

**FINLAY
EDICIONES**



BOLETÍN

VACCIENCIA

No. 12 (16 JUNIO - 22 JUNIO/2020)



...vacunar es prevenir.

Análisis bibliométrico sobre vacunas recombinantes

Fuente de información utilizada:



Estrategia de búsqueda:

TITLE: ("recombinant vaccine") 2613 records

Periodo de estudio 1999-2020

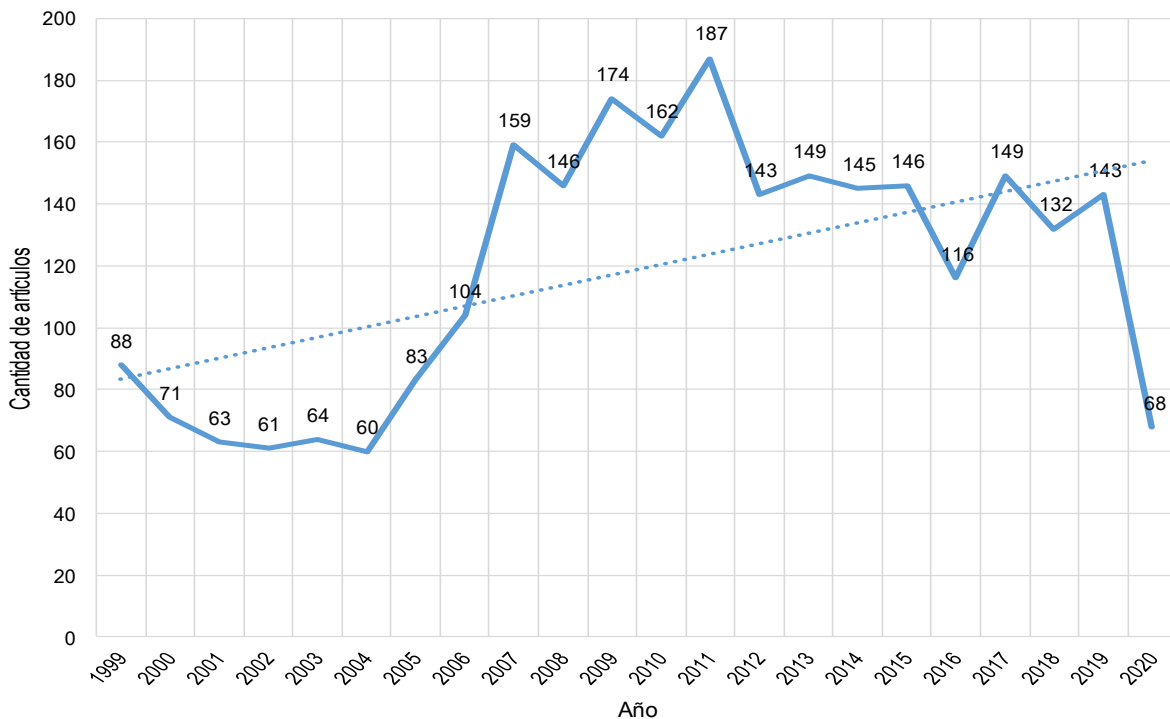
Las variables utilizadas en el análisis fueron:

- ⇒ 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.

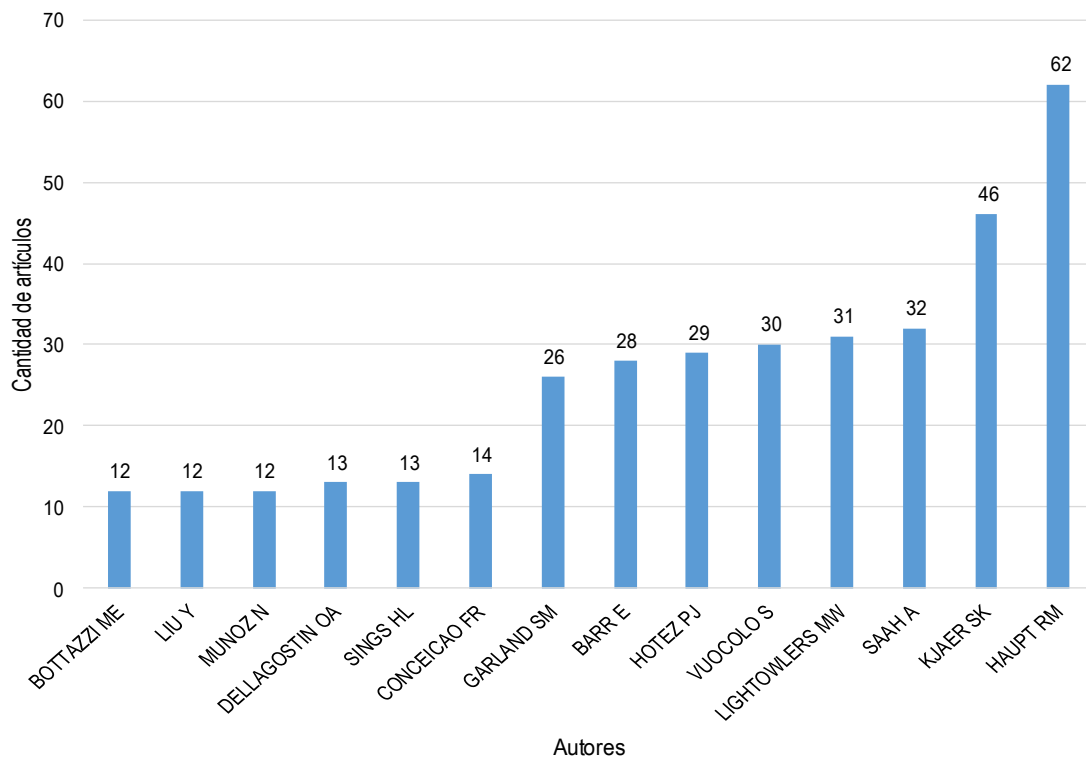
EN ESTE NÚMERO

- * Análisis bibliométrico vacunas recombinantes
- * Noticias en la Web sobre vacunas
- * Artículos científicos más recientes Medline sobre vacunas
- * Patentes más recientes PatentScope sobre vacunas
- * Patentes más recientes USPTO sobre vacunas

Productividad científica por año



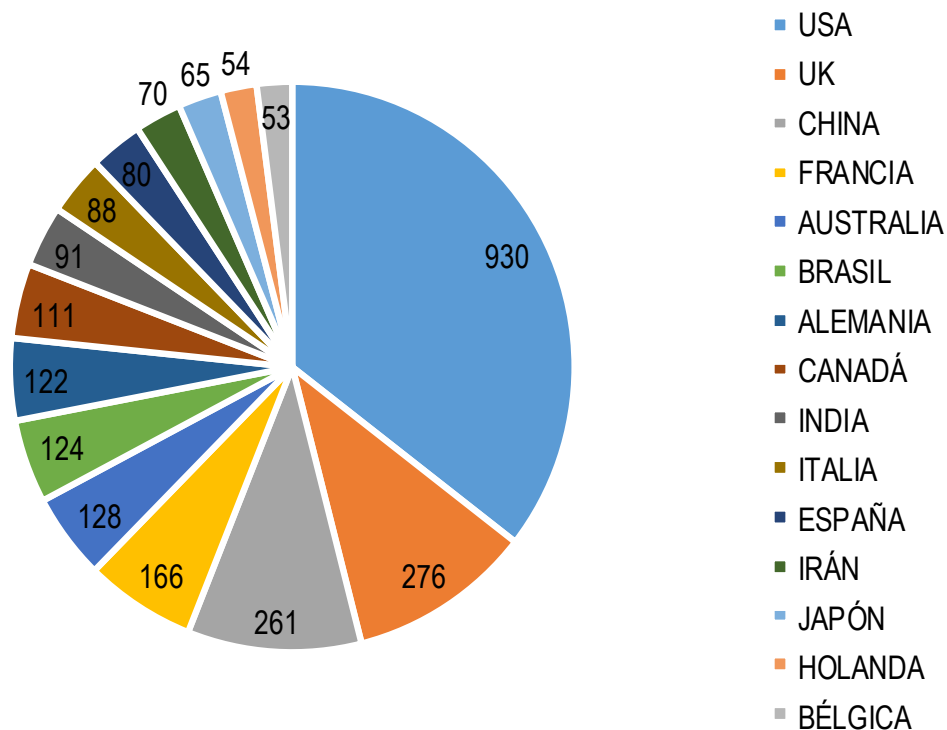
Autores con mayor productividad científica



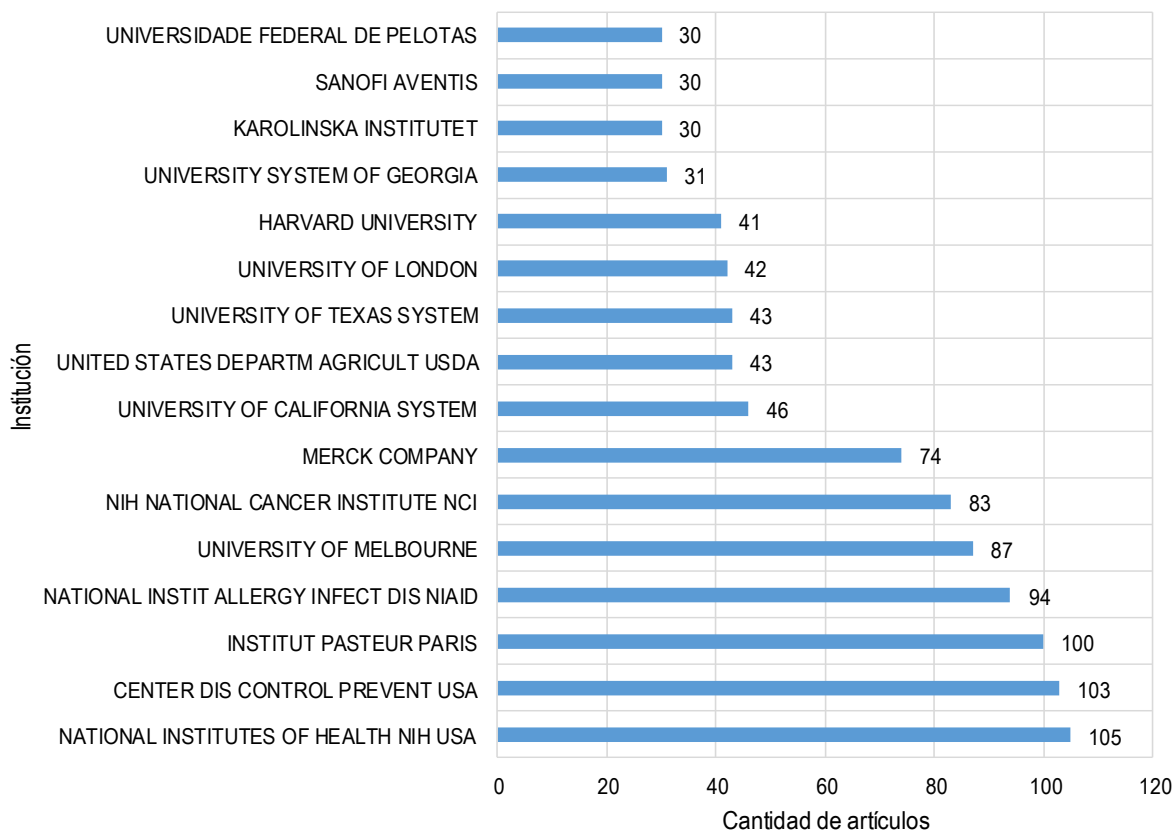
Revistas científicas que han publicado sobre el tema



Producción científica por países registrada en Web of Science



Instituciones que han trabajado el tema de estudio



Noticias en la Web

Cleveland Clinic develops coronavirus prediction model

16 jun. Cleveland Clinic researchers have developed one of the first risk prediction models that allows healthcare providers to predict a patient's likelihood of contracting COVID-19, and how likely the patient is to recover.

Risk for the coronavirus was reduced in those who had a pneumococcal polysaccharide or influenza vaccine, or were on melatonin, paroxetine, or carvedilol, according to Clinic research.

The research was recently published in the peer-reviewed medical journal *Chest*.

The Clinic's risk prediction model, called a nomogram, shows how age, race, gender, socioeconomic status, vaccination history and current medications affects a person's risk of contracting COVID-19. The risk prediction model provides a more scientific approach to COVID-19 testing, the Clinic said in a statement.

Clinic researchers developed the risk prediction model using

data from nearly 12,000 patients enrolled in the health system's COVID-19 registry before April 2.

The registry includes electronic medical record data from all patients tested at Clinic for the disease, whether they test positive or negative, the Clinic said.

"The ability to accurately predict whether or not a patient is likely to test positive for COVID-19, as well as potential outcomes including disease severity and hospitalization, will be paramount in effectively managing our resources and triaging care," Dr. Lara Jehi, the Clinic's chief research information officer, said in the statement.

