



EN ESTE NÚMERO

VacCiencia es una publicación dirigida a investigadores y especialistas dedicados a la vacunología y temas afines, con el objetivo de serle útil. Usted puede realizar sugerencias sobre los contenidos y de esta forma crear una retroalimentación que nos permita acercarnos más a sus necesidades de información.

- Noticias más recientes en la Web sobre vacunas.
- Artículos científicos más recientes de Medline sobre vacunas.
- Patentes más recientes en Patentscope sobre vacunas.

Noticias en la Web

La vacuna nasal contra la COVID-19 detiene la infección en ensayos con animales

1 ago. Una vacuna nasal de próxima generación contra COVID-19 parece hacer lo que las vacunas inyectables no pueden: en realidad detener la propagación del virus de persona a persona.

Los hámsters que recibieron la vacuna nasal no transmitieron el virus a otros si se infectaban, rompiendo el ciclo de transmisión, informaron los investigadores en la edición del 31 de julio de la revista Science Advances.

El estudio con animales proporciona más evidencias de que las vacunas administradas en la nariz o la boca podrían ser la clave para controlar la propagación de infecciones respiratorias como la influenza y COVID, dijeron los investigadores.

"Para prevenir la transmisión, hay que mantener baja la cantidad de virus en las vías respiratorias superiores", señaló el investigador sénior, Jacco Boon, profesor de medicina, microbiología molecular y patología e inmunología de la Facultad de Medicina de la Universidad de Washington, en St. Louis.

"Para empezar, cuanto menos virus haya, menos probabilidades hay de infectar a otra persona si tose o estornuda, o incluso si simplemente respira sobre ellos", añadió Boon en un comunicado de prensa de la universidad. "Este estudio muestra que las vacunas contra la mucosa son superiores a las vacunas inyectables en términos de limitar la replicación viral en las vías respiratorias superiores y prevenir la propagación al siguiente individuo".

"En una situación epidémica o pandémica, este es el tipo de vacuna que se va a querer", concluyó Boon.

Aunque las primeras vacunas contra el COVID lograron reducir la enfermedad grave y la muerte, las inyecciones no pudieron prevenir la propagación del virus. Una persona vacunada levemente enferma aún podría transmitir el virus a otra persona.

Los virus como la gripe, COVID y VRS se multiplican rápidamente en la nariz, lo que permite que se propaguen de persona a persona a través de la tos, los estornudos e incluso la respiración, apuntaron los investigadores.

Las vacunas inyectables tradicionales son mucho menos potentes en la nariz que en el torrente sanguíneo, lo que deja a la nariz relativamente desprotegida contra los virus que se multiplican rápidamente y se propagan fácilmente.

Los investigadores han teorizado durante mucho tiempo que una vacuna administrada por la nariz o la boca a través de aerosoles o gotitas podría reducir la transmisión de enfermedades al desencadenar una respuesta inmunitaria justo donde más se necesita.

Para este estudio, los investigadores utilizaron un proceso de dos pasos para probar una vacuna nasal contra COVID utilizada en la India contra la vacuna inyectable de Pfizer en un grupo de hámsters.

Los hámsters son naturalmente susceptibles a COVID, a diferencia de los ratones, lo que los convierte en un mejor animal de laboratorio con el que estudiar la transmisión del virus, explicaron los investigadores.

Después de dar a los hámsters vacunados unas semanas para desarrollar una respuesta inmunitaria



completa, los investigadores infectaron a otros hámsters con COVID y los juntaron a todos durante ocho horas.

La mayoría de los hámsters vacunados se infectaron, y el coronavirus se encontró después en las narices de 12 de 14 que recibieron la vacuna nasal y 15 de 16 que recibieron la inyección, dijeron los investigadores.

Sin embargo, los hámsteres que recibieron la vacuna nasal tenían niveles de virus en las vías respiratorias que eran de 100 a 100,000 veces más bajos que los de los hámsteres vacunados con una inyección, muestran los resultados.

En el segundo paso, los investigadores tomaron hámsters vacunados que se infectaron y los colocaron con otros hámsteres sanos durante ocho horas.

Ninguno de los hámsters expuestos a hámsters vacunados por la nariz se infectó, independientemente de si los nuevos hámsters habían sido vacunados o no, según muestran los resultados.

Por otro lado, alrededor de la mitad de los nuevos hámsters expuestos a los vacunados por inyección se infectaron, apuntaron los investigadores.

La vacunación por vía nasal rompió el ciclo de transmisión, concluyeron los investigadores.

"Las vacunas contra las mucosas son el futuro de las vacunas contra las infecciones respiratorias", dijo Boon. "Históricamente, el desarrollo de este tipo de vacunas ha sido un reto. Todavía hay mucho que no sabemos sobre el tipo de respuesta inmunitaria que necesitamos y cómo provocarla. Creo que vamos a ver muchas investigaciones muy interesantes en los próximos años que podrían conducir a grandes mejoras en las vacunas contra las infecciones respiratorias".

Fuente: infobae. Disponible en <https://acortar.link/L0KtAo>

Swissmedic authorises dengue fever vaccine

Aug 2. Qdenga (powder and solvent for solution for injection in a pre-filled syringe) can be placed on the market in Switzerland.

To date, no vaccine against dengue fever was authorised in Switzerland. At the end of July 2024, Swissmedic authorised the vaccine Qdenga, from manufacturer Takeda Pharma AG, after assessing its efficacy, safety and quality.

The vaccine is authorised for people aged 4 years and older who are travelling to regions in which dengue fever is prevalent. These primarily include subtropical and tropical regions in Central Africa, Latin America, India and Southeast Asia. The disease is also present in the south of the USA (Texas).

Qdenga helps to protect against dengue fever. The vaccine can be administered to adults, adolescents and children aged 4 years and older. It contains attenuated (weakened) versions of dengue virus variants 1, 2, 3 and 4. They cannot cause the disease but trigger the immune system (the body's natural defences) to defend the body against the virus. When a person receives the vaccine, their immune system recognises the attenuated variants as being foreign and forms antibodies against them. When they come into contact with the virus again, the body rapidly produces more antibodies to neutralise it before the person contracts dengue fever.

Dengue fever is a viral disease that is spread by infected mosquitoes (mainly yellow fever mosquitoes and Asian tiger mosquitoes). It usually develops between four and seven days after the person is bitten.

Symptoms include fever, headache, pain behind the eyes, muscle and joint pain, nausea or vomiting, swollen glands and rash.

Fuente: Swissmedic. Disponible en <https://acortar.link/QHSzqW>

Ante la preocupación por una nueva pandemia, un laboratorio argentino desarrollará la vacuna contra la gripe aviar

3 ago. El laboratorio argentino Sinergium Biotech comenzó la investigación para el desarrollo de una vacuna contra la gripe aviar dentro de un programa de la Organización Mundial de la Salud (OMS) que permitirá prepararse contra una posible pandemia. El proyecto trabaja con la tecnología del ARN mensajero (ARNm), usada contra la COVID-19, y se inicia en un contexto de crecientes contagios de virus H5N1 (de gripe aviar) en mamíferos. En junio se confirmó la primera muerte humana, un hombre en México sin antecedentes de contacto directo con aves silvestres.

Esta semana, la OMS anunció el nuevo proyecto de desarrollo y accesibilidad de vacunas experimentales de ácido ribonucleico mensajero (ARNm) contra la gripe aviar (H5N1) para los fabricantes de países de ingresos medianos y bajos. Forma parte del programa de Transferencia de Tecnología de ARNm de la OMS y el Medicines Patent Pool (MPP) y el laboratorio Sinergium Biotech desarrollará la vacuna para después compartir el conocimiento con otras empresas para agilizar la fabricación ante una posible pandemia. Este laboratorio forma parte del grupo Insud, fundado por Hugo Sigman y Silvia Gold. Otra empresa del grupo, mAbxience, estuvo a cargo de la producción del principio activo de la vacuna contra la COVID-19 de AstraZeneca, desarrollada por la Universidad de Oxford, para América Latina.

“Tenemos 14 diseños de candidatos vacunales, estamos en la etapa de descubrimiento y prueba de concepto. Son ensayos en animales en curso y ensayos in vitro. Los resultados vienen muy bien”, le dijo a elDiarioAR Germán Sánchez, gerente de I+D de la empresa argentina.

Una de las preocupaciones de los investigadores es la extensión acelerada del virus, no solo en aves sino también en mamíferos. En Estados Unidos se registraron brotes en vacas de ocho estados y se detectó el virus en leche no pasteurizada, que derivó en trabajadores con la enfermedad. En 2022, la gripe produjo mortalidad en lobos y elefantes marinos de Perú, Chile y Argentina. También está presente en la Antártida. Si bien en Europa la gripe aviar se mantiene solo en aves silvestres, están en alerta por la rapidez con la que se mueve el virus.

“Realmente lo que está ocurriendo a nivel animales es un brote, una pandemia a nivel zoonótico. Hay mucha potencialidad de que eso pueda seguir avanzando y el problema es cuando la influenza se empieza a mezclar. Principalmente porque este virus se puede llegar a mezclar con un virus estacional y de esa forma adquirir capacidades de ingresar al humano. Hoy en día no ingresa fácilmente al humano a menos que tenga algún evento de recombinación con otro virus como puede ser un virus estacional”, detalló Sánchez.

En junio de este año, la OMS confirmó la primera muerte por gripe aviar H5N2. Se trata de una persona de 59 años que murió el 24 de abril en México después de tener fiebre, dificultad respiratoria, diarrea, náuseas y malestar general. Se trata del primer caso humano confirmado en un laboratorio de una infección por el subtipo A(H5N2) de la gripe aviar notificado en todo el mundo. La víctima no tenía antecedentes de exposición a aves de corral u otros animales.

“Venimos detectando el comportamiento desde el 2002 y ahora se ha extendido a mamíferos. Esto nos

alentó a seguir incursionando en el desarrollo de la vacuna. Teniendo experiencia en la tecnología de la ARN, buscamos las secuencias de los genomas de los virus que estaban circulando. Lo que hicimos fue modificar la secuencias, extraer unas porciones del genoma del virus de influenza, de una proteína de superficie del virus que se llama hemaglutinina”, explicó Sánchez.

El laboratorio estima que para el 2025 tendrán los estudios preclínicos y toxicológicos y en lo que resta de este año buscan reducir los candidatos vacunales, que hoy son 14, en función de la inmunogenicidad y de los rendimientos de producción. El desarrollo de la vacuna podría estar listo para el año 2026, aunque los tiempos dependen también de la construcción de una nueva planta de tecnología de ARN que Sinergium Biotech comenzó en Argentina, una de las primeras en la región. “Es una vacuna que está pensada inicialmente para humanos, pero no quita que con pequeñas modificaciones se pueda llegar a vacunar animales. Arrancamos con H5N1 y después este programa va a migrar a otros tipos de influenza estacionales, como la vacuna de gripe”, anticipó el investigador.

