

**FINLAY**  
**EDICIONES**



# BOLETÍN VACCIENCIA

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 **IFV** INSTITUTO  
FINLAY DE  
VACUNAS

*...vacunar es prevenir.*

## Análisis bibliométrico sobre vacunas de ácidos nucleicos

Fuente de información utilizada:



Estrategia de búsqueda:

TOPIC: ("Nucleic acid vaccines") 153 records

Periodo de estudio 1999-2020

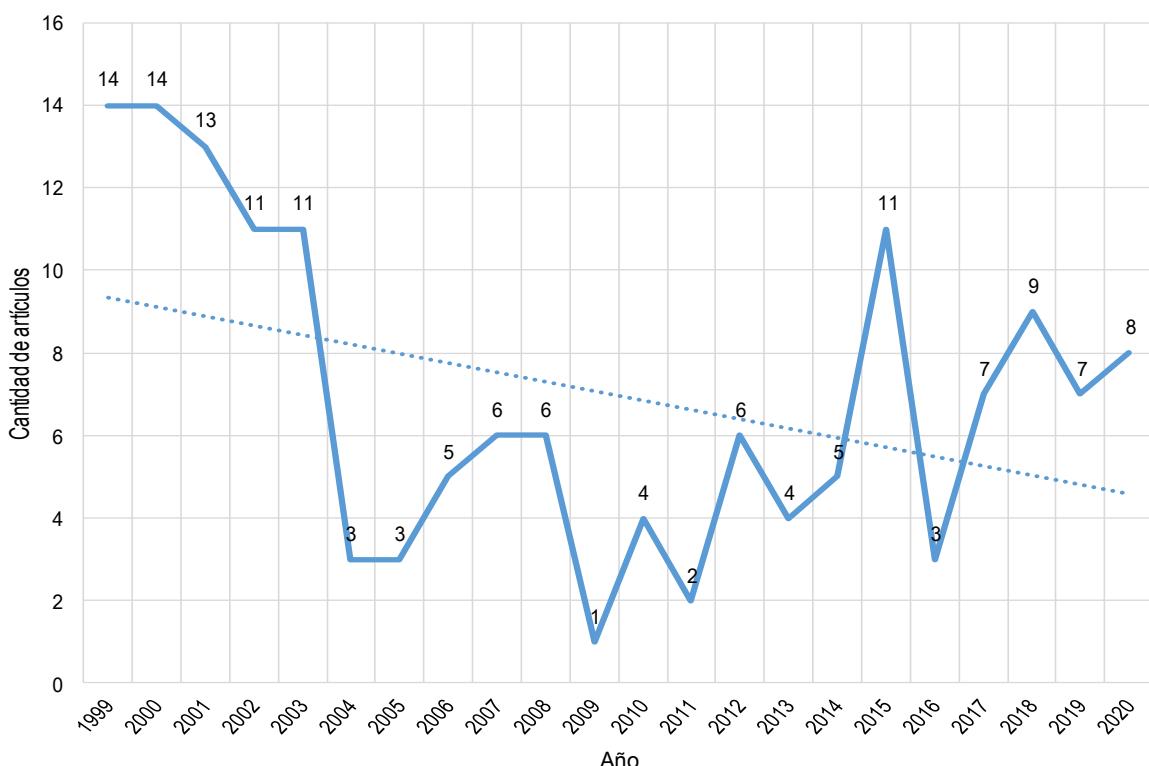
Las variables utilizadas en el análisis fueron:

- ⇒ Productividad científica por año.
- ⇒ Autores con mayor productividad científica.
- ⇒ Revistas con mayor número de publicaciones sobre el tema.
- ⇒ Instituciones que han trabajado el tema de estudio.
- ⇒ Países a la vanguardia sobre el tema.

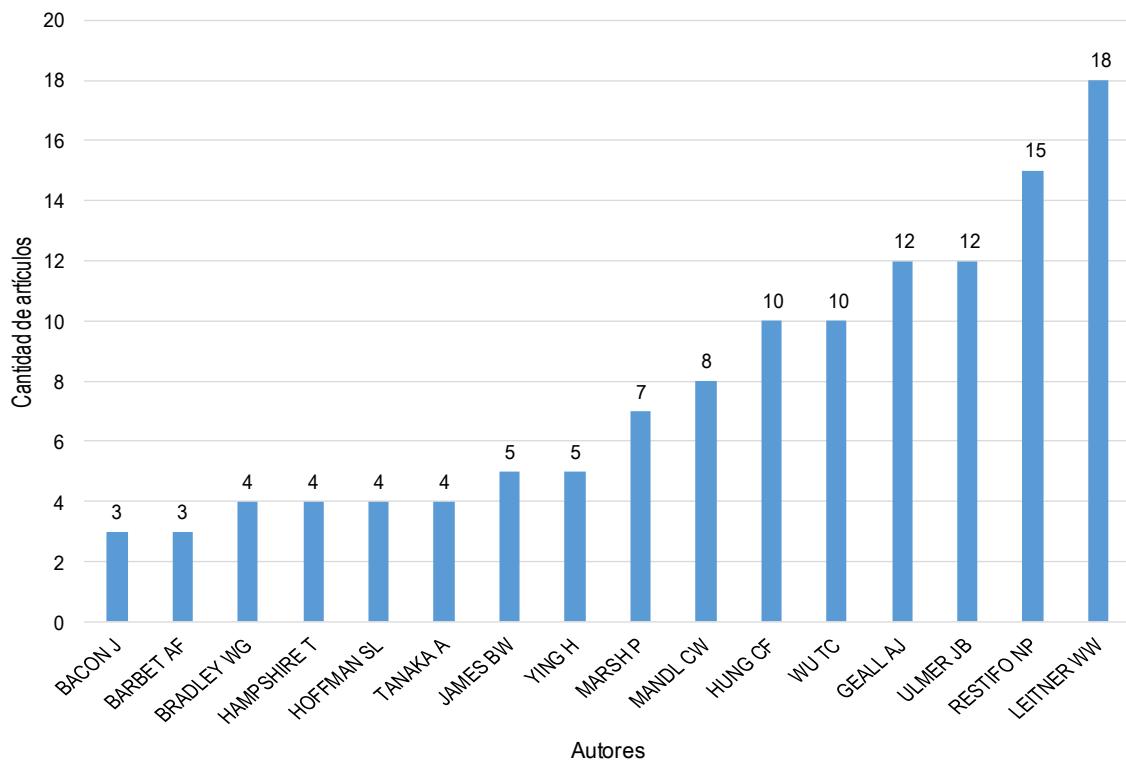
### EN ESTE NÚMERO

- \* Análisis bibliométrico sobre vacunas de ácidos nucleicos
- \* Noticias en la Web sobre vacunas
- \* Artículos científicos más recientes Medline sobre vacunas
- \* Patentes más recientes en PatentScope sobre vacunas
- \* Patentes más recientes en USPTO sobre vacunas

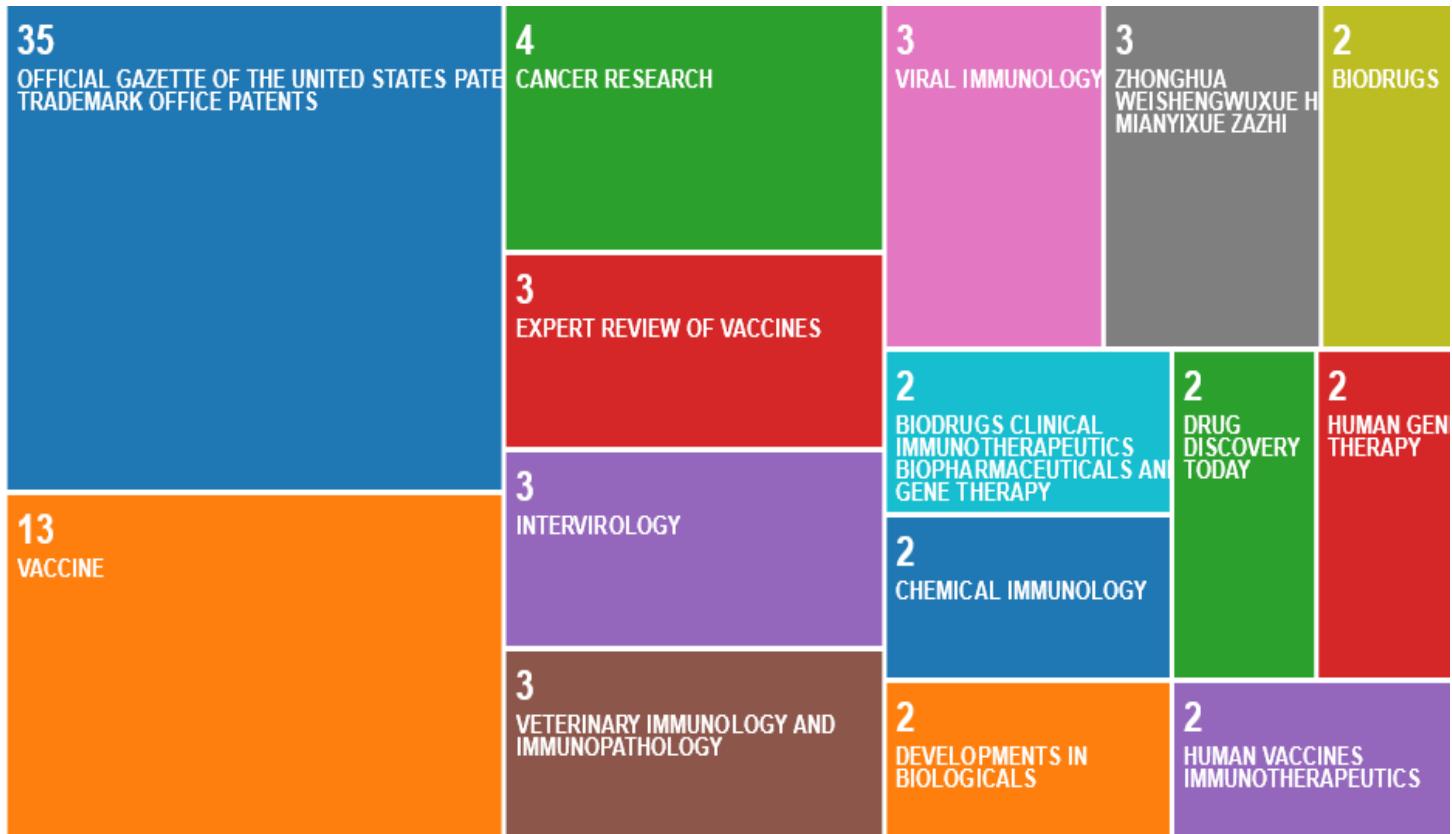
Productividad científica por año



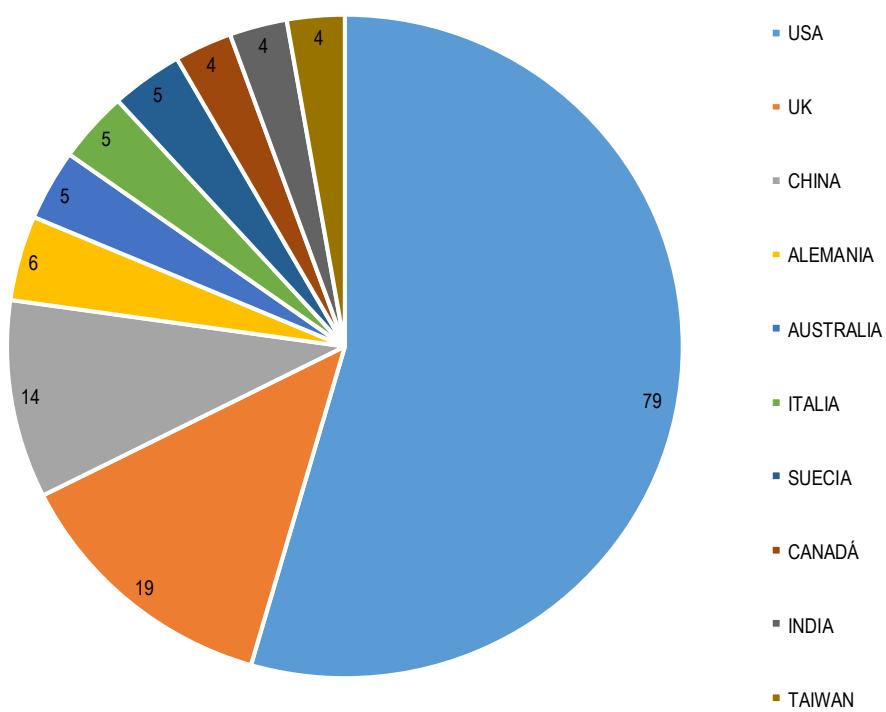
### Autores con mayor productividad científica