Jehi is a corresponding author on the study. Michael Kattan, chair of Lerner Research Institute's Department of Quantitative Health Sciences, is a co-author.

The risk calculator is available here.

The risk calculator study found that patients who received the pneumococcal polysaccharide vaccine (PPSV23) and flu

vaccine were less likely to test positive for COVID-19 than those who didn't receive the vaccinations.

Pneumococcal polysaccharide vaccine (PPSV23) protects against 23 types of pneumococcal bacteria and helps prevent meningitis and bacteremia, according to the Centers for Disease Control and Prevention.

Those who take melatonin as an over-the-counter sleep aid, the heart medication carvedilol or the anti-depressant paroxetine are less likely to test positive than patients who don't take those medications, the study suggested.

Patients of low socioeconomic status (as measured in this study by zip code) are more likely to test positive than patients of a higher socioeconomic category, and Asian patients are less likely than white patients to test positive, the study said.

"Further validation and research are needed into these initial insights, but these correlations are extremely intriguing," Jehi said.

Fuente: Cleveland. Disponible en <https://bit.ly/38kFxyA>

Vacuna china para Covid-19 muestra resultados alentadores en pruebas en humanos

16 jun. China National Biotec Group (CNBG) dijo el martes que su vacuna experimental contra el coronavirus ha desencadenado anticuerpos en pruebas clínicas y que la

compañía planea realizar pruebas avanzadas en humanos en otros países.

No se ha demostrado fehacientemente que ninguna vacuna sea capaz de proteger eficazmente

contra el virus que ha matado a más de 400,000 personas, pero hay múltiples candidatas que se encuentran en diversas etapas de desarrollo en todo el mundo.

Se descubrió que la vacuna, desarrollada por un instituto de investigación con sede en Wuhan afiliado a la empresa matriz de CNBG, Sinopharm, había inducido la producción de anticuerpos en todas las personas inoculadas sin efectos secundarios adversos, según datos preliminares de una prueba clínica en la que participaron 1,120 personas sanas.

CNBG dijo que está buscando proactivamente oportunidades para pruebas de fase 3 en el extranjero en etapas tardías y a gran escala.

“Hemos garantizado la intención de cooperar con empresas e institutos de muchos países”, dijo la empresa en un comunicado.

Los medios de comunicación estatales informaron que la candidata a vacuna, junto con una inyección experimental diferente desarrollada por la unidad de Sinopharm, se ha ofrecido a empleados chinos de firmas estatales que viajan al extranjero ya que los desarrolladores buscan más datos sobre su eficacia.

China tiene cinco vacunas candidatas para Covid-19 en pruebas con humanos, la mayor cantidad en cualquier país.

El fabricante de vacunas chino Sinovac Biotech (Sinovac) publicó durante el fin de semana resultados preliminares positivos de su potencial candidato a vacuna, que se espera que sea probada en un ensayo de fase 3 en Brasil.

Fuente: Forbes México. Disponible en <https://bit.ly/2VCRnze>

Gran Bretaña inicia ensayos de posible vacuna de coronavirus

16 jun. Científicos del Imperial College de Londres empezarán a inmunizar a gente en Gran Bretaña esta semana con su vacuna experimental contra el coronavirus. En tanto, la farmacéutica Sanofi y el gobierno francés anunciaron una inversión de más de 800 millones de euros (890 millones de dólares) como parte de la carrera mundial para obtener una vacuna eficaz.

Según un comunicado del gobierno británico, 300 personas sanas recibirán dos dosis de la posible vacuna contra la COVID-19 desarrollada en el centro, que ha recibido 41 millones de libras (51 millones de dólares) de fondos gubernamentales.

Por el momento, la vacuna desarrollada por el Imperial College ha sido probada únicamente en animales y en el laboratorio, donde produjo



niveles de anticuerpos mucho más altos de los habituales en personas infectadas.

Muchos científicos han advertido que la pandemia sólo podría detenerse con una vacuna efectiva, que normalmente tarda años en desarrollarse.

El gobierno británico dijo en un comunicado que se inmunizará a 300 personas con dos dosis de la vacuna experimental contra la COVID-19 desarrollada por el

Imperial College, la que ha recibido 41 millones de libras (51 millones de dólares) en fondos oficiales.

Robin Shattock, que encabeza la investigación en el Imperial, dijo que la ventaja de su vacuna es que requiere una cantidad pequeña: la dosis es la centésima parte de la que ensayan en Estados Unidos los Institutos Nacionales de la Salud (NIH por sus siglas en inglés) y Moderna Inc. Esto significa que se podrían fabricar millones de dosis en

relativamente poco tiempo.

“Si el gobierno del Reino Unido quiere comprar suficientes vacunas para la población del país, ya tenemos montada la infraestructura para entregarlas en los primeros dos trimestres del año próximo”, dijo.

En tanto, Sanofi trabaja en una vacuna que espera ensayar en seres humanos en los próximos meses y obtener la aprobación el año entrante.

La empresa prometió el martes invertir 610 millones de euros (680 millones de dólares) en una planta de producción de vacunas y un centro de investigaciones en Francia para poder producir en mayor escala para futuros riesgos de pandemia”.

El presidente francés Emmanuel Macron visitó un laboratorio de Sanofi junto con el CEO de la empresa, Paul Hudson, y anunció que su gobierno invertirá 200 millones de euros para reducir la dependencia francesa de otros países en materia de vacunas y otros medicamentos.

Macron ha propuesto que se considere a las vacunas un “bien común” de la humanidad no sujeto a las presiones del mercado.

La vacuna del Imperial emplea filamentos sintéticos de código genético basados en el virus. Una vez inyectada en el músculo, las propias células del cuerpo reciben instrucciones para realizar copias de una proteína en el coronavirus. Esto debería desencadenar una respuesta inmune para que el cuerpo pueda combatir cualquier futura infección con COVID-19.

El doctor Doug Brown, presidente de la Sociedad Británica de Inmunología, dijo que en teoría la tecnología empleada por el Imperial College debería conducir a la inmunidad a largo plazo contra el coronavirus, pero que por ahora se la debe someter a pruebas rigurosas. Brown no participó de las pruebas.

La Universidad de Oxford inició recientemente un estudio avanzado con 10.000 voluntarios y Estados Unidos se apresta a realizar en julio estudios con 30.000 personas en quienes se pondrán a prueba diversas posibles vacunas, incluidas la de Oxford y la de los INH y Moderna Inc.

Los científicos nunca han creado tan rápidamente vacunas a partir de cero y dista de estar claro si alguna

de ellas resultará segura y efectiva. Con todo, Gran Bretaña, Francia, Holanda, Alemania, Estados Unidos y otros países han pedido millones de dosis por adelantado.

Shattock dijo que, en caso de ser eficaz, la licencia para fabricar la vacuna del Imperial no será otorgada a un laboratorio farmacéutico en particular, sino que varios socios en el mundo recibirán permiso para producirla como parte de un “negocio social” sin fines de lucro.

“No podemos proveerla sin costo porque su fabricación cuesta dinero”, dijo. “La proveeremos al costo con un pequeño porcentaje y ese pequeño porcentaje variará de acuerdo con la situación económica del país”.

La Organización Mundial de la Salud observó el lunes que se han reportado 100.000 infecciones nuevas cada día durante las últimas dos semanas y que la mitigación de las restricciones en muchos países ha provocado un nuevo pico.

Fuente: Lancaster Online. Disponible en <https://cutt.ly/loEgQuu>

Científico hondureño Salvador Moncada participa en hallazgo de nuevo tratamiento para COVID-19

16 jun. El científico hondureño Salvador Moncada, junto a un equipo investigador, han descubierto un nuevo tratamiento enfocado en proteger y fortalecer los pulmones para hacer frente al virus que provoca la COVID-19.

El sitio web STAT, especializado en noticias de salud, publicó el 10 de junio un artículo con entrevistas a varios científicos, entre ellos Moncada, para indicar el posible uso de la enzima ACE2 como tratamiento de la COVID-19.

El trabajo en la revista científica *Circulation* explica todo el mecanismo que usaron los investigadores para llegar a la conclusión del nuevo tratamiento.

El artículo indicó que cuando una persona se infecta del virus,

los síntomas más graves aparecen en los pulmones debido a que ingresa a través de ACE2.

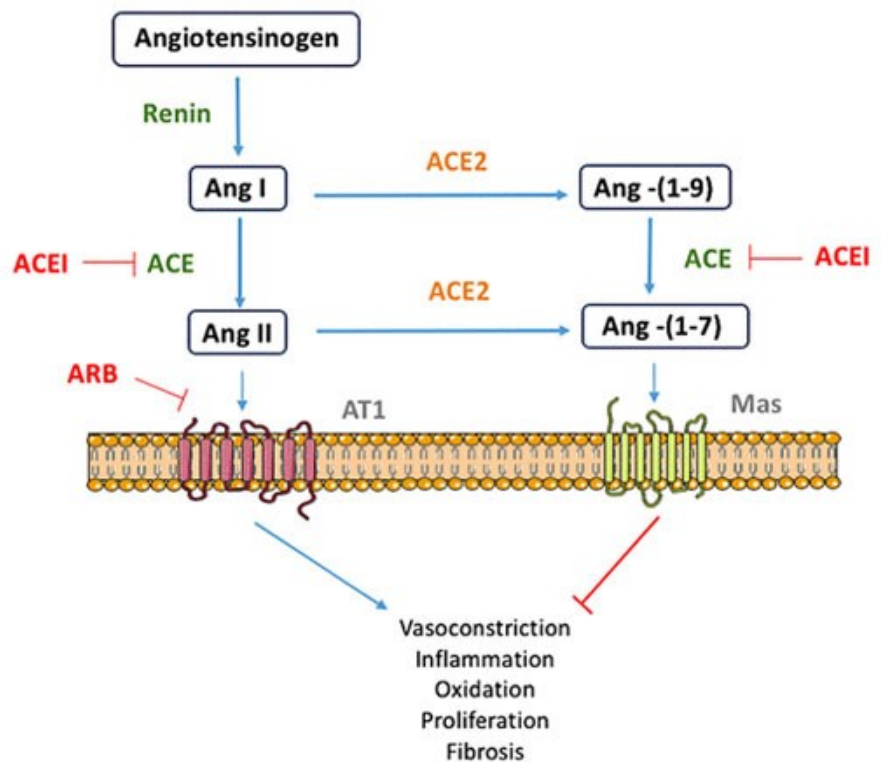
Los científicos presumen que la ACE2 podría indicar el camino hacia un tratamiento necesario para la pandemia debido que la enzima sirve como un árbitro en el sistema renina-angiotensina que sirve como regulador de la presión arterial.

“Cuando la angiotensina II, un péptido que constriñe los vasos sanguíneos, se eleva demasiado, ACE2 interviene y la convierte en angiotensina- (1-7), un galante en su Goofus que relaja los vasos y reduce la inflamación”, cita el artículo.

El científico hondureño, Salvador Moncada, señala en la publicación que "el sistema parece funcionar en un equilibrio en el que la angiotensina- (1-7) es protectora y la angiotensina II es el malo".

A criterio de la profesora de farmacología de la Universidad Autónoma de Madrid, Concepción Peiró, la angiotensina- (1-7) ha demostrado proteger contra la formación de coágulos sanguíneos, la oxidación y la muerte celular prematura.

La publicación menciona que Moncada y Peiró coescribieron una carta a la revista *Circulation* argumentando la hipótesis que la angiotensina- (1-7) puede proteger a los pulmones de los peores síntomas de la COVID-19.



Moncada recibió un correo electrónico del CEO de la compañía Constant Therapeutics, Rick Franklin, explicándole que desarrollaron el medicamento intravenoso “TXA127”, es una versión farmacéutica del péptido natural angiotensina- (1-7), que fue desarrollado como un tratamiento para enfermedades en las que el sistema renina-angiotensina no funcionaba.

Al momento de la declaración del COVID como pandemia, Rick Franklin vio el TXA127 podría ser un tratamiento para el huésped y no para el virus, dando tiempo a los pacientes al replicar los efectos de ACE2 según la publicación hasta que la enzima salga del coronavirus

"No fue ningún genio de nuestra parte reconstruir esta historia. Es solo que lo habíamos estado mi-

rando durante tanto tiempo que se hizo evidente para nosotros”, dijo Franklin.

La compañía Constant comenzó a escuchar de centros académicos de todo el mundo con la esperanza de estudiar la droga en el virus.

El primer ensayo patrocinado por la Universidad de Columbia, está programado para comenzar este mes inscribiendo a 100 pacientes con COVID-19 moderado y comparando una dosis diaria de TXA127 con placebo.

Los objetivos principales son la seguridad y la prevención de la insuficiencia pulmonar, con medidas secundarias de supervivencia, inflamación y la necesidad de asistencia respiratoria.

La compañía mapea un ensayo de fase 2 propio que involucraría aproximadamente siete sitios e

inscribiría a más de 200 pacientes con el virus pero que aún no requieren cuidados intensivos.

