



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.

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Noticias en la Web

Neumonía: Uno de los retos de vacunación a vencer en la post pandemia

1 jul. De acuerdo con datos de la Secretaría de Salud de México, durante los años 2015 y 2016 se alcanzaron coberturas adecuadas de vacunación. Sin embargo, los actuales datos recopilados por el Observatorio Mexicano de Vacunación (OMEVAC) en noviembre de 2021, revelan un panorama preocupante para los menores de edad. Sólo 20.7% de los niños de dos años, 31.7% de los de tres años y 44.5% de los infantes de cuatro años han completado el esquema básico de vacunación, evidenciando así el riesgo que enfrentan.



Específicamente hablando de la vacuna antineumocócica, se estima que la cobertura global de la tercera dosis es de 49%. Esta cifra sugiere que un gran número de niños están expuestos a un riesgo latente de contraer enfermedades neumocócicas. Estos padecimientos incluyen afecciones graves como neumonía, meningitis y bacteriemia febril, así como otras condiciones como otitis media, sinusitis y bronquitis.

En los últimos años los índices de vacunación han disminuido por diferentes factores, por ejemplo, la pandemia de COVID-19 alteró el sistema de salud provocando que alrededor de 23 millones de niños no recibieran vacunas en 2020. De manera alarmante, el número de niños que no recibieron ninguna vacuna aumentó en 3.4 millones durante ese mismo año.

“Ante este panorama, es importante resaltar que la vacunación contra el neumococo es la mejor forma de reducir el riesgo de contraer neumonía, por ello es importante inmunizar a los niños y adultos mayores, ya que, si no se detecta a tiempo, esta enfermedad puede ser fatal. La introducción de las vacunas antineumocócicas conjugadas ha representado un gran avance en la prevención de este grupo de enfermedades, estas vacunas han evolucionado a través de los años para ajustarse a los serotipos predominantes y hacer frente a la problemática de resistencia antimicrobiana, puntualiza el doctor Marcelo Díaz Conde, Líder Médico de Vacuna Antineumocócica en Pfizer México.

Reducir la incidencia de la neumonía

“En la actualidad, sabemos que aproximadamente 70% por ciento de las infecciones en niños por *S. pneumoniae* que ocurren en el mundo son causadas por entre 6 y 11 de sus serotipos más comunes. Por fortuna, hay dos vacunas conjugadas disponibles contra el neumococo que han demostrado un sustancial impacto contra las neumonías”, añade el especialista.

Neumonía en México y el mundo

En México, la neumonía, por su alta incidencia, se coloca como la séptima causa de mortalidad, de acuerdo con datos del INEGI 2023. Por su parte, la Organización Mundial de la Salud (OMS) registra anualmente alrededor de 450 millones de casos a nivel global que derivan en unos 4 millones de muertes, con mayor impacto en menores de cinco años y mayores de 75.

Especialistas de la Organización Panamericana de la Salud (OPS) han identificado al neumococo *S. pneumoniae* como el segundo agente causante de neumonía adquirida en la comunidad (NAC) que requiere hospitalización. Los menores de cinco años, principalmente en países de ingresos medios o bajos, son los más afectados por este tipo de infecciones, pues se calcula que más del 95% de todos los episodios de neumonía clínica y más de 99% de muertes por neumonía recaen en este sector de la población.

Los adultos de 65 años y más tienen 3.8 veces más posibilidades de enfermarse de neumonía por neumococo en comparación con los adultos sanos. Este peligro aumenta si además padecen alguna enfermedad crónica como diabetes mellitus (2.8 veces más), algún trastorno cardiovascular (3.8 veces más), asma (5.9 veces más) y el EPOC (7.7 veces más). Además, tienen 10 veces más probabilidad de ser hospitalizados por esta causa.

“Las primeras manifestaciones clínicas tras la infección por *S. pneumoniae* demoran entre 1 y 3 días en aparecer. Cabe destacar que los síntomas que experimente el paciente dependerán de la parte del cuerpo afectada. En el caso de la neumonía neumocócica, derivada de la infección pulmonar, incluye dolor torácico, tos, fiebre con escalofríos, respiración rápida o dificultad para respirar”, detalla el doctor Marcelo Díaz Conde.

Fuente: Vértigo Político. Disponible en <https://acortar.link/xR5zAw>

La FDA aprueba la vacuna V116 de MSD para la prevención de la enfermedad neumocócica invasiva

2 jul. La compañía Merck Sharp and Dohme (MSD) ha anunciado que la Administración de Alimentos y Medicamentos de EE.UU. (FDA) ha aprobado V116 en Estados Unidos y que se encuentra en proceso de revisión por la Agencia Europea de Medicamentos (EMA). La aprobación se produce tras la ‘revisión prioritaria’ de la FDA solicitada por MSD.

“La aprobación de hoy es un testimonio de nuestra estrategia específica para la población detrás de V116, que mostró una inmunogenicidad sólida en una variedad de poblaciones adultas y está impulsada por una comprensión profunda de la enfermedad neumocócica”, ha dicho el Dr. Dean Y. Li, presidente de MSD Research Laboratories.

V116 es la vacuna antineumocócica conjugada 21-valente indicada para la inmunización activa para la prevención de la enfermedad invasiva y la neumonía en adultos de 18 años o más. Esta terapia está específicamente diseñada para ayudar a hacer frente a los serotipos de *Streptococcus pneumoniae* predominantemente responsables de la enfermedad neumocócica invasiva (ENI) en adultos.

La enfermedad neumocócica es una infección causada por una bacteria llamada *Streptococcus pneumoniae*. Existen unos 100 tipos diferentes, denominados serotipos, de bacterias neumocócicas, que pueden afectar de forma diferente a los adultos que a los niños. La enfermedad neumocócica puede ser invasiva o no invasiva. La neumonía neumocócica es un tipo de neumonía bacteriana, que constituye la presentación clínica más frecuente de esta patología en adultos.

“Las complicaciones de la enfermedad neumocócica invasiva pueden provocar hospitalización, daños orgánicos e incluso la muerte. Muchos casos de enfermedad en adultos están causados por serotipos no incluidos en otras vacunas antineumocócicas conjugadas aprobadas”, indica el Dr. Walter Orenstein, profesor

"Estamos orgullosos de ofrecer V116 como una nueva opción específicamente diseñada para ayudar a proteger frente a la mayoría de los serotipos invasivos causantes de la enfermedad neumocócica en adultos."

emérito de medicina, epidemiología, salud global y pediatría de la Universidad de Emory y miembro del Comité Científico Asesor de MSD.



"V116 está diseñada para incluir los serotipos que causan la mayoría de los casos de enfermedad neumocócica invasiva en adultos, ayudando a proteger a los adultos contra la ENI y la neumonía neumocócica", añade Orenstein.

Esta indicación para la prevención de la neumonía causada por los serotipos indicados con anterioridad, está aprobada

en EE.UU. por la FDA bajo aprobación acelerada basada en las respuestas inmunitarias medidas por la actividad opsonofagocítica (OPA). La continuación de la aprobación para esta indicación puede depender de la verificación y descripción del beneficio clínico en un ensayo confirmatorio.

"V116 está diseñada para incluir los serotipos que causan la mayoría de los casos de enfermedad neumocócica invasiva en adultos"

Se espera que el Comité Asesor sobre Prácticas de Inmunización de los Centros para el Control y la Prevención de Enfermedades de Estados Unidos (CDC por sus siglas en inglés) se reúna este mes para discutir y hacer recomendaciones sobre el uso de V116 en adultos.

Según los datos del CDC de 2018-2021, los serotipos cubiertos por V116 son responsables de más casos de enfermedad neumocócica invasiva en adultos en comparación con PCV20 (vacuna neumocócica conjugada 20-valente). Por un lado, en adultos de 50 años o más, V116 cubre los serotipos responsables de aproximadamente el 84% de los casos de ENI, frente al 52% que cubre PCV20. Y en segundo lugar, en adultos de 65 años o más, V116 cubre los serotipos responsables de aproximadamente el 85% de los casos de ENI, en comparación con aproximadamente el 51% cubierto por PCV20.

Estos valores se basan en datos epidemiológicos del CDC y no reflejan la eficacia de las respectivas vacunas. Actualmente no hay estudios que comparen la eficacia de CAPVAXIVE y PCV20.

V116 incluye ocho serotipos únicos no cubiertos por otras vacunas antineumocócicas actualmente aprobadas; esos serotipos fueron responsables de aproximadamente el 27% de los casos de ENI en adultos de 50 años o más y de aproximadamente el 30% en adultos de 65 años o más, según los mismos datos del CDC.

Entre los datos clínicos que respaldan la aprobación se encuentran los resultados del estudio pivotal de Fase 3 STRIDE-3 (NCT05425732), que evaluó V116 en comparación con PCV20 en adultos de 18 años o más que no habían recibido previamente una vacuna neumocócica. La aprobación también se basa en los resultados de los estudios de Fase 3 STRIDE-5 (NCT05526716) y STRIDE-6 (NCT05420961), que evaluaron V116 en adultos con y sin experiencia previa en vacunación.

Fuente: ConSalud.es. Disponible en <https://acortar.link/lfpOzc>

First qualitative research study conducted in Turkmenistan focuses on HPV vaccination

Jul 4. Within the framework of a WHO–European Union joint project on immunization in central Asia, the WHO Country Office in Turkmenistan and the Ministry of Health and Medical Industry of Turkmenistan jointly conducted the country's first qualitative research study.

The project aimed at identifying factors influencing parents' decisions related to human papillomavirus (HPV) vaccination for their children. Consisting of focus-group discussions and in-depth interviews, the research provided an understanding of parents' attitudes and beliefs about HPV as well as barriers to HPV vaccination.

The results of the research conducted over 3 weeks in late 2023 will serve as the basis for activities to increase public awareness about HPV and sustain confidence in HPV vaccination in the future.

HPV vaccination in Turkmenistan

Turkmenistan included the HPV vaccine in its routine immunization schedule starting in 2016, for boys and girls at 9 years of age. Although national vaccination coverage remains high, a slight downward trend has been observed in both urban and rural areas: from 99.2% in 2021 to 98.5% in 2023.

With a relatively young population increasingly turning to the internet for information, it is important that evidence-based answers to potential questions about vaccination are readily available. However, official online information about vaccines remains limited, creating an opportunity for misinformation to spread with the potential to decrease vaccination uptake in the coming years.

The Ministry asked WHO to conduct a qualitative research study to identify what parents know about HPV, the diseases it can cause, the effectiveness of vaccination in preventing these diseases, and especially what questions or concerns they have on HPV vaccination that need to be addressed in a transparent and accessible manner.

The study, conducted jointly by experts from the Ministry and WHO, aimed to develop targeted interventions to better inform the public and health-care workers about HPV vaccines. Focus groups and in-depth interviews with health-care providers, parents and staff of public organizations were conducted to identify participants' knowledge, attitudes and behavioural determinants for uptake of HPV vaccine and childhood



vaccines in general.