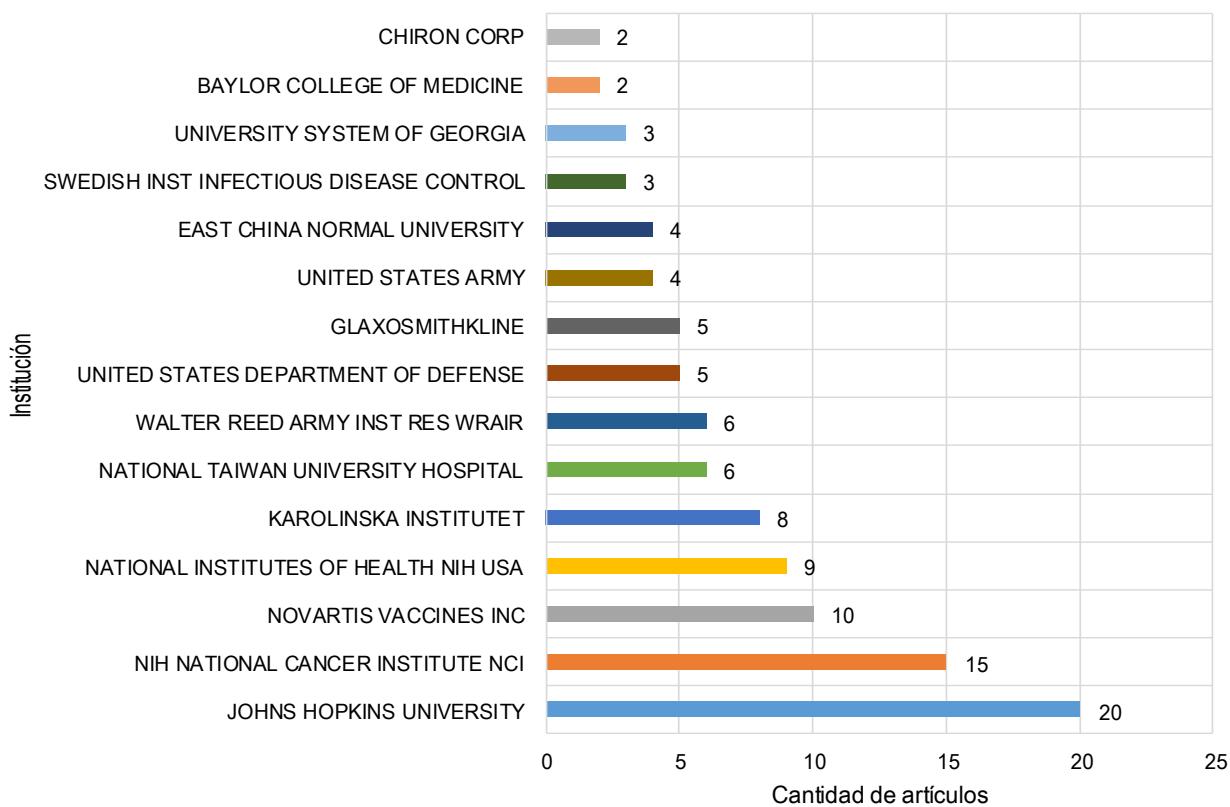
### Revistas científicas que han publicado sobre el tema (2019-2020)



### Producción científica por países registrada en Web of Science (1999-2020)



### Instituciones que han trabajado el tema de estudio



## Noticias en la Web

### ¿Qué hay de cierto en que la mutación del nuevo coronavirus lo volvería más infeccioso?

**1 jul.** Científicos identifican la mutación del virus SARS-CoV-2 que hoy se presenta con una mayor proporción y afecta a los alveolos pulmonares con más fuerza. Lo explica el doctor José Fernández, académico de la Universidad de Miami.

La estructura molecular del SARS-CoV-2, el virus causante de la COVID-19, cambió a su llegada a Estados Unidos y Europa. Uno de sus aminoácidos cambió al salir de China, es decir, mutó. ¿Lo hace más peligroso esta alteración? El doctor Elmer Huerta comenta al respecto.

Un virus es en esencia una gran molécula central de ácido nucleico que puede ser de ADN o de ARN (el nuevo coronavirus es de este tipo), el cual está protegido por una cápsula compuesta por grasas, azúcares y proteínas.

Esta cápsula es muy importante en el ciclo evolutivo del virus porque en ella se encuentran las moléculas o llaves que le permiten entrar a las células.

Tengamos presente las imágenes sobre el nuevo coronavirus. Esas proyecciones como patitas o antenitas se llaman espigas y son las que

usa el virus para ingresar a las células que tienen los receptores ACE2.

En otras palabras, las espigas del nuevo coronavirus son como llaves que van tras las cerraduras o receptores celulares ACE2 para ingresar, buscar el aparato reproductor y multiplicarse.

Lo interesante, y ya entrando al tema de las mutaciones del virus, es que esa espiga es en realidad una proteína, cuya cadena está compuesta por aproximadamente 1.300 aminoácidos, los cuales tienen un orden muy específico.

La noticia, publicada en The Washington Post, es que el aminoácido que ocupa el lugar 614 en la cadena, que cuando apareció en China era el aminoácido D o ácido aspártico, al llegar a Europa y luego a Estados Unidos, cambió al aminoácido G o glicina. En la actualidad, el 95 % de los virus secuenciados en el laboratorio de un investigador en Estados Unidos, tiene esa nueva mutación.

Ese simple cambio, de aminoácido D a aminoácido G, que en el lenguaje de los virólogos se conoce como mutación D614G, es lo que preocupa a los expertos, porque



de acuerdo con algunos experimentos, y cuyas publicaciones aun no han sido revisadas por pares, haría que el virus tenga hasta 10 veces más facilidad para contagiarse.

Sin embargo, si hay algo en lo que son muy enfáticos los investigadores, y que quiero repetir para no levantar alarma, es que esa mutación D614G no hace que el virus sea un supervirus, más letal o mortífero, sino que hace que el virus pueda contagiarse con más facilidad.

Al parecer, aunque hay otras teorías, esa mutación D614G haría que la espiga del virus sea más estable y no se rompa al tratar de penetrar la célula. Esto hace que la infección de las células sea más eficiente, lo cual explicaría, según varios investigadores, la rápida diseminación de la enfermedad por Europa, Estados Unidos y ahora América Latina.

Fuente: CNN en Español. Disponible en <https://cutt.ly/lpG71Mt>

*...vacunar es prevenir.*

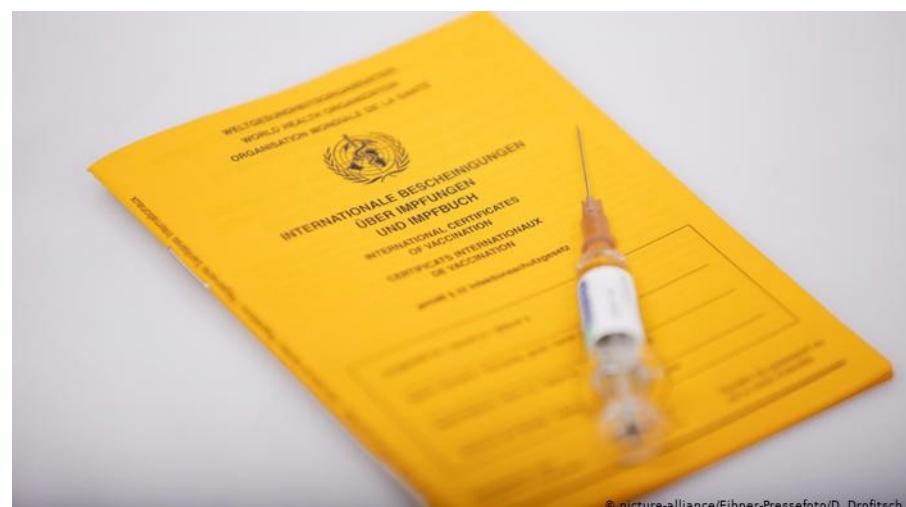
## OMS advierte que ninguna vacuna contra el COVID-19 está suficientemente avanzada

**3 jul.** Experto de la organización llamó, por ello, a no bajar los brazos y a adoptar medidas incluso de confinamiento. "Debemos iniciar el combate ahora, parar a este virus ahora", sostuvo.

El director de emergencias sanitarias de la Organización Mundial de la Salud (OMS), Mike Ryan, advirtió este viernes (03.07.2020) que ninguna de las decenas de vacunas que se están investigando para frenar el avance del COVID-19, o de las 17 que están en ensayos clínicos, está lo suficientemente avanzada como para pronosticar cuándo podría empezar a producirse una inoculación eficaz y segura.

"Sería poco inteligente predecir cuándo tendremos una vacuna lista", sostuvo Ryan, quien, sin embargo, estima que para finales de este año se podrían tener resultados sobre la eficacia de las vacunas candidatas. En ese caso se podría empezar con vacunaciones a principios del próximo año, pero ello dependerá de que haya una capacidad de producción suficiente, agregó.

Por lo mismo, el experto pidió a los países no bajar los brazos en el combate contra la pandemia, en momentos en que en distintos



© picture-alliance/Eibner-Pressefoto/D. Drofitsch

lugares del mundo comienzan a surgir rebrotos tras reaperturas prematuras. "La OMS comprende perfectamente que hay buenas razones para que los países quieran relanzar sus economías, pero no se puede ignorar tampoco el problema, no va a desaparecer como por acto de magia", señaló. Confinar más, si es necesario Ryan sostuvo que el desafío estará en reforzar la capacidad de producción al mismo tiempo que se avanza con los ensayos clínicos, lo que -confirmó- varios grupos farmacéuticos planean hacer. Los especialistas también están buscando tratamientos que permitan reducir la tasa de mortalidad. "Estamos viendo con nuevos antivirales o con combinaciones (en tratamientos)

de antivirales con antiinflamatorios para aumentar la respuesta inmunitaria", comentó Ryan.

De cualquier modo, el representante de la OMS hizo un llamado a las autoridades. "Ya es hora de que los países miren las cifras. Por favor, no ignoren lo que les dicen los números. La gente debe despertar. Las cifras no mienten y la situación en el terreno no miente", añadió. Subrayó que "nunca es demasiado tarde, en una epidemia, para tomar el control. Debemos iniciar el combate ahora. Debemos parar este virus ahora".

Ryan dijo que los países "deben absolutamente romper las cadenas de transmisión", inclusive adoptando medidas de confinamiento "si no hay alternativa".

Fuente: DW. Disponible en <https://cutt.ly/ja14upl>

## China's COVID-19 complex to produce over 100M vaccine

**3 jul.** China has completed the construction of a research laboratory and workshop complex in the city of Wuhan for producing vaccines to combat COVID-19 or coronavirus

pandemic, the state-run media said on Friday.

