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EDICIONES**



BOLETÍN

VACCIENCIA

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...vacunar es prevenir.

Análisis bibliométrico sobre vacunas contra el virus del papiloma humano (VPH)

Fuente de información utilizada:



Estrategia de búsqueda:

TITLE: ("papillomavirus vaccine") 1001 records

Periodo de estudio 1999-2020

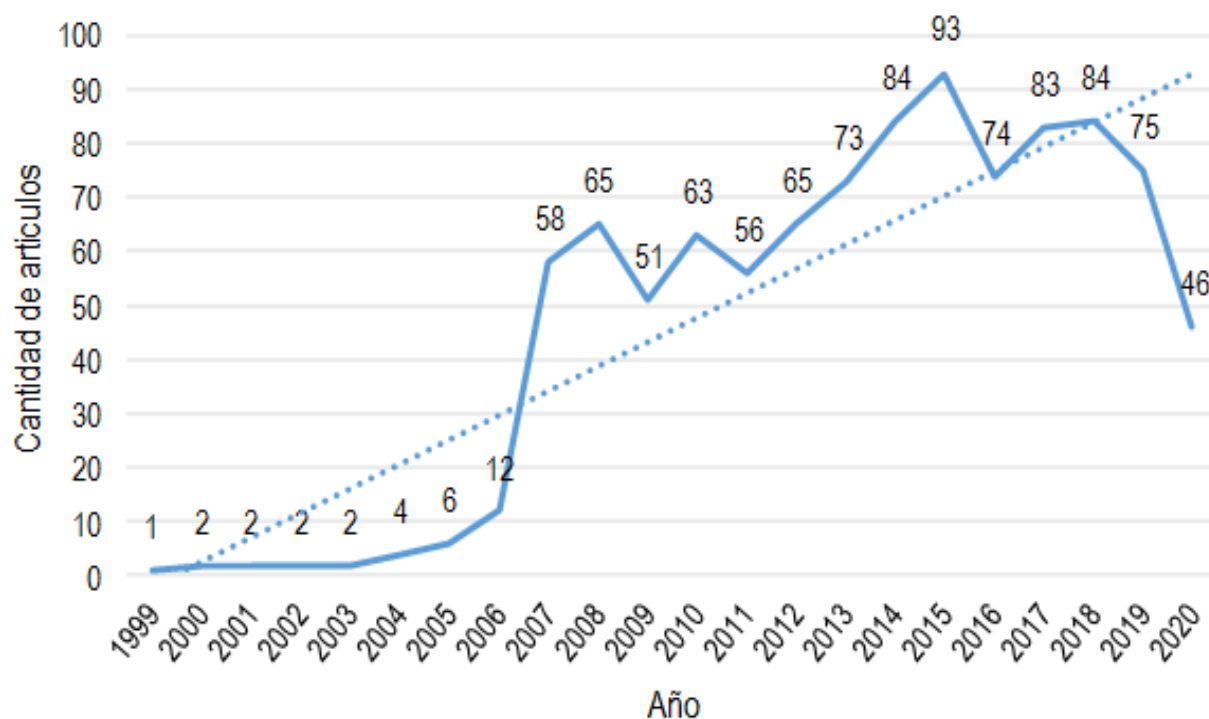
Las variables utilizadas en el análisis fueron:

- ⇒ 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.

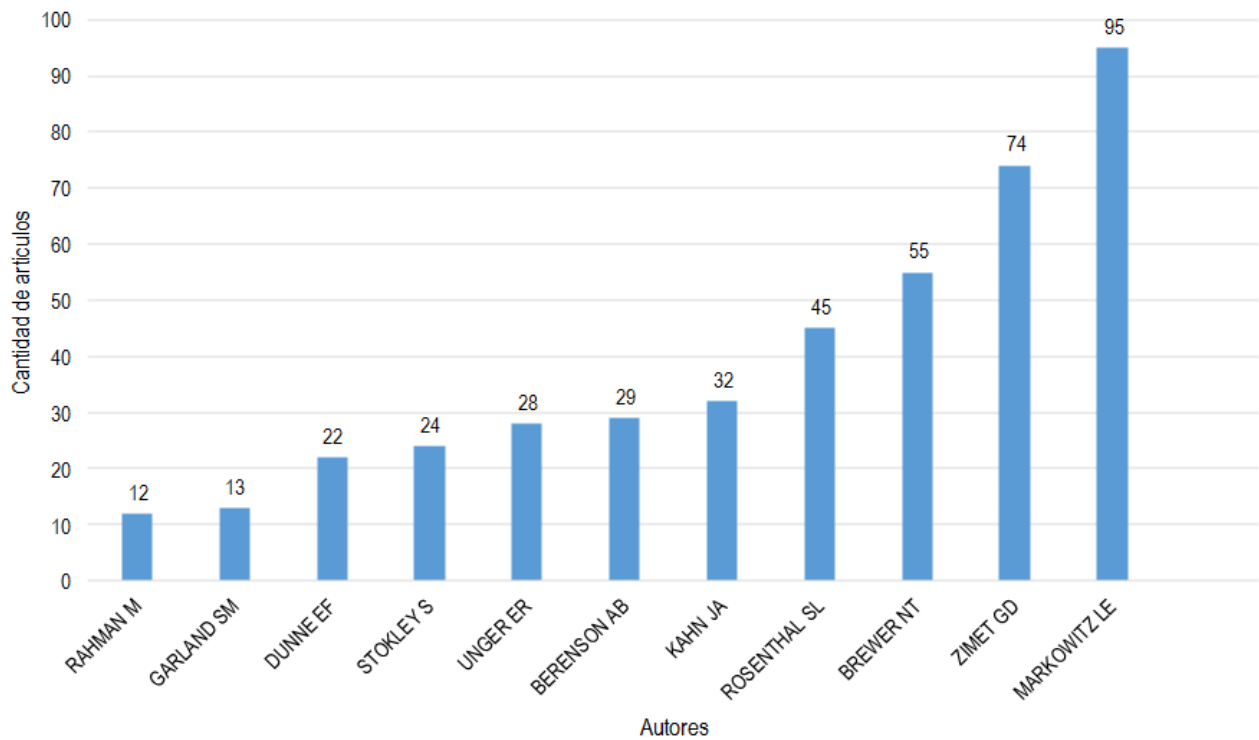
EN ESTE NÚMERO

- * Análisis bibliométrico vacunas contra el virus del papiloma humano (VPH)
- * Noticias en la Web sobre vacunas
- * Artículos científicos más recientes Medline sobre vacunas
- * Patentes más recientes USPTO sobre vacunas
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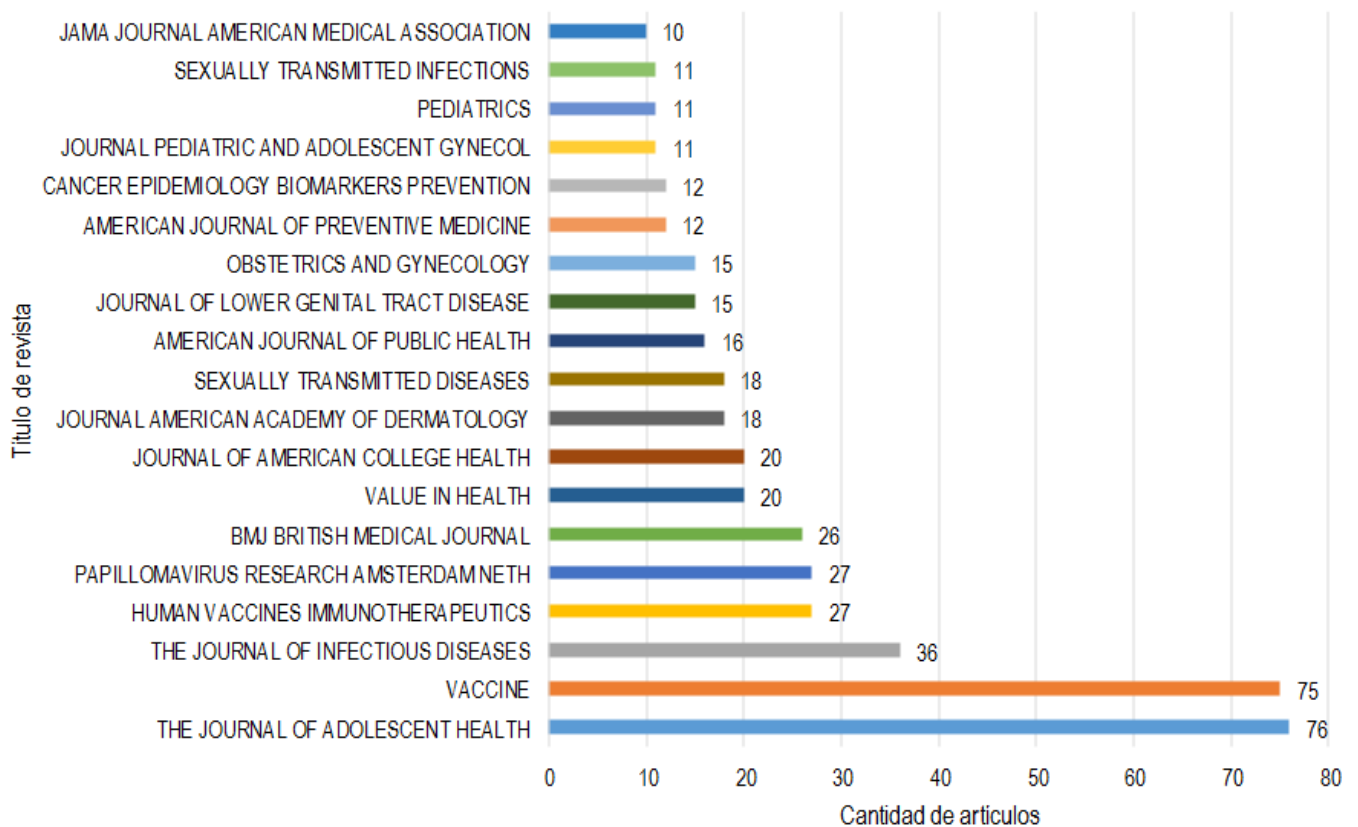
Productividad científica por año



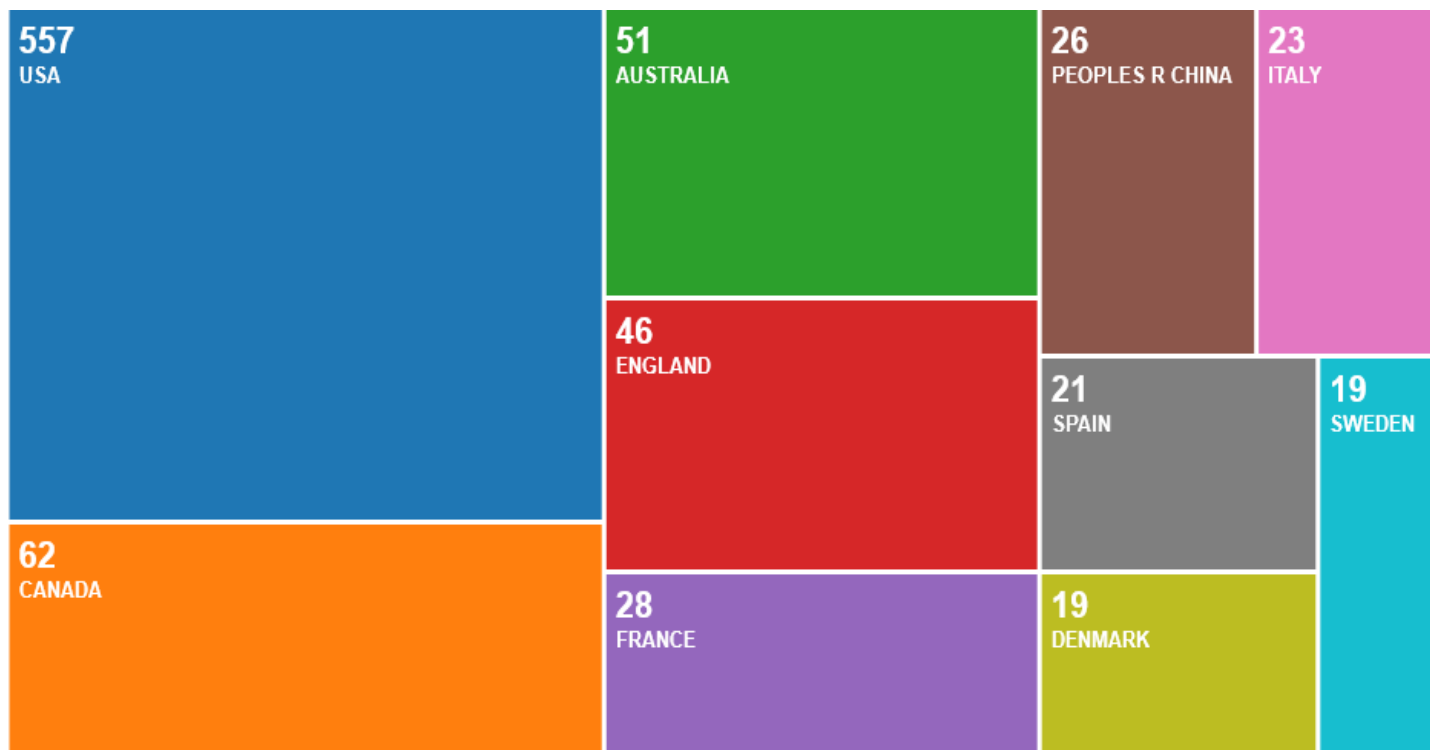
Autores con mayor productividad científica



