

VacCiencia

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EN ESTE NÚMERO

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

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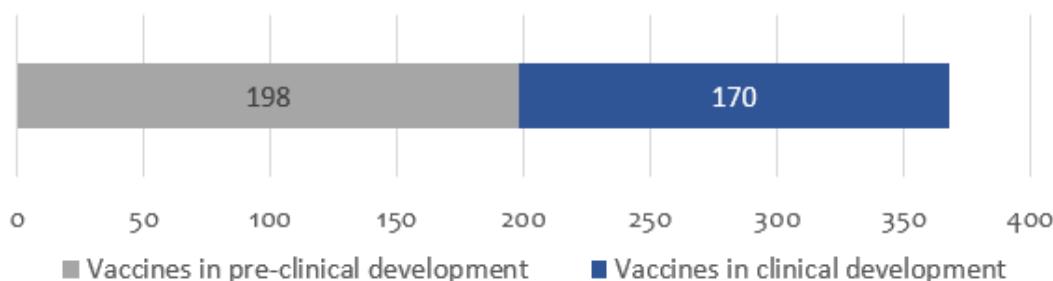
Resumen de la información publicada por la OMS sobre los candidatos vacunales contra la COVID-19 en desarrollo a nivel mundial

Última actualización por la OMS: 9 de agosto de 2022.

Fuente de información utilizada:



170 Vacunas en evaluación clínica y 198 en evaluación preclínica



Candidatos vacunales en evaluación clínica por plataforma

Platform		Candidate vaccines (no. and %)	
PS	Protein subunit	54	32%
VVnr	Viral Vector (non-replicating)	21	12%
DNA	DNA	16	9%
IV	Inactivated Virus	22	13%
RNA	RNA	41	24%
VVr	Viral Vector (replicating)	4	2%
VLP	Virus Like Particle	6	4%
VVr + APC	VVr + Antigen Presenting Cell	2	1%
LAV	Live Attenuated Virus	2	1%
VVnr + APC	VVnr + Antigen Presenting Cell	1	1%
BacAg-SpV	Bacterial antigen-spore expression vector	1	1%
		170	

Candidatos vacunales mucosales en evaluación clínica

Desarrollador de la vacuna/fabricante/país	Plataforma de la vacuna	Vía de administración	Fase
University of Oxford/Reino Unido	Vector viral no replicativo	Intranasal	1
CanSino Biological Inc./Beijing Institute of Biotechnology/China	Vector viral no replicativo	Inhalación	4
Vaxart/Estados Unidos	Vector viral no replicativo	Oral	2
Univ. Hong Kong, Xiamen Univ./Beijing Wantai Biol. Pharm./China	Vector viral replicativo	Intranasal	3
Symvivo/Canadá	ADN	Oral	1
ImmunityBio, Inc./Estados Unidos	Vector viral no replicativo	Oral o SL	1/2
Codagenix/Serum Institute of India	Virus vivo atenuado	Intranasal	3
Center for Genetic Engineering and Biotechnology (CIGB)/Cuba	Subunidad proteica	Intranasal	1/2
Razi Vaccine and Serum Research Institute/India	Subunidad proteica	IM e IN	3
Bharat Biotech International Limited/India	Vector viral no replicativo	Intranasal	3
Meissa Vaccines, Inc./Estados Unidos	Virus vivo atenuado	Intranasal	1
Laboratorio Avi-Mex/México	Virus inactivado	IM o IN	2/3
USSF + VaxForm/Estados Unidos	Subunidad proteica	Oral	1
CyanVac LLC/Estados Unidos	Vector viral no replicativo	Intranasal	1
DreamTec Research Limited/Hong Kong	BacAg-SpV	Oral	NA
Sean Liu, Icahn School of Medicine at Mount Sinai	Vector viral replicativo	IN/IM	2/3
Hannover Medical School/Alemania	Vector viral no replicativo	Inhalación	1
ACM Biolabs/Singapur	Subunidad proteica	IN/IM	1

Candidatos vacunales más avanzados a nivel global

Desarrollador de la vacuna/fabricante/país	Plataforma de la vacuna	Fase
Sinovac/China	Virus Inactivado	4
Sinopharm/Wuhan Institute of Biological Products/China	Virus Inactivado	4
Sinopharm/Beijing Institute of Biological Products/China	Virus Inactivado	4
University of Oxford/AstraZeneca/Reino Unido	Vector viral no replicativo	4
CanSino Biological Inc./Beijing Institute Biotechnology/China	Vector viral no replicativo	4
CanSino Biological Inc./Beijing Institute Biotechnology/China	Vector viral no replicativo (IH)	4
Gamaleya Research Institute/Rusia	Vector viral no replicativo	3
Janssen Pharmaceutical Companies/Estados Unidos	Vector viral no replicativo	4
Novavax/Estados Unidos	Subunidad proteica	3
Moderna/NIAID/Estados Unidos	ARN	4
Pfizer/BioNTech Fosun Pharma/Estados Unidos	ARN	4
Anhui Zhifei Longcom Biopharmac./Inst. Microbiol, Chin Acad Sci/China	Subunidad proteica	3
CureVac AG/Alemania	ARN	3
Institute of Medical Biology/Chinese Academy of Medical Sciences	Virus inactivado	3
Research Institute for Biological Safety Problems, Kazakhstan	Virus inactivado	3
Inovio Pharmac. + Intern. Vacc Inst. + Advaccine Biopharm Co., Ltd	ADN	3
Zydus Cadila Healthcare Ltd./India	ADN	3
Bharat Biotech International Limited/India	Virus Inactivado	3
Sanofi Pasteur + GSK/Francia/Gran Bretaña	Subunidad proteica	3
Shenzhen Kangtai Biological Products Co., Ltd./China	Virus Inactivado	3
Clover Biopharmaceuticals Inc./GSK/Dynavax/China/Reino Unido/EE.UU	Subunidad proteica	3
Vaxine Pty Ltd. + CinnaGen Co./Australia, Irán	Subunidad proteica	3
Medigen Vaccine Biol./Dynavax/NIAID/Taiwán/EE.UU	Subunidad proteica	4
Instituto Finlay de Vacunas/Cuba	Subunidad proteica	3
Federal Budget Res Inst State Res Cent Virol Biotechnol "Vector"/Rusia	Subunidad proteica	3
West China Hospital + Sichuan University/China	Subunidad proteica	3
Vaxxinity/EE.UU	Subunidad proteica	3
Univ. Hong Kong, Xiamen Univ. & Beijing Wantai Biological Pharm./China	Vector viral replicativo	3
Acad Milit Sci (AMS) Walvax Biotechnol, Suzhou Abogen Biosci/China	ARN	3
Medicago Inc./Canadá	Particula similar a virus	3
Codagenix/Serum Institute of India	Virus vivo atenuado	3
Center for Genetic Engineering and Biotechnology (CIGB)/Cuba	Subunidad proteica	3
Valneva, National Institute for Health Research, Reino Unido	Virus inactivado	3
Biological E. Limited/India	Subunidad proteica	3
Nanogen Pharmaceutical Biotechnology/Vietnam	Subunidad proteica	3
Shionogi/Japón	Subunidad proteica	3
Erciyes University/Turquía	Virus inactivado	3
SK Bioscience Co., Ltd./CEPI/Corea del Sur/Noruega	Subunidad proteica	3
Razi Vaccine and Serum Research Institute/Irán, India	Subunidad proteica	3
Bharat Biotech International Limited/India	Vector viral no replicativo (IN)	3
Jiangsu Rec-Biotechnology/China	Subunidad proteica	3
Radboud University/Holanda	Particula similar a virus	3
Livzon Pharmaceutical/China	Subunidad proteica	3
KM Biologics Co., Ltd./Japón	Virus inactivado	2
Bagheiat-allah University of Medical Sciences/AmitisGen/Irán	Subunidad proteica	3
Laboratorios Hipra, S.A.	Subunidad proteica	3
Arcturus Therapeutics, Inc./Estados Unidos	ARN	3
Sinocelltech Ltd./China	Subunidad proteica	3
Chumakov Federal Scientific Center for Research/Rusia	Virus Inactivado	3
PT Biofarma/Indonesia	Subunidad proteica	3
AIM Vaccine and LiveRNA Therapeutics/China	ARN	3
CanSino Biologics Inc./China	ARN	3
Moderna TX/Estados Unidos	ARN	3
China National Biotec Group Company Limited	Virus inactivado	3