While the laboratory is capable to research and study pathogenic virus vaccines, the workshop will produce over 100 million doses

of the COVID-19 vaccine annually, reported Xinhua news agency, quoting the China National Pharmaceutical Group (Sinopharm). The complex was earlier hit by the coronavirus pandemic, which first

emerged in Wuhan city located in Central China's Hubei province last December.

China National Biotec Group has also built another workshop in the capital Beijing which will also produce the anti-COVID-19 vaccines.

"The total annual production capacity of inactivated COVID-19 vaccines is expected to exceed 200 million doses, which will help ensure adequate supply," the report cited Yang Xiaoming, president of the group, as saying.

Fuente: AA Asia-Pacific. Disponible en <https://cutt.ly/ya15luP>

Wuhan Institute of Biological Products had started the clinical trials of the vaccine to combat COVID-19 in April. So far it has been tested on 1,120 volunteers, aged between 18-59.

The report claimed, citing no officials, that the results of the trials "showed a good safety record" as "no severe adverse reactions were found."

It noted that the vaccine receivers "were inoculated two injections under different procedures and doses."

"For those receiving two injections at an interval of 28 days, the seroconversion rate of neutralizing antibodies reached 100 percent," claimed the report. According to China's National Health Commission, the country reported five new cases, including three imported and two indigenous, both from capital Beijing as the city has seen another wave of infections since the second week of June. China so far recorded 83,542 pandemic cases with 4,634 deaths. As many as 78,499

## Consecuencias del coronavirus: la "siniestra" transformación que el SARS-CoV-2 provoca en las células humanas infectadas

**3 jul.** Desde el inicio de la pandemia de COVID-19 una pregunta que ha obsesionado a varios científicos alrededor del mundo es: ¿cómo este coronavirus invade y reprograma a las células humanas para provocar la infección y causar la muerte?

Conocer la respuesta es crucial en la búsqueda de medicamentos capaces de frenar al virus antes de que lleve a cabo esos procesos.

Un equipo internacional de científicos que ha estado explorando esta interacción descubrió varias claves de cómo el SARS-CoV-2 infecta las células.

El hallazgo más sorprendente -que lograron comprobar con extraordinarias imágenes- es que las células humanas infectadas por el coronavirus sufren una "siniestra" transformación.



Las células, siguiendo las instrucciones del virus, desarrollan largos filamentos, similares a tentáculos, que, se cree, podrían ayudar a la rápida propagación por el organismo.

"Lo que descubrimos es que el virus induce a la célula a crear estas protuberancias, que son como largas ramas o tentáculos", le dijo a BBC Mundo uno de los autores del

estudio, el profesor Pedro Beltrao, investigador del Instituto Europeo de Bioinformática del Laboratorio Europeo de Biología Molecular (EBI-EMBL), en Cambridge, Inglaterra.

"En otros virus se ha visto que (estas protuberancias) desempeñan un papel en la rápida propagación de la

infección porque le ayudan al virus a invadir células cercanas", agrega el investigador.

El estudio, en el que también participaron investigadores de la Universidad de California, San Francisco, y la Escuela Icahn de Medicina de Monte Sinaí, Nueva York, ambas en Estados Unidos, el Instituto Pasteur en Francia y la Universidad de Friburgo en Alemania, encontró también que varios medicamentos existentes podrían ser buenos candidatos para frenar la infección.

Estos medicamentos, muchos de los cuales fueron diseñados como tratamientos para cáncer, parecen bloquear las señales químicas que activan la creación de estas protuberancias.

### Replicación

Pero los investigadores también encontraron que el virus, además de provocar la creación de estos "tentáculos", lleva a cabo otras conductas dentro de la célula infectada.

"La finalidad principal del estudio fue tratar de encontrar fármacos que puedan evitar que el virus lleve a cabo cambios en la célula humana", explica Pedro Beltrao.

"Pero para lograr eso, necesitábamos primero entender cómo el virus toma control de los mecanismos de la célula para poder llevar a cabo su propia replicación", agrega.

El principal objetivo de un virus en

el organismo humano es crear copias de sí mismo para para poder propagar la infección.

Pero el virus no puede crear estas copias por sí solo. Necesita entrar a una célula, tomar el control de la maquinaria celular y manipularla para reproducirse.

"El virus no se puede replicar solo porque tiene un número muy pequeño de proteínas, así que tiene que tomar control de las proteínas de la célula humana", explica el investigador.

Entre estas proteínas hay varias que son clave, las llamadas enzimas quinasas, que son capaces de llevar a cabo modificaciones a otras proteínas que ya se ha producido.

Entonces el virus toma control de estas enzimas quinasas para llevar a cabo modificaciones en la célula y regular la actividad de esas enzimas.

Al alterar los patrones de las proteínas celulares, el virus puede promover su propia transmisión a otras células y avanzar su propagación.

### Filopodios

Tras analizar las modificaciones que el virus lleva a cabo, los científicos encontraron tres conductas principales en la célula infectada.

"Una de estas conductas es la creación de las protuberancias, los largos tentáculos", le dice a BBC Mundo Pedro Beltrao.

Estas protuberancias, llamadas filopodios, no son muy comunes pero ya se han visto que ocurren con otros virus, como el de Marburgo, explica el investigador.

En el pasado se ha visto que otros virus las utilizan tanto para salir de la célula afectada como para infectar otras células cercanas y acelerar así la infección.

Aunque en este estudio no se demostró cuál es la función de los filopodios, los investigadores creen que hay una "alta probabilidad" de que el SARS-CoV-2 también esté usando estos tentáculos para acelerar su propagación.

Lo que sí logró esta investigación es producir unas imágenes impresionantes de la célula infectada, que la muestran como nunca se había visto antes, donde se ven las extrañas estructuras de los filopodios creados por el coronavirus.

Las fotografías, captadas por la doctora Elizabeth Fischer de la Unidad de Microscopía de los Laboratorios Rocky Mountain en Estados Unidos, y científicos de la Universidad de Friburgo, Alemania, revelan cómo el virus brota de los filopodios que se expanden en múltiples ramificaciones.

Pero además de la creación de filopodios, el virus provoca otras modificaciones importantes en la célula infectada.

"Otra conducta que vimos es que la célula deja de dividirse en cierto punto particular del ciclo de división y pensamos que esto crea un

ambiente propicio para que el virus se replique", le explica a BBC Mundo Pedro Beltrao.

"Y la tercera conducta que detectamos es una mayor producción de citoquinas, responsables de la respuesta inflamatoria".

"Esto es importante, porque creemos que este es uno de los factores que pueden estar causando la exagerada inflamación en las etapas avanzadas de la enfermedad de COVID-19", agrega el investigador.

#### **Las principales responsables: las quinasas**

Los científicos descubrieron que las enzimas quinasas son las principales responsables de estas modificaciones en la célula.

El estudio investigó cómo el coronavirus reprograma a las células

humanas para provocar la infección.

Y la buena noticia, dicen los investigadores, es que ya existen muchos fármacos que podrían regular la actividad de las quinasas y, por lo tanto, podrían ser utilizados para tratar la COVID-19.

Los científicos probaron cerca de 70 fármacos existentes e identificaron siete, principalmente tratamientos anticancerosos y antiinflamatorios, que demostraron tener un efecto para inhibir la actividad de las quinasas.

En pruebas de laboratorio con líneas celulares, los fármacos lograron evitar la respuesta inflamatoria y detener la replicación del virus.

Ahora los investigadores esperan poder empezar ensayos clínicos

para probar los tratamientos en humanos.

El estudio, publicado en la revista Cell, demostró una vez más cómo la pandemia de coronavirus está acercando a investigadores de todo el mundo para trabajar juntos.

Y cómo la necesidad para entender más sobre este virus está acelerando los hallazgos científicos.

"Para mí, en lo personal, fue un proyecto científico fantástico, porque no todos los días puedes colaborar con tantos investigadores brillantes de diferentes partes del mundo", le dice a BBC Mundo Pedro Beltrao.

"Este fue un proyecto que en otra época hubiera tomado entre tres y cinco años, y se hizo en tres meses. Esto, para mí, fue algo increíble", agrega el científico.

Fuente: BBC News. Disponible en <https://cutt.ly/9a0o9xu>

## **Brasil autorizó ensayos de una vacuna contra la COVID-19 producida por una empresa china**

**4 jul.** Unos nueve mil brasileños participarán de los ensayos del producto de la compañía Sinovac. Anteriormente, autoridades locales habían habilitado las pruebas con la potencial vacuna de la Universidad de Oxford.

Brasil, epicentro latinoamericano de la pandemia, aprobó los ensayos de la potencial vacuna contra el coronavirus desarrollada por la empresa china Sinovac y que se aplicará a 9.000 voluntarios en el país.

La Agencia Nacional de Vigilancia Sanitaria (Anvisa), vinculada al Ministerio de Salud, indicó en una nota que las pruebas de la potencial vacuna, elaborada a partir de "cepas inactivas" del patógeno, servirán para "evaluar su seguridad y eficacia" en la inmunización contra la COVID-19.

Los ensayos se realizarán, según la previsión inicial, a 9.000 personas en los estados de Sao Paulo, el más azotado de Brasil por la pandemia, Rio Grande do

Sul, Minas Gerais y Paraná, además de en Brasilia, la capital del país.

El centro de investigación Instituto Butantan de Sao Paulo coordinará los estudios tras llegar a un acuerdo con el laboratorio chino para ejecutar la tercera y última fase de pruebas clínicas de la posible vacuna.

Esta es la segunda vez que las autoridades sanitarias brasileñas permiten probar la eficacia de

una vacuna candidata contra el nuevo coronavirus en el país, tras autorizar en junio pasado los ensayos de la fabricada por la Universidad de Oxford junto con la farmacéutica AstraZeneca.

Los test de esta última ya han empezado en un grupo de 2.000 personas y son conducidos por la Universidad Federal de São Paulo (Unifesp), con el apoyo financiero de la Fundación Lemann, del multimillonario brasileño Jorge Paulo Lemann.

El Ministerio de Salud anunció la semana pasada un acuerdo con la Universidad de Oxford y AstraZeneca que le permitirá producir esa vacuna.

El acuerdo compromete a Brasil con un desembolso de 288 millones de dólares por adquirir 100 millones de dosis y la transferen-

cia de tecnología para su producción de forma autónoma en el país.

La cartera reconoció que se trata de una inversión de riesgo debido a que la vacuna, aunque es una de las más prometedoras, aún está en su tercera fase de pruebas clínicas y su eficacia y seguridad aún no han sido comprobadas.

Brasil, con 63.174 muertes, de las que 1.290 se registraron el último día, y 1,5 millones de casos, es el segundo país más afectado en el mundo, después de Estados Unidos, y el epicentro latinoamericano de la pandemia.

El viernes, la OMS advirtió que ninguna de las decenas de vacunas para el COVID-19 que se están investigando o de las 17 que están en etapa de ensayos clínicos está lo suficientemente

avanzada como para pronosticar cuando podría empezar a producirse una vacuna eficaz y segura.

“Sería poco inteligente predecir cuando una vacuna estará lista”, dijo el director de Emergencias Sanitarias de la OMS, Mike Ryan, quien, sin embargo, estima que para finales de este año se podrían tener resultados sobre la eficacia de las vacunas candidatas.

En ese caso se podría empezar con vacunaciones a principios del próximo año, pero ello dependerá de que haya una capacidad de producción suficiente, agregó.

Fuente: infobae. Disponible en <https://cutt.ly/fa0Ozg2>

## Cuba cuenta con varios candidatos vacunales contra la COVID-19

**4 jul.** Tenemos varios candidatos vacunales y un grupo multidisciplinario que trabaja de forma acelerada para encontrar la vacuna contra la COVID-19, afirmó hoy el doctor Eduardo Martínez, presidente de BioCubaFarma.