Salvador Moncada menciona en el artículo que tiene “sentido biológico” como tratamiento de COVID-19, el TXA127 en ensayos clínicos puede ser un arduo proceso.

“Un porcentaje significativo de personas que contraen la enfermedad se recuperan sin desarrollar síntomas pulmonares graves. Eso significa que, para demostrar el potencial de TXA127, Constant necesitará inscribir a un gran número de pacientes o descubrir

cómo detectar los casos de COVID-19 que probablemente se beneficien del medicamento”, comentó.

El enfoque de la compañía parece prometedor, dijo Ankit Patel, nefrólogo del Hospital Brigham and Women en Boston. “Pero ACE2 hace más que solo crear angiotensina- (1-7), y replicar sus efectos beneficiosos puede requerir más que simplemente infundir más péptido”, dijo.

“Definitivamente parece que existe esta angiotensina II sin oposición en COVID-19, y tratar de revertirla parece ser una

buena técnica para abordar parte de la patología pulmonar que vemos”, añadió Patel.

El ceo de Constant considera un diseño de prueba que reclutaría pacientes que necesitan oxígeno pero que aún no están en la UCI, apostando que TXA127 puede diferenciarse del placebo cuando se trata de prevenir el daño pulmonar.

El papel de ACE2 en COVID-19 es ampliamente aceptado, y el potencial de la angiotensina- (1-7) es claro para los expertos de todo el mundo, expresó Franklin.

Fuente: Proceso Digital. Disponible en <https://cutt.ly/VoEiEXV>

Pneumonia vaccine price drops dramatically for lower-income countries thanks to the Gavi pneumococcal Advance Market Commitment

16 jun. Lower-income countries across the world will now be able to access life-saving pneumococcal conjugate vaccines, which protect against the leading cause of pneumonia, for US\$ 2 per dose, thanks to a new supply agreement between UNICEF, Gavi’s procurement partner, and the Serum Institute of India (SII). The new price represents a 43 per cent reduction from the Gavi price of US\$3.50 at the start of the Advance Market Commitment (AMC). The supply agreement is the eighth to take place under the Vaccine Alliance’s Advance Market Commitment (AMC) mechanism, and the first to



include a developing country manufacturer. Under the agreement, the Indian manufacturer will provide ten million PCV doses to Gavi-supported countries each year for the next ten years. The

pneumococcus bacterium is the leading cause of severe pneumonia and is a major cause of morbidity and mortality worldwide. Most of these deaths occur in lower-income countries and include a disproportionate

number of children under the age of two.

The AMC, which is set to end this year, was launched by Gavi in 2009 – with the support of donors Italy, the United Kingdom, Canada, Russia, Norway and the Bill & Melinda Gates Foundation – to solve a clear example of market failure: complex vaccines like PCV would normally reach low-income countries, where the disease burden is often highest, ten to 15 years after their introduction in industrialised countries. The AMC mechanism incentivised research and development (R&D), particularly for vaccines suitable for developing country epidemiological contexts, and an expansion in manufacturing capacity to meet demand from lower-income countries while maintaining an affordable price per dose.

A decade later the results are clear: the PCV vaccine has now been introduced in 60 lower-income countries, where coverage rates, at 48%, are now higher than the global average of 47%. Estimates indicate that more than 225 million children will have been vaccinated, and that over 700,000 deaths will have been prevented by the end of 2020. In that time, the AMC has catalysed a steady decrease in PCV prices and thanks to this eighth supply agreement will be closing this year

having facilitated the entry of a new manufacturer to the market as well as a record-setting low price for Gavi-supported countries that will result in an estimated millions of dollars in savings for both Gavi and lower income countries' vaccine budgets.

“This is fantastic news, which will make this highly effective, lifesaving vaccine even more affordable for the world’s poorest countries, and is a testament to the success of Gavi’s Pneumococcal AMC mechanism, which as now achieved everything it set out to do,” said Aurelia Nguyen, Gavi’s Managing Director for Vaccines and Sustainability. “Thanks to this visionary model we have a healthy PCV market that is producing enough vaccines to supply both rich and poor countries and, as a result, hundreds of millions of children are now protected against one of the world’s deadliest diseases.”

“Pneumonia is the biggest single killer of children, claiming the life of a child every 39 seconds. By being able to provide this quality-assured pneumococcal conjugate vaccine at an such an affordable price, we can save millions of children’s lives,” said Etleva Kadilli, Director of UNICEF’s supply and procurement headquarters. “Our innovative approach has incentivized industry to bring this vaccine at

scale to the market. The result means more countries – including middle-income countries – that have not yet introduced this vaccine into their routine immunization due to previously prohibitive pricing, will have an affordable option to accelerate access for all children.”

“It is incredible to see that Gavi’s commitment to developing innovative mechanisms – and our donors’ commitment to supporting them – has had such a dramatic and positive impact on the lives of the most vulnerable children around the world,” said Marie-Ange Saraka-Yao, Gavi’s Managing Director for Resource Mobilisation, Private Sector Partnerships & Innovative Finance. “This successful mechanism will now be used as a model to help ensure the world’s poorest countries get access to COVID-19 vaccines at the same time as the wealthiest, making it integral to the global effort to defeat this pandemic.”

At last week’s Global Vaccine Summit, Gavi launched the Gavi Advance Market Commitment for COVID-19 Vaccines (Gavi Covax AMC), a new financing instrument aimed at incentivising vaccine manufacturers to produce sufficient quantities of eventual COVID-19 vaccines, and to ensure access for developing countries.

Singapore to start human trials of Covid-19 vaccine in August

16 jun. Singapore scientists testing a Covid-19 vaccine from US firm Arcturus Therapeutics plan to start human trials in August after promising initial responses in mice. More than 100 vaccines are being developed globally, including several already in human trials from the likes of AstraZeneca and Pfizer, to try and control a disease that has infected more than 8 million people and killed over 430,000 worldwide.

The vaccine being evaluated by Singapore's Duke-NUS Medical School works on the relatively-untested Messenger RNA (mRNA) technology, which instructs human cells to make specific coronavirus proteins that produce an immune response.

"The fact that it replicates and triggers a very balanced immune response, both in terms of the antibody and killer cells - those are welcome properties," Ooi Eng

Eong, deputy director of the school's emerging infectious diseases programme, told Reuters on Tuesday.

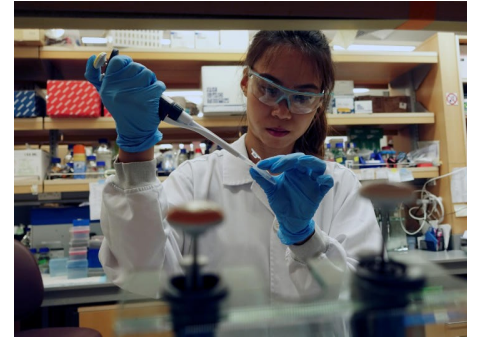
Antibodies stick to the virus and prevent it from infecting cells, while killer cells, another arm of the immune system, recognise infected cells and destroy them, he said.

The mRNA approach has not yet been approved for any medicine so its backers, which also include US biotech firm Moderna, are treading uncharted territory.

Because of that, Ooi said longer studies were needed to ensure its safety.

"The most optimistic case is that it's about this time next year, that we will have a vaccine," Ooi said.

Ooi is also working on a monoclonal antibody treatment for Covid-19 and will begin safety trials on healthy people this week, before testing on Covid-19 patients in the coming months.



Ooi said potential deployment of the treatment could be faster than the vaccine, without giving an exact timeline.

Antibodies are generated in the body to fight off infection. Monoclonal antibodies mimic natural antibodies and can be isolated and manufactured in large quantities to treat diseases.

Tiny city-state Singapore has one of the highest infection tallies in Asia, with more than 40,000 cases, largely due to mass outbreaks in dormitories for its migrant workers.

Fuente: Bangkok Post. Disponible en <https://cutt.ly/NoEQaol>

Identifican anticuerpos neutralizantes contra el SARS-CoV-2

16 jun. Tres nuevos estudios describen varios anticuerpos humanos recientemente descubiertos que se dirigen al virus SARS-CoV-2, aislado de los sobrevivientes de infección por SARS-CoV-2 y SARS-CoV.

Varios de estos anticuerpos mostraron capacidades protectoras y neutralizantes, ofreciendo pistas terapéuticas prometedoras,

y se descubrió que ocho anticuerpos de un análisis reaccionan de forma cruzada con un coronavirus específico de murciélago relacionado, con implicaciones para la identificación de anticuerpos ampliamente neutralizantes para proteger contra un posible coronavirus nuevo brotes en el futuro.

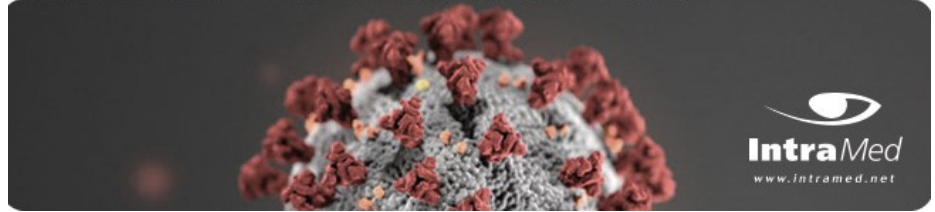
Philip Brouwer y sus colegas aislaron 403 anticuerpos

monoclonales de 3 pacientes convalecientes de COVID-19, lo que demuestra que los pacientes tenían fuertes respuestas inmunes contra el pico viral, un complejo de proteínas que se une al receptor ACE2 para facilitar la entrada en las células huésped humanas.

Un subconjunto de estos anticuerpos neutralizó el virus

atacando diversos epítomos en la espiga, con los dos más potentes dirigidos al dominio que une el receptor del huésped. En otro estudio, Thomas Rogers y sus colegas utilizaron una tubería de alto rendimiento para aislar y caracterizar los anticuerpos monoclonales de donantes convalescentes, seleccionando los anticuerpos que se unen al pico viral. Varios de los anticuerpos aislados se unieron al dominio de unión al receptor (RBD) y demostraron capacidades neutralizantes, y los más potentes se unieron en un sitio que se superpone al sitio de unión de ACE2. Dos de estos anticuerpos neutralizantes dieron protección contra la infección por SARS-CoV-2 cuando se probaron en hámsters sirios.

COVID-19 en IntraMed



En un tercer estudio para identificar anticuerpos ampliamente reactivos de reacción cruzada, Anna Wec y sus colegas aislaron y caracterizaron cientos de anticuerpos contra el pico viral de SARS-CoV-2 de las células B de memoria de un sobreviviente de SARS-CoV.

Ambos virus estrechamente relacionados dependen de la espiga para obtener la entrada de la célula huésped al unirse al receptor ACE2. De nueve anticuerpos que mostraron una fuerte neutralización cruzada de ambos virus, ocho

se dirigen al dominio que se une al receptor ACE2, y también neutralizaron una especie estrechamente relacionada de coronavirus de murciélago.

Tomados en conjunto, el trío de estudios ofrece varios anticuerpos humanos nuevos para ayudar a informar el diseño de medicamentos terapéuticos y vacunas contra el SARS-CoV-2, así como el diseño de vacunas ampliamente protectoras contra una variedad de coronavirus relacionados.

Fuente: IntraMed. Disponible en <https://cutt.ly/voESIW>

Médicos ecuatorianos tienen criterios diferentes sobre uso de dexametasona

17 jun. En contagiados muy graves, la nueva cepa de coronavirus produce una inflamación total del cuerpo, conocida como tormenta de citoquinas. Esta reacción puede provocar la muerte. Frente a eso, científicos probaron con dosis de dexametasona, para combatir este tipo de reacción. El fármaco fue estudiado por un grupo de investigadores de la Universidad de Oxford, en Reino Unido. Concluyeron que actúa como unas hormonas que sirven para desinflamar y que están presentes en el cuerpo humano.



El estudio se hizo con 2 000 pacientes en estado grave. Los primeros resultados dieron cuenta de

que el medicamento redujo las muertes en el 35%; mientras que la mortalidad bajó a la quinta parte

entre quienes recibían oxígeno. “Creo que en Ecuador debemos ser pacientes, esperar y no precipitarnos. Hay que recordar lo ocurrido con la hidroxiclороquina; ha provocado más revisiones de pares”, comentó Edgar Samaniego, autor de un libro sobre Fundamentos de la Farmacología y quien por décadas dio esa cátedra en la Universidad Central. Además, el médico y exrector, reiteró que hay que pedirle a la población “no cometer la barbaridad de probar este medicamento antiguo y muy barato. Los investigadores de Oxford advirtieron que lo aplican únicamente con quienes están muy graves, hospitalizados”. La dexametasona se usa contra enfermedades autoinmunes, para evitar rechazo de trasplante de

órganos, artritis reumatoidea y procesos alérgicos agudos, entre otras, indicó Edgar Samaniego. Entre otros efectos indeseables, dijo, puede producir, retención de líquidos, es decir edema; alteraciones de distribución de la grasa, que causan la cara de luna llena o hinchazón. Descarta que provoquen a los tratados por covid-19 osteoporosis o insuficiencia suprarrenal aguda pues eso pasa cuando se usa a largo plazo. En Oxford lo utilizan por 10 días. Por otro lado, Enrique Terán, médico especialista en farmacología, sí utilizaría este medicamento que pertenece al grupo de los glucocorticoides solo en pacientes que estén en condición de gravedad, en cuidados intensivos. Lo haría con “mucho cuidado”, ya que es importante disminuir la inflamación producto del

SARS-CoV-2. Pero -alerta- los médicos deben monitorear constantemente al paciente para descartar infecciones bacterianas. Esto debido a que el medicamento produce inmunosupresión, es decir, una baja de las defensas, por lo que la persona puede contagiarse con otros microorganismos que le pueden provocar la muerte. Una de las más comunes es la pseudomona, una bacteria intrahospitalaria. En el Consenso Interino Multidisciplinario Informado en la Evidencia sobre el Tratamiento de covid-19, que se actualizó en mayo y estuvo a cargo de gremios médicos y el Ministerio de Salud, no se indica el uso de estos fármacos para cualquier paciente, ya que aún no se ha probado su efectividad para neumonía grave.