Fidel Por Siempre

El legado de Fidel en la Ciencia Cubana. Breves apuntes

Desde que comenzó la pandemia en el mundo, provocada por el impacto del nuevo coronavirus, y su enfrentamiento en nuestro país, han sido reiteradas las expresiones de reconocimiento a la visión de largo alcance presente en las concepciones del líder histórico de la Revolución Cubana, Fidel Castro Ruz, sobre el papel de la ciencia en la solución de los principales problemas que habría de enfrentar la nación en el complejo camino hacia un desarrollo económico y social que garantizara el bienestar del pueblo.



El punto de referencia hoy para tales expresiones, lo constituye, evidentemente, la encomiable labor de nuestros científicos en la búsqueda primero, y en la materialización después, de cinco candidatos vacunales, cien por ciento cubanos, de los cuales dos ya han alcanzado la categoría de vacunas, como solución clínica ante la nueva enfermedad, con un probado impacto positivo, reconocido, incluso, a nivel internacional.

Y no nos falta razón a quienes afirmamos que en dichos resultados se transparenta la impronta de Fidel. Ello es innegable. El valor de sus concepciones, y de la obra que forjó, han sido decisivas para el enfrentamiento a la pandemia en Cuba.

Y es, precisamente, esa obra la que es indispensable develar y divulgar, esencialmente en las actuales y futuras generaciones, pues, aunque en la conciencia cotidiana, la vacuna sea, como ya se ha expresado, el punto referencial del justo reconocimiento, lo cierto es que las concepciones de Fidel Castro acerca de la ciencia, no se limitan al campo de la medicina, ni solo al aspecto técnico-material del desarrollo científico. Es una concepción multifacética e integral, fraguada desde los mismos albores de la Revolución.

No es posible, en tan poco espacio, reseñar el legado de Fidel en el campo de la ciencia, forjado en medio de enormes dificultades por las que ha atravesado el proceso revolucionario cubano, en condiciones de economía subdesarrollada y bajo un asedio sin precedentes por parte de sucesivos gobiernos de Estados Unidos.

En tales circunstancias, uno de los grandes méritos de Fidel Castro fue, desde un inicio, jerarquizar el valor del conocimiento para garantizar el futuro. De ahí su empeño en lograr el acceso masivo del pueblo a la educación y a la cultura, concebidos no solo como derechos humanos fundamentales, sino además como componentes esenciales de una concepción integral de la sociedad, en la que educación, cultura y ciencia conforman un todo único e interrelacionado y base para nuestra

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soberanía, independencia y desarrollo económico y social. A ello se suman su capacidad para analizar los contextos históricos, su voluntad para plantearse audaces metas y para trabajar en su consecución, así como su confianza ilimitada en las potencialidades intelectuales y éticas de los cubanos. Todo ello condicionó una inédita hazaña: lograr que un pequeño país subdesarrollado, bloqueado y amenazado por la principal potencia imperialista mundial, a solo 90 millas de sus costas, se adentrara con solidez por los complejos caminos de la ciencia.

Una de las primeras pautas trazadas por Fidel Castro, respecto a la definición de la política científica de la Revolución, - con un alcance hasta el presente y para el futuro-, fue su intervención el 15 de enero de 1960, en la Sociedad Espeleológica de Cuba, ocasión en la que se le otorgó el título de Socio de Honor de esa entidad científica, una de las pocas que existían en el país.

Esto tuvo lugar, en momentos en que la radicalización de la Revolución ya era palpable, lo que provocaba las acciones del enemigo, incluyendo la incitación al éxodo masivo de profesionales, además del inicio de la cruenta guerra económica impuesta por Estados Unidos. En este contexto, Fidel proyectó su concepción estratégica e integral sobre el rol de la ciencia, el pensamiento y la inteligencia para el desarrollo del país. Sentenciaba entonces el Comandante en Jefe: "El futuro de nuestra patria tiene que ser necesariamente un futuro de hombres de ciencia, tiene que ser un futuro de hombres de pensamiento, porque precisamente es lo que más estamos sembrando; lo que más estamos sembrando son oportunidades a la inteligencia; ya que una parte considerabilísima de nuestro pueblo no tenía acceso a la cultura, ni a la ciencia".

El futuro de nuestra Patria
tiene que ser necesariamente
un futuro de hombres de
ciencia.

A handwritten signature in black ink, appearing to read "Fidel Castro", is written over two curved lines forming an oval. The signature is fluid and cursive, with a prominent "F" at the beginning.

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Resulta interesante el hecho de que este discurso tiene lugar con anterioridad a sus célebres Palabras a los Intelectuales y al despliegue de la Campaña de Alfabetización en 1961, que convirtió a Cuba en el primer país de América Latina libre de analfabetismo.

