gratuita para menores de hasta 18 años."

La medida se oficializó a través de la Resolución Ministerial N° 440-2025/MINSA, publicada el pasado 18 de julio.

“La decisión permitirá que niñas, niños y adolescentes hasta los 18 años reciban la vacuna de forma gratuita en colegios, centros de salud y campañas comunitarias”, indicó la directora de la asociación de pacientes Esperantra, Karla Ruiz De Castilla.

Según dijo, la decisión del Minsa marca un avance fundamental para reducir la desigualdad en prevención oncológica. “Garantizar que la vacuna llegue a quienes más la necesitan es clave para salvar vidas”, recordó.



Con la incorporación de esta vacuna, De Castilla precisó que el país refuerza su estrategia de prevención primaria contra el cáncer de cuello uterino, avanzando hacia una cobertura más equitativa y efectiva para las poblaciones más vulnerables.

UN PASO CLAVE FRENTE A LA PRINCIPAL CAUSA DE MUERTE ONCOLÓGICA EN MUJERES

El cáncer de cuello uterino sigue siendo la primera causa de muerte por cáncer en mujeres peruanas, superando al cáncer de mama. De acuerdo con el informe “Reducir la brecha de equidad: atención oncológica para la mujer en Perú”, de Economist Impact, en el año 2022 se registraron 2,545 fallecimientos por este tipo de cáncer, en comparación con 1,951 por cáncer de mama.

El estudio en mención advierte que la enfermedad afecta con mayor frecuencia a mujeres jóvenes, de bajos ingresos y residentes en zonas rurales o comunidades indígenas, donde la vacunación enfrenta mayores dificultades de acceso.

Gracias a su inclusión en el Fondo Rotatorio de la Organización Panamericana de la Salud (OPS), De Castilla resaltó que el Perú podrá adquirir la vacuna nonavalente a menor costo y con disponibilidad garantizada, ya que todas las dosis cuentan con certificación de calidad de la OMS.

Fuente: Gestión Perú. Disponible en <https://n9.cl/9xztj>

Global Virus Network reaffirms support for mRNA vaccines and collaborative vaccine research

Aug 11. The Global Virus Network (GVN), a coalition of leading human and animal virologists from 80+ Centers of Excellence and Affiliates in more than 40 countries dedicated to advancing pandemic preparedness, is unequivocally committed to the continued development and deployment of mRNA vaccines and the global scientific collaboration that makes such innovation possible.

Vaccination remains one of public health's greatest achievements, preventing an estimated 4.4 million deaths globally each year. Building on this foundation, mRNA vaccine platforms have emerged as one of the most significant biomedical innovations of the 21st century, fundamentally reshaping our ability to respond rapidly and effectively to emerging viral threats. The real-world impact of mRNA vaccines has been striking. Between 2020 and 2024, COVID-19 vaccines are estimated to have prevented approximately 7.5 million deaths globally. In the United States alone, over 298 million mRNA vaccine doses were administered within the first

six months of rollout, with fewer than 1% of recipients experiencing significant adverse reactions or complications. Furthermore, clinical data showed that between March 2021 and January 2022, mRNA vaccines reduced the death rate by about 90%, and were 94% effective against severe disease during the Omicron wave.

Unlike traditional vaccines, which use weakened or inactivated viruses or parts of the pathogen to stimulate immunity, mRNA vaccines deliver synthetic genetic instructions that teach cells to produce a harmless viral protein. This approach primes the immune system without exposing individuals to the actual pathogen, offering a safe and adaptable platform with broad potential. Although propelled into the global spotlight by COVID-19, mRNA technology has been under development for decades, with early applications targeting rabies, influenza, and Zika, and more recently, showing promise in cancer immunotherapy.

"mRNA vaccine technology has fundamentally changed our ability to respond quickly and precisely to emerging viral threats," said Johan Neyts, PhD, Director of the GVN Center of Excellence at KU Leuven in Belgium, and a global leader in antiviral drug and vaccine development. "Various members of the GVN are working across continents to accelerate innovation in mRNA-based vaccines, not only for coronaviruses but also for dengue, Zika, Lassa fever, and other high-consequence pathogens."

The GVN emphasizes that scientific transparency, public health engagement, and a global commitment to research must remain at the heart of mRNA vaccine deployment strategies. "Global acceptance of vaccines, especially new platforms like mRNA, depends not only on scientific rigor but also on building and sustaining trust within communities," said Heidi Larson, PhD, Founding Director of the Vaccine Confidence Project and an internationally recognized expert in vaccine confidence and risk communication. "The GVN serves a critical function in this effort by convening trusted scientific voices around the world and advancing transparent, culturally informed engagement that helps bridge the gap between innovation and public confidence."

Around the world, governments and research institutions are advancing mRNA development and infrastructure as part of their pandemic preparedness and public health strategies. From South Africa to South Korea, Brazil to Belgium, GVN Centers are playing pivotal roles in these efforts.

"South Africa is leading a transformative moment in global health by building sustainable mRNA vaccine research and manufacturing ecosystems," said Quarraisha Abdool Karim, PhD, Co-Director of the GVN Center of Excellence at CAPRISA in South Africa and an internationally esteemed epidemiologist advancing health equity and scientific capacity across Africa. "By harnessing mRNA innovation locally, we're not only responding to today's needs but also empowering future generations of African scientists."

The GVN stands committed to advocating for a globally coordinated approach to the development and deployment of mRNA vaccines that includes:

- ♦ Expanding mRNA research and manufacturing capacity in low- and middle-income countries to close gaps in access and build regional resilience.
- ♦ Supporting next-generation mRNA innovation, including thermostable and self-amplifying formulations, to improve global distribution and longevity.
- ♦ Combating vaccine misinformation through collaborative initiatives with educators, journalists, and community leaders to ensure scientifically accurate, culturally sensitive messaging.

As a science-driven organization, the GVN is dedicated to advancing pandemic preparedness by fostering the exchange of knowledge, building global training capacity, accelerating translational research, and supporting effective and equitable responses to viral threats. mRNA vaccines are not just a product of biomedical ingenuity; they are a testament to what global collaboration can achieve when rooted in evidence and transparency.

About the Global Virus Network (GVN)

The Global Virus Network (GVN) is a worldwide coalition comprising 80+ Virology Centers of Excellence and Affiliates across 40+ countries, whose mission is to facilitate pandemic preparedness against viral pathogens and diseases that threaten public health globally. GVN advances knowledge of viruses through (i) data-driven research and solutions, (ii) fostering the next generation of virology leaders, and (iii) enhancing global resources for readiness and response to emerging viral threats. GVN provides the essential expertise required to discover and diagnose viruses that threaten public health, understand how such viruses spread illnesses, and facilitate the development of diagnostics, therapies, and treatments to combat them. GVN coordinates and collaborates with local, national, and international scientific institutions and government agencies to provide real-time virus informatics, surveillance, and response resources and strategies. GVN's pandemic preparedness mission is achieved by focusing on Education & Training, Qualitative & Quantitative Research, and Global Health Strategies & Solutions. The GVN is a non-profit organization. For more information, please visit www.gvn.org



Fuente: EurekAlert. Disponible en <https://n9.cl/8rw2y>

Pfizer introduces 20-valent Pneumococcal Conjugate Vaccine (PCV20) for adults in India

Aug 11. Pfizer has announced the launch of its next-generation 20-valent pneumococcal conjugate vaccine (PCV20) for adults in India. With broader serotype coverage, Pfizer's vaccine marks a significant advancement in protection against pneumococcal disease in adults.



image credit- shutterstock

Pfizer's vaccine helps protect against clinically relevant 20 serotypes responsible for a majority of invasive and non-invasive pneumococcal disease. It enables timely and proactive protection for all adults, including those living with chronic conditions. PCV20 will be available as a single shot vaccine and those vaccinated with PCV20 may not need a second dose.