Fuente: EL COMERCIO. Disponible en <https://cutt.ly/noRyht1>

La dexametasona es prometedora contra la COVID-19, pero solo para pacientes muy concretos

18 jun. Esta semana, la Universidad de Oxford (Reino Unido) aseguró mediante una nota de prensa que la dexametasona reduce el riesgo de muerte en pacientes de COVID-19 con complicaciones respiratorias. Aunque todavía no se ha publicado estudio alguno, el anuncio despertó expectación por formar parte del ensayo RECOVERY, cuyo objetivo es evaluar la eficacia de diferentes fármacos contra el coronavirus.

La dexametasona pertenece a la familia de los corticoides (o corticosteroides), que tienen propiedades antiinflamatorias, inmuno-

supresoras y antialérgicas. Estos fármacos son esenciales en el manejo de patologías como el asma, la enfermedad pulmonar obstructiva crónica (EPOC), el distrés respiratorio agudo, las alergias broncopulmonares, nasales, cutáneas y oculares. También se usan para los choques anafilácticos, la artritis reumatoide, la esclerosis múltiple, el edema cerebral e incluso reducir los efectos secundarios de la quimioterapia, entre otros trastornos.

La historia de los corticoides se remonta al año 1843, cuando el médico inglés Thomas Addison

describió por primera vez algunos casos de insuficiencia suprarrenal en pacientes que presentaban “un estado general de languidez y debilidad, desfallecimiento en la acción del corazón, irritabilidad en el estómago y un cambio peculiar en la piel”. Estos, inevitablemente, fallecían.

Posteriormente, en 1935, Edward Calvin Kendall y sus colaboradores de la Clínica Mayo (EE. UU.) aislaron, a partir de glándulas suprarrenales de buey, seis sustancias desconocidas. Una de ellas fue denominada como “compuesto E”: para no confundirla con la vitamina E, cambiaron su nombre por

“cortisona”. Este fue el primer corticoide de origen natural.

El descubrimiento les valió el Premio Nobel de Fisiología o Medicina en 1950. Durante los años siguientes se obtuvieron otros compuestos sintéticos, derivados de la cortisona, mediante cambios en su estructura química. Así se mejoraron sus propiedades farmacocinéticas y, sobre todo, su potencia.

¿Qué propiedades aporta la dexametasona?

Uno de estos corticoides sintéticos es la dexametasona. Esta tiene una gran potencia, pero sin muchos de los efectos adversos de corticoides naturales como la cortisona. Es muy liposoluble, lo que aumenta su absorción en el tubo digestivo y su penetración en los tejidos, lo que mejora su eficacia terapéutica. Por todo ello, supuso una novedad en su momento.

La dexametasona actúa como un potente antiinflamatorio e inmunosupresor: disminuye o elimina la respuesta de los tejidos a la inflamación. Sin embargo, aunque reduce los síntomas asociados a este proceso, no trata la causa que lo genera.

En otras palabras, este fármaco impide la acumulación de células inflamatorias como macrófagos y leucocitos, la fagocitosis, la liberación de enzimas lisosomales y de mediadores de la inflamación. El tiempo de acción es prolongado y su efecto es 7,5 veces superior al de otros corticoides como la prednisona y la prednisolona, y 30 veces mayor que el de la hidrocortisona.

COVID-19: la tormenta perfecta

La inflamación es un mecanismo que se desencadena ante una amenaza, infecciosa o no, y cuya finalidad es mantener la homeostasis de nuestro cuerpo. Sin embargo, es necesario que esta respuesta sea regulada de forma precisa, tanto en intensidad como en duración, para que sea beneficiosa.

En caso contrario puede surgir el “síndrome de liberación de citoquinas”. Esta “tormenta de citoquinas” es causada por una respuesta inflamatoria sistémica aguda, mediada por unas sustancias naturales proinflamatorias que fabrica nuestro organismo; las citoquinas. Puede desencadenarse por una amplia variedad de factores, como infecciones y reacciones a algunos medicamentos.

En pacientes afectados por la COVID-19, cuando la respuesta del sistema inmune no es capaz de controlar eficazmente al coronavirus, como puede suceder en personas mayores, el virus se propaga de forma más agresiva. Esto produce daño en los tejidos pulmonares, lo que activa a los macrófagos y granulocitos y conduce a la liberación masiva de citoquinas proinflamatorias.

Todo este proceso inflamatorio puede complicarse, lo que da lugar a la “tormenta de citoquinas” observada de forma frecuente en pacientes graves de COVID-19. También aumentan los neutrófilos y se ve reducido el número de linfocitos totales. Otros marcadores inflamatorios también han sido detectados en sangre a niveles ele-

vados en pacientes de COVID-19.

¿Por qué los corticoides?

Dada su potencia antiinflamatoria, los corticoides son muy utilizados para el tratamiento de la COVID-19 en sus estadios más graves. Pero no hay que olvidar que suprimen el funcionamiento del sistema inmunológico, por lo que no se pueden emplear en las etapas iniciales de la enfermedad: solo son útiles en la etapa inflamatoria.

Así, la desametaxona se usa en estos pacientes para detener parte del daño producido cuando el sistema inmunológico se sobreactiva en esta tormenta de citoquinas, mientras el organismo intenta luchar contra el coronavirus.

El uso de corticoides en pacientes COVID-19 positivos ya se ha planteado en protocolos de diferentes países, incluida España, sobre todo en pacientes adultos graves ingresados en UCI. Sin embargo, aún no se ha estandarizado una dosis ni una pauta terapéutica concreta. De forma habitual se utiliza en dosis bajas y durante cortos periodos de tiempo, con el objetivo de minimizar el riesgo de efectos adversos.

Aunque la evidencia científica es escasa, los primeros datos aportados por la Universidad de Oxford son muy alentadores. Parecen confirmar que la dexametasona, en dosis de 6 mg una vez al día durante diez días, reduce la mortalidad en un tercio de los pacientes que necesitaron ventilación artificial y en un quinto de los pacientes que recibieron oxígeno.

Sin embargo, no se han apreciado beneficios entre aquellos pacientes

que no requirieron asistencia respiratoria. Según estos resultados, la dexametasona podría evitar una muerte de cada ocho pacientes tratados que requieran ventilación, y una muerte de cada 25 entre aquellos que reciben oxígeno.

Aunque se trata de datos preliminares, los investigadores trabajan en la obtención de los datos definitivos de este ensayo en el menor tiempo posible.

España también investiga los corticoides

Los datos aportados, aunque provisionales y sin publicar, son muy

prometedores, tal como han comunicado recientemente responsables de la Organización Mundial de la Salud. Parecen confirmar una notable reducción de la mortalidad y de las complicaciones asociadas a la infección.

También en España se investiga el uso de corticoides en ensayos clínicos ya autorizados y en fase de reclutamiento de pacientes en estadios avanzados.

La dexametasona puede convertirse en el estándar de atención en este grupo concreto de pacientes de COVID-19 pues, además, es muy económica, su disponibilidad

es absoluta y se puede usar de inmediato para salvar vidas en todo el mundo.

Pero no se debe confundir el hecho de que este fármaco pueda ser prometedor para pacientes concretos con su uso generalizado en todos ellos, ni para tratar o prevenir esta enfermedad. El empleo de corticoides sin control médico es sumamente peligroso, debido al desarrollo de efectos adversos graves, como la inmunosupresión y la insuficiencia adrenal aguda por la supresión brusca del tratamiento.

Fuente: HUFF POST. Disponible en <https://cutt.ly/QoRo3K7>

El enigma de la vacuna contra el coronavirus: fechas que ya barajan la OMS, China y Europa

20 jun. La llegada de la vacuna contra el coronavirus que asola al mundo es la noticia esperada por todos. La única solución sanitaria real a una pandemia que deja millones de infectados y que se ha cobrado cientos de miles de vidas humanas en todo el mundo. Dar una respuesta a cuándo llegará comienza a ser una obsesión para la que no hay una respuesta concreta, aunque sí cada vez más clara.

La vacuna que China está desarrollando para hacer frente a la Covid-19 podría tardar más de un año en estar disponible para la venta debido a la falta de nuevos infectados con los que realizar ensayos clínicos, tal y como ha informado la agencia de noticias CNS.

Actualmente se están realizando más de una decena de ensayos

clínicos de posibles vacunas en diversos países, pero ninguna de ellas ha pasado aún a la tercera fase de pruebas, que requiere la participación de miles de personas para garantizar su efectividad.

China, donde se detectó el virus por vez primera a finales de 2019, registró menos de diez casos de coronavirus al día a lo largo del mes de mayo, lo que ha dificultado los ensayos.

"Esperamos que podamos cooperar a nivel internacional y realizar una fase tres a nivel clínico de forma múltiple para lograr así que la vacuna llegue al mercado", ha explicado el vicepresidente del Grupo Nacional de Biotecnología Chino (CNBG), Zhang Yutao.

"La vacuna no llegará al mercado al menos hasta el año que viene tal y como avanzan las investigaciones", ha aseverado en una

entrevista a la citada agencia. Un nuevo brote ha dejado numerosos infectados en Pekín, la capital china, si bien Yang ha aseverado que el número de pacientes en comparación con la densidad poblacional es demasiado bajo como para propiciar un "ambiente de ensayo ideal".

La previsión de la OMS, algo más optimista

La Organización Mundial de la Salud (OMS) ha informado por su parte de que espera que a finales de año ya estén disponibles "millones de dosis" de vacunas y que en 2021 estén disponibles otros 2.000 millones.

Actualmente hay unas 300 vacunas en ensayos y tres ya están cerca de comenzar la fase final de las pruebas con personas, la de la Universidad de Oxford, la vacuna RNA de la compañía

Moderna y otra más que se está desarrollando en China.

No obstante, la jefa de científicos de la OMS, Soumya Swaminathan, ha avisado de que esta hipótesis "no son certeras" ya depende del resultado final de los ensayos, si bien ha recordado que el organismo de Naciones Unidas trabaja con esos supuestos para la adquisición, distribución y reparto justo.

"Tengo esperanza, soy optimista, pero el desarrollo de una vacuna es una tarea compleja, viene con mucha incertidumbre. Lo bueno de esto es que tenemos muchas vacunas y plataformas para que, incluso si la primera o la segunda falla no perdamos la esperanza y no nos rindamos. Si tenemos suerte habrá una o dos vacunas exitosas a finales de año", ha detallado la experta.

La estrategia de la UE con la vacuna

La Comisión Europea ha desvelado esta semana su estrategia para garantizar en todo el bloque vacunas "seguras,

eficaces y de calidad" para la Covid-19 en un plazo de entre 12 y 18 meses y que estará basada en un sistema centralizado de compras anticipadas a fabricantes con capacidad de producción en la UE.

La comisaria de Salud, Stella Kyriakides, ya trasladó las características principales de esta estrategia a los ministros de Sanidad de los Veintisiete en la reunión telemática que tuvo lugar hace una semana. El ministro español, Salvador Illa, mostró horas después su apoyo a la iniciativa.

La estrategia persigue un triple objetivo: garantizar vacunas seguras, eficaces y de calidad", asegurar un "acceso rápido" a las mismas por parte de los Estados miembros y sus poblaciones y conseguir que además sea "justo y asequible".

El plan de Bruselas se basa además en dos pilares y el primero de ellos pasa por adaptar el marco legislativo vigente a la "urgencia actual" y utilizar toda la flexibilidad posible para "acelerar el desarrollo, la autorización y la disponibilidad de vacunas" mientras se

mantienen los estándares de calidad y seguridad.

El segundo pilar de la estrategia representa el núcleo de la misma y persigue garantizar la producción de las futuras vacunas en el bloque y un "suministro suficiente" para todos los socios comunitarios. El Ejecutivo comunitario sugiere hacer a través de Acuerdos de Compra Anticipados (APA, por sus siglas en inglés).

