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

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

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- Patentes más recientes en Patentscope sobre vacunas.

Noticias en la Web

Delivering next-generation medicines and vaccines

Feb 17. Pharmaceutical companies have been at the forefront of advances in medical innovation that are leading to real improvements in the health of people worldwide. These advances are also supporting healthcare systems as they grapple with the rise of chronic diseases, ageing populations, the impact of climate change, and emerging infectious diseases, according to Dr. David Reddy, Director General of the International Federation of Pharmaceutical Manufacturers and Associations (IFPMA).



Over the past 20 years, the industry has launched over 940 novel active substances globally, addressing cancer, cardiovascular diseases, neurology, and infectious diseases, among others. There are over 12,700 medicines in various stages of clinical development globally, half of them biologics and the other half small molecules. By 2035, it is estimated that 700 new medicines could be launched that can prevent, slow, or stop disease progression.

"From my own experience working on mRNA technology decades ago, it has been hugely rewarding to see the acceleration of new mRNA vaccines after their use in the COVID-19 response. Today, 286 vaccines are under development, targeting a wide array of diseases beyond infectious ones, including cancer, allergies and even Alzheimer's," said Dr. Reddy, who holds a PhD in Cellular and Molecular Biology and has post-doctoral research experience in molecular neurobiology.

Dr. Reddy has more than 30 years of experience in the development and commercialization of medicines, including 13 years as the head of the Medicines for Malaria Venture (MMV). During his tenure as CEO of MMV, the organization saw 15 malaria medicines launched, 1.2 billion treatments distributed, saving more than 15 million lives, and raised around \$900 million in funding.

He said that precision medicines that are already transforming cancer treatment now hold potential for chronic diseases as well. Advances such as CRISPR — a technology that research scientists use to selectively modify the DNA of living organisms — are likewise paving the way for potential cures for previously untreatable genetic disorders, with cell therapies diversifying treatment options across a range of diseases.

Dr. Reddy noted that this progress is underpinned by the pharmaceutical industry's commitment to investing in the research and development that makes advances in medical innovation possible. The top 50 pharmaceutical companies alone are estimated to have spent a total of \$167 billion in R&D in 2022. Moreover, the industry's R&D spending is rising, increasing by almost 60% in the 10 years from 2012 to 2022.

"This R&D investment has a significant impact on health outcomes, but also strengthens healthcare systems and economies worldwide," Dr. Reddy said.

Data from an IFPMA-commissioned analysis carried out by the independent economic research institute WifOR demonstrated that the industry contributed \$2,295 billion to global GDP in 2022 through direct, indirect, and induced effects.

For every dollar generated by pharmaceutical activities, an additional \$2.04 was created along the global supply chain. On top of the industry as a whole directly employing 7.8 million people worldwide, a further 44.7 million jobs were supported indirectly, and 22.4 million jobs were supported through induced effects in the supply chain.

Dr. Reddy pointed out that this data does not take into account the significant economic benefits that medicines and vaccines provide healthcare systems and more broadly to global economies. Just one study demonstrated that adult vaccination programs return 19 times their initial investment.

HIV is one area that clearly shows the remarkable impact pharmaceutical innovation has had on global health. Since the virus that causes AIDS was discovered, more than 30 medicines have been approved to treat HIV infection. With time, medicines have improved in tolerability, efficacy, and convenience for patients. A report from UNAIDS, based on data from 204 countries and territories, found that HIV infections decreased by 22% — from 2.11 million to 1.65 million — between 2010 and 2021, while HIV-related deaths decreased by 40% during the same period, from 1.19 million to 718,000.

A recent study suggests the early impact of HPV vaccination on cervical cancer deaths, observing a substantial reduction in mortality — a 62% drop in cervical cancer deaths among women under age 25 over the last decade in the US, where the HPV vaccine is recommended since 2006.

To sustain innovation for a healthier future, Dr. Reddy cautioned against taking the continued translation of scientific progress into the next generation of medicines and vaccines for granted. He said that underscoring the importance of creating a system that encourages investment and collaboration is needed for without which such innovation would simply not happen.

Fuente: Business World. Disponible en <https://acortar.link/5rl4PL>

Penmenvy FDA Approval History

Feb 17. Penmenvy (meningococcal groups A, B, C, W, and Y vaccine) is a vaccine indicated for active immunization to prevent invasive disease caused by *Neisseria meningitidis* serogroups A, B, C, W, and Y in individuals 10 through 25 years of age.

- ◆ FDA Approved: Yes (First approved February 14, 2025).
- ◆ Brand name: Penmenvy.
- ◆ Generic name: meningococcal groups A, B, C, W, and Y vaccine.
- ◆ Dosage form: Lyophilized Powder for Injection.
- ◆ Company: GlaxoSmithKline.
- ◆ Treatment for: Meningococcal Disease Prophylaxis.



Penmenvy combines the antigenic components of GSK's two well-established meningococcal vaccines: Bexsero (MenB vaccine) and Menveo (MenACWY vaccine). Penmenvy is a pentavalent MenABCWY vaccine that provides broad serogroup coverage in one vaccine, reducing the total number of injections required for protection against invasive meningococcal disease.

Invasive Meningococcal Disease (IMD) is an uncommon but serious illness that can lead to death for up to one in six of those who contract it in as little as 24 hours from onset, despite treatment. Approximately one in five survivors may experience long-term consequences such as brain damage, amputations, hearing loss, and nervous system problems. Adolescents and young adults between the ages of 16 and 23 years are at

high risk due to common behaviors that help transmit the bacteria that cause IMD such as living in close quarters like college dormitories.

Penmenvy protects against invasive meningococcal disease by complement-mediated antibody-dependent killing of *Neisseria meningitidis* serogroups A, B, C, W, and Y.

FDA approval of Penmenvy was supported by positive results from two phase III trials (NCT04502693 and NCT04707391) which evaluated the vaccine's safety, tolerability, and immune response in over 4,800 participants aged 10-25 years.- Bexsero was first approved in 2015 for the prevention of IMD caused by *Neisseria meningitidis* serogroup B in individuals aged 10 through 25 years.- Mencevo was first approved in 2010 for the prevention of IMD caused by *Neisseria meningitidis* serogroups A, C, Y, and W in individuals aged from 2 months through 55 years of age.

Penmenvy is administered by intramuscular injection as two doses, spaced 6 months apart.

Warnings and precautions associated with Penmenvy include syncope (fainting).

Commonly reported ($\geq 10\%$) solicited adverse reactions after Dose 1 and Dose 2, respectively in individuals aged 10 through 25 years include pain at the injection site (92% and 88%), fatigue (51% and 42%), headache (42% and 36%), myalgia (15% and 12%), nausea (15% and 10%), erythema (13% and 12%), and swelling (13% and 12%). Commonly reported ($\geq 10\%$) solicited adverse reactions after Dose 1 and Dose 2, respectively in MenACWY conjugate vaccine-experienced individuals aged 15 through 25 years include pain at the injection site (80% and 74%), headache (41% and 33%), fatigue (40% and 33%), myalgia (15% and 13%), and nausea (15% and 12%).

Fuente: Drugs.com. Disponible en <https://acortar.link/Pk3nHE>

Vaccination against human papillomavirus to be carried out for the first time in Cuba

Feb 18. Cuba is today strengthening the preparation of health sector specialists to support the introduction in the country, for the first time, of the vaccine against the human papillomavirus (HPV).

Initially, it will be applied to a universe of 68,524 nine-year-old girls, taking into account the following scheme: one dose of 0.5 milliliters intramuscularly, and two doses in girls diagnosed with an immunodeficient disease, explained Dr. Lena Lopez Ambron, head of the National Immunization Program of the Ministry of Public Health (Minsap).

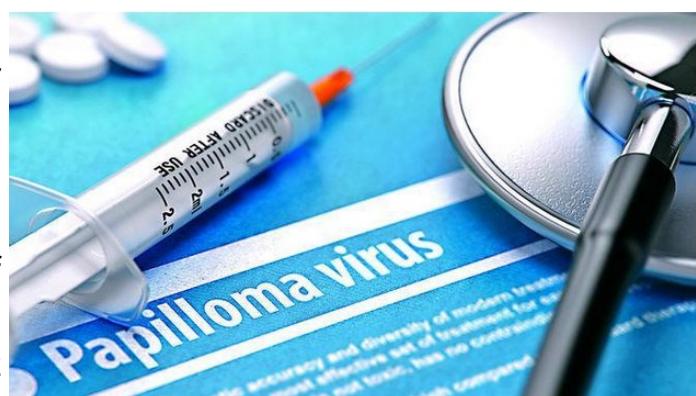


Photo: Prensa Latina

According to a report by Prensa Latina, the coverage objective is equal or higher than 95%, and the goal is to achieve the elimination of HPV-related cervical cancer, with less than four cases in 100,000 women per year, she explained.

The arrival of the immunobiological will be possible thanks to the joint efforts of the Global Alliance for Vaccines and Immunization, the Minsap, and the Pan American Health Organization/World Health Organization (PAHO/WHO), according to the national workshop on cervical cancer control held recently.

According to Dr. Miguel González, advisor at the PAHO/WHO Representation on the island, vaccination against HPV is one of the measures that are part of the global initiative for the elimination of cervical cancer,

together with early detection and access to treatment.

By 2030, the global goal is to vaccinate 90% of girls, detect lesions early in 70% of women, and treat 90% of patients, he said.

Data from the National Cancer Registry show that, in Cuba, one out of every five people dies of cancer, and one out of every three people who die between 30 and 69 years of age is due to this cause.

According to that source, more than 53,000 cases are diagnosed annually, and the trend is increasing, while the number of deaths from this cause, although it has not increased, has not decreased over the years.

Fuente: Granma. Disponible en <https://acortar.link/oaxuig>

Challenges mount for vaccine makers

Feb 19. End-of-year earnings reports reveal drugmakers are grappling with declining vaccination rates and an unpredictable Trump administration.

Vaccine makers are up against unique headwinds. On the one hand, technologies like mRNA have offered breakthroughs for tackling infectious diseases. But at the same time, market uptake and sales growth for several new shots have been sluggish in the last year.

Now, with Robert F. Kennedy Jr., a long-time vaccine critic, sworn in as the new head of the U.S. Department of Health and Human Services, drugmakers are also facing fresh regulatory uncertainty.

Recent earnings reports from the space's top players underscore this challenging environment.

But while drugmakers face issues like increased vaccine skepticism, they're also keeping the R&D wheels turning and gearing up for potential wins in 2025. Here's what end-of-year reports from some of the largest vaccine makers reveal about the industry's shifting winds.

The challenging RSV and flu landscape

Vaccination rates are falling broadly across disease areas.

Among young children, the divide in vaccination rates has fallen along political lines, with kindergartners from states where President Trump won the popular vote showing higher rates of vaccine exemptions, The New York Times reported in January.

Flu vaccination rates for the 2024-2025 season are also well below those from the previous three years, according to the Centers for Disease Control and Prevention data. The trend is happening while the U.S. faces its worst flu seasons since 2009, with at least 29 million illnesses, according to the CDC.

Yet, not every company has been hit by the downturn in flu shots. Sanofi, the largest influenza vaccine maker, reported that global sales for its flu jab were slightly down for the year, although 2024 was a particularly strong year for comparison, CEO Paul Hudson said during the company's earnings call.



Sanofi's vaccine portfolio was also buoyed by uptake in what's become a complex market for RSV.

Overall, the company's vaccine sales rose 13.5% during the year, mostly driven by European sales of Beyfortus, its RSV antibody approved for babies.

The new shot is among a swath of new RSV drugs that have debuted over the past few years, including a handful approved in the U.S. since 2023. But the breakthrough hasn't led to runaway sales for drugmakers amid the declining vaccination trend.

GSK's Arexvy, which became the first RSV vaccine approved in 2023 and protects older adults, reached about \$734 million in sales in 2024, down 52% from the previous year, according to the company's year-end earnings report.

GSK placed some of the blame on the CDC after the agency's Advisory Committee on Immunization Practices changed its recommendations on who should get vaccinated against RSV last year, potentially reducing the patient population.

Still, the pharma company was bullish on Arexvy's future, with revaccination and age cohort expansion expected down the road.

"We really are in the foothills of this vaccine," said Emma Walmsley, CEO of GSK, during the fourth quarter earnings call.

In addition, Arexvy dominated the RSV space in 2024, holding onto about 58% of market share for the year, said Luke Miels, GSK's chief commercial officer.

"I think we are happy with that," he said. "The key point is to preserve value and position ourselves for the future, when we do think ACIP ultimately will move to expand this population."

GSK is aiming to hold onto its market control over Pfizer, which also saw sales of its RSV vaccine for older adults, Abrysvo, decline last year. In fact, Abrysvo sales plummeted 62% year over year in the fourth quarter alone. But Pfizer executives also noted during the company's year-end earnings call that the drug notched a 13% increase in market share during the year.