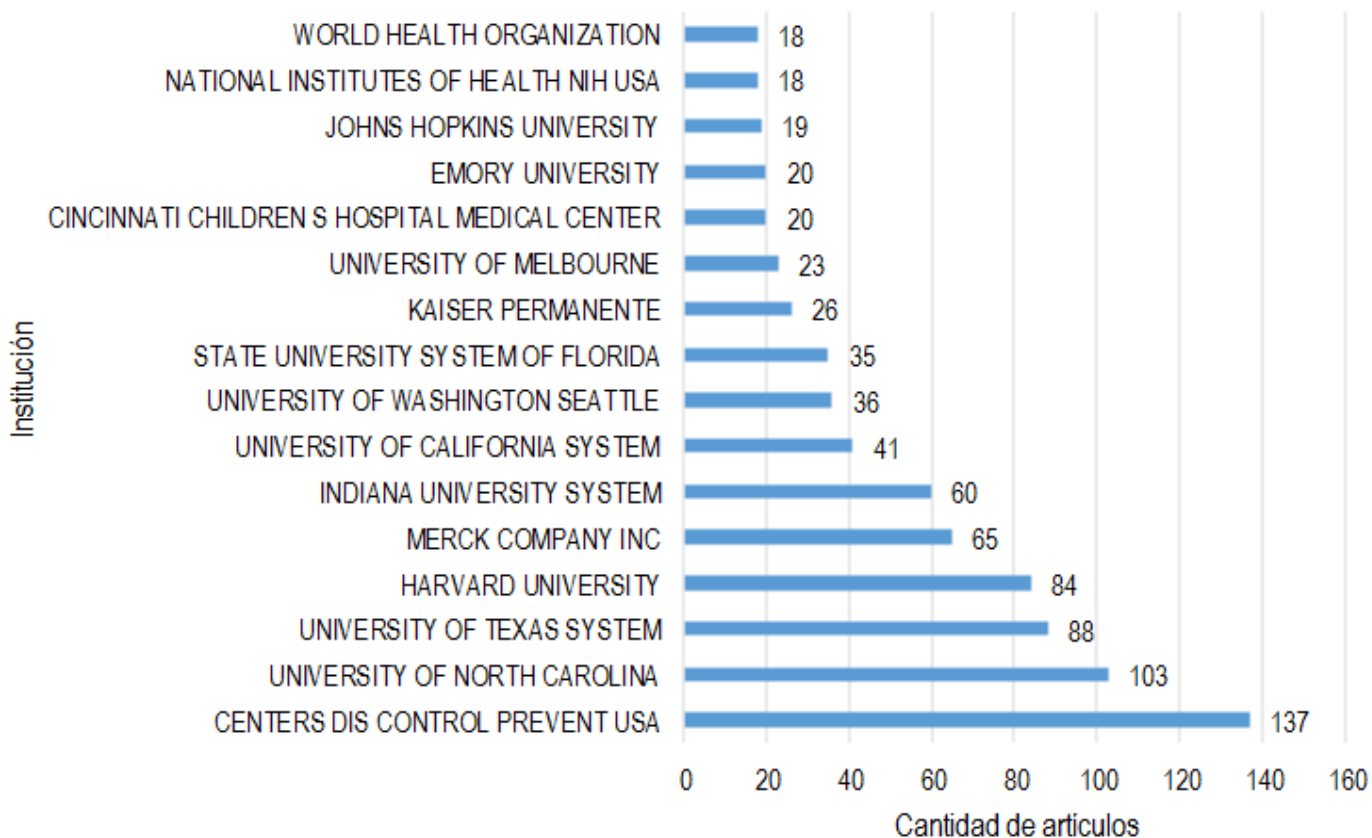
Revistas científicas que han publicado sobre el tema (2019-2020)



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



Instituciones que han trabajado el tema de estudio



Noticias en la Web

El poder de la vacuna contra la tuberculosis: ¿revacunación protegería frente a la Covid-19?

9 jun. Desde hace tiempo se conocen algunos estudios en los que se muestra la capacidad protectora de esta vacuna frente a otros patógenos que no son la bacteria de la tuberculosis, incluso, frente algunos virus. Nuevos estudios en desarrollo sugieren que esta inoculación reactiva la memoria inmunológica de las personas, protegiendo contra el coronavirus.

Un grupo de investigadores de la Universidad Católica de Chile y del Instituto Milenio de Inmunología e Inmunoterapia, IMII, liderado por el Alexis Kalergis, e integrado por los académicos de la Universidad Católica: Susan Bueno, Camila Covián y Angello Retamal, recientemente publicaron dos trabajos en la revista especializada *Frontiers in Immunology*. En ellos los especialistas sugerirían un posible efecto beneficioso de los programas de inmunización con la vacuna BCG (*Bacillus Calmette-Guérin*) -utilizada para combatir los casos de tuberculosis- como protección frente al coronavirus humano. Los análisis presentados en este trabajo de investigadores chileno sugieren la importancia de evaluar científicamente el posible efecto benéfico de la BCG contra el Sars-CoV-2 por medio de estudios clínicos. Además, las observaciones presentadas en esta publicación por los investigadores



nacionales, se basan en datos públicos que sugieren que los países que utilizan la BCG presentan, en general, un menor número de casos por millón de habitantes, en comparación con aquellos que no la utilizan.

Este fenómeno podría deberse a la inmunidad entrenada que induce esta vacuna en las personas y que consiste en la capacidad de las células inmunes innatas para actuar con cierta programación inmunológica. Por lo tanto, los científicos plantean que este tipo de inmunidad podría ser favorable como estrategia de inmunización contra el Sars-CoV-2.

Sin embargo, los académicos plantean que otros factores podrían también influir en los índices que se discuten, como por ejemplo la frecuencia de los diagnósticos, el nivel socioeconómico y la tasa de contagio en los

países analizados.

“Estudiamos el número de casos positivos y fatalidades en diferentes países y los correlacionamos con la inclusión de la vacuna BCG al nacer en sus programas nacionales de inmunización. Curiosamente, aquellos donde esta se administra reportan una tasa de contagio más baja y una menor cantidad de fallecimientos relacionadas con Covid-19”, explica Alexis Kalergis.

El académico de la Universidad Católica agrega que “esto sugiere que esta vacuna puede inducir inmunidad entrenada, la cual podría otorgar cierto nivel de protección para el Sars-CoV-2 y disminuir la probabilidad de infección. Se ha demostrado que este tipo de inmunidad confiere protección contra una amplia variedad de patógenos, incluidas bacterias, hongos, virus y protozoos”. Sin embargo, esta hipótesis debe ser evaluada

científicamente por medio de estudios clínicos. Como consecuencia, existen actualmente más de diez estudios clínicos en el planeta para evaluar si acaso la BCG común puede proteger contra el Covid-19.

Estos datos sugieren un papel crucial para esta vacuna en el desarrollo de memoria inespecífica contra otros virus respiratorios, como Sars-CoV-2. Sin embargo, el fenotipo “entrenado” dura un tiempo limitado, lo que sugiere que este tipo de inmunidad desarrollada al nacer podría no ser capaz de proteger a los adultos contra infecciones posteriores. Por lo tanto, los investigadores del Departamento de Genética Molecular y Microbiología, Facultad de Ciencias Biológicas de la Universidad Católica sugieren que la vacunación con BCG podría reactivar la inmunidad entrenada de una manera más fuerte en comparación con la primera inducción.

Modelos experimentales

El Dr. Alexis Kalergis comenta que, en un modelo experimental de infección viral por fiebre amarilla, la inducción de inmunidad entrenada reduce los niveles de viremia o ingreso de un virus en el torrente sanguíneo.

“Además, se ha demostrado que la vacuna BCG es efectiva para prevenir infecciones agudas del

tracto respiratorio superior en los ancianos y se asocia con una reducción del asma y la atopia en adultos”, resalta Kalergis.

“La vacuna BCG ha demostrado su inmunogenicidad y seguridad, por lo que se ha utilizado durante casi 100 años en humanos. Además, puede conferir protección independiente del antígeno contra una amplia variedad de patógenos”, destaca Kalergis. Dos ensayos clínicos diferentes, agrega, respaldan la idea de que la revacunación con BCG induce una activación más fuerte de la protección cruzada no específica asociada a esta vacuna.

El primer ensayo clínico realizado entre 1935 y 1947, evidenció que la revacunación de los niños disminuía progresivamente su mortalidad general. La primera vacuna lo redujo solo en un 3%, pero lograron un 47% de reducción de la mortalidad en los niños después de la tercera revacunación.

Otro ensayo clínico realizado en Guinea-Bissau también demostró una disminución de la mortalidad en niños revacunados, con una reducción del 64%.

Tratamiento antiviral efectivo

Las vacunas estimulan la activación de la respuesta inmune adaptativa y el desarrollo de la memoria inmunológica.

Sin embargo, uno de los mayores

problemas de la pandemia producida por el SARS-CoV-2 es la ausencia de un tratamiento antiviral efectivo o una vacuna, el cual pueda contrarrestar la respuesta inflamatoria e incluso el daño agudo severo a los pulmones.

Para el desarrollo de una vacuna, es necesario conocer la estructura del patógeno contra el cual está diseñada la formulación, así como los componentes inmunogénicos, como los adyuvantes. Sin embargo, el desarrollo de una nueva formulación y ensayos preclínicos y clínicos puede llevar una cantidad significativa de tiempo.

Tomando en cuenta su seguridad como vacuna en grandes poblaciones, BCG podría considerarse por su amplia disponibilidad y bajo costo como una buena estrategia para el desarrollo de inmunidad entrenada y protección contra nuevos patógenos.

Actualmente, la capacidad de inmunidad entrenada inducida por BCG para proteger contra Covid-19 se está evaluando en más de diez ensayos clínicos. Uno se está llevando a cabo en Holanda, involucrando a 1,500 participantes y 147 trabajadores de la salud que serán vacunados, y otro en Australia con 4.000 participantes y 148 voluntarios que serán vacunados.

Fuente: La Tercera. Disponible en <https://bit.ly/319wHSV>



Científicos identifican 47 medicamentos existentes que podrían ayudar a combatir la COVID-19

9 jun. En lugar de pensar en fármacos de nuevo diseño, científicos de la Universidad de California buscaron entre los 2 mil medicamentos ya aprobados para humanos, algunos que pudieran tratar o indicar un camino terapéutico para la COVID-19.

El equipo multidisciplinario de UCSF identificó 69 medicamentos y componentes con potencial para tratar la COVID-19. A medida que lo hacía, enviaba muestras de esas sustancias al Instituto Pasteur, en París, y a la Escuela de Medicina Icahn de Mount Sinai, en Nueva York, para someterlos a pruebas. Hasta el momento, 47 mostraron “fuertes pautas para el tratamiento” y se identificaron “dos mecanismos separados” sobre el modo en que interceptarían al SARS-CoV-2, según sintetizaron en un primer estudio.

En principio, esas pautas y mecanismos derivados del cruce entre el mapa del coronavirus y el catálogo de la FDA no señalaban más que interacciones potenciales: “No sabíamos si las drogas que identificamos harían que una persona fuera más resistente al virus o más susceptible, si harían algo o nada”. Para eso hacía falta probarlas en muestras vivas del causante de la COVID-19 y en células, que se tomaron del mono verde porque

reaccionan de manera muy similar a las humanas.

Los investigadores de UCSF identificaron principalmente dos grupos de sustancias que afectan al virus de dos maneras diferentes, una de las cuales no había sido descrita antes.

El primer grupo interrumpe la traducción del mensaje del ARN, que es uno de los pasos finales que da el coronavirus para multiplicarse. Cuando ingresa a una célula, el SARS-CoV-2 se apropia de los mecanismos naturales de ella para hacer copias de sí mismo. Primero se replica, luego transcribe las instrucciones para hacer eso numerosas veces gracias a la capacidad propia de la célula y por último traduce ese mensaje: en ese punto las proteínas hacen nuevas copias que pasan a infectar a otras células. Y el proceso vuelve a comenzar.

Actualmente está en estudio una molécula similar a la ternatina-4, el medicamento Aplidin, un anti-tumoral de origen español cuyo principio activo es la plitidespina. Según dijo Luis Mora, director de la empresa que lo produce, PharmaMar, sus efectos contra la COVID-19 podrían ser “mil veces superiores a los que consigue el remdesivir”, el antiviral de Gilead aprobado por la FDA.

En cambio, la zotatifina apunta a otra proteína.

El equipo de investigación de UCSF trabaja actualmente con el laboratorio que la produce eFFECTOR Therapeutics, para comenzar cuanto antes los ensayos clínicos. Según explicó el director ejecutivo de la firma, Steve Worland, a diferencia de los otros antivirales esta molécula, originalmente creada contra el cáncer, no apunta al virus sino que actúa sobre las células que el SARS-CoV-2 secuestra.

Los receptores celulares, que se encuentran tanto en el interior como en la superficie de las células, “actúan como interruptores especializados”, comparó el investigador: “Cuando una molécula específica se une a un receptor específico, le indica así a la célula que haga una tarea específica. Con frecuencia los virus utilizan receptores para infectar las células”.

El mapa original había mostrado que dos receptores importantes en los tratamientos farmacológicos, SigmaR1 y SigmaR2, podían participar en el combate contra la COVID-19. “Las pruebas confirmaron nuestras sospechas”, siguió Krogan.

China inicia la etapa 1 de las pruebas clínicas del anticuerpo recombinante JS016 para tratar la COVID-19

9 jun. Según informes, es la primera prueba clínica a nivel mundial del anticuerpo en un participante humano saludable después de terminar las pruebas en primates no humanos, indicó la comisión municipal de ciencia y tecnología de Shanghai.

El primer sujeto del Hospital Huashan de la Universidad de Fudan, en Shanghai, recibió hoy por la mañana una inyección de JS016, un anticuerpo recombinante, completamente humano y monoclonal contra la COVID-19.

El anticuerpo, desarrollado de forma conjunta por la compañía biofarmacéutica Junshi Biosciences, el Instituto de Microbiología de la Academia de Ciencias de China y otros, entró a la etapa 1 de la prueba clínica después de recibir la aprobación de la Administración Nacional de Productos Médicos.



La prueba aleatoria, a doble ciego y controlada con placebo busca evaluar la tolerabilidad, seguridad, características farmacocinéticas e inmunogenicidad de JS016 entre la población china, para sentar bases para posteriores estudios clínicos del anticuerpo.

La prueba fue dirigida por Zhang Jin, subdirector del Instituto de Antibióticos del hospital, y Zhang Wenhong, jefe del Centro de Enfermedades Contagiosas del hospital.

Se espera que la terapia con anticuerpo neutralizador sea la primera opción de tratamiento en el combate a la COVID-19, dijo Zhang Wenhong, quien señaló que el anticuerpo puede atacar con precisión el coronavirus e inhibir la replicación del virus.

Fuente: sinembargo. Disponible en <https://bit.ly/3fUVwWR>

La expansión del coronavirus depende del estilo de vida

9 jun. Una serie de variables que caracterizan la organización y el estilo de vida de un país (desde el número de visitantes que recibe, a su consumo eléctrico o sus emisiones de CO₂), predicen el número total de contagiados y de muertos por Covid-19 que va a tener ese país con mucha más exactitud que otras variables, como el desarrollo de su sanidad o su nivel

de riqueza y progreso. Esta es la conclusión de una investigación desarrollada por científicos de la Facultad de Veterinaria de la Universidad Complutense de Madrid y publicada en la sección de COVID-19 SARS-CoV-2 preprints de MedRxiv: analiza qué variables pueden predecir mejor el número de infecciones y muertes en todo el mundo por SARS-CoV-2.

Los autores estudian docenas de variables en 50 países diferentes, desde los indicadores generales del país (área, superficie, porcentaje de población urbana...), a sus parámetros demográficos (tasa de nacimiento, esperanza de vida, población mayor de 65 años...), sus estimadores económicos (tasa de inflación, tasa de

paro...), su gasto en I+D y en educación, sus indicadores de salud (camas de hospital por habitante, médicos por habitante...), su consumo eléctrico, sus índices de contaminación ambiental (emisiones de CO₂, óxido nítrico, metano...), etc.

Además, analizaron cómo la eficacia de estas variables predictivas varía con el tiempo: hay análisis independientes para los meses de marzo, abril y mayo de 2020 (y los autores pretenden seguir su estudio durante los meses siguientes).