The study was conducted in cities, including the capital, as well as in rural sites in 2 regions. Data collection and analysis were conducted using the COM-B framework, which looks at 3 key components: capability, opportunity and motivation for behaviour change.

Study outcomes

Study findings revealed that attitudes toward HPV were generally positive, partially due to positive attitudes toward vaccination in general but also due to preparatory steps taken by health authorities prior to introducing the HPV vaccine in 2016.

These steps included informing and training health workers to administer and answer questions about the vaccine and to inform parents and children about the benefits of HPV vaccination in preventing HPV infection, emphasizing its role in preventing the spread of the virus rather than only in preventing cervical cancer.

Despite high levels of knowledge and trust in vaccination, study participants did reveal certain gaps in knowledge and potential vulnerability to misinformation. Based on the findings, researchers proposed several measures, including:

making up-to-date information on childhood vaccination available through a single online portal to ensure accessibility and availability for the public;

training health workers to increase their capacity to effectively communicate with parents on HPV and other vaccines in the routine immunization schedule; and

using existing facility-level data and ongoing activities to conduct local, community-based interventions to effectively engage the minority of parents delaying or rejecting vaccination.

Based on these recommendations, the Ministry is developing an action plan that will include regular training for health workers and provision of information to parents via online resources and individual consultations.

With an eye to sustaining high demand for vaccination in the future, the Ministry is also planning to pilot an education module for 10–12-year-olds called Immune Patrol in several schools. WHO developed Immune Patrol to increase health literacy, resistance to misinformation, and knowledge about the immune system and immunization. WHO will provide technical support to the Ministry to implement the action plan and to pilot the Immune Patrol package in 2024 and beyond.

Fuente: World Health Organization. Disponible en <https://acortar.link/j7sghl>

Vacuna contra COVID-19 que provoca una respuesta inmunitaria superior a las demás

5 jul. Nuevos avances para tratar el coronavirus. La farmacéutica Moderna ha anunciado que su vacuna combinada contra la influenza (gripe) y la COVID-19 provoca una respuesta inmunitaria superior a las demás. El nuevo fármaco ha conseguido pasar la fase 3 del ensayo y ha cumplido con sus criterios de valoración primarios, provocando una respuesta inmunitaria más alta que las vacunas de comparación autorizadas utilizadas en el estudio.

"Las vacunas combinadas tienen el potencial de reducir la carga de los virus respiratorios en los sistemas de salud y las farmacias, así como de ofrecer a las personas opciones de vacunación más convenientes que

podrían mejorar el cumplimiento y brindar una mayor protección contra las enfermedades estacionales", dijo Stéphane Bancel, director ejecutivo de Moderna.

Al igual que las vacunas pioneras contra el coronavirus, esta se basa en el ARNm, que son aquellas en las que se emplea ácido ribonucleico para lograr el desarrollo de una respuesta inmune. El rápido éxito de Moderna demuestra que el ARN puede ayudar a superar algunas de estas dificultades, afirma James Thaventhiran, inmunólogo clínico de la Universidad de Cambridge (Reino Unido), "es un gran ejemplo de por qué la tecnología es apasionante", afirma, y añade que las vacunas combinadas que utilizan ARNm son "sólo el principio" de la tecnología del ARN.



La vacuna combinada contra la influenza y la COVID-19 de Moderna / Juan Manuel Serrano Arce - Europa Press - Archivo.

¿Cómo funcionan?

Las vacunas basadas en ARNm inyectan ARNm en las células para que hagan copias de los antígenos que el sistema inmunitario debe reconocer. Así, en lugar de tener que fabricar un montón de componentes diferentes, este tipo de vacunas simplemente envuelven un conjunto de instrucciones en una capa de lípidos y luego las envían al cuerpo para que las células produzcan sus propios antígenos. "Una sola inyección bastará para protegerse tanto de la gripe como de COVID", agrega Thaventhiran. En general, el desarrollo de vacunas combinadas de ARNm demuestra "que el ARN tiene un futuro positivo. No es solo una casualidad".

Fuente: El Periódico de España. Disponible en <https://acortar.link/sjFZZP>

Presidente de Cuba felicita a creadores de nueva vacuna infantil

6 jul. El presidente de Cuba, Miguel Díaz-Canel, felicitó al equipo científico del Instituto Finlay de Vacunas (IFV) que desarrolló el inyectable Quimi-Vio contra enfermedades respiratorias. En su perfil en la red social X el mandatario escribió: La ciencia cubana lo volvió a hacer: ¡tenemos nueva vacuna para nuestros niños! Se trata de Quimi-Vio, que protege contra neumonías, meningitis, otitis y otras enfermedades causadas por neumococo. Resultado de quince años de estudio. Gracias al equipo que hizo realidad el sueño, apuntó.

La vacuna antineumocócica cubana Quimi-Vio recibió la víspera el registro sanitario por parte el Centro para el control Medicamentos, Equipos y Dispositivos Médicos de la República de Cuba, luego de concluir todos los ensayos clínicos requeridos, que demostraron su calidad, seguridad y eficacia.



De acuerdo con la directora de Investigaciones del IFV, Dagmar García, se trata de una vacuna muy compleja, que permitirá mejorar significativamente los indicadores de salud de la infancia, en términos de morbilidad de las enfermedades respiratorias, y de mortalidad infantil por infecciones.

La científica detalló recientemente que el primer estudio con Quimi-Vio se llevó a cabo en la central provincia de Cienfuegos en 2023, donde se vacunó a más del 90 por ciento de los niños entre uno y cinco años (aproximadamente, 11 mil 600 pequeños).

También allí realizaron un ensayo clínico comunitario, en el que se inyectó al 93 por ciento de todos los niños entre uno y cinco años de edad.

Al año de haberlos inmunizado se observó un 63 por ciento de reducción de las tasas de hospitalizaciones por enfermedad respiratoria; y un 73 por ciento de reducción de la enfermedad neumocócica invasiva, por los serotipos que están contenidos en la vacuna, señaló.

En junio de este año, se anunció que un estudio de intervención contra la enfermedad neumocócica en lactantes menores de un año (que tengan dos, cuatro y 11 meses) tendrá lugar con Quimi-Vio también en territorio cienfueguero.

Según la investigadora principal del IPK, María Eugenia Toledo, Quimi-Vio protegerá a los infantes menores de un año de 11 serotipos que producen la enfermedad neumocócica.

Fuente: Prensa Latina. Disponible en <https://acortar.link/WH8T9N>

Africa surpasses global averages in vaccination

Jul 7. The breadth of vaccine protection in Gavi-supported countries, many of which are in Africa, has surpassed global averages, achieving 56 percent coverage compared to the worldwide average of 53 percent, says a recent report by Gavi, the Vaccine Alliance.

In a remarkable milestone for global health, at least 54 vaccine programmes originally introduced with Gavi funding are now self-financed by countries as of 2022, up from 40 in 2018.

Countries contributed a record \$162 million towards the co-financing of Gavi-supported vaccines in 2022, demonstrating strong ownership and financial sustainability.

Despite facing fiscal challenges, climate change, conflict and instability, Gavi-supported countries maintained or increased domestic resources for vaccine co-financing in 2022.

This commitment brought their total contribution to \$1.5 billion since the introduction of the co-financing policy in 2008. Notably, 2.6 million more children received basic routine immunisations in 2022 compared to 2021.

The Vaccine Alliance has successfully vaccinated over 1 billion children through routine immunisation programmes from 2000 to 2022.



PHOTO | SHUTTERSTOCK

In 2022, Gavi supported 40 vaccine introductions and preventive campaigns, in addition to 40 outbreak response vaccination campaigns.

This achievement highlights the substantial impact of efforts in combating some of the world's deadliest diseases. In 2022 alone, more than 68 million children were vaccinated.

Gavi plays a crucial role in immunising more than half of the world's children.

From 2000 to 2022, the Alliance facilitated over 1.8 billion vaccinations through preventive campaigns and averted more than 17.3 million future deaths.

The COVID-19 Vaccine Global Access (Covax) initiative further prevented over 2.7 million deaths across participating low- and middle-income countries.

Fuente: The East African. Disponible en <https://acortar.link/hdmS0k>

Un estudio alerta de que aumentan los casos de enfermedad neumocócica en España

9 jul. Un equipo del Laboratorio de Referencia de Neumococos del Centro Nacional de Microbiología (CNM) del Instituto de Salud Carlos III (ISCIII) en colaboración con el Centro de Investigación Biomédica en Red (CIBER) ha publicado una investigación que analiza la situación de la enfermedad neumocócica invasiva en España entre los años 2019 y 2023 en población pediátrica y adulta, caracterizando el impacto que ha tenido la pandemia por COVID-19. El estudio también ha analizado el impacto que podrían tener las nuevas vacunas conjugadas recientemente aprobadas, así como las que están en fase de desarrollo clínico para la prevención de los nuevos serotipos emergentes aparecidos tras la pandemia.



'*Streptococcus pneumoniae*' o neumococo.

El trabajo, que se ha publicado en la revista *Journal of Infection*, demuestra que el uso de medidas no farmacológicas como el uso de mascarillas, lavado de manos, el confinamiento, así como el distanciamiento social provocaron una disminución importante de casos de enfermedad neumocócica en los dos primeros años de pandemia. Sin embargo, tras el levantamiento y flexibilización de estas medidas, la enfermedad neumocócica repuntó y actualmente existen niveles incluso superiores a los que había antes de la pandemia por la COVID-19.

El equipo responsable del trabajo está liderado por el investigador del CNM-ISCIII José Yuste. También participan en la investigación Julio Sempere, Covadonga Pérez-García, Samantha Hita, Aída Úbeda, Erick Joan Vidal, Joaquín Llorente y Mirian Domenech, del Laboratorio de Referencia de Neumococos del Instituto. Además, colaboran investigadores e investigadoras de las Áreas de Enfermedades Respiratorias (CIBERES) y de Epidemiología y Salud Pública (CIBERESP) del CIBER-ISCIII, de la Dirección General de Salud Pública del Ministerio de Sanidad, y de diversas universidades y hospitales españoles.

Entre los serotipos que más han aumentado, según el trabajo, se encuentra el serotipo 3, que es uno de los relacionados con una mayor mortalidad, y que afecta principalmente a población menor de 5 años y a

personas adultas por encima de 65 años. También destaca la presencia del serotipo 4, ligado a infecciones en adultos jóvenes. Por otro lado, el serotipo 24F en población pediátrica, y el serotipo 8 en población adulta, siguen teniendo también importancia en la incidencia de la enfermedad.