Moderna launched its RSV vaccine mResvia last year, marking the biopharma's second approved product. However, full-year sales were just \$25 million. The vaccine is the third RSV shot approved for older adults, and like Pfizer and GSK, the company is bullish on its prospects.

"While early RSV sales were limited, we see long-term opportunity to expand our presence in this market both in the U.S. and internationally," CFO Jamey Mock said during the earnings call this month.

Regulatory upheaval

Now at the helm of HHS, Kennedy has wide-ranging ability to interfere with standardized vaccine policies, such as changing the role of ACIP, which advises the CDC on the use of vaccines.

Pfizer CEO Albert Bourla said he was "cautiously optimistic" about Kennedy's leadership position during the earnings call and noted he was looking forward to working on areas of agreement, such as chronic disease.

"The president introduced me to him, and we had dinner all three together, and we tried to understand his view," he said. "Do I expect that we will agree on everything on vaccines? I don't know. But I think probably ... he will have a way more tempered view on how to interact with the vaccines. I think there are a lot of opportunities that probably outweigh the risks that we have with the radical change that... we are seeing now with the Trump administration."

Moderna CEO Stephane Bancel also offered some insight into how the company views the administration's new leadership during the year-end earnings call.

"We look forward to working with the new team as they get confirmed by the Senate," Bancel said. "Vaccines are a very important piece of keeping people healthy, and we look forward to having those discussions as people get confirmed."

Vaccine developments in the coming year

Looking ahead, drugmakers are intent on expanding their RSV base and are still funneling R&D dollars into new vaccines.

Moderna is on the cusp of potentially snagging approvals for three new vaccines this year, including its next-gen COVID vaccine, an RSV jab for high-risk adults between 18-59 and a flu-COVID combo shot for people 50 and older.

With upcoming PDUFA dates in May and June for the new COVID and RSV shots, the biopharma has a lot on the line over the next several months. Moderna is also in a pivotal phase 3 study for a cytomegalovirus vaccine candidate and a two-season pivotal phase 3 study for a norovirus vaccine.

Elsewhere, Sanofi reported it moved forward with six new vaccine studies during the fourth quarter of 2024, including a phase 3 study for a candidate in pneumococcal in children. And GSK noted its pipeline includes plans to expand its shingles vaccine label during the first half of 2025.

Fuente: Pharma Voice. Disponible en <https://goo.su/zahzhLH>

PMA questions delay in dengue vax approval in Philippine

Feb 20. The Philippine Medical Association (PMA) has urged President Ferdinand Marcos Jr. to intervene and expedite the approval of new-generation dengue vaccines amid the surge of dengue cases in the country. The appeal was made by the country's doctors at the launching of the Empowering Networks to Defeat Dengue or End Dengue Coalition — founded by PMA, in partnership with healthcare organizations, the academe, researchers, and advocates — on Tuesday in Quezon City, with the collective goal of achieving zero dengue deaths by 2030.



December 04, 2017 Empty vials of dengue vaccine Dengvaxia injected to the elementary students of Manila are collated at the Manila Central Vaccine storage room in Sta Cruz Manila. Sanofi Pasteur Philippines producer of Dengvaxia says that the dengue vaccine does not contain viruses that can make people ill with dengue or severe dengue. —File photo by Edwin Bacasmas.

In a letter addressed to Marcos read during the event by Dr. Erica Tania Davillo, chair of the PMA's ad hoc committee on dengue advocacy, the organization acknowledged that the safety of dengue vaccines was of "utmost concern" because of the country's previous experience with Dengvaxia.

"[But] with recent advances in vaccine technology, there are new-generation dengue vaccines in the market or in Phase 3 clinical trials, which showed promise in terms of vaccine effectiveness and safety," it added.

Next-gen vaccines

The PMA specifically cited Qdenga of Japanese pharmaceutical company Takeda, which is approved in 40 countries and one of the only two dengue vaccines approved by the World Health Organization (WHO) to prevent dengue in highly endemic countries like the Philippines. (The other WHO-approved dengue vaccine is Dengvaxia of French pharmaceutical giant Sanofi Pasteur.)

"Having witnessed the devastating impact of dengue on our patients and their families, we strongly urge the government to grant access to these new-generation dengue vaccines and allow Filipinos the right to protect themselves from this dreadful disease," the PMA said.

Dr. Lulu Bravo, executive director of the Philippine Foundation for Vaccination, expressed concern over the delay of the approval of the certificate of public registration of Qdenga, noting that vaccine licensing typically takes about a year.

"If you recall, Dengvaxia was licensed within six months, but Qdenga has been in the process for two years with no approval yet," she added, noting that studies on the safety and efficacy of new-generation vaccine have been ongoing for over a decade.

Food and Drug Administration (FDA) spokesperson Pamela Sevilla declined to comment on the status of Takeda's application for Qdenga, which it filed in 2023.

"The FDA cannot disclose any personal or sensitive information pertaining to any brand or product unless otherwise allowed through a letter by the company owner. Rest assured that we will inform the public of any outcome soon," she said in a message to the Inquirer.

Last month, the Union of Local Authorities of the Philippines president and Quirino Gov. Dax Cua also expressed impatience over why Qdenga was still not approved by the FDA.

"The dengue vaccine, which is already being used by our neighboring countries, would be a great help in addressing the threat of dengue. That's why I hope we can approve its use as soon as possible," Cua said.

Health Secretary Teodoro Herbosa previously explained that even if Qdenga is cleared by the FDA, it could not yet be used for mass vaccination by the government without approval from the Health Technology Assessment Council.

WHO-cleared

The WHO in May last year cleared Qdenga for use in children aged 6 to 16 in areas with high infection rates. The prequalification made it eligible for procurement through United Nations agencies like the UN Children's Fund.

The vaccine, which contains weakened versions of the four dengue virus strains, is recommended as a two-dose schedule with a minimum interval of three months between doses.

Fuente: INQUIRER.NET. Disponible en <https://goo.su/YUjwl4F>

VaxLab, Duke-NUS Study Vaccine Challenges in Asia

Fab 21. Asian countries that graduate from donor funding programs face unique challenges in introducing new vaccines to national immunization programs, a new report from the Innovation Lab for Vaccine Delivery Research(VaxLab) finds.

The report, which is available on the website of the Asia Pacific Observatory on Health Systems and Policies (APO), highlights the disparities in vaccine coverage and inclusion of vaccines in national immunization programs in 13

Asian countries. While vaccine coverage rates were high in all countries, middle-income countries that are no longer eligible for donor funding through organizations such as GAVI, the Global Vaccine Alliance, covered the fewest vaccine-preventable diseases in their national immunization programs.

Policymakers, researchers and leaders of international organizations from across Asia discussed the report's findings at a workshop on Feb. 10 in Singapore, co-hosted by the SingHealth Duke-NUS Global Health Institute and the Asia Pacific Immunization Coalition.

"Funding remains a major concern for sustaining immunization efforts," said Xinyu Zhang, Ph.D., a research assistant professor of global health at Duke Kunshan University and former postdoctoral researcher at the Duke Global Health Institute, who was the lead author of the report. "GAVI-eligible countries rely heavily on external funding, while GAVI-ineligible countries depend on domestic financing. Countries that graduate from GAVI support face a 'funding cliff,' which poses a significant challenge to maintaining immunization rates."

While most countries have achieved over 90 percent coverage for routine vaccinations, newer vaccines such as human papillomavirus (HPV) vaccine, pneumococcal conjugate vaccine and rotavirus vaccine show lower coverage rates, according to the report.

"The findings from this report underscore the urgent need for sustainable immunization systems and stronger regional collaboration in Asia," said Shenglan Tang, M.D., Ph.D., director of VaxLab and co-director of the Global Health Research Center at Duke Kunshan University.

To address these challenges, workshop participants discussed key areas for strengthening national immunization programs, emphasizing the need for sustainable financing and increased domestic investment. Innovative financing mechanisms and regional collaboration on cost-sharing and procurement were highlighted as essential to maintaining long-term immunization efforts.

Participants stressed the importance of building public trust through transparent communication and involving communities as active partners. The workshop also underscored the need to invest in robust monitoring and evaluation systems to support evidence-based decision-making and efficient resource allocation.

Fuente: Duke Global Health. Disponible en <https://goo.su/zApzwSK>



Participants at a Feb. 10 workshop in Singapore to discuss challenges in vaccine coverage across 13 Asian countries.

Study reveals the benefits of new vaccine

Feb 23. A new study has revealed that there has been a 62% reduction in respiratory syncytial virus (RSV) related hospitalisations among the eligible age group following the introduction of the vaccine.

The study, conducted by Public Health Scotland (PHS) in collaboration with the University of Strathclyde, and published in *The Lancet Infectious Diseases*, concludes that the RSV vaccine is highly effective in reducing hospitalisations in older adults.

RSV is a common and highly infectious respiratory virus that affects the breathing system and can be very serious for those who are at the highest risk of serious illness, including older adults.



Scotland's new RSV vaccination programme was launched last August, with local health boards inviting adults aged 75-79, including those turning 75 before July 2025, to come forward for their free vaccine ahead of winter. By the end of November, uptake of the vaccine in this older adult population had reached 68%.

This study is the first to evidence the positive impact of the vaccination programme in reducing hospitalisations and underlines the importance of older adults coming forward for their vaccine. One dose offers multi-year protection and the study's results show that, in the first year alone, the programme has reduced serious illness among older adults.

Dr Sam Ghebrehewet, head of immunisation and vaccination at PHS, said: "The success of the RSV programme marks another significant step in protecting the population of Scotland against preventable diseases.

"Public Health Scotland continues to work closely with local health boards to ensure as many people as possible receive their vaccine. As well as being offered to older adults, the vaccine is also offered during pregnancy. Getting vaccinated is the best and simplest thing you can do to protect yourself, or your newborn baby, against serious illness caused by RSV." Neil Gray, Cabinet Secretary for Health and Social Care, added: "We were pleased to be the first nation in the UK to introduce the new RSV vaccine in time to maximise the benefit to the more vulnerable ahead of winter. This research demonstrates just how many people avoided ending up in hospital as a result."

Fuente: Angus County World. Disponible en <https://goo.su/l4vwHL>

Cervical Cancer Elimination Continues in the Americas in 2025

Feb 24. The Pan American Health Organization (PAHO) and the Spanish Agency for International Development Cooperation recently formalized a memorandum of understanding to reinforce their shared commitment to improving public health across the Americas.

Announced on February 21, 2025, the memorandum covers various areas of cooperation aligned with the United Nations' Sustainable Development Goals, with a key focus on eliminating cervical cancer. Each year, cervical cancer claims the lives of approximately 40,000 women in the Americas.

PAHO Director Jarbas Barbosa emphasized the significance of this collaboration for the region.

"Spain's support in the fight against cervical cancer and other public health areas is critical for strengthening the health systems. With AECID's support, we can make progress toward the elimination of this cancer, which disproportionately impacts women in the most vulnerable situations," said Dr. Barbosa in a press release.

PAHO is leading efforts to eliminate cervical cancer in the Americas, aligning its actions with the World Health Organization's 90-70-90 targets.

These aim for 90% of girls fully vaccinated with the HPV vaccine by age 15, 70% of women screened using a high-performance test by age 35 and 45, 90% with pre-cancer treatment, and 90% with invasive cancer managed.

In the Americas, the PAHO recommends vaccinating against human papillomavirus (HPV). However, only 48 countries have introduced an HPV vaccine, and coverage rates vary widely.

Few countries have reached 90% HPV vaccination coverage, while others remain below 10%.

The PAHO's Revolving Fund announced on February 7, 2025, that the general availability of the 9-valent HPV vaccine will be easier and more affordable for Latin American countries. Health agencies are debating HPV dosage protocols.

The PAHO stated it is working to close these gaps, ensuring access to safe and affordable vaccines and diagnostic tools through its Regional Revolving Funds.

Fuente: VAX BEFORE TRAVEL. Disponible en <https://n9.cl/778jva>

BTIG mantiene la calificación de Compra para las acciones de Vaxcyte, con un objetivo de 160 dólares

Feb 24. BTIG reafirmó su postura positiva sobre las acciones de Vaxcyte (NASDAQ:PCVX), manteniendo una calificación de Compra y un precio objetivo de 160,00 dólares. La empresa, con una capitalización bursátil de aproximadamente 10.000 millones de dólares, goza de un fuerte apoyo de Wall Street, con objetivos de los analistas que oscilan entre 135 y 163 dólares. Según datos de InvestingPro, Vaxcyte mantiene una calificación de consenso de los analistas muy favorable de 1,2, lo que indica fuertes recomendaciones de compra en general.