Meenakshi Nevatia, Managing Director, Pfizer Ltd., India, said, "We are excited to launch PCV20, our 20-valent pneumococcal conjugate vaccine, in India. We believe this vaccine, with its broad coverage of 20 pneumococcal disease serotypes, will address the increasing need for adult immunisation in our country."

Adults over 50 years of age, as well as individuals with comorbidities such as asthma, COPD (Chronic Obstructive Pulmonary Disease), chronic kidney disease, and diabetes, face a higher risk of developing pneumococcal infections, which can result in complications, hospitalisation, and even death.

Timely immunisation is critical to reducing the risk of severe illness, hospitalisation, and mortality caused by pneumococcal disease. Vaccination with PCVs are considered one of the most effective public health tools for reducing the burden of pneumococcal disease.

The introduction of PCV20 in India strengthens Pfizer's longstanding commitment to preventive health and builds on 25 years of leadership in pneumococcal vaccine innovation.

Fuente: BioSpectrum India. Disponible en <https://n9.cl/yi95y>

SK bioscience takes on Pfizer, MSD with late-stage pneumococcal vaccine

Aug 13. SK bioscience has entered the final stage of clinical trials for its pneumococcal vaccine and is accelerating preparations for commercialization. The company is expanding its production facilities in partnership with Sanofi and has also entered the Chinese market, one of the largest in Asia.

Although the company posted an operating loss of 31.2 billion won (\$22.5 million) in the second quarter of this year based on separate financial statements, continuing its trend of losses, it is maintaining its aggressive R&D investment policy. Notably, SK bioscience has invested 33.7 billion won to expand its vaccine portfolio, with more than half of its total R&D expenses dedicated to commercializing its vaccines.

Among the vaccines currently under development, the product closest to commercialization is GBP410, a pneumococcal vaccine for children. Since signing a joint development agreement with Sanofi in 2014, the company has been conducting global phase 3 clinical trials and is on the verge of commercialization.

GBP410 has demonstrated equivalent immunogenicity and safety to existing approved vaccines through multinational phase 2 clinical trials, and phase 3 trials are underway in about 7,700 children and adolescents worldwide.



Upon launch, it is expected to compete with Pfizer's Prevenar 20 (20-valent vaccine) and MSD's Vaxneuvance (15-valent vaccine), and is being developed to cover 20 or more serotypes.

Production capacity is also being expanded in line with commercialization. SK bioscience has expanded its vaccine production facility at L House in Andong, North Gyeongsang Province, securing approximately 4,200 square meters of new space. The facility is expected to obtain certification from the U.S. FDA for current good manufacturing practices (cGMP) in mid-2026.

The company is also moving forward with its entry into the Chinese market. Clinical trials in China are mandatory for local sales, and SK bioscience recently received approval from the Center for Drug Evaluation (CDE) under the China's National Medical Products Administration (NMPA) for its phase 1 and 3 clinical trial applications (CTAs) for GBP410. The goal is to quickly complete phase 1 in China and then proceed to phase 3.

The company plans to actively utilize its cooperation network with Sanofi to penetrate the Chinese market. Sanofi has experience in launching various vaccines and immunotherapeutics in China, including the respiratory syncytial virus (RSV) preventive antibody therapy "Beyfortus" (nirsevimab), and this know-how is expected to contribute to the commercialization of GBP410.

"GBP410 is scheduled to complete global phase 3 trials by 2027 and then initiate the regulatory approval process, with commercialization expected as early as 2028," an SK bioscience official said.

Fuente: Korea Biomedical Review. Disponible en <https://n9.cl/m5ce9>

Grupo IHP participa en el ensayo europeo de una nueva vacuna contra la meningitis en lactantes desde las seis semanas de vida

13 ago. Grupo IHP ha formado parte del ensayo clínico internacional MET58, cuyos resultados muestran avances significativos en la protección frente a la meningitis en bebés. La investigación, recientemente publicada en la revista *Infectious Diseases and Therapy*, ha analizado por primera vez la administración de la vacuna tetravalente MenACYW-TT (MenQuadfi®) desde las seis semanas de edad, frente a la indicación actual a partir de los 12 meses.

El estudio ha comparado esta vacuna con la ya comercializada MCV4-TT (Nimenrix), administrándolas junto a las vacunas habituales del calendario (hexavalente, neumococo y triple vírica). En total, han participado 1.660 bebés de entre seis semanas y 18 meses, repartidos en 33 centros de siete países europeos.

En representación del grupo pediátrico andaluz, el doctor Ignacio Salamanca, coordinador de la Unidad de Investigación de Grupo IHP, ha sido uno de los investigadores del ensayo. "Hasta ahora, la vacuna sólo estaba indicada a partir del año de vida, dejando desprotegidos a los menores de esa edad. Este estudio demuestra que se puede iniciar la vacunación desde los 42 días sin comprometer la respuesta inmunitaria ni la eficacia del resto de vacunas coadministradas", ha afirmado.

Los resultados son muy positivos: más del 95 % de los lactantes alcanzaron niveles de protección



adecuados tras la dosis de recuerdo. La vacuna mostró una eficacia igual o superior frente a los serotipos C, W e Y, y una respuesta suficiente frente al serogrupo A, sin interferencias con otras vacunas infantiles. Además, no se registraron efectos adversos relevantes, sólo reacciones leves como fiebre, irritabilidad o dolor en la zona de inyección.

“Este avance abre la posibilidad de contar con una nueva herramienta de prevención frente a la enfermedad meningocócica invasora en lactantes, el grupo de mayor riesgo, con tasas de incidencia cuatro veces superiores a las del resto de la población”, ha subrayado el doctor Salamanca.

Fuente: Diario de Sevilla. Disponible en <https://n9.cl/7deae>

With scrapped mRNA funding, a vaccine maker finds a silver lining

Aug 14. When the HHS pulled \$500 billion in funding from the development of mRNA vaccines earlier this month, Secretary Robert F. Kennedy Jr. said in a statement that the Make America Healthy Again agenda would enable a shift to “safer, broader vaccine platforms.”

Experts, however, pointed to the unprecedented speed with which mRNA vaccines were developed and distributed in response to the COVID-19 pandemic that would not have been possible using traditional immunization methods. University of Minnesota infectious disease expert Mike Osterholm tweeted, “I haven’t seen a more dangerous decision in public health in my 50 years in the business.”

Beyond implications for future pandemic preparedness, the funding pullback also revealed a dynamic within the pharma industry between existing and emerging technologies, particularly as it pertains to public health and government funding for vaccines.