En virtud de este mecanismo, la Comisión negociará y cerrará acuerdos con fabricantes de vacunas en nombre de los Estados miembros. Bruselas financiará con 2.400 millones de euros parte de los costes en los que incurran estas compañías a través del presupuesto del Instrumento de Ayuda de Emergencia (ESI).

Estos pagos se concebirán como un anticipo por la compra de dosis de vacunas que posteriormente tendrán que pagar los países de la UE como los productores que hayan sido seleccionados por el Ejecutivo comunitario cuando exista una vacuna.

Fuente: 20 minutos. Disponible en <https://cutt.ly/joRBcVd>

Estudian la destrucción del material genético del SARS-COV-2 mediante la herramienta CRISPR

20 jun. El profesor de la Universidad Pablo de Olavide, Miguel Ángel Moreno Mateos, investigador Ramón y Cajal en el Centro Andaluz de Biología del Desarrollo (CABD), co-lidera un proyecto de investigación financiado por el Consejo Superior de Investigaciones Científicas (España) cuyo objetivo es destruir el

genoma del coronavirus SARS-CoV-2 empleando la herramienta de edición genética de última generación CRISPR-Cas13d.

El SARS-CoV-2 es un virus cuyo genoma está formado por una única cadena de ácido ribonucleico (ARN), una molécula que, al igual que el ácido desoxirribonucleico (ADN), es esencial para la vida de

los organismos. Está repleto de instrucciones genéticas para hacer millones de copias de sí mismo y estas instrucciones están codificadas en 30.000 'letras' de ARN que la célula infectada lee y traduce a las distintas proteínas virales que promueven la replicación viral.

"La tecnología CRISPR-Cas es

una revolución comparable a la acontecida en los años 70 y 80 con el DNA recombinante y los inicios de la ingeniería genética. Básicamente, ahora podemos ir de manera dirigida a cualquier lugar de un genoma (de ADN o de ARN como es el del SARS-CoV-2) y manipularlo a nuestro antojo”, afirma el profesor Miguel Ángel Moreno, experto en esta tecnología. Esta herramienta tiene muchas variantes y en este proyecto el equipo de investigación emplea la proteína Cas13d, que, como explica el investigador de la UPO, “es capaz de encaminarse de manera precisa y dirigida al ARN viral, lo que es el cerebro del virus de donde salen todas sus instrucciones, y cortarlo, como si fuera unas tijeras, llevándolo a su eliminación y evitando su propagación”.

El investigador de la UPO lidera la primera fase del proyecto, que consiste en la selección de las formulaciones CRISPR-Cas13d más eficientes para el SARS-CoV-2 y otros virus de RNA relacionados con éste, empleando embriones de pez cebra como sistema modelo *in vivo*. “Somos pioneros en la optimización de la tecnología CRISPR-Cas13 *in vivo*, y lo que estamos haciendo es preparar las formulaciones más eficientes con unos modelos artificiales, usando determinados RNAs del SARS-CoV-2 y otros virus similares pero de forma independiente, que no

generan ningún peligro ni necesitan un laboratorio de alta seguridad. Estas formulaciones se probarán después en modelos reales”, explica Miguel Ángel Moreno.

Tras evaluar su funcionalidad y su no toxicidad, estos reactivos se testarán en el Centro Nacional de Biotecnología. Las formulaciones más eficientes serán entonces probadas en modelos celulares de infección *in vitro* donde se podrá analizar la capacidad de eliminación por parte de dichas formulaciones de virus similares al SARS-CoV-2 pero menos peligrosos. De este modo, se optimizarán las condiciones de eliminación antiviral con las formulaciones más eficientes previamente probadas. Finalmente, en un laboratorio con la bioseguridad adecuada, se probarán en modelos de infección de SARS-CoV-2 *in vitro* e *in vivo* usando todas las optimizaciones anteriores.

“Podríamos tener una terapia muy flexible y podría, por ejemplo, ser adaptada fácilmente a distintas versiones mutantes del SARS-CoV-2 y otros virus de RNA de forma relativamente sencilla, aunque debemos tener cautela porque estamos empezando. Como se suele decir, la paciencia es la madre de la ciencia”, señala Moreno.

El proyecto agrupa a investigadores de perfiles diversos pero complementarios: Miguel Ángel Moreno Mateos (UPO-CABD),

biólogo del desarrollo y experto en la optimización de sistemas CRISPR-Cas *in vivo*; Dolores Rodríguez (CNB-CSIC), viróloga experta en el manejo y caracterización de diferentes tipos de virus; y Almudena Fernández (CIBER-ISCIII) y Lluís Montoliu (CNB-CSIC), genetistas, expertos en el uso de las herramientas CRISPR de edición genética para la generación de modelos animales de enfermedades raras.

“Contar con un equipo multidisciplinar para este proyecto ha sido crítico. La ciencia sin una colaboración fluida no va a ningún sitio”, afirma Miguel Ángel Moreno Mateos. Actualmente las colaboraciones y el equipo continúan creciendo y también se han incorporado recientemente Manuel Collado y Pablo del Pino, desde Instituto de Investigaciones Sanitarias de Santiago (IDIS) y la Universidad de Santiago de Compostela, respectivamente.

El estudio está financiado por el CSIC a través de la Plataforma de Salud Global, que engloba más de 200 grupos de investigación de diferentes especialidades, desde biotecnología y nanotecnología hasta demografía e inteligencia artificial, para abordar los retos que plantea la epidemia del coronavirus SARS-CoV-2 con el objetivo de plantear soluciones a corto, medio y sobre todo largo plazo. (Fuente: CSIC/DICYT)

Fuente: NCYT Noticias de la Ciencia y la Tecnología. Disponible en <https://cutt.ly/GoR0L6J>

...vacunar es prevenir.

IMSS desarrolla vacuna contra coronavirus a partir de anticuerpos de pacientes

20 jun. El Instituto Mexicano del Seguro Social (IMSS) desarrolla una vacuna basada en proteínas del coronavirus SARS CoV-2, a partir del estudio de los anticuerpos producidos por sus primeros 300 pacientes covid-19 y mediante el uso de un potenciador de la respuesta inmunitaria ya patentado por la institución. Lo anterior lo reveló Constantino López Macías, jefe de la Unidad de Investigación Médica e Inmunoquímica (UIMI) de la Unidad Médica de Alta Especialidad (UMAE) Hospital de Especialidades del Centro Médico Nacional Siglo XXI.

“Para saber cómo inducir protección necesitamos saber qué significa esa protección, con los conocimientos modernos de inmunología vemos cómo es que el organismo se protege contra la infección y ya que sabemos cuál es la respuesta de anticuerpos necesaria de linfocitos T y la inmunidad innata, es decir, los componentes de la protección, entonces sabemos cómo podemos inducirlos”, señaló Constantino López Macías en entrevista exclusiva con MILENIO.

Un protocolo clínico desarrollado en esa unidad médica en el que se estudió la respuesta inmune de 300 pacientes covid-19 y sus contactos, permitió identificar las proteínas virales para diseñar el antígeno de la vacuna del IMSS

que funciona con un adyuvante - un potenciador de la respuesta inmune- desarrollado hace algunos años en la misma unidad de investigación con base en proteínas de salmonella, y que ya está patentando por el IMSS. “Lo que se hizo fue identificar cuáles son los fragmentos del virus que reconoce el sistema inmune y que pueden usarse como vacuna, es vacuna de subunidades la que estamos haciendo en el IMSS--; sin embargo, estas subunidades no son tan inmunogénicas, hay que ponerle algún potenciador -se llaman adyuvantes- y se mezclan con sustancias que ayudan a potenciar la respuesta inmune, como es el caso del adyuvante que se desarrolló en el IMSS que lo vamos a ocupar para esta vacuna”.

En términos simples, el investigador médico explicó que al infectarse, el organismo humano emprende una batalla para defenderse del virus, por lo que al identificar a los anticuerpos vencedores, “a los que ganaron esa guerra les preguntamos: cómo le hiciste para ganarle, y entonces con esa información, lo usamos para entrenar al sistema inmune de los demás”, pero requieren de un adyuvante para ayudar a reclutar más células. ¿Cómo funciona la vacuna? En la primera entrevista que concede, el doctor Constantino López explicó que al ingresar al organismo, el antígeno diseñado para la vacuna estimula

la respuesta inmune innata a través de células dendríticas, éstas son las células presentadoras de antígeno más potentes que existen y con capacidad de activar los linfocitos T, que son células del sistema inmune que juegan un papel como mediadores de la respuesta inmune contra el virus.

“El antígeno es captado por estas células dendríticas y es presentado a los linfocitos T para producir citocinas y por otro lado el antígeno es liberado y es reconocido por los linfocitos B y producen anticuerpos. Esta es una manera en la que el sistema innato y los linfocitos T y B son entrenados a través de la vacuna para producir esos anticuerpos y células T que son de los principales actores en la defensa, de esa manera se entrenan y cuando llega el virus que causa la enfermedad ya están entrenados para reconocerlo, atacarlo y destruirlo”, explicó. Además, dijo que el adyuvante ayuda a que se recluten más de las células de la respuesta innata, que “haya un ambiente que promueva la activación más eficiente en las células T y B para producir estas células con características adecuadas para que se induzca una respuesta para que no solo actúe en un tiempo corto, sino que actúe a larga plazo”.

¿En qué fase se encuentra el desarrollo de la vacuna?

López Macías dijo que se encuentran en la fase de producción de los antígenos para comenzar los ensayos preclínicos en ratones; explicó que tras las pruebas en laboratorio, el siguiente paso es producirla y garantizar que puede escalarse; de ahí que la importancia de formar parte del grupo de

científicos que representa a México ante la Coalición para la Innovación en la Preparación de Epidemias (CEPI).

“Estamos haciendo los estudios de producción y escalamiento y en el grupo CEPI están convergiendo entidades académicas que tiene plantas piloto que permiten hacer los escalamientos y de ahí ya se

puede hacer a que se produzcan a nivel industrial. Estamos colaborando con la UNAM y el IPN, entonces estaríamos haciendo esto muy probablemente en los próximos meses para que estemos evaluando estos candidatos en los modelos preclínicos”.

Fuente: MILENIO. Disponible en <https://cutt.ly/RoRLKyZ>

Clover lanza sexto ensayo de una vacuna china para la COVID-19

20 jun. Clover Biopharmaceuticals se convirtió el viernes en el sexto desarrollador chino de una posible vacuna para la COVID-19 en pasar a ensayos en humanos, lanzando un estudio en Australia que probará su fórmula de inmunización con refuerzos.

China y Estados Unidos se han enfrentado a la crisis de salud y están impulsando esfuerzos paralelos para lograr que las vacunas se aprueben y se produzcan a gran escala a finales de este año.

Actualmente no hay vacunas o tratamientos aprobados para la enfermedad causada por el nuevo coronavirus, pero alrededor de

una docena de vacunas se están probando en todo el mundo.

El virus surgió en la ciudad china de Wuhan a fines del año pasado y desde entonces se ha extendido a nivel mundial, matando a más de 450.000 personas.

Un aumento en los casos en la capital de China, Pekín, ha obligado a las autoridades a estudiar el genoma de una posible nueva cepa. [nL8N2DW11K]

Las pruebas de Clover, que está reclutando a unos 150 pacientes adultos y ancianos, evaluará dos refuerzos diferentes, o adyuvantes, de la británica GSK y la estadounidense Dynavax en com-

binación con su vacuna candidata, SCB-2019, dijo la compañía china.

La también china CanSino Biological, Sinopharm y el Instituto de Productos Biológicos Wuhan se encuentran entre los que ya están probando vacunas, al igual que AstraZeneca, Moderna y CureVac de Alemania.

Clover dijo que los datos iniciales de seguridad de su estudio se esperan para agosto, y que su objetivo sería comenzar estudios más amplios para fin de año. Su vacuna se basa en proteínas llamadas antígenos que se tomarán en combinación con los adyuvantes.

Fuente: infobae. Disponible en <https://cutt.ly/roRZ9F0>

Científicos reclaman más investigación sobre el SARS-CoV-2 en animales para minimizar riesgos

22 jun. A medida que aumenta la evidencia de la posibilidad de que el SARS-CoV-2 infecte a varios animales, los científicos del University College de Londres (UCL) dicen que se necesita un esfuerzo global para reducir el riesgo de que el virus regrese más tarde a las personas.

En un comentario para 'The Lancet Microbe', los investigadores escriben que si el virus se vuelve común en una población animal que vive cerca de personas, como mascotas o ganado, existe el riesgo de que ocurra otro brote incluso si el virus se erradica en las personas en el área.

Los autores piden más investigación sobre qué animales son susceptibles al SARS-CoV-2, el virus que causa la enfermedad de COVID-19, y sugieren implementar programas de vigilancia para evaluar regularmente a los animales que podrían presentar los mayores riesgos de transmisión.