La firma se centra en los próximos datos provisionales del ensayo de Fase 2 de VAX-24, el candidato a vacuna neumocócica de Vaxcyte para bebés. Aunque no se espera que se revelen datos en la conferencia de resultados trimestrales del martes, los analistas consideran que este ensayo es una evaluación significativa de la plataforma de Vaxcyte. El análisis de InvestingPro muestra que la empresa mantiene una puntuación global JUSTA de salud financiera, con puntuaciones particularmente fuertes en impulso de precio y gestión de flujo de caja. Los suscriptores pueden acceder a 6 ProTips adicionales y métricas financieras completas para evaluar mejor el potencial de inversión de Vaxcyte.

Los analistas prevén que VAX-24 pueda superar a las vacunas existentes al fallar en menos serotipos después de las tres primeras dosis administradas a los niños. El rendimiento después de tres dosis es particularmente importante porque indica el nivel de protección que tienen los niños antes de recibir una cuarta dosis de refuerzo. En comparación, PREVNAR20, una vacuna actualmente aprobada, falló en seis serotipos después de tres dosis, mientras que se espera que VAX-24 falle en dos como máximo, con la posibilidad de no fallar en ninguno.

La dirección de Vaxcyte ha indicado que la proporción de serotipos en VAX-31 para bebés refleja la de la formulación para adultos, lo que sugiere una fuerte traducción de la inmunogenicidad de adultos a bebés. Pequeños ajustes en las proporciones de serotipos en VAX-31 tienen como objetivo mejorar la eficacia de la vacuna aprovechando la respuesta inmunitaria de los serotipos más fuertes.

De cara al futuro, Vaxcyte planea iniciar un estudio de no inferioridad de Fase 3 en adultos a mediados de 2025, con datos principales previstos para 2026 y una posible entrada en el mercado en 2027. La sólida posición financiera de la empresa se evidencia por su robusto ratio de liquidez de 17,88 y un mínimo ratio de deuda sobre capital de 0,01, lo que sugiere recursos suficientes para financiar sus programas de desarrollo clínico. El candidato VAX-31 de la empresa ha demostrado una alta inmunogenicidad y un perfil de seguridad favorable, posicionándolo como un fuerte competidor para el mercado de vacunas neumocócicas para adultos una vez aprobado.

La reciente recomendación del Comité Asesor sobre Prácticas de Inmunización (ACIP) de reducir la edad recomendada para las vacunas conjugadas neumocócicas a 50 años o más podría ampliar significativamente el mercado total al que puede dirigirse VAX-31.

Se espera que los datos finales de las tres primeras dosis de VAX-24 estén disponibles a finales del primer trimestre de 2025, con resultados de la dosis de refuerzo previstos para finales de año. Además, se proyectan resultados principales del estudio de Fase 2 de VAX-31 en bebés para mediados de 2026, seguidos de datos de inmunogenicidad de la dosis de refuerzo nueve meses después.

En otras noticias recientes, Vaxcyte ha anunciado el avance a la etapa final de su estudio de Fase 2 para VAX-31, un candidato a vacuna conjugada neumocócica. Esta etapa evaluará aún más la seguridad, tolerabilidad e inmunogenicidad de la vacuna en bebés sanos, y la empresa planea publicar datos principales de la serie de inmunización primaria a mediados de 2026. Además, Goldman Sachs ha iniciado la cobertura de Vaxcyte con una calificación de Compra, citando fuertes datos clínicos para sus programas principales como un indicador positivo para el futuro de la empresa. La firma ha establecido un precio objetivo de 135,00 dólares, destacando el potencial de VAX-31 y otros activos en desarrollo para impulsar el crecimiento a largo plazo.

La plataforma tecnológica patentada de Vaxcyte ha sido fundamental para superar los desafíos tradicionales en el desarrollo de vacunas, posicionando a la empresa como un actor significativo en el mercado de vacunas neumocócicas. Los esfuerzos de la empresa para abordar los desafíos de las vacunas de próxima generación han sido bien recibidos, y Goldman Sachs señala la reducción de riesgos de VAX-31 como un factor clave en su potencial regulatorio y comercial. Estos desarrollos subrayan el compromiso de Vaxcyte de avanzar en su cartera de vacunas y expandir su presencia en el mercado.

Fuente: Investing.com. Disponible en <https://n9.cl/0m474c>

IMUNON Announces New Immunogenicity Data from Phase 1 Clinical Trial of Its DNA Vaccine in Treatment of COVID-19

Feb 26. IMUNON, Inc. (NASDAQ: IMNN), a clinical-stage company focused on developing non-viral DNA-mediated immunotherapy and evaluating an adaptation of the platform's potential as a next-generation vaccine, today announced new safety and immunogenicity data from ongoing analyses of results from the Company's first Phase 1 proof-of-concept clinical trial of IMNN-101, its investigational DNA plasmid vaccine based on the Company's proprietary PlaCCine® technology platform. The Phase 1 study was conducted in 24 healthy volunteers as a seasonal COVID-19 vaccine, targeting the SARS-CoV-2 Omicron XBB1.5 spike antigen. IMNN-101 was administered as a single dose vaccine without a booster dose in study participants who were previously vaccinated against the Omicron XBB1.5 variant. Results demonstrated that IMNN-101 is safe and well-tolerated with no serious adverse effects. IMNN-101 induced a persistent 2- to 4-fold increase in serum neutralizing antibody (NAb) titers from baseline through Week 4, further increasing NAb titers between Week 2 and Week 4. The immune response was observed against the XBB1.5 variant and many newer variants following treatment, demonstrating the IMNN-101 vaccine's cross-reactivity.

"We have strong evidence of vaccine immunogenicity based on the neutralizing antibody response against the Omicron XBB.1.5 strain in this trial, and expect partnering interest in our proof-of-concept data from the PlaCCine platform," said Stacy Lindborg, Ph.D., president and chief executive officer of IMUNON. "These data demonstrate that our first-in-human vaccine based on our PlaCCine platform is safe and immunogenic and is well-suited to developing vaccine candidates for protecting the population against a potential future exposure to a pathogen or controlling a rising pathogen. Given proof of immunogenicity, early indications of durability of protection, and competitive advantages in the stability of our vaccine at workable temperatures compared with available mRNA vaccines, we believe that IMNN-101 has significant potential as a superior next-generation vaccine and will seek potential partners for further development."

The participants in the Phase 1 trial had high baseline immune characteristics presumably from prior infection and multiple previous vaccinations against COVID-19 and ongoing infection as evidenced by the rise in viral nucleocapsid antigen during the study period. Modest increases in T cell responses were observed in this setting of trial participants having received multiple immunizations prior to the study.

"Data from this trial is of high quality and show that IMUNON's DNA vaccine is immunogenic in humans. Following immunization, participants' NAb titers increased through Week 4 with a 2- to 4-fold increase from baseline, a clear and convincing response to the vaccination," said Ai-ris Collier, M.D., Co-Director of the Clinical Trials Unit, Center for Virology and Vaccine Research Center, Beth Israel Deaconess Medical Center.

The Phase 1 clinical data of IMNN-101 is consistent with strong evidence of immunogenicity and protection for the PlaCCine platform in rodents and non-human primates, with prior preclinical results showing that protection exceeded 95% in non-human primates, which is comparable to mRNA vaccines. The robust immunogenicity profile, expected durability of protection, comparative ease of manufacturing, and stability at workable temperatures (up to one year at 4°C and one month at 37°C) suggest that our vaccine based on the PlaCCine technology platform may be a potential viable alternative to available messenger RNA (mRNA) vaccines.

About PlaCCine® and IMNN-101

IMNN-101 utilizes the company's PlaCCine® technology platform, a proprietary composition of a DNA

plasmid that regulates the expression of key pathogen antigens and a novel synthetic DNA delivery system. The plasmid-based expression vector accommodates single or multiple antigens through its flexible vector design, offers manufacturing flexibility compared to with viral or other DNA or protein vaccines, and the synthetic delivery system protects DNA from degradation and facilitates DNA uptake after injection with acceptable safety.

About the Phase 1 PoC Clinical Trial

This U.S. Phase 1 proof-of-concept (PoC) study inoculated 24 participants to evaluate three escalating doses of IMNN-101 with eight participants at each dose. All participants were treated at DM Clinical Research in Philadelphia. For this study, IMNN-101 has been designed to protect against the SARS-CoV-2 Omicron XBB1.5 variant, in accordance with the FDA's Vaccines and Related Biological Products Advisory Committee's June 2023 announcement of the framework for updated COVID-19 doses. The primary objectives of the study are to evaluate safety and tolerability in healthy adults. Secondary objectives include evaluating IMNN-101's ability to elicit neutralizing antibody responses, cellular responses and their associated durability.

About IMUNON

IMUNON is a clinical-stage biotechnology company focused on advancing a portfolio of innovative treatments that harness the body's natural mechanisms to generate safe, effective and durable responses across a broad array of human diseases, constituting a differentiating approach from conventional therapies. IMUNON is developing its non-viral DNA technology across its modalities. The first modality, TheraPlas®, is developed for the coding of cytokines and other therapeutic proteins in the treatment of solid tumors where an immunological approach is deemed promising. The second modality, PlaCCine®, is developed for the delivery of DNA-coded viral antigens that can elicit a strong immunological response. This technology may represent a promising platform for the development of vaccines in infectious diseases.

The Company's lead clinical program, IMNN-001, is a DNA-based immunotherapy for the localized treatment of advanced ovarian cancer currently in Phase 2 development. IMNN-001 works by instructing the body to produce safe and durable levels of powerful cancer-fighting molecules, such as interleukin-12 and interferon gamma, at the tumor site. Additionally, the Company has entered a first-in-human study of its COVID-19 booster vaccine (IMNN-101). We will continue to leverage these modalities and to advance the technological frontier of plasmid DNA to better serve patients with difficult-to-treat conditions. For more information on IMUNON, visit www.imunon.com.

Fuente: First Word PHARMA. Disponible en <https://n9.cl/gm1du>

Brasil anuncia su primera vacuna contra el dengue y proyecta producir 60 millones de dosis anuales

26 feb. El Gobierno de Brasil anunció este martes el desarrollo de la primera vacuna contra el dengue producida íntegramente en el país, un avance clave en la lucha contra esta enfermedad que ha alcanzado cifras récord en los últimos años. A partir de 2026, la producción alcanzará las 60 millones de dosis anuales, destinadas a ser distribuidas en la red de salud pública.

La vacuna, de dosis única y con eficacia comprobada contra los cuatro serotipos del virus del dengue, será elaborada por el Instituto Butantan, un prestigioso centro de investigación biomédica en Brasil, en alianza con la empresa china WuXi Biologics. El proyecto cuenta con una inversión inicial de 1.260 millones de reales (aproximadamente 221 millones de dólares).

El anuncio fue realizado en una ceremonia oficial encabezada por el presidente brasileño, Luiz Inácio Lula da Silva, y la ministra de Salud, Nísia Trindade. La funcionaria presentó el proyecto en un contexto político complejo, marcado por especulaciones sobre una posible reforma del gabinete en la que podría perder su cargo.

Una vacuna clave para enfrentar la crisis sanitaria

El dengue es una enfermedad viral transmitida por el mosquito *Aedes aegypti* y se ha convertido en un problema de salud pública en Brasil y otros países tropicales. En 2024, Brasil alcanzó una cifra récord de 6,65 millones de casos probables y registró 6.022 muertes, la cantidad más alta de decesos desde que se tiene registro.

Si bien en lo que va de 2025 se ha registrado una reducción del 30 % en los casos probables con respecto al mismo período del año anterior (402.000 casos hasta la fecha), algunas regiones continúan en situación crítica. Es el caso del estado de São Paulo, que declaró emergencia sanitaria tras reportar 200.000 casos probables entre enero y febrero, con 102 muertes confirmadas y otras 225 en investigación.

La crisis del dengue no se limita a Brasil. En países vecinos como Perú, la situación también es alarmante. En 2023, el país registró más de 270.000 casos y 423 muertes, convirtiéndose en la peor epidemia de dengue en su historia reciente.

El cambio climático, con temperaturas más altas y lluvias intensas, ha contribuido a la proliferación del mosquito transmisor, afectando especialmente al norte del país. La llegada de la nueva vacuna brasileña podría representar una oportunidad para Perú de fortalecer su estrategia de inmunización y mitigar los efectos de futuros brotes.

Un esfuerzo de inmunización sin precedentes

Para combatir la propagación del dengue, Brasil implementó en 2024 una campaña de vacunación con la vacuna desarrollada por la farmacéutica japonesa Takeda, conocida como Qdenga. Sin embargo, debido a la disponibilidad limitada de dosis, solo 3,3 millones de niños entre 10 y 14 años pudieron ser inmunizados.

Este año, el Ministerio de Salud ha adquirido nueve millones de dosis adicionales de la vacuna japonesa, con la posibilidad de incrementar la cantidad en el segundo semestre. No obstante, la producción nacional de la nueva vacuna de Butantan permitirá a Brasil contar con una fuente propia y masiva de inmunización, lo que representa un paso decisivo en la lucha contra el dengue.