The technology behind mRNA vaccines has been celebrated in scientific circles as a major step forward in immunization, allowing for a timely response to new viruses, but the industry hasn’t put all of its eggs in that one basket. Older technologies like live-attenuated vaccines and viral vectors, for example, are still used in many kinds of shots, and companies that use these traditional methods could find that less funding for mRNA gives their own vaccines a boost — for now.

“In general for us, I see it as something that bodes positively,” said David Dodd, CEO of GeoVax, a biotech that uses a viral vector platform called modified vaccinia ankara in the clinic for COVID and head and neck cancers, as well as preclinical programs in mpox, Ebola, Marburg, Zika and more. “For the last five years, we were at a point where if you’re not doing mRNA, then you’re out of date and no one was going to support you, and that was a little bit too dramatic because there’s no one technology or platform that covers everything you need.”

Still, the mRNA platform holds plenty of value and should play a role in any robust public health system, Dodd said. Nothing compares to the manufacturing speed and access mRNA provides. But the technology also has limitations, particularly because it targets a specific antigen when a broader, multi-antigen response could be more fully protective against a disease in the long run.

There are pros and cons for each type, and different vaccine technologies don’t have to — and shouldn’t — be mutually exclusive, Dodd said.

Fuente: PharmaVoice. Disponible en <https://n9.cl/g5dqq>

New Study to Assess Immunogenicity of Butantan-DV Vaccine in ARDs

Aug 15. A new study will evaluate the safety and immunogenicity of the live attenuated tetravalent Butantan-Dengue vaccine (Butantan-DV) in individuals with granulomatosis with polyangiitis (GPA)—a form of antineutrophil cytoplasmic antibody (ANCA)-associated vasculitis (AAV)—among other autoimmune rheumatic diseases (ARDs).

The study, sponsored by the University of São Paulo General Hospital in São Paulo, Brazil, will test the vaccine in patients aged 7 to 59 years as well as in healthy age- and sex-matched volunteers.

“The Butantan-DV vaccine is a single-dose, tetravalent, live-attenuated formulation derived from all four dengue serotypes. Phase 2-3 data in the general population have demonstrated an overall efficacy of ~80 % and an acceptable safety profile, with clear operational advantages over other licensed vaccines (broader age indication and single-dose schedule),” the researchers wrote. “However, its immunogenicity and safety have not been prospectively examined in ARD patients under low-grade or no immunosuppression—precisely the subgroup for whom live vaccines may be permissible but still pose theoretical risks.”

The open-label, phase 3b prospective study expects to enroll 318 patients with an ARD and 159 age- and sex-matched controls, all of whom live in dengue-endemic areas of Brazil.

The participants will receive a single subcutaneous 0.5 mL Butantan-DV dose on day 1, with follow-up visits on days 7, 14, and 42. At these visits, they will have a clinical examination, laboratory testing, and a review of an adverse-event diary.

Long-term follow-up of the participants will continue to day 400 to monitor safety signals and to assess antibody persistence.

Two substudies, one on cellular immunity and another on viremia, will also be performed.

The primary outcome measure will be seroconversion rates in low-grade immunosuppressed patients with ARDS compared with healthy controls at day 42. Additional outcomes include common and serious adverse events, viremia kinetics, T-cell immunity, and durability of the antibody responses.



The study aims to inform dengue vaccine policy for individuals from 7 to 59 years old living with autoimmune rheumatic diseases in dengue endemic areas.

Fuente: Rare Disease Advisor. Disponible en <https://n9.cl/qavw0>

Ab&B Bio-Tech's RSV Vaccine Gains Approval in China and U.S.

Aug 17. Ab&B Bio-Tech CO., LTD. JS announced that its Investigational New Drug application for a self-developed Recombinant RSV vaccine (CHO cell) (Adjuvant) has been approved by both the China Center for Drug Evaluation and the U.S. Food and Drug Administration. This vaccine, targeting the highly contagious Respiratory Syncytial Virus, shows promising preclinical results with higher expression levels, better thermal stability, and superior immunogenicity compared to existing vaccines. The approval marks a significant step forward for the company in addressing a major health concern, potentially enhancing its market position and offering a new preventive solution for RSV infections.



- ◆ Ab&B Bio-Tech's RSV vaccine IND approved in China and U.S.
- ◆ The vaccine shows superior preclinical results, enhancing market position.

More about Ab&B Bio-Tech CO., LTD. JS Class H

Ab&B Bio-Tech CO., LTD. JS is a China-based vaccine company focused on the research, development, manufacturing, and commercialization of innovative and traditional vaccines using new technical methods. The company aims to replace traditional and imported vaccines in China and expand its presence in international markets. Its product range includes the quadrivalent subunit influenza vaccine and lyophilized human rabies vaccine candidate, among others.

Fuente: Tip Ranks. Disponible en <https://n9.cl/2qlew>

New mRNA treatment could protect us from all viruses

Aug 17. Columbia University researchers have created a new mRNA-based therapy that could help protect against all sorts of viruses.

Inspired by a rare genetic mutation that makes some people naturally virus-resistant, the team—led by immunologist Dusan Bogunovic—found a way to mimic this superpower in others.

How the treatment works

They designed a nasal drip using mRNA to temporarily boost 10 key antiviral genes in your body. Delivered with tiny lipid particles, this treatment kicked off just enough immune response to block viruses like flu, COVID-19, and Zika in lab animals—without causing harmful inflammation.

Potential game-changer in pandemic preparedness

If this works in humans, it could mean quick, broad protection against new viral outbreaks—even before vaccines are ready.

The researchers are still fine-tuning things for people, but this approach could seriously level up how we fight future pandemics.

Fuente: NewsBytes. Disponible en <https://n9.cl/9m2gp>



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Estrategia de búsqueda: (Vaccine) AND DP:([01.08.2025 TO 17.08.2025]) as the publication date 52 records.

1. [WO/2025/169951](#) METHOD FOR DETERMINING [VACCINE](#)-INDUCED IMMUNE RESPONSE

WO - 14.08.2025

Clasificación Internacional [C12Q 1/6827](#)Nº de solicitud PCT/JP2025/003724 Solicitante KEIO UNIVERSITY Inventor/a NAMKOONG, Ho

The purpose of the present invention is to provide a new marker that indicates the relevance between a [vaccine](#) and the immunity of a subject inoculated with the [vaccine](#). More specifically, the purpose of the present invention is to provide, for example, a method for determining the waning of [vaccine](#) effect using a new marker correlated with a tendency of [vaccine](#) effect to wane. The problem is solved by providing a method for determining the waning of [vaccine](#) effect, the method comprising a step for detecting a mutation in an immunoglobulin heavy-chain region in DNA contained in a sample from a subject, wherein, when the mutation is detected in the immunoglobulin heavy-chain region, the [vaccine](#) effect in the subject is determined to have a high tendency to wane.