El principal problema es que, cuando el SARS-CoV-2 empezó su expansión por el mundo, había una absoluta falta de conocimientos: era un nuevo virus del que no se sabía nada.

Muchos asumieron que se comportaría como el virus de la gripe. Otros que lo haría de forma similar a otros coronavirus anteriores (SARS-CoV-1 o MERS). Pero, en realidad no había conocimientos suficientes para saberlo.

Nadie sabía cómo acertar

Muchos países -por ejemplo, España- se fiaron de la supuesta robustez de su sistema sanitario. Recordemos que al principio de la pandemia de la Covid-19, el director del Centro de Coordinación de Alertas y Emergencias Sanitarias previó que el SARS-CoV-2 tendría “poco efecto en España” pues teníamos “la mejor sanidad del mundo”.

No sirvió de nada: Al principio de desatarse la pandemia, el número de muertos de cada país tan solo dependió del número de contagiados que tenía ese país. Daba igual que tuviesen una sanidad extraordinaria o que su sanidad fuera endeble. Poco importó que tuviesen muchos o pocos médicos por habitante y muchas o pocas camas hospitalarias.

Al inicio de la pandemia, tanto los países con una sanidad excelente como los países con una sanidad deficiente mostraron un comportamiento prácticamente idéntico: en todos ellos se moría el mismo porcentaje de infectados.

Este resultado era de esperar: Como no se sabía nada del virus, no había vacuna ni fármacos eficaces y se desconocían los protocolos de tratamiento: los grandes recursos sanitarios todavía servían de poco.

En esos momentos, la clave estaba en impedir que la gente se contagiase.

Y en el número de contagios, la organización de los distintos países resultó esencial: en aquellos que recibían un mayor número de turistas y viajeros (casos de Italia, España, Francia, Estados Unidos...) se desató la catástrofe.

Y eso que en las fechas en las que se inició la pandemia no era temporada alta (lo que hubiese tenido un resultado desolador). Pero el que un país esté más o

menos volcado en el turismo condiciona su estilo de vida: un estilo de vida que resulta fatal en la expansión de las pandemias.

La prevención inicial es esencial

A los países que adoptaron, cuanto antes, estas estrategias preventivas controlando rigurosamente las fronteras y manteniendo, de una forma u otra, un riguroso aislamiento social, les fue muy bien, con independencia de su grado de desarrollo.

Países desarrollados como Taiwán o Nueva Zelanda, que primaron las medidas preventivas, acertaron. Países ricos como Suecia, que confiando en su sistema sanitario relajaron la prevención, se equivocaron (y recientemente sus responsables lo reconocieron).

Se dio la paradoja de que países como Vietnam (una nación pobre, con una elevada población -95 millones de habitantes-, una sanidad precaria y 1400 kilómetros de frontera con China) apenas resultó afectado por el coronavirus (a finales de abril aún no había tenido ni un solo muerto por Covid-19). Uruguay, que adoptó pronto medidas de confinamiento, no tuvo que enfrentarse a la catástrofe sanitaria de su vecino Brasil.

También la contaminación influye

Además, otra serie de variables permiten predecir con gran exactitud las cifras de muertes y contagios por Covid-19:

Entre ellas destacan las emisiones de CO₂ y de óxido nítrico a la atmósfera.

En la gran mayoría de los casos, basta conocer si un país tiene muchas emisiones de estos gases contaminantes para acertar con el número de contagiados y muertos por Covid-19 que ha tenido.

Sin embargo, esto no quiere decir que sean causa y efecto: que haya más CO₂ atmosférico no necesariamente significa que el CO₂ favorezca los contagios por SARS-CoV-2. Lo que significa es que los países que liberan más CO₂ tienen mayor número de contagiados y de muertos por Covid-19 (y mientras más CO₂ liberan, tienen más contagiados).

Probablemente tenga que ver con su estilo de vida: buena parte del CO₂ y del óxido nítrico liberados a la atmósfera provienen del transporte (en buena parte de los automóviles y los aviones).

Otras variables indirectamente relacionadas con la contaminación, como el consumo eléctrico, también presentan una correlación significativa con el número de muertos por Covid-19. Y con ellas ocurre lo mismo: solo son variables predictivas, no causa y efecto.

Cambio global, más pandemias

El cambio global está acelerando las pandemias que asolan a la humanidad.



Tan solo en lo que va de siglo la humanidad padeció epidemias y pandemias como el SARS (2002-2004), la Gripe A (2009-2010), el MERS (2012), el Chikungunya (2013-2014), el Ébola (2014, 2018), el Zika (2015-2016), el Cólera (2016-2017) y el Dengue (año 2019).

Seguimos con una elevada tasa de SIDA, y gracias a las antivacunas se están produciendo rebrotes preocupantes de enfermedades que se creían erradicadas como el Sarampión, la Polio o la Difteria. Buena parte de ellas son zoonóticas (tienen un origen en los animales salvajes tal y como ocurrió con el SARS-CoV-2).

Vendrán más pandemias. No tardarán mucho. Y serán peores que la del Covid-19, concluyen los investigadores.

Por eso, cada vez se piensa más en el concepto de One Health (Una Salud). Nuestra salud y nuestro bienestar se basa en tres pilares fundamentales: la salud de los ecosistemas, la salud de los animales (salvajes y domésti-

cos) y la salud humana. Tenemos que cuidar estas tres patas.

Estilo de vida insostenible

La gestión de la salud de los animales puede enseñarnos mucho acerca de la gestión de las pandemias humanas.

Hay muchos más animales de abasto que seres humanos. Viven muchísimo más hacinados. Están sometidos a muchas más pandemias potenciales que las personas.

Y, por razones económicas, no se puede gastar mucho dinero en el tratamiento de pandemias. Por eso la gestión veterinaria de las pandemias se basa en la prevención y en tomar medidas de aislamiento inmediato.

Unas medidas que hubiesen funcionado excelentemente impidiendo que la Covid-19 llegase a ser una pandemia mundial, concluyen los investigadores.

EMASESA desarrolla sistema de alerta temprana de SARS-CoV-2 en aguas residuales

9 jun. El nuevo escenario mundial ocasionado por la aparición de un nuevo coronavirus, ha cambiado nuestra forma de vida hasta extremos insospechados. Este nuevo virus ha dado lugar a una nueva enfermedad humana denominada COVID-19, que afecta principalmente a las vías respiratorias y ha generado una pandemia mundial que no entiende de fronteras. La emergencia mundial ha provocado que la comunidad científica se vuelque en realizar estudios sobre el nuevo virus y buscar soluciones a nuestro alcance para luchar contra esta pandemia.

En la sociedad en su conjunto, existe un miedo generalizado por la aparición de nuevos brotes. La dinámica del virus hace que se produzca un desfase entre el contagio y la aparición de los primeros síntomas.

Por ello es importante poder contar con herramientas de detección precoz de población portadora, para la prevención de nuevos casos.

Distintas líneas de investigación en España y Europa han constatado que en las aguas residuales se pueden detectar un aumento de los niveles de circulación del Covid-19 o partes de sus fragmentos genéticos en las aguas residuales, tanto en los casos en que haya un foco de población contagiada o, cuando el foco de la población sólo haya presentado los primeros síntomas o sean asintomáticos, por lo que el

análisis de las aguas residuales se convierte en una herramienta de anticipación ante nuevos brotes y muy útil para los servicios médicos y la administración.

Ante estas evidencias científicas y la irrupción de la pandemia por la COVID-19 y la preocupación por la crisis sanitaria ocasionada, EMASESA lleva semanas trabajando en la detección del actual coronavirus una vez que las aguas residuales llegan a sus EDAR, Estación Depuradora de Aguas Residuales (El Coper, Norte, Tablada, Ranilla, Mairena-El Viso). EMASESA, como responsable de la gestión de las aguas residuales, cuenta con una extensa red de saneamiento de 2.980 km. de longitud, que recoge las aguas de más de un millón de personas de Sevilla y su área metropolitana (Alcalá de Guadaíra, Dos Hermanas, La Rinconada, Camas, San Juan de Aznalfarache, Coria del Río, La Puebla del Río, Mairena del Alcor, Alcalá del Río, El Garrobo y El Ronquillo), transportándolas hasta las EDAR.

EMASESA viene trabajando en su protocolo habitual en la toma y análisis permanente de muestras por ello tenemos muestras congeladas desde semanas previas a la pandemia.

Con relación al COVID-19 nos hemos ido adaptando a los protocolos de este nuevo virus a través del Proyecto: "Determinación de SARS-CoV-2 en aguas residuales

como Sistema de Alerta Temprana para el estudio epidemiológico de la pandemia" Hoy somos capaces y estamos en condiciones de afirmar si existe COVID-19 en nuestras aguas residuales. Tal es así que se ha constatado que en días en los que aún había pocos contagios se encontraron restos de COVID-19 en la red de saneamiento.

La información obtenida es muy sensible y relevante por ello, partiendo de esta dinámica de muestreo y análisis, se está trabajando en tramificar y sectorizar la red de saneamiento para facilitar la detección de procedencia de la muestra y ser capaces de aislar la información, si se produjera, un posible brote en una zona concreta de cualquier municipio. Por ejemplo, una barriada o un distrito tanto en Sevilla como en su área metropolitana.

Podemos confirmar que, a día de hoy, con la aplicación de este sistema de alerta, todas las muestras analizadas han dado resultados negativos en COVID-19.

Este método nace con vocación de ampliarse a otros patógenos y virus; y de ser una herramienta complementaria para la de otros profesionales a nivel sanitario o epidemiológico.

Así, podremos poner a disposición de las autoridades sanitarias una herramienta de estudio que complementa a la información que ya maneja

permitiéndoles localizar zonas y posibles focos en los que puedan producirse nuevos brotes y posibilitar su temprana mitigación. Así como detectar casos asintomáticos, ya que como adelantábamos, en las muestras se puede detectar el virus antes de que aparezcan los síntomas de contagio en las personas afectadas.

La presencia de restos de virus en las aguas residuales no es

perjudicial para la salud. Se analizan muestras de restos de virus para cuantificar su presencia. Las EDAR son instalaciones seguras, ya que los tratamientos que reciben las aguas residuales permiten la eliminación de estos restos de virus, devolviéndose al cauce el agua “límpia” es decir, en óptimas condiciones.

Para Emasesa, como empresa pública de un servicio esencial, ha

sido una preocupación la calidad y seguridad del agua, así como la protección de la salud de las personas y la conservación del medio ambiente. De ahí el esfuerzo por ofrecer nuevos servicios y trabajar con instalaciones más seguras, como el nuevo laboratorio que incorpora novedades tecnológicas y científicas para estar al día en el control de calidad.

Fuente: RETEMA Revista Técnica de Medio Ambiente. Disponible en <https://bit.ly/2CuRvd9>

Congress to closely monitor pneumonia vaccine tender

10 jun. A party-list lawmaker on Wednesday said Congress would closely monitor the tender for the anti-pneumonia vaccine for children to ensure cost-effectiveness and viability of the product choice, in Manila, Philippine.

PHILRECA Rep. Presley de Jesus said a resolution has been filed to encourage an open, fair, and competitive bidding for the pneumococcal conjugate vaccines, PCV 10 and PCV 13.

He said the procurement of vaccines should not be delayed to prevent further disease outbreaks, particularly vaccine-preventable diseases (VPDs), amid the coronavirus disease 2019 (Covid-19) pandemic.

“As a policy, we should not delay the procurement of vaccines for VPDs especially as we have seen a surge of VPDs in several pockets of community outbreaks. While the DOH (Department of Health)

promised a swift review of the PCV tender, we at Congress should make sure the viability of all types of pneumonia vaccines,” de Jesus said.

House Resolution 906 urges the DOH to ensure continued safe implementation of the mandated National Immunization Program for children despite the challenges posed by the pandemic.

During a virtual forum hosted by the Samahang Plaridel on Tuesday, Pediatric Infectious Diseases Society of the Philippines (PIDSP) vice president Dr. Mary Ann Bunyi the two PCVs are comparable in performance based on new evidence gathered by global health experts.

“We have been given updates on both PCV10 and PCV13. These updates have been reviewed and assessed by the immunization committee and we see that both are comparable,” Bunyi said.

“So, the updates that were given to

us by both companies we have forwarded to the Department of Health and we leave it to the DOH, which pneumococcal vaccine will be made available for public use. Especially given for free to the vulnerable children,” he added.

Meanwhile, Health Undersecretary Dr. Maria Rosario Vergeire said the government is spending PHP4.9 billion on the procurement of pneumococcal vaccines

The DOH previously requested the Health Technology Assessment Committee to review the National Immunization Program (NIP), particularly the Pneumococcal Vaccination Program for children, in light of new 2017 and 2019 evidence from the World Health Organization (WHO).

In February 2019, WHO reaffirmed this earlier position saying that the two available PCVs in the market, PCV10 and PCV13, are equally effective in preventing overall pneumococcal diseases in children.

Vergeire said the result of the review is yet to be released, but a public health forum regarding the matter is set to happen this week. "We have a public forum on Thursday based on the assessment that has been done by the HTAC so that we can address the different queries coming from our stakeholders. We can also consult on the different aspects of this assessment," she said.

Dr. Wilda Silva, National Immunization Program Manager of the DOH, reported on the high reproduction number of vaccine-preventable diseases, such as polio, diphtheria, pertussis, and measles.

"Although there is a big problem with SARS-CoV-2, we should not forget other diseases which have vaccines as a bullet to control these vaccine-preventable diseases," Silva said.

Silva stressed the importance of continued immunization services, saying that "vaccine gives our children a good start at life."

"...ALTHOUGH THERE IS A BIG PROBLEM WITH SARS-CoV-2, WE SHOULD NOT FORGET OTHER DISEASES WHICH HAVE VACCINES AS A BULLET TO CONTROL THESE VACCINE-PREVENTABLE DISEASES ."

Fuente: Philippine Canadian Inquirer. Disponible en <https://bit.ly/387C1b1>

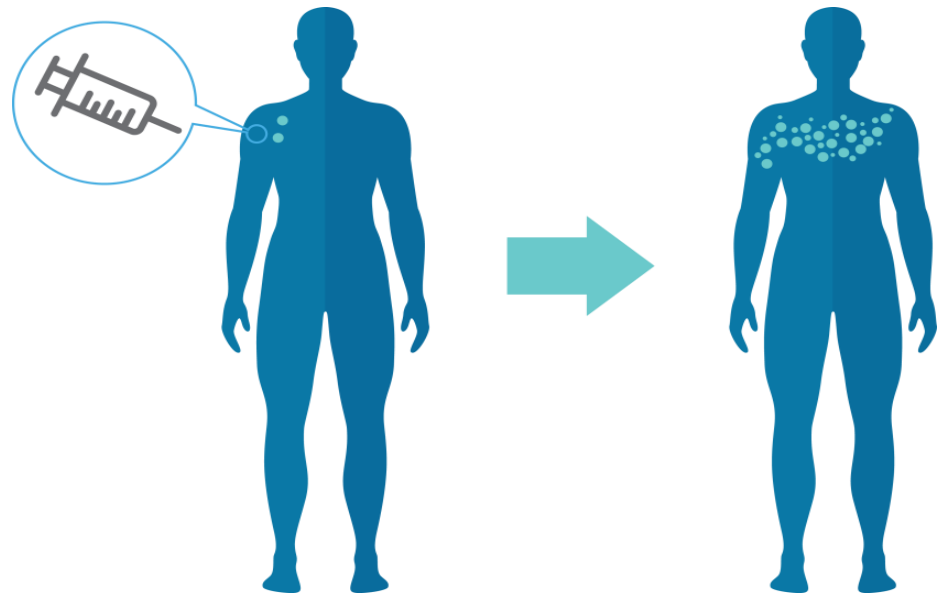
Biología de San Diego ayudará con ensayo de vacuna inteligente contra covid-19

10 jun. TriLink Biotechnologies se asoció con científicos del Reino Unido para probar si la vacuna es segura y efectiva.