La coautora, la profesora Joanne Santini, del UCL Structural & Molecular Biology, recuerda que "cada vez hay más pruebas de que algunos animales pueden contraer el SARS-CoV-2 de las personas y posteriormente pueden transmitirlo a otras personas, pero no sabemos exactamente cómo esto es un gran riesgo, ya que es un área de estudio que aún no ha sido priorizada".

"Necesitamos desarrollar estrategias de vigilancia para asegurarnos de que no nos tome por sorpresa un gran brote en animales, que podría representar una amenaza no solo para la salud animal sino también para la salud humana", añade.

A su juicio, "la transmisión del virus en poblaciones animales podría volverse irreversible si no se controla, y puede amenazar el éxito de las medidas de salud pública existentes si las personas continúan contagiando el virus de una población infectada de animales".

Los autores escriben que la inmensa escala de la pandemia de COVID-19 aumenta la posibilidad de que suficientes animales se conviertan en 'reservorios' del virus, lo que podría ser más probable que en epidemias pasadas, como el brote de SARS-CoV-1 más contenido en 2002-2003.

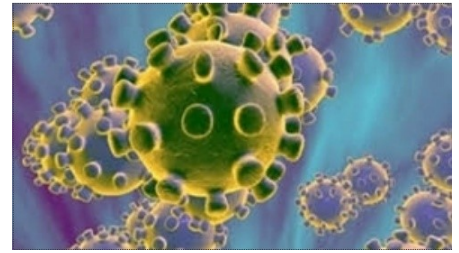
El profesor Santini y la coautora profesora Sarah Edwards, del

UCL Science & Technology Studies, revisaron la evidencia de estudios de casos, experimentos que prueban la infección en pequeños grupos de animales, así como estudios de laboratorio y modelos que describen mecanismos probables de infección.

Los estudios de modelado y de laboratorio sugieren que el SARS-CoV-2 podría, en teoría, transmitirse a numerosos animales, según los hallazgos de que la proteína espiga del virus se adhiere a las células huésped, utilizando una proteína que se encuentra en muchas especies diferentes.

El documento de investigación informa de que, una vez que los científicos identifiquen qué animales podrían infectarse, deben averiguar si se sentirán mal o si no serán asintomáticos, y si una persona infectada puede transmitir el virus a otros animales o incluso a los humanos.

En particular, ha habido casos recientes en los Países Bajos de visón de granja infectado con SARS-CoV-2, lo que llevó a dos personas a contraer el virus de estos animales, en un brote que ha provocado la eliminación de miles de visones. Los investigadores dicen que este ejemplo resalta no solo el riesgo para la salud humana, sino también las preocupaciones sobre el bienestar animal y la posible pérdida de medios de vida en el sector agrícola.



El profesor Edwards comenta que "existe una necesidad urgente de una vigilancia generalizada, mediante el análisis de muestras, preferentemente no invasivas, de un gran número de animales, en particular mascotas, ganado y fauna silvestre que se encuentran muy cerca de las poblaciones humanas".

A su juicio, "es poco probable que más experimentos de laboratorio con pequeñas cantidades de animales nos den las pruebas necesarias para estar seguros de que ciertas especies son totalmente seguras, lo que hace que el trabajo de vigilancia importante sea la única opción real en este caso".

"Necesitamos más información, al mismo tiempo que tomamos medidas de precaución simples, especialmente con las especies que tienen el potencial de propagar rápidamente el virus en el medio silvestre --continúa--. Una evaluación de riesgos robusta también requeriría revisar nuestra capacidad para manejar un brote en esos animales, es decir, nuestra capacidad para aislar, proteger o contener diferentes animales".

Fuente: alicante plaza. Disponible en <https://cutt.ly/joR4NmT>

...vacunar es prevenir.

Pfizer Announces Start of Four Phase 3 Clinical Trials for Investigational Vaccines

22 jun. First subjects recently administered immunizations in two studies of 20-valent pneumococcal conjugate vaccine candidate in infants; a pentavalent meningococcal vaccine candidate in adolescents; and a respiratory syncytial virus vaccine candidate in pregnant women.

Two studies (NCT04382326 and NCT04379713) of the 20-valent pneumococcal polysaccharide conjugate vaccine candidate, 20vPnC, evaluating a four-dose series in infants starting at 2 months of age. Both studies will expand the data on the safety and tolerability of the investigational vaccine in infants and include a control group of Prevnar 13® (Pneumococcal 13-valent Conjugate Vaccine [Diphtheria CRM197 Protein]). Study NCT04382326 has the goal of determining immunologic noninferiority between 20vPnC and Prevnar 13®, a critical requirement for vaccine licensure.

One study (NCT04424316) of the respiratory syncytial virus (RSV) vaccine candidate, RSVpreF, in pregnant women to evaluate the safety and efficacy of RSVpreF in infants born to immunized pregnant women as compared to placebo.

One study (NCT04440163) of the pentavalent meningococcal vaccine candidate, MenABCWY, in adolescents and young adults to assess the safety, tolerability, and immunogenicity of the

MenABCWY vaccine candidate compared to licensed meningococcal vaccines, with the goal of determining immunologic noninferiority.

“The start of four Phase 3 studies across our portfolio of investigational vaccines is a testament to the talented and dedicated colleagues working throughout Pfizer, and the continued commitment to unlock the potential promise and value that vaccines hold for our world,” said Kathrin U. Jansen, Ph.D., Senior Vice President and Head of Vaccine Research & Development at Pfizer Inc. “If approved, all three vaccine candidates could help prevent serious, possibly deadly infectious diseases that negatively impact millions of people of all ages globally.”

About 20vPnC Pediatric

Approximately 3,500 infants will be enrolled in total for these two studies. In both studies, infants will be vaccinated with either 20vPnC or Prevnar 13® (13vPnC) at 2, 4, 6, and 12-15 months of age, along with other routine infant vaccines according to the current CDC recommended schedule. Additional information can be found at www.clinicaltrials.gov under the identifiers NCT04382326 and NCT04379713. The results of the descriptive Phase 2 infant study with 20vPnC (NCT03512288) have been submitted for presentation at ID Week 2020.



In May 2017 the FDA granted Fast Track status for a pediatric indication for 20vPnC.1

Global Burden of Pneumococcal disease

Pfizer's 20vPnC vaccine candidate includes 13 serotypes already included in Prevnar 13® (13vPnC). Together, the 20 serotypes included in 20vPnC are responsible for the majority of currently circulating pneumococcal disease in the U.S. and globally.2,3,4,5,6,7,8

About RSVpreF

The Phase 3 trial of RSVpreF is a global, double-blind, placebo-controlled study that will enroll 6,900 pregnant women ages 18 through 49 and their infants. Additional information about the study can be found at www.clinicaltrials.gov under the identifier NCT04424316.

In April 2020, positive top-line results were achieved for a Phase 2b proof-of-concept study of RSVpreF, which evaluated the safety, tolerability and immunogenicity of RSVpreF in vaccinated pregnant women ages 18 through 49 and their infants. Detailed results from the study will be shared at a future medical conference. In November 2018, the

FDA granted Fast Track status to RSVpreF for prevention of RSV-associated lower respiratory tract illness in infants by active immunization of pregnant women.

Global Burden of RSV

RSV is a virus that can cause severe respiratory disease in infants and older adults.^{9,10} Globally, there are an estimated 33 million cases of RSV annually in children less than 5 years of age, with about 3 million hospitalized and approximately 120,000 dying each year from complications associated with the infection. Nearly half of these pediatric hospitalizations and deaths occur in infants less than 6 months of age.¹¹ The medical community is limited to offering only supportive care for those with the illness.

Fuente: Pfizer. Disponible en <https://cutt.ly/hoBwxg6>

About MenABCWY

The Phase 3 trial will enroll approximately 2,413 adolescents and young adults (10 through 25 years of age) from the United States and Europe. Additional information about the study can be found at www.clinicaltrials.gov under the identifier NCT04440163.

Initiation of the Phase 3 trial is based on positive results from a proof-of-concept study (NCT03135834) in 543 adolescents and young adults. Detailed results from the proof-of-concept study have been submitted for presentation at ID Week 2020.

Pfizer's pentavalent meningococcal vaccine candidate combines its two approved meningococcal vaccines, Nimenrix™ (meningococcal group A, C, W-

135, and Y conjugate vaccine) and Trumenba® (meningococcal group B vaccine). Approvals of Nimenrix™ and Trumenba® vary by country.

Global Burden of Meningococcal Disease

Meningococcal disease is an uncommon but serious disease that can attack without warning^{12,13} and lead to meningitis and serious blood infections.^{14,15} The majority of invasive meningococcal disease cases worldwide can be attributed to five *Neisseria meningitidis* groups (A, B, C, W and Y).¹⁶ Together, these meningococcal groups account for 96% of all invasive meningococcal disease (IMD), with group B accounting for the majority of disease in adolescents and young adults in the U.S. and Europe.

Vacuna contra coronavirus COVID-19: La propuesta israelí pasa la prueba con roedores

22 jun. A raíz de la crisis de sanidad que estamos viviendo en la actualidad, varios grupos de científicos han tomado la labor de buscar una vacuna que pueda prevenir que una enfermedad que se propaga tan rápido como el COVID-19 vuelva a tener un brote que sea tan devastador como el actual.

Entre este grupo de científicos, este fin de semana del 21 de junio destacaron los israelíes y los chinos por haber anunciado

avances importantes en las vacunas trabajadas desde que empezó la pandemia.

El Instituto de Israel para la Investigación Biotecnológica probó con éxito una posible vacuna en roedores, cuyos experimentos son clave para el desarrollo de la medicina dado que si funcionan, pueden continuar con otros animales y en última instancia con humanos, según los investigadores en su informe por el medio digital Sin Embargo.

Hubo dos grupos de hámsters infectados dentro del experimento: los que recibieron la vacuna con "dosis única" y los que no. Los roedores inmunizados dejaron de perder peso corporal, mientras que los pulmones de aquellos sin vacunar sufrieron daños extensos en sus tejidos y vías respiratorias, a diferencia del otro grupo cuyos efectos colaterales fueron menores...

Fuente: Yahoo finanzas. Disponible en <https://cutt.ly/YoBuPFt>



VacciMonitor es una revista con más de 25 años de difundir los resultados científicos sobre vacunas de instituciones nacionales e internacionales y así coadyuvar a la visibilidad de este sector de la ciencia en Cuba y otros países, principalmente de Hispanoamérica. <http://vaccimonitor.finlay.edu.cu>

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Patentes registradas en PatentScope

Estrategia de búsqueda: *Vaccine in the title or abstract AND 20200616:20200622 as the publication date*

38 records

1. [20200188508](#) PORCINE EPIDEMIC DIARRHEA VIRUS S PROTEIN AND SUBUNIT VACCINE THEREOF AS WELL AS METHOD FOR PREPARING SUBUNIT VACCINE AND APPLICATION THEREOF

US - 18.06.2020

Int.Class [A61K 39/215](#) Appl.No 16600334 Applicant NOVO BIOTECH CORP. Inventor Hong QIAN

The disclosure discloses a porcine epidemic diarrhea virus S protein and a subunit vaccine thereof as well as a method for preparing the subunit vaccine and application thereof. The vaccine contains 30~220 µg of a recombinant porcine epidemic diarrhea virus S protein and a pharmaceutically acceptable ISA 201 VG adjuvant. A method for preparing the subunit vaccine comprises the following steps: (1) cloning the recombinant porcine epidemic diarrhea virus S protein; (2) expressing and purifying the recombinant porcine epidemic diarrhea virus S protein; (3) preparing the recombinant porcine epidemic diarrhea virus S protein prepared in (2) into a water phase; (4) emulsifying the water phase and the ISA 201 VG adjuvant in a volume ratio of 46:54 to obtain a vaccine. The vaccine is high in safety, good in immunogenicity, stable in batches, low in production cost and strong in immunogenicity. On the other hand, CHO cell strains suspending and stably and efficiently expressing the PEDV-S protein are successfully constructed and screened for the first time. The CHO cell strain can express the PEDV-S protein in high yield, the yield can reach 1 g/L, and the expressed PEDV-S protein is easy to purify.

2. [20200188507](#) INFLUENZA VIRUS VACCINE AND METHOD OF MAKING

US - 18.06.2020

Int.Class [A61K 39/145](#) Appl.No 16608662 Applicant Mississippi State University Inventor Xiu-Feng Wan

A composition for increasing vaccine yields due to increased virus growth in mutation comprising a vaccine strain bearing the Y161F mutation in hemagglutinin (HA). Y161F in HA increases HA thermostability without changing its original antigenic properties and enhances its binding affinity in the vaccine production platforms used in influenza vaccine manufacturing. A method for optimizing preparation of influenza vaccine seed strains which can further lower the cost of vaccines and increase profits for the vaccine companies, and also maintain antigenic stability during vaccine deliveries.

3. [WO/2020/123759](#) SUBUNIT VACCINE CONSTRUCTS FOR FLAVIVIRUSES

WO - 18.06.2020

Int.Class [A61K 39/12](#) Appl.No PCT/US2019/065886 Applicant REGENTS OF THE UNIVERSITY OF MINNESOTA Inventor DAVID, Sunil A.