Una vez que el laboratorio obtenga los datos preclínicos suficientes, va a compartir la información sobre técnicas y materiales con otros fabricantes que integran el Programa de Transferencia de Tecnología de ARNm para que puedan agilizar la fabricación de esas vacunas y reforzar la preparación contra una posible pandemia. Son empresas públicas, privadas o mixtas de países de ingresos medianos y bajos del Hemisferio Sur como Egipto, Kenia, Nigeria, Senegal, Túnez, Bangladesh, Indonesia, India, Pakistán, Serbia, Sudáfrica, Ucrania y Vietnam. La idea es que tengan la capacidad para producir vacunas de ARNm no solo para COVID-19, sino también para la producción de otras inmunizaciones críticas para el sistema de salud.

“Esta iniciativa ilustra a la perfección el motivo por el que la OMS creó el programa, que no es otro que el de impulsar la investigación, el desarrollo y la producción en países de ingresos medianos y bajos con el fin de prepararnos mejor para actuar eficaz y equitativamente en el caso de que se declare una pandemia”, dijo el director General de la OMS, Tedros Adhanom Ghebreyesus, durante la presentación del proyecto de gripe aviar.

“El programa intenta democratizar esta tecnología ARN, que en la pandemia estaba segmentada en Estados Unidos o en Alemania, con compañías que no transfirieron su tecnología. Lo que se vio es que así se logran respuestas más rápidas en pandemia. Tiene un rol muy importante a nivel social. Mientras más rápido sacás la vacuna, más vidas terminás salvando”, explicó Sánchez.

Fuente: el Diario AR. Disponible en <https://acortar.link/JgpzSu>

SK Bioscience's Investments Charge Momentum for Vaccine Production

Aug 5. SK Bioscience, which acquired Germany's IDT Biologika in June, announced on July 25th that it has entered into a conditional equity purchase agreement (SAFE) to invest approximately \$2.1 million in the U.S. biotech company Sunflower Therapeutics. This partnership is expected to create significant technological synergies between the two companies.



Founded in 2018, Sunflower Therapeutics is a biotech firm that has developed the 'Yeast Expression System,' a protein manufacturing technology essential for the development of antigens and antibodies.

This system simplifies the vaccine production process, reducing the time required and enhancing overall efficiency, thereby lowering manufacturing costs.

Through this SAFE investment, SK Bioscience aims to optimize its vaccine production processes using Sunflower's technology. The company anticipates that incorporating Sunflower's Yeast Expression System into the vaccine production line at its Andong L House facility could improve yield by up to 7.7 times compared to current methods. This enhancement is projected to reduce production costs per dose by 88.7%.

The two companies first established their relationship in 2023 while collaborating on the research and development of a human papillomavirus (HPV) vaccine. SK Bioscience plans to explore various strategies to maximize the value of its investment during Sunflower's potential future IPO and third-party mergers and acquisitions, while also expanding their technological collaboration.

An SK Bioscience representative stated, "We have been continuously collaborating with Sunflower since last year. Due to the potential changes in our equity stake depending on Sunflower's IPO and other related events, we cannot disclose the specific percentage of our holdings in Sunflower."

Meanwhile, in June, SK Bioscience invested approximately \$250 million to acquire IDT Biologika, a German contract development and manufacturing organization (CDMO) for biopharmaceuticals. IDT Biologika has an impressive track record recognized by regulatory agencies in the U.S., Europe, and more than ten other key pharmaceutical markets.

At a press conference on June 27th, Ahn Jae-yong, CEO of SK Bioscience, stated, "From SK Bioscience's perspective, the acquisition of IDT Biologika is a 'perfect fit.' There are complementary aspects between the company's Andong L House vaccine plant and IDT Biologika. By combining the software and hardware of both companies, we aim to create immediate synergies and expand our revenue."

SK Bioscience has expanded its production capacity with the acquisition of IDT Biologika and aims to optimize its vaccine processes through its investment in Sunflower Therapeutics. This move is seen as an effort to seek synergies in the field of vaccine development.

An SK Bioscience representative commented, "The investment in Sunflower considers the potential synergies with the acquisition of Germany's IDT. Regardless of the synergies from the IDT acquisition, various technologies from Sunflower can be utilized in multiple aspects at the Andong L House vaccine plant."

Fuente: Hit News. Disponible en <https://acortar.link/D7arta>

Las vacunas COVID reducen el riesgo de infarto y ACV, concluyó un estudio que evaluó a 46 millones de personas

5 ago. Las vacunas son un hito histórico en el sistema de salud global y cada año salvan entre 3,4 y 5 millones de vidas en todo el mundo. Al vacunarse, el sistema inmunológico de una persona se activa, por lo que se reduce el riesgo de contraer enfermedades prevenibles por inmunización.

Además, la inmunización brinda beneficios integrales para la salud, al proteger al organismo de diversas patologías



asociadas. En ese sentido, una nueva investigación mostró evidencia contundente sobre el papel de las vacunas contra el SARS-CoV-2 como escudo protector contra las enfermedades del corazón.

La incidencia de los infartos de miocardio y accidentes cerebrovasculares (ACV) ha disminuido significativamente tras la vacunación contra la COVID-19, según el estudio publicado en la revista *Nature Communications*. El trabajo fortalece la evidencia sobre los beneficios de la vacunación, no solo en términos de prevención de COVID-19, sino también en la reducción de incidentes cardiovasculares.

La investigación, que fue dirigida por las universidades británicas de Cambridge, Bristol y Edimburgo, junto con Centro de Ciencia de Datos de la Fundación Británica del Corazón (BHF, por sus sigla en inglés), analizó un vasto conjunto de datos pertenecientes a 46 millones de personas en Inglaterra -casi toda la población adulta del país- durante el periodo comprendido entre el 8 de diciembre de 2020 y el 23 de enero de 2022.

De acuerdo al análisis, la incidencia de trombosis arteriales, incluyendo infartos de miocardio y accidentes cerebrovasculares (ACV), se redujo hasta en un 10% en las 24 semanas posteriores a la administración de la primera dosis de la vacuna contra la COVID-19.

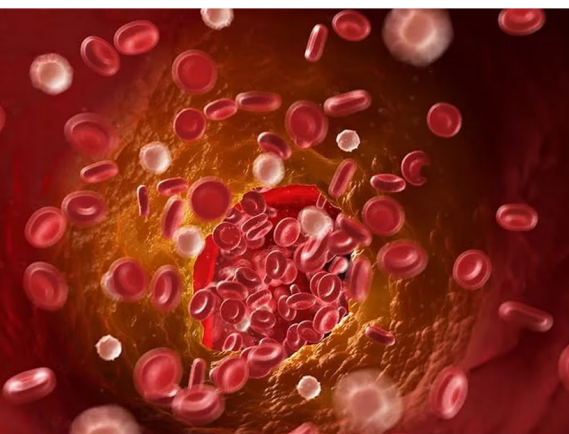
Este descenso fue aún más pronunciado tras la segunda dosis, alcanzando una disminución del 27% con la vacuna de AstraZeneca y del 20% con la vacuna de Pfizer/Biotech. “La vacunación demostró ser una herramienta clave no sólo en la prevención de la enfermedad grave por COVID-19, sino también en la reducción de complicaciones cardiovasculares”, afirmaron los investigadores en el mencionado estudio.



Además, el estudio analizó la incidencia de episodios trombóticos venosos, como la embolia pulmonar y la trombosis venosa profunda de las extremidades inferiores, trastornos que siguieron un patrón similar de reducción post-vacunación. “Estos hallazgos subrayan la eficacia del programa de vacunación en la protección integral contra diversas complicaciones de salud”, añadieron los investigadores.

El equipo de investigación del Centro de Ciencia de Datos de la Fundación Británica del Corazón (BHF) utilizó datos no identificados de consultas médicas, ingresos hospitalarios y registros de defunciones, asegurando que el análisis se realizara dentro de un entorno de datos seguro.

Complicaciones cardiovasculares raras post vacunación



Algunos estudios previos habían identificado una incidencia mayor de complicaciones cardiovasculares raras en ciertas vacunas contra la COVID-19, como la miocarditis y pericarditis (inflamación del corazón) asociada a las vacunas basadas en ARNm o la trombocitopenia trombotica inducida por vacunas basadas en adenovirus, como la de AstraZeneca.

Los científicos del estudio remarcaron que estas complicaciones fueron eventuales, confirmaron que los beneficios de la vacunación superan los riesgos asociados a efectos secundarios extremadamente raros y a las secuelas que deja en el organismo la infección por COVID-19.

En ese mismo sentido, la investigación publicada en *Nature Communications* suma tranquilidad al no identificar nuevas afecciones cardiovasculares adversas vinculadas a la vacunación. Este estudio refuerza la percepción de que los beneficios de la vacunación superan ampliamente los riesgos, incluso en grupos con preexistencias cardiovasculares.

Además, uno de los puntos clave del trabajo es la observación de que la incidencia de enfermedades cardiovasculares es mayor tras sufrir COVID-19, especialmente en los casos graves. Esta podría ser una de las razones por las que la vacunación resulta en una menor incidencia de infartos de miocardio y accidentes cerebrovasculares comparado con las personas no vacunadas.

El profesor Nicholas Mills, coautor del estudio y profesor de Cardiología en la Universidad de Edimburgo, manifestó: “Nuestros hallazgos confirman que la vacunación contra COVID-19 no solo es crucial para prevenir la enfermedad en sí, sino que también tiene un impacto positivo significativo en la reducción de complicaciones cardiovasculares”.

Por otro lado, el doctor Steven Liu de la Universidad de Cambridge, remarcó: “Este enorme estudio nos proporciona la evidencia más robusta disponible hasta la fecha sobre la relación entre la vacunación y la reducción de riesgos cardiovasculares”.

El especialista de Cambridge dijo que el trabajo representó un gran esfuerzo que mostró la importancia de las iniciativas de datos a gran escala para comprender mejor los efectos secundarios de las vacunas y resaltar sus beneficios adicionales. “Analizar datos de casi toda la población adulta de Inglaterra nos permite hacer observaciones muy amplias y representativas sobre los efectos de la vacunación”, afirmó el doctor Liu.

Los investigadores esperan que estos resultados incrementen la confianza pública en las vacunas y ayuden a mitigar cualquier reticencia frente a la vacunación.

El profesor William Whiteley, director asociado del Centro de Ciencias de Datos de BHF y profesor de Neurología y Epidemiología de la Universidad de Edimburgo, dijo: “El lanzamiento del programa de vacunación COVID-19 comenzó con fuerza en el Reino Unido, con más del 90% de la población mayor de 12 años vacunada con al menos una dosis en enero de 2022. Este estudio realizado en toda Inglaterra ofrece a los pacientes la tranquilidad de que la primera, segunda y dosis de refuerzo de las vacunas contra la COVID-19 son seguras para el sistema cardiovascular. Demuestra que los beneficios de la segunda dosis y de las dosis de refuerzo, con menos eventos cardiovasculares comunes (infartos de miocardio y accidentes cerebrovasculares) después de la vacunación, superan las complicaciones cardiovasculares muy poco frecuentes”.