Encontrar una vacuna que funcione contra covid-19 no será suficiente. Los científicos necesitarán hacer rápidamente una vacuna suficiente para cientos de millones, tal vez miles de millones, de personas.

¿Una solución de la biología de San Diego para este desafío de fabricación? Una vacuna que haga más de sí misma.

TriLink Biotechnologies se asoció con investigadores del Imperial College London para probar dicha vacuna en un ensayo que comenzará a mediados de junio. Si tiene éxito, el enfoque podría ayudar a llevar una vacuna contra covid-19 a una amplia franja de la población rápidamente, dice Anton McCaffrey, director



de ciencia e innovación emergente de TriLink.

"En este momento, todo el mundo quiere ir a la velocidad de la luz", dijo McCaffrey. "Es necesario saber que se puede hacer (una vacuna) a la escala necesaria para vacunar a una parte sustancial de la población".

Una vacuna de imitación

La vacuna de TriLink utiliza genes

que los virus normalmente utilizan para copiar su material genético. Solo que esta vez, esos genes ayudan a copiar una vacuna que centra la atención del sistema inmunológico en la superficie del nuevo coronavirus. Después de copiarse a sí misma en el curso de un par de semanas, la vacuna eventualmente sería eliminada del sistema de una persona, según McCaffrey.

Los genes virales que permiten que la vacuna se copie a sí misma también la hacen más grande y difícil de producir, pero los científicos no necesitarían hacer tanto.

La compañía estima que el enfoque reduce la cantidad de vacuna que cada persona necesitaría entre 25 y 50 veces.

Debido a que una vacuna autoreplicante se copia a sí misma de la misma manera que un virus, activaría las mismas alarmas provocadas por la infección.

Eso podría ser algo bueno, dice McCaffrey, ya que una respuesta antiviral llevaría a un contraataque inmunológico más fuerte.

Eso tendrá que ser demostrado por ensayos clínicos.

Los investigadores del Imperial College comenzarán un ensayo clínico de fase 1 para probar la seguridad de la vacuna a mediados de junio.

Si eso va bien, los científicos del Reino Unido llevarán a cabo un ensayo más amplio para probar si la vacuna protege contra covid-19.

TriLink puede producir suficientes vacunas para los ensayos clínicos. Pero McCaffrey dice que necesitaría construir nuevas instalaciones o licenciar su tecnología para hacer suficientes vacunas para uso global. TriLink Biotechnologies emplea a unas 200 personas y fue fundada en San Diego en 1996.

La biotecnológica de San Diego Arcturus Therapeutics está explorando una estrategia similar para la vacuna contra covid-19 en asociación con la autoridad sanitaria nacional de Singapur.

Fuente: The San Diego Union Tribune. Disponible en <https://bit.ly/2BiYd3>

La OMS descarta que haya vacuna contra el coronavirus en 2020

10 jun. La directora del Departamento de Salud Pública y de Ambiente en la Organización Mundial de la Salud (OMS), María Neira, ha asegurado que la vacuna contra el covid-19, la enfermedad que provoca el nuevo coronavirus, no va a estar disponible este año, si bien ha asegurado que en pocas semanas habrá "buenas noticias" sobre los tratamientos.

Durante los encuentros 'Conversaciones 2020', organizados por el Economista, Neira ha señalado que, "tal vez", a principios de 2021 ya se pueda comercializar una vacuna contra el nuevo coronavirus, aunque ha avisado de que hay que ser "realistas" y ver que sólo quedan

seis meses para terminar el año y que no va a dar tiempo a desarrollarla.

"Este año lo veo francamente difícil por mucha aceleración que haya y se tenga el máximo apoyo por parte de todos los países y de la OMS para obtenerla, pero hay muchos procesos que hay que seguir para asegurarse de cuando la vacuna salga al mercado sea segura", ha recalcado la dirigente de la OMS.

Ahora bien, respecto a los tratamientos, y después de que el organismo de Naciones Unidas haya retomado las investigaciones con el fármaco hidroxiquina, tras suspenderlas temporalmente por un posible



aumento del riesgo de muerte y enfermedades cardíacas, Neira ha asegurado que en unas pocas semanas se sabrá si aporta o no beneficio para los pacientes.

Finalmente, la directora del Departamento de Salud Pública y de Ambiente ha asegurado que los tratamientos que se están utilizando en la actualidad, y que están siendo valorados por la OMS, van a dar "buenas noticias" en un par de semanas.

Fuente: EL ESPAÑOL. Disponible en <https://bit.ly/2BfdGiB>



Triple golpe al coronavirus: pruebas, vacunas y medicamentos

10 jun. La lucha contra el coronavirus tiene tres frentes: diagnóstico, prevención y tratamiento. Los científicos rusos comentaron los avances del país en cada una de estas áreas en la reunión organizada por el Ministerio de Educación Superior y Ciencia de Rusia, la Academia Rusa de Ciencias y el portal Nauchnaya Rossiya (Rusia Científica).

El diagnóstico implica la introducción de sistemas de pruebas para detectar la presencia del coronavirus SARS-CoV-2 y la prueba de anticuerpos. La profilaxis consiste en desarrollo de diferentes variantes de vacunas contra este virus. El tratamiento incluye la creación de fármacos recombinantes de cuerpos específicos que neutralizan el virus, así como el desarrollo de fármacos antivirales con una actividad anti-SARS-CoV-2.

Diagnóstico avanzado

Un sistema de prueba para la determinación de inmunoenzimas de anti-SARS-CoV-2 fue creado por los científicos rusos del Instituto Engelhardt de Biología Molecular junto con el Instituto de Hematología del Ministerio de Sanidad.

El inmunógeno era la proteína S, que permite al coronavirus SARS-CoV-2 entrar en las células, específica para el SARS-CoV y el SARS-CoV-2. La proteína recombinante fue desarrollada en células de mamíferos.

"Los anticuerpos detectados son anticuerpos neutralizantes, y esto es muy importante", aclaró

Vladimir Chejonin, vicepresidente de la Academia Rusa de Ciencias.

Precisó que esta prueba permite detectar la inmunoglobulina G con una sensibilidad del 95% y una especificidad del 98%. La prueba ya se usa en la práctica.

Vacunas prometedoras

Rusia ya ha registrado ocho vacunas candidatas en fase de ensayo clínico que incluyen vacunas vectorizadas, inactivadas, de ADN y ARN.

Además en la etapa preclínica se encuentran 26 vacunas vectorizadas, 43 vacunas subunidades, 10 de ADN, cinco inactivadas y algunas otras.

Asimismo, según el académico, los científicos rusos desarrollan una inusual vacuna contra el coronavirus que se administra como un yogur.

"La vacuna mucosal contra SARS-CoV-2 se está desarrollando en el Instituto de la Medicina Experimental de San Petersburgo", señaló.

Según Chejonin, por métodos de ingeniería genética los científicos insertaron la proteína S, que permite al coronavirus SARS-CoV-2 entrar en las células, en los apéndices pilosos de las bacterias probióticas.

Chejonin precisó que esta vacuna puede ser inyectada como un producto de leche fermentada e informó que están en curso los estudios preclínicos de la vacuna.

Fármacos únicos

Los especialistas de la Academia Rusa de Ciencias han desarrollado

un fármaco que permite prevenir la penetración del coronavirus en el cuerpo. Los médicos militares de San Petersburgo ya llevan a cabo ensayos preclínicos.

Al mismo tiempo, los científicos intentan crear las drogas que bloquearían la interacción del nuevo coronavirus con las células epiteliales, mencionó Chejonin.

Estos fármacos han pasado las normas de laboratorio, y ahora se llevan a cabo ensayos preclínicos de esta droga.

Previsión innovadora

En Rusia se desarrolló un modelo descriptivo y predictivo de la pandemia COVID-19 basado en métodos de análisis inteligente de datos.

Se elaboraron modelos de aprendizaje automático sobre la base de los datos obtenidos de la vigilancia de la situación epidemiológica en San Petersburgo, informó.

Los científicos pudieron calcular la probabilidad de los picos de la enfermedad, el número de pacientes atendidos y dados de alta. Además, se elaboraron escenarios detallados teniendo en cuenta la intensidad de las medidas de contención, los modelos de mortalidad y la duración del tratamiento del paciente.

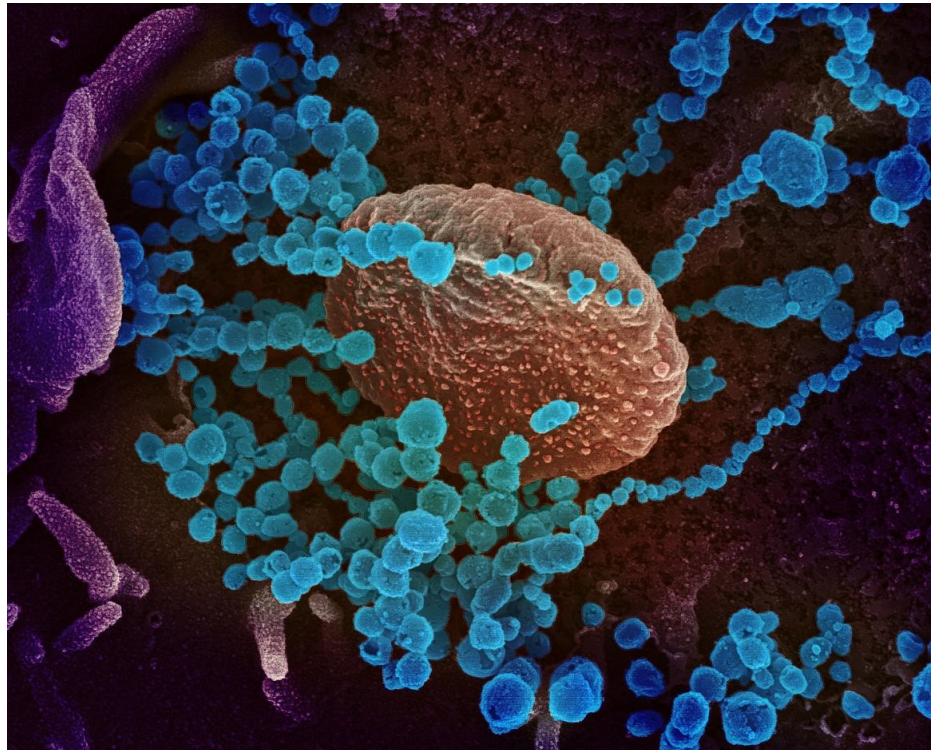
"Este es un trabajo muy importante, que permite hoy en día no solo predecir, sino también tomar decisiones sobre la atención médica de los pacientes y la construcción de nuevas instalaciones para su terapia", concluyó Chejonin.

Eli Lilly comienza la dosificación en el estudio de fase uno del segundo tratamiento potencial con anticuerpos contra la COVID-19

10 jun. Eli Lilly and Company (Indianápolis, Ind, EUA) anunció que su socio Shanghai Junshi Biosciences Co. Ltd. (Shanghái, China) dosificó al primer voluntario sano en un estudio sobre un posible tratamiento de anticuerpos neutralizantes diseñado para combatir la COVID-19.

La medicina en investigación, denominada JS016, es desarrollada conjuntamente por Junshi Biosciences y Lilly. JS016 es un anticuerpo neutralizante monoclonal completamente humano recombinante, que se ha modificado para disminuir la función efectora. JS016 se une específicamente al dominio de unión al receptor de la proteína spike de la superficie del SARS-CoV-2 y puede bloquear efectivamente la unión de los virus al receptor en la superficie de la célula huésped de ACE2.

Junshi Biosciences lidera su desarrollo en la Gran China, mientras que Lilly tiene derechos exclusivos en el resto del mundo y comenzará a dosificar pacientes en un estudio complementario de Fase 1 en los Estados Unidos en los próximos días. Ambos estudios de Fase 1 tienen como objetivo evaluar la seguridad, tolerabilidad, farmacocinética e inmunogenicidad de JS016 en participantes sanos que no han sido diagnosticados con COVID-19.



Este es el segundo anticuerpo neutralizante de Lilly en iniciar ensayos clínicos, después de LY-CoV555 que ingresó recientemente en la Fase 1 y actualmente está en prueba en pacientes hospitalizados con COVID-19.

Lilly planea un programa de desarrollo clínico que incluye una cartera de monoterapia y combinaciones de regímenes de anticuerpos (este último a menudo denominado “cócteles” de anticuerpos) para comprender cuál proporción la mejor eficacia y tolerabilidad en los pacientes. Estos cócteles incluirán JS016, LY-CoV555, así como anticuerpos adicionales actualmente en desa-

rollo preclínico. JS016 y LY-CoV555 se unen a diferentes epítopos en la proteína spike y, por lo tanto, expanden la diversidad de opciones para lograr la eficacia y evitar la resistencia.

“Hay muchas cosas que aún no sabemos sobre la COVID-19”, dijo Daniel Skovronsky, M.D., Ph.D., director científico de Lilly y presidente de Lilly Research Laboratories. “El mejor camino para aprender más sobre el potencial para neutralizar los anticuerpos, ya sea en monoterapia o en combinación, es a través de ensayos aleatorios cuidadosamente controlados. Esperamos los resultados de dichos ensayos en los próximos meses”.

Early Data From Coronavirus Vaccine Trials Are Encouraging, But There's Still A Long Way To Go

11 jun. Virologist Dave Wessner explains what we know about two promising coronavirus vaccines so far - and how much more we still need to know before they come onto the market.

In May, biotechnology company Moderna described positive preliminary results from a Phase I trial of its potential vaccine (mRNA-1273) for the Covid-19 virus. Just days later, researchers from CanSino Biologics published data about its own potential vaccine (Ad5-nCoV). In both cases, the news was met with great initial fanfare. However, although the news is encouraging, it doesn't mean that an effective vaccine will be available anytime soon. Here are the answers to some of the most pressing questions about the vaccines.

Do these two vaccines work the same?

No. As the name indicates, mRNA-1273 is an mRNA-based therapeutic. A molecule of mRNA encoding the novel coronavirus spike protein is encapsulated in a lipid nanoparticle. Upon injection, the mRNA enters that person's cells and is read by ribosomes, your cell's protein factories, in a similar fashion to reading code in a computer program. Having been "programmed," the ribosomes being making copies of the viral protein.