2. [WO/2025/166964](#) METHOD FOR REDUCING OR ELIMINATING ALLOGRAFT REJECTION BY USING THYMUS [VACCINE](#)

WO - 14.08.2025

Clasificación Internacional [A61K 39/00](#)Nº de solicitud PCT/CN2024/099608 Solicitante TONGJI UNIVERSITY Inventor/a ZHANG, Xiaoqing

Provided is a method for reducing or eliminating allograft rejection by using a thymus vaccine. The method comprises the following steps: expressing major histocompatibility complex (MHC) from a donor and/or grafting a thymus epithelial cell from a donor in the thymus tissue of a recipient by means of the thymus vaccine. The thymus vaccine includes a thymic gene vaccine and/or a thymocyte vaccine. The thymic gene vaccine is used for expressing an MHC antigen from the donor in the thymus of the recipient, and comprises: (1) a gene expression vector; and (2) polynucleotides separately encoding MHC class I molecules and MHC class II molecules of the donor. The thymocyte vaccine is prepared by the following steps: sorting by using an anti-Epcam antibody to give thymus epithelial cells. By establishing a convenient donor source, the limitations of allograft caused by MHC mismatch are overcome, thus helping solve the clinical problem of limited donor sources.

3. WO/2025/162490 PREPARATION OF NEW MRNA VACCINE, AND USE THEREOF IN PHARMACEUTICAL COMBINATION FOR TREATING HEPATITIS B VIRUS DISEASE

WO - 07.08.2025

Clasificación Internacional A61K 39/29Nº de solicitud PCT/CN2025/075833 Solicitante JIANGSU CELL TECH MEDICAL RESEARCH INSTITUTE CO., LTD. Inventor/a YANG, Meijia

Provided in the present invention are a preparation of a new mRNA vaccine, and the use thereof in a pharmaceutical combination for treating the hepatitis B virus disease. Specifically, provided in the present invention is a therapeutic mRNA vaccine against chronic hepatitis B infection, which vaccine contains an mRNA encoding a large protein (L protein) region of a hepatitis B virus surface antigen, and also contains two mRNAs respectively encoding a middle protein (M protein) region of the hepatitis B virus surface antigen and encoding a small protein (S protein) region thereof, or contains an mRNA encoding the small protein (S protein) region. The therapeutic mRNA vaccine of the present invention can break the immune tolerance state in chronic hepatitis B infection, and has the potential to functionally cure chronic hepatitis B infection clinically.

4. 4598953 AUF DC ABZIELENDER IMPFSTOFF GEGEN NIPAH-VIRUSINFEKTION

EP - 13.08.2025

Clasificación Internacional C07K 16/28Nº de solicitud 23783875 Solicitante INST NAT SANTE RECH MED Inventor/a LEVY YVES

Nipah virus (NiV) is a recently emergent, highly pathogenic, zoonotic paramyxovirus. The inventors now designed an anti-CD40 mAb associated either with the Niv G ectodomain protein (Generation-1 vaccine), or the Niv G ectodomain protein and down-selected epitopes from the Niv F and N proteins (Generation-2 vaccine). Quality controls were performed on vaccine batches. The immunogenicity of both vaccines has been tested in hCD40Tg mice. A dose-dependent IFN γ T cell response to the antigen was observed. Three weeks post-boost, specific IgG were detectable in groups immunized with 10 μ g of vaccine. B cell responses were markedly improved 1 week post-boost. All samples at week -4 showed a neutralization with an average titer at 1:500. The inventors also demonstrated the potency of an innovative DC-targeting vaccine candidate to prevent NiV-B infection in challenge experiments in an AGM model. Responses were showed to cross-neutralize multiple strains of NiV, but also HeV. Targeting Nipah virus antigens to professional APCs can be efficiently used as a prophylactic means against a Nipah virus challenge at a lethal dose. Accordingly, the present invention relates to antibodies that are directed against a surface antigen of an antigen presenting

cell wherein the heavy chain and/or the light chain is conjugated or fused to the Nipah virus antigenic polypeptides.

5. WO/2025/170389 **VACCINE** COMPOSITION COMPRISING NOROVIRUS GII MRNA

WO - 14.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/KR2025/001935Solicitante K-BIOCELL INC.Inventor/a OH, Myung Ryurl

The present invention relates to a **vaccine** composition comprising mRNA encoding structural protein VP1 of a norovirus GI genotype. An mRNA **vaccine** composition was prepared by deriving a consensus sequence of norovirus GI genotype VP1 antigen, and the immunogenicity of the **vaccine** composition was confirmed. Therefore, the **vaccine** composition can be effectively used to prevent and treat norovirus infection.

6. 20250249087 CONSTRUCTION OF NOVEL HUMAN INFLUENZA VIRUS **VACCINE** AND APPLICATION

US - 07.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud 19002606Solicitante Wuhan JinYiTai Biological Co., Ltd.Inventor/a Yeping TAN

The present disclosure discloses a method and an application of a human influenza virus **vaccine**. The method and a non-replicating recombinant adenovirus named Ad-JYT-CMV-recHA promoter to control E1B19K or E1B19K-E1B55K fusion protein gene, and recombinant oncolytic adenoviruses named Ad-EE1A-hTERT-recHA and Ad-HTE1A-hTERT-recHA, respectively is constructed by optimizing a nucleotide sequence for influenza virus hemagglutinin HA, and thus the recHA gene is overexpressed in mammalian cells and the recHA is anchored on the cell surface and used for preventing influenza virus cold; human TERT promoter is used to control recHA gene, and a novel influenza recHA **vaccine** is introduced into an oncolytic adenovirus therapy as a tumor-heterologous artificial antigen for treating a tumor; furthermore, the recHA glycoprotein is anchored on the tumor cell surface as a tumor-specific artificial target, which is expected to combine with a CART-HA cell for treating a solid tumor.

7. 4598943 NEUE CHIMÄRE REKOMBINANTE IMPFSTOFFANTIGENE AUF MULTIPROTEINBASIS ZUR PRÄVENTION VON LYME-KRANKHEIT BEI TIEREN UND MENSCHEN

EP - 13.08.2025

Clasificación Internacional C07K 14/20Nº de solicitud 23875683Solicitante UNIV VIRGINIA COMMONWEALTHInventor/a MARCONI RICHARD T

A **vaccine** formulation for humans or other mammals (including dogs, horses, and cats) is provided. The **vaccine** formulation includes two chimeric proteins designed to elicit antibodies that bind to several targets on the surface of Lyme disease spirochetes during their residence in ticks and in mammals, and act synergistically to kill the bacteria through both antibody-mediated complement dependent and complement-independent mechanisms.

8. WO/2025/166642 HIGH-BIOACTIVITY PLASMID DNA, PREPARATION AND STORAGE THEREOF, **VACCINE**, KIT AND USE THEREOF

WO - 14.08.2025

Clasificación Internacional C12N 15/10Nº de solicitud PCT/CN2024/076716Solicitante CAPITAL MEDICAL UNIVERSITYInventor/a SUN, Wenzhi

The present invention relates to the field of biomedicine. Particularly provided are a high-bioactivity plasmid DNA, preparation and storage thereof, a **vaccine**, a kit, and a use thereof. The preparation method for the high-bioactivity plasmid DNA comprises: lysing bacterial cells after expansion culture and collecting a supernatant to obtain a crude solution containing a plasmid; purifying the crude solution by means of two-step liquid chromatography to obtain a purified plasmid DNA; precipitating the purified plasmid DNA by means of an alcohol and/or a polyol to concentrate and activate the plasmid DNA; and storing the plasmid DNA as a precipitate. The provided preparation method meets medical-grade standards, shortens the plasmid DNA purification time and reduces the material cost, improves the plasmid DNA yield and activity, and facilitates long-term storage. In addition, the plasmid DNA has higher in-vivo transfection efficiency and expression stability, thereby facilitating efficient introduction of a target gene into an organism. Thus, the plasmid DNA obtained by means of the method is suitable for use as an active component of a gene therapy drug or a DNA **vaccine**.