La coautora principal, la Dra. Venexia Walker, investigadora asociada de la Universidad de Bristol, afirmó: “Dado el papel fundamental que desempeñan las vacunas contra la COVID-19 en la protección de las personas contra la COVID-19, es importante que sigamos estudiando los beneficios y los riesgos asociados a ellas. La disponibilidad de datos de toda la población nos ha permitido estudiar diferentes combinaciones de vacunas contra la COVID-19 y considerar complicaciones cardiovasculares poco frecuentes. Esto no habría sido posible sin la gran cantidad de datos a los que tenemos el privilegio de acceder y nuestras estrechas colaboraciones entre instituciones”.

Fuente: Infobae. Disponible en <https://acortar.link/hfvEKs>

Los casos de COVID-19 van al alza en el mundo y existe el riesgo de aparición de nuevas variantes

7 ago. La agencia sanitaria mundial alerta del aumento de contagios y estima poco probable que la oleada disminuya en el corto plazo. Asimismo, expresa preocupación por la baja en la cobertura de vacunación, sobre todo ante la posibilidad de que surja una cepa más virulenta que pudiera causar enfermedad grave.

Las infecciones por COVID-19 están aumentando en todo el mundo y es poco probable que disminuyan en el corto plazo, advirtió este martes la Organización Mundial de la Salud (OMS), que también destacó el riesgo de que pronto aparezcan variantes más graves del coronavirus.

"La COVID-19 todavía está muy presente" y circula en todos los países, dijo la directora de Prevención de Pandemias de la OMS, la doctora Maria Van Kerkhove.

En conferencia de prensa en Ginebra, la epidemióloga indicó que los datos del sistema de vigilancia de la agencia sanitaria, basados en centinelas en 84 países, informan en este momento que el porcentaje de pruebas positivas para SARS-CoV-2 ha aumentado durante varias semanas.

"Las pruebas con resultado positivo rebasan el 10%, pero la cifra fluctúa según la región. En Europa, ese porcentaje es superior al 20%", detalló.

Además, se han registrado nuevas olas de infección en América, Europa y el Pacífico occidental.

La vigilancia de las aguas residuales sugiere que la circulación del SARS-CoV-2 es de dos a 20 veces mayor que las cifras documentadas.

Circulación atípica durante el verano

Las tasas de circulación de infecciones tan elevadas en los meses de verano boreal son atípicas para los virus respiratorios, que tienden más bien a propagarse principalmente en temperaturas frías.

Sin embargo, en los últimos meses, muchos países han experimentado oleadas de COVID-19, independientemente de la temporada.

Esto está ocurriendo ahora mismo en los Juegos Olímpicos, "donde al menos 40 atletas dieron positivo", dijo Van Kerkhove.

Riesgo creciente de nuevas cepas

La experta subrayó que a medida que el virus continúa evolucionando y propagándose, existe el riesgo creciente de que aparezca una cepa más grave del virus que pueda evadir los sistemas de detección y no responder a la intervención médica.

"Estoy preocupada", enfatizó la especialista de la OMS, argumentando que con una cobertura de vacunación tan baja y una circulación tan grande, "si tuviéramos una variante que fuera más virulenta, la susceptibilidad de las poblaciones en riesgo a desarrollar una enfermedad grave sería enorme", acotó.

Fortalecer las campañas de vacunación

Si bien las admisiones hospitalarias, incluidas las de cuidados intensivos, siguen siendo mucho menores que durante el pico de la pandemia, la OMS instó a los gobiernos a fortalecer las campañas de vacunación, garantizando que los grupos de mayor riesgo reciban las vacunas al menos una vez cada doce meses.

"Como individuos, es importante tomar medidas para reducir el peligro de infección y enfermedad grave, incluyendo asegurarse de haber recibido una dosis de la vacuna contra el COVID-19 en los últimos doce

meses, especialmente si se pertenece a un grupo de riesgo”, enfatizó Van Kerkhove.

La OMS detalló que la disponibilidad de vacunas ha disminuido sustancialmente en los últimos 18 meses porque el número de productores de vacunas contra el coronavirus se ha reducido.

La doctora Van Kerkhove señaló que es muy difícil para los fabricantes mantener el ritmo de producción, al margen de que no hace falta que mantengan el paso de 2021 y 2022. “Pero seamos muy claros: existe un mercado para las vacunas contra la COVID-19 que ya están hechas.

En cuanto a las vacunas nasales, precisó que aún están en desarrollo y explicó que podrían abordar la transmisión, reduciendo así el riesgo de más variantes, infecciones y enfermedad grave.

Fuente: reliefweb. Disponible en <https://acortar.link/sIVALx>

VAX-31 is the Broadest Spectrum Pneumococcal Conjugate Vaccine in the Clinic Today

Aug 7. Vaxcyte, Inc. today announced financial results for the second quarter ended June 30, 2024, and provided a business update.

“We continue to make significant strides toward building the potentially best-in-class pneumococcal conjugate vaccine (PCV) franchise and expect to announce the VAX-31 adult Phase 1/2 study topline safety, tolerability and immunogenicity data in September,” said Grant Pickering, Chief Executive Officer and Co-founder of Vaxcyte, in a press release on August 6, 2024.

“Our clinical program assessing VAX-31, the broadest-spectrum PCV in the clinic today, will provide significant insights into the full potential of this vaccine candidate across the adult population.”

“Following the VAX-31 adult data readout, we plan to advance either VAX-24 or VAX-31 into Phase 3 clinical development in adults.”

Mr. Pickering continued, “Additionally, we look forward to delivering the topline data from the primary immunization series of the VAX-24 infant Phase 2 study by the end of the first quarter of 2025, followed by topline data from the booster dose by the end of 2025.”

“We believe VAX-24 has a potential best-in-class profile for this vital population and is designed to cover more serotypes than any infant pneumococcal vaccine on-market today.”

The Company also confirmed cash, cash equivalents, and investments were \$1,851.9 million as of June 30, 2024, compared to \$1,242.9 million as of December 31, 2023.

Fuente: Precision Vaccinations. Disponible en <https://acortar.link/tHxKr5>

MSD Korea introduces Vaxneuvance: The first pneumococcal vaccine with proven immunogenicity

Aug 7. “Vaxneuvance, a pneumococcal protein conjugate vaccine introduced after 13 years, has demonstrated either superior immunogenicity or at least comparable effectiveness to the existing PCV13 vaccine in all 15 serotypes, including the newly added 22F and 33F.”



MSD Korea, the Korean branch of the multinational pharmaceutical company MSD (Merck, USA), hosted a media seminar on August 6 at Seoul Square in Jung-gu District, Seoul. The event, themed “Vaxneuvance: Opening the Era of Pneumococcal Vaccine Immunogenicity,” highlighted the vaccine’s strengths. Key speakers included Cho Min-hee, Head of the External Cooperation Division of MSD Korea, Cho Jae-yong, Executive Director of the Vaccine Business Division of MSD Korea, and Professor Kang Hyun-mi of the Department of Pediatrics at the Catholic University of Korea Seoul St. Mary’s Hospital.

By adding the two important serotypes, 22F and 33F, identified as major serotypes causing pneumococcal diseases worldwide, into the 13 serotypes, Vaxneuvance has expanded its scope of protection. The vaccine has shown immunogenicity that aligns with all World Health Organization (WHO) standards for each of the 15 pneumococcal serotypes (1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F, 22F, 23F, 33F).

At the seminar, Professor Kang Hyun-mi of the Department of Pediatrics at the Catholic University of Korea Seoul St. Mary’s Hospital delivered a presentation on “A Time of Great Change in the Pneumococcal Vaccine Market, Vaccine Selection Criteria, and the Clinical Value of Vaxneuvance.” She noted that 34% of Korean invasive pneumococcal disease cases were caused by serotypes 19A, 19F, and 3, while 66% were caused by 23A. Although the incidence of invasive pneumococcal disease in Korea has significantly decreased since the introduction of PCV13, the incidence of disease caused by 23A has remained unchanged.



"When developing a new PCV, there is a risk that including additional serotypes could reduce the vaccine’s immunogenicity against the common ones," Kang stated. This suggests that the effectiveness of the vaccine in preventing invasive pneumococcal disease might be reduced.

In fact, an analysis of the main phase 3 clinical trial results revealed that immunogenicity for common serotypes decreased as the number of serotypes in the PCV increased. Consequently, the WHO defines 'immunogenicity' as the capacity of a vaccine to produce a measurable immune response.

Vaxneuvance, which includes 15 pneumococcal serotypes, has been proven safe and immunogenic in over 20,000 subjects. In particular, the phase 3 clinical trial (PNEU-PED) demonstrated Vaxneuvance’s non-inferior immunogenicity for the 13 common serotypes and stronger immunogenicity for serotypes 3, 22F, and 33F.

Since April, Vaxneuvance has been part of the National Immunization Program (NIP) for children, allowing free administration at hospitals and clinics nationwide. The NIP targets all children aged 2 months to under 5 years, including those who have not yet been vaccinated with the pneumococcal conjugate vaccine or have not completed their vaccination schedule.

Clinical studies on children have confirmed that Vaxneuvance can be cross-vaccinated with the existing PCV13. Thus, children who have received at least one dose of PCV13 can cross-vaccinate with Vaxneuvance for the remaining recommended vaccinations. "Since its inclusion in the NIP, Seoul St. Mary’s Hospital has started vaccinating with the 15-valent vaccine," Professor Kang expressed. "The vaccination rate is high," she added.

Fuente: THE BIO NEWS. Disponible en <https://acortar.link/azUISE>

El VRS suele preceder a las infecciones bacterianas por neumococo

El virus respiratorio sincitial (VRS) y el neumococo son dos de los patógenos respiratorios (el primero, un virus; el segundo, una bacteria) más comunes que afectan tanto a niños como a adultos mayores. Su interacción es muy relevante en el contexto de las infecciones respiratorias, pues, tal y como destaca Federico Martín-Torres, jefe de Pediatría y director de Pediatría Traslacional y Enfermedades Infecciosas en el Hospital Clínico Universitario de Santiago de Compostela, en declaraciones a GM, “el VRS suele preceder a las infecciones bacterianas por neumococo.

“Este fenómeno se debe a que el VRS puede dañar el epitelio respiratorio, lo que facilita la adhesión y colonización del neumococo”, explica. De hecho, a nivel molecular, se ha demostrado su interacción directa. “El VRS aumenta la expresión de receptores en las células epiteliales respiratorias que el neumococo utiliza para adherirse e invadir los tejidos; la coinfección es frecuente y puede conducir a complicaciones severas como la neumonía y la sepsis”, apunta Martín-Torres.

Interacción entre VRS y neumococo

En este sentido, la interacción entre este virus y esta bacteria quedó patente durante la pandemia de COVID-19. El experto afirma que, al dejar de circular el VRS y, por tanto, reducirse las infecciones a consecuencia del virus, también cayó la incidencia de la enfermedad neumocócica. Eso sí, Martín-Torres aclara que, aunque se redujeron los casos de enfermedad, no disminuyó la transmisión de neumococo de persona a persona, es decir, la “tasa de portadores de neumococo era similar a la pre-pandemia”.

Estas interacciones entre VRS y neumococo “varían con la edad”.