Así mismo, con su extraordinaria sensibilidad humanista, desde entonces se comprometió a revertir el olvido al que estaba sometida la ciencia en la república neocolonial, y reconoció que lograr el desarrollo científico formaba parte de los objetivos priorizados de la Revolución, no solo desde el punto de vista socioeconómico, sino en materia de justicia social. A la vez, hizo patente su compromiso de promover políticas para el cultivo de las inteligencias y para el desarrollo de la ciencia y del pensamiento: A tal efecto, afirmaba: “¡Cuántas inteligencias se habrán desperdiciado en ese olvido! ¡Cuántas inteligencias se habrán perdido! Inteligencias que hoy se incorporarán a la vida de su país; inteligencias que hoy se incorporarán a la cultura y a la ciencia, porque para eso estamos convirtiendo las fortalezas en escuelas; para eso estamos construyendo ciudades escolares; para eso estamos llenando la isla de maestros, para que en el futuro la patria pueda contar con una pléyade brillante de hombres de pensamiento, de investigadores y de científicos”.

Esos fueron los inicios de un largo y complejo camino que llega hasta el presente y se enrumba con optimismo hacia el futuro, pues en consecuencia con dicha estrategia, a lo largo de todo el proceso revolucionario, se han dedicado múltiples esfuerzos y recursos al desarrollo de la ciencia en Cuba y su implementación en la práctica social, en cuyo empeño es preciso –y justo– reconocer la titánica labor realizada por los miles de hombres de ciencia, y de muchos otros que, sin serlo, han contribuido significativamente en dicha obra, formados todos, precisamente, en los principios de la política científica nacional, fundada por Fidel Castro.

No es fácil sintetizar el legado de Fidel en el campo de la ciencia y en el despliegue de proyectos tecnológicos y de innovación en las diversas ramas de la economía y del desarrollo social en general. Pero sin lugar a dudas sus aportes y logros son palpables en los altos índices de desarrollo científico que hoy Cuba puede exhibir.

Vale la pena, por tanto, recordar algunos hitos que dan fe de su invaluable obra a favor del desarrollo simultáneo e integral de la cultura, la educación y la ciencia en la Cuba revolucionaria.

Es una obra sembrada durante el primer lustro de la Revolución, en las propias raíces de la construcción del socialismo en Cuba. No es casual que bajo su guía e impulso, desde 1959, se iniciara una profunda revolución educacional y que en 1961 se desplegara la inédita Campaña de Alfabetización a partir de sus convicciones sobre el importante rol de la participación popular y de los jóvenes en el proceso de transformación revolucionaria de la sociedad cubana.

Su visión de futuro también fue decisiva cuando la nueva institucionalidad creada para el desarrollo de la ciencia se concibió con un sentido inclusivo de la tecnología, la innovación y la protección de los recursos naturales. Un hito en ese empeño fue la creación de la Academia de Ciencias de Cuba en 1962, bajo una concepción integradora de todas las esferas y disciplinas de la ciencia, a lo que se unió el despliegue de la reforma universitaria y la creación del Centro Nacional de Investigaciones Científicas (CENIC) en 1965, matriz de otras instituciones que posteriormente fueron desarrollándose hasta conformar toda una potente y eficaz infraestructura para el desarrollo de la ciencia en el país.

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Desde esos pilares, levantados por Fidel, la ciencia dejó de constituir una actividad de élites o de científicos aislados, para convertirse en patrimonio del pueblo a partir de la universalización de la educación. Ello se hizo patente, con singular fuerza, a lo largo de los años 70 y 80, en los que, junto con la creación de universidades y de múltiples centros de investigación, surgieron entidades como el Fórum de Ciencia y Técnica, las Brigadas Técnicas Juveniles, el Movimiento de Innovadores y Racionalizadores y otras, vinculadas con el movimiento obrero y sindical, que habrían de desempeñar un papel protagónico en la ardua tarea de contrarrestar las nefastas consecuencias derivadas del subdesarrollo y del bloqueo económico impuesto por los Estados Unidos.

Mención especial merece el impulso, diseño estratégico y presencia fundacional de Fidel en el surgimiento de diversas entidades de investigación en el campo de las ciencias biomédicas y agropecuarias, entre otras, con una proyección interdisciplinaria y colossal visión de futuro. En ese marco sobresale la creación del Sector Biotecnológico a partir de 1981, cuando su despliegue era monopolizado por países del llamado primer mundo. Ahí radica uno de los más importantes antecedentes de la fortaleza científica cubana que ha permitido hoy el enfrentamiento exitoso al nuevo coronavirus.



Lo que Fidel generó no se limita al rescate de inteligencias, ni a la creación de instituciones científicas, aunque esto solo ya sería un gran mérito. Su gran aporte en este campo ha sido generar una política de desarrollo de la ciencia y la tecnología impregnada de valores éticos, con un sentido humanista y de trabajo colectivo, de colaboración interinstitucional, de solidaridad internacional y de promoción de los diversos campos de la investigación científica, incluyendo las ciencias básicas, las ciencias técnicas y nucleares, así como la no menos importante esfera de las ciencias sociales y humanísticas.

Esa concepción tuvo su prueba de fuego en la década de los años 90 del pasado siglo, ante la necesidad de potenciar una economía basada en las ciencias para enfrentar los negativos impactos del derrumbe del socialismo en Europa del Este y la URSS y del recrudecimiento del bloqueo económico de Estados Unidos contra Cuba. A la vez, para garantizar la independencia y soberanía del país, la supervivencia de la Revolución y las bases para el desarrollo económico en nuevas y más complejas condiciones nacionales e internacionales.

En aquel contexto la capacidad de previsión de Fidel resultó decisiva al proclamar que la independencia del país dependía del desarrollo de la ciencia y la tecnología. Ante la escasez de recursos de todo tipo, sobre todo los energéticos, estimuló y potenció la producción de la inteligencia y el conocimiento, consciente de que estos factores habrían de desempeñar un rol estratégico en el desarrollo y futuro de la nación.

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Esta concepción fue validada por acciones concretas que elevaron el nivel de desarrollo científico de la nación, particularmente la organización de los polos científicos a partir de 1991, cuya integración generó capacidades para potenciar los recursos científicos, tecnológicos y organizativos con que contaba el país y atender programas priorizados que dieran solución a múltiples problemas de la sociedad.