In contrast, Ad5-nCoV is a recombinant adenovirus-based vaccine. Basically, the gene encoding the SARS-CoV-2 spike protein is

added to an adenovirus (one of the types of viruses that cause the common cold) that's been genetically modified so it can't make you sick. Again, following intramuscular injection, the viral spike protein should be produced. In both cases, our immune system should recognize this foreign protein and begin manufacturing antibodies.

Hopefully, antibodies produced against the SARS-CoV-2 spike protein would inhibit the actual virus, if a person became infected, thereby preventing disease. However, as Phase I trials, neither of these studies addressed that crucial point.

What is a Phase I trial?

To address this question, let's look at what a Phase I study is not. Generally, Phase I trials do not examine efficacy. In other words, we won't know from these trials if the vaccines work. That's another question for another trial. Instead, researchers conducting these studies asked three more basic questions. Is the vaccine safe? Do participants develop antibodies to the Covid-19 virus? Can these antibodies inhibit the virus in the laboratory? For both the mRNA-1273 and Ad5-nCoV vaccines, the preliminary answer to all three questions appears to be, "yes."

It should be noted, though, that data from the mRNA-1273 trial have not been released. We have Moderna's press release, but little else. In contrast, data from the Ad5-nCoV trial have been published. It also should be noted that the most

encouraging part of the Moderna report is based on an early analysis of just eight "sentinel" participants. The trial has not yet been completed and a final analysis is not available.

What do we know about the clinical trials so far?

The Moderna study initially enrolled 45 participants between the ages of 18 and 55, whereas the CanSino Biologics study included 108 participants between the ages of 18 and 60. In both cases, the participants received one of three different doses of the candidate vaccine. For both vaccines, few of the participants experienced unusual adverse side effects. That's good news. And all of the participants developed antibodies. That's also good news. Perhaps most importantly, both groups reported evidence of participants developing neutralizing antibodies. In the Moderna study, all eight sentinel participants developed neutralizing antibodies. In the CanSino Biologics study, 50% of participants who received the low or intermediate vaccine dose and 75% of participants who received the high dose developed neutralizing antibodies.

What are neutralizing antibodies?

When exposed to a virus, our immune system typically produces a variety of antibodies that recognize different structural determinants, or antigens, associated with that virus. However, only certain antigens are used by the virus to

initiate an infection. Often, antibodies that bind to these determinants inhibit viral replication and, as a result, are called neutralizing antibodies. In contrast, non-neutralizing antibodies may bind to the virus, but do not inhibit its replication.

So, participants in both studies produced neutralizing antibodies. Does that mean that these individuals now are protected from future SARS-CoV-2 infections? No. To detect neutralizing antibodies, the scientists used *in vitro* assays. SARS-CoV-2 was mixed with serum from a participant or a control solution. The treated virus then was added to mammalian cells grown in the laboratory. The number of cells that became infected was determined. If fewer

infected cells were detected when the virus was mixed with serum from a participant, then that participant was deemed to have made neutralizing antibodies.

But what happens in the lab does not necessarily mirror what happens in the body.

Although the identification of neutralizing antibodies is encouraging, questions remain. Do these same antibodies offer protection? How long-lasting is this protection? Will antibodies elicited by this vaccine recognize related coronaviruses or mutants of this virus that may emerge? We know from our long history of trying to develop an HIV vaccine that the road can be bumpy. And a vaccine against SARS-CoV-2 may present additional challenges.

Animals given a candidate vaccine for the closely-related SARS virus exhibited a severe immunopathology after being exposed to the virus. In other words, rather than being protected by the vaccines, the animals' immune systems went into overdrive, making them sicker than they might have been otherwise. Before a vaccine for the Covid-19 virus becomes available, subsequent Phase II and Phase III trials will be needed to measure how well the vaccine works, to shed light on these important safety questions, and to identify any unforeseen challenges.

For now, the news is encouraging, but still preliminary.

Fuente: Forbes. Disponible en <https://bit.ly/2Ba1CXj>

Can old vaccines from science's medicine cabinet ward off coronavirus?

11 jun. Researchers think tuberculosis and polio vaccines could rev up the body's innate immune system against a new pathogen.

The old vaccines are oddities among the cutting-edge and targeted technologies being developed to combat the novel coronavirus. New vaccines aim to teach the body's immune system to recognize and destroy the coronavirus, but scientists are only now beginning to test them in people. Vaccines developed against TB and polio have already been used in millions of people and could offer a low-risk way to rev up the body's first line of defense — the

innate immune system — against a broad array of pathogens, including the coronavirus.

"This is the only vaccine in the world that can be given to combat covid-19 right now," said Jeffrey D. Cirillo, a professor of microbial pathogenesis and immunology at Texas A&M Health Science Center, who is leading a trial of the tuberculosis vaccine, called bacillus Calmette-Guérin and known by the shorthand BCG. The BCG vaccine, Cirillo noted, is already approved by the Food and Drug Administration and has a lengthy record of being used safely.

Scientists are betting on an un-

derappreciated facet of the body's immune system. Vaccines are designed to teach it to develop a memory of a particular pathogen. But over the years, vaccines that use live, weakened pathogens have been shown to have potent off-target effects, activating other components of the immune response to beat back other infections, including respiratory diseases.

The idea isn't necessarily that those vaccines could altogether prevent covid-19, the disease caused by the novel coronavirus, but that they might lessen the severity of disease and prepare the

innate immune system to fight off the virus for a short period of time.

Research comparing rates of coronavirus infections in countries that widely use the tuberculosis vaccine against those that do not initially drew attention to the idea that the inoculation could offer protection, spurring ongoing trials in the United States, the Netherlands and Australia.

A group of prominent researchers working to raise money to test the oral polio vaccine in 11,000 people described their ambitions in a paper published Thursday in the journal Science.

If shown effective, those vaccines could potentially provide protection against the second wave of coronavirus, which is likely to crest before a covid-specific vaccine is widely available.

Azra Raza, a professor of medicine at Columbia University Medical Center, said BCG can improve people's ability to fight off other pathogens, even for patients who are given the vaccine for another approved use, against bladder cancer.

Fuente: The Washington Post. Disponible en <https://wapo.st/3dJPQxg>

Surfacen, producto cubano eficaz en el tratamiento de adultos con COVID-19

11 jun. El surfacen, producto cubano para tratar el síndrome de dificultad respiratoria, ha sido utilizado, de manera satisfactoria, en el protocolo cubano a pacientes adultos con la COVID-19 en estado crítico o grave.

Nivian Montes de Oca Martínez, directora general del Centro Nacional de Sanidad Agropecuaria (Censa), explicó a la Agencia Cubana de Noticias que hoy la institución, gracias a la eficacia del producto en adultos con el nuevo coronavirus, ha aportado a varios foros internacionales de discusión científica artículos sobre sus beneficios, y cuándo y cómo utilizarlo.

El Surfacen se ha suministrado a pacientes intubados en la Unidad de Cuidados Intensivos, pero de manera muy temprana, lo que constituye un resultado significativo de la ciencia cubana, explicó la directiva.

Nunca un producto es el responsable máximo de la evolución positiva de los ingresados, sino



que la combinación del arsenal de medicamentos, la asistencia médica, así como los protocolos de manejo han arrojado estos buenos resultados, aclaró Montes de Oca Martínez.

Para obtener este surfactante se precisa de cerdos pequeños que son criados con una tecnología específica y diferenciados de aquellos que son para la producción de carne para el consumo de la población, pues solo deben alcanzar los 25 kilogramos de peso, refirió Odalys Uffo Reinoso, subdirectora general de Internacionali-

zación y Gestión Empresarial.

Estos animales son sacrificados y de sus pulmones se realiza un lavado, lo que constituye la materia prima fundamental para la elaboración del surfacen, un producto, que al ser ciento por ciento natural presenta la desventaja de que disminuye la rapidez con que se pueden obtener las dosis.

Las producciones de surfacen no recaen solo sobre el Censa, por ejemplo, intervienen el Ministerio de Agricultura, con unidades que son donantes de cerdos, y el

Centro de Biopreparados en la parte final del proceso: formulación y envasado de productos, una industria con altos estándares de calidad.

Al Censa le corresponde la obtención del instituto farmacéutico activo que son los lavados pulmonares con las condiciones necesarias como la concentración adecuada; además es labor del centro la comercialización de los lotes, destacó Uffo Reinosá.

El surfacen está registrado en el país desde 1995 y es aplicado principalmente en las terapias intensivas de los hospitales a pacientes en edad pediátrica, a niños con síndromes de dificultad

respiratoria y a neonatos con bajo peso; este año el Ministerio de Salud Pública tiene una demanda mínima de siete mil dosis.

Sin embargo, hasta el año pasado se presentaron dificultades con las materias primas fundamentales que se precisan para realizar los lavados pulmonares, situación que con el apoyo del Minsap y de BioCubaFarma ha tenido solución.

Al ser este un fármaco de gran interés para el país y de producción nacional, lo que garantiza soberanía tecnológica, se logró importar, a través de un sistema diferenciado, la materia prima suficiente para asegurar la elaboración de las dosis en el 2020.

Las producciones de surfacen deben mantenerse, e incluso en un futuro ampliarse puesto que se trabaja para su exportación y se conoce que, en su formulación actual, puede ser un transportador para que otras moléculas puedan llegar al pulmón y tratar otras afecciones.

Se evalúa, además, su eficacia para el asma bronquial y otras enfermedades que se producen en el sistema respiratorio, por lo que es necesario pensar con más fuerza en el encadenamiento productivo, sobre todo en la transferencia de tecnologías tan complicadas como la animal, señaló Montes de Oca Martínez.

Fuente: tv avileña. Disponible en <https://bit.ly/2Zf6nEs>

Sinovac and Butantan Join Efforts to Advance the Clinical Development of An Inactivated Vaccine for COVID-19 to Phase III

11 jun. Sinovac Biotech Ltd., (“Sinovac” or the “Company”) (NASDAQ: SVA), a leading provider of biopharmaceutical products in China, and Instituto Butantan, a leading Brazilian producer of immunobiologic products, today announced the signing of a clinical development collaboration agreement to advance the clinical trials of CoronaVac, Sinovac’s inactivated vaccine candidate against COVID-19 to Phase III.

Sinovac has made significant progress in the development of CoronaVac. Promising preclinical results regarding CoronaVac were recently published in the peer-reviewed academic journal

Science in an article stating that the vaccine candidate is safe and provides protection to rhesus macaques (monkeys) through an animal challenge study. Sinovac has received approval from China’s National Medical Products Administration (NMPA) to conduct Phase I/II human clinical trials in China to determine the vaccine candidate’s safety, tolerance, dosage and immunization schedule. In addition, Sinovac is constructing a commercial vaccine production plant in China that is expected to manufacture up to 100 million doses of CoronaVac annually.

Through this collaboration, Instituto Butantan will sponsor Phase III

clinical trials in Brazil. This is the first in a series of agreements expected to be completed between the parties to establish extensive collaboration that includes technology licensing, market authorization and commercialization of CoronaVac. In this way, Instituto Butantan can ensure that the Brazilian population has access to this vaccine.

Dr. Dimas Covas, Director of Instituto Butantan commented, “This pandemic is having a tragic impact worldwide and this distinguished alliance with Sinovac to conduct the last phase of the clinical trials will bring hope to have a vaccine in the short term. Butantan expects to support not only on the

clinical development, but also commercialization and manufacturing activities of CoronaVac in Brazil."

Dr. Ricardo Palacios, Clinical Research Medical Director of Instituto Butantan added, "Sinovac's CoronaVac is based on a well-known, reliable technology suited to being incorporated into existing public health immunization programs in Brazil. Current epidemiology in

Brazil and the experience of Butantan in clinical development will complement Sinovac's efforts allowing accelerated progress toward development of a safe and effective immunization against COVID-19."

Mr. Weidong Yin, Chairman, President and CEO of Sinovac, commented, "We are proud to take part in the fight against COVID-19, and we look forward to working with Instituto Butantan to help the

people of Brazil. Through this partnership, Sinovac will be able to further the unprecedented speed of developing CoronaVac without compromising our high safety standards and procedures. We also welcome this opportunity to further our commitment to developing vaccines for global use and to our mission of supplying vaccines to eliminate human diseases."

Fuente: DAILY JOURNAL. Disponible en <https://bit.ly/387EMcl>

Científicos de Florida descubren que mutación hace más eficaz al SARS-CoV-2

13 jun. Un equipo del Instituto de Investigación Scripps de Florida ha descubierto que una leve mutación genética en el coronavirus SARS-CoV-2 incrementa de manera significativa su capacidad para infectar las células, según un comunicado de la institución.

La viróloga Hyeryun Choe, autora principal del estudio, dijo que pudieron determinar en los sistemas de cultivo celular que "los virus con esa mutación son mucho más contagiosos que los que no la tienen".

Lo que hace la mutación D614G es incrementar hasta 4 ó 5 veces el número o la densidad de "espigas" funcionales existentes en la superficie viral y a la vez hacerlas más flexibles.

Las espigas, que le dan al virus su aspecto de corona, son precisamente las que le hacen capaz de infectar las células, apuntando

a los receptores celulares ACE2.

"Nuestros datos son muy claros, el virus se hace mucho más estable con la mutación", dijo Choe.

Según el comunicado del Instituto de Investigación Scripps, que tiene sede en Jupiter (sureste de Florida), la variante del SARS-CoV-2 que circuló en los primeros brotes no tenía la mutación D614G, que es ahora la variante dominante en gran parte del mundo.

Según Michael Farzan, coautor de la investigación y copresidente del Departamento de Inmunología y Microbiología de Scripps, ninguna de las secuencias del SARS-CoV-2 depositadas en la base de datos GenBank tenía la mutación.

En marzo ya aparecía en una de cada 4 muestras y en mayo en el 70 % de las muestras, señaló.

Choe y Farzan, que realizaron su investigación con virus inocuos diseñados para producir proteínas

claves del coronavirus, advierten que se necesitan estudios epidemiológicos adicionales para determinar si lo que ellos han comprobado sobre una mayor efectividad para contagiar las células a causa de la mutación sucede también en "el mundo real".

Ambos científicos han estudiado los coronavirus desde hace casi 20 años, desde que se registró el primer estallido de SARS, y en 2003 fueron los primeros en descubrir que el SARS apuntaba a los receptores ACE2 de las células, como hace el SARS-CoV-2.

Además de Choe y Farzan, también trabajaron en esta investigación respaldada por el Centro Nacional de Salud, los científicos Lizhou Zhang, Cody Jackson, Huihui Mou, Amrita Ojha, Erumbi Rangarajan y Tina Izzard, todos ellos del Instituto Scripps.