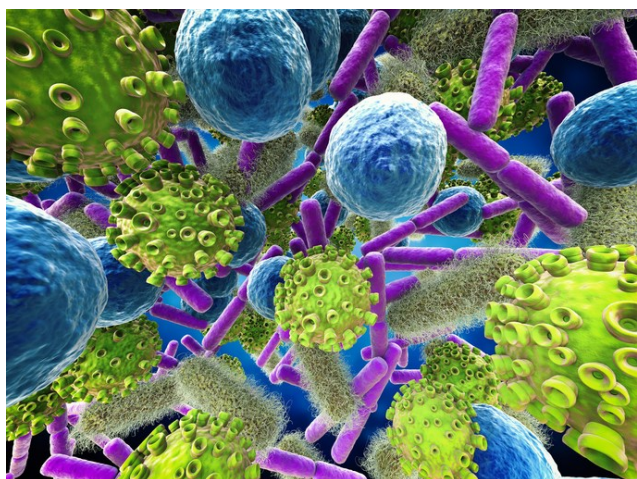
En niños, las infecciones son “más frecuentes y severas” debido a la “inmadurez” de su sistema inmunológico, mientras que en los adultos, especialmente en aquellos que padecen comorbilidades o tienen una edad avanzada, “la severidad de las infecciones puede ser alta” por la disminución de la respuesta inmunitaria y la presencia de enfermedades crónicas como la diabetes o la enfermedad pulmonar obstructiva crónica (EPOC).

“La diferencia en la respuesta inmunitaria se debe principalmente a la ontogenia del sistema inmunológico en los niños y la inmunosenescencia en los adultos mayores”, resume Martín-Torres. “Durante la pandemia, se observó que las infecciones respiratorias en adultos mayores tenían peores desenlaces, resaltando la necesidad de entender las diferencias inmunológicas por edad, entre otros factores”, agrega.

La importancia de la inmunización

Ante este hecho, el experto apunta a la importancia de la inmunización, “herramienta clave para prevenir infecciones respiratorias”. “La vacuna neumocócica, especialmente la vacunación conjugada, ha reducido significativamente la incidencia de enfermedades invasivas por neumococo en todas las edades”, asegura. También recuerda que está disponible la inmunización frente al VRS en lactantes con el anticuerpo monoclonal nirsevimab o protección pasiva a través de vacunas VRS para la gestante, así como en adultos con “tres vacunas disponibles que han demostrado una gran eficacia”.

Respecto a una posible inmunización cruzada, Martín-Torres es precavido: “Es pronto para sacar



conclusiones sobre su interacción, sin embargo, sabemos que la vacunación frente a neumococo, reduce la incidencia de infecciones respiratorias de cualquier etiología y específicamente de infecciones virales graves". "Y en el caso del nirsevimab, hemos visto también una reducción de las infecciones respiratorias de cualquier causa, y no solo del VRS", añade.

"De algún modo esto apunta a la interrelación que existe entre virus y bacterias y más específicamente entre neumococo y VRS, reforzando el papel de protección sinérgica que estas medidas de prevención pueden tener", concluye el experto.

Además, la prevención es más importante si cabe en el contexto de la conocida como 'triple epidemia', es decir, circulación simultánea de VRS, gripe y COVID-19. "Para abordar esta situación, se recomienda una estrategia integral en la que, sin duda, la vacunación es un pilar fundamental", asevera. Porque, "al margen de las interacciones específicas entre estos patógenos y con otros patógenos, no debemos olvidar que la infección secuencial también tiene consecuencias muy graves", explica Martín-Torres.

De este modo, de cara a la próxima temporada de infecciones respiratorias, la vacunación es "la clave más potente". "Es esencial tener el calendario vacunal al día, siguiendo las recomendaciones oficiales e independientemente de la edad", destaca el experto, quien insiste especialmente en los más vulnerables y susceptibles a estos tres virus y también frente al neumococo: personas de edad avanzada, personas con enfermedades de base cardíacas, respiratorias o metabólicas, inmunodeprimidos, etc.

"Los cuatro (VRS, gripe, COVID-19 y neumococo) explican la gran mayoría de las infecciones respiratorias graves que podemos padecer y hoy son prevenibles mediante vacunación en la mayor parte de los casos", concluye.

Fuente: Gaceta Médica. Disponible en <https://acortar.link/DUN0jw>

V116 Vaccine Supported for Adult Pneumococcal Disease Prevention in Phase 3 Trial

Aug 9. For adult patients, V116, an investigational 21-valent pneumococcal conjugate vaccine (PCV21) is noninferior to PCV20 for serotypes shared by both vaccines and superior to PCV20 for all serotypes unique to V116, aside from 15C, according to study results published in *The Lancet Infectious Diseases*.

Researchers conducted a randomized, double-blind, active comparator controlled, international phase 3 trial (ClinicalTrials.gov Identifier: NCT05425732) to evaluate the safety, tolerability, and immunogenicity of V116. Adults with no history of pneumococcal vaccination were eligible for the analysis and enrolled across 2 cohorts. Cohort 1 was stratified by age (50-64, 65-74, 75-84, and ≥ 85 years) and comprised patients aged 50 years and older who were randomly assigned 1:1 to receive 1 dose of either V116 or PCV20. Cohort 2 comprised patients aged between 18 and 49 years who were randomly assigned 2:1 to receive 1 dose of either V116 or PCV20. The primary safety outcomes were the percentage of patients with solicited injection-site and systemic adverse events (AEs) through day 5 and vaccine-related severe AEs up to 6 months after vaccination. The primary immunogenicity outcomes were as follows:

- ⇒ Noninferiority of V116 to PCV20 was tested using serotype-specific opsonophagocytic activity geometric mean titers (GMT) ratios for serotypes common to both vaccines (cohort 1);
- ⇒ Superiority of V116 to PCV20 was tested for opsonophagocytic GMT ratios for the serotypes unique to V116 (cohort 1);

- ⇒ Superiority of V116 to PCV20 was tested by the percentage of patients with at least a 4-fold rise from day 1 to day 30 for serotypes unique to V116 (cohort 1); and
- ⇒ Immunobridging for all 21 serotypes in V116 for adults aged 18 to 49 and 50 to 64 years (cohort 2).

A total of 2663 patients (women, 58.7%; resident of US, 47.4%) were included in the analysis and received V116 (cohort 1, n=1181; cohort 2, n=201) or PCV20 (cohort 1, n=1181; cohort 2, n=100). Among patients in the first cohort, 1179 (99.8%) and 1177 (99.7%) received V116 and PCV20, respectively. Among patients in the second cohort, 200 (99.5%) and 100 (100%) received V116 and PCV20, respectively.

In cohort 1, V116 vs PCV20 met noninferiority criteria for all 10 serotypes common to both vaccines (all $P < .0001$), as well as superiority criteria for 10 of the 11 serotypes unique to V116 ($P < .0001$ for all unique serotypes except 15C, which was $P = .41$).

Further, V116 vs PCV20 met superiority criteria for 10 of 11 serotypes unique to V116 based on the percentage of patients with at least a 4-fold rise in opsonophagocytic activity from day 1 to day 30 ($P < .0001$ for all unique serotypes except 15C, which was $P = .67$). Moreover, immune responses among V116 recipients were noninferior between those aged 18 to 49 vs 50 to 64 years for all V116 serotypes (all $P < .0001$).

“Real-world evidence will be needed to assess vaccine effectiveness for the prevention of vaccine-type pneumococcal disease.”

In regard to safety, 685 (58.2%) V116 recipients and 778 (66.2%) PCV20 recipients in cohort 1 and 164 (82.0%) V116 recipients and 79 (79.0%) PCV20 recipients in cohort 2 reported at least 1 AE.

In cohort 1, mortality occurred among 4 patients who received V116 and 2 patients who received PCV20. No deaths were considered related to vaccination. For V116 recipients, mortality was attributed to sepsis (n=1), cerebrovascular accident (n=1), myocardial infarction (n=1), and hepatic cirrhosis plus hepatic encephalopathy (n=1). For PCV20 recipients, mortality was attributed to cardiac arrest (n=1) and abdominal abscess (n=1). There were no vaccine-related severe AEs reported.

Study limitations include the lack of data on frailty status, the inability to assess the effect of V116 vaccination on nasopharyngeal pneumococcal carriage, and the exclusion of older patients (≥ 75 years), those with a history of pneumococcal vaccination, and those with immunocompromising conditions.

“Real-world evidence will be needed to assess vaccine effectiveness for the prevention of vaccine-type pneumococcal disease,” the researchers concluded.

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Fuente: Infectious Disease Advisor. Disponible en <https://acortar.link/LJZBMj>

22nd Country Authorizes Second-Gen Dengue Vaccine

Aug 11. At the end of July 2024, Swissmedic authorized Takeda Pharma AG's QDENGAR[®] (TAK-003) Tetravalent Vaccine (Live, Attenuated) vaccine after assessing its efficacy, safety, and quality.

To date, Switzerland has not authorized a dengue fever prevention vaccine. However, about 21 other countries have issued QDENGAR authorization.

According to Swissmedic's statement on August 2, 2024, QDENGAR is authorized for people aged four years and older who travel to regions where dengue fever is prevalent. These areas primarily include subtropical and tropical regions in Central Africa, Latin America, India, and Southeast Asia.

The U.S. Centers for Disease Control and Prevention (CDC) recently announced that the global incidence of dengue this year is the highest in recent history, with about 100 countries reporting higher-than-usual dengue cases.

In 2024, dengue infections, both travel-related and locally transmitted, were confirmed in the south of the USA, in states such as Florida (335) and Texas (18).

Swissmedic says QDENGAR cannot cause the disease, but vaccination triggers the immune system to defend the body against the virus. When a person receives the vaccine, their immune system recognizes the attenuated variants as foreign and forms antibodies against them. When they come into contact with the virus again, the body rapidly produces more antibodies to neutralize it before the person contracts dengue fever.

Dengue fever is a viral disease spread by infected mosquitoes. It usually develops between four and seven days after the person is bitten. Symptoms include fever, headache, pain behind the eyes, muscle and joint pain, nausea or vomiting, swollen glands, and rash.

As of August 11, 2024, QDENGAR is unavailable in the USA.

Fuente: Precision Vaccinations. Disponible en <https://acortar.link/PGQFwO>

New vaccine for respiratory disease rolls out in Scotland

Aug 12. A new vaccination programme aimed at protecting newborn babies and older adults against a dangerous respiratory disease is now being rolled out in Scotland.

The Respiratory Syncytial Virus (RSV) immunisation programme begins on Monday morning, and will be offered in the other UK nations from September.

RSV is common and highly infectious. It affects the breathing system and can cause severe illness in vulnerable groups, including infants and older people.

It is the leading cause of emergency respiratory admissions to hospital in infants.

In 2022-23, more than 1,500 infants under the age of one and more than 500 people aged 75 and over were



hospitalised with RSV, according to Public Health Scotland.

Across the UK as a whole it results in 25-30 infant deaths each year.

While for many the symptoms are mild, the infection is easily spread and 90% of children will catch it within the first two years of their lives.