De igual forma, y como expresión de la materialización de esta línea de pensamiento de Fidel, se crea, en 1994, el Ministerio de Ciencia, Tecnología y Medio Ambiente, como herramienta institucional para la proyección y concreción de la política científico-tecnológica nacional y la protección del medio ambiente, en sintonía con las ideas expuestas por él en la Conferencia de Naciones Unidas Sobre Medio Ambiente y Desarrollo, celebrada en Río de Janeiro en 1992, y que tanto impacto tuvieron en la comunidad internacional.

Desde entonces diversos programas continuaron enriqueciendo el quehacer científico nacional, varios de ellos asociados a la Batalla de Ideas que, bajo la concepción y dirección de Fidel, se desplegó en los primeros años del actual siglo.

Sus esfuerzos en defensa del papel de la ciencia en el desarrollo del país y de la solución de los problemas que afectan a la sociedad, fueron palpables hasta los últimos años de su vida, particularmente en el impulso de alternativas, con la aplicación del conocimiento científico, para enfrentar las dificultades en los ámbitos de la salud y la alimentación del pueblo.

Resumiendo todo lo anterior, que apenas constituye una apretada síntesis de las concepciones y la obra de Fidel Castro acerca de la ciencia y su papel en la vida social, se puede afirmar que ésta tiene un singular significado, muchas veces protagónico y decisivo, para la actividad y desarrollo de la comunidad científica cubana de nuestros días, no solo por lo que hizo, sino por lo que su legado aporta al presente y al futuro de nuestro país.

Los que de una u otra forma trabajamos en esa comunidad, independientemente de la rama de la ciencia de que se trate, somos parte de los agradecidos por su obra, y frutos de la promoción y siembra de inteligencias que proyectó desde los mismos albores del proceso revolucionario cubano.

Hoy, en Cuba, la ciencia constituye un arma poderosa, porque está en manos de miles de profesionales y trabajadores integrados a nuestro pueblo, plagado de hombres y mujeres de vasta experiencia, formando equipo con jóvenes educados y talentosos capaces de llevar adelante la Revolución como proceso integral de liberación nacional, antiimperialista y socialista.



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De ahí que, en el actual contexto de enfrentamiento a la pandemia del coronavirus, el reconocimiento de la impronta y de la huella del pensamiento y la obra de Fidel Castro, con un profundo sentido ético, a favor de una ciencia para el mejoramiento humano, para la paz y no para la guerra, para y con el pueblo, y con nobles compromisos para el despliegue de solidaridad e internacionalismo, brote de manera espontánea en los hijos de este pueblo.

Fuente: CUBAHORA. Disponible en <https://bit.ly/3dwyx8o>



Noticias en la Web

La vacuna de Medigen supera a la AstraZeneca en estudio de fase 3 en Paraguay

1 ago. El estudio de fase 3 de la vacuna contra la COVID-19 del laboratorio taiwanés Medigen, desarrollado en Paraguay, demostró que es "segura, tolerable y que funciona", dijo a Efe el investigador principal, Julio Torrales, sobre los resultados de este trabajo divulgados este lunes en Asunción.

El estudio, que duró 8 meses, comparó el biológico MVC-COV1901 elaborado por Medigen Vaccine Biologics Corporation (MVC) con la vacuna AZD1222 del laboratorio AstraZeneca.

"Lo que buscábamos era determinar la seguridad en términos de efectos adversos, la tolerabilidad de nuestra vacuna y su inmunogenicidad, que quiere decir la capacidad de generar anticuerpos", explicó el también catedrático.

La investigación mostró que la vacuna candidata causó "menos" efectos adversos locales y sistémicos, como cefalea, dolor en el sitio de la inyección y mialgia, que la AstraZeneca.

En total, participaron 1.030 personas divididas en dos grupos que recibieron uno de los biológicos. Ni los participantes, los investigadores o patrocinadores fueron informados sobre cuál fue el biológico administrado durante la evaluación.

El experto detalló que el grupo que recibió la vacuna candidata desarrolló 2,6 veces más anticuerpos específicos (o antiespiga) que los inmunizados con AstraZeneca.

"Lo que nosotros estamos informando al Gobierno paraguayo es que tenemos otra opción: tenemos una vacuna segura, tolerable, muy inmunogénica que podría formar parte del bagaje de la lucha contra la covid-19", agregó Torrales.

Señaló que el estudio tuvo lugar en las localidades de San Lorenzo (a unos 15 kilómetros de Asunción) y Ciudad del Este (sureste), en la frontera con Brasil y Argentina.

En febrero pasado, la Dirección Nacional de Vigilancia Sanitaria (Dinavisa) de Paraguay concedió la autorización para el uso de emergencia de la vacuna de Medigen en personas mayores de 18 años.

Se recomienda la aplicación intramuscular de dos dosis, con un intervalo de 4 semanas entre cada una.

Las vacunas, desarrolladas con la colaboración del Instituto Nacional de Salud de Estados Unidos (NIH, por su sigla en inglés), requieren una temperatura de conservación de entre 2 y 8 grados centígrados.

En julio del año pasado, la Administración de Alimentos y Medicamentos de Taiwán otorgó la autorización de emergencia a esta vacuna.

Fuente: SWI swissinfo.ch. Disponible en <https://bit.ly/3AoOkzv>



Fuente de la imagen: Sitio web www.medigenvac.com

Bruselas comprará 250 millones de vacunas para la COVID-19 a la española Hipra

2 ago. "Dado el nuevo aumento de casos en Europa, este acuerdo pondrá la vacuna de Hipra a disposición de los países participantes, tan pronto como la vacuna haya recibido una evaluación positiva por parte de la Agencia Europea de Medicamentos", ha señalado este martes la Comisión Europea para anunciar la adquisición de vacunas a la farmacéutica española Hipra.

La firma del contrato recoge la compra para el próximo otoño de 250 millones de dosis de la vacuna contra la covid-19 desarrollada por esta empresa, cuyo fármaco está aún pendiente de aprobación por la Agencia Europea de Medicamentos (EMA). Asimismo, según informó la CE en un comunicado recogido por Efe, en la adquisición participan catorce Estados miembros y países, que pueden solicitar hasta 250 millones de dosis.

El Ejecutivo comunitario valora que la vacuna proteica de Hipra añade "una opción más" que complementa la cartera de vacunas comunitaria, declaró la comisaria de Salud y Seguridad Alimentaria, Stella Kyriakides.

El contrato de adquisición con Hipra, con sede en Girona, se suma a los firmados por Bruselas con las farmacéuticas AstraZeneca, Sanofi-GSK, Janssen Pharmaceutica, BioNtech-Pfizer, Moderna, Novavax y Valneva.