9.WO/2025/168744A **VACCINE** FOR PROTECTING A SWINE AGAINST AFRICAN SWINE FEVER

WO - 14.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2025/053180Solicitante INTERVET INTERNATIONAL B.V.Inventor/a VAN DEN BORN, Erwin

The invention pertains to a live attenuated African swine fever virus Georgia 2007 (ASFV-G) Δ 9GL/ Δ UK/ Δ EP153R strain for use in a **vaccine** for protecting a swine against an infection with African swine fever virus (ASFV) by administering the **vaccine** comprising the live attenuated ASFV-G- Δ 9GL/ Δ UK/ Δ EP153R strain to the swine.

10.4598588SUPERANTIGENIMPFSTOFFKONJUGAT ZUR BEHANDLUNG VON KREBS

EP - 13.08.2025

Clasificación Internacional A61K 47/68Nº de solicitud 23875747Solicitante MUSC FOUND FOR RES DEVInventor/a DOLLOFF NATHAN G

The present disclosure provides compositions comprising **vaccine** conjugates with a SMEZ-2 carrier. Further provided are methods for treating cancer comprising administering the **vaccine** conjugates provided herein.

11.20250255951VACCINES AGAINST CHLAMYDIA SP.

US - 14.08.2025

Clasificación Internacional A61K 39/118Nº de solicitud 19197254Solicitante STATENS SERUM INSTITUTInventor/a Frank Follmann

The present invention describes an efficient **vaccine** against a *Chlamydia trachomatis* (Ct). The **vaccine** is based on recombinant fusion molecules that are capable of generating a high titered neutralizing antibody response that is protective against various Ct serovars. Our invention furthermore describe the combination of these antibody promoting fragments with Ct antigens that are targets for T cells with the aim to provide a **vaccine** that activate both arms of the immune system.

12. WO/2025/171182 TREATMENT OF CANCER PATIENTS WITH TUMOR INFILTRATING LYMPHOCYTE THERAPIES IN COMBINATION WITH CANCER **VACCINE**

WO - 14.08.2025

Clasificación Internacional A61K 40/11Nº de solicitud PCT/US2025/014867 Solicitante IOVANCE BIOTHERAPEUTICS, INC. Inventor/a NATARAJAN, Arvind

Provided are methods for treating cancer in a patient in need thereof with TILs in combination with a cancer **vaccine**.

13. 20250255949 STREPTOCOCCUS SUIS **VACCINE** COMPOSITION COMPRISING IMMUNOGENIC FUSION POLYPEPTIDES

US - 14.08.2025

Clasificación Internacional A61K 39/09Nº de solicitud 18857553 Solicitante Intervacc AB Inventor/a Sara Frosth

The present disclosure relates to immunogenic fusion polypeptides, immunogenic compositions and **vaccine** compositions comprising said fusion polypeptides and use thereof for immunization of mammals susceptible to *Streptococcus suis* infection. The disclosure also relates to methods for preparing, formulating and administering such compositions.

14. 4593877 IMPFSTOFFKONSTRUKT UND VERWENDUNGEN DAVON

EP - 06.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23801865 Solicitante CEVA HAMPTON LTD Inventor/a THOMPSON IAN

The present disclosure relates generally to **vaccine** constructs comprising a polynucleotide encoding a salmon alphavirus-like particle and uses thereof, including for inducing an immune response against salmonid alphavirus.

15. 20250259721 DIGITAL **VACCINE** SYSTEM, METHOD AND DEVICE

US - 14.08.2025

Clasificación Internacional G16H 10/60Nº de solicitud 19193575 Solicitante VYDIANT, INC Inventor/a James Kaput

A digital **vaccine** system, method and device that maintains a health knowledge base, inputs user characteristics, generates health scores based on the user characteristics and provides pathogen risk recommendations based on the user characteristics, health scores and knowledge base, wherein the recommendations are indicated by the knowledge base to be likely to improve the user's health.

16. 20250255957 VACCINES HAVING AN ANTIGEN AND INTERLEUKIN-21 AS AN ADJUVANT

US - 14.08.2025

Clasificación Internacional A61K 39/39Nº de solicitud 19184899 Solicitante The Trustees of the University of Pennsylvania Inventor/a David B. Weiner

Disclosed herein is a **vaccine** comprising an antigen and IL-21. Also disclosed herein are methods for increasing an immune response in a subject. The methods may comprise administering the **vaccine** to the subject in need thereof.

17. WO/2025/166439 ANTIGENIC PEPTIDE, IMMUNOGENIC COMPOSITION AGAINST SCHISTOSOMIASIS CONTAINING THE PEPTIDE, AND USES

WO - 14.08.2025

Clasificación Internacional C12P 21/00Nº de solicitud PCT/BR2025/050048 Solicitante INSTITUTO BUTANTAN Inventor/a VERJOVSKI-ALMEIDA, Sergio

The present invention relates to a new **vaccine** candidate against schistosomiasis. This **vaccine** candidate is composed of peptides from different proteins of the parasite *Schistosoma mansoni*. Additionally, the present invention relates to use of the peptides for preparing a medicament for the prevention and/or treatment of schistosomiasis and to a method for preventing and/or treating schistosomiasis.

18. 20250255944 IMMUNOACTIVATOR, **VACCINE** ADJUVANT, AND METHOD FOR INDUCING IMMUNITY

US - 14.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18844599 Solicitante TEIKYO UNIVERSITY Inventor/a Ryo SUZUKI

Disclosed is an immunoactivator that is capable of enhancing induction of both humoral immunity and cellular immunity and contains an active ingredient derived from cell walls (derived from a natural product); a **vaccine** adjuvant; and a method for inducing immunity including administering the immunoactivator. The immunoactivator contains, as an active ingredient, particles having a maximum diameter within a range of 1 to 800 nm, wherein the particles comprise cell-wall-derived polysaccharides.

19. WO/2025/161943 LIPID COMPOUND AND LIPID NANOPARTICLE FOR DELIVERY

WO - 07.08.2025

Clasificación Internacional C07C 229/16Nº de solicitud PCT/CN2025/072219 Solicitante RINUAGENE BIOTECHNOLOGY CO., LTD. Inventor/a DONG, Yijie

Disclosed in the present application are a compound having a structural formula as shown in formula (I), and a pharmaceutically acceptable salt thereof or a stereoisomer thereof. Further disclosed in the present application is a nanoparticle composition containing the compound or the pharmaceutically acceptable salt thereof or the stereoisomer thereof. The nanoparticle of the present application can efficiently deliver a drug and a **vaccine** into a cell, thereby exerting the therapeutic or prophylactic purpose of the drug and the **vaccine**.