Streptococcus pneumoniae o neumococo es la principal causa de las neumonías comunitarias de etiología bacteriana, así como de sepsis y meningitis. Este estudio liderado desde el ISCIII confirma el aumento de casos por cepas invasivas, algunas de ellas asociadas altos niveles de resistencia a los antibióticos, lo que supone una importante preocupación para la salud pública. En este sentido, este mismo equipo investigador ya señaló en 2022, en un estudio publicado en *Lancet Microbe*, del posible aumento de casos por cepas resistentes al principio de la pandemia. Además, otro estudio previo, publicado en *Clinical Infectious Diseases* con datos de los años 2009-2019, había permitido generar un 'mapa' de la enfermedad neumocócica invasiva en España en la última década, señalando un aumento de casos.

El equipo liderado por el doctor Yuste concluye que el uso de las nuevas vacunas conjugadas, con más espectro de protección y/o inmunogenicidad que las que había anteriormente y que se han autorizado recientemente en España, podrían ayudar a prevenir muchos de estos casos, incluyendo nuevos serotipos emergentes. Según apunta, es importante recordar que, en población adulta, la vacunación frente a neumococo no es estacional y que, por tanto, se puede administrar en cualquier momento del año; a diferencia de las vacunas contra la gripe, que al modificarse todos los años hay que administrarse cada año, la vacuna frente al neumococo se administra generalmente una sola vez en la vida.

Fuente: Agencia Iberoamericana para la Difusión de la Ciencia y la Tecnología.

Disponible en <https://acortar.link/QUyAmi>

Pfizer recibe el visto bueno de la Comisión Europea para una nueva vacuna contra la COVID-19

10 jul. La farmacéutica estadounidense y la biotecnológica BioNTech han desarrollado una vacuna actualizada contra el coronavirus adaptada al linaje Ómicron JN.1. El compuesto se aprueba para personas a partir de seis meses de edad.

Pfizer y BioNTech persisten en el mercado de las vacunas contra la COVID-19. Ambas compañías han anunciado que la Comisión Europea (CE) ha concedido la aprobación de la vacuna monovalente adaptada a ómicron JN.1 para la inmunización activa frente al coronavirus en personas a partir de los seis meses de edad.



Esta nueva adaptación se basa en las recomendaciones del Grupo Consultivo Técnico sobre la Composición de las Vacunas frente a la COVID-19 de la Organización Mundial de la Salud (OMS) y del Grupo de Trabajo de Emergencia (ETF) de la Agencia Europea del Medicamento (EMA, por sus siglas en inglés) para actualizar las vacunas frente a la variante JN.1 en la campaña 2024 y 2025.

Esta autorización sigue a la reciente opinión positiva por parte del Comité de Medicamentos de Uso Humano (Chmp, por sus siglas en inglés) de la EMA. La autorización es válida en los 27 Estados miembros de la Unión Europea (UE), además de Islandia, Liechtenstein y Noruega.

Pfizer y BioNTech también han presentado una solicitud ante la Administración de Alimentos y Medicamentos de Estados Unidos

Tras la decisión de la CE, la vacuna actualizada ya está disponible para su envío inmediato a los estados miembros de la UE para su uso conforme a las recomendaciones oficiales de cada uno de ellos. Pfizer y BioNTech han desarrollado la vacuna monovalente frente a la COVID-19 adaptada a Ómicron JN.1 para garantizar la disponibilidad del suministro de cara a la próxima temporada de otoño e invierno, momento del año en el que se espera que aumente la demanda de vacunación frente al virus.

La aprobación de la CE se basa en todo el conjunto de resultados clínicos, preclínicos y en vida real anteriores que respaldan la seguridad y la eficacia de las vacunas frente a la COVID-19 de Pfizer y BioNTech.

Pfizer y BioNTech también han presentado una solicitud ante la Administración de Alimentos y Medicamentos de Estados Unidos (FDA, por sus siglas en inglés) solicitando la aprobación de la vacuna monovalente frente a la COVID-19 adaptada a ómicron KP.2 para personas a partir de seis meses de edad.

Fuente: Planta Doce. Disponible en <https://acortar.link/V54klb>

OMS alerta sobre menor vacunación contra COVID-19; aún registran 1,500 muertes semanales

11 jul. El director general de la OMS aseguró que se registra un grave descenso en la vacunación de grupos vulnerables contra la COVID-19.

Datos de la Organización Mundial de la Salud (OMS) indican un descenso en la vacunación contra COVID-19 incluso en grupos de riesgo, pese a que esta enfermedad sigue causando unas 1,500 muertes semanales, advirtió este jueves el director general de la agencia, Tedros Adhanom Ghebreyesus.

“Nuestros datos muestran que la cobertura de vacunación ha descendido entre trabajadores sanitarios y personas mayores de 60, que son dos de los principales grupos de riesgo”, alertó el experto etíope en su rueda de prensa semanal para repasar los principales temas sanitarios a nivel global.

“Recomendamos que las personas de estos grupos de riesgo sigan recibiendo vacunas contra la covid-19 doce meses después de su última dosis”, agregó Tedros.

Fuente: Forbes México. Disponible en <https://acortar.link/pGzYMN>



WHO publishes guidance for HPV vaccine developers to help tackle cervical cancer

Jul 11. In more than nine out of ten cases, the HPV virus leads to cervical cancer, the fourth most common cancer in women globally.

The World Health Organization (WHO) has published a new report to guide vaccine developers when creating innovative vaccines to treat human papillomavirus (HPV) infections in adults.

Currently the main cause of cervical cancer – the fourth most common cancer in women globally – HPV is a very common group of viruses that leads to cervical cancer in more than nine out of every ten cases.

Despite already-existing WHO-prequalified vaccines, including Merck & Co's – known as MSD outside the US and Canada – Gardasil 9 (HPV 9-valent, recombinant) and GSK's Cervarix (HPV bivalent (types 16, 18) vaccine, recombinant), to prevent HPV infections, there are currently no vaccines that can treat them.

However, new therapeutic vaccines are being developed using the latest immunotherapy advances to clear HPV or treat precancerous cells in people already exposed to the virus.

The WHO Preferred Product Characteristics for Therapeutic HPV Vaccines report calls on developers and outlines how they can ensure that their products meet global public health needs and how to quickly roll out vaccines if approved, while considering medical indications for potential use of such a vaccine, target populations, safety and efficacy criteria, as well as practical considerations such as delivery strategy, storage and dosage schedules, and affordability.

Currently, there are more than 20 therapeutic HPV vaccine candidates that are at different stages of development, with several in clinical trials, that are likely to be most beneficial for adult women who did not receive the HPV vaccination prior to contracting the virus.

In addition, these new potential therapeutic vaccines align with WHO's Cervical Cancer Elimination Initiative, which aims to vaccinate 90% of girls with preventive HPV vaccines, screen 70% of women with a high-performance test and treat 90% of women who have precancerous cells in the cervix or cervical cancer by 2030.

Dr Sami Gottlieb, medical doctor and epidemiologist, department of sexual and reproductive health and research, WHO, commented: "Therapeutic HPV vaccines could be a catalytic innovation to complement these efforts, increasing options for the millions of women who have already acquired HPV and reducing their risks of developing life-threatening cancer in the future."

Fuente: PM Live. Disponible en <https://acortar.link/ImweYC>



Belarusian Ministry of Health grants registration to Cuban vaccine Cimavax

Jul 12. The Ministry of Health of the Republic of Belarus reports that the Center for Expertise and Testing in Health Care has granted registration to the Cuban vaccine Cimavax, it was learned here today.

Developed by the Cuban Center for Molecular Immunology, it is the world's first officially patented and registered therapeutic vaccine against lung cancer.

The Cuban Ambassador to the Slavic nation, Santiago Pérez, commented that the decision of the regulatory agency of this European country to grant registration to Cimavax is an undeniable achievement of Cuban biotechnology in a country with such high standards and requirements as Belarus.

Pérez recalled that Belarus was the first country on the continent to register the Cuban vaccine Soberana, developed against COVID-19.

The Cuban diplomat said the registration is a sign of progress in bilateral relations, especially in the economic commercial field, which would allow an increase in Cuba's exports.

Likewise, it shows the progress made in cooperation between the scientific communities of the two countries, and perhaps it is the beginning of a more dynamic process of mutual registration of medicines both in Belarus and Cuba, said Perez. (Source: PL)

Fuente: RADIO HABANA CUBA. Disponible en <https://acortar.link/zrTt2o>



China Proceeds with mRNA RSV Vaccine Candidate

Jul 13. The board of directors of CSPC Pharmaceutical Group Limited announced on July 11, 2024, that the mRNA Respiratory Syncytial Virus (RSV) vaccine candidate (SYS6016) has obtained approval from the National Medical Products Administration of the People's Republic of China to conduct human clinical trials in China.

Currently, there is no vaccine available in China that protects people from RSV infection.

In preclinical studies, SYS6016 translated into the prefusion conformation F-protein in vivo and induced high titers of long-lasting neutralizing antibodies.

CSPC wrote that this vaccine candidate exhibits good protection against RSV-A and RSV-B subtype viral strains and has a good safety profile.

CSPC confirmed it would endeavor to advance the clinical research and market SYS6016 as soon as possible to create value for society and shareholders.

As of July 13, 2024, three RSV vaccines and one monoclonal antibody for infants (Beyfortus) were approved for use in the United States.

Fuente: Precision Vaccinations. Disponible en <https://acortar.link/Wi5O3V>

Oxford scientists testing two new respiratory vaccines

Jul 14. Scientists at the University of Oxford have launched a new study to test a two-in-one respiratory vaccine.

The phase one study will test a vaccine targeting both respiratory syncytial virus (RSV) and human metapneumovirus (hMPV), and another for RSV alone in five to eight-month-old infants.

RSV is a common virus affecting children across the globe. Within their first two years, up to 90 per cent of children contract the virus and it frequently reinfects older children and adults.

Although RSV usually causes mild respiratory illness, some infants experience severe bronchiolitis, which leads to the inflammation of the small airways, sometimes causing extreme breathing difficulties.

In the UK, about one in 50 children under the age of one are admitted to hospital as a result of RSV infections.

Closely related to RSV, hMPV is a respiratory pathogen, associated with a range of illnesses from mild infection to severe bronchiolitis and pneumonia.

These potential jabs are being developed by pharmaceutical company Moderna.

Dr Simon Drysdale, consultant in paediatric infectious diseases and immunology at the Oxford Vaccine Group, and principal investigator on the study, said: "RSV and hMPV are two of the leading causes of respiratory infections in children, the elderly and immunocompromised patients worldwide.

"This trial is an important early step in the development of a paediatric vaccine against RSV and hMPV."

The jabs are messenger RNA vaccines. They work by instructing cells in the body to produce a specific protein, triggering an immune response.

This prompts the immune system to remember the protein, enabling a rapid and effective reaction if the individual contracts the actual virus in the future.

The Rhyme study is the first attempt to create a vaccine that could simultaneously protect against RSV and hMPV.

It aims to enrol approximately 60 paediatric participants between five and eight months old across seven UK sites.



Over the course of around 24 months, each participant will receive three injections at 56-day intervals.