En esta línea, se han conseguido unos 4.200 millones de dosis en el marco de la Estrategia de Vacunas de la UE y los países participantes podrían decidir donar las vacunas a países de renta baja y media o redirigirlas a otros países europeos. "Ante el aumento de las infecciones por la covid-19 en Europa, tenemos que garantizar la máxima preparación para los meses de otoño e invierno", añadió Kyriakides.

Sánchez aplaude el acuerdo de Hipra con UE

Al respecto se ha pronunciado el presidente del Gobierno, Pedro Sánchez, que ha celebrado el contrato de compra conjunta porque demuestra la fortaleza de la innovación en la industria farmacéutica en España. "Por fin hemos logrado esta vacuna", se ha felicitado Sánchez en una rueda de prensa en el palacio de La Almudaina (Palma) tras despachar con el rey.

Para el Gobierno español, en palabras del líder del Ejecutivo, el contrato supone una "prueba magnífica de lo bien que puede funcionar la colaboración público-privada en una nueva política industrial", que hace que hoy España "sea referente" a nivel de vacunación y también por esta aportación "frente a nuevas variantes". De ahí que haya querido trasladar la satisfacción y compromiso del Ejecutivo con esta industria. "Es una muy buena noticia", ha concluido.

La vacuna de Hipra contra la covid-19 está basada en dos proteínas recombinantes estructuralmente similares, una correspondiente a la variante alfa y otra a la variante beta, que se unen formando una estructura única llamada dímero, y que está acompañada de un adyuvante que incrementa la respuesta inmunológica. Una de sus ventajas es que se puede adaptar a las variantes, según la farmacéutica.

La profilaxis, que actualmente está siendo objeto de revisión por parte de la EMA, se está desarrollando como dosis de refuerzo en personas previamente inmunizadas de 16 años o más y si recibe la autorización de comercialización, los países participantes podrán adquirirla a través del contrato vigente.

Fuente: Público. Disponible en <https://bit.ly/3AfjsRK>



Padecer COVID-19 en múltiples oportunidades incrementa el riesgo de sufrir COVID-19 prolongada, indica estudio

3 ago. La reinfección de COVID-19 es un tema de conversación no solo en medios. Entre nuestro círculo social hemos escuchado de alguien que ha padecido la enfermedad hasta 3 o 4 veces.

El Dr. Elmer Huerta analiza dos estudios en donde se indican los efectos de esas reinfecciones y uno de esos tienen que ver con los síntomas persistentes de la enfermedad, conocido como COVID-19 prolongada.

Anteriormente, escuchamos lo que se llama COVID-19 persistente o *long-covid*, y como esa condición —al igual que muchas otras en la historia de la medicina— tuvo que luchar por ser reconocida por los profesionales de la salud.

No hay duda que con la llegada de la variante Ómicron y sus diferentes sublinajes, la pandemia ha cambiado drásticamente.

Ómicron, la variante que cambió el curso de la pandemia

Si antes de Ómicron se esgrimía el concepto de inmunidad de rebaño como una de las armas de control de la pandemia, la llegada de esa familia del SARS-CoV-2 cambió completamente las reglas del juego.

Eso es porque una de las características más importantes de la familia Ómicron es que es capaz de evadir los anticuerpos neutralizantes que se producen como consecuencia de la infección previa y de la vacunación.

Eso hace que una persona pueda infectarse dos, tres y hasta cuatro veces, con lo que el concepto de inmunidad de rebaño ya no es posible.

Recordemos que la inmunidad de rebaño se define como la protección que adquiere una comunidad como consecuencia de que una cierta proporción de la población se infecte con determinado virus. Dicha definición tiene su fundamento en que una persona no puede reinfecarse con el virus en cuestión.

Al ocurrir millones de reinfecciones, el concepto de inmunidad de rebaño se convierte entonces en una situación relativamente irrelevante.

¿Múltiples infecciones de COVID-19, mayor protección?

Si se acepta entonces que una persona puede infectarse más de una vez, una pregunta muy pertinente es la que se refiere al significado de esa, o de esas reinfecciones: ¿le brinda algún beneficio, o al revés, lo predispone a complicaciones?

Dos recientes estudios, hechos por investigadores de la Universidad Washington en St. Louis, así como del sistema de Atención de Veteranos en St. Louis y la Universidad de St. Louis, en Missouri, Estados Unidos, brindan respuestas a esas preguntas.

En el primero, publicado en *Nature Medicine* del 25 de mayo, los investigadores compararon lo que sucedió con la salud de casi 34.000 personas que tuvieron por lo menos una reinfección después de haber sido vacunadas contra COVID-19, con la de tres grupos de personas, llamados grupos de control.

El primero, llamado contemporáneo, compuesto de casi 5 millones de personas que vivieron durante la pandemia, pero sin historia de infección por el SARS-CoV-2.

El segundo, un grupo de casi 5 millones y medio de personas que vivieron antes de la pandemia y el tercero, de casi 3 millones de personas vacunadas contra COVID-19.

Los resultados muestran que a pesar de que los síntomas de COVID-19 prolongada, incluyendo un mayor riesgo de muerte, pueden presentarse también en personas vacunadas que se reinfectan, su frecuencia es menor en este último grupo.

Los autores concluyen que, aunque la vacunación puede reducir parcialmente los riesgos de muerte y COVID-19 prolongada, se requiere que las políticas de salud pública sigan alentando la prevención primaria de la infección con el uso de mascarillas, distancia social y ventilación de espacios cerrados.

La COVID-19 prolongada tras una reinfección

En el segundo estudio, aún una prepublicación no revisada por pares, los autores amplían su estudio para saber si el tener dos o más infecciones aumenta el riesgo de sufrir de síntomas de COVID-19 prolongada.

Para eso, comparan tres grupos:

El primero de más de 257.000 personas que tuvieron una sola infección del SARS-CoV-2,

El segundo, de casi 39.000 personas que tuvieron dos, tres o cuatro infecciones por SARS-CoV-2,

El tercero, de más de 5 millones y medio de personas sin historia de infección.

Como dato interesante, en el grupo de casi 39.000 reinfecciones se vio que:

- Más de 36.000 lo tuvieron dos veces,
- Más de 2.200 lo tuvieron tres veces y
- 246 se infectaron cuatro veces.

Con respecto al tiempo entre infección e infección, se vio que la media entre la primera y la segunda infección fue de 79 días (con un intervalo entre 48 y 119 días), y entre la segunda y tercera infección fue de 65.