Durante su comparecencia en el programa radio-televisivo de la Mesa Redonda, el especialista explicó que en la actualidad hay 205 vacunas registradas en el mundo, 21 de ellas en ensayos clínicos.

Agregó que en el caso de Cuba al llegar la pandemia con mayor

retraso que en naciones como China, se comenzó a trabajar en la vacuna después y para ello debían tener la información genética del virus y la secuencia.

En el país -dijo- existen personas con una amplia experiencia y se está trabajando de forma acelerada para cumplir todos los pasos, y estamos creando condiciones para producirla a gran escala, explicó. Añadió que a lo mejor no somos los primeros en tener la vacuna a nivel mundial, pero si vamos a ser de los primeros en lograr una

amplia cobertura de vacunación a la población, estamos convencidos que va a ocurrir así, enfatizó Martínez.

### Varias instituciones trabajan con candidatos vacunales

A su vez mencionó que el Instituto Finlay de Vacunas, el Centro de Inmunología Molecular y el Centro Ingeniería Genética y Biotecnología son los que llevan la delantera en el país en esa labor, aunque se irán incorporando otras instituciones para brindar su apoyo en esa importante tarea.

En otro momento de su intervención, el presidente de BioCubaFarma habló sobre las causas multifactoriales que inciden en la escasez de medicamentos, no solo en Cuba, sino a nivel mundial.

Aquí en el país -expresó- se nos adicionan problemas para adquirir materias primas, reactivos y piezas de repuesto, sobre todo debido al cruel bloqueo económico, comercial y financiero de Estados Unidos.

No obstante buscamos alternativas y podemos decir que nosotros también exportamos medicamen-

tos y los ingresos son aprovechados en la propia producción de medicinas muy necesarias para el pueblo, destacó.

### **Situación actual de los medicamentos en Cuba**

Asimismo se refirió a las afectaciones con las navieras habituales, provocando excesivos tiempos logísticos; mientras puso como ejemplo que un barco demoró dos meses en llegar al país con 22 tipos de medicamentos.

Explicó que se nos ha ido agotando la materia prima y eso ha influido igualmente en el desabastecimiento existente en los meses

recientes. Además, resaltó que los medicamentos por lo general requieren de más de 10 componentes y si solo falta uno se dificulta la elaboración de estos.

Al referirse a los de mayor afectación, habló de los antiasmáticos, antihipertensivos y antialérgicos; así como los antibióticos que han estado muy afectados por falta de materias primas traídas de China y la India, puntualizó.

Agregó que se sigue trabajando buscando de manera permanente nuevos proveedores y tratando de agilizar la llegada de materias primas.

Fuente: tele pinar. Disponible en <https://cutt.ly/Ea2eJg7>

## **Los misterios sin respuesta del coronavirus y la covid-19**

**6 jul.** Hace seis meses el mundo comenzó a conocer la aparición del nuevo coronavirus SARS-CoV-2 en Wuhan (China), una ciudad de 11 millones de habitantes. Este virus fue relacionado de manera inmediata con el ya conocido SARS, que en el 2003 causó la muerte de 800 personas.

Para entonces, pocos dimensionaban que este nuevo agente que causa la enfermedad COVID-19 fuera el generador de la más grave crisis de salud pública en más de 100 años y que a la fecha, después de convertirse en una pandemia, ha matado a más de medio millón de personas y ha comprometido la salud de al menos 10 millones de personas en todo el mundo.

Y si bien durante este tiempo el virus ha impulsado procesos de

investigación para tratar de comprender la dinámica viral, sus efectos y sus posibles tratamientos, aun quedan preguntas sin responder que inquietan a la ciencia y sin un reto para el conocimiento acumulado que tiene como objetivo librarse de sus negativos efectos lo más pronto posible.

La revista científica Nature acaba de publicar un artículo en el que recoge y analiza los principales cuestionamientos sobre el SARS-CoV-2 y sobre los cuales los investigadores no tienen respuesta todavía.

### **¿Por qué las personas responden de manera diferente al virus?**

Las estadísticas demuestran que los afectados por COVID-19 tienen marcadas diferencias en los efectos de la enfermedad. De hecho,

algunas nunca desarrollan síntomas mientras que otras, incluso aparentemente sanas, producen neumonías graves y hasta mueren.

De acuerdo con Kari Stephansson, director ejecutivo de DeCODE Genetics en Islandia, las diferencias dramáticas pueden estar fundamentadas en variaciones determinadas por los genes humanos que no han sido analizadas en profundidad en razón a que aún no se cuenta con la suficiente cantidad de análisis.

Sin embargo, hay orientaciones que demuestran, por ejemplo, cómo en Italia quienes desarrollaron insuficiencia respiratoria grave tenían la posibilidad de portar una o dos variantes genéticas particulares al compararlos con quienes no tenían la

enfermedad. Una de estas variantes estaría en el genoma que determina el tipo de sangre ABO. Hay que decir, en todo caso, que no es el tipo de sangre el que determina la severidad o no sino las características generales de las personas con ciertos grupos de sangre.

La otra variación estaría cerca de varios genes que codifican una proteína que interactúa con el receptor que facilita la entrada del virus a las células; y otros que determinan la respuesta inmunitaria contra los patógenos, agrega Stephansson en la revista *Nature*.

Hoy este tema se sigue investigando a través de búsquedas con análisis de genomas completos en personas sanas y que hayan tenido casos graves.

### **¿Existe la inmunidad contra el coronavirus?**

La respuesta a esta pregunta tiene el acelerador puesto en todos los grupos de investigación inmunológica en razón a que la duración de las defensas que deja el virus en el cuerpo es determinante no solo para definir la evolución de la pandemia sino también para la potencial elaboración de tratamientos o vacunas. Los estudios han encontrado que los niveles de anticuerpos contra el SARS-CoV-2 permanecen altos durante algunas semanas después de la infección, pero luego empiezan a disminuir. Llama la atención que estas defensas pueden permanecer

altas por más tiempo en personas que habían padecido enfermedades graves. "Mientras más virus, más anticuerpos que más durarán", dice en *Nature* el inmunólogo George Kassiotis, del Instituto Francis Crick de Londres, sobre un patrón que se ha observado en otras infecciones virales como las del Sars.

En esos casos se demostró que la mayoría de las personas perdieron sus anticuerpos en los primeros años, pero quienes tuvieron las formas más graves los mantuvieron incluso después de 12 años. Y si bien los investigadores aún desconocen las reacciones inmunológicas específicas contra el SARS-CoV-2, se cree que la inmunidad va más allá de los anticuerpos y están mediadas también por células (linfocitos-T), importantes en las defensas a largo plazo, aunque aún no hay un marcador claro y medible.

### **¿El nuevo coronavirus ha mutado de manera preocupante?**

Todos los virus mutan a medida que infectan a las personas, y el SARS-CoV-2 no es la excepción, recuerda *Nature*. De hecho, los investigadores moleculares han rastreado estas mutaciones para seguir la propagación del virus a nivel mundial, pero además se buscan cambios estructurales sustanciales que puedan generar linajes más agresivos o con mayor capacidad de transmisión.

David Robertson, biólogo computacional de la Universidad de Glasgow, le dijo a esta revista que

al tratarse de un virus nuevo es importante saber si se torna más peligroso. Todo esto porque dichos cambios tienen la posibilidad de disminuir la efectividad de las vacunas, algo que complicaría la situación.

Por ahora, la mayoría de las mutaciones encontradas no tienen impacto, por lo que los investigadores buscan si hay cambios en otra dirección. En ese sentido, *Nature* aclara que las variaciones encontradas en algunos lugares y que en un comienzo fueron clasificadas como más letales no tienen consistencia académica.

### **¿Cómo actuaría la vacuna?**

De acuerdo con *Nature*, las vacunas efectivas podrían ser la única forma para salir de la pandemia. Actualmente hay 200 proyectos en desarrollo y 20 ya se encuentran en ensayos clínicos.

Ya hay algunos datos en animales y humanos sobre estas etapas tempranas y estos sugieren que las vacunas podrían ser efectivas para prevenir la infección pulmonar, pero no en otras partes. De hecho, la vacuna de la Universidad de Oxford podría prevenir el desarrollo de las formas graves, pero no la propagación del virus, según datos en monos.

Y aunque los datos en humanos son escasos y se ha encontrado que algunas promueven la creación de potentes anticuerpos, no se tiene claro si son suficientemente altos para detener nuevas infecciones y si persisten en el tiempo.

Lo cierto es que, según Dave OconCon, virólogo de la Universidad de Wisconsin, citado por Nature, se podrían tener vacunas clínicas útiles dentro de 12 o 18 meses, que tendrían que mejorarse progresivamente.

#### ¿Cuál es el origen del virus?

Nature indica que la mayoría de los investigadores coinciden que el SARS-CoV-2 puede estar relacionado con murciélagos, específicamente en la especie heradura, que alberga dos coronavirus estrechamente relacionados con este nuevo virus: el RATG-13

y el RmYM02, que comparten el 93 por ciento de la secuencia.

La revista aclara que después de analizar 1.200 coronavirus en murciélagos de China se sugiere que probablemente su origen sea en murciélagos de la provincia de Yunnan, pero no de otros países.

Los investigadores también aislaron coronavirus en pangolines de Malasia y estos comparten el 92 por ciento de su genoma con el SARS-CoV-2, aunque no se ha comprobado que haya saltado de estos animales a los humanos. *Nature* insiste que para rastrear

inequívocamente el paso de los animales a las personas es necesario encontrar una especie que albergue una versión genética que comparta más del 99 por ciento del genoma del SARS-CoV-2, algo que no por ahora no se considera fácil.

Zhang Zhigang, microbiólogo evolutivo de la Universidad de Yunnan, dice que con base en esta premisa las investigaciones han aislado virus de animales domésticos y silvestres en todo el sudeste asiático. Ese sigue siendo otro de los misterios alrededor del coronavirus.

Fuente: EL TIEMPO. Disponible en <https://cutt.ly/ra2fqFG>

## Pneumonia vaccine indirectly strengthens body vs COVID-19, says expert

**8 jul.** Vaccination against pneumonia helps the body indirectly strengthen itself against the coronavirus disease 2019 (COVID-19), according to a pediatric specialist.

Dr. John Ong urged the Philippines to continue using the pneumococcal conjugate vaccine (PCV) 13 as the country grapples with the coronavirus menace.

"If you are inoculated for pneumonia, you are indirectly

strengthening your body's defenses against COVID-19," he said in a radio interview.

Pneumonia is one of the potential complications of COVID-19.

Ong said the Philippines must stick to the broad-spectrum PCV 13, which combats 13 pneumococcal strains, rather than the PCV 10 because it is "more cost-effective."

He also cited a Department of

Health study showing that, with PCV 13, the country had "fewer cases of pneumonia, we had fewer deaths."

"So, our experience shows that it's still better if we use the more effective, rather than the cheaper, vaccine," Ong added.

A health expert from El Salvador earlier urged the Philippines to continue using the "superior" PCV13.