20. WO/2025/170904 INJECTABLE VACCINES AGAINST CANINE RESPIRATORY DISEASES

WO - 14.08.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/US2025/014445 Solicitante ZOETIS SERVICES LLC Inventor/a KLINK, Holly Annette

Provided is an injectable **vaccine** comprising: a heat treated purified whole cell extract of *B bronchiseptica*, wherein this extract has been heated for 10 to 60 minutes at 45°C to 65°C; and p68 pertactin protein. Methods of making and methods of using the **vaccine** are also provided.

21. WO/2025/168804A **VACCINE** FOR PROTECTING A PREGNANT SWINE AGAINST AFRICAN SWINE

WO - 14.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/EP2025/053301Solicitante INTERVET INTERNATIONAL B.V.Inventor/a VAN DEN BORN, Erwin

The present invention pertains to the use of a live attenuated African swine fever virus Georgia 2007-Δ9GL/ΔUK/ΔEP153R (ASFV-G-Δ9GL/ΔUK/ΔEP153R) strain for protecting a pregnant swine against an infection with African swine fever virus (ASFV) by administering a **vaccine** comprising the live attenuated ASFV-G-Δ9GL/ΔUK/ΔEP153R strain to the pregnant swine.

22. 20250255952SELF-AMPLIFYING MRNA **VACCINE** BASED ON Z7 GENOME

US - 14.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 19050556Solicitante Fengwei BaiInventor/a Fengwei Bai

The present invention relates to a self-amplifying mRNA **vaccine**, including: a self-amplifying backbone comprising a live attenuated virus having a hairpin loop insert, and wherein at least one or more structural genes of the live attenuated virus have been replaced with at least one target antigen-encoding nucleic acid sequence.

23. WO/2025/171256COMPOSITIONS AND METHODS FOR TREATING MALARIA

WO - 14.08.2025

Clasificación Internacional C07K 16/20Nº de solicitud PCT/US2025/015008Solicitante CASE WESTERN RESERVE UNIVERSITYInventor/a KING, Christopher L.

An anti-PvAMA1 antibody, antibody fragment, antigen binding fragment thereof or **vaccine** for use in treating and/or preventing malaria as described herein.

24. 4596712VERFAHREN ZUR IDENTIFIZIERUNG UND VERWENDUNG VON BIOMARKERN

EP - 06.08.2025

Clasificación Internacional C12Q 1/6806Nº de solicitud 24155636Solicitante WOBBLE GENOMICS LTDInventor/a KUO RICHARD IZEN

Methods for producing an RNA **vaccine** for a subject with a disease and producing a database of disease biomarkers are provided. Methods for discovering disease biomarkers and diagnosing disease are also provided. The present invention enables frequent monitoring of subjects and adjustment of treatment. The present invention also provides methods for preparing processed nucleic acid samples with a more uniform distribution of sequences, which can be more readily analysed. The methods may be combined to further improve the ability to produce RNA vaccines and discover disease biomarkers.

25. WO/2025/165727 METHODS AND COMPOSITIONS FOR PROMOTING ANTI-TUMOR IMMUNITY AND ENHANCING THE IMMUNE FUNCTION

WO - 07.08.2025

Clasificación Internacional A61K 39/39Nº de solicitud PCT/US2025/013344 Solicitante NEW YORK UNIVERSITY Inventor/a PACOLD, Michael

Various methods and compositions for the activating or augmenting an immune response, and for augmenting an antibody production are presented herein. Also presented herein are methods of augmenting an immune response to a vaccine. Also presented herein are methods of preventing or delaying age-related decline in physical strength or exercise tolerance. Further presented herein are methods and compositions for improving wound healing. Yet further presented herein are methods for inhibiting growth of a tumor.

26. 20250255954 STABILIZED SPIKE PROTEIN AND METHOD OF USE THEREOF AS A CORONAVIRUS DISEASE 2019 (COVID-19) VACCINE

US - 14.08.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18856345 Solicitante Yuanhan Wu Inventor/a Yuanhan Wu

Disclosed herein is a new method referred to as Conformational Shifting by Distance and Volume Analysis (CS-DVA) which can be employed to change the dynamics of multi-state glycoproteins for altered immune responses. Also disclosed are stabilized spike antigens and methods of use thereof for SARS-COV-2 vaccines.

27. WO/2025/163087 METHODS OF IDENTIFYING AND USING BIOMARKERS

WO - 07.08.2025

Clasificación Internacional C12Q 1/6806Nº de solicitud PCT/EP2025/052431 Solicitante WOBBLE GENOMICS LIMITED Inventor/a KUO, Richard Izen

Methods for producing an RNA vaccine for a subject with a disease and producing a database of disease biomarkers are provided. Methods for discovering disease biomarkers and diagnosing disease are also provided. The present invention enables frequent monitoring of subjects and adjustment of treatment. The present invention also provides methods for preparing processed nucleic acid samples with a more uniform distribution of sequences, which can be more readily analysed. The methods may be combined to further improve the ability to produce RNA vaccines and discover disease biomarkers.

28. 4598571 GEFLÜGELIMPFSTOFFE UND VERFAHREN ZUM SCHUTZ VON GEFLÜGEL

EP - 13.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23801214 Solicitante ZOETIS SERVICES LLC Inventor/a AOKI SERGIO MORAES

A method of protecting an offspring of a hen against infectious bronchitis infection is provided, the method comprising administering to said hen a vaccine comprising an inactivated infectious bronchitis virus and adjuvanted with water-in-oil emulsion and an immunostimulatory oligonucleotide.

29. 4598578 STABILISIERUNG EINES VIRUSBASIERTEN THERAPEUTISCHEN MITTELS

EP - 13.08.2025

Clasificación Internacional A61K 47/26Nº de solicitud 23874154 Solicitante ELAREX
 INC Inventor/a IWASHKIW JEREMY ANDREW

A virus-based active agent is mixed with trehalose and water and dried. The mixture may also contain one or more of pullulan and albumin. The mixture may be dried to a moisture content of 0.1-10%. The drying may be under vacuum sufficient to produce a foam. Some or all of the drying may be at a temperature in the range of 15-40°C, or at a temperature in the range of 1-15°C, or both. The active agent may be based on a vesicular stomatitis virus (VSV) or an adenovirus (AdV). The dried mixture may be stored at a temperature in the range of 1-55°C. A composition includes a virus, which may be a derived or modified form of a virus such as VSV or AdV. The composition also includes trehalose and optionally one or more of pullulan, and albumin. The composition may be used for a virus-based **vaccine**.

30. 2637987 METHOD OF DETERMINING VACCINATIONS AND/OR MEDICATION REQUIRED BY AN INDIVIDUAL

GB - 13.08.2025

Clasificación Internacional G16H 10/60Nº de solicitud 202401808 Solicitante PERSONALISED
 DIAGNOSTICS LTD Inventor/a CLIVE MINIHAN

Computer method 300 for determining vaccinations and / or medication required by an individual. Method 300 is based on obtaining indications of the individual's: risk 301; medical history 305 and what vaccinations and/or medication are available 307. Determining what vaccinations and / or medication the individual requires 309 is based on the obtained indications 301, 305, 307. Other embodiments include making this determination by obtaining a list of all diseases present in a region and filtering it based on the individual's travel plans. Alternatively, a list of all available medication and /or vaccines is obtained and filtered based on travel plans. Further embodiments include a method for adding a new disease to a database; methods for obtaining up-to-date information by periodically crawling travel advice websites to identify any change(s). In the event of change(s) an alert is provided for assessment by a clinician to determine whether a change in medication and /or vaccination are required. Alternatively, a machine learning model may be applied to determine the nature of the change(s) and whether a change in medication and /or **vaccine** is required.