La introducción de esta vacuna refuerza la estrategia de salud pública del país para frenar la incidencia de la enfermedad, reducir la mortalidad y mitigar el impacto del dengue en el sistema sanitario.

Con este avance, Brasil se posiciona a la vanguardia en la producción de vacunas contra enfermedades tropicales y fortalece su autonomía en la fabricación de inmunizantes clave para su población. Además, este desarrollo puede beneficiar a otros países de la región, como Perú, que podrían acceder a las dosis brasileñas para enfrentar su propia crisis sanitaria y mejorar la respuesta ante brotes futuros.



Fuente: Salud con Lupa. Disponible en <https://n9.cl/eyv57>

La vacuna contra la COVID-19 del CSIC que se cedió a la OMS aún no se ha probado en humanos

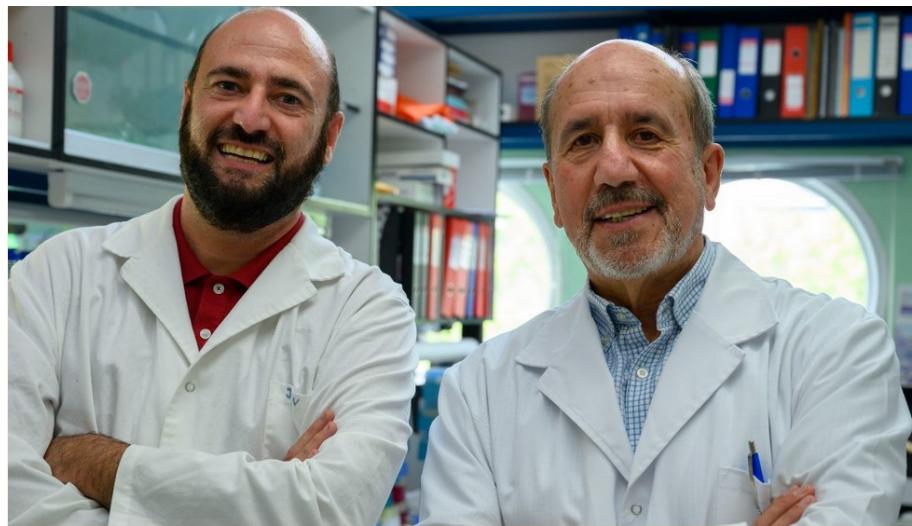
27 feb. En agosto de 2023, el Consejo Superior de Investigaciones Científicas (CSIC) transfirió a la Organización Mundial de la Salud (OMS) la patente de su prototipo de vacuna contra la COVID-19. Se trata de la inyección desarrollada por los prestigiosos investigadores Juan García Arriaza y Mariano Esteban, del Centro Nacional de Biotecnología (CNB-CSIC). Los estudios preclínicos demostraron una eficacia del 100% en modelos animales, previniendo la replicación del virus en pulmones y cerebro y evitando daños asociados. Sin embargo, un año y medio después de la cesión a la OMS y de cinco del inicio de la pandemia del coronavirus, desde el CSIC indican a THE OBJECTIVE que esta aún no ha iniciado sus ensayos clínicos en humanos.

En marzo de 2020, el Gobierno concedió 4,5 millones de euros al CSIC para el desarrollo de vacunas contra el coronavirus. En total, fueron tres inyecciones las que los científicos del Consejo Superior de Investigaciones Científicas pusieron en marcha. La de los investigadores Arriaza y Esteban es la única de las tres vacunas para la cual se llegó a pedir autorización a la Agencia Española del Medicamento (Aemps) para iniciar ensayos en humanos, en 2021. De hecho, en mayo de ese año, el Gobierno otorgó al CSIC 2,4 millones de euros para llevar a cabo los ensayos clínicos de fase I/II combinado en los que participarían varios hospitales españoles. Sin embargo, la autorización de la Aemps para el comienzo de estos estudios médicos en personas nunca llegó y, finalmente, en agosto de 2023, el CSIC –organismo dependiente del Ministerio de Ciencia e Innovación– decidió transferir su vacuna, basada en el virus vaccinia MVA como vector, a la OMS.

Ahora, según ha explicado el propio Mariano Esteban a THE OBJECTIVE, la inoculación «está en proceso de consideración por las autoridades suizas para iniciar un ensayo clínico en 2025 como parte de la concesión de un proyecto europeo (2024-2028), cuyo objetivo es comparar cuatro vacunas frente al SARS-CoV-2 (dos aprobadas por la EMA y las otras dos, una francesa y la nuestra española) desde el punto de vista celular, molecular e inmune, para de esta forma conocer en profundidad las diferencias y ventajas de cada una de las vacunas». La autorización corresponde a las autoridades sanitarias suizas y francesas, ya que el ensayo clínico se realizará en París y Lausanne.

¿Por qué la Aemps no autorizó sus ensayos en humanos?

Es muy confuso por qué la Aemps rechazó autorizar el inicio de los ensayos clínicos en humanos. Sin una versión oficial sobre los motivos que paralizaron el proceso, hubo información contradictoria sobre sus ensayos con macacos. Se publicó que uno de los animales utilizados en las pruebas había sufrido daños; sin embargo, los científicos desmintieron esta información y afirmaron que la vacuna era totalmente segura en los tres modelos animales en que se ha habido probado: ratones, hámsteres y macacos. «La vacuna confiere



Los investigadores Juan García Arriaza y Mariano Esteban. | CSIC

una protección del 100% contra la infección causada por el SARS-CoV-2 en los tres modelos animales, controlando la replicación del virus en las vías respiratorias y en los pulmones, la patología pulmonar y previniendo la tormenta de citoquinas», subrayaron Arriaza y Esteban en un comunicado.

De esta forma, tras no conseguir el espaldarazo de las autoridades sanitarias españolas, el CSIC firmó la cesión de esta vacuna para que llegue a países en desarrollo. La transferencia de esta tecnología se realizó de forma gratuita a través de la iniciativa COVID-19 Technology Access Pool (C-TAP) de la OMS y la organización Medicines Patent Pool (MPP), con el objetivo de facilitar la producción y distribución de la vacuna en países de ingresos bajos y medios.

El CSIC acordó no cobrar regalías por la explotación de la vacuna siempre que se fabrique para estos países.

¿Qué ha pasado con las otras dos vacunas del CSIC?

Las otras dos vacunas contra la COVID-19 desarrolladas por el CSIC son la del químico y virólogo Luis Enjuanes y la del médico e investigador Vicente Larraga. Con respecto a la primera, el propio Enjuanes, en una entrevista con el Colegio Oficial de Médicos de Ciudad Real el pasado mes de enero, indicó que la inyección podrá estar lista a finales de este año. «Tenemos que asegurar que en personas no se producen efectos secundarios. En marzo empezaríamos los ensayos clínicos de fase 1 y 2 y 3 en personas», explica el investigador, que agrega que están preparando dos versiones de la vacuna intranasal e intramuscular para ver cuál es más segura y eficaz.

«La aplicación intranasal tiene gran recelo en las agencias evaluadoras. Aunque nuestra preferencia es esa», indica el experto, que asegura están «acelerando los procesos» debido a que han empezado a colaborar con una gran multinacional. A diferencia de las vacunas que hay ahora, la de Enjuanes es esterilizante, es decir, que bloquearía al virus en la vía de entrada, las mucosas nasales. Las de ahora impiden que la infección avance y la enfermedad se agrave. La suya evitaría la propia infección.

En el caso de la inoculación de Larraga, en 2022 completó con éxito sus pruebas en dos especies animales y se preparaba para iniciar ensayos preclínicos en macacos, con la esperanza de obtener autorización para pruebas en humanos hacia finales de ese año. Sin embargo, hasta la fecha no se han publicado actualizaciones adicionales sobre el avance de esta vacuna en ensayos clínicos o su disponibilidad para el público.

Fuente: THE OBJECTIVE. Disponible en <https://n9.cl/fmpis>

Bavarian Nordic Receives Marketing Authorization in Europe for Chikungunya Vaccine for Persons Aged 12 and Older

Feb 28. Bavarian Nordic A/S (OMX: BAVA) from Denmark, announced that the European Commission has granted marketing authorization in Europe for VIMKUNYA® for active immunization for the prevention of disease caused by chikungunya virus in individuals 12 years and older.

The virus-like particle (VLP) single-dose vaccine is the first chikungunya vaccine approved in Europe for persons as young as 12 years old. The approval, valid in all EU member states, as well as in Iceland, Liechtenstein, and Norway, marks the second approval of VIMKUNYA, following the approval by the U.S. Food and Drug Administration (FDA) earlier this month. Bavarian Nordic also recently submitted a Marketing Authorization Application (MAA) to the UK Medicines and Healthcare products Regulatory Agency (MHRA) with potential approval of the chikungunya vaccine in the UK in the first half of 2025.

Bavarian Nordic will launch VIMKUNYA in key European markets in the first half of 2025.

"We are highly encouraged by the European Commission's accelerated decision to approve our chikungunya vaccine in Europe, which offers a differentiated profile for travelers, including those as young as 12 years," said Paul Chaplin, President and CEO of Bavarian Nordic. "As we expand our presence across Europe, this vaccine will help to further consolidate our leading position in travel vaccines, and we look forward to making the vaccine available in key markets during the first half of 2025."

The marketing authorization was granted by the European Commission upon recommendation by the Committee for Medicinal Products for Human Use (CHMP) of the European Medicines Agency (EMA) in January 2025 and was based on results from two phase 3 clinical trials which enrolled more than 3,500 healthy individuals 12 years of age and older. The studies met their primary endpoints, with results showing that 21 days after vaccination, the vaccine induced neutralizing antibodies in up to 97.8% of the vaccinated individuals and demonstrated a rapid immune response starting to develop within one week.

About chikungunya

Chikungunya is a mosquito-borne disease caused by the chikungunya virus (CHIKV). In the past 20 years, the virus has emerged across several regions in Asia, Africa, and the Americas, including many popular travel destinations, often causing large unpredictable outbreaks. Since its discovery, CHIKV has been identified in more than 110 countries, with evidence of transmission confirmed in more than 50 countries over the past five years¹. Chikungunya typically presents with acute symptoms, including fever, rash, fatigue, headache, and often severe and incapacitating joint pain. Most patients recover, but 30-40% of those affected may develop chronic symptoms that can last for months or even years². In 2024, approximately 480,000 cases of chikungunya and over 200 deaths were reported worldwide³. Recent data suggest that chikungunya is severely underreported and often misdiagnosed as dengue fever due to a similar symptom profile⁴.

About VIMKUNYA® Chikungunya vaccine (recombinant, adsorbed)

VIMKUNYA is an adjuvanted VLP recombinant protein vaccine for active immunization for the prevention of disease caused by chikungunya virus (CHIKV) in individuals 12 years and older. Because VLPs contain no virus genetic material, the vaccine cannot infect cells, reproduce or cause disease. VIMKUNYA will be available as a suspension for injection in a pre-filled syringe.

While the mechanism of action of CHIKV VLP vaccine still needs to be further characterised, it is thought that the vaccine can induce protection from CHIKV infection by inducing neutralising antibodies against certain CHIKV proteins resulting in neutralisation of live virus. An adjuvant is added to increase the magnitude of vaccine-mediated immune responses. The most common side effects are injection site pain, fatigue, headache, and myalgia.

Full product information will be available from:

<https://www.ema.europa.eu/en/medicines/human/EPAR/vimkunya>

About Bavarian Nordic

Bavarian Nordic is a global vaccine company with a mission to improve health and save lives through innovative vaccines. We are a preferred supplier of mpox and smallpox vaccines to governments to enhance public health preparedness and have a leading portfolio of travel vaccines.

For more information, visit www.bavarian-nordic.com

Fuente: First Word PHARMA. Disponible en <https://n9.cl/le92z>

Infant PCV13 Immune Response Blunted by Vaccination in Respiratory Viral Seasons

Feb 28. A blunted immune response was observed among young infants who received their first dose of a pneumococcal 13-valent conjugate vaccine (PCV13) during peak respiratory viral seasons, according to study findings published in *Clinical Infectious Diseases*.



"Infants who receive their first pneumococcal conjugate vaccine (PCV) dose during a respiratory viral season may have increased susceptibility to respiratory viral immune blunting with higher carrier-load PCVs."

Researchers conducted a post hoc analysis of a large, double-blinded, randomized study that compared outcomes of PCV13 vs PCV7 administration in infants at 2, 4, 6, and 12 months of age. The analysis occurred from February 2008 to July 2009 and comprised infants in Israel who received their first PCV dose at 7 to 9 weeks of age either during or outside of a respiratory viral season. The researchers used local epidemiologic data to define the respiratory viral season as December through April and collected blood samples from infants at 7 and 13 months of age (1 month after vaccination) to assess serum anticapsular-binding immunoglobulin (Ig) G antibodies for each serotype included in PCV13. Geometric mean serotype-specific IgG concentrations (GMC) were calculated for statistical analysis.