Fuente: Listín Diario. Disponible en <https://bit.ly/2BRwxEX>



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358. [Suspected adverse reaction to erysipelas vaccine in sheep.](#)
Bidewell C, Carson A, Diesel G, Floyd T. *Vet Rec*. 2020 Jun 13;186(18):610-611. doi: 10.1136/vr.m1482.
PMID: 32527900

Patentes registradas en Patent Google

Estrategia de búsqueda: *Vaccine in the title or abstract AND 20200609:20200615 as the publication date*

[Dose reduced povlio virus vaccine compositions and methods for its production](#)

WO EP US CN JP KR AU BR CA CU EA MX PE PH SG [EP3663396A1](#) Rajeev Mhalasakant DHERE Serum Institute Of India Private Limited

Priority 2014-10-07 • Filed 2015-10-06 • Published 2020-06-10

[Pertussis booster vaccine](#)

[WO2020117618A1](#) Nicolas Burdin Sanofi Pasteur Inc.

Priority 2018-12-05 • Filed 2019-11-29 • Published 2020-06-11

[Hepatitis b nanoparticle-based vaccine for influenza virus](#)

WO EP CN EP3661549A1 Audray K. HARRIS The U.S.A. As Represented By The Secretary, Department Of Health And Human ...

Priority 2017-08-02 • Filed 2018-08-02 • Published 2020-06-10

[Swine mycoplasmal pneumonia attenuated live vaccine and use thereof](#)

WO EP US [EP3098301B1](#) Guoqinq SHAO Jiangsu Academy of Agricultural Sciences

Priority 2014-01-26 • Filed 2014-01-26 • Granted 2020-06-10 • Published 2020-06-10

[A vaccine for protection against streptococcus suis](#)

WO EP CN EP3661546A1 Antonius Arnoldus Christiaan Jacobs Intervet International B.V.

Priority 2017-08-03 • Filed 2018-08-02 • Published 2020-06-10

[A vaccine comprising a pcv2 orf2 protein of genotype 2b](#)

WO EP EP3661548A1 Melanie SNO Intervet International B.V.

Priority 2017-08-03 • Filed 2018-08-02 • Published 2020-06-10

[Vaccine composition](#)

EP US CN JP KR CA IN RU [EP3662927A2](#) Daisuke Asari Nitto Denko Corporation

Priority 2013-02-05 • Filed 2014-01-29 • Published 2020-06-10

[A recombinant koi herpesvirus \(khv\) and a diva vaccine for preventing and/or ...](#)

EP [EP3662929A1](#) Walter Fuchs IDT Biologika GmbH

Priority 2018-12-07 • Filed 2018-12-07 • Published 2020-06-10

[Nanoparticle platform for antibody and vaccine delivery](#)

WO EP CA EP3661968A1 Jean-Philippe Julien The Hospital for Sick Children

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

[Cross-protective arenavirus vaccines and their method of use](#)

WO EP US CN JP KR AU CA ES [EP3662935A1](#) Kate Broderick Inovio Pharmaceuticals, Inc.

Priority 2011-07-11 • Filed 2012-07-11 • Published 2020-06-10

A DNA plasmid **vaccine** of any one of claims 1 to 4 for use in a method of inducing a protective immune response against LASV in a subject. The DNA plasmid **vaccine** for use as claimed in claim 5 wherein the DNA **vaccine** is administered by electroporation. The DNA plasmid **vaccine** for use as claimed in ...

[Mesoporous silica compositions for modulating immune responses](#)

WO EP US CN JP AU CA DK HK HU LT PT [EP3662896A1](#) Jaeyun Kim President and Fellows of Harvard College

Priority 2012-04-16 • Filed 2013-04-16 • Published 2020-06-10

A composition comprising mesoporous silica rods comprising an immune cell recruitment compound and an immune cell activation compound, and optionally comprising an antigen such as a tumor lysate. The composition is used to elicit an immune response to a **vaccine** antigen.

[Induction of protective immunity against antigens](#)

WO EP CN EP3661547A1 Roy Curtiss Iii University of Florida Research Foundation, Incorporated

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

85. Clifton-Hadley FA, Breslin M, Venables LM, Sprigings KA, Cooles SW, Houghton S, Woodward MJ. 2002. A laboratory study of an inactivated bivalent iron restricted Salmonella enterica serovars Enteritidis and Typhimurium dual **vaccine** against Typhimurium challenge in chickens. Vet Microbiol 89: 167 ...

[Compositions comprising jak inhibitors and haart drugs for use in the ...](#)

WO EP US CN AU BR CA MX [EP2785184B1](#) Christina GAVEGNANO Emory University

Priority 2011-11-30 • Filed 2012-11-30 • Granted 2020-06-10 • Published 2020-06-10

d) AZT, and e) a NNRTI or a protease inhibitor, optionally wherein the NNRTI is Sustiva or the protease inhibitor is Kaletra, or wherein the JAK inhibitor, HAART, and **vaccine** or immunostimulatory compound are administered in combination or alternation. The medicament for use according to claim 3, ...

[Surrogate therapeutic endpoint for anti-ctla-4 based immunotherapy of disease](#)

WO EP US CN JP AU CA IL MX NO NZ ZA [EP1639010B1](#) Israel Lowy E. R. Squibb & Sons, L.L.C.

Priority 2003-05-30 • Filed 2004-05-28 • Granted 2020-06-10 • Published 2020-06-10

). In these instances, non-immunogenic tumors, such as the B16 melanoma, have been rendered susceptible to destruction by the immune system. The tumor cell **vaccine** may also be modified to express other immune activators such as IL2, and costimulatory molecules, among others. The study of gene ...

[Novel peptides and combination of peptides for use in immunotherapy against ...](#)

WO EP US CN JP KR AU BR CA CL CO CR EA GB HK IL MA MX PE PH ZA [EP3322717B1](#) Heiko Schuster immatics biotechnologies GmbH

Priority 2015-07-15 • Filed 2016-07-14 • Granted 2020-06-10 • Published 2020-06-10

In an especially preferred embodiment, the pharmaceutical compositions comprise the peptides as salts of acetic acid (acetates), trifluoro acetates or hydrochloric acid (chlorides). Preferably, the medicament of the present invention is an immunotherapeutics such as a **vaccine**. It may be ...

[Formulation and delivery of modified nucleoside, nucleotide, and nucleic acid ...](#)

WO EP US [EP2971010B1](#) Stephen G. HOGGE ModernaTX, Inc.

Priority 2013-03-14 • Filed 2014-03-14 • Granted 2020-06-10 • Published 2020-06-10

An mRNA for use in a therapeutic method of increasing the duration of protein expression from said mRNA in a mammalian cell or tissue of a mammal, said method comprising administering said mRNA via electroporation; wherein administration comprises the steps of (a) injecting the mammal with mRNA, ...

Combination therapies

WO EP US CN JP KR AU BR CA EA MX [EP3662903A2](#) Zhu Alexander CAO Novartis AG

Priority 2014-10-03 • Filed 2015-10-02 • Published 2020-06-10

Nestle, F. et al. (1998) Nature Medicine 4: 328-332). DCs may also be transduced by genetic means to express these tumor antigens as well. DCs have also been fused directly to tumor cells for the purposes of immunization (Kugler, A. et al. (2000) Nature Medicine 6:332-336). As a method of ...

Means and methods for the prediction of treatment response of a cancer patient

WO EP US [EP2619576B1](#) Niels Grabe Niels Grabe

Priority 2010-09-24 • Filed 2011-09-20 • Granted 2020-06-10 • Published 2020-06-10

tositumomab-iodine 131; treosulfan; tretinoin; trilostane; trimetrexate; triptorelin; tumor necrosis factor alpha natural; ubenimex; bladder cancer vaccine; Maruyama vaccine; melanoma lysate vaccine; valrubicin; verteporfin; virulizin; zinostatin stimalamer; abarelix; AE 941 (Aeterna); ambamustine;

Compositions comprising bacterial strains

WO EP US CN JP KR AU CA CL CO DK EA ES HR HU IL LT MD MX PE PL PT RS SG SI TW
[EP3662917A1](#) Alex STEVENSON 4D Pharma Research Limited

Priority 2015-06-15 • Filed 2016-06-15 • Published 2020-06-10

21. A food product comprising the composition of any preceding embodiment, for the use of any preceding embodiment. 22. A vaccine composition comprising the composition of any preceding embodiment, for the use of any preceding embodiment. 23. A method of treating or preventing a disease or ...

Human anti-tau antibodies

WO EP US CN JP KR AU BR CA EA HK IL MX PH SG ZA [EP2935326B1](#) Paul H. Weinreb Biogen MA Inc.

Priority 2012-12-21 • Filed 2013-12-20 • Granted 2020-06-10 • Published 2020-06-10

In the wake of the success of Abeta-based **immunization** therapy in transgenic animal models, the concept of active immunotherapy was expanded to the tau protein. Active vaccination of wild type mice using the tau protein was however found to induce the formation of neurofibrillary tangles, axonal ...

[Aav vectors targeted to oligodendrocytes](#)

WO EP US CN JP AU CA [EP2900686B1](#) Thomas MCCOWN The University of North Carolina At Chapel Hill

Priority 2012-09-28 • Filed 2013-09-27 • Granted 2020-06-10 • Published 2020-06-10

The virus vectors of the invention can further be administered to a subject to elicit an immunogenic response (e.g., as a **vaccine**). Typically, vaccines of the present invention comprise an effective amount of virus in combination with a pharmaceutically acceptable carrier. Optionally, the dosage is ...

[Continuous processing methods for biological products](#)

WO EP US CN JP KR BR [EP2649016B1](#) Thomas C. Ransohoff Pall Corporation

Priority 2010-12-06 • Filed 2011-12-06 • Granted 2020-06-10 • Published 2020-06-10

... , Worcester, MA (US)), have made it possible to convert separation process steps that were traditionally operated in batch mode into SMB processes that can be run continuously, taking a clarified feed and purifying in a continuous fashion a target molecule such as a monoclonal antibody or a **vaccine** ...

[Methods and compositions for production of vaccinia virus](#)

WO EP US CN JP KR AU BR CA [EP2739293B1](#) David Kirn SillaJen Biotherapeutics, Inc.

Priority 2011-08-05 • Filed 2012-08-03 • Granted 2020-06-10 • Published 2020-06-10

P. D. Ellner, Infection 26, 263 (Sep-Oct, 1998). 12. F. Fenner, Prog Med Virol 23, 1 (1977). 13. A. W. Artenstein et al., **Vaccine** 23, 3301 (May 9, 2005). 14. T. P. Monath et al., Int J Infect Dis 8 Suppl 2, S31 (Oct, 2004). 15. Z. S. Guo, D. L. Bartlett, Expert Opin Biol Ther 4, 901 (Jun, 2004 ...

[Methods for the treatment of ocular disease in human subjects](#)

WO EP US CN JP KR AU BR CA EA HK IL MX PH SG [EP2916827B1](#) Vladimir ZARNITSYN Clearside Biomedical Inc.

Priority 2012-11-08 • Filed 2013-11-08 • Granted 2020-06-10 • Published 2020-06-10

In one embodiment, the drug is selected from a suitable oligonucleotide (e.g ., antisense oligonucleotide agents), polynucleotide (e.g ., therapeutic DNA), ribozyme, dsRNA, siRNA, RNAi, gene therapy vectors, and/or **vaccine**. In a further embodiment, the drug is an aptamer (e.g ., an oligonucleotide or ...

[Compositions and methods for immunotherapy](#)

WO EP US CN JP KR AU BR CA HK IL MX PH RU SG [EP2961831B1](#) Renier J. BRENTJENS Memorial Sloan Kettering Cancer Center

Priority 2013-02-26 • Filed 2014-02-26 • Granted 2020-06-10 • Published 2020-06-10

). Augmenting the immunogenicity of cancer cells through the CD40L/CD40 pathway has been shown to induce an endogenous anti-tumor response in previously published **vaccine** studies using the infusion of autologous CLL tumor cells transduced with an adenovirus vector encoding CD40L (Ad-CD40L CLL cells) ...

[Magnet recycling to create nd-fe-b magnets with improved or restored magnetic ...](#)

WO EP US CN JP AU BR HK TW [EP3011573B1](#) Miha Zakotnik Urban Mining Technology Company, LLC

Priority 2013-06-17 • Filed 2014-06-17 • Granted 2020-06-10 • Published 2020-06-10

Magnet coatings may be completely removed by cleaning the surfaces with an abrasive jet that removes a portion, or all, of the surface of the magnets by ablation. In some implementations, the abrasive can be steel **shot**, tungsten carbide, ceramic, or steel grit. The size of the particles may be ...

[Activation of resident memory t cells for the treatment of cancer](#)

WO EP CA EP3661550A2 Vaiva D. VEZYS Regents of the University of Minnesota

Priority 2017-08-03 • Filed 2018-08-03 • Published 2020-06-10

1 1. The method of any one of the preceding claims, wherein the composition comprises at least one antigenic peptide from a **vaccine**. 12. The method of claim 1 1, wherein the **vaccine** is selected from the group consisting of a chickenpox **vaccine**, a polio **vaccine**, a German measles **vaccine**, a mumps ...

[Crystalline forms of ethyl \(\(s\)-\(\(\(\(2r,5r\)-5-\(6-amino-9h-purin-9-yl\)-4-fluoro- ...](#)

WO EP US AR TW EP3661937A1 Olga Viktorovna Lapina Gilead Sciences, Inc.

Priority 2017-08-01 • Filed 2018-07-30 • Published 2020-06-10

[0203] Examples of HIV vaccines include peptide vaccines, recombinant subunit protein vaccines, live vector vaccines, DNA vaccines, CD4-derived peptide vaccines, **vaccine** combinations, rgp120 (AIDSVAX), ALVAC HIV (vCP1521)/AIDSVAX B/E (gp120) (RV144), monomeric gp120 HIV-1 subtype C **vaccine**, Remune, ...

[Anti-il-17 antibodies](#)

WO EP US CN JP KR AU BR CA CY DK EA ES HK HR HU IL LT LU NO PL PT RS SI [EP1963368B3](#)
Barrett Allan Eli Lilly And Company

Priority 2005-12-13 • Filed 2006-12-05 • Granted 2020-06-10 • Published 2020-06-10

Other suitable methods of producing or isolating antibodies of the invention, including human or artificial antibodies, can be used, including, for example, methods which select a recombinant antibody (e.g., single chain Fv or Fab) from a library, or which rely upon **immunization** of transgenic ...

[Human antigen binding proteins that bind beta-klotho, fgf receptors and ...](#)

WO EP US CN JP KR AR AU BR CA CL CR CY DK EA ES HR HU IL LT MA ME MX NZ PE PL PT RS SG
SI TN TW UA UY ZA [EP3202787B1](#) Shaw-Fen Sylvia Hu Amgen Inc.

Priority 2009-12-07 • Filed 2010-12-03 • Granted 2020-06-10 • Published 2020-06-10

In one embodiment, all of the variable and constant domains are derived from human immunoglobulin sequences (a fully human antibody). These antibodies can be prepared in a variety of ways, examples of which are described below, including through the **immunization** with an antigen of interest of a ...