Winter pressures on NHS

The vaccine is being administered on the advice of the UK's Joint Committee on Vaccination and Immunisation (JCVI).

Doses are being offered to women from 28 weeks into their pregnancies, to protect newborns, as well as those aged 75 and as a one-off catch up for those aged 75 to 79.

Dr Sam Ghebrehewet, head of immunisation and vaccination at Public Health Scotland, said: "RSV can be very serious for those who are more vulnerable, such as newborns, infants and older adults.

"If you are eligible, getting vaccinated is the best and simplest thing you can do to protect yourself or your newborn baby from RSV."

First Minister John Swinney said Scotland's early rollout of the vaccination programme could help relieve winter pressures on the NHS.



He added: "It is equally important that those aged 75-79 take up their offer of this vaccine."

What are the symptoms of RSV?

Symptoms of RSV usually start within a few days of getting infected.

According to the NHS, most people only get cold-like symptoms, such as:

- ⇒ a runny or blocked nose
- ⇒ sneezing
- ⇒ a cough
- ⇒ tiredness
- ⇒ a high temperature

Babies with RSV may also be irritable and feed less than usual.

If RSV leads to a more serious infection (such as bronchiolitis) it may also cause:

- ⇒ a cough that gets worse
- ⇒ faster breathing or long gaps between breaths
- ⇒ difficulty feeding or eating
- ⇒ noisy breathing (wheezing)

Fuente: BBC. Disponible en <https://acortar.link/NTMXyt>



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EP - 07.08.2024

Clasificación Internacional [A61K 39/00](#)^N de solicitud 22873232 Solicitante CHA [VACCINE](#) RES INSTITUTE CO LTD Inventor/a YUM JUNG SUN

The present invention pertains to: an anti-cancer [vaccine](#) composition comprising [peptides derived from a tumor-associated antigen (TAA)] and [an adjuvant consisting of a lipopeptide and an immunoactive substance]; and a use thereof. Specifically, the peptides derived from a tumor-associated antigen specifically bind to a human leukocyte antigen (HLA), a combination of the peptides having the above characteristics is mixed with the adjuvant in an optimal ratio to prepare a [vaccine](#) composition, and the [vaccine](#) composition is used for preventing or treating cancer.

2. [4410305](#) IMPFSTOFFZUSAMMENSETZUNG GEGEN KREBS MIT HSP90-ANTIGENPEPTID UND VERWENDUNG DAVON

EP - 07.08.2024

Clasificación Internacional [A61K 39/00](#)^N de solicitud 22876850 Solicitante ASTON SCI INC Inventor/a KANG JIN HO

The present disclosure relates to an anticancer [vaccine](#) composition including a peptide of SEQ ID NO: 1 and a peptide of SEQ ID NO: 2, which are epitopes of HSP90, and the [vaccine](#) composition according to the present disclosure may effectively inhibit the growth of tumors in an animal model of tumor cell line transplantation without severe adverse effects, and thus may be useful for treating cancer or preventing cancer recurrence.

3. [4408483](#) COVID19-MRNA-IMPFSTOFF

EP - 07.08.2024

Clasificación Internacional [A61K 48/00](#)^N de solicitud 22877397 Solicitante UNIV TEXAS Inventor/a HU HAITAO

A solution has been discovered that provides a more effective Coronavirus [vaccine](#). The solution is an mRNA [vaccine](#) encoding a SARS-CoV-2 nucleoprotein (N) (mRNA-N) in combination with an mRNA [vaccine](#) encoding SARS-CoV-2 spike protein (S) (mRNA-S). Chemically modified mRNA-N (pseudouridine) and/or chemically modified mRNA-S (pseudouridine) can be synthesized and packaged in lipid nanoparticles (LNP). In mouse and hamster models, it was shown that mRNA-N alone is immunogenic and can significantly diminish viral loads in the mouse lung after prime-boost intramuscular immunization. In addition, the combinatorial mRNA-N/mRNA-S vaccination induces substantially stronger protection against SARS-CoV-2 than vaccination with mRNA-S alone.

4. [WO/2024/161361](#) MODIFIED BCG [VACCINE](#)

WO - 08.08.2024

Clasificación Internacional [A61K 39/04](#)^N de solicitud PCT/IB2024/050966 Solicitante UNIVERSITY OF THE WITWATERSRAND, JOHANNESBURG Inventor/a KANA, Bavesh Davandra

The present invention relates to a recombinant Mycobacterium bovis BCG strain comprising a plasmid having a short guide RNA (sgRNA) target sequence for knockdown of the Mb3739 gene. Also provided are [vaccine](#) compositions for eliciting an immune response against Mycobacterium tuberculosis comprising the recombinant Mycobacterium bovis BCG strain engineered to activate the NOD-1 pathway. The recombinant Mycobacterium bovis BCG strain or the [vaccine](#) compositions may be useful in methods of eliciting an immune response against Mycobacterium tuberculosis. The invention also relates to methods of obtaining the recombinant Mycobacterium bovis BCG strain.

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5. [20240261387](#) RESPIRATORY SYNCYTIAL VIRUS MRNA [VACCINE](#)

US - 08.08.2024

Clasificación Internacional [A61K 39/155N](#)° de solicitud 18394486 Solicitante Vernagen, LLC Inventor/a Baek KIM

Provided herein are a respiratory syncytial virus (RSV) [vaccine](#) composition including a messenger ribonucleic acid (mRNA) including an open reading frame (ORF) encoding RSV mutant F B strain protein, and optionally a mRNA including an ORF encoding RSV mutant F A strain protein, and a method of inducing immune response against RSV by administering an effective amount of the RSV [vaccine](#) composition to a subject in need thereof. Provided herein are also a respiratory syncytial virus (RSV) and human metapneumovirus virus (hMPV) [vaccine](#) composition including a mRNA including an ORF encoding RSV mutant F A strain protein, a mRNA including an ORF encoding RSV mutant F B strain protein, and a mRNA including an ORF encoding hMPV F protein, and a method of inducing immune response against RSV and hMPV by administering an effective amount of the RSV and hMPV [vaccine](#) composition to a subject in need thereof.

6. [WO/2024/163092](#) RESPIRATORY SYNCYTIAL VIRUS MRNA [VACCINE](#)

WO - 08.08.2024

Clasificación Internacional [A61K 39/12N](#)° de solicitud PCT/US2023/085663 Solicitante VERNAGEN, LLC Inventor/a KIM, Baek

Provided herein are a respiratory syncytial virus (RSV) [vaccine](#) composition including a messenger ribonucleic acid (mRNA) including an open reading frame (ORF) encoding RSV mutant F B strain protein, and optionally a mRNA including an ORF encoding RSV mutant F A strain protein, and a method of inducing immune response against RSV by administering an effective amount of the RSV [vaccine](#) composition to a subject in need thereof. Provided herein are also a respiratory syncytial virus (RSV) and human metapneumovirus virus (hMPV) [vaccine](#) composition including a mRNA including an ORF encoding RSV mutant F A strain protein, a mRNA including an ORF encoding RSV mutant F B strain protein, and a mRNA including an ORF encoding hMPV F protein, and a method of inducing immune response against RSV and hMPV by administering an effective amount of the RSV and hMPV [vaccine](#) composition to a subject in need thereof.

7. [WO/2024/156912](#) ANTI-ABETA [VACCINE](#) THERAPY

WO - 02.08.2024

Clasificación Internacional [A61K 39/00N](#)° de solicitud PCT/EP2024/052002 Solicitante AC IMMUNE SA Inventor/a STREFFER, Johannes

A liposomal [vaccine](#) composition comprising a !-amyloid ($A\beta$)-derived peptide antigen displayed on the surface of the liposome that comprises, consists essentially of or consists of amino acids 1-15 of $A\beta$, a peptide comprising a universal T-cell epitope and an adjuvant comprising monophosphoryl lipid A (MPLA) is used for inducing an anti- $A\beta$ immune response in a human subject without inducing a serious adverse event. The !-amyloid ($A\beta$)-derived peptide antigen (SEQ ID NO: 1) is administered in an amount of 300-2000 μ g. The liposomal [vaccine](#) composition is administered intramuscularly or subcutaneously.

8. [WO/2024/163918](#) MRNA [VACCINE](#) FOR HERPES SIMPLEX VIRUS

WO - 08.08.2024

Clasificación Internacional [A61K 39/12N](#)° de solicitud PCT/US2024/014282 Solicitante UNIVERSITY OF HOUSTON SYSTEM Inventor/a ZHANG, Shaun

This invention relates to an mRNA-based [vaccine](#) composition for strains of herpes simplex virus (HSV), such as herpes simplex virus-2 (HSV-2). The mRNA [vaccine](#) composition comprises a combination of herpesvirus glycoprotein D (gD) and gB-pf mRNA, either alone or in combination with gE and/or gC mRNA.

9. [20240252606](#) PHOSPHORYLATED POLYPEPTIDE ANTIGEN [VACCINE](#), PREPARATION METHOD THEREFOR AND APPLICATION THEREOF

US - 01.08.2024

Clasificación Internacional [A61K 39/00N](#)° de solicitud 17059665Solicitante CHANGCHUN BCHT BIOTECHNOLOGY CO.Inventor/a Wei KONG

The present invention discloses a phosphorylated polypeptide antigen [vaccine](#), comprising at least two polypeptide fragments or conservatively modified variants thereof from human full-length Tau protein, wherein the polypeptide fragments or conservatively modified variants thereof contain phosphorylation sites. The present invention also discloses a complex [vaccine](#) formed by coupling a phosphorylated polypeptide antigen [vaccine](#) with a carrier. The polypeptide antigen [vaccine](#) and the complex [vaccine](#) can be used for preventing and/or treating tauopathy comprising Alzheimer's disease (AD).

10.[20240261398](#)ADJUVANT AND [VACCINE](#) CONTAINING ADJUVANT

US - 08.08.2024

Clasificación Internacional [A61K 39/39N](#)° de solicitud 18634012Solicitante The University of TokyoInventor/a Yoshihiro KAWAOKA

The present invention is intended to provide an adjuvant having high safety to living bodies and an action to sufficiently reinforce immune function, and a [vaccine](#) comprising the adjuvant. Specifically, the present invention relates to 34 novel adjuvant candidate compounds, which have been identified by screening 145 food additives and 51 injection additives, using, as indicators, an increase in the antibody titer against influenza virus and a protective effect against infection with influenza virus, and then selecting those having the function of increasing the antiviral antibody titer in blood and the protective effect against viral infection. In addition, the present invention also relates to a [vaccine](#) comprising these adjuvant candidate compounds.

11.[20240261383](#)RECOMBINANT FUSION PROTEIN [VACCINE](#) CONTAINING CLOSTRIDIODES DIFFICILE FLIC AND FLID

US - 08.08.2024

Clasificación Internacional [A61K 39/08N](#)° de solicitud 18390057Solicitante University of South FloridaInventor/a Xingmin Sun

A novel [vaccine](#) and methods of preventing and treating *C. difficile* infection in a patient is described. The [vaccine](#) is comprised of a fusion protein (denoted FliCD) comprised of the FliC and FliD from *C. difficile* and joined by a linker sequence. Administration of the [vaccine](#), as well as anti-FliCD serum, has been shown to prevent *C. difficile* infection as well as treat existing infections.