Los resultados del estudio indican que, al contrario de lo que la gente cree, tener dos o más infecciones no brinda mayor protección o resistencia a desarrollar síntomas, sino que, al revés, aumenta la posibilidad de presentar síntomas persistentes.

En ese sentido, se vio que el riesgo de presentar nuevos síntomas fue mayor cuando la persona se reinfectaba, viéndose que los síntomas duraban por lo menos seis meses. Ese mayor riesgo de desarrollar síntomas fue observado independientemente de si alguien había sido vacunado o no, y fue directamente proporcional al número de infecciones, observándose que aumentaba con cada infección posterior.

Síntomas más comunes que se presentan en una reinfección

Los síntomas más comunes después de las reinfecciones incluyeron:

- ⇒ Dolor de pecho
- ⇒ Ritmos cardíacos anormales
- ⇒ Ataques cardíacos
- ⇒ Inflamación del músculo cardíaco o del saco que rodea el corazón
- ⇒ Insuficiencia cardíaca y coágulos sanguíneos

Los problemas pulmonares comunes incluyeron:

- ⇒ Dificultad para respirar
- ⇒ Bajo nivel de oxígeno en la sangre
- ⇒ Enfermedad pulmonar

Acumulación de líquido alrededor de los pulmones

En resumen, la aparición de la variante Ómicron cambió completamente el panorama de la pandemia, haciendo que las reinfecciones sean mucho más frecuentes que antes, las que —al contrario de lo que uno pudiera creer— no es que refuerzan el sistema de defensa causando enfermedades más leves, sino que nos ponen en un mayor riesgo de desarrollar síntomas de COVID-19 persistente.

Si bien es cierto que la vacunación puede proteger en parte contra esas complicaciones, esa protección no sería completa, por lo que es importante que —mientras persista la pandemia— hagamos todo el esfuerzo para evitar la infección.

Fuente: CNN en español. Disponible en <https://cnn.it/3bJB8eZ>

Over 6.42m Belarusians now fully vaccinated against COVID-19

Aug 4. Over 6.512 million people in Belarus have got the first shot of a COVID-19 vaccine, of them over 6.424 million have completed the vaccination process, BelTA learned from the press service of the Belarusian Healthcare Ministry.

Over 892,300 people completed the full vaccination series in Brest Oblast, over 789,900 in Vitebsk Oblast, over 973,800 in Gomel Oblast, over 693,500 in Grodno Oblast, over 691,000 in Mogilev Oblast, over 983,700 in Minsk Oblast, and over 1.28 million in the city of Minsk.

Over 111,000 employees of organizations and institutions have completed the vaccination process.

Over 150,700 teenagers aged 12-17 have received one shot of the vaccine, of them more than 145,500 children have been fully vaccinated.

Thus, 70.4% of the country's population have received one dose of the vaccine, 69.4% have been fully vaccinated.

Belarusians are also getting booster shots of the vaccine, with 26.1% of the population having completed it so far.

Fuente: BELTA. Disponible en <https://bit.ly/3SMayCk>

Tinnitus, nuevo efecto secundario de la vacuna de AstraZeneca contra la COVID-19: qué es y cómo tratarlo

5 ago. A medida que va pasando el tiempo, se van conociendo cada vez más cosas sobre la COVID-19, algo en lo que también ha influido la llegada de las nuevas variantes, cuyos cambios en los tiempos de incubación, síntomas o duración del virus, han dado un giro de 180° a todo lo que ya sabíamos de la enfermedad.

Aunque esté menos presente, el virus sigue estando ahí y siguen existiendo contagios e ingresos hospitalarios.



Hasta mediados de julio, las nuevas variantes BA.4 y BA.5 de Ómicron supusieron más del 75% de las nuevas infecciones, tal y como reflejó el último informe de la situación epidemiológica hecho público por el Ministerio de Sanidad este martes.

Nuevos efectos secundarios

¿Y qué hacer ante la permanente presencia de un virus que va mutando y del que surgen más variantes? Vacunarse continúa siendo la mejor opción para prevenir tanto la enfermedad como su gravedad en caso de contagiarnos.

Los nuevos conocimientos del virus que mencionábamos anteriormente también se aplican a las vacunas, ya que a medida que va pasando el tiempo y cada vez más población está inoculada, se van descubriendo nuevos efectos adversos de las mismas. Uno de ellos es el Tinnitus, un viejo conocido para muchos que han padecido la COVID-19.

¿Por qué? Porque según el último informe de Farmacovigilancia sobre las vacunas frente a la COVID-19 publicado por la Agencia Española de Medicamento y Productos Sanitarios (AEMPS) se han detectado nuevos efectos secundarios de las vacunas, tanto de las más como de las menos utilizadas.

El dossier revela que en Moderna se ha identificado la inflamación extrema de la extremidad vacunada; en AstraZeneca, el Tinnitus, la parestesia o la hipoestesia; y en Nuvaxovid, la anafilaxia, la parestesia y la hipoestesia. Sin embargo, aún se está evaluando si los sangrados menstruales abundantes son un posible efecto adverso con Pfizer y Moderna.

¿Qué es el Tinnitus?

Según la Clínica Mayo, se conoce como Tinnitus a aquella sensación de pitido u otros ruidos en uno o ambos oídos. No es un sonido externo, por lo que otras personas no pueden escucharlo. Afecta a entre un 15% y un 20% de la población, sobre todo en adultos mayores.

Este zumbido es un problema de salud muy frecuente en nuestra sociedad y puede empeorar notablemente la calidad de vida de quienes lo padecen.

Las infecciones virales pueden generar síntomas relacionados con la audición. Esto se debe a que las membranas mucosas tienden a congestionarse y acumulan líquido detrás de los timpanos. Por eso, estos virus pueden dañar las células sensoriales del oído interno. No es de extrañar entonces que la enfermedad infecciosa causada por el virus SARS-COV-2 esté relacionada con el Tinnitus.

Principales síntomas

El especialista del Hospital Universitario Central de Asturias, Faustino Núñez Batalla, precisa que se define como 'persistente' aquel tinnitus que dura más de 6 meses, al tiempo que remarca que puede ocurrir en uno o ambos lados de la cabeza y ser notado como proveniente de dentro o de fuera de la misma.

"Más frecuentemente ocurre en el lado donde coexiste con una sordera, en particular en aquellos pacientes que describen una sensación más intensa", aprecia el especialista.

Por otro lado explica que se califica como 'tinnitus primario' aquel que no tiene una causa clara (idiopático) y que se puede asociar o no con una sordera; mientras que se define como 'tinnitus secundario' aquel que se asocia con una causa subyacente, con una enfermedad orgánica.