[Anti-influenza antibody](#)

WO EP US CA [EP2931747B1](#) Xavier Saelens Vib Vzw

Priority 2012-12-11 • Filed 2013-12-11 • Granted 2020-06-10 • Published 2020-06-10

Example 2: **Immunization** and VHH phage library construction N1rec was next used as an immunogen for the generation and selection of NA-specific VHH. An alpaca (Vicunia pacos) was immunized at day 0 with 125 µg of N1rec, followed by 6 weekly boosts. One week after the last **immunization**, blood was ...

[Heterocyclic compound and application thereof](#)

WO EP [EP3663281A1](#) Tatsuhiko Fujimoto Takeda Pharmaceutical Company Limited

Priority 2017-08-03 • Filed 2018-08-02 • Published 2020-06-10

Examples of the antibody drug and **vaccine** preparation include **vaccine** preparation against angiotensin II, **vaccine** preparation against CETP, CETP antibody, antibody against TNF α antibody and other cytokines, amyloid β **vaccine** preparation, **vaccine** for type 1 diabetes (e.g., DIAPEP-277 of Peptor), ...

[Genetic products which are differentially expressed in tumours and use thereof](#)

WO EP US JP AU CA DE ES [EP3095791B1](#) Özlem TÜRECI BioNTech SE

Priority 2003-09-10 • Filed 2004-09-10 • Granted 2020-06-10 • Published 2020-06-10

(1) In the first case, peptides conjugated to KLH (keyhole limpet hemocyanin) (length: 8-12 amino acids) are synthesized using a standardized in vitro method and these peptides are used for **immunization**. As a rule, 3 **immunizations** are carried out with a concentration of 5-1000 μ g / **immunization**.

[Nuclear transport modulators and uses thereof](#)

WO EP US CN JP KR AU BR CA CL CO CY DK EA ES HK HR HU IN LT MX NZ PE PL PT RS SG SI UA [EP3663291A1](#) Vincent P. SANDANAYAKA Biogen MA Inc.

Priority 2012-05-09 • Filed 2013-05-09 • Published 2020-06-10

... . Contemporary methods for generating an immune response against tumors include intravesicular BCG immunotherapy for superficial bladder cancer, prostate cancer **vaccine** Provenge, and use of interferons and other cytokines to induce an immune response in renal cell carcinoma and melanoma patients.

[Means and methods for determining t cell recognition](#)

WO EP US JP AU CA [EP3152569B1](#) Carsten LINNEMANN AIMM Therapeutics B.V.

Priority 2014-06-05 • Filed 2015-06-05 • Granted 2020-06-10 • Published 2020-06-10

Preferably, T cells are used that are from an individual who has been exposed to said pathogen before, so that memory T cells will be present. If one or more test peptides appear to be recognized by T cells, these peptides are candidates for a **vaccine** against said pathogen. Again, the sensitivity ...

[Virucidal disinfection composition](#)

EP PL [EP1685854B2](#) Andreas Arndt B. Braun Medical AG

Priority 2005-01-28 • Filed 2006-01-27 • Granted 2020-06-10 • Published 2020-06-10

) and is effective within a minute against adenovirus type 5, strain Adenoid 75, papovavirus [Simianvirus 40 (SV40), strain 777], poliovirus (polio **vaccine** strain type I, strain LSc-2ab) and vaccinia virus (strain Elstree). In addition, it was found that the agents according to the invention are ...

[Method of culturing segmented filamentous bacteria in vitro](#)

WO EP US CN JP AU CA [EP3237602B1](#) Gérard EBERL Institut Pasteur

Priority 2014-12-23 • Filed 2015-12-23 • Granted 2020-06-10 • Published 2020-06-10

Said SFB strain can be in the form of a filament, an intracellular offspring or a spore, preferably a spore. The terms "immunogenic composition" and "**vaccine** composition" are used interchangeably herein. In some embodiments, the present invention relates to the genetically modified SFB strain as ...

[Antibody binding active alpha-synuclein](#)

WO EP CN KR AU CA EP3661961A1 Ariel Louwrier Stressmarq Biosciences Inc.

Priority 2017-08-02 • Filed 2018-08-02 • Published 2020-06-10

WHAT IS CLAIMED IS: 1. A monoclonal antibody 2F11 as deposited under ATCC PTA-124174. 2. A composition comprising the monoclonal antibody 2F11 defined in claim 1, in a pharmaceutically acceptable carrier, adjuvant, vehicle or excipient. 3. A **vaccine** comprising the monoclonal antibody 2F11 defined ...

[Pd-1 promoter methylation in cancer](#)

WO EP US CN JP KR AR AU CA IL MX [EP3303619B1](#) Edward KADEL H. Hoffnabb-La Roche Ag

Priority 2015-05-29 • Filed 2016-05-27 • Granted 2020-06-10 • Published 2020-06-10

). In the hybridoma method, a mouse or other appropriate host animal, such as a hamster, is immunized as herein described to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the protein used for **immunization**. Alternatively, lymphocytes may be ...

[System and method for peer referencing in an online computer system](#)

WO EP US CA [EP3663999A1](#) Qin YE Medversant Technologies, LLC

Priority 2010-02-05 • Filed 2011-02-07 • Published 2020-06-10

FIG. 4 is an exemplary screen **shot** of a screen listing peers that are available for inviting into a user's peer network according to one embodiment of the invention. According to one embodiment, the resulting peers are peers who are in the user's virtual professional network who are deemed to be ...

[Unmanned aircraft system with swappable components](#)

WO EP US AU EP3661847A1 Paul Perry Zipline International Inc.

Priority 2017-08-01 • Filed 2018-07-31 • Published 2020-06-10

For instance, if the payload is a **vaccine**, then the payload inventory management system may provide a refrigerated storage container for **vaccine** doses and may monitor and report the temperature in the storage container and the number of doses stored in the container to the logistics system 2008. ...

[Novel sulfonamide carboxamide compounds](#)

WO EP CN KR AU CA IL SG TW UY EP3661925A1 Matthew Cooper Inflazome Limited

Priority 2017-07-07 • Filed 2018-07-04 • Published 2020-06-10

modification (e.g. protection of the 2'-OH with a methyl group or replacement of the 2'-OH by -F or -N 3). In some embodiments, the one or more cancer vaccines are selected from an HPV **vaccine**, a hepatitis B **vaccine**, Oncophage, and/or Provenge. In some embodiments, the one or more immunomodulatory ...

[Microfluidic systems with capillary pumps](#)

WO EP EP3661649A1 Jaroslav Belotserkovsky Katholieke Universiteit Leuven

Priority 2017-08-04 • Filed 2018-08-06 • Published 2020-06-10

Figure 54: A pump activation and (micro)needle application "button". In one embodiment, the (iSIMPLE) microneedle drug or **vaccine** delivery device is designed such that the physical force, such as a finger push, that is applied used to activate the propulsion pump is also useful to provide the ...

[Antibodies binding to tumour associated carbohydrate antigens, pharmaceutical ...](#)

WO EP US CN JP KR AU BR CA CL IL PH RU SG TW ZA [EP3662928A1](#) Jiann-Shiun Lai OBI Pharma Inc.

Priority 2014-04-10 • Filed 2015-04-10 • Published 2020-06-10

An "effective amount," as used herein, refers to a dose of the **vaccine** or pharmaceutical composition that is sufficient to reduce the symptoms and signs of cancer, such as weight loss, pain and palpable mass, which is detectable, either clinically as a palpable mass or radiologically through ...

[Therapeutic agent or prophylactic agent for dementia](#)

WO EP US CN JP KR AR AU BR CA DK ES HK HR HU IL LT MX NZ PH PL PT RS RU SG SI TW ZA
[EP3662931A1](#) Hiroshi Mori Osaka City University

Priority 2012-05-31 • Filed 2013-05-30 • Published 2020-06-10

and Fig. 19 . To summarize, passive **immunization** with anti-pSer413 monoclonal antibody improved memory impairment in the model mice to a level of 50% or greater compared to non-Tg. The results of (4-1) and (4-2) confirmed a drug effect for pSer413 epitope monoclonal antibody even at a dose of 0.1 mg ...

[Biomimetic nanomaterials and uses thereof](#)

WO EP EP3661563A1 Yizhou Dong The Ohio State Innovation Foundation

Priority 2017-07-31 • Filed 2018-07-31 • Published 2020-06-10

In certain embodiments, the drug is an antibiotic, anti-viral agent, anesthetic, steroidal agent, anti-inflammatory agent, anti-neoplastic agent, anti-cancer agent, antigen, **vaccine**, antibody, decongestant, antihypertensive, sedative, birth control agent, progestational agent, anticholinergic, ...

[Methods and systems for authenticating and tracking objects](#)

WO EP US CN BR CA IN MX [EP2761538B1](#) Nabil M. Lawandy Spectra Systems Corporation

Priority 2011-09-29 • Filed 2012-09-11 • Granted 2020-06-10 • Published 2020-06-10

The combination of authentication technology and QR codes can track and authenticate pharmaceuticals and other packaged products anywhere in the world. Figures 6A and 6B show a syringe that is preloaded with a **vaccine**. In Figure 6A the syringe has been coded by exposing it to the code forming ...

[Bicyclic peptide ligands specific for cd137](#)

WO EP CN EP3661948A1 Lihong CHEN BicycleTx Limited

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

... , where cross-reactivity with homologues or paralogues needs to be carefully controlled. In some applications, such as **vaccine** applications, the ability to elicit an immune response to predetermined ranges of antigens can be exploited to tailor a **vaccine** to specific diseases and pathogens.

[Direct expression of antibodies](#)

WO EP US [EP3663315A1](#) Andrew Geall Novartis AG

Priority 2014-10-29 • Filed 2015-10-28 • Published 2020-06-10

The present invention relates to methods exogenous nucleic acid molecules, such as RNA, that encode an antibody or antigen-binding fragment of an antibody, and optionally encode a cellular modulation factor and/or a potentiation factor. The cellular modulation factor is a factor that results in ...

[Method for classification of a sample on the basis of spectral data and ...](#)

WO EP US CN JP IN NL RU [EP2836958B1](#) René Raymond PARCHEN BiosparQ B.V.

Priority 2012-04-10 • Filed 2013-04-10 • Granted 2020-06-10 • Published 2020-06-10

The above example illustrates the problems of conventional methods of analyzing spectral data. These methods are incapable of dealing with single particle spectra directly, since they do not take the above **shot-to-shot** variations into account. Furthermore, they are incapable of dealing with ...

[Pharmaceutical product comprising mite allergen extract\(s\) and a method for the ...](#)

WO EP US CN JP KR AU CA MX RU ZA [EP3662916A1](#) Heather Michelle Webster ALK-Abelló A/S

Priority 2010-06-03 • Filed 2011-06-03 • Published 2020-06-10

The invention relates to a pharmaceutical product comprising an allergen extract or an allergoid thereof for the treatment and/or prevention of allergy and allergic asthma caused by house dust mites, which extract comprises at least one extract of mite bodies selected from the following groups a)-b) ...

[Device for simulating a mortar](#)

WO EP US KR AU BR CA DE SG [EP3017267B1](#) Ernst Christians Rheinmetall Defence Electronics GmbH

Priority 2013-07-03 • Filed 2014-07-03 • Granted 2020-06-10 • Published 2020-06-10

... GNSS antennas can also be swiveled around at least one axis and can be fixed in a preferred position. The operation of the original weapon for firing the **shot** is not changed. The firing of the **shot** is detected on the mortar firing pin, the operation does not differ from the real firing of the **shot**.

[Medical imaging apparatus and control method thereof](#)

WO EP US KR [EP3197363B1](#) Min-Cheol Park Samsung Electronics Co., Ltd.

Priority 2014-09-26 • Filed 2015-09-25 • Granted 2020-06-10 • Published 2020-06-10

A control method for a medical imaging apparatus (200), the control method comprising: radiating (1210) X-rays onto an object and onto a calibration phantom (215), according to a first irradiating condition for a pre-**shot**; and detecting (1220) the X-rays having passed through the object and through ...

[A method for diagnosing or monitoring kidney function](#)

WO EP US CN JP CA DK ES HK PL RU TR [EP3361260B1](#) Andreas Bergmann sphingotec GmbH

Priority 2012-10-02 • Filed 2013-10-01 • Granted 2020-06-10 • Published 2020-06-10

All antibodies bound the MRPENK peptide, comparable to the peptides which were used for **immunization**. Except for NT-MRPENK-antibody (10% cross reaction with EEDDSLANSDDLK), no antibody showed a cross reaction with MR-PENK fragments not used for **immunization** of the individual antibody. Pro- ...

[Medical hollow needle assembly and method for manufacturing hollow needle](#)

WO EP US CN JP [EP3045194B1](#) Tetsuya Ooyauchi Terumo Kabushiki Kaisha

Priority 2013-09-11 • Filed 2014-09-10 • Granted 2020-06-10 • Published 2020-06-10

illustrates a syringe 1 with a fixed needle according to an embodiment of the present invention (medical hollow needle assembly). The syringe 1 with a fixed needle is used as a medical syringe for injecting a drug solution such as **vaccine** into a living body such as a human body, and includes a ...

[Compositions and methods for modulating an immune response](#)

WO EP US JP CA [EP3663763A1](#) Richard S. Blumberg The Brigham and Women's Hospital, Inc.

Priority 2013-11-26 • Filed 2014-11-25 • Published 2020-06-10

... nodules was carried out following lung inflation and fixation in 10% formalin. For DC **immunization** experiments, DC were isolated by collagenase digestion from the lung and draining lymph nodes of metastasis-bearing donor mice and 1×10^6 DC were injected s.c. into the hind footpad of recipient mice.

[Large diameter slag wool, composition and method of making same](#)

WO EP US CA MX TW [EP3475233B1](#) Wenqi Luan USG Interiors, LLC

Priority 2016-06-22 • Filed 2017-06-19 • Granted 2020-06-10 • Published 2020-06-10

A slag wool produced using the composition of any one of claims 1 to 3, wherein the slag wool has a fiber diameter in a range of 4.0 microns to 10.0 microns. The slag wool of claim 4, wherein the slag wool has a **shot** content of 50% or less, 45% or less, 40% or less. A method for the manufacture of ...

[BENZO\[b\]THIOPHENE STING AGONISTS FOR CANCER TREATMENT](#)

WO EP AU CA EP3661498A1 Saso CEMERSKI Merck Sharp & Dohme Corp.

Priority 2017-08-04 • Filed 2018-07-30 • Published 2020-06-10

Adjuvants, such as aluminum hydroxide or aluminum phosphate, can be added to increase the ability of the **vaccine** to trigger, enhance, or prolong an immune response. Additional materials, such as cytokines, chemokines, and bacterial nucleic acid sequences, like CpG, a toll-like receptor (TLR) 9 ...

[Bcma monoclonal antibody-drug conjugate](#)

WO EP US CN KR AU CA CO SG TW EP3661963A1 Krista KINNEER MedImmune, LLC

Priority 2017-08-01 • Filed 2018-07-31 • Published 2020-06-10

EXAMPLE 1 [0061] This example describes the generation of a monoclonal antibody directed against B- cell maturation antigen (BCMA). [0062] Following the RIMMS **immunization** regime described in Kilpatrick et al. , Hybridoma, 16(4): 381-389 (1997), six week old female Ablexis transgenic mice (Ablexis, ...