The sites participating in the trial are Norfolk and Norwich University Hospitals NHS Foundation Trust, Alder Hey Children's NHS Foundation Trust, Imperial College Healthcare NHS Trust and St George's University Hospitals NHS Foundation Trust.

Also taking part are Oxford University Hospitals NHS Foundation Trust, University Hospital Southampton NHS

Foundation Trust and University Hospitals Bristol and Weston NHS Foundation Trust, with the research being sponsored by Moderna.

The study is supported by the National Institute for Health and Care Research Clinical Research Network Thames Valley and South Midlands.

It has been reviewed and approved by the Health and Social Care Research Ethics Committee.

Fuente: Oxford Mail. Disponible en <https://acortar.link/Dlvef6>

La vacunación infantil en el mundo se estanca, alerta la ONU

15 jul. Las tasas de vacunación infantil en el mundo se están estancando y aún no han recuperado los niveles previos a la pandemia de COVID-19, alertó la ONU el lunes.

Comparado con el nivel de 2019, 2,7 millones de niños más no fueron vacunados o lo fueron de manera incompleta en 2023, según un comunicado conjunto de UNICEF y la OMS.

"Las últimas tendencias demuestran que muchos países continúan desatendiendo demasiados niños", lamentó la directora ejecutiva de UNICEF, Catherine Russell.



En 2023 solo 84 % de los niños, es decir 108 millones, recibieron tres dosis de la vacuna contra la difteria, el tétanos y la tos ferina (DTAP), siendo la tercera dosis un indicador clave de la cobertura global de vacunación, según datos publicados por las agencias de salud y de infancia de la ONU.

Este porcentaje no cambió desde el año pasado, lo que significa que los modestos avances observados en 2022, después de la fuerte caída debido a la COVID-19, se detuvieron, señalan las organizaciones.

Esta tasa era del 86 % en 2019, antes de la pandemia.

"Estamos retrasados", admitió ante la prensa Kate O'Brien, responsable de vacunación en la OMS.

En 2023, 14,5 millones de niños en el mundo eran considerados "cero dosis", es decir, no habían recibido ninguna dosis de vacuna, una cifra en aumento desde los 13,9 millones en 2022 y los 12,8 millones en 2019, según los datos publicados el lunes.

La mitad de los niños no vacunados en el mundo viven en 31 países afectados por conflictos.

6,5 millones de niños en el mundo no recibieron su tercera dosis de la vacuna DTAP, necesaria para asegurar la protección contra enfermedades en lactantes y niños pequeños.

Estas disparidades en la cobertura de vacunación favorecen el desarrollo de ciertas enfermedades como el sarampión.

"Los brotes de sarampión son una señal de advertencia, que revelan lagunas existentes en la vacunación y golpean a los más vulnerables", declaró el director general de la OMS, Tedros Adhanom Ghebreyesus, en el comunicado.

En 2023, 83 % de los niños en el mundo recibieron su primera dosis de la vacuna contra el sarampión a

través de los servicios de salud básicos, el mismo nivel que en 2022, pero aún por debajo del 86 % antes de la pandemia.

Más de 300.000 casos de sarampión fueron registrados en 2023, casi tres veces más que el año anterior, destacó Ephrem Lemango, responsable de vacunación en UNICEF.

Al menos 103 países experimentaron brotes en los últimos cinco años.

En cambio, 91 países con una sólida cobertura de vacunación contra el sarampión no registraron brote alguno.

Fuente: FRANCE 24. Disponible en <https://acortar.link/7wmibN>

Ministry of Health and Wellness receives 50,000 doses of Pediatric Pneumococcal Vaccine (PCV) Vaccine from Brazil

Jul 16. On July 15, 2024, the Belizean Ministry of Health & Wellness received 50,000 doses of the Pediatric Pneumococcal Vaccine (PCV) donated by the Government of Brazil. The PCV vaccine will prevent pneumonia, meningitis and other complications caused by pneumococcal disease in children under five years of age.

The Ministry noted that Vaccinated children will benefit from less hospital visits and less hospital stay (Hospitalization); their growth and development will not be compromised due to pneumococcal infection.

The Ministry of Health & Wellness extended gratitude and appreciation to the Government and Ambassador of Brazil for the donation of the Pneumococcal Pediatric Vaccine, and encourages parents to get their children five years and under vaccinated. The vaccine is available at all public health facilities and mobile clinics.

Fuente: Breaking Belize News BBN. Disponible en <https://acortar.link/K0ey9L>



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Patentes registradas en Patentscope

Estrategia de búsqueda: (Vaccine) AND DP:([01.07.2024 TO 16.07.2024]) as the publication date 58 records.

1. [WO/2024/144272](#) **VACCINE** COMPOSITION COMPRISING NOROVIRUS GII MRNA

WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/KR2023/021767 Applicant K-BIOCELF INC. Inventor OH, Myung Ryurl
The present invention relates to a **vaccine** composition comprising mRNA encoding the structural protein VP1 of a norovirus GI genotype. The mRNA **vaccine** composition was prepared by deriving the consensus sequence of a norovirus GI genotype VP1 antigen, and the immunogenicity of the **vaccine** composition was confirmed. Therefore, the **vaccine** composition can be effectively used to prevent and treat norovirus infection.

2. [WO/2024/144267](#) **VACCINE** COMPOSITION COMPRISING HIV-1 MRNA

WO - 04.07.2024

Int.Class [A61K 39/21](#) Appl.No PCT/KR2023/021756 Applicant K-BIOCELF INC. Inventor OH, Myung Ryurl
The present invention relates to a **vaccine** composition comprising an HIV-1 MRNA. The mRNA **vaccine** composition is prepared by deriving the common sequence of an HIV-1 antigen, and the immunogenicity of the **vaccine** composition is identified, and thus the present invention can be effectively used in the prevention and treatment of HIV infection.

3. [WO/2024/144262](#) **VACCINE** COMPOSITION COMPRISING NOROVIRUS GI MRNA

WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/KR2023/021746 Applicant K-BIOCELF INC. Inventor OH, Myung Ryurl
The present invention relates to a **vaccine** composition comprising mRNA encoding the structural protein VP1 of the norovirus genotype GI. An mRNA **vaccine** composition was prepared by deriving a common sequence of a norovirus genotype GI VP1 antigen, and through identification of the immunogenicity of the **vaccine** composition, the composition can be usefully employed to prevent and treat norovirus infection.

4. [20240226256](#) COMBINATION THERAPY TUMOUR CELL **VACCINE**

US - 11.07.2024

Int.Class [A61K 39/00](#) Appl.No 18561772 Applicant Queen's University at Kingston Inventor Kyle Seaver
A cancer **vaccine** includes at least one tumour associated antigen (TAA), at least one Toll-like receptor (TLR) agonist, at least one cytokine, and a pharmaceutically acceptable vehicle. The at least one TAA may be provided by dead tumour cells, such as γ -irradiated tumour cells or lysis and UV treated tumour cells, the at least one TLR agonist may comprise 5 CpG-1826 and the at least one cytokine may comprise IL-27. When administered to a mammalian subject the cancer **vaccine** prevents, inhibits, or slows tumour development in the subject, and the **vaccine** may provide a long-term T cell activation and memory against tumour development in the subject.

5. [20240226282](#) NANT COVID **VACCINE** CROSS REACTIVITY

US - 11.07.2024

Int.Class [A61K 39/215](#) Appl.No 18620192 Applicant ImmunityBio, Inc. Inventor Patrick Soon-Shiong

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Recombinant SARS-CoV2 **vaccine** compositions and methods are presented that have unexpected cross-reactivity against a variety of other coronaviruses, and particularly against SARS-CoV1, MERS-CoV, OC43-CoV, and HKU1-CoV in addition to significant reactivity against SARS-CoV2A. Moreover, the **vaccine** compositions presented herein also produced cross-reactive memory B cells as well as cross-reactive memory T cells with cross-reactivity spanning a relatively wide range of different coronaviruses.

6. [WO/2024/143748](#) COMPOSITION FOR INCREASING STABILITY OF ANIMAL **VACCINE** OR DIAGNOSTIC ANTIGEN, CONTAINING AMINO ACIDS AS ACTIVE INGREDIENT, AND USE THEREOF
WO - 04.07.2024

Int.Class [A61K 39/00](#) Appl.No PCT/KR2023/012012 Applicant CHOONGANG **VACCINE** LABORATORIES CO., LTD (CAVAC) Inventor YOON, In-joong

The present invention relates to a composition for increasing the stability of an animal **vaccine** or a diagnostic antigen, comprising amino acids as an active ingredient, and use thereof. The stability-increasing composition according to the present invention can effectively maintain a prophylactically-effective virus titer without loss of infectivity titer in virus vaccines, and can facilitate transport and storage of vaccines, and thus can be effectively used in related fields.

7. [WO/2024/146068](#) USE OF TRIPTERINE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF IN PREPARATION OF **VACCINE** ADJUVANT
WO - 11.07.2024

Int.Class [A61K 39/39](#) Appl.No PCT/CN2023/098745 Applicant NATIONAL **VACCINE** AND SERUM INSTITUTE (NCSI) Inventor WANG, Zhibiao

The present invention provides a use of tripterine or a pharmaceutically acceptable derivative thereof in the preparation of a **vaccine** adjuvant, and a **vaccine** adjuvant containing the tripterine or the pharmaceutically acceptable derivative thereof. The present invention finds that the immune response of an organism to an antigen can be enhanced by using the tripterine or the pharmaceutically acceptable derivative thereof as the **vaccine** adjuvant. Compared with metal salt adjuvants such as an aluminum adjuvant that is not easy to metabolize in vivo and has a risk of accumulation, the tripterine or the pharmaceutically acceptable derivative thereof in the present invention is a small molecule compound, is easy to degrade and excrete, is high in safety, and has a better immunogenicity enhancement effect than that of the aluminum adjuvant.

8. [20240216498](#) LARGE-SCALE FLAVIVIRAL **VACCINE** PRODUCTION AND MANUFACTURE
US - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No 18523331 Applicant Takeda Vaccines, Inc. Inventor Joseph David Santangelo

The present invention provides methods for large-scale flaviviral **vaccine** production and manufacture. The methods provided herein are specifically contemplated for large-scale production and manufacture of live, attenuated flaviviral vaccines such as live, attenuated, dengue virus vaccines. Further, the methods provided herein pertain to formulation of live, attenuated, monovalent, divalent, trivalent, or tetravalent viral **vaccine** products.