¿Cómo tratarlo?

En algunos casos, podrá ser tratado si la causa es una afección de salud subyacente. En este caso, el médico podrá reducir los síntomas tratando la causa. Por ejemplo:

- ⇒ Eliminar la cera de los oídos.
- ⇒ Tratamiento de una afección de los vasos sanguíneos.
- ⇒ Audífonos.
- ⇒ Cambios de medicamentos.

Según explica la Clínica Mayo, en muchos casos el Tinnitus no tiene cura, pero se pueden realizar tratamientos para hacer que los síntomas sean menos perceptibles, como por ejemplo, un dispositivo electrónico para inhibir el ruido:

- ⇒ Máquinas de ruido blanco: reproducen sonidos similares a los ambientales como la caída de la lluvia o las olas del mar. También, ventiladores, humidificadores y aire acondicionado pueden ayudar.
- ⇒ Dispositivos de enmascaramiento: se colocan en el oído y son similares a los audífonos.

Cuando nada de esto sirve, sólo queda el asesoramiento psicológico a través del tratamiento conductora para ayudar al paciente a vivir con el Tinnitus.

Fuente: ONDA CERO. Disponible en <https://bit.ly/3QiVKTQ>

Las vacunas anti-Covid-19 de Cuba, más allá de sus fronteras

5 ago. Además de Soberana Plus, ha llegado al Viejo Continente otra de las vacunas cubanas, Soberana 02.

Soberana 02, diseñada en el Instituto Finlay de Vacunas (IFV) de la isla caribeña, al igual que Soberana Plus; recibió luz verde en abril para ser producido parcialmente en Italia gracias a un acuerdo entre dicha entidad y la empresa ADIENNE Pharma & Biotech y la Agencia de Intercambios Económicos y Culturales con Cuba (AICEC) -ambas italianas-.

El memorándum suscrito en el congreso BioHabana 2022 permitirá formular y envasar en el país europeo el inmunógeno de la isla.

En esa ocasión, el director general del IFV, Vicente Verez, indicó que el acuerdo estaba cerrado pensando en un potencial momento de entrada de esta vacuna en Europa o, incluso, en América del Norte. Ello necesita de alianzas para compartir la ciencia".

Verez también ha explicado que la intención es, en una segunda fase, poder valorar si se podrían producir íntegramente en esta empresa italiana las vacunas Soberana 02 y otras de IFV.

¿HASTA DÓNDE HAN LLEGADO LAS VACUNAS CUBANAS CONTRA LA COVID-19?

SOBERANA

ITALIA
Soberana 02 será producida en Italia gracias a un acuerdo con la empresa ADIENNE Pharma & Biotech y la Agencia para el intercambio económico y cultural con Cuba (AICEC).

NICARAGUA y VENEZUELA
Soberana 02 y Soberana Plus, cuen-

IRÁN
Soberana 02 se produce también en Irán bajo el nombre PastoCorona.

BELARÚS
Belarús se convirtió en el primer país de Europa en registrar el fármaco anti-covid-19 de Cuba: Soberana Plus.

REPÚBLICA ÁRABE SAHARAUI DEMOCRÁTICA
Cuba donó un lote de 458 mil dosis de

Con Italia, los hilos que unen los productos anti-Covid-19 de IFV son más fuertes. En noviembre de 2021, alrededor de 30 personas de entre 19 y 59 años viajaron desde este país a Cuba para participar como voluntarias en el estudio clínico Soberana Plus Turín.

El objetivo del ensayo fue evaluar el inmunógeno en convalecientes de COVID-19 y sujetos sin antecedentes de la enfermedad, pero inmunizados con otras vacunas.

En el marco de este análisis colaborativo, en el hospital “Amadeo di Savoia” de la ciudad de Turín, se evalúan sueros de voluntarios cubanos vacunados con Soberana Plus desde julio del año pasado.

Los resultados demostraron la capacidad de Soberana Plus para inducir anticuerpos neutralizantes contra las variantes alfa, beta y delta del coronavirus SARS-CoV-2 que causa la enfermedad.

Primera en el mundo, especialmente diseñada para repotenciar la inmunidad previamente inducida por otras vacunas anti-Covid-19 o por infección natural, Soberana Plus también ha sido utilizada en Cuba como dosis de refuerzo en el régimen de vacunación anti-Covid-19, diseñado por IFV a partir de dos dosis de su otro producto Soberana 02.

La combinación de ambos ha demostrado en ensayos clínicos una eficacia del 92,4 % frente a la enfermedad sintomática, según el informe de resultados finales del estudio de fase III de Soberana 02.

Este fue el cronograma utilizado para la vacunación contra la COVID-19 en niños de 2 a 18 años en el país, la primera campaña en el mundo que llegó a más de 1,7 millones de infantes.

Las autoridades cubanas calificaron de “éxito total” la vacunación en la edad pediátrica; Pues se han prevenido alrededor de 70 000 casos y desde 2021 no ha muerto ningún niño por la COVID-19 en Cuba.

No es sólo en Europa donde se conocen los “soberanos”. Soberana 02 llegó a Irán a principios de 2021 cuando aún era solo un candidato vacunal, con el objetivo de completar la fase III de sus ensayos clínicos.

“Como parte de la colaboración con otros países en el desarrollo de vacunas contra la COVID-19, se han enviado al Instituto Pasteur de Irán 100.000 dosis de Soberana02, que serán utilizadas en ensayos clínicos en este país”, anunció el grupo de empresas BioCubaFarma.

A fines de junio del mismo año, Soberana 02 recibió la autorización de uso de emergencia en la República Islámica de Irán. La autorización fue concedida al Instituto Pasteur de Irán (IPI), que comercializará la vacuna en territorio iraní, en el marco de un acuerdo de colaboración suscrito con el Instituto Finlay de Vacunas.

De esta forma, el país persa se convierte en el primero del mundo en producir vacunas cubanas contra la COVID-19.

En este país se inauguró un centro de producción masiva de la inyección de la vacuna PastoCorona, resultado de la transferencia, al Instituto Pasteur, de la tecnología de la vacuna cubana Soberana 02.



De este lado del mundo, en Nicaragua y Venezuela; Soberana 02 y Soberana Plus están autorizados para uso de emergencia.

Soberana 02 se encuentra entre las vacunas aprobadas por las autoridades sanitarias venezolanas para su uso en niños mayores de dos años y a principios de 2022 llegó al país bolivariano un envío de un millón de dosis de Soberana Plus.

Las autoridades sanitarias han aclarado que utilizarán este último medicamento para brindar protección contra la reinfección con la variante Ómicron.