Among 1058 infants included in the final analysis, 533 received PCV13 and 525 received PCV7. Of 404 PCV13 recipients and 405 PCV7 recipients tested at post-infant series (post-primary) visits, 179 and 188, respectively, were vaccinated during respiratory viral seasons, while 225 and 217, respectively, were not. Across all 4 groups, patient sex, ethnicity, and age at receipt of each dose were similar.

In the PCV13 group, the researchers observed lower GMCs for all serotypes 1 month after completion of the 3-dose series in patients who received the first dose during vs outside of respiratory viral seasons. The between-group difference reached statistical significance for 10 of the 13 serotypes. Similar results were observed among subgroups of PCV13 and PCV7 recipients who received the first 2 vaccine doses during vs outside of respiratory viral seasons.

"The current study provides a unique opportunity to demonstrate both the influence of seasonality and increased carrier load on the immune response of infants to PCVs."

Further analysis of patients in the PCV13 group showed that GMCs after receipt of a booster dose at 13 months of age were higher than those following completion of the 3-dose series, indicating a robust booster response. However, the GMC ratio of PCV administration during vs outside of respiratory viral seasons was lower for 11 of the 13 serotypes and reached statistical significance for 2 serotypes. The researchers noted similar findings in a subgroup analysis of patients who received the first 2 vaccine doses during vs outside of the respiratory viral season.

In the PCV7 group, patients vaccinated during vs outside of respiratory viral seasons exhibited no significant differences in immune response following completion of the 3-dose series at 7 months of age. Moreover, vaccination during a respiratory viral season was associated with higher GMCs following receipt of a booster dose at 13 months of age.

Study limitations include the lack of data regarding pathogen exposure and the inability to rule in or rule out the occurrence of asymptomatic transmission.

According to the researchers, "The current study provides a unique opportunity to demonstrate both the

influence of seasonality and increased carrier load on the immune response of infants to PCVs."

Disclosure: This research was supported by Pfizer Inc. One study author declared affiliations with biotech, pharmaceutical, and/or device companies. Please see the original reference for a full list of disclosures.

Fuente: Infectious Disease ADVISOR. Disponible en <https://n9.cl/zqhib>

Zydus Lifesciences introduces WHO-recommended flu vaccine in 2025

Feb 28. Zydus Lifesciences, a leading, discovery-based, global pharmaceutical company is ready to launch the season's first India's Flu protection as per WHO recommended composition of quadrivalent influenza virus vaccines for use in the 2025 southern hemisphere. The company's Quadrivalent Inactivated Influenza vaccine VaxiFlu-4 will offer protection against:

- ⇒ A/Victoria/4897/2022 (H1N1) pdm09-like virus,
- ⇒ A/Croatia/10136RV/2023 (H3N2)-like virus,
- ⇒ B/Austria/1359417/2021 (B/Victoria lineage)-like virus,
- ⇒ B/Phuket/3073/2013 (B/Yamagata lineage)-like virus.



Zydus Lifesciences quadrivalent vaccine, by covering strains of both influenza A and influenza B, provides a broader protection and significantly reduces the risk of vaccine mismatch. The vaccine has been cleared by the Central Drug Laboratory (CDL).

VaxiFlu-4 is being marketed by Zydus Vaxxicare-a division of the group focussing on preventives. The Quadrivalent Inactivated Influenza vaccine has been developed at the Vaccine Technology Centre (VTC) in Ahmedabad which has proven capabilities in researching, developing, and manufacturing of safe and efficacious vaccines.

Development of Zydus Lifesciences

Speaking on the development Dr. Sharvil Patel, Managing Director, Zydus Lifesciences Limited said, "Preventives are the key to public health in both the developing and the developed world and vaccines have the potential to improve the quality of life. In India, there is a pressing need for access to affordable, high-quality vaccines that can address healthcare challenges. With vaccines like VaxiFlu-4 we are serving the cause of public health through annual immunization and preventing flu outbreaks."

Because of annual and occasional outbreaks, the control of influenza has become a major public health challenge. Annual Influenza (flu) vaccination is the best way to prevent flu and its potentially serious complications. Influenza is a contagious respiratory illness caused by influenza viruses which spreads from person to person, mainly through airborne respiratory droplets generated from coughing and sneezing or direct contact with an infected surface or individual.

It can cause illnesses that range in severity and at times lead to hospitalization and death-with the latter occurring mainly in high-risk groups, such as under-five children, the elderly, and people with immunosuppressive and chronic medical conditions. According to the World Health Organization (WHO), seasonal influenza results in 290,000-650,000 deaths every year.

Fuente: HEALTHCARERADIUS. Disponible en <https://n9.cl/fv9m6>

Moderna Inc (MRNA) Secures UK Approval for RSV Vaccine mRESVIA

Mar 1. Moderna Inc (MRNA, Financial) announced that the UK's Medicines and Healthcare products Regulatory Agency (MHRA) has granted marketing authorization for mRESVIA (mRNA-1345). This vaccine is designed for active immunization to prevent lower respiratory tract disease (LRTD) caused by respiratory syncytial virus (RSV) in adults aged 60 and older. This approval marks Moderna's second product authorized in the UK, underscoring the company's commitment to combating respiratory diseases through mRNA technology.

Positive Aspects

- * mRESVIA is Moderna's second approved product in the UK, highlighting the company's expanding portfolio.
- * The vaccine targets a significant health issue, as RSV is a leading cause of respiratory infections in the elderly.
- * Phase 3 clinical trials showed positive results with no serious safety concerns.
- * The vaccine will be produced at the Moderna Innovation and Technology Centre in Oxfordshire, enhancing local manufacturing capabilities.

Negative Aspects

- * The approval is limited to adults aged 60 and older, potentially restricting the market size.
- * There are inherent risks and uncertainties associated with forward-looking statements regarding vaccine efficacy and safety.

Financial Analyst Perspective

From a financial standpoint, the approval of mRESVIA in the UK represents a strategic expansion of Moderna's product offerings, potentially increasing revenue streams. The successful Phase 3 trial results and the absence of serious safety concerns are likely to bolster investor confidence. However, the market is limited to a specific age group, which may impact the overall financial impact. Investors should also consider the risks associated with forward-looking statements and regulatory challenges in other markets.

Market Research Analyst Perspective

The authorization of mRESVIA in the UK positions Moderna as a key player in the respiratory vaccine market, particularly for the elderly demographic. The high incidence of RSV-related health issues in the UK underscores the demand for effective vaccines. Moderna's focus on local manufacturing at the Oxfordshire facility could enhance supply chain efficiency and responsiveness to market needs. However, the competitive landscape and regulatory hurdles in other regions remain critical factors to monitor.

Fuente: Gurufocus. Disponible en <https://n9.cl/g8p8t6>



VacciMonitor es una revista dedicada a la vacunología y temas afines como Inmunología, Adyuvantes, Infectología, Microbiología, Epidemiología, Validación, Aspectos regulatorios, entre otros. Arbitrada, de acceso abierto y bajo la Licencia Creative Commons está indexada en:

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Filters activated: (vaccine[Title/Abstract]) AND (("2025/02/15"[Date - Publication] : "2025/02/28"[Date - Publication])) records.

Monkeypox Vaccine.

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COVID-19 Vaccines.

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RNA neoantigen vaccines prime long-lived CD8+ T cells in pancreatic cancer.

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Patentes registradas en Patentscope

Estrategia de búsqueda: (Vaccine) AND DP:([15.02.2025 TO 28.02.2025]) as the publication date 63 records.

1. 20250064919A SARS-COV-2 HUMAN PARAINFLUENZA VIRUS TYPE 3-VECTORED VACCINE

US - 27.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18726909Solicitante Board of Regents, The University of Texas SystemInventor/a Alexander Bukreyev

The disclosure is directed to a new human parainfluenza virus (HPIV)/SARS-CoV-2 vaccine or vaccine construct/polynucleotide. Specifically, the HPIV is a Human parainfluenza virus type 3 (HPIV3), and the vaccine construct encoding a SARS-CoV-2 spike protein (S protein), that is a SARS-CoV-2 S protein S1 subunit and/or a SARS-CoV-2 S protein receptor binding domain (RBD), in certain aspects the vaccine or vaccine construct is administered via intranasal administration.

2. WO/2025/035187VACCINE ANTIGEN

WO - 20.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud PCT/AU2024/050878Solicitante MACFARLANE BURNET INSTITUTE FOR MEDICAL RESEARCH AND PUBLIC HEALTH LIMITEDInventor/a DRUMMER, Heidi

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

3. WO/2025/042223METHOD FOR PREDICTING TRANSLATION EFFICIENCY OF MESSENGER RNA VACCINE AND ANALYSIS DEVICE

WO - 27.02.2025

Clasificación Internacional G16B 40/20Nº de solicitud PCT/KR2024/012535Solicitante IUCF-HYU (INDUSTRY-UNIVERSITY COOPERATION FOUNDATION HANYANG UNIVERSITY)Inventor/a NAM, Jin-Wu

A method for predicting translation efficiency of an mRNA vaccine comprises the steps in which: an analysis device receives sequence information of a target mRNA vaccine; the analysis device extracts, from the

sequence information, an input sequence including a 25 nt sequence to which a primer binds in a 5' untranslated region (UTR), a 50 nt sequence of the 5' UTR immediately before a coding sequence (CDS) site, and a 30 nt sequence after a start codon of a coding region; the analysis device calculates secondary structure information for the input sequence; and the analysis device inputs the input sequence and the secondary structure information into a pre-trained deep learning model so as to predict the translation efficiency of the corresponding mRNA vaccine.

4.20250057943 FOWL ADENOVIRUS SUBUNIT VACCINE AND PRODUCTION METHOD THEREOF

US - 20.02.2025

Clasificación Internacional A61K 39/235Nº de solicitud 18726065Solicitante VETERINÄRMEDIZINISCHE UNIVERSITÄT WIENInventor/a MICHAEL HESS

The present invention provides a fowl adenovirus (FAdV) subunit vaccine, comprising at least a chimeric FAdV fiber protein and an adjuvant. This vaccine may be used to ameliorate or prevent adenoviral gizzard erosion (AGE), inclusion body hepatitis (IBH) or hepatitis-hydropericardium syndrome (HHS) in birds. The invention further relates to a method of producing an FAdV subunit vaccine, comprising the steps of expressing a chimeric FAdV fiber protein in an expression system, purifying the fiber protein, and combining the fiber protein with an adjuvant to obtain the FAdV subunit vaccine.

5.20250057934 STABLE EMULSIONS OF ANTIGENS

US - 20.02.2025

Clasificación Internacional A61K 39/135Nº de solicitud 18723268Solicitante INTERVET INC.Inventor/a Martin PIESL

The present invention relates to the field of vaccinology, more specifically of veterinary vaccinology. In particular, the invention relates to an adjuvant composition comprising an emulsion of water, tocopherol or a pharmaceutically acceptable ester thereof, and a polyethoxyethylene cetostearyl ether as an emulsifier. Said adjuvant composition can be used for formulating a vaccine, particularly an emulsion vaccine, comprising a bacterial, parasitic, or viral antigen. The resulting vaccine composition can be used in a method of protecting a human or animal target against infection and/or disease caused by a pathogen, particularly caused by a bacterium, parasite or virus. The invention further relates to methods for the manufacture of such adjuvant compositions and for the manufacture of such vaccine composition.

6.20250057939 FREEZE-DRIED VIRAL COMBINATION VACCINE COMPOSITIONS AND PROCESS FOR PREPARATION THEREOF

US - 20.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18690297Solicitante Serum Institute Of India Private LimitedInventor/a Rajeev Mhalasakant DHERE

The present disclosure relates to field of lyophilized/freeze-dried viral combination composition/formulation and methods for manufacturing and obtaining the composition comprising at least three live attenuated virus selected from a group of Coronavirus, Measles virus and Rubella virus; and stabilizers comprising of at least one carbohydrate, at least one amino acid and at least one hydrolyzed protein. The said lyophilized/freeze-dried viral combination composition/formulation is a vaccine composition that preserves the desired characteristics of each virus, including stability and immunogenicity. The composition can be safely

administered subcutaneously as a combination **vaccine** composition such that the immunogenicity of each of the measles, rubella and SARS-CoV-2 is not inferior to that observed for each of the three viruses when administered as individual vaccines and is found to be equivalent or improved as compared to immunogenicity of SARS-CoV-2 **vaccine** given intranasally. The purification process is devoid of chromatography steps.