Fuente: GMA NEWS ONLINE. Disponible en <https://cutt.ly/3a2hD03>



*...vacunar es prevenir.*



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## Patentes registradas en PatentScope

Estrategia de búsqueda: Vaccine in the title or abstract AND 20200701:20200708 as the publication date

50 records

1. [WO/2020/134822](#) VACCINE CONTAINER HAVING FAILURE INDICATOR AND FAILURE INDICATION METHOD THEREOF  
WO - 02.07.2020

Clasificación Internacional [G01K 11/16](#) N° de solicitud PCT/CN2019/121406 Solicitante SHAANXI DINGWEI GEOTECHNICAL ENGINEERING CO., LTD. Inventor/a LIU, Yongqi

A vaccine container having a failure indicator (3) and a failure indication method thereof, the vaccine container comprising a vaccine bottle (1), the failure indicator (3) and a failure indicator box (4). The failure indicator box (4) has a sealed cavity structure, and the failure indicator (3) is placed in a cavity of the failure indicator box (4); and the failure indicator box (4) is fixedly connected to the vaccine bottle (1). The failure indicator (3) is obtained by uniformly mixing a component A and a component B; a raw material of the component A comprises a phase change heat storage material, an alkali solution and an emulsifier; a raw material of the component B comprises the same phase change heat storage material as the component A, a phenolphthalein indicator and an emulsifier; and the phase change temperature of the phase change heat storage material is 2-8°C. The vaccine container having the failure indicator (3) may absorb heat in the environment, control the temperature rise of a vaccine, and improve the stability of a vaccine transportation process, and when the temperature exceeds 8°C, the failure indicator (3) turns red, so that whether the vaccine fails or not may be indicated, and the safety of the vaccination is thus ensured.

## 2.[WO/2020/139308](#) INACTIVATED STAPHYLOCOCCAL LIQUID VACCINE

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000162 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The invention relates to vaccines for the treatment and prophylaxis of staphylococcal infection and can be used for a specific immunotherapy. The vaccine comprises: Staphylococcus aureus – 15 strains, Staphylococcus haemolyticus – 3 strains, Staphylococcus epidermidis – 3 strains. In accordance with the manufacturing method, an agar-grown bacterial culture is washed off the surface of a solid growth medium with pyrogen-free distilled water and placed into sterile containers; to remove growth medium impurities, the bacterial strains are washed with distilled water and subjected to centrifugation, following which the supernatant is discarded; the resulting pellet consisting of the biomass of the individually grown bacteriological strain culture is suspended in pyrogen-free distilled water for injection and standardised, wherein microorganism strains are taken in equal quantities; the standardised strain suspension is inactivated by autoclaving; equal quantities of a preparation of an embryonic origin or saline or water for injection are added to the containers with the inactivated bacteria. The vaccine is administered subcutaneously in the area of the inferior angle of the scapula, alternately in the right and left sides, the dose being increased at each subsequent injection. The invention makes it possible to increase the immunogenic effect of the vaccine and to enhance the immunological and clinical effects following vaccination while avoiding allergic reactions, and to reduce time required for producing the vaccine.

## 3.[WO/2020/139309](#) PSEUDOMONAS INACTIVATED VACCINE "PSEUDOPRIMAVAC" AGAINST PSEUDOMONAS

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000163 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The inventions relate to medical microbiology and the pharmaceutical industry, in particular to vaccine preparations, namely vaccines against Pseudomonas infection, methods for the production of said vaccines, and methods of treatment and prophylaxis, and can be used for therapeutic and prophylactic purposes in acute and chronic bacterial diseases of Pseudomonas etiology. A vaccine comprises at least 12 clinical pathogenic

strains of four types of bacteria of genus Pseudomonas, namely: 9 of *Pseudomonas aeruginosa*, 1 of *Pseudomonas maltophilia*, 1 of *Pseudomonas alcaligenes* and 1 of *Pseudomonas stutzeri*, isolated in various acute and chronic inflammatory diseases of bacterial etiology. The inventions increase the immunogenic effect of the vaccine, strengthen the immunological and clinical effect after vaccination without allergic reactions and provide a wide spectrum of application.

4. [WO/2020/139312](#) KLEBSIELLA- AND PROTEUS-BASED VACCINE “KLEPROPRIMAVAC” AGAINST KLEBSIELLA AND PROTEUS

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000166 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The inventions relate to medical microbiology, the pharmaceutical industry, and more particularly to vaccine preparations, namely vaccines against Klebsiella and Proteus, methods for production thereof and methods of treatment and prophylaxis, and can be used for the treatment and prophylaxis of localised and systemic, acute and chronic diseases of bacterial aetiology in children and adults. The vaccine comprises Klebsiella pneumonia, Klebsiella oxytoca, Proteus vulgaris, Proteus mirabilis, and Proteus peneri bacterial strains, no less than 2 Klebsiella species and 3 Proteus species in total, including 21 original bacterial strains, more particularly: Klebsiella pneumonia - 9, Klebsiella oxytoca - 3, Proteus vulgaris - 5, Proteus mirabilis - 3, and Proteus peneri – 1 of a specific location, which are isolated from various acute and chronic inflammatory diseases of bacterial aetiology in children and adults, which have various degrees of antibiotic resistance and include hospital strains having 100% antibiotic resistance. The inventions make it possible to increase the immunogenic effect of the vaccine, to enhance the immunological and clinical effects following the vaccination while avoiding allergic reactions, and to broaden the range of applications.

5. [WO/2020/139313](#) INACTIVATED STREPTOCOCCAL LIQUID VACCINE

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000167 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Artem Ihorovych

The invention relates to vaccines against Streptococcus and Staphylococcus and can be used for treatment of diseases of bacterial aetiology. The vaccine comprises no less than 4 Streptococcus species and 3 Staphylococcus species, including 21 Streptococcus and Staphylococcus strains. In accordance with the manufacturing method, an agar-grown bacterial culture is washed off the surface of a solid growth medium with pyrogen-free distilled water and placed into sterile containers; to remove growth medium impurities, the bacterial strains are washed with distilled water and subjected to centrifugation, following which the supernatant is discarded; the resulting pellet consisting of the biomass of the individually grown bacteriological strain culture is suspended in pyrogen-free distilled water for injection and standardised, wherein microorganism strains are taken in equal quantities; the standardised strain suspension is inactivated by autoclaving; equal quantities of a preparation of an embryonic origin or saline or water for injection are added to the containers with the inactivated bacteria; in order to control sterility, vaccine samples are plated on a nutrient broth, the plates are incubated in a thermostat, and in the absence of growth, a preservative is added and the vaccine is dispensed into sterile vials. The vaccination course consists of 10 to 12 injections which are administered every other day.

The invention makes it possible to increase the immunogenic effect of the vaccine and to enhance the systemic and local immunological and clinical effects following vaccination while avoiding allergic reactions.

6.[2579856](#)MHC Class I associated peptides for prevention and treatment of hepatitis B virus infection

GB - 08.07.2020

Clasificación Internacional [A61K 39/29](#) Nº de solicitud 201820614 Solicitante EMERGEX VACCINES HOLDING LTD Inventor/a RAMILA PHILIP

A vaccine composition comprising a peptide comprising a CD8+ T-cell epitope as set out in SEQ ID No.s 1-43 as define herein. The CD8+ T cell epitope may be capable of interacting with two different HLA super-types. The vaccine composition may comprise two or more hepatitis B viral (HBV) peptides, each comprising a different CD8+ T cell epitope. The peptide or peptides may be attached to a nanoparticle. The vaccine composition may comprise a peptide comprising a CD4+ epitope. The CD4+ T cell epitope may be capable of interacting with all HLA class II subtypes. The vaccine composition may be used to treat a HBV infection of serotype adr, adw, ayr, or ayw. The vaccine composition may be used to treat a HBV infection of genotype A, B, C, D, E, F, G, H, I, or J.

7.[20200206340](#)A VACCINE COMPRISING A PCV2 ORF2 PROTEIN OF GENOTYPE 2B

US - 02.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 16634756 Solicitante Intervet Inc. Inventor/a Melanie SNO

The present invention pertains to a vaccine comprising an ORF2 encoded protein of porcine circo virus 2 (PCV2) and a pharmaceutically acceptable carrier, for use in a method to protect a pig against an infection with porcine circo virus type 2 by administering the vaccine to the pig, wherein the vaccine comprises less than 20 µg per dose of the ORF2 encoded protein, the protein being of a porcine circo virus of genotype 2b.

8.[3673916](#)IMPFSTOFF GEGEN LAWSONIA INTRACELLULARIS UND PORCINES CIRCOVIRUS2

EP - 01.07.2020

Clasificación Internacional [A61K 39/02](#) Nº de solicitud 20153113 Solicitante INTERVET INT BV Inventor/a JACOBS ANTONIUS ARNOLDUS CHRISTIAAN

The present invention pertains to a vaccine comprising in combination killed whole cellLawsonia intracellularisbacteria and porcine circo virus 2 (PCV2) ORF2 protein for use in protecting a pig against an infection with Lawsonia intracellularis and PCV2 by an intradermal administration of the vaccine. The invention also pertains to a method to protect a swine against an infection withLawsonia intracellularisbacteria and PCV2.

9.[WO/2020/139311](#)"ECOPRIMAVAC" ESCERICHIA- AND ENTEROCOCCUS-BASED INACTIVATED LIQUID VACCINE AGAINST E. COLI AND ENTEROCOCCI, METHOD FOR PRODUCTION THEREOF, AND METHOD OF TREATMENT AND PROPHYLAXIS USING SAME

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000165 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The group of inventions relates to medical microbiology and the pharmaceutical industry, and more particularly to vaccine preparations, and even more particularly to vaccines against intestinal enterobacteriaceae and enterococci, methods for production thereof and methods of treatment and prophylaxis of localised and systemic, acute and chronic diseases of bacterial aetiology in children and adults. The EcoPrimavac Escherichia- and Enterococcus-based inactivated liquid vaccine against *E. coli* and Enterococcus comprises *Escherichia coli*, *Enterococcus faecalis*, *Enterococcus faecium* and *Enterococcus durans* bacterial strains, with no less than 22 bacterial strains in total, and more particularly: *Escherichia coli* - 9, *Enterococcus faecalis* - 8, *Enterococcus faecium* - 3, *Enterococcus durans* - 2 from specific locations, isolated from various acute and chronic inflammatory diseases of bacterial aetiology in children and adults, which have various degrees of antibiotic resistance and include hospital strains having 100% antibiotic resistance, inter alia, bacterial strains deposited at the Depository of the Institute of Microbiology and Virology of the National Academy of Sciences of Ukraine.

10.[3673069](#)BUNYAVIREN-VAKZIN

EP - 01.07.2020

Clasificación Internacional [C12N 15/67](#) Nº de solicitud 18759922 Solicitante CUREVAC AG Inventor/a PETSCH BENJAMIN

The present invention is directed to an artificial nucleic acid, particularly to an artificial RNA, and to polypeptides suitable for use in treatment or prophylaxis of an infection with a virus of the order Bunyavirales, particularly Severe fever with thrombocytopenia syndrome virus (SFTSV), Rift Valley fever virus (RVFV), or Crimean-Congo hemorrhagic fever virus (CCHFV), or a disorder related to such an infection. The present invention further concerns a Bunyavirales vaccine, particularly a SFTSV, RVFV, or CCHFV vaccine. The present invention is directed to an artificial nucleic acid, polypeptides, compositions and vaccines comprising the artificial nucleic acid or the polypeptides. The invention further concerns a method of treating or preventing a disorder or a disease, first and second medical uses of the artificial nucleic acid, polypeptides, compositions and vaccines. Further, the invention is directed to a kit, particularly to a kit of parts, comprising the artificial nucleic acid, polypeptides, compositions and vaccines.

11.[20200206336](#)A VACCINE FOR PROTECTION AGAINST STREPTOCOCCUS SUIS

US - 02.07.2020

Clasificación Internacional [A61K 39/09](#) Nº de solicitud 16634768 Solicitante Intervet Inc. Inventor/a Antonius Arnoldus Christiaan JACOBS

The present invention pertains to a vaccine comprising an IgM protease antigen of *Streptococcus suis*, for use in a method for protecting piglets having maternally derived anti-*Streptococcus suis* antibodies against *Streptococcus suis*, by administering the vaccine to the piglets at an age of at most 28 days, preferably before the piglets are weaned.