12.[WO/2024/163138](#)RECOMBINANT FUSION PROTEIN [VACCINE](#) CONTAINING CLOSTRIDIODES DIFFICILE FLIC AND FLID

WO - 08.08.2024

Clasificación Internacional [A61K 38/16N](#)° de solicitud PCT/US2024/010819Solicitante UNIVERSITY OF SOUTH FLORIDAInventor/a SUN, Xingmin

A novel [vaccine](#) and methods of preventing and treating *C. difficile* infection in a patient is described. The [vaccine](#) is comprised of a fusion protein (denoted FliCD) comprised of the FliC and FliD from *C. difficile* and joined by a linker sequence. Administration of the [vaccine](#), as well as anti-FliCD serum, has been shown to prevent *C. difficile* infection as well as treat existing infections.

13.[WO/2024/159517](#)HEPATITIS B MRNA AND [VACCINE](#) AND USE THEREOF

WO - 08.08.2024

Clasificación Internacional [C12N 15/51N](#)° de solicitud PCT/CN2023/074384Solicitante SOUTHERN UNIVERSITY OF SCIENCE AND TECHNOLOGYInventor/a WANG, Peng

Provided are a hepatitis B mRNA and a [vaccine](#) and use thereof. A protein encoded by an mRNA molecule contains at least one of a hepatitis B PreS1 antigen protein, a hepatitis B core antigen protein, a hepatitis B polymerase, and a polymerase T cell epitope enriched fragment. The use of an mRNA-LNP [vaccine](#) for immunizing healthy and HBV model mice can induce strong humoral immune response and wide cellular immune response, and can significantly improve viral serological indicators of HBV model mice, thus indicating that the [vaccine](#) has the potential of clinical functional cure of HBV.

14.[20240252627](#)MESENCHYMAL STEM CELLS AS [VACCINE](#) ADJUVANTS AND METHODS FOR USING THE SAME

US - 01.08.2024

Clasificación Internacional [A61K 39/39N](#)° de solicitud 18629427Solicitante Longeveron Inc.Inventor/a Joshua M. HARE

The present invention provides a method of enhancing an immune response to a [vaccine](#) by administering a [vaccine](#) and a population of isolated allogeneic human mesenchymal stem cells. The present invention also provides kits comprising a [vaccine](#) in a first container and a population of isolated allogeneic human mesenchymal stem cells in a second container.

15. [20240261386](#)COXSACKIEVIRUS B3 [VACCINE](#)

US - 08.08.2024

Clasificación Internacional [A61K 39/125](#)Nº de solicitud 18165762Solicitante KING FAISAL UNIVERSITYInventor/a JAWHAR GHARBI

The Coxsackievirus B3 (CVB3) [vaccine](#) including a mutant strain of Coxsackievirus B3 (CVB3) (SEQ ID NO: 1) has specific double mutations introduced in the Internal Ribosome Entry Segment (IRES) region of the wild type Coxsackievirus B3 (CVB3) genome in the nucleotide positions 473 (in which uracil is substituted for cytosine) and 475 (in which cytosine is substituted for uracil). The resulting double mutant (SEQ ID NO: 1) demonstrates a significant decrease in its replicative capacity and a drastic decrease in its translation efficiency compared to the wild-type Coxsackievirus B3 (CVB3) strain.

16. [20240252616](#)LIVE-ATTENUATED VIRUS [VACCINE](#)

US - 01.08.2024

Clasificación Internacional [A61K 39/215](#)Nº de solicitud 18016137Solicitante GRIFFITH UNIVERSITYInventor/a Surendran Mahalingam

This invention relates to a codon deoptimized severe acute respiratory syndrome coronavirus 2 (SARS-COV-2) genome. In particular, embodiments of the invention concern a [vaccine](#) comprising live attenuated SARS-COV-2 comprising a partly codon deoptimized viral genome, SARS-COV-2 comprising a partly codon deoptimized viral genome, as well as their use in methods of treatment and prevention of viral infection. The ORF1a region of the viral genome has been codon deoptimized.

17. [20240261391](#)TLR7 AGONIST CONJUGATED PEPTIDE-BASED NOVEL CORONAVIRUS NANOEMULSION [VACCINE](#) AND PREPARATION THEREOF

US - 08.08.2024

Clasificación Internacional [A61K 39/215](#)Nº de solicitud 18565013Solicitante SHANGHAI INSTITUTE OF MATERIA MEDICA, CHINESE ACADEMY OF SCIENCESInventor/a Xinxin ZHANG

The present invention relates to a novel coronavirus [vaccine](#) using a TLR7 agonist conjugated peptide as an antigen and an emulsion as an adjuvant. An antigen polypeptide of the conjugated peptide is a polypeptide derived from an S protein of SARS-COV-2, and the adjuvant is an oil-in-water nanoemulsion containing squalene. The conjugated peptide nanoemulsion [vaccine](#) preparation of the present invention is thermally stable, and can induce a high level of protective humoral immune response in a cynomolgus monkey, and the neutralizing antibody titer of antiserum after immunization of cynomolgus monkey is high, such that invasion of wild-type strain and mutant novel coronavirus can be blocked. The [vaccine](#) of the present invention has a nearly complete protection effect on the upper and lower respiratory tracts of the cynomolgus monkey in a cynomolgus monkey SARS-COV-2 challenge test. The nanoemulsion [vaccine](#) of the present invention is fast and convenient to prepare, and can realize large-scale production in a short term for coping with the novel coronavirus outbreak.

18. [20240252620](#)COMBINED AGONIST ADJUVANT FOR CORONAVIRUS [VACCINE](#)

US - 01.08.2024

Clasificación Internacional [A61K 39/215](#)Nº de solicitud 18564263Solicitante THE REGENTS OF THE UNIVERSITY OF MICHIGANInventor/a Pamela Wong

The disclosure is directed to compositions and methods for inducing an immune response against a coronavirus, which involve a coronavirus [vaccine](#) and an adjuvant composition. The adjuvant composition comprises a nanoemulsion, an agonist of retinoic acid-inducible gene I (RIG-I), and/or an agonist of a toll-like receptor.

19. [WO/2024/160088](#)USE OF AESCIN AND/OR SALT COMPOUND THEREOF AS ADJUVANT IN [VACCINE](#)

WO - 08.08.2024

Clasificación Internacional [A61K 39/39](#)Nº de solicitud PCT/CN2024/073565Solicitante SICHUAN UNIVERSITYInventor/a SUN, Xun

The present invention provides a use of aescin and/or an aescin salt compound as an adjuvant in a [vaccine](#). The results of animal experiments show that the combination of aescin and an antigen can increase the antibody levels of sera IgG, IgG1, and IgG2a, i.e., enhance both humoral and cellular immune responses, and can be used as an adjuvant in a [vaccine](#).

20. [WO/2024/160956](#) ANTI-TUBERCULOSIS [VACCINE](#) TARGETING SELECTED MYCOBACTERIUM TUBERCULOSIS PROTECTIVE ANTIGENS TO DENDRITIC CELLS

WO - 08.08.2024

Clasificación Internacional [A61K 39/04N](#) de solicitud PCT/EP2024/052500 Solicitante INSTITUT NATIONAL DE LA SANTÉ ET DE LA RECHERCHE MÉDICALE Inventor/a LEVY, Yves

There is an urgent need for an efficient therapeutic [vaccine](#) against tuberculosis (TB), which remains a major public health issue. Current "classic" strategies under development failed or are not optimal, and to reach the World Health Organization's 2035 End TB Strategy, more efficacious vaccines are needed. The inventors have generated a post-exposure/therapeutic TB [vaccine](#) candidate (CD40.TB), consisting in an antibody directed against a surface antigen (i.e., CD40) of an antigen presenting cell (i.e., dendritic cell) wherein the heavy chain is conjugated to 3 pertinent Mycobacterium tuberculosis (Mtb) antigens, and prone to induce strong anti-TB humoral and cellular immunity.

21. [20240261390](#) [VACCINE](#) FOR PREVENTION OR TREATMENT OF VIRAL INFECTION

US - 08.08.2024

Clasificación Internacional [A61K 39/215N](#) de solicitud 18563019 Solicitante LEMONEX INC. Inventor/a CHEOL HEE WON

A nucleic acid molecule of RBD-(L)n-X sequence, wherein, RBD is a sequence of at least a partial region including the receptor-binding domain of the spike protein, L is a linker sequence, n is 0 or 1, and X is the nucleotide sequence of SEQ ID NO: 1 may be used in a [vaccine](#) composition against various viral infections.

22. [20240261393](#) SARS-COV-2 RNA [VACCINE](#) COMPOSITIONS AND METHODS OF USE

US - 08.08.2024

Clasificación Internacional [A61K 39/215N](#) de solicitud 18612698 Solicitante HDT Bio Corp. Inventor/a Steven Gregory Reed

The disclosure provides compositions, methods of treatment, and methods of making and using compositions to deliver a nucleic acid to a subject. Methods of using the compositions as a COVID-19 [vaccine](#) for the treatment of a coronavirus infection are also provided.

23. [20240252622](#) NUCLEIC ACID [VACCINE](#) AGAINST THE SARS-COV-2 CORONAVIRUS

US - 01.08.2024

Clasificación Internacional [A61K 39/215N](#) de solicitud 18615472 Solicitante INSTITUT PASTEUR Inventor/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.

24. [4408459](#) DNA-IMPfstoff zur Verwendung bei der therapeutischen und/oder prophylaktischen Behandlung von Tumorerkrankungen

EP - 07.08.2024

Clasificación Internacional [A61K 39/00N](#) de solicitud 22793599 Solicitante UNIV DEGLI STUDI DI TORINO Inventor/a NOVELLI FRANCESCO

The invention relates to a recombinant expression vector suitable for use as a prophylactic or therapeutic [vaccine](#) against tumor diseases. In addition to a promoter and any additional transcription regulatory elements, the recombinant expression vector of the invention comprises a nucleotide sequence encoding an immunogenic synthetic peptide resulting from the fusion of two or more of the amino acid sequences SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:8, or SEQ ID NO:9 of the human EN01

protein, excluding peptides that correspond to fragments of the native human EN01 protein. The recombinant vector of the invention, or the immunogenic synthetic peptides encoded thereby, are useful as a prophylactic or therapeutic [vaccine](#) against tumor diseases.

25. [WO/2024/163912](#) [VACCINE](#) AND THERAPEUTIC PROTEIN DELIVERY COMPOSITIONS COMPRISING FUSION PROTEINS

WO - 08.08.2024

Clasificación Internacional [A61K 39/145](#)Nº de solicitud PCT/US2024/014273 Solicitante THE CHILDREN'S MEDICAL CENTER CORPORATION Inventor/a MCCARTHY, Kevin R.