[Generating an environment map around a mobile object with high accuracy](#)

WO EP JP EP3662446A1 Takuto Motoyama Sony Corporation

Priority 2017-08-02 • Filed 2018-07-30 • Published 2020-06-10

<Superimposition of group of 3D points as measurement result of LiDAR on image> A measurement result of the LiDAR 301 is information indicating a group of points in a 3D space, and the measurement result is superimposed on an image **shot** by the camera 302 thereby to generate a distance image.

[Medical device and method for limiting the use of the medical device](#)

WO EP US CN JP [EP2780059B1](#) Ilona Eggert Sanofi-Aventis Deutschland GmbH

Priority 2011-11-18 • Filed 2012-11-15 • Granted 2020-06-10 • Published 2020-06-10

wherein in one embodiment the pharmaceutically active compound has a molecular weight up to 1500 Da and/or is a peptide, a proteine, a polysaccharide, a **vaccine**, a DNA, a RNA, an enzyme, an antibody or a fragment thereof, a hormone or an oligonucleotide, or a mixture of the above-mentioned ...

[Compact high mechanical energy storage and low trigger force actuator for the ...](#)

WO EP AU CA EP3661587A1 Pierre Armand Vincent LEMAIRE Vaxxas Pty Limited

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

[0229] In an alternate embodiment of the present applicator devices of the invention the microprojections of the microprojection array may be uncoated and the membrane may be coated by a substance such as a **vaccine**. In this embodiment the applicator pushes the microprojections of the ...

[Veterinary product](#)

WO EP CN AU CA EP3661603A1 Jeronimo Carnes Sanchez Laboratorios LETI, S.L. Unipersonal

Priority 2017-07-31 • Filed 2018-07-30 • Published 2020-06-10

There is provided a **vaccine** obtainable according to the process of the first aspect of the present invention. The **vaccine** may be for sub-cutaneous, sub-lingual or epicutaneous use. There is provided the use of a **vaccine** according to the present invention in the treatment of mite allergy or in the ...

[Transient silencing of argonaute1 and argonaute4 to increase recombinant ...](#)

WO EP CA EP3662067A1 Kiva FERRARO Plantform Corporation

Priority 2017-08-03 • Filed 2018-08-02 • Published 2020-06-10

19. The plant or plant cell of claim 16 or 17, wherein the recombinant protein is a therapeutic enzyme, optionally butyrylcholinesterase. 20. The plant or plant cell of claim 16 or 17, wherein the recombinant protein is a **vaccine** or a Virus Like Particle. 21 . An isolated nucleic acid molecule ...

[Hiv antibody compositions and methods of use](#)

WO EP US CN JP KR AU BR CA IL MX PH RU [EP3662930A1](#) Francois Vigneault AbViro LLC

Priority 2015-09-24 • Filed 2016-09-24 • Published 2020-06-10

According to another embodiment, the present invention provides a **vaccine** comprising at least one antibody of the invention and a pharmaceutically acceptable carrier. According to one embodiment, the **vaccine** is a **vaccine** comprising at least one antibody described herein and a pharmaceutically ...

[Thiazolopyridine derivatives as adenosine receptor antagonists](#)

WO EP CN KR AU CA SG TW EP3661941A1 Eva-Maria TANZER Merck Patent GmbH

Priority 2017-08-01 • Filed 2018-07-02 • Published 2020-06-10

Immunomodulators Interferon Dexosome therapy (Anosys) Oncophage (Antigenics) Pentrix (Australian Cancer GMK (Progenies) Technology) Adenocarcinoma **vaccine** JSF-154 (Tragen) (Biomira) Cancer **vaccine** (Intercell) CTP-37 (AVI BioPharma) Norelin (Biostar) JRX-2 (Immuno-Rx) BLP-25 (Biomira) PEP-005 (...

[Compositions and methods for inhibition of mica/b shedding](#)

WO EP US EP3661552A1 Lucas FERRARI DE ANDRADE Dana-Farber Cancer Institute, Inc.

Priority 2017-05-22 • Filed 2018-05-22 • Published 2020-06-10

An antigen may be MICA and/or MICB, or a fragment thereof. An antigen may also be a tumor antigen, against which protective or therapeutic immune responses are desired, e.g., antigens expressed by a tumor cell (e.g., in a **vaccine** in combination with an anti-MICA and/or anti-MICB antibody). [00135] ...

[Methods for activating immune cells](#)

WO EP CN KR AU CA SG EP3661545A1 Phillip S. KIM Trutino Biosciences Inc.

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

[0060] The term "**vaccine**" refers to a biological composition that, when administered to a subject, has the ability to produce an acquired immunity to a particular pathogen or disease in the subject. Typically, one or more antigens, or fragments of antigens, that are associated with the pathogen or ...

[Drug compound and purification methods thereof](#)

WO EP US CN KR AU CA SG EP3661522A2 Rajashree Joshi-Hangal Otsuka Pharmaceutical Co., Ltd.

Priority 2017-08-03 • Filed 2018-08-02 • Published 2020-06-10

[00223] In some embodiments, the lyophilized pharmaceutical composition described herein can be used in combination with a cancer **vaccine**, for example a cancer **vaccine** selected from a CTA cancer **vaccine**, such as a **vaccine** based on a CTA antigen selected from: NY- ESO-1, LAGE-1, MAGE-A1, -A2, -A3, - ...

[Methods for identifying and separating pro-allergic specific t cells](#)

WO EP EP3662057A1 Eric WAMBRE Benaroya Research Institute at Virginia Mason

Priority 2017-08-01 • Filed 2018-07-31 • Published 2020-06-10

... , such as allergen-specific immunotherapy (ASIT; also referred to as allergen **vaccine** therapy). The theory of ASIT is that exposure to gradually increasing allergen exposure will decrease the population of reactive pathogenic T cells and increase the population of T cells that promote tolerance.

[Device and method for multiple changes in the composition of a fluid](#)

DE DE102018009597A1 Martin Leuthold Sartorius Stedim Biotech Gmbh

Priority 2018-12-07 • Filed 2018-12-07 • Published 2020-06-10

) reach. In addition, by using an ultrafiltration membrane with a pore size of 2 to 100 nm as the second filter medium (5 , 15) a **vaccine** contained in the fluid as a product is retained by the ultrafiltration membrane. Components of the fluid that the second filter medium (5 , 15) can pass (e.g ...

[Purification process for biological molecules such as plasmid dna using anionic ...](#)

EP [EP3663401A1](#) Andreas Zurbriggen Lonza Limited

Priority 2019-02-28 • Filed 2019-02-28 • Published 2020-06-10

... pharmaceutically acceptable excipients. Plasmid DNA purified by the method of the present invention may be used as DNA-based **vaccine** or be used as a "prodrug" wherein the plasmid will then involve the cell's transcription and translation apparatus to biosynthesize the therapeutic entity in situ.

[Shot peening shot peening](#)

DE DE102018131530A1 Katrin Heider Schaeffler Technologies AG & Co. KG

Priority 2018-12-10 • Filed 2018-12-10 • Published 2020-06-10

Pressure piece for a synchronization gearbox with a heat-treated housing and cleaned by means of cleaning jets, characterized in that the pressure piece housing is treated by the cold-forming process of the **shot peening**. Method for increasing the strength of a housing of a pressure piece used in ...

[Methods for treating active eosinophilic esophagitis](#)

EP CN KR AU CA EP3661551A1 Allen Radin Regeneron Pharmaceuticals, Inc.

Priority 2017-08-04 • Filed 2018-08-03 • Published 2020-06-10

(4) Treatment with a live (attenuated) **vaccine** (Chickenpox (varicella), FluMist-Influenza, Intranasal influenza, Measles (rubeola), Measles-mumps-rubella combination, Measles-mumps-rubella-varicella combination, Mumps, Oral polio (Sabin), Oral typhoid, Rubella, Smallpox (vaccinia), Yellow fever, ...

[Composition for gene therapy of the central nervous system, process of ...](#)

WO EP BR [EP3662934A1](#) Roselena SILVESTRI SCHUH Universidade Federal Do Rio Grande Do Sul

Priority 2017-07-31 • Filed 2018-07-12 • Published 2020-06-10

... , especially due to their mucus adhesive properties, especially when the target is nasal administration aiming for the treatment of disorders of the central nervous system (Khatri, K. et al. Surface modified liposomes for nasal delivery of DNA **vaccine**. **Vaccine**, 2008, V. 26(18), p. 2225-33).

[Use of immune cell-specific gene expression for prognosis of prostate cancer ...](#)

WO EP AU CA EP3662082A2 Elai Davicioni Decipher Biosciences, Inc.

Priority 2017-08-04 • Filed 2018-08-02 • Published 2020-06-10

Methods, systems, and kits for the diagnosis, prognosis and the determination of cancer progression of prostate cancer in a subject are disclosed. In particular, the disclosure relates to the use of immune cell-specific gene expression in determining prognosis and identifying individuals in need ...

[Drive-in machine](#)

WO EP CN [EP3663049A1](#) Toshinori Yasutomi Koki Holdings Co., Ltd.

Priority 2017-07-31 • Filed 2018-06-29 • Published 2020-06-10

When determining "YES" at the step S2, the microcomputer 63 determines whether the operator is selecting the continuous **shot** mode or not at a step S3. When determining "YES" at the step S3, the microcomputer 63 performs the regular driving operation corresponding to the continuous **shot** mode at a ...

[Method and computer program product for detecting objects in night shots and ...](#)

DE DE102018221313A1 Mark Schutera Zf Friedrichshafen Ag

Priority 2018-12-10 • Filed 2018-12-10 • Published 2020-06-10

Method for detecting objects in night shots (x1) by means of a detector (O2) which is trained to detect objects in day shots (x2) of an environment of an automatically operated vehicle (40), comprising the method steps: Providing a night **shot** (x1) (V1), Mapping the night **shot** (x1) onto a feature ...

[Holding device for a vehicle and method for mounting a holding device](#)

DE DE102017007977B4 Martin Schoch Audi Ag

Priority 2017-08-23 • Filed 2017-08-23 • Granted 2020-06-10 • Published 2020-06-10

52 introduced. The clip 41 locked with the second **shot** 52 in a rest position. In a further step, the blocking agent 42 in the first **shot** 21st and in the second **shot** 52 inserted and locks the bracket 41 in the rest position. In the illustrated embodiment, the first **shot** 21st and the second **shot** 52

[Injection molding machine and injection molding information processing device](#)

EP CN JP [EP3381647B1](#) Masahiro Abe Sumitomo Heavy Industries, Ltd.

Priority 2017-03-31 • Filed 2018-03-29 • Granted 2020-06-10 • Published 2020-06-10

The reference **shot** may be a past **shot** of other shots to be compared and may be selected on the operation screen. The reference **shot** may be a **shot** at the time of shipment inspection at the manufacturer of the injection molding machine or may be a **shot** at the time of the injection molding at the ...

WO EP AU EP3661734A1 Owen Matthews Tyre Tuft Pty Ltd

Priority 2017-08-02 • Filed 2018-08-02 • Published 2020-06-10

18. The plug, kit or method of claim 15, wherein when the sealant is selected from the group consisting of Stans tire sealant, Tyre **Shot** sealant, Orange Seal sealant, Slime tire sealant, Continental sealant, Shimano sealant, Effetto Mariposa sealant, Tufo sealant, Mitas tyre sealant, ...

[Methods and compositions for viral-based gene editing in plants](#)

WO EP US AR CA EP3661354A1 Xingpeng LI R. J. Reynolds Tobacco Company

Priority 2017-07-31 • Filed 2018-07-30 • Published 2020-06-10

These expression vectors have proven versatile with demonstrated expression of numerous heterologous proteins, including cytokines, interferon, bacterial and viral antigens, growth hormone, **vaccine** antigens, single chain antibodies and monoclonal antibodies (mAbs). These expression levels support ...

[COMBINATIONS OF PD-1 ANTAGONISTS AND BENZO\[b](#)

WO EP AU CA EP3661499A1 Saso CEMERSKI Merck Sharp & Dohme Corp.

Priority 2017-08-04 • Filed 2018-07-30 • Published 2020-06-10

Adjuvants, such as aluminum hydroxide or aluminum phosphate, can be added to increase the ability of the **vaccine** to trigger, enhance, or prolong an immune response. Additional materials, such as cytokines, chemokines, and bacterial nucleic acid sequences, like CpG, a toll-like receptor (TLR) 9 ...

[Compositions and methods for targeting masas to treat cancers with spliceosome ...](#)

WO EP EP3662064A2 William Brian DALTON The Johns Hopkins University

Priority 2017-07-31 • Filed 2018-07-31 • Published 2020-06-10

[0393] As part of the process necessary to generate antibodies, which would be one therapeutic modality for targeting MASAs, immunogens representing the cryptic cell surface proteins to be targeted must be themselves created for **immunization** of animals and eventual screening of antibody ...

[Bispecific antibodies and uses thereof](#)

WO EP KR EP3661555A1 Yue Liu Ab Studio Inc.

Priority 2017-08-01 • Filed 2018-08-01 • Published 2020-06-10

NY. Acad Sci. 569:86-103; Flexner et al, 1990, **Vaccine**, 8: 17-21; U.S. Pat. Nos. 4,603,112, 4,769,330, and 5,017,487; WO 89/01973; U.S. Pat. No. 4,777,127; GB 2,200,651 ; EP 0,345,242; WO 91/02805; Berkner-Biotechmques, 6:616-627, 1988; Rosenfeld et al., 1991, Science, 252:431-434; Kolls et al, ...

[Virus vaccinal pour thérapie par prodroque activée par des enzymes exprimées ...](#)

WO EP US JP CA GB [EP3010518B1](#) Richard Marais The Institute of Cancer Research: Royal Cancer Hospital

Priority 2013-06-19 • Filed 2014-06-18 • Granted 2020-06-10 • Published 2020-06-10

Virus de la **vaccine** à utiliser dans le traitement d'une tumeur selon l'une quelconque des revendications précédentes, dans lequel le procédé comprend l'administration du virus de la **vaccine** par injection intraveineuse ou intratumorale. Virus de la **vaccine** à utiliser dans le traitement d'une tumeur ...

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

Results of Search in US Patent Collection db for: (ABST/vaccine AND ISD/20200526->20200601),

7 resultados.

PAT. NO.	Title
1 10,676,511	Coronaviruses epitope-based vaccines
2 10,675,345	Recombinant influenza virus vaccines for influenza
3 10,675,344	Multi-CBV vaccine for preventing or treating type I diabetes
4 10,675,343	Vaccines and methods for creating a vaccine for inducing immunity to all dengue virus serotypes
5 10,675,341	Parenteral norovirus vaccine formulations
6 10,675,338	Peptides and combination of peptides for use in immunotherapy against breast cancer and other cancers
7 10,675,249	Vaccine delivery systems using yeast cell wall particles

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