12.[WO/2020/138761](#)CHIMERIC VIRUS OF PORCINE REPRODUCTIVE AND RESPIRATORY SYNDROME VIRUS, AND VACCINE USING SAME

WO - 02.07.2020

Clasificación Internacional [C07K 14/005](#) Nº de solicitud PCT/KR2019/017319 Solicitante BIOPOA, INC.  
Inventor/a CHO, Sun-Hee

The present invention relates to a chimeric virus of porcine reproductive and respiratory syndrome (PRRS) virus, the chimeric virus being usable as a vaccine. A PRRSV chimeric virus of the present invention is more attenuated than the parent strain thereof, and thus promotes the secretion of neutralizing antibodies while having low pathogenicity and high stability, thereby being usable as a vaccine in the effective prevention and treatment of PRRS diseases.

13. [WO/2020/139310](#) "CANDOPRIMAVAC" INACTIVATED LIQUID VACCINE AGAINST CANDIASIS, METHOD FOR PRODUCTION THEREOF AND METHOD OF TREATMENT AND PROPHYLAXIS USING SAME

WO - 02.07.2020

Clasificación Internacional [A61K 36/06](#) Nº de solicitud PCT/UA2019/000164 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The group of inventions relates to medical microbiology and the pharmaceutical industry, and more particularly to vaccine preparations and to methods for the production and use thereof, and can be used for the treatment and prophylaxis of localised and systemic Candida-associated lesions. Moreover, the vaccine comprises no less than 14 strains of 3 species of Candida fungi, namely: Candida albicans - 7, Candida krusei - 4, Candida glabrata - 3 of different locations, isolated from various fungal lesions on the skin, mucosa and internal organs, inter alia, Candida fungi strains with various degrees of resistance to antifungal drugs, and which are deposited at the Depository of the Institute of Microbiology and Virology of the National Academy of Sciences of Ukraine.

14. [3675891](#) KOMBINATIONSKREBSTERAPIE

EP - 08.07.2020

Clasificación Internacional [A61K 38/17](#) Nº de solicitud 18851102 Solicitante MICROVAX LLC Inventor/a DEISSEROTH ALBERT B

A method and combination for treating a cancer patient by combining two distinct immuno-therapy solutions for administration to a patient within a common time period, comprising a checkpoint inhibitor antibody component such as a PD-1 or PD-L1 antibody administered by infusion, and a TAA/ecdCD40L vaccine component administered subcutaneously, wherein an initial antibody component administered is followed by at least several successive antibody boosts and an initial vaccine component administered is followed by at least several successive vaccine boosts, both the initial and boosts of each administered within at least said common time period, wherein the combined administration of said two distinct immuno-therapy solutions provides for an enhanced therapeutic effect, over that of the therapeutic effect of either of the two distinct immuno-therapy component solutions when administered alone as monotherapy.

15. [WO/2020/139298](#) CELLULAR AND HUMORAL IMMUNITY AUGMENTER MULTIPLE PHASE EMULSION VACCINE ADJUVANT AGAINST ANIMAL DISEASES

WO - 02.07.2020

Clasificación Internacional [A61K 39/39](#) Nº de solicitud PCT/TR2019/051208 Solicitante REAKIM ENDUSTRIYEL PERFORMANS KATKILARI KIMYEVI MADDELER SANAYI VE TICARET LTD. STI. Inventor/a BUYUKBAYRAM, Muhammet

The invention relates to vaccine adjuvant used with the purpose of protection of animal health in the field of veterinary. The referred adjuvant shows cellular and humoral immunity augmenter effect in its use. The adjuvant being the subject of the invention also increases the shelf life of vaccines, and provides a much longer life cycle for the vaccines due to its multiple phase emulsion. The referred vaccine adjuvant contained pharmaceutical mineral oil, surface active agents, and purified lyophilized additives from plants.

16. [2020204008](#) Gastrointestinal site-specific oral vaccination formulations active on the ileum and appendix  
AU - 02.07.2020

Clasificación Internacional [A61K 9/48](#) Nº de solicitud 2020204008 Solicitante Therabiome, LLC Inventor/a  
The invention provides oral vaccine formulations which deliver an antigen in the vicinity of the distal ileum and the area of the ileal Brake and/or the appendix. These vaccines are useful in the treatment and/or prevention of variety of disorders, including viral and bacterial infections and cancers. Related methods of treatment which use the oral vaccine formulations of the invention are also provided.

17. [WO/2020/138217](#) PREPARATION INCLUDING VACCINE ADJUVANT

WO - 02.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud PCT/JP2019/050947 Solicitante SUMITOMO DAINIPPON PHARMA CO., LTD. Inventor/a ONITA, Maiko

Provided is a composition that is useful as a vaccine adjuvant and has excellent storage stability and immunostimulatory activity. Specifically provided is a freeze-dried preparation that has high storage stability, said preparation containing a (4E, 8E, 12E, 16E, 20E)-N-{2-[{4-[(2-amino-4-[(3S)-1-hydroxyhexane-3-yl]amino)-6-methylpyrimidine-5-yl)methyl]benzyl}(methyl)amino]ethyl}-4,8,12,17,21,25-hexamethylhexacos-4,8,12,16,20,24-hexaenamide, squalene, a hydrophilic surfactant, and an oleophilic surfactant, and being characterized by containing an ascorbic acid-based antioxidant and an excipient.

18. [WO/2020/135898](#) DERIVADOS DE OLIGOSACÁRIDOS SINTÉTICOS COMO VACUNA CONTRA BORDETELLA PERTUSSIS

WO - 02.07.2020

Clasificación Internacional [C07H 15/04](#) Nº de solicitud PCT/CU2019/050001 Solicitante INSTITUTO FINLAY DE VACUNAS Inventor/a VÉREZ BENCOMO, Vicente Guillermo

La presente invención proporciona fragmentos de oligosacáridos sintéticos provenientes del pentasacárido terminal del lipooligosacárido de Bordetella pertussis, un método para obtener los fragmentos de oligosacáridos sintéticos y los conjugados a partir de los mismos. También proporciona las composiciones vacunales que contienen tales glicoconjungados y que inducen una respuesta inmune capaz de reducir la colonización nasofaríngea de Bordetella pertussis.

19. [WO/2020/136683](#) ADAPTATION OF ENTEROVIRUS TO VERO CELLS AND VACCINE FORMULATIONS THEREOF

WO - 02.07.2020

Clasificación Internacional [A61K 39/125](#) Nº de solicitud PCT/IN2019/050960 Solicitante BHARAT BIOTECH INTERNATIONAL LIMITED Inventor/a RAYCHOUDHURI, Amit

The present invention discloses Enterovirus D68 adapted to propagate to high titers in Vero cells and method of adaptation thereof. The present invention also discloses suitable vaccine composition comprising inactivated Enterovirus D68 antigen.

20. [3672630](#) HERSTELLUNG EINES GRIPPEIMPFSTOFFES IN MYCELIOPHTHORA THERMOPHILA

EP - 01.07.2020

Clasificación Internacional [A61K 39/145](#) Nº de solicitud 18848087 Solicitante DYADIC INT INC Inventor/a EMALFARB MARK

Recombinant expression of influenza virus surface proteins in the fungus *Myceliophthora thermophila* strain C1 is provided. The recombinant proteins are for use in influenza vaccine compositions.

21. [WO/2020/139316](#) POLYVALENT, COMBINED, INACTIVATED LIQUID VACCINE "UROPRIMAVAK", AND METHOD FOR THE PRODUCTION THEREOF

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000170 Solicitante MARKOV, Ihor Semenovich Inventor/a MARKOV, Ihor Semenovich

The group of inventions relates to medical microbiology and the pharmaceutical industry, in particular to: vaccine preparations, more particularly to vaccines against bacterial and fungal agents; methods for the production thereof; and methods for treatment and prevention, and can be used to treat and prevent acute and chronic inflammatory diseases of the genitourinary system in children and adults, in men and women, caused by non-specific agents of bacterial and/or fungal etiology. A course of vaccinations consists of 10-12 injections which are given every other day, wherein on the first day a dose of 0.1 ml is given intracutaneously in the inner surface of the forearm to form "peau d'orange" skin, and the following injections are given subcutaneously in a cycle alternating between the right upper arm/thigh and the left thigh/upper arm with gradually increasing doses of the preparation. An enduring, positive clinical effect, and a more prolonged general and local immunostimulative effect are provided.

22. [WO/2020/139315](#) "PNEUMOPRIMAVAC" POLYVALENT COMBINED INACTIVATED LIQUID VACCINE

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000169 Solicitante MARKOV, Ihor Semenovich Inventor/a MARKOV, Ihor Semenovich

The inventions relate to medical microbiology, and more particularly to vaccines against bacterial and fungal causative agents of bronchial and pulmonary diseases, as well as to methods for manufacturing same and to methods of treatment and prophylaxis, and can be used for the treatment and prophylaxis of local and systemic, acute and chronic diseases of bacterial and fungal origin in children and adults. The present vaccine contains strains of Klebsiella pneumoniae, Escherichia coli, Staphylococcus aureus, Staphylococcus haemolyticus, Streptococcus pyogenes, Streptococcus pneumonia, Streptococcus viridians, Enterococcus faecalis,

Enterococcus faecium, Enterococcus durans, Klebsiella oxytoca, Enterobacter aerogenes, Enterobacter cloacae, Proteus vulgaris, Proteus mirabilis, Citrobacter freundii, Alcaligenes faecalis, Morganella morganii, Pseudomonas aeruginosa, Pseudomonas alcaligenes, Candida albicans, Candida krusei, Candida glabrata, giving a total of no less than 20 species of bacteria and three species of Candida fungi, including not less than 38 strains of bacteria and five strains of fungi of the genus Candida, specifically: Staphylococcus aureus - 5, Staphylococcus haemolyticus - 3, Streptococcus pyogenes - 4, Streptococcus pneumonia - 2, Streptococcus viridians - 1, Enterococcus faecalis - 3, Enterococcus faecium - 1, Enterococcus durans - 1, Escherichia coli - 3, Klebsiella pneumonia - 2, Klebsiella oxytoca - 1, Enterobacter aerogenes - 2, Enterobacter cloacae - 1, Proteus vulgaris - 1, Proteus mirabilis - 1, Citrobacter freundii - 1, Alcaligenes faecalis - 1, Morganella morganii - 1, Pseudomonas aeruginosa - 3, Pseudomonas alcaligenes - 1, Candida albicans - 2, Candida krusei - 2, Candida glabrata - 1, isolated in the case of different acute and chronic inflammatory diseases of bacterial and fungal origin.

23. [20200208134](#) MEANS AND METHODS FOR PRODUCING PHOSPHATE CONTAINING CAPSULAR POLYSACCHARIDES

US - 02.07.2020

Clasificación Internacional [C12N 15/03](#) Nº de solicitud 16632998 Solicitante MEDIZINISCHE HOCHSCHULE HANNOVER (MHH) Inventor/a Timm Fiebig

The present invention relates to a host cell, which comprises under the control of a heterologous promoter a polynucleotide comprising a nucleotide sequence encoding a polypeptide capable of synthesizing a polysaccharide consisting of a dimeric repeating unit as well as to a vaccine composition comprising such host cell. Furthermore, either such host cell or a polypeptide expressed by such host cell is used for the production of a polysaccharide consisting of a dimeric repeating unit which may be used as a glycoconjugate vaccine.

24. [20200206329](#) NUCLEOTIDE SEQUENCE EXPRESSING AN EXOSOME-ANCHORING PROTEIN FOR USE AS VACCINE

US - 02.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud 16341042 Solicitante ISTITUTO SUPERIORE DI SANITA' Inventor/a Maurizio Paolo Maria FEDERICO

The present invention concerns a nucleotide sequence expressing a fusion protein, said fusion protein comprising or consisting of an exosome-anchoring protein fused at its C-terminus with an antigen, or a DNA expression vector comprising said nucleotide sequence, for use as vaccine.