Provided herein are compositions comprising a polypeptide molecule, or a nucleic acid encoding such a polypeptide molecule, comprising an ectodomain polypeptide of interest and an ectodomain liberation sequence (ELS) that, when cleaved, separates the polypeptide of interest into a soluble ectodomain and a transmembrane domain. Methods of use of such polypeptides or nucleic acids encoding them are also described, including, for example generation and/or delivery of [vaccine](#) antigens and, for example, delivery of therapeutic polypeptides.

26. [4410969](#) REKOMBINANTER LEBENDER ABGESCHWÄCHTER RSV-IMPfstoffSTAMM UND HERSTELLUNGSVERFAHREN DAFÜR

EP - 07.08.2024

Clasificación Internacional [C12N 7/00](#)Nº de solicitud 22876926 Solicitante SK BIOSCIENCE CO LTD Inventor/a SEO KI-WEON

The present invention provides a recombinant attenuated respiratory syncytial virus (RSV) comprising i) a nucleic acid encoding an F protein of a stabilized pre-fusion respiratory syncytial virus (RSV) or its analogue, variant, or fragment; or ii) a nucleic acid encoding a G protein of vesicular stomatitis Indiana virus (VSV) or its analogue, variant, or fragment, and provides a genome of the recombinant RSV, and a recombinant vector comprising the genome. The recombinant attenuated RSV can be provided as a live [vaccine](#) strain which is safe and has excellent stability while maintaining infectiousness.

27. [WO/2024/162788](#) ANTIGEN FOR SEVERE FEVER WITH THROMBOCYTOPENIA SYNDROME VIRUS, AND HIGH-EFFICACY [VACCINE](#) COMPOSITION COMPRISING SAME

WO - 08.08.2024

Clasificación Internacional [C07K 14/005](#)Nº de solicitud PCT/KR2024/001524 Solicitante KOREA RESEARCH INSTITUTE OF CHEMICAL TECHNOLOGY Inventor/a KIM, Seong Jun

In the present invention, an antigen exhibiting higher antigenicity by being fused with a human LRRC24 protein-derived transmembrane domain peptide (LRRC24p) has been discovered, and it has also been identified that a neutralizing antibody is more well-formed even in actual animal experiments. It has been identified that, after inoculation with a recombinant antigen, excellent protective immunity is induced in all mice in a post-immunization challenge test in which SFTSV infection occurs. By using the recombinant antigen exhibiting these results, a [vaccine](#) for prevention of SFTSV, which exhibits lethal risk, can be developed.

28. [WO/2024/159030](#) HUMAN PAPILLOMAVIRUS, VARICELLA-ZOSTER VIRUS, AND RABIES VIRUS ANTIGENS AND USES THEREOF IN CANCER IMMUNOTHERAPY

WO - 02.08.2024

Clasificación Internacional Nº de solicitud PCT/US2024/012976 Solicitante THE UNITED STATES OF AMERICA, AS REPRESENTED BY THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES Inventor/a CUBURU, Nicolas

Provided herein are varicella-zoster virus [vaccine](#), human papillomavirus [vaccine](#), and Rabies [vaccine](#) antigens, compositions thereof, and uses thereof in cancer immunotherapy and cancer treatment.

29. [20240252670](#) ANTIBODY DRUG CONJUGATE, PREPARATION METHOD THEREFOR AND APPLICATION THEREOF

US - 01.08.2024

Clasificación Internacional [A61K 47/68N](#)° de solicitud 18560288Solicitante TSINGHUA UNIVERSITYInventor/a Xuebin LIAO

Disclosed are an antibody drug conjugate, a preparation method therefor and an application thereof, which are in particular, a conjugate of an anti-PD-L1 antibody and a TLR7 and/or TLR8 agonist, a pharmaceutical composition thereof, a preparation method therefor and an application thereof. In the present invention, a modified anti-PD-L1 antibody having mutated cysteine is obtained by means of gene editing, basically retains the structure of the original antibody, and may be used for the construction of antibody drug conjugates. By means of anti-tumor experiments, it has been discovered that the obtained antibody drug conjugate has good activity, such as strong anti-tumor activity, which may significantly improve the survival rate of tumor-bearing animals, and significantly reduce toxicity. Moreover, the antibody drug conjugate is less burdensome on the bodies of test animals, which greatly reduces the minimum effective dose of small molecular drugs when used alone, expands the therapeutic window thereof, is expected to be used in the development of therapeutic drugs for various diseases (such as tumors, viral diseases such as hepatitis B, etc.), and has good application prospects and value.

30. [20240262876](#)RECOMBINANT **VACCINE** AGAINST HELMINTHS IN PICHIA PASTORIS AND METHODS FOR PRODUCING AND PURIFYING PROTEINS FOR USE AS VACCINES AGAINST HELMINTHS

US - 08.08.2024

Clasificación Internacional [C07K 14/435N](#)° de solicitud 18443739Solicitante Fundacao Oswaldo CruzInventor/a Miriam TENDLER

The present invention is related to the recombinant production of proteins by using a synthetic gene for high protein expression in *Pichia pastoris*. More specifically, the invention describes the production of Sm14 *Schistosoma mansoni* recombinant protein, where a synthetic gene was created to promote high expression of such protein, a gene which was cloned under control of two types of *Pichia pastoris* promoters: methanol-inducible promoter (AOXI) and constituent promoter (GAP). With these constructions, *Pichia pastoris* strains were genetically manipulated to efficiently produce **vaccine** antigen Sm14. The processes to produce and purify this protein from *P. pastoris* cells, which can be escalated for their industrial production, were also improved.

31. [WO/2024/159105](#)METHODS OF TREATING CANCER COMPRISING ADMINISTRATION OF INTRATUMORAL DCS IN COMBINATION WITH SYSTEMIC IGG MONOCLONAL ANTIBODY

WO - 02.08.2024

Clasificación Internacional [A61K 35/15N](#)° de solicitud PCT/US2024/013120Solicitante H. LEE MOFFITT CANCER CENTER AND RESEARCH INSTITUTE, INC.Inventor/a CZERNIECKI, Brian

Disclosed are combination therapies comprising a semaphoring **vaccine** and dendritic cell **vaccine** and methods of their use in the treatment of cancer.

32. [20240252608](#)PEPTIDE ANALOGS CAPABLE OF ENHANCING STIMULATION OF A GLIOMA-SPECIFIC CTL RESPONSE

US - 01.08.2024

Clasificación Internacional [A61K 39/00N](#)° de solicitud 18342884Solicitante University of Pittsburgh - Of the Commonwealth System of Higher EducationInventor/a Hideho OKADA

The invention provides a peptide derived from the interleukin-13 receptor $\alpha 2$, which serves as a HLA-A2-restricted cytotoxic T lymphocyte (CTL) epitope. The invention can be used as a **vaccine** for glioma and can be formulated into compositions for medical or veterinary use. In addition, the invention provides the use of a peptide derived from the Eph family of tyrosine kinase receptors which can be also used as a **vaccine** for glioma and can be formulated into compositions for medical or veterinary use.

33. [WO/2024/157221](#)PHARMACEUTICAL COMPOSITIONS FOR DELIVERY OF HERPES SIMPLEX VIRUS GLYCOPROTEIN C, GLYCOPROTEIN D, AND GLYCOPROTEIN E ANTIGENS AND RELATED METHODS

WO - 02.08.2024

Clasificación Internacional [A61K 39/12N](#)° de solicitud PCT/IB2024/050751Solicitante BIONTECH SEInventor/a GÜLER, Alptekin

The present disclosure provides pharmaceutical compositions for delivery of HSV antigens (e.g., an HSV **vaccine**) and related technologies (e.g., components thereof and/or methods relating thereto).

34. [WO/2024/156908](#) LIPOSOMAL CONSTRUCT

WO - 02.08.2024

Clasificación Internacional [A61K 9/00N](#)° de solicitud PCT/EP2024/051996 Solicitante AC IMMUNE SA Inventor/a DI BONAVENTURA, Ivan

A liposomal construct comprises a liposome, at least one adjuvant; and a peptide containing at least one T-cell epitope which comprises, consists essentially of, or consists of the amino acid sequence of SEQ ID NO: 4 or an analogue thereof that retains alanine at position 2 and does not contain any methionine residues. The peptide preferably has the amino acid sequence of SEQ ID NO: 6. The liposomal construct may be used to generate a liposomal [vaccine](#) composition that additionally comprises at least one antigenic peptide displayed on the surface of the liposome. The liposomal [vaccine](#) compositions are useful in therapy. Methods of manufacture are also described.

35. [WO/2024/160126](#) TANDEM DESIGN METHOD AND APPARATUS FOR MULTI-EPITOPE VACCINES, DEVICE, AND STORAGE MEDIUM

WO - 08.08.2024

Clasificación Internacional [G16B 30/10N](#)° de solicitud PCT/CN2024/074020 Solicitante SHENZHEN NEOCURNA BIOTECHNOLOGY CORPORATION Inventor/a WAN, Ji

The present invention relates to the technical field of bioinformatics. Disclosed are a tandem design method and apparatus for multi-epitope vaccines, a device, and a storage medium. The method comprises: acquiring multiple sequence alignment data and a sequence feature matrix of candidate [vaccine](#) sequences; performing calculation according to the multiple sequence alignment data and the sequence feature matrix to obtain an initial PSSM for feature encoding, so as to obtain action feature information; at the same time, performing local feature extraction on the sequence feature matrix to obtain local feature information; obtaining an enhanced PSSM on the basis of the local feature information and the action feature information; then calculating predicted lysis probabilities of sites of the candidate [vaccine](#) sequences according to the enhanced PSSM and the sequence feature matrix; and finally constructing a mixed integer linear programming problem for optimization solution to obtain multiple multi-epitope vaccines. Thus, local hidden patterns and features of adjacent amino acid residues of the candidate [vaccine](#) sequences can be extracted, and the initial PSSM is corrected to obtain a more accurate enhanced PSSM to participate in subsequent epitope tandem design, thereby improving the accuracy and reliability of the designed multi-epitope vaccines.

36. [WO/2024/159101](#) METHODS FOR IDENTIFYING DISSEMINATED CANCER CELLS IN BREAST CANCER PATIENTS

WO - 02.08.2024

Clasificación Internacional [A61K 35/15N](#)° de solicitud PCT/US2024/013114 Solicitante H. LEE MOFFITT CANCER CENTER AND RESEARCH INSTITUTE, INC. Inventor/a CZERNIECKI, Brian

Disclosed are methods for the detection and isolation of disseminated cancer cells and the used of tumor antigen pulsed type 1 dendritic cell [vaccine](#) for the treatment and prevention of metastasis and abscopal tumors.

37. [4408462](#) PANCORONAVIRUS-IMPfstOFFE

EP - 07.08.2024

Clasificación Internacional [A61K 39/12N](#)° de solicitud 22877654 Solicitante GRITSTONE BIO INC Inventor/a GITLIN LEONID

Disclosed herein are [vaccine](#) compositions that include Pancorona receptor binding domain (RBD) encoding cassettes and/or MHC epitope-encoding cassettes. Also disclosed are nucleotides, cells, and methods associated with the compositions including their use as vaccines.