This disclosure describes a subunit vaccine for a flavivirus, methods of making the vaccine, and methods of using the vaccine. The flavivirus may include, is a mosquito-borne flavivirus, for example, Zika virus (ZIKV), dengue virus (DENV), Yellow Fever (YF) virus, and West Nile Virus (WNV). The subunit vaccine may be administered with an adjuvant.

4. [3664838](#) DIFFERENTIAL COATING OF MICROPROJECTIONS AND MICRONEEDLES ON ARRAYS
EP - 17.06.2020

Int.Class [A61K 39/145](#) Appl.No 18844031 Applicant VAXXAS PTY LTD Inventor JUNGER MICHAEL CARL
The present invention relates to devices and methods for coating microprojection or microneedle arrays including arrays that contain vaccine formulations, more specifically to multivalent vaccine formulations where components of the multivalent vaccine might be incompatible. The present invention further relates to stable vaccine formulations for administration via a microprojection array in which the microprojections are densely packed and in which the vaccine formulations are sprayed on to the microprojections such that the formulations dry quickly.

5. [WO/2020/123777](#) RECOMBINANT MUMPS VIRUS VACCINE EXPRESSING GENOTYPE G FUSION AND HEMAGGLUTININ-NEURAMINIDASE PROTEINS
WO - 18.06.2020

Int.Class [C12N 7/00](#) Appl.No PCT/US2019/065926 Applicant THE UNITED STATES OF AMERICA, AS REPRESENTED BY THE SECRETARY, DEPARTMENT OF HEALTH AND HUMAN SERVICES Inventor RUBIN, Steven A.

A recombinant, attenuated mumps virus is described. The recombinant virus is based on the genotype A Jeryl Lynn vaccine strain, but is modified to express genotype G consensus fusion (F) and hemagglutinin-neuraminidase (HN) proteins. The recombinant virus optionally includes a mutation that prevents expression of viral protein V. The recombinant mumps virus can be used as a vaccine to inhibit mumps virus infection and the development of mumps disease.

6. [WO/2020/121983](#) VACCINE FOR PREVENTING OR TREATING CONGENITAL INFECTION WITH CYTOMEGALOVIRUS
WO - 18.06.2020

Int.Class [A61K 39/245](#) Appl.No PCT/JP2019/047966 Applicant KM BIOLOGICS CO., LTD. Inventor TORIKAI Masaharu

The problem addressed by the present invention is to provide an effective vaccine for preventing or treating congenital infection with CMV. This vaccine for preventing or treating congenital infection with cytomegalovirus (CMV) contains a CMV envelope glycoprotein B (gB protein) antibody and a pentamer antibody.

7. [WO/2020/122734](#) PANCREAS DISEASE VIRUS MARKERS
WO - 18.06.2020

Int.Class [C12Q 1/70](#) Appl.No PCT/NO2019/050274 Applicant PATOGEN AS Inventor DEVOLD, Magnus Andreas

The present invention relates to methods for detection of the Pancreatic Disease (PD) virus used for production of PD vaccines, use of primers and probes to differentiate PD virus used in a vaccine and infectious virus, and a kit for detection of PD virus used in a vaccine. The method includes nine SNP's and related methods that can be used to identify the virus in the vaccine.

8. [202047022653](#) FILOVIRUS VACCINE AND METHODS OF USE
IN - 19.06.2020

Int.Class [A61K 39/12](#) Appl.No 202047022653 Applicant HAWAII BIOTECH, INC. Inventor CLEMENTS, David E.

The data reported herein describe the production and evaluation of a recombinant subunit filovirus vaccine using insect cell expressed surface glycoprotein (GP) and a highly effective adjuvant. The vaccine provides protection in humans against filovirus infection, including Ebola virus and Marburg virus.

9.[20200188509](#)ORF VIRUS-BASED PLATFORM FOR VACCINE DELIVERY

US - 18.06.2020

Int.Class [A61K 39/275](#) Appl.No 16086894 Applicant South Dakota Board of Regents Inventor Diego G. Diel

The present invention is directed to novel vaccine delivery platform based on the Orf virus (ORFV) genome, which carry heterologous antigens, methods of making and methods of using the same for prevention of infections, diseases, and other conditions in animals.

10.[20200188513](#)COMPOSITIONS OF VACCINES AND ADJUVANTS AND METHODS FOR THE TREATMENT OF URINARY TRACT INFECTIONS

US - 18.06.2020

Int.Class [A61K 39/39](#) Appl.No 16689027 Applicant Sequoia Vaccines, Inc Inventor Gary Eldridge

This invention describes novel adjuvant compositions and formulations with excellent stability at refrigerated and room temperatures and up to and about 37° C. that can be produced at remarkably low costs. This invention describes novel vaccine compositions and formulations to treat and prevent urinary tract infections caused by gram-negative bacteria including *Escherichia coli* and multi-drug resistant *E. coli*. This invention also describes methods of administration of said novel vaccine compositions and formulations and methods of treatment to prevent and treat urinary tract infections caused by gram-negative bacteria including *E. coli* and multi-drug resistant *E. coli*.

11.[3106175](#)Hidtil ukendt immunterapi mod flere forskellige tumorer, herunder nerve- og hjernetumorer

DK - 22.06.2020

Int.Class [A61K 39/00](#) Appl.No 16179163 Applicant Immatix Biotechnologies GmbH Inventor HILF, Norbert

The present invention relates to peptides, 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 cytotoxic T cell (CTL) peptide epitopes, alone or in combination with other tumor-associated peptides that serve as active pharmaceutical ingredients of vaccine compositions that stimulate anti-tumor immune responses. The present invention relates to 30 peptide sequences and their variants derived from HLA class I and class II molecules of human tumor cells that can be used in vaccine compositions for eliciting anti-tumor immune responses.

12.[WO/2020/123341](#)HEPATITIS B VIRUS VACCINE AND USES THEREOF

WO - 18.06.2020

Int.Class [A61K 39/29](#) Appl.No PCT/US2019/065151 Applicant SU, Zhuang Inventor SU, Zhuang

A hepatitis B virus (HBV) vaccine particle is described, including a recombinant HBV surface antigen including L surface protein; optionally M surface protein; and optionally S surface protein; wherein the molar percentage of L surface protein to the sum of L, M, and S surface proteins is at least about 1 mole %, 8 mole

%, 10 mole %, 20 mole %, 30 mole %, 40 mole %, or 50 mole %. Methods of making the same and methods of treating or preventing HBV infection in a subject using the same are also described.

13. [3120870](#) Hidtil ukendt immunterapi mod flere forskellige tumorer, herunder nerve- og hjernetumorer
DK - 22.06.2020

Int.Class [A61K 39/00](#) Appl.No 16179226 Applicant Immatics Biotechnologies GmbH Inventor Schoor, Oliver
The present invention relates to peptides, 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 cytotoxic T cell (CTL) peptide epitopes, alone or in combination with other tumor-associated peptides that serve as active pharmaceutical ingredients of vaccine compositions that stimulate anti-tumor immune responses. The present invention relates to 30 peptide sequences and their variants derived from HLA class I and class II molecules of human tumor cells that can be used in vaccine compositions for eliciting anti-tumor immune responses.

14. [20200188519](#) REDUCED FOAMING VACCINE COMPOSITIONS
US - 18.06.2020

Int.Class [A61K 47/26](#) Appl.No 16802089 Applicant Abic Biological Laboratories Ltd. Inventor Noel Yves Henri Jean Genin
The present invention relates to novel stable compressed vaccine composition comprising at least one anhydrous antigenic component comprising a stabilizer susceptible to foaming when the composition is mixed with liquid diluent; and an effective amount of a sugar alcohol.

15. [3157549](#) Hidtil ukendt immunterapi mod flere forskellige tumorer i blodet, navnlig kronisk lymfatisk leukæmi (CLL)
DK - 22.06.2020

Int.Class [A61K 39/00](#) Appl.No 15730135 Applicant immatics biotechnologies GmbH Inventor STICKEL, Juliane
The present invention relates to peptides, 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 cytotoxic T cell (CTL) peptide epitopes, alone or in combination with other tumor-associated peptides that serve as active pharmaceutical ingredients of vaccine compositions that stimulate anti-tumor immune responses. The present invention relates to several novel peptide sequences and their variants derived from HLA class I and HLA class II molecules of human tumor cells that can be used in vaccine compositions for eliciting anti-tumor immune responses.

16. [3120868](#) Hidtil ukendt immunterapi mod flere forskellige tumorer, herunder nerve- og hjernetumorer
DK - 22.06.2020

Int.Class [A61K 39/00](#) Appl.No 16179174 Applicant Immatics Biotechnologies GmbH Inventor HILF, Norbert
The present invention relates to peptides, 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 cytotoxic T cell (CTL) peptide epitopes, alone or in combination with other tumor-associated peptides that serve as active pharmaceutical ingredients of vaccine compositions that stimulate anti-tumor immune responses. The present invention relates to 30 peptide sequences and their variants

derived from HLA class I and class II molecules of human tumor cells that can be used in vaccine compositions for eliciting anti-tumor immune responses.

17. [WO/2020/123296](#) PREDICTING IMMUNOGENIC PEPTIDES USING STRUCTURAL AND PHYSICAL MODELING

WO - 18.06.2020

Int.Class [G06F 19/16](#) Appl.No PCT/US2019/064959 Applicant UNIVERSITY OF NOTRE DAME DU LAC Inventor BAKER, Brian

Disclosed herein are methods for predicting immunogenicity of a candidate peptide. The method comprises obtaining a three-dimensional candidate structural representation of the candidate peptide bound to an antigen presenting molecule; obtaining a plurality of candidate measurements; and predicting, with an electronic processor, the immunogenicity of the candidate peptide based upon the plurality of candidate measurements. Further disclosed herein are methods for producing vaccines. The method for producing a vaccine comprises predicting immunogenicity of one or more candidate peptides using the methods described herein, and producing a vaccine comprising one or more peptides predicted to be immunogenic.

18. [WO/2020/123912](#) INHIBITION OF ASPH EXPRESSING TUMOR GROWTH AND PROGRESSION

WO - 18.06.2020

Int.Class [A61K 39/00](#) Appl.No PCT/US2019/066174 Applicant RHODE ISLAND HOSPITAL Inventor WANDS, Jack, R.

Disclosed are compositions and methods for an immunotherapy in a subject containing a vaccine construct for an immunization against a purified tumor antigen and a checkpoint inhibitor for treating a tumor in the subject, in which the tumor is characterized as comprising a low frequency of neoantigen expression and the composition potentiates an anti-tumor immune response without inducing autoimmunity in the subject. A pharmaceutical composition containing the composition as an active component and a pharmaceutically acceptable carrier, and a combinatorial composition containing a vaccine construct for an immunization against a purified tumor antigen and an immune checkpoint inhibitor, in which the tumor is characterized as comprising a low frequency of neoantigen expression, are also described.

19. [201811047503](#) A RECOMBINANT PEPTIDE, AND APPLICATIONS THEREOF AGAINST ESCHERICHIA-RELATED INFECTIONS

IN - 19.06.2020

Int.Class [A61K /](#) Appl.No 201811047503 Applicant CHAIRMAN, DEFENCE RESEARCH & DEVELOPMENT ORGANISATION Inventor MANOHARAN, Renuka Ramalingam

The present disclosure discloses a recombinant DNA comprising a nucleic acid sequence as set forth in SEQ ID NO: 1, a recombinant protein encoded by the recombinant DNA, a recombinant vector comprising the recombinant DNA, and arecombinant host cell comprising the recombinant vector. The present disclosure furtherdiscloses a vaccine composition comprising the recombinant protein of the presentdisclosure against Escherichia-related infections. Also disclosed are the methods forproducing the recombinant protein and for preparing the vaccine composition of thepresent disclosure.

20. [3666286](#) UNIVERSAL INFLUENZA VACCINE

EP - 17.06.2020

Int.Class [A61K 39/145](#) Appl.No 19215483 Applicant CAMBRIDGE TECH LLC Inventor HAUSE BEN
Immunogenic compositions for inducing a universal immune response to influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies which provide protection against heterologous influenza infection. Compositions comprising recombinant baculovirus expression vectors expressing neuraminidase in cultured insect cells dispersed in a pharmaceutically-acceptable carrier comprising insect cell culture media, and optional adjuvant. Methods of inducing immune responses against influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies in a host animal susceptible to infection.

21. [WO/2020/123964](#) LABYRINTHIN-BASED PEPTIDES FOR CANCER IMMUNOTHERAPIES AND USES THEREOF

WO - 18.06.2020

Int.Class [C07K 14/47](#) Appl.No PCT/US2019/066264 Applicant LABYRX IMMUNOLOGIC THERAPEUTICS (USA) LIMITED Inventor RADOSEVICH, James A.