7.WO/2025/036230 PROTEIN, ADENOVIRUS AND VACCINE AGAINST INFECTION OF SUBTYPE OF SARS-COV-2 OMICRON MUTANT STRAIN XBB

WO - 20.02.2025

Clasificación Internacional C07K 19/00Nº de solicitud PCT/CN2024/110575Solicitante WESTVAC BIOPHARMA CO., LTD.Inventor/a WEI, Xiawei

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

8.20250057938 STABILISED LIQUID VACCINES OF LIVE VIRUSES

US - 20.02.2025

Clasificación Internacional A61K 39/155Nº de solicitud 18723259Solicitante INTERVET INC.Inventor/a Edwin KETS

The present invention relates to liquid **vaccine** composition comprising a live virus and a natural deep-eutectic solvent (NADES) as the carrier. The carrier additionally comprises an additive selected from methionine and (hydroxy) ectoine. The additive is able to reduce the loss of virus titre over time, upon storage in a NADES-based liquid **vaccine** composition having up to 50% w/w of water. Such compositions are less viscous which is favourable for the manufacture of the carrier, and the formulation and use of the liquid **vaccine**.

9.WO/2025/040174 METHOD FOR CHARACTERIZING INHALATION VACCINE PERFORMANCE

WO - 27.02.2025

Clasificación Internacional C12N 15/861Nº de solicitud PCT/CN2024/114309Solicitante CANSINO BIOLOGICS INC.Inventor/a ZHAO, Xiaolong

A method for characterizing inhalation **vaccine** performance. The method comprises quantitatively linking an NGI cascade impactor, quantitative real-time polymerase chain reaction (QPCR), and droplet digital PCR (ddPCR) to establish a complete performance characterization method for inhalation vaccines. A testing environment with stable results and high repeatability has been obtained by in-depth analysis of the various influencing factors that the NGI cascade impactor has in respect of inhalation **vaccine** characterization. Meanwhile, the optimal characterization sample dosage has been determined on the basis of multiple

repeated experiments. The present application innovatively proposes combining NGI, QPCR and ddPCR; the method allows for accurate testing that takes little time, and testing of large batch samples can be accomplished, ensuring scalable application in industry.

10. WO/2025/041889 RECOMBINANT INFLUENZA VIRUS VECTOR EXPRESSING FOREIGN ANTIGEN AND VACCINE COMPOSITION COMPRISING SAME

WO - 27.02.2025

Clasificación Internacional C12N 15/86Nº de solicitud PCT/KR2023/012567Solicitante SUNGSHIN WOMEN'S UNIVERSITY INDUSTRY-ACADEMIC COOPERATION FOUNDATIONInventor/a SONG, Jae Min

The present invention relates to a recombinant influenza virus vector expressing a foreign antigen and a vaccine composition comprising same. The vector has a partially deleted NS1 gene of the influenza virus vector, resulting in the expression of an NS1 protein with a shortened C-terminal, which attenuates the virus and enables vaccine production. Additionally, the insertion of a foreign gene induces antibodies against the foreign protein (antigen), thereby achieving cross-immunogenicity.

11. 20250066771 STREAMLINED SELECTION OF NATURALLY CIRCULATING, ANTIGENIC MATCH AND HIGH-YIELD VACCINE VIRUSES FOR SEASONAL INFLUENZA VACCINE PRODUCTION

US - 27.02.2025

Clasificación Internacional C12N 15/10Nº de solicitud 18792336Solicitante The Curators of the University of MissouriInventor/a Xiufeng Henry Wan

A method of predicting preferred candidate viral strains for vaccine production is described herein. A machine learning framework for predicting preferred candidate viral strains is provided that emphasizes antigen-matching and yield predictions that correlate with features of viral proteins.

12. WO/2025/038171 A BACTERIAL VESICLE-BASED PNEUMOCOCCAL VACCINE AGAINST INFLUENZA-MEDIATED SECONDARY STREPTOCOCCUS PNEUMONIAE PULMONARY INFECTION

WO - 20.02.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/US2024/034124Solicitante ALBANY MEDICAL COLLEGEInventor/a SUN, Wei

A *Yersinia pseudotuberculosis* strain (designated as YptbS46) was tailored with an Asd⁺ plasmid pSMV92 that can synthesize high amounts of the Spn pneumococcal surface protein A (PspA) antigen and monophosphoryl lipid A as an adjuvant. The recombinant strain produced outer membrane vesicles (OMVs) enclosing a high amount of PspA protein (designated as OMV-PspA). A prime-boost intramuscular immunization with 30 pg of OMV- PspA induced both memory adaptive and innate immune responses in co-infected mice, reduced the viral and bacterial burden, and provided complete protection against secondary Spn infection. Also, the OMV-PspA immunization afforded significant cross-protection against the secondary Spn A66.1 infection and long-term protection against the secondary Spn D39 challenge. An OMV vaccine delivering Spn antigens can thus be a new pneumococcal vaccine candidate.

13. WO/2025/038983 INACTIVATED WHOLE VIRION VACCINE AGAINST INFLUENZA AND METHODS OF USE THEREOF

WO - 20.02.2025

Clasificación Internacional C12N 7/06Nº de solicitud PCT/US2024/042824Solicitante COLORADO STATE UNIVERSITY RESEARCH FOUNDATIONInventor/a GOODRICH, Raymond

Provided herein are methods for inactivating an influenza particle, the methods comprising contacting the influenza particle with UV light in the presence of riboflavin such that the integrity of the antigenic proteins of the influenza particle are preserved. Vaccine compositions comprising inactivated influenza particles, and methods of use thereof, are also provided

14.2025200780 COMPOSITIONS AND METHODS FOR PERSONALIZED NEOPLASIA VACCINES

AU - 20.02.2025

Clasificación Internacional N° de solicitud 2025200780Solicitante Dana-Farber Cancer Institute, Inc.Inventor/a Fritsch, Edward F.

The invention provides a method of making a personalized neoplasia vaccine for a subject diagnosed as having a neoplasia, which includes identifying a plurality of mutations in the neoplasia, analyzing the plurality of mutations to identify a subset of at least five neo antigenic mutations predicted to encode neo-antigenic peptides, the neo-antigenic mutations selected from the group consisting of missense mutations, neoORF mutations, and any combination thereof, and producing, based on the identified subset, a personalized neoplasia vaccine.

15.20250057936 VACCINE COMPOSITION

US - 20 02 2025

Clasificación Internacional A61K 39/145 N° de solicitud 18910182 Solicitante Lionel Scott Inventor/a Lionel Scott

Respiratory virus vaccine compositions for nasal administration to a mammal comprising a hygroscopic gel-forming material, at least one isolated bioactive respiratory virus immunogen, and an adjuvant, kits, receptacles, uses therefor, and methods of manufacture thereof.

16.20250057928 VACCINE COMPOSITION FOR INDUCING ANTI-IL-23 ANTIBODY

US - 20 02 2025

Clasificación Internacional A61K 39/00Nº de solicitud 18717944Solicitante OSAKA
UNIVERSITYInventor/a Hironori NAKAGAMI

The present invention provides a vaccine composition containing a complex of a T cell receptor antigen peptide and a B cell receptor antigen peptide and capable of inducing the production of an antibody against IL-23, wherein the B cell receptor antigen peptide is represented by the following formula (I):

X1-X2-X3-X4-X5-X6-X7-X8 (1)

wherein

- - X1 is S, A, G, T, K, or R,
 - X2 is P, A, G, S, T, K, or R,
 - X3 is S, A, G, T, K, or R,
 - X4 is Q, A, G, T, or N,
 - X5 is P, A, G, S, T, Q, or N,
 - X6 is W, A, Y, or E,
 - X7 is Q, A, G, T, or N, and
 - X8 is R, A, G, or K.

17.20250057942 NUCLEIC ACID VACCINE AGAINST THE SARS-COV-2 CORONAVIRUS

US - 20.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18941809Solicitante INSTITUT
PASTEURInventor/a Etienne SIMON-LORIERE

The invention relates to an immunogenic or vaccine composition against the 2019 novel coronavirus (SARS-CoV-2), comprising a nucleic acid construct encoding a SARS-CoV-2 coronavirus Spike(S) protein antigen or a fragment thereof comprising the receptor-binding domain, wherein the nucleic acid construct sequence is codon-optimized for expression in 5 human.

18.4507722 UNIVERSELLER INFLUENZAIMPFSTOFF IN NANOEMULSION

EP - 19.02.2025

Clasificación Internacional A61K 39/145Nº de solicitud 23788836Solicitante BLUEWILLOW BIOLOGICS
INCInventor/a GANESAN SHYAMALA

The present invention relates to methods for inducing a broadly reactive immune response to multiple strains of influenza in a subject comprising intranasally administering a nanoemulsion vaccine composition comprising a computationally optimized influenza immunogen or protein.

19.WO/2025/042248 METHOD OF SELECTING NEOANTIGEN FOR DEVELOPMENT OF PERSONALIZED CANCER VACCINE

WO - 27.02.2025

Clasificación Internacional G16B 40/20Nº de solicitud PCT/KR2024/012654Solicitante LG CHEM,
LTD.Inventor/a JEONG, Seihwan

Provided are a method of selecting a tumor-specific neoantigen (immunogenic peptide), and a use of the selected tumor-specific neoantigen for preparing a personalized cancer vaccine.

20.WO/2025/036474 RESPIRATORY SYNCYTIAL VIRUS ANTIGENIC POLYPEPTIDE, NUCLEIC ACID, VACCINE, AND USE THEREOF

WO - 20.02.2025

Clasificación Internacional C07K 14/135Nº de solicitud PCT/CN2024/112630Solicitante SHENZHEN RHEGEN BIOTECHNOLOGY CO., LTD.Inventor/a HU, Yong

A respiratory syncytial virus antigenic polypeptide, a nucleic acid, a vaccine, and a use thereof. Provided is a respiratory syncytial virus antigenic polypeptide or an immunogenic fragment thereof. The respiratory syncytial virus antigenic polypeptide has at least 26-th to 470-th or full-length amino acid sequences shown in SEQ ID NO:1, or derivative amino acid sequences formed by replacing, deleting, or adding one or several amino acids on the basis of these amino acid sequences and having equivalent functions. An increase in the pre-F immunogenicity can be achieved and/or an increase in the protein expression level can be facilitated.

21.WO/2025/037284 GENETICALLY MODIFIED BACTERIA FOR MULTI-MODAL SECRETION OF A NEOANTIGEN

WO - 20.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/IB2024/058038Solicitante BACCINE LTD.Inventor/a STRAUSSMAN, Ravid

A vaccine and methods of treatment thereof, wherein the vaccine comprises a recombinant Gram-negative bacteria genetically modified to express a first antigen fusion peptide comprising a neoantigen or series thereof, said neoantigen or series thereof associated with a first secretion signal from a double membrane-spanning secretion system and a second antigen fusion peptide comprising a homologous neoantigen or series thereof, associated with a second secretion signal from an outer membrane-spanning secretion system. The Gram-negative bacteria may be further modified for quadmodal transport. Specifically, the fusion peptides include signal peptides are each associated with a Type III (T3SS) and a Type V (T5SS) secretion system.

22.WO/2025/043210 VACCINE COMPOSITIONS, COMPONENTS, AND METHODS OF USE

WO - 27.02.2025

Clasificación Internacional A61K 39/08Nº de solicitud PCT/US2024/043720Solicitante THE REGENTS OF THE UNIVERSITY OF MICHIGANInventor/a SUN, Duxin

The present disclosure provides vaccine components and compositions, and methods of use thereof. More specifically, the disclosure provides compositions comprising one or more T cell epitopes(e.g., one or more tumor-associated T cell epitopes and/or one or more T cell epitopes derived from a microorganism), or a nucleic acid encoding thereof, and one or more B cell epitopes, or a nucleic acid encoding thereof. The disclosure also provides RNA molecules (e.g., mRNA molecules) encoding B cell epitopes to mediate B cell antigen presentation, magnetic nanoparticles comprising antigen clusters having B cell and/or T cell epitopes, and compositions and vaccines (e.g., cancer vaccines) thereof.

23.WO/2025/037838 METHOD FOR DESIGNING PERSONALIZED CANCER VACCINE USING B CELL-REACTIVE NEOANTIGEN

WO - 20.02.2025

Clasificación Internacional G16B 40/20Nº de solicitud PCT/KR2024/011714Solicitante KOREA ADVANCED INSTITUTE OF SCIENCE AND TECHNOLOGYInventor/a CHOI, Jung Kyoon

The present disclosure relates to a method for designing a personalized cancer vaccine, using a B cell-reactive neoantigen, the method comprising the steps of: receiving a peptide sequence extracted from cancer tissue and allelic sequence data of B cell receptors; calculating the preferential binding strength between amino acids in the peptide sequence and allelic sequences of B cell receptors; and calculating B cell epitope prediction scores by inputting the preferential binding strength between amino acids to a trained neural network model.

24.4511059STREPTOCOCCUS SUIS IMPFSTOFFZUSAMMENSETZUNG MIT IMMUNOGENEN FUSIONSPOLYPEPTIDEN

EP - 26.02.2025

Clasificación Internacional A61K 39/09Nº de solicitud 23717588Solicitante INTERVACC ABInventor/a FROSTH SARA

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 administrating such compositions.