9. [WO/2024/139647](#) HUMAN PAPILLOMAVIRUS TYPE 16 DNA **VACCINE** AND USE THEREOF
WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/CN2023/128073 Applicant INSTITUTE OF BASIC MEDICAL SCIENCES, CHINESE ACADEMY OF MEDICAL SCIENCES Inventor XU, Xuemei

Disclosed are a human papillomavirus type 16 DNA **vaccine** and use thereof, which relate to the technical field of biology. An antigen gene of the present invention is an optimized human papillomavirus type 16 E6 protein and E7 protein fusion gene, and a human heat shock protein 70 gene is further fused downstream thereof. A **vaccine** plasmid further comprises a replication origin and a CpG motif-based immunostimulatory sequence. The human papillomavirus type 16 DNA **vaccine** disclosed can induce an HPV16 E6 and E7 antigen specific

cellular immune response by means of intramuscular injection electroporation immunization. The DNA vaccine can be used for preventing and treating HPV16 infection, and treating infection-related precancerous lesions and malignant tumors.

10. [2024204248](#) COMBINATION THERAPY WITH NEOANTIGEN VACCINE

AU - 04.07.2024

Int.Class [C07K 16/28](#) Appl.No 2024204248 Applicant Dana-Farber Cancer Institute, Inc. Inventor Fritsch, Edward F.

The present invention relates to neoplasia vaccine or immunogenic composition administered in combination with other agents, such as checkpoint blockade inhibitors for the treatment or prevention of neoplasia in a subject.

11. [20240216499](#) PROTEIN-BASED NANOPARTICLE VACCINE FOR METAPNEUMOVIRUS

US - 04.07.2024

Int.Class [A61K 39/145](#) Appl.No 18461862 Applicant Icosavax, Inc. Inventor Andrew Lawrence Feldhaus

Provided are virus-like particle vaccines for human metapneumovirus (hMPV) in which the ectodomain of hMPV F protein is linked to, and thereby displayed on, a symmetric protein-based virus-like particle. For example, the vaccine antigen may be a N-terminal fusion of the ectodomain of hMPV F protein to a protein having a multimerization domain for a one- or two-component virus-like particle, such as a two-component icosahedral virus-like particle. Further provided are vaccine compositions, methods of manufacturing, and methods of use, e.g., immunizing a subject to generate a protective immune response to hMPV.

12. [4396198](#) NEW CORONAVIRUS VACCINE AND METHOD FOR DESIGNING AND OBTAINING A VIRUS VACCINE

EP - 10.07.2024

Int.Class [C07K 14/005](#) Appl.No 22772886 Applicant MAX DELBRUECK CENTRUM FUER MOLEKULARE MEDIZIN HELMHOLTZ GEMEINSCHAFT Inventor DE LA ROSA KATHRIN

The present invention relates to a mutant receptor-binding domain (mRBD) of a coronavirus (mRBD-CORONA) or a fragment thereof and mutant spike protein of the coronavirus (CORONA-mSpike) or a fragment thereof comprising the CORONA-mRBD or the fragment thereof. Furthermore, the present invention relates to a polypeptide or protein comprising the mRBD-CORONA or the fragment thereof or CORONA-mSpike or the fragment thereof and a nucleic acid comprising a nucleotide sequence encoding for the mRBD-CORONA or the fragment thereof or the CORONA-mSpike or the fragment thereof. Furthermore, the present invention relates to a vaccine composition comprising one or more CORONA-mRBDs or fragments thereof, one or more CORONA-mSpikes, one or more polypeptides or proteins and/or one or more nucleic acids according to the present invention. Furthermore, the present invention relates to the one or more CORONA-mRBDs or fragments thereof, the one or more CORONA-mSpikes, the one or more polypeptides or proteins, the one or more nucleic acids and/or the vaccine composition according to the present invention for use in the prevention and/or treatment of diseases caused by coronaviruses in a subject. Furthermore, the present invention relates to a method for designing and/or obtaining an active ingredient for a vaccine composition and to a VIRUS-mRBD or a fragment thereof designed and/or obtained by the method for obtaining the VIRUS-mRBD according to the present invention.

13. [WO/2024/144254](#) VACCINE COMPOSITION COMPRISING ZIKA VIRUS INACTIVATED BY EPIGALLOCATECHIN-3-GALLATE AS ACTIVE INGREDIENT

WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/KR2023/021729 Applicant GREENVAX INC. Inventor SEONG, Baik Lin

The present invention relates to a vaccine composition comprising Zika virus inactivated by Epigallocatechin-3-gallate (EGCG) as an active ingredient, wherein it is confirmed that the Zika virus vaccine inactivated by

EGCG not only has a superior **vaccine** effectiveness due to a neutralizing antibody compared to existing Zika virus vaccines inactivated by formaldehyde which is a virus inactivating material, but also significantly improves a cytotoxic immune effect caused by an antibody, and thus, EGCG alone can be effectively utilized as a Zika virus **vaccine** composition without a separate inactivating material or combination with an adjuvant.

14. [4396333](#) UTILIZATION OF MICRO-RNA FOR DOWNREGULATION OF CYTOTOXIC TRANSGENE EXPRESSION BY MODIFIED VACCINIA VIRUS ANKARA (MVA)

EP - 10.07.2024

Int.Class [C12N 7/00](#) Appl.No 22772951 Applicant BAVARIAN NORDIC AS Inventor HAUSMANN JÜRGEN

The invention relates to a recombinant Modified Vaccinia Virus Ankara (MVA) comprising a series of miRNA target sequences arranged in a miRblock that is linked to a transgene, wherein each miRNA target sequence corresponds to a miRNA in a eukaryotic MVA producer cell. The present invention also relates to medical uses of the recombinant MVA.

15. [WO/2024/144348](#) **VACCINE** COMPOSITION COMPRISING IONIC COMPLEX OF CATIONIC MOLECULAR TRANSPORTER AND MRNA

WO - 04.07.2024

Int.Class [A61K 39/00](#) Appl.No PCT/KR2023/021981 Applicant BIOPHARMA CORP. Inventor RHEE, Jinseol

The present invention relates to an mRNA-based **vaccine** composition comprising mRNA encoding an antigen peptide or protein, a liposome, and a cationic molecular transporter (SG6). The **vaccine** composition of the present invention has no cytotoxicity, high mRNA delivery efficiency, excellent stability and safety, and excellent antigen peptide or protein expression efficiency, and thus can be effectively used as an mRNA-based **vaccine** composition. Meanwhile, the present invention was made with the support of the Korea Health Industry Development Institute's health and medical technology research and development project by funding from the Ministry of Health and Welfare (Project identification number: HQ21C0274).

16. [WO/2024/138836](#) HEAT-RESISTANT PROTECTIVE AGENT FOR LIVE **VACCINE**, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 04.07.2024

Int.Class [A61K 47/26](#) Appl.No PCT/CN2023/075359 Applicant JINYUBAOLING BIO-PHARMACEUTICAL CO., LTD Inventor LI, Chao

A heat-resistant protective agent for a live **vaccine**, a preparation method therefor, and use thereof. The protective agent for a live **vaccine** comprises: an inulin, gelatin, glycine, polyvinylpyrrolidone, bovine serum albumin, proteolyzed casein, D-sorbitol, a water-soluble phospholipid, pollen pini, and a tocopherol, wherein the mass ratio of pollen pini to the tocopherol is (1-2):1. The components of the protective agent are simple and effective, the raw materials are safe and readily available, and the preparation method is convenient and fast. When the protective agent is used to prepare a live Mycoplasma bovis **vaccine**, the protective agent can effectively reduce the loss rate of live mycoplasma bacteria during lyophilization and extend the shelf life, and meanwhile the **vaccine** can induce a long-lasting, efficient immune response in an organism.

17. [WO/2024/144193](#) **VACCINE** COMPOSITION COMPRISING GOLD-NANOPARTICLE-CARRIER HAVING DOUBLE-STRANDED DNA BOUND THERETO

WO - 04.07.2024

Int.Class [A61K 39/385](#) Appl.No PCT/KR2023/021565 Applicant NES BIOTECHNOLOGY CO., LTD. Inventor LEE, Kangseok

The present invention relates to a **vaccine** composition containing double-stranded DNA transported into cells by means of gold nanoparticles and, more particularly, to a **vaccine** composition characterized in that the double-stranded DNA is derived from viruses, bacteria, or cancer genes and expresses an antigen to induce an immune response.

18. [4395818](#) STABILIZATION OF ADJUVANTED [VACCINE](#) COMPOSITIONS AND THEIR USE

EP - 10.07.2024

Int.Class [A61K 39/00](#) Appl.No 22865813 Applicant VAXCYTE INC Inventor GRAINGER CHRISTOPHER LAIN
The present disclosure provides stabilized [vaccine](#) compositions that resist the formation of unsuitable adjuvant flocculant or aggregates. These compositions comprise an aluminium adjuvant, a non-aluminium phosphate salt, sodium chloride and a polypeptide-polysaccharide conjugate comprising at least two non-natural amino acids. The present disclosure further provides methods of using such compositions to induce immune responses against infections in subjects.

19. [WO/2024/147556](#) COMPOSITION FOR ANTICANCER [VACCINE](#) COMPRISING K-RAS MUTANT MULTIPLE EPITOPE POLYPEPTIDE AS ACTIVE INGREDIENT

WO - 11.07.2024

Int.Class [A61K 39/00](#) Appl.No PCT/KR2023/021882 Applicant MYONGJI MEDICAL FOUNDATION Inventor LEE, Wang Jun

The present invention relates to a composition for an anticancer [vaccine](#), the composition comprising, as an active ingredient, a K-ras mutant multiple epitope polypeptide consisting of the amino acid sequence of SEQ ID NO: 1. Using the composition for an anticancer [vaccine](#) according to the present invention enhances the immune surveillance function against cancer cells expressing mutations in the K-ras protein such that cancer cells are detected and killed early through immune response mechanisms and cancer cells that have already formed are selectively killed, thus making it possible to prevent and treat cancer.

20. [WO/2024/145150](#) PSEUDORABIES VIRUS LIVE ATTENUATED [VACCINE](#) FOR PIGS, COMPRISING A DELETION OF GENE UL23

WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/US2023/085345 Applicant ZOETIS SERVICES LLC Inventor KONG, YiBo
This disclosure provides an attenuated suid herpesvirus 1 (a Pseudorabies virus) wherein the TK, gl and gE genes thereof are modified relative to a parent field strain, such that the resultant virus is safe and effective for use as a live [vaccine](#) that protects swine animals from challenge with a virulent Pseudorabies virus.

21. [20240226270](#) NOVEL [VACCINE](#) COMPOSITIONS FOR PORCINE EPIDEMIC DIARRHEA VIRUS AND PORCINE DELTACORONAVIRUS

US - 11.07.2024

Int.Class [A61K 39/215](#) Appl.No 17177724 Applicant Zoetis Services LLC Inventor Jacqueline Gayle Marx
The present invention is directed to novel immunogenic compositions that protect swine from disease caused by porcine epidemic diarrhea virus (PEDV). The present invention is also directed to novel immunogenic compositions that protect swine from disease caused by porcine deltacoronavirus (PDCoV), alone or as combination [vaccine](#) to protect against PEDV. The compositions of the invention provide killed viruses whose effectiveness is enhanced by the selection of preferred adjuvants. Novel culture methods are also employed to increase reproducible yield of cultured viruses. Live vaccines are also provided from the Calaf14 PEDV isolate.