31. WO/2025/163760 ATTENUATED BACTERIAL STRAIN, LIVE ATTENUATED **VACCINE**, CONTROL METHOD, AND PRODUCTION METHOD

WO - 07.08.2025

Clasificación Internacional C12N 15/09Nº de solicitud PCT/JP2024/002908 Solicitante THE NATIONAL
 RESEARCH AND DEVELOPMENT AGENCY, JAPAN FISHERIES RESEARCH AND EDUCATION
 AGENCY Inventor/a MATSUURA Yuta

Provided by the present invention is an attenuated bacterial strain of *Nocardia seriolae*, in which there is a mutation in at least a part of the MtrA gene and the amino acid 179 Val is substituted by Gly in at least part of the amino acid sequence encoded by the MtrA gene.

32. [WO/2025/165132](#) RANKL MUTANT AND VACCINE COMPOSITION COMPRISING SAME

WO - 07.08.2025

Clasificación Internacional [C07K 14/705](#)Nº de solicitud PCT/KR2025/001531 Solicitante ARKGEN BIOSCIIONS CO.,LTD. Inventor/a LIM, Wonbong

An RANKL protein mutant according to the present invention can inhibit RANK activity through inhibition of the ability of RANKL to bind to both OPG and RANK, thereby suppressing differentiation into osteoclasts. Therefore, the RANKL protein mutant of the present invention can be effectively used for the prevention or treatment of bone metabolic diseases.

33. [WO/2025/171082](#) PREFUSION-STABILIZED HSV2 GB PROTEINS

WO - 14.08.2025

Clasificación Internacional [C07K 14/035](#)Nº de solicitud PCT/US2025/014710 Solicitante BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM Inventor/a MCLELLAN, Jason

Provided herein are engineered HSV2 gB polypeptides. In some aspects, the engineered gB polypeptides exhibit enhanced conformational stability and/or antigenicity. Methods are also provided for use of the engineered gB polypeptides as diagnostics, in screening platforms, and/or in vaccine compositions.

34. [20250255828](#) LYOPHILIZED LIPID NANOPARTICLES AND METHODS OF THEIR USE

US - 14.08.2025

Clasificación Internacional [A61K 9/51](#)Nº de solicitud 19047781 Solicitante Ultragenyx Pharmaceutical Inc. Inventor/a ISAIAS PRADO

The invention is directed to the field of therapeutic formulations, in particular to lyophilization of a therapeutic cargo molecule, such as RNA. The invention provides a method for lyophilization of a molecule. The present disclosure further describes a lyophilized composition obtainable by the inventive method, a pharmaceutical composition, a vaccine, a therapeutic and a kit or kit of parts. Moreover, the disclosure herein provides a novel lyophilization excipient that protects the composition from degrading when, for example, lyophilizing RNA. The use of the inventive method further includes the manufacture of a composition that can be used after lyophilization with equivalent therapeutic effect and composition integrity.

35. [WO/2025/166067](#) SARS-COV-2 VACCINE COMPOSITIONS AND METHODS

WO - 07.08.2025

Clasificación Internacional [C07K 14/32](#)Nº de solicitud PCT/US2025/013877 Solicitante HELIX NANOTECHNOLOGIES, INC. Inventor/a RAJANIEMI, Hannu

Disclosed herein are fusion polypeptides comprising: (i) a fragment antigen comprising an epitope of SARS-CoV-2; and (ii) a complement binding polypeptide. The disclosure also provides polynucleotides (e.g., mRNA) encoding the same. Also disclosed herein are methods of making and using the fusion polypeptides and fusion polynucleotides of the present disclosure.

36. [20250249086](#) PROTEIN-SACCHARIDE CONJUGATION WITH SODIUM CYANOBOROHYDRIDE

US - 07.08.2025

Clasificación Internacional A61K 39/095Nº de solicitud 18854372Solicitante Sanofi Pasteur Inc.Inventor/a Maryalice Ginley

Methods and uses of conjugating saccharides to protein carriers are disclosed herein. Exemplary conjugates prepared according to those methods and uses are also disclosed. Additionally, methods for quantifying the amount of sodium borohydride in a sodium cyanoborohydride reagent are disclosed herein. **Vaccine** compositions as well as related methods and uses are also disclosed herein.

37.20250255946LOW-DOSE NEOANTIGEN **VACCINE** THERAPY

US - 14.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18891398Solicitante Gritstone bio, Inc.Inventor/a Karin Jooss

Disclosed herein are compositions that include antigen-encoding nucleic acid sequences and/or antigen peptides. Also disclosed are nucleotides, cells, and methods associated with the compositions including their use as vaccines, including vectors and methods for a heterologous prime/boost vaccination strategy.

38.4600256IMPFSTOFF GEGEN DAS RESPIRATORISCHE SYNZYTIALVIRUS SOWIE HERSTELLUNGSVERFAHREN DAFÜR UND VERWENDUNG DAVON

EP - 13.08.2025

Clasificación Internacional C07K 14/135Nº de solicitud 24810137Solicitante LIVERNA THERAPEUTICS INCInventor/a LI JIANGLONG

The present disclosure relates to an immunological composition comprising or encoding a human respiratory syncytial virus antigen, and the immunological composition is selected from a nucleic acid immunological composition, a polypeptide immunological composition or a viral immunological composition.

39.20250250306IMMUNOGENIC PROTEINS AND NUCLEIC ACIDS ENCODING THE SAME

US - 07.08.2025

Clasificación Internacional C07K 14/005Nº de solicitud 19061222Solicitante International AIDS **Vaccine** Initiative, Inc.Inventor/a Jordan WILLIS

The invention relates to proteins and nucleic acids for immunization regimens, modifications thereof, and/or development of nanoparticles, and/or development of membrane-anchored immunogens, and methods of making and using the same. The invention also encompasses cell surface trimers that bind to the broadly neutralizing antibodies and/or nucleic acids encoding the same.

40.20250257141COMBINATION THERAPY FOR THE TREATMENT OF CANCER COMPRISING A FAS AXIS ANTAGONIST AND A T-REG CELL DEPLETING AGENT ANTAGONIST

US - 14.08.2025

Clasificación Internacional C07K 16/28Nº de solicitud 18859717Solicitante Hoffmann-La Roche Inc.Inventor/a Maria Amann

The present disclosure is directed to the combination of a Fas axis antagonist, such as an anti-FasL antibody, and a Treg cell depletion therapy, for example an anti-CD25 antibody, optionally with a cancer vaccine, for use in the treatment of cancer.