25.WO/2025/040802NEW SCHEME OF ADMINISTRATION OF A MULTIVALENT VACCINE AGAINST SWINE INFECTIONS

WO - 27.02.2025

Clasificación Internacional A61K 39/08Nº de solicitud PCT/EP2024/073717Solicitante CEVA SANTE ANIMALEInventor/a FACHINGER, Vicky

The present disclosure relates to a multivalent vaccine composition and its use in the protection against swine infections in a female pig and its progeny with a new scheme of administration.

26.20250057945VACCINE ADJUVANT AGENT CONTAINING POLYACRYLIC ACID POLYMER AND USE OF SAME

US - 20.02.2025

Clasificación Internacional A61K 39/39Nº de solicitud 18721547Solicitante TOKO YAKUHIN KOGYO CO., LTD.Inventor/a Taizou KAMISHITA

The present disclosure provides a vaccine adjuvant agent comprising a polymer in which a constitutional unit is derived from acrylic acid, wherein the polymer has a carboxyl group content of 60.0 to 62.5% by mass.

27.4511492VETERINÄRE IMPFSTOFFE UND VERFAHREN ZUR BEHANDLUNG VON PASTEURELLA MULTOCIDA-INFektIONEN BEI TIEREN

EP - 26.02.2025

Clasificación Internacional C12N 15/63Nº de solicitud 23790816Solicitante ENG ANTIGENS INCInventor/a MORAES TREVOR

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

28.4511493VETERINÄRE IMPFSTOFFE UND VERFAHREN ZUR BEHANDLUNG VON PASTEURELLA MULTOCIDA-INFektIONEN BEI TIEREN

EP - 26.02.2025

Clasificación Internacional C12N 15/63Nº de solicitud 23790817Solicitante ENG ANTIGENS INCInventor/a MORAES TREVOR

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

29.WO/2025/036985A VACCINE FOR THE TREATMENT OR PREVENTION OF BURKHOLDERIA SPECIES INFECTION IN A SUBJECT.

WO - 20.02.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/EP2024/073032Solicitante UNIVERSITY COLLEGE DUBLINInventor/a MCCLEAN, Siobhan

A composition comprising BpPA26 is provided. The composition is for use in vaccine therapy to treat or prevent pathogenic Burkholderia species infection in a mammal, typically a human or an equine mammal. The species include *Burkholderia cepacia* complex, *Burkholderia mallei*, and *Burkholderia pseudomallei*.

30.WO/2025/040792MULTIVALENT VACCINE AGAINST SWINE INFECTIONS

WO - 27.02.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/EP2024/073694Solicitante CEVA SANTE ANIMALEInventor/a FACHINGER, Vicky

The present disclosure relates to a multivalent vaccine composition and its use in the protection against *Escherichia coli* (*E.coli*), *Clostridium* spp., porcine parvovirus and *Erysipelothrix rhusiopathiae* (*E. rhusiopathiae*) infections in a pig and its progeny, preferably with a new scheme of administration for passive and active immunization of a pig and its progeny.

31.20250057947USE OF TRIMANGANESE TETRAOXIDE PARTICLES IN PREPARATION OF VACCINE ADJUVANT

US - 20.02.2025

Clasificación Internacional A61K 39/39Nº de solicitud 18763977Solicitante Guangzhou Realbenefitspot Pharmaceutical Co., Ltd.Inventor/a Yaling WANG

Disclosed is a use of trimanganese tetraoxide particles in preparation of a vaccine adjuvant. The adjuvant is a particle adjuvant, the particle adjuvant is trimanganese tetraoxide particles externally wrapped with or

without an excipient, and the particle size of the particle adjuvant is 5 nm to 3000 nm. The trimanganese tetraoxide particle adjuvant provided in the present invention can be effectively combined with a single-stranded nucleotide adjuvant and can effectively carry an immune antigen, and a more excellent immunotherapy effects can be achieved when a fewer antigen dose and a relatively low injection amount are used; immune cells are efficiently activated, and body fluid balance and cellular immunity are achieved.

32.20250057940 IMMUNE MEMORY ENHANCED PREPARATIONS AND USES THEREOF

US - 20.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18752524Solicitante Torigen Pharmaceuticals, Inc.Inventor/a Mark Suckow

Formulations and preparations having immune memory enhanced properties are disclosed that provide for enhancing immune response against a tumor growth, cancer, infectious agent, bacteria, virus or other infectious or non-infectious agent. The vaccine formulation includes an immune memory invoking component, such as an antigen of an infectious agent, virus (e.g., Rabies), bacteria, prion, neo-antigen or other moiety antigen, and a targeted antigen (e.g., a harvested tumor tissue (B-cell, T-cell, epitopes)). The vaccine formulation/preparations may comprise a target infectious agent protein/peptide component (such as a SARS-CoV-2 spike protein epitope) mixed with, or fused to (or otherwise conjugated) an immune-memory associated viral antigen (such as Rabies, polio, or other peptide/protein antigen or peptide or fragment thereof).

33.20250064910 A MAPS VACCINE TARGETING SALMONELLA ENTERICA SEROVARS

US - 27.02.2025

Clasificación Internacional A61K 39/112Nº de solicitud 18715610Solicitante THE CHILDREN'S MEDICAL CENTER CORPORATIONInventor/a Richard MALLEY

Technologies for the prevention and/or treatment of *Salmonella* infections.

34.20250064915 NUCLEIC ACID MOLECULES AND USES THEREOF

US - 27.02.2025

Clasificación Internacional A61K 39/125Nº de solicitud 18940501Solicitante CureVac SEInventor/a Susanne RAUCH

The present invention is directed to an artificial nucleic acid and to polypeptides suitable for use in treatment or prophylaxis of an infection with Norovirus or a disorder related to such an infection. In particular, the present invention concerns a Norovirus vaccine. The present invention is directed to an artificial nucleic acid, polypeptides, compositions and vaccines comprising the artificial nucleic acid or the polypeptides. The invention further concerns a method of treating or preventing a disorder or a disease, first and second medical uses of the artificial nucleic acid, polypeptides, compositions and vaccines. Further, the invention is directed to a kit, particularly to a kit of parts, comprising the artificial nucleic acid, polypeptides, compositions and vaccines.

35.20250057935 ADENOVIRAL VECTOR SYSTEM FOR GENE DELIVERY

US - 20.02.2025

Clasificación Internacional A61K 39/145Nº de solicitud 18749120Solicitante Purdue Research FoundationInventor/a Suresh Kumar Mittal

Disclosed herein a unique cell line system to generate a novel bovine adenovirus vector that provides more gene insertion capabilities and better immunogenicity for inserted antigens. The unique cell line is used for generating and growing of the new adenovirus vectors for gene delivery or recombinant vaccine production.

36.4507732PROTEIN-SACCHARID-KONJUGATION MIT Natriumcyanoborohydrid

EP - 19.02.2025

Clasificación Internacional A61K 47/64Nº de solicitud 23720209Solicitante SANOFI PASTEUR INCInventor/a GINLEY MARYALICE

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.WO/2025/038868LIPID NANOPARTICLE FORMULATIONS AND USES THEREOF

WO - 20.02.2025

Clasificación Internacional A61K 47/54Nº de solicitud PCT/US2024/042532Solicitante ICAHN SCHOOL OF MEDICINE AT MOUNT SINAIInventor/a DONG, Yizhou

The present disclosure relates to polynucleotide formulations containing lipid nanoparticles and methods of administration, particularly in vaccine formulations.

38.20250059238RECOMBINANT HIV ENV POLYPEPTIDES AND THEIR USES

US - 20.02.2025

Clasificación Internacional C07K 14/005Nº de solicitud 17299062Solicitante Dennis R. BURTONInventor/a Dennis R. BURTON

The present disclosure relates to recombinant HIV Env polypeptides and their use in the treatment and prevention of HIV/AIDS.

39.4512817NEOANTIGENIDENTIFIZIERUNG, HERSTELLUNG UND VERWENDUNG

EP - 26.02.2025

Clasificación Internacional C07K 7/08Nº de solicitud 24205415Solicitante GRITSTONE BIO INCInventor/a BULIK-SULLIVAN BRENDAN

Disclosed herein is a system and methods for determining the alleles, neoantigens, and vaccine composition as determined on the basis of an individual's tumor mutations. Also disclosed are systems and methods for obtaining high quality sequencing data from a tumor. Further, described herein are systems and methods for identifying somatic changes in polymorphic genome data. Further, described herein are systems and methods for selecting a subset of patients for treatment. A utility score indicating an estimated number of neoantigens presented on the surface of tumor cells is determined for each patient based on one or more neoantigen candidates identified for the patient. The subset of patients are selected based on the determined utility

scores. The selected subset of patients can receive treatment, such as neoantigen vaccines or checkpoint inhibitor therapy. Finally, described herein are unique cancer vaccines.

40.4511058ZUSAMMENSETZUNGEN, VORRICHTUNGEN, SYSTEME UND VERFAHREN IM ZUSAMMENHANG MIT IMPFUNG UND STERILEM SCHUTZ GEGEN MALARIA

EP - 26.02.2025

Clasificación Internacional A61K 39/015Nº de solicitud 23935577Solicitante MALARVX INC.Inventor/a AVRIL MARION

Systems, compositions, devices, methods, etc., provide improved anti-malaria immunological responses comprising making, providing and administering vaccines comprising specific RNA molecules such as self-replicating replicon RNA (repRNA) encoding proteins from Plasmodium such as the *P. yoelii* (Py) CS protein (CSP), including in some embodiments substantially target proteins encoding target antigens, for example a whole or substantially whole CSP in the repRNA. The prime-and-trap intervals for the administration of the vaccine can comprise administration of only a single dose of a repRNA-Non-encapsulating oil-in-water emulsion nanocarriers (e.g., LION™) component followed by administration of as few as 3 or 2 doses, or even just a single dose, of the WO component (e.g., RAS or genetically attenuated WO) at 0 day (same day), or 1, 2, 3, 4, 5, 10, 14, 15 days or 28 days later.

41.WO/2025/042889STABLE SALMONELLA VACCINE FORMULATIONS

WO - 27.02.2025

Clasificación Internacional A61K 39/02Nº de solicitud PCT/US2024/043036Solicitante ELANCO US INC.Inventor/a BUCK, Niklas

The present disclosure provides formulations, kits, and vaccines directed to immunization of animals against Salmonella. Methods of using the formulations, kits, and vaccines for protection of avians against one or more species of Salmonella are also provided.

42.WO/2025/038896BROAD SPECTRUM CONJUGATE VACCINE TO PREVENT KLEBSIELLA PNEUMONIAE AND PSEUDOMONAS AERUGINOSA INFECTIONS

WO - 20.02.2025

Clasificación Internacional A61K 39/108Nº de solicitud PCT/US2024/042585Solicitante UNIVERSITY OF MARYLAND, BALTIMOREInventor/a CROSS, Alan, S.

The present invention is drawn to conjugates comprising a glycosylated native FlaA flagellin protein of *Pseudomonas* and *Klebsiella* surface polysaccharide antigens, such as *Klebsiella pneumoniae* O polysaccharides from serovars O1, O2a, O2a,c, O3, O4, O5, O7, O8 and O12. The present invention also provides pharmaceutical compositions comprising the same and methods of inducing an immune response in subjects by administering the compositions.

43.20250057949T CELL THERAPY WITH VACCINATION AS A COMBINATION IMMUNOTHERAPY AGAINST CANCER

US - 20.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18720347Solicitante The United States of America, as represented by the Secretary, Department of Health and HumanInventor/a Sri Krishna

Disclosed are methods of treating or preventing cancer in a mammal, the method comprising: (a) isolating T cells from a tumor sample from the mammal, wherein the isolated T cells are one or both of exhausted and differentiated, and the isolated T cells have antigenic specificity for a tumor-specific antigen expressed by the tumor sample from the mammal, wherein the tumor-specific antigen is a tumor-specific neoantigen or an antigen with a tumor-specific driver mutation; and optionally expanding the numbers of isolated, tumor antigen-specific T cells; and (b) administering to the mammal (i) the isolated T cells of (a) and (ii) a vaccine which specifically stimulates an immune response against the tumor-specific antigen for which the isolated T cells have antigenic specificity.