38. [20240252554](#) TREATMENT AND PREVENTION OF NEUROPATHOLOGY ASSOCIATED WITH NEURODEGENERATIVE DISEASES

US - 01.08.2024

Clasificación Internacional [A61K 35/74N](#) de solicitud 18633352 Solicitante ILIAD Biotechnologies, LLC Inventor/a Keith Rubin

Administering a live, attenuated *Bordetella pertussis*-based vaccine to a subject at risk for developing a neurodegenerative disease featuring A β brain plaques can prevent or reduce the amount of A β brain plaques that would have developed in the subject without such treatment.

39. [WO/2024/164014](#) RSV F VACCINE FORMULATIONS

WO - 08.08.2024

Clasificación Internacional [C07K 14/135N](#) de solicitud PCT/US2024/014509 Solicitante NOVAVAX, INC. Inventor/a PATEL, Nita

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

40. [WO/2024/163969](#) MODULAR EVALUATION OF IMMUNOGENICITY USING MULTI-PLATFORM HUMAN IN VITRO SYSTEMS

WO - 08.08.2024

Clasificación Internacional [A61K 31/739N](#) de solicitud PCT/US2024/014350 Solicitante THE CHILDREN'S MEDICAL CENTER CORPORATION Inventor/a VAN HAREN, Simon, D.

Provided herein are methods for characterizing population-specific immunogenicity of immunomodulatory agents including small molecule adjuvants and adjuvanted vaccines. The provided methods are useful for determining the activity and mechanism of action of such agents and may be used, for example, to further characterize previously known vaccine adjuvants, facilitate discovery and development of new immunomodulatory agents, and define biomarkers of their safety and efficacy.

41. [WO/2024/158722](#) SARS-COV-2 VACCINE CONSTRUCTS

WO - 02.08.2024

Clasificación Internacional [A61K 39/12N](#) de solicitud PCT/US2024/012464 Solicitante RUTGERS, THE STATE UNIVERSITY OF NEW JERSEY Inventor/a ANDERSON, Stephen

The present disclosure describes, inter alia, fusion polypeptides comprising a SARS-CoV-2 Spike polypeptide fragment comprising at least a portion of the N-terminal domain, domains CD1, RBM, and CD2, and at least a portion of CTD1, wherein the N- or C-terminus of the Spike polypeptide fragment is fused to a heterologous N- or C-terminal tag comprising at least two, at least three, or at least four amino acids, as well as polynucleotides and vectors expressing such fusion polypeptides, pharmaceutical compositions comprising the polypeptides or polynucleotides encoding them, host cells for their production, and methods of using such pharmaceutical compositions as vaccines or for generation of antibodies.

42. [WO/2024/161017](#) AVIAN-SPECIFIC VIRAL VACCINE VECTORS

WO - 08.08.2024

Clasificación Internacional [A61K 39/12N](#) de solicitud PCT/EP2024/052645 Solicitante HENNRICH, Alexandru Adrian Inventor/a HENNRICH, Alexandru Adrian

The present invention relates to a nucleic acid molecule comprising, in expressible form, a glycoprotein (G) gene deleted genome of a negative single-strand RNA virus of the order Mononegavirales, preferably a VSV rhabdovirus, wherein said genome comprises, as additional elements, an (Envelope) Env gene of an avian leukosis virus (ALV) and one or more sequences encoding one or more immunogenic proteins or peptides.

43. [20240252621](#) VIRUS-LIKE PARTICLE VACCINE FOR CORONAVIRUS

US - 01.08.2024

Clasificación Internacional [A61K 39/215N](#) de solicitud 18565728 Solicitante Icosavax, Inc. Inventor/a Niranjan Kanasa-Thanan

The present disclosure relates to targeting SARS-CoV-2, in particular, prevalent strains of SARS-CoV-2, and methods of using such vaccines to induce neutralizing antibody levels against SARS-CoV-2.

44. [WO/2024/159313](#) **VACCINE** FOR STREPTOCOCCUS EQUI SUBSP. ZOOEPIDEMICUS

WO - 08.08.2024

Clasificación Internacional [A61K 39/09](#)Nº de solicitud PCT/CA2024/050111 Solicitante UNIVERSITY OF SASKATCHEWAN Inventor/a COSTA, Matheus

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

45. [WO/2024/163508](#) METHODS AND COMPOSITIONS FOR QUADRIVALENT INFLUENZA **VACCINE**

WO - 08.08.2024

Clasificación Internacional [A61K 39/145](#)Nº de solicitud PCT/US2024/013595 Solicitante ARCTURUS THERAPEUTICS, INC. Inventor/a SULLIVAN, Brian

Provided herein are RNA molecules encoding viral replication proteins and antigenic proteins or fragments thereof. Also provided herein are compositions that include RNA molecules encoding viral replication proteins and antigenic proteins or fragments thereof, and lipids. RNA molecules and compositions including them are useful for inducing immune responses.

46. [4408886](#) BEHANDLUNG MIT NICHTIMMUNOGENER RNA ZUR ANTIGENIMPFUNG UND PD-1-ACHSEN-BINDENDEN ANTAGONISTEN

EP - 07.08.2024

Clasificación Internacional [C07K 16/24](#)Nº de solicitud 22800119 Solicitante BIONTECH SE Inventor/a SAHIN UGUR

The present disclosure relates to methods and agents for antigen vaccination and inducing effective antigen-specific immune effector cell responses such as T cell responses. These methods and agents are, in particular, useful for the treatment of diseases characterized by diseased cells expressing an antigen the immune effector cells are directed to. In some embodiments, the present disclosure relates to methods comprising administering to a subject (i) non-immunogenic RNA encoding a peptide or polypeptide comprising an epitope for inducing an immune response against an antigen in the subject, i.e., non-immunogenic RNA encoding **vaccine** antigen; and (ii) a PD-1 axis binding antagonist such as an anti-PD-1 antibody and/or an anti-PD-L1 antibody.

47. [20240252610A](#) BROADLY PROTECTIVE PROPHYLACTIC **VACCINE** AGAINST PSEUDOMONAS AERUGINOSA

US - 01.08.2024

Clasificación Internacional [A61K 39/104](#)Nº de solicitud 18562170 Solicitante UNIVERSITY OF KANSAS Inventor/a Wendy L. PICKING

Disclosed are compositions comprising a fusion polypeptide comprising i) a fusion of a needle tip protein or an antigenic fragment thereof and/or a translocator protein or an antigenic fragment thereof from a Type III secretion system (T3SS) of a Gram negative bacteria and ii) the A1 subunit of the labile toxin (LTA1) from enterotoxigenic *Escherichia coli* or cholera toxin, and methods of their use.

48. [20240262869](#) POLYPEPTIDE FRAGMENTS, IMMUNOGENIC COMPOSITION AGAINST SARS-COV-2, AND IMPLEMENTATIONS THEREOF

US - 08.08.2024

Clasificación Internacional [C07K 14/005](#)Nº de solicitud 18004065 Solicitante Indian Institute of Science Inventor/a Raghavan VARADARAJAN

The present disclosure discloses the polypeptide fragment having an amino acid sequence with at least 95% identity to the amino acid sequence selected from the group consisting of SEQ ID NO: 2, SEQ ID NO: 4, and SEQ ID NO: 6. The present disclosure also discloses

nucleic acid fragment encoding the polypeptide fragment as described herein. Moreover, the present disclosure also discloses recombinant construct, recombinant vector and recombinant host cells. Also disclosed herein is an immunogenic composition comprising the polypeptide fragment as described herein, and a method for preparing the said immunogenic composition. The immunogenic composition is in form of **vaccine**. The polypeptide fragment and/or immunogenic composition is capable of eliciting protection against severe acute respiratory syndrome coronavirus 2. A kit comprising the polypeptide, or the immunogenic composition as described herein is also disclosed.

49. [20240262891](#) HYBRIDOMA CELL LINE FOR SECRETING ANTI-RABIES VIRUS M PROTEIN MONOCLONAL ANTIBODY AND APPLICATION THEREOF

US - 08.08.2024

Clasificación Internacional [C07K 16/10N](#)° de solicitud 17294414 Solicitante ZHEJIANG UNIVERSITY Inventor/a Jiyoung ZHOU

The present invention discloses a hybridoma cell line for secreting monoclonal antibody against rabies virus M protein and application thereof, relates to the field of biotechnology. A classification of the hybridoma cell line is named as hybridoma cell line 4A1, and the hybridoma cell line was deposited on Apr. 1, 2019 in the China Center for Type Culture Collection, Wuhan University, Wuhan, China, with a deposit number CCTCC NO: C201947. The monoclonal antibody prepared by the hybridoma cell line has high titer, good specificity and excellent biological characteristics. The present invention identifies the variant antigen epitope recognized by the RABV M protein, the hybridoma cell line can be used to distinguish Flury strain and other RABV strains, prepare kit for detecting rabies virus RABV, detect RABV infection and differential diagnosis **vaccine** Flury strain and other RABV strains.

50. [20240261394](#) COMPOSITIONS AND METHODS FOR INDUCING IMMUNE RESPONSES AGAINST CLASS I FUSION PROTEIN VIRUSES

US - 08.08.2024

Clasificación Internacional [A61K 39/225N](#)° de solicitud 17924963 Solicitante University of Virginia Patent Foundation Inventor/a Steven L. Zeichner

Provided are modified bacteria and derivatives thereof that express nucleotide sequence encoding an antigen of a viral family selected from the group comprising Retroviridae (e.g., HIV, including a HIV Fusion Peptide antigen), Orthomyxoviridae, Paramyxoviridae, Arenaviridae, 5 Filoviridae, and/or Coronaviridae (e.g., an SARS-CoV, SARS-CoV-2 Fusion Peptide, and/or PEDV). In some embodiments, the bacterium has a reduced genome and induces an enhanced immune response against the viral antigen of interest when administered to a subject. In some embodiments, the viral (e.g., SARS-CoV, 10 SARS-CoV-2, PEDV, and/or HIV) antigen is expressed on a surface of a bacterium. Also provided are method for producing antibodies against viral antigens, **vaccine** compositions, methods for vaccinating subjects, methods for treating viral infections in subjects, and expression vectors for expressing viral antigens including but not limited to coronavirus (e.g., SARS-CoV, SARS-CoV-2, and/or PEDV) antigens and/or HIV antigens on the surface of reduced 15 genome bacteria.

51. [WO/2024/157026](#) RH5-INTERACTING PROTEIN (RIPR) EGF DOMAIN (RIPR EGF) ANTIGEN-BASED MALARIA **VACCINE**

WO - 02.08.2024

Clasificación Internacional [A61K 39/015N](#)° de solicitud PCT/GB2024/050210 Solicitante OXFORD UNIVERSITY INNOVATION LIMITED Inventor/a DRAPER, Simon J

The present invention relates to antigens, antibodies and vaccines for treatment or prevention of malaria.

52. [2024204962](#) METHODS FOR ENHANCING EFFICACY OF A **VACCINE** BY ADMINISTERING AN IL-4R ANTAGONIST

AU - 01.08.2024

Clasificación Internacional N° de solicitud 2024204962 Solicitante Regeneron Pharmaceuticals, Inc. Inventor/a Evans, Robert

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