25. [3677276](#) IMPFSTOFFZUSAMMENSETZUNG

EP - 08.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud 18849767 Solicitante UNIV TOHOKU Inventor/a SATO YASUFUMI

The present invention provides a vaccine composition for treating or preventing cancer expressing VASH2, containing a peptide including an amino acid sequence represented by SEQ ID NO: 4.

26.[WO/2020/139314](#) ANTIBACTERIAL POLYVALENT COMBINED INACTIVATED LIQUID VACCINE

WO - 02.07.2020

Clasificación Internacional [A61K 35/74](#) Nº de solicitud PCT/UA2019/000168 Solicitante MARKOV, Ihor Semenovych Inventor/a MARKOV, Ihor Semenovych

The inventions relate to medical microbiology, and more particularly to vaccines against bacterial and fungal causative agents, as well as to methods for manufacturing same and to methods of treatment and prophylaxis, and can be used for the treatment and prophylaxis of local and systemic, acute and chronic diseases of bacterial and fungal origin in children and adults. The present vaccine contains strains of Klebsiella pneumoniae, Escherichia coli, Staphylococcus aureus, Staphylococcus haemolyticus, Streptococcus pyogenes, Streptococcus pneumonia, Streptococcus viridians, Enterococcus faecalis, Enterococcus faecium, Enterococcus durans, Klebsiella oxytoca, Enterobacter aerogenes, Enterobacter cloacae, Proteus vulgaris, Proteus mirabilis, Citrobacter freundii, Alcaligenes faecalis, Morganella morganii, Pseudomonas aeruginosa, Pseudomonas alcaligenes, Candida albicans, Candida krusei, Candida glabrata, giving a total of no less than 20 species of bacteria and three species of Candida fungi, including not less than 38 strains of bacteria and five strains of fungi of the genus Candida, specifically: Staphylococcus aureus - 5, Staphylococcus haemolyticus - 3, Streptococcus pyogenes - 4, Streptococcus pneumonia - 2, Streptococcus viridians - 1, Enterococcus faecalis - 3, Enterococcus faecium - 1, Enterococcus durans - 1, Escherichia coli - 3, Klebsiella pneumonia - 2, Klebsiella oxytoca - 1, Enterobacter aerogenes - 2, Enterobacter cloacae - 1, Proteus vulgaris - 1, Proteus mirabilis - 1, Citrobacter freundii - 1, Alcaligenes faecalis - 1, Morganella morganii - 1, Pseudomonas aeruginosa - 3, Pseudomonas alcaligenes - 1, Candida albicans - 2, Candida krusei - 2, Candida glabrata - 1, isolated in the case of different acute and chronic inflammatory diseases of bacterial and fungal origin.

27.[3675902](#) ZUSAMMENSETZUNGEN UND VERFAHREN ZUR BEHANDLUNG VON KREBS UND INFektIONEN UNTER VERWENDUNG VON BAKTERIOPHAGEN UND DEREN MUTANTEN

EP - 08.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 18850295 Solicitante UNIV MICHIGAN STATE Inventor/a HUANG XUEFEI

Provided herein are vaccine composition comprising an antigen conjugated to a capsid, wherein the capsid comprises wild type or native sequence. Provided herein are also vaccine composition comprising an antigen conjugated to a capsid, wherein said capsid comprises at least one mutation, such as a non-natural mutation. Such compositions are useful in the treatment and prevention of pathogenic infections, inflammatory diseases, and neurodegenerative disease, and cancer, among others.

28.[WO/2020/138482](#) MICRONEEDLE ARRAY FOR BCG VACCINE

WO - 02.07.2020

Clasificación Internacional [A61M 37/00](#) Nº de solicitud PCT/JP2019/051571 Solicitante COSMED PHARMACEUTICAL CO., LTD. Inventor/a QUAN, Ying-shu

Because the needle tips do not come out of the tube in a conventional BCG vaccine tube needle, strength and technique were necessary and it was difficult to easily, evenly and reliably insert needles and perform a vaccination, which resulted in the problems that it was not possible to achieve the sufficient number of needle

marks that shows the inoculation effect and that the child being vaccinated experienced pain and distress. These problems are solved by adopting a microneedle patch case of a thermoplastic plastic as the BCG vaccination tube needle, and setting the needle length to 0.2-1.0 mm.

29.[3672625](#)SYNTHEtISCHER IMPFSTOFF

EP - 01.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud 18755823 Solicitante MEDIZINISCHE HOCHSCHULE HANNOVER Inventor/a WIRTH THOMAS

The present invention relates to a pharmaceutical combination of compositions for use in the treatment or prevention of a disease having cells bearing a target antigen as a vaccine and to a method for vaccination of a mammal, especially of a human for raising a cellular immune response directed against cells of the mammalian recipient, especially human recipient, which cells express a target antigen. The target antigen can e.g. be an autoantigen like a malignant antigen, i.e. a tumour-specific antigen. The pharmaceutical combination of compositions comprises a first composition and a second composition, wherein the second composition is for administration to recipient subsequent to the administration of the first composition, e.g. 2 to 10 days after the first composition. The pharmaceutical combination of compositions has the advantage of raising an effective antigen-specific T-cell response against cells bearing a target antigen that can be a malignant autoantigen, e.g. for raising an antigen-specific T-cell response against cells bearing a tumour-antigen. A further advantage is that the pharmaceutical combination of compositions can raise an antigen-specific T-cell response within a comparatively short time.

30.[3676285](#)IMPFSTOFF ZUR VERWENDUNG BEI DER PROPHYLAXE UND/ODER BEHANDLUNG EINER KRANKHEIT

EP - 08.07.2020

Clasificación Internacional [C07K 14/005](#) Nº de solicitud 18762507 Solicitante INPROTHER APS Inventor/a HOLST PETER

The present invention relates to an adenoviral vector capable of encoding a virus-like particle (VLP), said VLP displaying an inactive immune-suppressive domain (ISD). The vaccine of the invention shows an improved immune response from either of both of the response pathways initiated by CD4 T cells or CD8 T cells.

31.[20200206342](#)ENGINEERED VARIANTS OF HIV-1 ENV FOR PRESENTATION OF QUARTENARY EPITOPEs

US - 02.07.2020

Clasificación Internacional [A61K 39/21](#) Nº de solicitud 16631275 Solicitante THE BOARD OF TRUSTEES OF THE UNIVERSITY OF ILLINOIS Inventor/a Erik PROCKO

Provided herein are HIV-1 Env proteins or fragments thereof comprising one or more amino acid mutations; and nucleic acid molecule encoding the same. Further provided is a method of screening a compound for binding to one or more mutant HIV-1 Env proteins; and methods for eliciting an immune response against an HIV-1 infected cell, comprising administering to a subject an amount of a mutant HIV-1 Env protein, a fragment thereof or a mutant HIV-1 Env trimeric complex, or portion thereof, effective to elicit an immune response in the

subject. Further provided is a pharmaceutical composition, such as a vaccine, comprising the mutant HIV-1 Env protein or fragment thereof.

32. [20200207831](#) PEPTIDES AND COMBINATION OF PEPTIDES FOR USE IN IMMUNOTHERAPY AGAINST LEUKEMIAS AND OTHER CANCERS

US - 02.07.2020

Clasificación Internacional [C07K 14/74](#) Nº de solicitud 16805351 Solicitante Immatics Biotechnologies GmbH Inventor/a Juliane Sarah WALZ

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

33. [20200208108](#) INTERFERON PRIMED PLASMACYTOID DENDRITIC CELLS

US - 02.07.2020

Clasificación Internacional [C12N 5/0784](#) Nº de solicitud 16612441 Solicitante AARHUS UNIVERSITET Inventor/a Martin ROELSGAARD JAKOBSEN

A method is provided for producing a plasmacytoid dendritic cells (pDCs), wherein hematopoietic stem and progenitor cells (HSPCs) are provided and incubated in a first medium comprising cytokines and growth factor whereby the HSPCs are differentiated into precursor-pDCs and then adding interferons (IFNs) to the first medium to obtain a second medium whereby said precursor-pDCs are differentiated into pDCs. Furthermore, a technique is provided for producing genetically modified pDCs, by initially genetically modifying HSPCs using transfection methods, including electroporation, to deliver sgRNA and Cas9 protein. Moreover, a pharmaceutical formulation and a vaccine is provided which comprises pDC or genetically modified pDCs obtained according to that method.

34. [20200206338](#) METHOD OF TREATING MAMMALS DISPLAYING SEVERE NEUROLOGICAL SYMPTOMS OF ADVANCED CANINE DISTEMPER VIRUS INFECTION USING NDV-INDUCED SERUM

US - 02.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 16236258 Solicitante DANCLAY Properties, LLC Inventor/a Anupama Y. FIELDS

This application discloses methods, compositions, and articles of manufacture for using NDV-induced serum to treat non-human mammals, including dogs, displaying severe neurological symptoms of advanced canine distemper virus (CDV) infection. An off-the-shelf chicken vaccine is injected into two or-more healthy mammals, of the same species as the sick animal, to provoke an immune response. After allowing several hours for the immune response to develop, blood is drawn from the healthy mammals and centrifuged to obtain serum. This "NDV-induced serum" can be injected into a mammal of the same species displaying severe neurological

symptoms of advanced CDV infection on a prescribed schedule to bolster the sick animal's immune responses, preferably leading to clearance of the virus without the need for a spinal tap, which is the standard treatment for advanced CDV infection in mammals.

35.[20200206268](#) PEPTIDES AND COMBINATION OF PEPTIDES OF NON-CANONICAL ORIGIN FOR USE IN IMMUNOTHERAPY AGAINST DIFFERENT TYPES OF CANCERS

US - 02.07.2020

Clasificación Internacional [A61K 35/17](#) Nº de solicitud 16814508 Solicitante Immatics Biotechnologies GmbH Inventor/a Heiko SCHUSTER

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

36.[20200206337](#) LIVE ATTENUATED VACCINES

US - 02.07.2020

Clasificación Internacional [A61K 39/104](#) Nº de solicitud 16684086 Solicitante SERVICIO GALEGO DE SAÚDE (SERGAS) Inventor/a Germán Bou Arévalo

The present invention refers to a method for the production of live attenuated bacterial strains, suitable as vaccine candidates, comprising the steps of:

- A. providing a bacterial strain capable of expressing glutamate racemase and possibly D-amino acid transaminase and comprising a peptidoglycan cell wall, and
- B. inactivating the gene or genes encoding for the glutamate racemase enzyme and, if needed, the gene or genes encoding for the enzyme D-amino acid transaminase in such way that the bacterial strain is no longer capable of expressing a functional glutamate racemase and/or a functional D-amino acid transaminase;  
wherein the inactivation of said genes causes said bacterial strain to be auxotrophic for D-glutamate.

37.[3673917](#) NOROVIRUS-IMPFSTOFF

EP - 01.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 18248201 Solicitante THEMIS BIOSCIENCE GMBH Inventor/a TAUBER ERICH

The present invention provides immunogenic compositions, nucleic acid molecules and VLPs suitable as Norovirus vaccine candidates. Further provided are host cells for producing the biological material as well as methods for producing and/or purifying the immunogenic compositions and VLPs. Further provided is an immunogenic composition for use in methods of preventing/treating a Norovirus infection in a subject.