44.4230209 PHARMACEUTICAL COMPOSITION, PHARMACEUTICAL COMBINED FORMULATION, AND COMBINED FORMULATION KIT FOR PREVENTION OR TREATMENT OF CHRONIC HEPATITIS B, EACH COMPRISING, AS ACTIVE INGREDIENT, ORAL ANTIVIRAL AGENT AND THERAPEUTIC VACCINE INCLUDING LIPOPEPTIDE AND POLY(I:C) ADJUVANT

PL - 24.02.2025

Clasificación Internacional A61K 31/513Nº de solicitud 21894866SolicitanteInventor/a JUNG SUN YUM

45.20250064914 NUCLEOSIDE-MODIFIED RNA FOR INDUCING AN ADAPTIVE IMMUNE RESPONSE

US - 27.02.2025

Clasificación Internacional A61K 39/12Nº de solicitud 18925561Solicitante The Trustees of the University of PennsylvaniaInventor/a Drew Weissman

The present invention relates to compositions and methods for inducing an adaptive immune response in a subject. In certain embodiments, the present invention provides a composition comprising a nucleoside-modified nucleic acid molecule encoding an antigen, adjuvant, or a combination thereof. For example, in certain embodiments, the composition comprises a vaccine comprising a nucleoside-modified nucleic acid molecule encoding an antigen, adjuvant, or a combination thereof.

46.4507723 SARS-COV-2-IMPFSTOFFZUSAMMENSETZUNGEN

EP - 19.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 23789109Solicitante MERCIA PHARMA INCInventor/a BLACKBURN PETER

The present disclosure provides compositions of adjuvanted SARS-CoV-2 vaccines and their use to prevent and manage Covid-19 infection, including host hyperinflammatory responses to infection, including long term symptoms associated with Covid infection.

47.20250064920 IBV VACCINE WITH HETEROLOGOUS DMV/1639 SPIKE PROTEIN

US - 27.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18790098Solicitante Boehringer Ingelheim Vetmedica GmbHInventor/a Grace Albanese

The present invention relates i.a. to an IBV (infectious bronchitis virus) encoding for a heterologous DMV S (spike) protein or fragment thereof. Further, the present invention relates to an immunogenic composition

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comprising said IBV encoding for a heterologous DMV S (spike) protein or fragment thereof. Furthermore, the present invention relates to methods for immunizing a subject comprising administering to such subject the immunogenic composition of the present invention. Moreover, the present invention relates to methods of treating or preventing clinical signs caused by IBV in a subject of need, the method comprising administering to the subject a therapeutically effective amount of an immunogenic composition according to the present invention.

48.4511479 BEHANDLUNG VON ENTZÜNDUNGSERKRANKUNGEN

EP - 26.02.2025

Clasificación Internacional C12N 7/04Nº de solicitud 23791796Solicitante CYN K BIO INCInventor/a UENO RYUJI

The present disclosure provides a method for treating an inflammatory condition, especially an age related inflammatory condition in a mammalian subject in need thereof, which comprises an effective amount of a virus like particle comprising a viral structural protein and a galectin-3 antigen, a composition or vaccine comprising for the purpose thereof.

49.20250057941 DYSREGULATION OF TRAUMA REGULATION PATHWAY TREATMENT AND MONITORING TECHNIQUES

US - 20.02.2025

Clasificación Internacional A61K 39/215Nº de solicitud 18939064Solicitante GE Precision Healthcare LLCInventor/a Christopher Michael Puleo

The subject matter of the present disclosure generally relates to techniques for addressing or correcting dysregulation of the trauma regulation pathway. The dysregulation may be associated with a physiological condition, such as a SARS-CoV-2 viral infection. In an embodiment, the techniques include treating dysregulation based on a renin-angiotensin pathway molecule or cell and/or a splenic pathway molecule or cell using targeted neuromodulation. In an embodiment, neuromodulation is used to regulate the immune system, e.g., as an energy-based adjuvant for a vaccine.

50.4512477 VERBESSERTE LAMPENKONSTRUKTIONEN

EP - 26.02.2025

Clasificación Internacional A61P 35/00Nº de solicitud 24201125Solicitante IMMUNOMIC THERAPEUTICS INCInventor/a HEILAND TERI

The present invention provides improved LAMP Constructs comprising specific fragments of the LAMP luminal domain to deliver antigens to immune cells for enhanced processing. These LAMP Constructs can be used for the treatment of disease and in particular, allergies, infectious disease, diabetes, hyperproliferative disorders and/or cancer. The improved LAMP Constructs allow for presentation of properly configured three dimensional epitopes for production of an immune response when administered to a subject. The improved LAMP Constructs can be multivalent molecules, and/or can be provided as part of a multivalent vaccine containing two or more LAMP Constructs. The improved LAMP Constructs as described herein can also be used to generate antibodies when administered to a non-human vertebrate.

51.20250066419 POLYPEPTIDE FOR DELIVERING ANTIGEN TO IMMUNE CELLS

US - 27.02.2025

Clasificación Internacional C07K 7/06Nº de solicitud 18724235Solicitante JW SHINYAK CORPORATIONInventor/a Yoon Jae JEON

The present invention relates to a polypeptide for delivering an antigen to immune cells and, specifically, to: a novel polypeptide comprising a cell membrane penetrating peptide and a peptide binding to a surface molecule on immune cells; a fusion polypeptide in which an antigen is coupled to the polypeptide; a nucleic acid coding for the polypeptide or the fusion polypeptide; an immune cell sensitized with the fusion polypeptide or the nucleic acid coding therefor; and an immunotherapeutic agent, antitumor or anticancer vaccine, and a composition for treating a tumor or cancer, each comprising the immune cell.

52.WO/2025/043016A UNIVERSAL VACCINE STRATEGY FOR CONFERRING PROTECTION AGAINST DIVERSE PATHOGENS

WO - 27.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud PCT/US2024/043282Solicitante THE BOARD OF TRUSTEES OF THE LELAND STANFORD JUNIOR UNIVERSITYInventor/a PULENDRA, Bali

Compositions and methods are provided for immunizing an individual or population of individuals to generate an immune response that protects against pathogen infection in an antigen-agnostic manner. The protective antigen-agnostic immune response is active for a period of time following immunization, from about 7 days, 10 days, 14 days following immunization, and may be active for at least 2 weeks, at least 3 weeks, at least 4 weeks, at least 5 weeks, or more.

53.WO/2025/039937USE OF COMPOUND IN TREATMENT OF LIVER DISEASES

WO - 27.02.2025

Clasificación Internacional A61K 31/519Nº de solicitud PCT/CN2024/111818Solicitante BEIJING SYNTHETIC VACCINE BIOSCIENCES CO., LTD.Inventor/a LIAO, Xuebin

Use of a compound in the treatment of liver diseases. Experimental results show that the compound has a relatively high liver-tissue-targeting property, very low toxicity to liver primary cells and high liver safety, has significant efficacy on liver diseases such as viral hepatitis (e.g., hepatitis B) and liver cancers (including primary liver cancers and metastatic liver cancers), and has relatively good physicochemical properties, metabolic stability, bioavailability and safety, thus having very good drug development prospects.

54.4512420ZUSAMMENSETZUNG ZUR REDUZIERUNG DER GRÖSSE ODER DES VOLUMENS VON ZIELGEWEBE ODER KIT DAMIT

EP - 26.02.2025

Clasificación Internacional A61K 39/12Nº de solicitud 23792262Solicitante SK BIOSCIENCE CO LTDInventor/a KIM EUN-SOM

The present invention provides a pharmaceutical composition for treating obesity, the composition including: one or more viruses selected from the group consisting of yellow fever virus, herpes zoster virus, and rubella virus; or a genetic material coding for a protein derived from these viruses. Preferably, the pharmaceutical

composition is a vaccine composition. The composition provides a reduction in target tissues, preferably tissues containing adipocytes, or an effect that leads to the death of adipocytes.

55.20250066740 NOVEL CHIMPANZEE ADENOVIRUS VECTOR, CONSTRUCTION METHOD THEREFOR, AND APPLICATION THEREOF

US - 27.02.2025

Clasificación Internacional C12N 7/00Nº de solicitud 18014647Solicitante JIAXING ANYU BIOTECHNOLOGY CO., LTDInventor/a Ping CHEN

A chimpanzee adenovirus vector from newly discovered and isolated chimpanzee adenovirus with unique HVR sequences and a method of its construction and application thereof, with higher viral titer, and provides a method for determining its viral titer. The chimpanzee adenovirus circular vector is constructed through a shuttle plasmid, and knocks out E1 and E3 to construct a replication-defective chimpanzee adenovirus vector. The chimpanzee adenovirus vector described in this article has no preexisting antibody in the population, and the knockout of E1 and E3 is safe. At the same time, it is significantly different from human adenovirus type 5 E1 in 293 cells, which can greatly avoid recovery mutation (RCA) and make it safer. The novel coronavirus vaccine has strong stability and does not cause mutations after multiple passages, and it can induce strong humoral immunity and cellular response in mouse models.

56.2998014 VACCINE FOR USE IN PROTECTING OFFSPRING OF A SOW AGAINST PORCINE ENDEMIC DIARRHEA VIRUS

ES - 18.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud 15762993Solicitante Intervet International B.V.Inventor/a BORN VAN DEN, Erwin

57.4509185 VERFAHREN ZUR ERHÖHUNG DER IMMUNOGENITÄT VON SCHWACH IMMUNOGENEN ANTIGENSPEZIFISCHEN IMPFSTOFFEN UNTER VERWENDUNG VON ORALEN HEFE-BETA-GLUCANEN

EP - 19.02.2025

Clasificación Internacional A61P 35/00Nº de solicitud 24223791Solicitante MEMORIAL SLOAN KETTERING CANCER CENTERInventor/a CHEUNG NAI-KONG

The present disclosure provides methods for enhancing the immunogenicity of a poorly immunogenic antigen-specific vaccine as well as methods for promoting diversification of the gut microbiome in a subject in need thereof comprising administering to the subject an effective amount of a beta-glucan extract derived from yeast. Kits for use in practicing the methods are also provided.

58.4511060 ENTEROCOCCUS FAECALIS IMPFSTOFF UND VERWENDUNGEN DAVON

EP - 26.02.2025

Clasificación Internacional A61K 39/09Nº de solicitud 23792699Solicitante VAXCYTE INCInventor/a FAIRMAN JEFFERY C

The present disclosure provides immunogenic compositions comprising at least one recombinant polypeptide antigen derived from an *Enterococcus* bacterium (e.g., *E. faecalis*, *E. faecium*, *E. durans*). The disclosure

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further provides methods, and uses of the immunogenic compositions, for protecting or treating a subject from infection by an *Enterococcus* bacterium. Such infections may cause, or worsen, conditions such as root canal failure, endocarditis, bacteremia, urinary tract infections, prostatitis, intraabdominal infection, cellulitis, dysbiotic gastrointestinal tract, prosthetic joint infection, or wound infections.

59.4512419POLYVALENTE PNEUMOKOKKEN-POLYSACCHARID-KONJUGAT-IMPFSTOFFKOMPONENTE UND ANWENDUNG DAVON

EP - 26.02.2025

Clasificación Internacional A61K 39/116Nº de solicitud 23791262Solicitante SHANGHAI REINOVAX BIOLOGICS CO LTDInventor/a ZHU XIANCHAO

The present invention relates to a polyvalent pneumococcal polysaccharide protein conjugate and immunogenicity thereof, and specifically provides an immunogenic composition containing capsular polysaccharides of streptococcus pneumoniae from different serotypes, and a carrier, the serotypes at least comprising 2, 8, 9N, 10A, 11A, 12F, 15B, 17F, 20, 22F and 33F. The immunogenic composition can improve the immunogenicity of polysaccharides of different serotypes, and may prevent invasive infection caused by pneumococci of various different serotypes.

60.3626262VEGFR-2 TARGETING DNA VACCINE FOR COMBINATION THERAPY

PL - 17.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud 19205420SolicitanteInventor/a HEINZ LUBENAU

61.2998010VACCINE FOR USE IN PROTECTING A PIG AGAINST PORCINE ENDEMIC DIARRHEA VIRUS

ES - 18.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud 15762624Solicitante Intervet International B.V.Inventor/a BORN VAN DEN, Erwin

62.20250057933RHINOVIRUS VACCINE

US - 20.02.2025

Clasificación Internacional A61K 39/125Nº de solicitud 18817094Solicitante IP2IPO INNOVATIONS LIMITEDInventor/a Sebastian Johnston

The invention relates to immunogenic compositions, and in particular, to immunogenic compositions for preventing, treating or ameliorating human rhinovirus (RV) infections. The invention is especially concerned with RV VP0 peptides (or proteins) and polynucleotides encoding such peptides, and their use in immunogenic compositions for eliciting an immune response and preventing rhinovirus infections.

63.20250064908MULTICOMPONENT CHEMICAL COMPOSITION OF A PEPTIDE-BASED NEOANTIGEN VACCINE

US - 27.02.2025

Clasificación Internacional A61K 39/00Nº de solicitud 18924824Solicitante AMAZON TECHNOLOGIES, INCInventor/a Frank Wilhelm Schmitz

Provided herein are immunogenic compositions comprising tumor-specific neoantigen long peptides, tumor-specific neoantigen short peptides, and adjuvant, optionally a helper peptide, and optionally a tumor-specific peptide. The disclosure also provides methods of using these immunogenic compositions for treating cancer.

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