22. [4393505](#) [VACCINE](#) FOR PREVENTING AFRICAN SWINE FEVER, COMPRISING AFRICAN SWINE FEVER VIRUS-DERIVED ANTIGEN PROTEIN

EP - 03.07.2024

Int.Class [A61K 39/187](#) Appl.No 22861612 Applicant BIOAPPLICATIONS INC Inventor KANG HYANGJU
The present invention relates to: a recombinant vector comprising a nucleotide sequence of antigen protein(s) Lectin, CD2v, p72, p54, p30, p15, p35, E199L, and/or F317L of an African swine fever virus; a transformant transformed by means of the recombinant vector; and a [vaccine](#) composition for preventing African swine

fever, comprising, as an active ingredient, African swine fever virus antigen protein(s) Lectin, CD2v, p72, p54, p30, p15, p35, E199L, and/or F317L, isolated from the transformant; and the like.

23. [4395820](#) NOVEL LIPID NANOPARTICLES FOR DELIVERY OF NUCLEIC ACIDS COMPRISING PHOSPHATIDYLSERINE

EP - 10.07.2024

Int.Class [A61K 39/39](#) Appl.No 22773176 Applicant CUREVAC SE Inventor BAUMHOF PATRICK

The invention relates to a [vaccine](#) composition comprising a) at least one nucleic acid encoding at least one antigen or fragment or variant thereof; and b) a carrier composition, wherein the carrier composition comprises the phospholipid phosphatidylserine. The present invention further relates to a pharmaceutical composition comprising the [vaccine](#) composition and a pharmaceutically acceptable carrier, diluent or excipient, and to the [vaccine](#) composition or pharmaceutical composition for use in the treatment or prophylaxis of (as well as a corresponding method of treatment thereof) infectious diseases; cancer or tumor diseases, disorders or conditions; specific liver diseases; allergies; or autoimmune disease, disorder or condition; in a subject. Still further, the present invention is concerned with a kit or kit of parts, comprising the [vaccine](#) composition or the pharmaceutical composition as well as a method of inducing an immune response in a subject. Finally, the present invention is concerned with a use of a [vaccine](#) composition or the pharmaceutical composition or the kit or kit of parts for (i) inducing an immune response and for (ii) inducing an antigen specific T-cell response in a subject.

24. [4392065](#) SHIGELLA [VACCINE](#)

EP - 03.07.2024

Int.Class [A61K 39/112](#) Appl.No 22761566 Applicant GLAXOSMITHKLINE BIOLOGICALS SA Inventor MICOLI FRANCESCA

The present invention relates to immunogenic compositions and their use in providing protection against illness caused by infection with Shigella. The immunogenic compositions comprise Shigella GMMA with particular doses of O-antigen.

25. [20240228981](#) ATTENUATED VIRUSES USEFUL FOR VACCINES

US - 11.07.2024

Int.Class [C12N 7/00](#) Appl.No 18512196 Applicant The Research Foundation for The State of University New York Inventor Eckard Wimmer

This invention provides an attenuated virus which comprises a modified viral genome containing nucleotide substitutions engineered in multiple locations in the genome, wherein the substitutions introduce synonymous deoptimized codons into the genome. The instant attenuated virus may be used in a [vaccine](#) composition for inducing a protective immune response in a subject. The invention also provides a method of synthesizing the instant attenuated virus. Further, this invention further provides a method for preventing a subject from becoming afflicted with a virus-associated disease comprising administering to the subject a prophylactically effective dose of a [vaccine](#) composition comprising the instant attenuated virus.

26. [4395746](#) STABILIZATION OF ANTIGENS FOR LONG TERM ADMINISTRATION IN TRANSDERMAL MICRONEEDLE PATCHES

EP - 10.07.2024

Int.Class [A61K 9/10](#) Appl.No 22865833 Applicant UNIV CONNECTICUT Inventor AGRAHARI VIVEK

Described herein are compositions and methods for stabilizing RNA and protein antigens for long-term storage and use in transdermal microneedle patches, methods for filling microneedles, and methods of use. A stabilized RNA [vaccine](#) composition comprises: a complex of RNA with one or more cationic polymers; and one or more cationic lipid entities. A method for stabilizing RNA comprises: forming a complex comprising the RNA with one or more cationic polymers; mixing the complex with one or more cationic lipid entities comprising

liposomes or lipid nanoparticles to form a lipid mixture; and drying the lipid mixture under vacuum. The compositions and methods may be employed in the preparation of **vaccine** medicaments.

27. [4392066](#) IMMUNOGENIC COMPOSITIONS AND THEIR USE

EP - 03.07.2024

Int.Class [A61K 39/145](#) Appl.No 22768820 Applicant OSIVAX Inventor LE VERT ALEXANDRE

The invention relates to immunogenic compositions and their use as a **vaccine** for the prevention of influenza disease in a human subject. More specifically, the invention relates to methods of use of an immunogenic composition as a **vaccine** or immunotherapy in the prevention or treatment of influenza disease in a human subject in need thereof, said immunogenic composition comprising: a fusion protein comprising (i) an influenza nucleoprotein antigen and, (ii) a carrier protein comprising a self-assembling polypeptide derived from C4bp oligomerization domain and a positively charged tail, wherein an amount of 180 µg, or more, of said fusion protein is administered to said human subject.

28. [20240216502](#) UNIVERSAL **VACCINE** PLATFORM

US - 04.07.2024

Int.Class [A61K 39/29](#) Appl.No 18504962 Applicant Hugh MASON Inventor Hugh MASON

The disclosure relates to vaccination compositions, for example, against human papillomavirus, Zika virus, and flu virus. The disclosure also relates to vectors for producing the virus-like particles and immune complex platforms of the vaccination compositions.

29. [4393955](#) ANTIBODY SPECIFICALLY BINDING TO CD47, RECOMBINANT ONCOLYTIC VIRUS THEREOF AND USE THEREOF

EP - 03.07.2024

Int.Class [C07K 16/30](#) Appl.No 22860533 Applicant SHANGHAI SINOBAY BIOTECHNOLOGY CO LTD Inventor XU JIANQING

Provided is an antibody capable of specifically binding to CD47 or an antigen binding fragment thereof. Also provided is a recombinant oncolytic virus, which is operably inserted with or comprises a gene coding sequence of an anti-CD47 antibody or a CD47 ligand, wherein the anti-CD47 antibody comprises an Fc mutant having A330L/I332E mutations (ALIE antibody), i.e., the anti-CD47 antibody is αCD47-Fc(ALIE). Also provided are a preparation method for the recombinant oncolytic virus, a use of the recombinant oncolytic virus in preparation of anti-tumor drugs, and a vaccinia virus Tiantan strain capable of efficiently expressing an αCD47-Fc(ALIE) gene.

30. [WO/2024/140767](#) POLYNUCLEOTIDE MOLECULE FOR PREVENTING OR TREATING HPV INFECTION-RELATED DISEASES

WO - 04.07.2024

Int.Class [C12N 15/62](#) Appl.No PCT/CN2023/142229 Applicant RINUAGENE BIOTECHNOLOGY CO., LTD. Inventor CEN, Shan

The present application relates to a polynucleotide molecule that can be used for preventing or treating HPV infection-related diseases, and a pharmaceutical product, a pharmaceutical composition, or an mRNA **vaccine** comprising said polynucleotide.

31. [4396197](#) VIRAL-LIKE PARTICLES FOR THE TREATMENT OR PREVENTION OF AN INFECTION BY A CORONAVIRIDAE VIRUS

EP - 10.07.2024

Int.Class [C07K 14/005](#) Appl.No 22765924 Applicant UNIV SORBONNE Inventor KLATZMANN DAVID

The invention pertains to new viral-like particles (VLPs), pharmaceutical compositions comprising the same and methods of using the same to prevent or treat an infection by a Coronaviridae virus. Advantageously, these VLPs can be used as a **vaccine** to be orally or nasally administrated.

32. [4392040](#) GLYCATED CHITOSANS FOR TREATMENT OF VIRAL INFECTIONS

EP - 03.07.2024

Int. Class [A61K 31/722](#) Appl. No 22862137 Applicant IMMUNOPHOTONICS INC Inventor LAM SIU KIT

Methods of treating, preventing and/or inhibiting one or more of the symptoms of a respiratory viral infection in a subject by administering a therapeutically effective amount of a glycosylated chitosan (GC) polymer with characteristics as disclosed herein or a [vaccine](#) composition comprising such GC polymer in combination with one or more viral antigens are provided. Methods of reducing morbidity and/or mortality of a respiratory viral infection in a subject, methods of inducing an innate immune response in mucosa of a subject, methods of generating mucosal secretory IgA antibodies and/or neutralizing serum IgG antibodies in a subject, and methods of vaccinating a subject against a respiratory viral infection, wherein these methods comprise administering to the subject a therapeutically effective amount of a glycosylated chitosan (GC) polymer as disclosed herein or a [vaccine](#) composition comprising such GC polymer in combination with one or more viral antigens are further provided.

33. [4392100](#) PRE-FILLED BLOW-FILL-SEAL INTRADERMAL INJECTION SYSTEM

EP - 03.07.2024

Int. Class [A61M 5/28](#) Appl. No 22790120 Applicant KOSKA FAMILY LTD Inventor KOSKA MARC

A pre-filled blow-fill-seal (BFS) IntraDermal (ID) medical agent injection system (600) assembled and configured to allow delivery of a single dose of a therapeutic agent (e.g., [vaccine](#), drug, medicament, etc.) from a BFS vial (610) to a patient in an auto-disable fashion.

34. [WO/2024/139848](#) NUCLEIC ACID DELIVERY CARRIER, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 04.07.2024

Int. Class [A61K 9/16](#) Appl. No PCT/CN2023/132937 Applicant CANSINO (SHANGHAI) BIOLOGICAL RESEARCH CO., LTD. Inventor WANG, Haomeng

A lipid nanoparticle composition, a preparation method therefor, and use thereof in nucleic acid delivery. An mRNA drug or [vaccine](#) can be prepared on the basis of the lipid nanoparticle composition. The starting materials of the lipid nanoparticle composition comprise an ionizable cationic lipid, an auxiliary phospholipid, a sterol compound, a lipid polyethylene glycol conjugate, and a buffer.

35. [WO/2024/147503](#) TROP2 IMMUNOGENIC PEPTIDE

WO - 11.07.2024

Int. Class [C07K 14/47](#) Appl. No PCT/KR2023/020268 Applicant ASTON SCIENCE INC. Inventor JUNG, Hun

The present invention relates to TROP2 immunogenic peptides and a use thereof. The TROP2 immunogenic peptide according to the present invention selectively binds to MHC class II to enhance only the immunogenicity of specific immune cells capable of killing cancer cells and thus can be advantageously used as an excellent cancer [vaccine](#) for the prevention and/or treatment of cancer by minimizing the immune escape mechanism of cancer cells.