41. WO/2025/162105 HUMAN PAPILLOMA VIRUS (HPV) MRNA VACCINES AND USES THEREOF

WO - 07.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud PCT/CN2025/073829 Solicitante RINUAGENE BIOTECHNOLOGY CO., LTD. Inventor/a DONG, Yijie

The invention provides a HPV mRNA lipid nanoparticle vaccine, comprising an open reading frame encoding a fusion protein of E2, E6 and E7 antigens of a high-risk HPV antigen. The invention also provides a method for treating and preventing HPV infection related diseases such as cervical cancer, cervical precancerous lesions, high-grade intraepithelial lesions of the cervix, high-grade squamous intraepithelial lesions (HSIL) or cervical intraepithelial neoplasia (CIN).

42. 3034089 VACCINE AGAINST PORCINE PARVOVIRUS

ES - 12.08.2025

Clasificación Internacional A61K 39/23Nº de solicitud 17801619 Solicitante Boehringer Ingelheim Vetmedica GmbH Inventor/a VAUGHN, Eric, Martin

43. 20250257102 PREFUSION-STABILIZED HMPV F PROTEINS

US - 14.08.2025

Clasificación Internacional C07K 14/08Nº de solicitud 19195969 Solicitante BOARD OF REGENTS, THE UNIVERSITY OF TEXAS SYSTEM Inventor/a Jason McLellan

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.

44. 20250255840 HIV TREATMENT COMPOSITIONS AND METHODS

US - 14.08.2025

Clasificación Internacional A61K 31/167Nº de solicitud 19175991 Solicitante NantCell, Inc. Inventor/a Kayvan Niazi

HIV treatment, and especially treatment of latent infected CD4 cells, can be significantly improved using a kick-and-kill approach that employs an immune stimulation component and/or HDAC inhibition as one treatment component, and that may also include a second component in which a vaccine composition, various NK cells, CAR-T cells, and/or broadly neutralizing antibodies are administered.

45. 20250249088 THERAPEUTIC AND VACCINE CANDIDATES AGAINST SARS-COV-2

US - 07.08.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18702303 Solicitante GENEONE LIFE SCIENCE, INC. Inventor/a Kar MUTHUMANI

Provided herein is an immunogenic composition comprising a synthetic antigen to COVID spike proteins, particularly the chimeric intermediate structure C-A Complex. Also disclosed herein is a method of preventing and/or treating a COVID infection in a subject in need thereof, by administering the immunogenic composition to the subject.

46. 4598547 VERFAHREN UND ZUSAMMENSETZUNGEN ZUR BEHANDLUNG VON BLASENKREBS
EP - 13.08.2025

Clasificación Internacional A61K 35/13Nº de solicitud 23875792Solicitante UNIV JEFFERSONInventor/a ANDREWS DAVID

The present disclosure relates to compositions and methods for treating bladder cancer. In some embodiments the compositions and methods involve using antisense (AS) nucleic acids directed against Insulin-like Growth Factor 1 Receptor (IGF-1R). The AS may be administered to the patients systemically, or may be used to produce an autologous cancer cell vaccine. In embodiments, the AS are provided in an implantable irradiated bioreactor chamber comprising tumor cells and an effective amount of the AS. The chambers are irradiated and implanted in the abdomen of subjects and stimulate an immune response that attacks tumors distally. The compositions and methods disclosed herein may be used to treat many different kinds of bladder cancer, including metastatic breast cancer.

47. 321533 METHODS OF TREATING PANCREATIC CANCER WITH A PD-1 AXIS BINDING ANTAGONIST AND AN RNA VACCINE

IL - 01.08.2025

Clasificación Internacional A61K 39/00Nº de solicitud 321533Solicitante GENENTECH, INC.Inventor/a

48. 20250255953 RECOMBINANT INFLUENZA VIRUSES WITH STABILIZED HA FOR REPLICATION IN EGGS

US - 14.08.2025

Clasificación Internacional A61K 39/145Nº de solicitud 19067428Solicitante The University of TokyoInventor/a Yoshihiro Kawaoka

Modified influenza virus neuraminidases are described herein that improve viral replication, thus improving the yield of vaccine viruses. Expression of such modified neuraminidases by influenza virus may also stabilize co-expressed hemagglutinins so that the hemagglutinins do not undergo mutation.

49. 4593874 VIRUSÄHNLICHE PARTIKEL MIT SARS-COV-2-ANTIGENEN ALS BOOSTER-IMPfstoffe UND VERWENDUNGEN DAVON

EP - 06.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23769188Solicitante BAVARIAN NORDIC ASInventor/a RAMBICHLER STEPHAN

The present invention relates to the use of vaccines comprising virus-like particles displaying at least one SARS-CoV-2 antigen, such as the receptor-binding domain (RBD) of the SARS-CoV-2 spike protein, as vaccine boosters. Antigens are displayed on virus-like particles (VLPs) and produce an immune response in vaccinated subjects. The invention also relates to methods of treatment using the recombinant VLPs as boosters to treat and/or prevent infection with SARS-CoV-2, and methods of preparation thereof.

50. 4599067 KÜNSTLICHE POLYNUKLEOTIDE ZUR EXPRESSION VON PROTEINEN

EP - 13.08.2025

Clasificación Internacional C12N 15/67Nº de solicitud 23785796 Solicitante CERTEST BIOTEC S
Inventor/a BROSET BLASCO ESTHER

The present invention provides a polynucleotide comprising, in the 5' to 3' direction, a 5' untranslated region (5'-UTR) and an open reading frame (ORF), wherein the 5'-UTR comprises at least two tandem repeats of sequence 5'-GCCNCC-3' operatively linked to the ORF, and wherein N is any nucleotide. The invention also provides a composition comprising a lipid nanoparticle and the polynucleotide and a pharmaceutical composition, and their use in medicine, particularly for use as a **vaccine** or for use in gene therapy.

51. 4599846 NEUE PROMOTOREN

EP - 13.08.2025

Clasificación Internacional A61K 39/12Nº de solicitud 25168096 Solicitante BOEHRINGER INGELHEIM
VETMEDICA GMBH Inventor/a MUNDT ALICE

The present invention relates to the field of (vector) vaccines, and especially to novel promoter sequences, expression cassettes and vectors, which are suitable to express genes of interest, especially antigen encoding sequences. The viral vectors of the present invention are useful for producing an immunogenic composition or **vaccine**.

52. 20250249090 NEW FELINE HERPES VIRUS **VACCINE**

US - 07.08.2025

Clasificación Internacional A61K 39/245Nº de solicitud 18252590 Solicitante Boehringer Ingelheim
Vetmedica GmbH Inventor/a Callie Ann Visek

The present invention relates i.a. to an EHV (Equine Herpesvirus) comprising a Feline Herpes Virus (FHV) Antigen encoding sequence inserted into ORF70 (US4) and/or ORF1/3. Furthermore, the present invention relates to methods for immunizing a feline comprising administering to such feline an immunogenic composition of the present invention. Moreover, the present invention relates to methods for the treatment or prophylaxis of clinical signs caused by Feline Herpes Virus in a feline.

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

Estrategia de búsqueda: *vaccine.ti. AND @PD>="20250801"<="20250817* 18 records

Document ID	Title	Inventor	Applicant Name
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US 20250255944 A1	IMMUNOACTIVATOR, VACCINE ADJUVANT, AND METHOD FOR INDUCING IMMUNITY	SUZUKI; Ryo et al.	TEIKYO UNIVERSITY
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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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