Antigenic compositions comprising one or more labyrinthin-derived peptides are described herein. In some embodiment, each peptide of the antigenic composition comprises a T-cell epitope and/or a B-cell epitope. In other aspects, the present disclosure provides, e.g., vaccine compositions comprising tin antigenic composition disclosed herein, including kits, medicines, and compositions (such as pharmaceutical compositions and unit dosages) thereof. Also provided are methods of using the compositions disclosed herein, such as methods of treatment thereof and methods of producing antibodies, and antibody compositions thereof, against the one or more labyrinthin-derived peptides or a portion thereof.

22. [WO/2020/121273](#) HETEROLOGOUS PRIME BOOST VACCINE COMPOSITIONS AND METHODS

WO - 18.06.2020

Int.Class [A61K 39/12](#) Appl.No PCT/IB2019/060766 Applicant GLAXOSMITHKLINE BIOLOGICALS SA Inventor CAPONE, Stefania

Simian adenoviral vectors and RNA molecules, each encoding an immunogen of interest, can be sequentially administered to provide potent and long-lasting immunity.

23. [WO/2020/123836](#) HERBOXIDIENE SPLICING MODULATOR ANTIBODY-DRUG CONJUGATES AND METHODS OF USE

WO - 18.06.2020

Int.Class [A61K 47/68](#) Appl.No PCT/US2019/066029 Applicant EISAI R&D MANAGEMENT CO., LTD. Inventor FISHKIN, Nathan

Linker-drug compounds and antibody-drug conjugates that bind to human oncology targets are disclosed. The linker-drug compounds and antibody-drug conjugates comprise a herboxidiene splicing modulator drug moiety. The disclosure further relates to methods and compositions for use in the treatment of neoplastic disorders by administering the antibody-drug conjugates provided herein. The herboxidiene itself is also claimed. Further claims are directed to its use, and to the use of a neoantigen, generated by the herboxidiene or its ADC, or a vaccine against this neoantigen.

24. [20200188501](#) BACTERIAL VACCINE COMPONENTS AND USES THEREOF

US - 18.06.2020

Int.Class [A61K 39/085](#) Appl.No 16749556 Applicant SOCPRA - SCIENCES ET GENIE, s.e.c. Inventor FRANCOIS MALOUIN

Agents, compositions, methods and kits useful for the treatment and diagnosis of Staphylococcal intramammary infection are disclosed. The agents, compositions, methods and kits are derived from genes expressed during Staphylococcal intramammary infection, and more particularly genes SACOL0442, based on the gene nomenclature from the *Staphylococcus aureus* COL (SACOL) genome.

25. [3666290](#)A TELOMERASE ENCODING DNA VACCINE

EP - 17.06.2020

Int.Class [A61K 48/00](#) Appl.No 19211063 Applicant INVECTYS Inventor LANGLADE DEMOYEN PIERRE

The invention provides a nucleic acid construct comprising a sequence that encodes a human telomerase reverse transcriptase (hTERT) protein which is devoid of telomerase catalytic activity and of a nucleolar localization signal. The construct is useful triggering an immune response in a subject, against cells that overexpress telomerase, preferably dysplasia cells or tumor cells.

26. [20200188497](#)TRANSFECTED T-CELLS AND T-CELL RECEPTORS FOR USE IN IMMUNOTHERAPY AGAINST CANCERS

US - 18.06.2020

Int.Class [A61K 39/00](#) Appl.No 16802431 Applicant Immatix Biotechnologies GmbH Inventor Dominik MAURER

The present description relates to T-cell receptors (TCRs) binding to tumor-associated antigens (TAAs) for targeting cancer cells, T-cells expressing same, methods for producing same, and methods for treating cancers using same. In particular, the present description relates to TCRs and their variants that bind to HLA class I or II molecules with a peptide, such as IGF2BP3-001 have the amino acid sequence of KIQEILTQV (SEQ ID NO:1). The present description further relates to peptides, proteins, nucleic acids and cells for use in immunotherapeutic methods. In particular, the present description relates to the immunotherapy of cancer. The present description 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.

27. [20200188496](#)UNIVERSAL CANCER VACCINES AND METHODS OF MAKING AND USING SAME

US - 18.06.2020

Int.Class [A61K 39/00](#) Appl.No 16617830 Applicant ARIZONA BOARD OF REGENTS ON BEHALF OF ARIZONA STATE UNIVERSITY Inventor Stephen Albert JOHNSTON

Disclosed herein are compositions comprising a universal cancer vaccine and methods of treating and preventing cancer using such compositions.

28. [20200188500](#)VACCINE COMPRISING CLOSTRIDIUM TOXOIDS

US - 18.06.2020

Int.Class [A61K 39/08](#) Appl.No 16620846 Applicant HIPRA SCIENTIFIC, S.L.U. Inventor Xavier GIBERT PÉREZ

The present invention relates to an immunogenic composition comprising one or more *C. difficile* toxoid for use in a medicament for animals. The invention also encompasses an immunogenic composition comprising one or more *C. difficile* A toxoid and one or more *C. difficile* B toxoid and one or more *C. perfringens* Type A toxoid. The invention also encompasses vaccines comprising said immunogenic compositions, vaccines for use in the treatment and/or prevention of disease caused by *C. difficile* and *C. perfringens*, and kits thereof.

29. [20200188504](#) HEPATITIS C VIRUS GENE SEQUENCES AND METHODS OF USE THEREFOR
US - 18.06.2020

Int.Class [A61K 39/12](#) Appl.No 16608308 Applicant VANDERBILT UNIVERSITY Inventor James E. CROWE, JR.

The present disclosure relates to compositions and methods for inducing an adaptive immune response against Hepatitis C virus (HCV) in a subject. In some embodiments, the present disclosure provides a composition comprising a nucleic acid molecule encoding a HCV antigen, an HCV antigen, an adjuvant, or a combination thereof. For example, in some embodiments, the composition comprises a vaccine comprising a nucleic acid molecule encoding a HCV antigen, an HCV antigen, an adjuvant, or a combination thereof.

30. [WO/2020/123757](#) UNIVERSAL INFLUENZA VACCINE
WO - 18.06.2020

Int.Class [A61K 39/145](#) Appl.No PCT/US2019/065883 Applicant CAMBRIDGE TECHNOLOGIES LLC
Inventor HAUSE, Ben

Immunogenic compositions for inducing a universal immune response to influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies which provide protection against heterologous influenza infection. Compositions comprising recombinant baculovirus expression vectors expressing neuraminidase in cultured insect cells dispersed in a pharmaceutically-acceptable carrier comprising insect cell culture media, and optional adjuvant. Methods of inducing immune responses against influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies in a host animal susceptible to infection.

31. [20200188505](#) UNIVERSAL INFLUENZA VACCINE
US - 18.06.2020

Int.Class [A61K 39/12](#) Appl.No 16711873 Applicant Cambridge Technologies LLC Inventor Ben Hause
Immunogenic compositions for inducing a universal immune response to influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies which provide protection against heterologous influenza infection. Compositions comprising recombinant baculovirus expression vectors expressing neuraminidase in cultured insect cells dispersed in a pharmaceutically-acceptable carrier comprising insect cell culture media, and optional adjuvant. Methods of inducing immune responses against influenza, and particularly influenza A, by eliciting anti-neuraminidase antibodies in a host animal susceptible to infection.

32. [20200190146](#) METHODS OF DETECTION AND REMOVAL OF RHABDOVIRUSES FROM CELL LINES
US - 18.06.2020

Int.Class [C07K 14/005](#) Appl.No 16662784 Applicant TAKEDA VACCINES, INC. Inventor Joel R. HAYNES
The present disclosure relates to compositions, methods, mixtures, and kits for detecting the presence of, and for removing, a virus from a product produced in an insect cell. The disclosure also relates to proteins, peptides, polypeptides, drug substances, biological products, vaccine antigens, and virus-like particles that

are produced in an insect cell and that are free or substantially free of a virus. The disclosure also relates to compositions, methods, assays, and kits for detecting a rhabdovirus in a sample.

33. [201811047504](#)A RECOMBINANT PEPTIDE, AND APPLICATIONS THEREOF AGAINST STAPHYLOCOCCUS AUREUS-RELATED INFECTIONS

IN - 19.06.2020

Int.Class [A61K](#) / Appl.No 201811047504 Applicant CHAIRMAN, DEFENCE RESEARCH & DEVELOPMENT ORGANISATION Inventor JAYAKRISHNAN, Achuth

The present disclosure discloses a recombinant DNA (comprising a nucleic acid sequence as set forth in SEQ ID NO: 1), a recombinant protein encoded by therecombinant DNA (comprising an amino acid sequence as set forth in SEQ ID NO: 2), a recombinant vector comprising the recombinant DNA, and a recombinant host cell comprising the recombinant vector. The present disclosure further discloses a vaccine composition comprising the recombinant protein of the present disclosure for targeting Staphylococcus-related infections. Also disclosed are the methods for producing therecombinant protein and for preparing the vaccine composition of the present disclosure.

34. [20200188438](#)IMMUNOTHERAPY WITH B*08 RESTRICTED PEPTIDES AND COMBINATION OF PEPTIDES AGAINST CANCERS AND RELATED METHODS

US - 18.06.2020

Int.Class [A61K 35/17](#) Appl.No 16717142 Applicant Immatics Biotechnologies GmbH Inventor Gisela SCHIMMACK

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.

35. [3664839](#)HEADLESS HEMAGGLUTIN INFLUENZA VACCINE

EP - 17.06.2020

Int.Class [A61K 39/145](#) Appl.No 18864814 Applicant GEORGIA STATE UNIV RESEARCH FOUNDATION INC Inventor WANG BAOZHONG

Disclosed are universal influenza based on a truncated influenza hemagglutinin (HA) protein lacking a head domain (hrHA). Also disclosed is a composition comprising a nanoparticle coated with a disclosed hrHA polypeptide. Also disclosed is a composition comprising a virus like particle (VLP) expressing on its surface a disclosed hrHA polypeptide.

36. [20200190525](#)Reducing the Toxicity of Agrobacterium Endotoxin

US - 18.06.2020

Int.Class [C12N 15/74](#) Appl.No 16617971 Applicant ARIZONA BOARD OF REGENTS ON BEHALF OF ARIZONA STATE UNIVERSITY Inventor Qiang Chen

The present invention relates to the fields of genetically modified *Agrobacterium* strains, vaccine adjuvants, and generally molecular biology and immunology. Provided herein are modified *Agrobacterium* strains that produce lipopolysaccharide (LPS) having reduced toxicity or detoxified lipopolysaccharide, and methods of obtaining such strains for plant-based production of biologics. Also provided herein are uses of reduced or detoxified LPS as adjuvants suitable for clinical use.

37. [20200190160](#) PEPTIDES AND COMBINATION THEREOF FOR USE IN THE IMMUNOTHERAPY AGAINST CANCERS

US - 18.06.2020

Int.Class [C07K 14/74](#) Appl.No 16799380 Applicant IMMATICS BIOTECHNOLOGIES GMBH Inventor 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.

38. [WO/2020/120910](#) PRODUCTION OF VIRAL VACCINES ON AN AVIAN CELL LINE

WO - 18.06.2020

Int.Class [A61K 39/12](#) Appl.No PCT/FR2019/053036 Applicant UNIVERSITE CLAUDE BERNARD LYON 1 Inventor ROSA-CALATRAVA, Manuel

The invention relates to the use of the immortalised cell line ECACC 09070703, filed on 7 July 2009 with the European Collection Of Cell Cultures (ECACC, Salisbury, United Kingdom) under the number 09070703, for the production of a viral vaccine consisting of an attenuated strain derived from a human metapneumovirus.

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

Results of Search in US Patent Collection db for: (ABST/vaccine AND ISD/20200616->20200622),

9 resultados.

PAT. NO.	Title
1 10,683,361	Anti-CD40 antibodies
2 10,683,337	Peptides, combination of peptides as targets and for use in immunotherapy against gallbladder cancer and cholangiocarcinoma and other cancers
3 10,682,426	Rabies vaccine

- 4 [10,682,407](#) [Constrained immunogenic compositions and uses therefor](#)
 - 5 [10,682,406](#) [Nucleic acid comprising or coding for a histone stem-loop and a poly\(A\) sequence or a polyadenylation signal for increasing the expression of an encoded pathogenic antigen](#)
 - 6 [10,682,404](#) [Targets of Acinetobacter baumannii](#)
 - 7 [10,682,402](#) [MSI-specific frameshift peptides \(FSP\) for prevention and treatment of cancer](#)
 - 8 [10,682,399](#) [Target peptides for colorectal cancer therapy and diagnostics](#)
 - 9 [10,682,398](#) [Yeast vaccine vector including immunostimulatory and antigenic polypeptides and methods of using the same](#)
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Edición: Annia Ramos Rodríguez aramos@finlay.edu.cu
Ma. Victoria Guzmán Sánchez mguzman@finlay.edu.cu
Randelys Molina Castro rmolina@finlay.edu.cu
Yamira Puig Fernández yamipuig@finlay.edu.cu
Rolando Ochoa Azze ochoa@finlay.edu.cu