36. [20240216493](#) CANCER THERAPY COMPOSITIONS AND USES THEREOF

US - 04.07.2024

Int. Class [A61K 39/00](#) Appl. No 18607029 Applicant HDT Bio Corp Inventor 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. Compositions described herein include lipid carriers, optionally including an inorganic particle, capable of admixing with nucleic acids. Nucleic acids provided herein include those encoding for cancer antigens (full length proteins or fragments) as well as antibodies. Methods of using the compositions as a therapeutic [vaccine](#) for the treatment of a cancer are also provided.

37. [WO/2024/148169](#) NANOPARTICLE COMPLEXES FOR ENHANCED SAFETY

WO - 11.07.2024

Int.Class [A61K 9/127](#)Appl.No PCT/US2024/010326Applicant HDT BIO CORP.Inventor KIMURA, Taishi

The disclosure provides compositions, methods of treatment, and methods of making and using compositions to deliver a nucleic acid to a subject that, optionally, have reduced reactogenicity and promotes a local innate immune response in the subject while promoting an adaptive immune response. Compositions described herein include nanoparticles, optionally including an inorganic particle, capable of admixing with nucleic acids encoding proteins, antibodies, or immunomodulators. Methods of using the compositions as a therapeutic [vaccine](#) for the treatment of an infection or cancer are also provided.

38.[20240226258](#)IN-VITRO TRANSCRIPT MRNA AND PHARMACEUTICAL COMPOSITION COMPRISING SAME

US - 11.07.2024

Int.Class [A61K 39/02](#)Appl.No 17905830Applicant ABION INC.Inventor Young Key SHIN

The present invention relates to an RNA in-vitro transcript mRNA for intracellular expression of a gene of interest and a pharmaceutical composition comprising same for vaccines. When injected into animal cells, the in-vitro transcript mRNA including the gene of interest according to the present invention allows the protein of interest to be expressed in the animal cell in large quantities, and, as such, can be used as a gene [vaccine](#) against autoimmune diseases, infectious diseases, cancer- or tumor-related diseases, inflammatory diseases, and so on.

39.[20240229143](#)FORMULATION OF PEPTIDE IMMUNOTHERAPIES

US - 11.07.2024

Int.Class [C12Q 1/6886](#)Appl.No 18256243Applicant IOGENETICS, LLCInventor Jane Homan

This invention provides a method for maximizing the immune response to mutated tumor specific proteins, either by means of stimulation of dendritic cells or T cells in vitro followed by administration of these cells to a patient, or by means of administration of a neoantigen [vaccine](#) in which de novo peptides, or their encoding nucleic acids, have been designed to ensure an appropriate level of binding affinity to a particular cancer patient's MHC alleles. This invention further provides for modulating the immune response in an immunopathology

40.[202370610](#)METHODS OF [VACCINE](#) ADMINISTRATION TO SALMONIDS

DK - 12.07.2024

Int.Class [A61K 39/02](#)Appl.No PA 2023 70610Applicant ZOETIS Services LLCInventor Adérito Luis MONJANE

The disclosure provides compositions and methods for salmonid vaccination against PMCV and at least one other pathogen

41.[4397318](#)IMMUNOSTIMULATORY FORMULATION, AND COSMETIC, FOOD, FEED ADDITIVE, AND QUASI-DRUG CONTAINING SAID IMMUNOSTIMULATORY FORMULATION

EP - 10.07.2024

Int.Class [A61K 47/64](#)Appl.No 21955905Applicant JAPAN ECO SCIENCE CO LTDInventor MIYAMOTO HIROKUNI

Without being limited to an epitope of a conventional antigen, it is possible to perform relatively simple formulation for activating a mucosal immune system and inducing highly diverse immunoglobulins by a non-invasive method. In a conventional [vaccine](#), it is difficult to select an epitope antigen, and it takes time to approve the [vaccine](#); therefore, it is assumed that it may be difficult to obtain an effect after approval for a pathogen target such as a virus that is likely to be mutated. In addition, since vaccination is not basically administered by the non-invasive method, suitable medical practice in a medical institution has been essential. Provided is an immunostimulatory preparation which is a protein obtained by fusing a bacteria-derived heat

shock protein (HSP: heat shock protein) and a viral peptide antigen, and enables induction of a highly diverse immunoglobulin by simultaneously stimulating nasal, respiratory, and oral administration. Provided are a feed, a feed additive, and an environmental symbiotic preparation for livestock, or a cosmetic, a food, and a quasi-pharmaceutical product for humans, which enable symbiosis with an environmental microorganism containing the immunostimulatory preparation. Provided are a preparation, a cosmetic for humans, a food, and a quasi-pharmaceutical product that enable symbiosis with environmental microorganisms utilizing livestock-derived IgA, IgG, and IgY among the immunoglobulins.

42. [20240228980](#) CRYPTIC PROTEINS EXPRESSED FROM DEFECTIVE VIRAL GENOMES INTERFERE WITH INFLUENZA VIRUS REPLICATION

US - 11.07.2024

Int.Class [C12N 7/00](#) Appl.No 18495338 Applicant Wisconsin Alumni Research Foundation Inventor Andrew Mehle

The disclosure provides for methods for making and using modified influenza gene products, alone or in combination, e.g., to inhibit wild-type influenza virus replication, to serve as an immunostimulatory agent, and/or as attenuated [vaccine](#) backbones. In one embodiment, the genomes of the DIPs provide for inhibitory activity, producing a dual effect in which both the RNA itself and the encoded protein coordinate to interfere with replication. Thus, the ability of DIPs to block replication of WT virus provides for a treatment for infection, use as an immunostimulatory agent, and as attenuated viruses for vaccination.

43. [4397676](#) NOVEL PEPTIDES AND COMBINATION OF PEPTIDES FOR USE IN IMMUNOTHERAPY AGAINST VARIOUS TUMORS

EP - 10.07.2024

Int.Class [C07K 14/47](#) Appl.No 24164808 Applicant IMMATICS BIOTECHNOLOGIES GMBH Inventor MAHR ANDREA

The present invention relates to a peptide, nucleic acid and cells for use in immunotherapeutic methods. In particular, the present invention relates to the immunotherapy of cancer. The present invention furthermore relates to a tumor-associated T-cell peptide epitope, alone or in combination with other tumor-associated peptides that can for example serve as an active pharmaceutical ingredient of [vaccine](#) compositions that stimulate anti-tumor immune responses, or to stimulate T cells ex vivo and transfer into patients. The peptide bound to molecules of the major histocompatibility complex (MHC), or the peptide as such, can also be the target of antibodies, soluble T-cell receptors, and other binding molecules.

44. [20240229007](#) INACTIVATED VACCINES & NUCLEIC ACID APTAMER THERAPEUTICS DERIVED THEREFROM

US - 11.07.2024

Int.Class [C12N 13/00](#) Appl.No 17313798 Applicant Universal Stabilization Technologies, Inc. Inventor Victor Bronshtein

Methods for producing dry thermostable inactivated vaccines with improved inactivation/killing without compromising antigenicity are disclosed. The methods provide for microbes to be irradiated with an ionizing radiation dose above 24 kGy without compromising the integrity of useful epitopes. To achieve enhanced inactivation, microorganisms (virions, bacterium, fungi, etc.) are first immobilized in a dry glassy matrix, then subsequently irradiated at elevated radiation dose while immobilized in the glass. Aptamers can be isolated from the inactivated microbes, such as inactivated SARS-COV-2, and modified or developed as therapeutics for neutralizing infectious disease. Additionally, non-neutralizing aptamers can be combined to produce an enhanced [vaccine](#).

45. [WO/2024/140251](#) PHARMACEUTICAL COMPOSITION FOR INDUCING IMMUNE RESPONSE OF MAMMAL AGAINST NOROVIRUS, PREPARATION METHOD THEREFOR, AND USE THEREOF

WO - 04.07.2024

Int.Class [A61K 39/12](#) Appl.No PCT/CN2023/138780 Applicant GRAND THERAVAC LIFE SCIENCES (NANJING) CO., LTD. Inventor LI, Jianqiang

Provided are a pharmaceutical composition for inducing an immune response of a mammal against norovirus, a preparation method therefor, and use thereof. The pharmaceutical composition is composed of the following components: i) a norovirus VLP protein, an active fragment of the protein, a variant of the protein, or a mixture of at least two thereof; ii) an aluminum adjuvant; iii) sodium chloride; and iv) water. Further provided are a [vaccine](#) and a kit comprising the pharmaceutical composition and uses thereof. The pharmaceutical composition is not only suitable for various norovirus valence types, but also has superior adsorption completeness, blocking antibody titer and specific IgG antibody titer.

46. [20240216491](#) NEOGLYCOCONJUGATES AS VACCINES AND THERAPEUTIC TOOLS

US - 04.07.2024

Int.Class [A61K 39/00](#) Appl.No 18433142 Applicant KORANEX CAPITAL Inventor Tze Chieh SHIAO

Neoglycoconjugates as immunogens and therapeutic/diagnostic tools are described herein. The neoglycoconjugates are produced by conjugating a carbohydrate antigen intermediate to a free amine group of a carrier material (e.g., carrier protein). The intermediate comprises a linker having a first end and a second end, the first end being conjugated to a carbohydrate antigen via a thio ether bond and the second end comprising a functional group reactable with a free amine group. Following coupling, the carbohydrate antigen becomes covalently bound to the carrier material via an amide, a carbamate, a sulfonamide, a urea, or a thiourea bond, thereby producing the neoglycoconjugate. Applications of the neoglycoconjugates as antigens, immunogens, vaccines, and in diagnostics are also described. Specifically, the use of (neo)glycoconjugates as [vaccine](#) candidates and other therapeutic tools against cancers, viruses such as SARS-CoV-2, and other diseases characterized by expression of aberrant glycosylation are also described.

47. [WO/2024/146614](#) HUMAN PAPILLOMAVIRUS TYPE 51 L1 PROTEIN MUTANT, METHOD FOR REDUCING DEGRADATION OF RECOMBINANT PROTEIN AND USE

WO - 11.07.2024

Int.Class [C07K 14/025](#) Appl.No PCT/CN2024/070680 Applicant BEIJING HEALTH GUARD BIOTECHNOLOGY, INC. Inventor WU, Shuming

A human papillomavirus type 51 L1 protein mutant, a method for reducing the degradation of a recombinant human papillomavirus type 51 L1 protein using same and the use thereof. The mutation of R at position 422 of the amino acid sequence of HPV51L1 into T or Q by means of a genetic engineering technique successfully solves a protein degradation problem thereof. Experiments show that the mutations both do not affect the expression of the corresponding mutated L1 protein, the degradation ratio of the modified L1 protein is obviously reduced, and the immunogenicity of the corresponding VLP is also not affected. By means of mutation of the human papillomavirus type 51 L1 protein, the protein degradation problem is avoided, so that the corresponding HPV51L1-VLP obtained by recombinant expression of the modified sequence or the assembly thereof is more suitable as a [vaccine](#) antigen protein to prevent this type of papillomavirus infection.