38.[WO/2020/136282](#)NOROVIRUS VACCINES

WO - 02.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud PCT/EP2019/087160 Solicitante THEMIS BIOSCIENCE GMBH Inventor/a TAUBER, Erich

The present invention provides immunogenic compositions, nucleic acid molecules and VLPs suitable as Norovirus vaccine candidates. Further provided are host cells for producing the biological material as well as methods for producing and/or purifying the immunogenic compositions and VLPs. Further provided is an immunogenic composition for use in methods of preventing/treating a Norovirus infection in a subject.

39.[20200207811](#)MALARIA VACCINE

US - 02.07.2020

Clasificación Internacional [C07K 14/02](#) Nº de solicitud 16634099 Solicitante OXFORD UNIVERSITY INNOVATION LIMITED Inventor/a Adrian V.S. HILL

The invention relates to a composition comprising a polypeptide comprising, or consisting of, the amino acid sequence of SEQ ID NO: 1, or a sequence having at least 80%, 85%, 90%, 95%, 98%, or 99% sequence identity to SEQ ID NO: 1 (R21), wherein said polypeptide is in the form of a virus-like particle (VLP), wherein said particle comprises less than 10% free hepatitis B surface antigen protein, for use in the immunisation of a human subject susceptible to *Plasmodium falciparum* infection, characterised in that said composition is administered in a dosage regimen of at least one dose of 1 µg to 20 µg R21 per administration for a subject at least 18 years old, or at least one dose of 0.5 µg to 10 µg R21 per administration for a subject less than 18 years old. The invention also relates to kits, methods and uses.

40.[20200206333](#)COMPOSITIONS FOR INDUCING AN IMMUNE RESPONSE

US - 02.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud 16708218 Solicitante President and Fellows of Harvard College Inventor/a Nisarg J. Shah

Acute myeloid leukemia (AML) is a clonal disorder of hematopoietic stem and progenitor cells. It is a devastating disease with a poor prognosis and an average 5-year survival rate of about 30%. Disclosed herein are composition and methods for treating leukemia with a biomaterial comprising a polymer scaffold, a dendritic cell activating factor, a dendritic cell recruitment factor, and at least one leukemia antigen. The biomaterial-based vaccine disclosed herein promotes a potent, durable and transferable immune response against acute myeloid leukemia to prevent cell engraftment and synergizes with chemotherapy to prevent relapse.

41.[20200207832](#)NOVEL PEPTIDES AND COMBINATION OF PEPTIDES FOR USE IN IMMUNOTHERAPY AGAINST OVARIAN CANCER AND OTHER CANCERS

US - 02.07.2020

Clasificación Internacional [C07K 14/74](#) Nº de solicitud 16815965 Solicitante immatics biotechnologies GmbH Inventor/a Heiko Schuster

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

42.[3677279](#)KONFORMATIONSSTABILISIERTE RSV-F-PRÄFUSIONSPROTEINE

EP - 08.07.2020

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 19213895 Solicitante CALDER BIOSCIENCES INC  
Inventor/a MARSHALL CHRISTOPHER PATRICK

In some embodiments, the present invention provides respiratory syncytial virus (RSV) F proteins, polypeptides and protein complexes that comprise one or more cross-links to stabilize the protein, polypeptide or protein complex in its pre-fusion conformation. In some embodiments the present invention provides RSV F proteins, polypeptides and protein complexes comprising one or more mutations to facilitate such cross-linking. In some embodiments the present invention provides compositions comprising such proteins, polypeptides or protein complexes, including vaccine compositions, and methods of making and using the same.

43.[3677593](#)MODIFIZIERTES HSV-GB-PROTEIN UND HSV-IMPFSTOFF DAMIT

EP - 08.07.2020

Clasificación Internacional [C07K 14/035](#) Nº de solicitud 18851447 Solicitante KM BIOLOGICS CO LTD  
Inventor/a MORI HIROAKI

A modified protein of a herpes simplex virus (HSV) envelope glycoprotein B (gB), in which at least one non-neutralizing antibody-inducing epitope (non-neutralizing epitope) present in domain IV and domain I of wild-type HSV gB is inactivated (de-epitoped).

44.[WO/2020/135471](#)MONOCLONAL ANTIBODY AGAINST HUMAN INTERLEUKIN-4 RECEPTOR ALPHA AND USE THEREOF

WO - 02.07.2020

Clasificación Internacional [C07K 16/28](#) Nº de solicitud PCT/CN2019/128156 Solicitante QYUNS THERAPEUTICS CO., LTD. Inventor/a QIU, Jiwan

Provided are antibodies and fragments against human interleukin-4 receptor alpha (hIL-4Ra) and uses thereof. The antibodies and fragments preferably have heavy chain complementary determining regions as set forth in SEQ ID NOs: 1-3 or 14-16 and light chain complementary determining regions as set forth in SEQ ID NOs: 4-6 or 17-19.

45.[20200206331](#)AN IMMUNOGENIC COMPOSITION HAVING IMPROVED STABILITY, ENHANCED IMMUNOGENICITY AND REDUCED REACTOGENICITY AND PROCESS FOR PREPARATION THEREOF

US - 02.07.2020

Clasificación Internacional [A61K 39/00](#) Nº de solicitud 16631965 Solicitante SERUM INSTITUTE OF INDIA PVT LTD. Inventor/a Rakesh KUMAR

An immunogenic composition comprising of Diphtheria toxoid antigen (D), tetanus toxoid (T) antigen, Hepatitis B surface antigen (HBsAg), inactivated whole-cell *B. pertussis* (wP) antigen, *Haemophilus influenzae* type B (Hib) capsular saccharide conjugated to a carrier protein, Inactivated Polio Virus (IPV) antigen and additionally one or more antigens and the method of preparing the same. A fully liquid combination vaccine, showing improved immunogenicity, reduced reactogenicity and improved stability. Improved methods of formaldehyde inactivation, improved adsorption profile of Diphtheria toxoid antigen (D), tetanus toxoid (T) antigen and Hepatitis B (HepB) surface antigen adsorbed individually onto aluminium phosphate adjuvant, minimum total aluminum content ( $\text{Al}^{3+}$ ) and optimized concentration of 2-phenoxyethanol (2-PE) as preservative.

46.[3677592](#) MODIFIZIERTES HSV-GD-PROTEIN UND IMPFSTOFF DAMIT

EP - 08.07.2020

Clasificación Internacional [C07K 14/035](#) Nº de solicitud 18849655 Solicitante KM BIOLOGICS CO LTD Inventor/a MORI HIROAKI

The modified HSV gD protein of the present invention is a modified protein of a herpes simplex virus (HSV) envelope glycoprotein D (gD), wherein the modified HSV gD protein is derived from a wild-type HSV gD by modification of at least one of B cell epitopes having low or no neutralizing antibody-inducing activity compared to a B cell epitope present in a receptor-binding domain (RBD) (decotopes) in the ectodomain of the wild-type HSV gD, so that the modified epitope does not function as an epitope.

47.[WO/2020/136235](#) M2-DEFECTIVE POXVIRUS

WO - 02.07.2020

Clasificación Internacional [A61K 39/275](#) Nº de solicitud PCT/EP2019/087063 Solicitante TRANSGENE SA Inventor/a KLEINPETER, Patricia

The present invention is in the field of oncolytic viruses. The invention provides new poxviruses which are engineered to be defective for the function encoded by the M2L locus (i.e., m2 function). Such poxviruses lack a functional m2 binding activity to at least one or both of CD80 and CD86 co-stimulatory antigens. Said oncolytic poxviruses are preferably vaccinia virus having a total or partial deletion of the M2L locus. The present invention also relates to cells and compositions comprising such poxviruses and their use for treating proliferative diseases such as cancers and for preventing diseases (vaccination, especially in veterinary field). More precisely, the invention provides an alternative to the existing oncolytic viruses which are largely used in virotherapy. The m2-defective poxviruses are particularly useful for the expression of immunomodulatory polypeptides such as anti-CTLA-4 antibodies with the purposes of stimulating or improve immune response.

48.[3675903](#) PARAMYXOVIRIDAE-EXPRESSIONSSYSTEM

EP - 08.07.2020

Clasificación Internacional [A61K 39/155](#) Nº de solicitud 18769224 Solicitante BOEHRINGER INGELHEIM VETMEDICA GMBH Inventor/a NIKOLIN VELJKO

The present invention relates to the field of (vector) vaccines, and especially to an enhanced arrangement of nucleotide sequences for expressing a Paramyxoviridae virus containing an exogenous gene of interest. The

present invention further concerns related expression cassettes and vectors, which are suitable to express genes of interest, especially antigen encoding sequences. The viral vectors of the present invention are useful for producing an immunogenic composition or vaccine.

49. [WO/2020/132770](#) PARTÍCULAS TIPO-VIRUS (VLP) DEL VIRUS DE LA ANEMIA INFECCIOSA DEL SALMÓN (ISAV) COMPRENDIENDO LA PROTEÍNA DE MATRIZ Y UNA O MÁS PROTEÍNAS ANTIGÉNICAS SE DICHO VIRUS; MÉTODO DE OBTENCIÓN, COMPOSICIÓN, VACUNA Y ALIMENTO PARA PECES BACULOVIRUS RECOMBINANTE; Y KIT DE VACUNACIÓN

WO - 02.07.2020

Clasificación Internacional [A61K 39/145](#) Nº de solicitud PCT/CL2019/050152 Solicitante UNIVERSIDAD DE SANTIAGO DE CHILE Inventor/a CORTEZ SAN MARTIN, Marcelo

El presente invento se refiere al campo de la medicina veterinaria, particularmente, con vacunas y sanidad animal en el ámbito de la acuicultura. El presente invento se relaciona con partículas tipo-virus (VLP, Virus Like Particles), formadas por complejos moleculares que comprenden proteínas del virus de la anemia infecciosa de salmón (ISAV) cuyos genes se expresan en célula de insectos, y que son utilizadas como vacunas en conjunto con cuerpos celulares inducidos como adyuvantes. Es también del ámbito del presente invento, las composiciones farmacéuticas, vacunas, alimento para peces y kit que comprenden los VLP, como también el uso de los VLP para preparar medicamentos.

50. [2020203971](#) NOVEL PEPTIDES AND COMBINATION OF PEPTIDES FOR USE IN IMMUNOTHERAPY AGAINST EPITHELIAL OVARIAN CANCER AND OTHER CANCERS

AU - 02.07.2020

Clasificación Internacional [C07K 14/47](#) Nº de solicitud 2020203971 Solicitante Immatics Biotechnologies GmbH Inventor/a

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

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

Results of Search in US Patent Collection db for: (ABST/vaccine AND ISD/20200701->20200708),

9 resultados.

PAT. NO.	Title
1 <a href="#">10,703,795</a>	<a href="#">Peptides and combination of peptides for use in immunotherapy against esophageal cancer and other cancers</a>
2 <a href="#">10,703,777</a>	<a href="#">Miniature protein scaffolds and methods for use thereof</a>
3 <a href="#">10,702,596</a>	<a href="#">Polysaccharide purification for vaccine production using lytic enzymes, tangential flow filtration, and multimode chromatography</a>
4 <a href="#">10,702,595</a>	<a href="#">Manufacture of vaccines and compositions for the prevention of <i>Salmonella</i> infections</a>
5 <a href="#">10,702,594</a>	<a href="#">Dried saponin liposomal composition</a>
6 <a href="#">10,702,593</a>	<a href="#">Peptides and combination of peptides for use in immunotherapy against NHL and other cancers</a>
7 <a href="#">10,702,592</a>	<a href="#">Peptides and combination of peptides for use in immunotherapy against NHL and other cancers</a>
8 <a href="#">10,702,591</a>	<a href="#">Therapeutic cancer vaccine targeted to HAAH (aspartyl-[asparaginyl]-beta-hydroxylase</a>
9 <a href="#">10,702,553</a>	<a href="#">Peptides and combination of peptides of non-canonical origin for use in immunotherapy against different types of cancers</a>

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