48. [4396206](#) METHODS FOR ISOLATION OF LIPID-DISC COMPOSITIONS AND USES THEREOF

EP - 10.07.2024

Int.Class [C07K 14/245](#) Appl.No 22769710 Applicant VIB VZW Inventor REMAUT HAN

The invention relates to the field of bacterial membrane protein structures. More specifically, the invention relates to lipid nanodiscs compartmentalized by SlyB protein oligomers isolated from the outer membrane of Gram-negative bacteria. More specifically, the invention provides for a SlyB nanodisc structure wherein the SlyB-oligomer forms the membrane scaffold protein belt, which is surrounded by outer saccharolipid moieties anchored to the SlyB proteins, and which encloses a lipid bilayer nanodomain containing one or more

phospholipid layers, wherein macromolecules such as (outer) membrane protein molecules may be captured and stabilized. More specifically, methods to produce and isolate chemically defined stable SlyB nanodisc particles are disclosed herein. Finally, the invention relates to the use of said SlyB nanodiscs as a self-adjuvanting vehicle, as part of an immunogenic composition, and provides for novel means for use in eliciting an immune response against macromolecules enclosed in said SlyB nanodiscs, or for use in a [vaccine](#) composition.

49. [WO/2024/146613](#) HUMAN PAPILLOMAVIRUS 56-TYPE L1 PROTEIN MUTANT, METHOD FOR REDUCING RECOMBINANT PROTEIN DEGRADATION, AND APPLICATION
WO - 11.07.2024

Int.Class [C07K 14/025](#) Appl.No PCT/CN2024/070679 Applicant BEIJING HEALTH GUARD BIOTECHNOLOGY, INC. Inventor CHEN, Xiao

A human papillomavirus (HPV) 56-type L1 protein mutant, a method for reducing the degradation of a recombinant HPV 56-type L1 protein, and an application. The modification of mutating the R at position 420 of an amino acid sequence of HPV56L1 into T or Q by using the genetic engineering technology successfully solves the problem of the degradation of the protein. The mutation does not affect the expression of a corresponding mutated L1 protein, the degradation ratio of a modified L1 protein is significantly reduced, and the immunogenicity of a corresponding VLP is not influenced. The HPV 56-type L1 protein is mutated and modified, so that the degradation problem is solved, the manufacturing difficulty is reduced, and the qualitative trait is improved. Thus, a corresponding HPV56L1-VLP obtained by recombinant expression of a modified sequence and assembly is more suitable as a [vaccine](#) antigen protein for preventing this type of papillomavirus infection.

50. [20240228547](#) POLYPEPTIDE FOR RESISTING NOVEL CORONAVIRUS AND APPLICATION THEREOF
US - 11.07.2024

Int.Class [C07K 14/005](#) Appl.No 18441563 Applicant INSTITUTE OF MICROBIOLOGY, CHINESE ACADEMY OF SCIENCES Inventor Fu GAO

The present application relates to a polypeptide for preventing or treating a novel coronavirus and an application thereof. The polypeptide is P3 polypeptide and any one of P3-1 polypeptide, P3-2 polypeptide, P3-3 polypeptide, P3-4 polypeptide, and P3-5 polypeptide derived from the P3 polypeptide. The amino acid sequences of the P3 polypeptide, the P3-1 polypeptide, the P3-2 polypeptide, the P3-3 polypeptide, the P3-4 polypeptide, and the P3-5 polypeptide are respectively shown in SEQ ID NOS: 1-6. The polypeptide of the present application has a strong inhibitory effect on original strains and a plurality of variant strains of the novel coronavirus, and can be used for preparing a drug or a [vaccine](#) for preventing and/or treating diseases caused by the novel coronavirus. The polypeptide is expected to also have the potential of preventing and/or treating new variant strains appearing in the future and sarbecovirus.

51. [313190](#) NASAL [VACCINE](#)-SPRAYING FORMULATION FOR SIMULTANEOUSLY TARGETING NASAL MUCOSA AND NASOPHARYNX
IL - 01.07.2024

Int.Class [A61K 35/76](#) Appl.No 313190 Applicant TOKO YAKUHIN KOGYO Co., LTD. Inventor KAMISHITA, Taizou

52. [WO/2024/139640](#) NOVEL CORONAVIRUS IGA ANTIBODY DETECTION IMMUNOPROBE, PREPARATION METHOD THEREFOR AND USE THEREOF
WO - 04.07.2024

Int.Class [G01N 33/558](#) Appl.No PCT/CN2023/127731 Applicant XIAMEN FORTUNE BIOTECH CO., LTD Inventor CHEN, Zhilong

A novel coronavirus IgA antibody detection immunoprobe, a preparation method therefor and a use thereof. The novel coronavirus IgA antibody detection immunoprobe comprises colloidal gold, wherein the colloidal gold adsorbs an anti-human IgA antibody protein or a new coronavirus spike protein. A novel coronavirus IgA antibody detection test strip comprises a bottom plate (7), and a sample pad (1), a gold pad (2), a chromatography membrane (5), and a water absorption pad (6) which are provided on the bottom plate (7) and arranged in sequence; the chromatography membrane (5) is a solid-phase nitrocellulose membrane composed of a test line (3) and a quality control line (4); the gold pad (2) is coated with an immunoprobe. When the immunoprobe coated on the gold pad (2) contains the anti-human IgA antibody protein, the test line (3) is coated with the new coronavirus spike protein; or when the immunoprobe coated on the gold pad (2) contains the new coronavirus spike protein, the test line (3) is coated with the anti-human IgA antibody protein; the quality control line (4) is coated with a quality control protein. The present invention is used for evaluating the level of IgA generated by nasal mucosa after a novel coronavirus nasal spray **vaccine** is inoculated.

53. **4396204** ORAL **VACCINE** VIA DENTAL BACTERIA AND EMITTED PEPTIDES TO PREVENT COVID-19 INFECTION

EP - 10.07.2024

Int. Class **C07K 14/005** Appl. No 22871020 Applicant KOTLYAR DAVID Inventor KOTLYAR DAVID

Disclosed is a pharmaceutical composition to prevent transmission of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the pharmaceutical composition comprising: genetically modified bacteria; sequences of small peptides; and pharmaceutical excipients, wherein the genetically modified oral bacteria are modified to translate, produce, and emit the sequences of small peptides which neutralize SARS-CoV-2 against COVID-19, wherein transgenic technology is used to modify the genetically modified oral bacteria to add genes in genetically modified oral bacteria that are transcribed to produce small peptides from the sequences of small peptides so added, wherein the sequences of small peptides show extreme binding and neutralization to SARS-CoV-2 but not to host proteins or processes, and wherein the pharmaceutical excipients aid the oral and/or nasal administration of the pharmaceutical composition.

54. **20240226281** ORAL **VACCINE** VIA DENTAL BACTERIA AND EMITTED PEPTIDES TO PREVENT COVID-19 INFECTION

US - 11.07.2024

Int. Class **A61K 39/215** Appl. No 18608668 Applicant David Kotlyar Inventor David Kotlyar

Disclosed is a pharmaceutical composition to prevent transmission of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), the pharmaceutical composition comprising: genetically modified bacteria; sequences of small peptides; and pharmaceutical excipients, wherein the genetically modified oral bacteria are modified to translate, produce, and emit the sequences of small peptides which neutralize SARS-CoV-2 against COVID-19, wherein transgenic technology is used to modify the genetically modified oral bacteria to add genes in genetically modified oral bacteria that are transcribed to produce small peptides from the sequences of small peptides so added, wherein the sequences of small peptides show extreme binding and neutralization to SARS-CoV-2 but not to host proteins or processes, and wherein the pharmaceutical excipients aid the oral and/or nasal administration of the pharmaceutical composition.

55. **20240226273** HIGHLY ATTENUATED REPLICATION-COMPETENT RECOMBINANT POXVIRUS AS A **VACCINE** PLATFORM AND METHODS OF USE

US - 11.07.2024

Int. Class **A61K 39/215** Appl. No 18283042 Applicant Arizona Board of Regents on behalf of Arizona State University Inventor Bertram Jacobs

Recombinant poxvirus expressing severe acute respiratory syndrome coronavirus 2 structural proteins and virus-like particles are described, along with methods of making and using the same.

56.[WO/2024/146109A](#) NOVEL MRNA **VACCINE** FOR THE TREATMENT AND PREVENTION OF HPV-ASSOCIATED LESIONS AND TUMORS

WO - 11.07.2024

Int.Class [A61K 39/12](#)Appl.No PCT/CN2023/107199Applicant VIROGIN BIOTECH (SHANGHAI)

LTD.Inventor ZHANG, Kuan

Immunogenic proteins are provided which have been derived from a strain of HPV which causes or is associated with causing cancer, which are mutated so that they do not cause increased cell proliferation. In various embodiments, the immunogenic protein is derived (e.g., mutated) from an HPV strain such as HPV 16, 18, 31, 33,35, 39, 45, 51, 52, 56, 58, 59, 66 or 68. Also provided are compositions (e.g., mRNA compositions) encoding such immunogenic proteins, as well as method for treating and preventing HPV infections.

57.[WO/2024/145863A](#) NOVEL MRNA **VACCINE** FOR THE TREATMENT AND PREVENTION OF HPV-ASSOCIATED LESIONS AND TUMORS

WO - 11.07.2024

Int.Class [C12N 15/62](#)Appl.No PCT/CN2023/070634Applicant VIROGIN BIOTECH (SHANGHAI)

LTD.Inventor ZHANG, Kuan

Immunogenic proteins are provided which have been derived from a strain of HPV which causes or is associated with causing cancer, which are mutated so that they do not cause increased cell proliferation. Within various embodiments of the invention the immunogenic protein is derived (e.g., mutated) from an HPV strain such as HPV 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66 or 68. Also provided are compositions (e.g., mRNA compositions) encoding such immunogenic proteins, as well as method for treating and preventing HPV infections.

58.[WO/2024/141791](#) COMBINED DELIVERY OF ANTIGENS AND TOLEROGENIC SIGNALS VIA DUAL-SIZED HYDROGEL SPHERES AND MOF COMPOSITES FOR TYPE-1

DIABETES **VACCINE** DEVELOPMENT

WO - 04.07.2024

Int.Class [A61K 47/69](#)Appl.No PCT/IB2023/000798Applicant THE HONG KONG UNIVERSITY OF SCIENCE AND TECHNOLOGYInventor CHUNG, Jin Teng

The subject invention pertains to a novel platform for the sustained release of Type-1 diabetes antigens. More specifically, the subject invention provides a combination of dual sized hydrogel particles for the delivery of Type-1 diabetes antigens and multi-component adjuvants to confer immune tolerance.

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