Por su parte, la Autoridad Reguladora de Nicaragua ha indicado desde octubre de 2021 que Soberana 02 se ofrece como una herramienta terapéutica de acceso seguro para reducir la transmisibilidad del COVID-19; particularmente en la población pediátrica de dos a 17 años de edad.

Soberana 02 también ha llegado a los brazos de los niños de la República Árabe Saharaui Democrática. En febrero, Cuba donó a ese país un lote de 458 000 dosis de esta vacuna para uso pediátrico.

Abdala también protege fuera de Cuba

Aunque los vacunas de la línea de Soberana ya son conocidos en varias naciones, lo mismo sucede con Abdala, vacuna diseñada en el Centro de Ingeniería Genética y Biotecnología de Cuba y la primera en este país y en América Latina diseñada contra el virus SARS-CoV-2.

A Vietnam, México, Venezuela y Nicaragua han llegado millones de dosis de esta vacuna que, aplicada en un esquema de tres dosis cada 14 días, ha demostrado una efectividad del 92,28% contra enfermedades sintomáticas. Además, ha demostrado una eficacia del 100% en la prevención de enfermedades sistémicas graves y muertes por COVID-19.

En septiembre de 2021, a pedido del Centro de Investigación y Producción de Vacunas y Productos Biológicos Médicos, el Ministerio de Salud de Vietnam aprobó la importación y uso de la vacuna Abdala en esta nación indochina.

Durante ese mismo mes llegó a ese país el primer lote de la vacuna cubana, compuesto por 900 000 dosis compradas por Vietnam y otras 150 000 donadas por la nación caribeña.

Durante el recibimiento de ese primer envío, el viceministro de Relaciones Exteriores Dang Hoang Giang agradeció a Cuba las dosis, de un total de 10 millones acordadas con la mayor de las Antillas y desde donde también se transferirá a Vietnam la tecnología para producir la vacuna en Ese pais.

En Nicaragua, Abdala está aprobado para uso de emergencia desde octubre de 2021; mientras que Venezuela firmó un contrato para suministrar 12 millones de unidades del inmunógeno con la isla caribeña el mismo año.

A finales de 2021, la Comisión Federal para la Protección contra Riesgos Sanitarios (Cofepris) del gobierno mexicano se pronunció sobre la autorización de uso de emergencia de Abdala.

Como autoridad reguladora nacional de referencia, calificada por la Organización Panamericana de la Salud; Las decisiones de la Cofepris son reconocidas por varios países de la región, por lo que es probable que las vacunas aprobadas se utilicen en otros países.



Como miembro de la Conferencia Internacional sobre Armonización de Requisitos Técnicos para el Registro de Productos Farmacéuticos de Uso Humano, todas las decisiones de esta autoridad se toman sobre la base de la evidencia técnico-científica presentada.

En diciembre del mismo año, el Ministerio de Salud de San Vicente y las Granadinas anunció en su cuenta de redes sociales de Facebook que Abdala estaba disponible en los puntos de vacunación de ese país.

Con este anuncio, esta nación se convierte en la primera de la Comunidad del Caribe (Caricom) en autorizar una de las vacunas de Cuba contra la Covid-19.

¿Y la certificación de la OMS?

A pesar de toda la evidencia presentada anteriormente, muchos escépticos dirán que las vacunas cubanas aún no son reconocidas por la Organización Mundial de la Salud (OMS); y no son sin razón; solo aclara eso; Aunque el problema ha durado más de lo esperado, cada país puede autorizar el uso de una vacuna, esté o no aprobada por este organismo internacional.

Sin embargo, esto no significa que Cuba esté detenida en el proceso.

Así lo confirmó recientemente el director general de la OMS, Tedros Adhanom, quien aseguró en mayo pasado que el organismo estaba siguiendo el proceso de certificación de las vacunas anti-Covid-19 creadas y desarrolladas en Cuba.

Desde marzo, el Grupo de las Industrias Biotecnológica y Farmacéutica de la Nación Caribeña (BioCubaFarma) ha detallado en su cuenta de Twitter que la OMS ha sido informada del documento finalizado para ser revisado por expertos e iniciar el proceso de reconocimiento internacional de la vacuna Abdala.

La entidad señaló en esta misma red social que el Centro de Ingeniería Genética y Biotecnología (CIGB) a cargo de esta vacuna inició durante este mes el intercambio formal con la entidad de salud.

A mediados de febrero de este año, el director de BioCubaFarma, Eduardo Martínez, explicó que la empresa estaba trabajando en el texto, compuesto por varios capítulos con resultados sobre la investigación clínica y preclínica, el desarrollo farmacéutico, así como todo lo referente a las instalaciones de producción, un aspecto al que se han hecho adaptaciones.

Sobre los procesos seguidos con Abdala para esta evaluación, Martínez señaló que se tomó la decisión de trasladar el sitio de producción a la recién inaugurada fábrica de Mariel, ubicada en este polo industrial, al oeste de La Habana.

En funcionamiento desde noviembre de 2021, el Parque Tecnológico Industrial CIGB-Mariel es considerado el más moderno de Cuba y uno de los más avanzados de América Latina y el Caribe con laboratorios de control de calidad y almacenes de materias primas y productos terminados.

También cuenta con fábricas para obtener el principio activo de vacunas e inmunógenos completos en formulaciones líquidas, en polvo y spray.

El objetivo es que los representantes de la OMS visiten también las plantas productivas de esta entidad donde realizarán la inspección necesaria para posteriormente obtener el permiso y su inclusión en la lista de productos reconocidos por el organismo de las Naciones Unidas, subrayó Martínez.

“Cuba siempre mantiene intercambios con la representación de la OMS/OPS sobre todo lo relacionado con la precalificación de vacunas anti-Covid-19”, dijo.

Y la afirmación no es menos cierta, desde septiembre del año pasado, el representante de la Organización Panamericana y Mundial de la Salud (OPS/OMS) en la isla caribeña, José Moya, informó a la agencia AP un encuentro virtual entre especialistas en La Habana, Ginebra y Washington para compartir información, coordinar documentación y establecer cronogramas.

La solidaridad de Cuba no sólo ha llenado de batas blancas al mundo; También llega a varios países en ampollas que pueden salvar millones de vidas. Las vacunas desarrolladas en esta isla cuentan con el compromiso, esfuerzo y capacidad de científicos con más de tres décadas de experiencia produciendo sus propios inmunógenos.

Fuente: Cuba ES Euro. Disponible en <https://bit.ly/3SQ6LE2>

Prestigiosa revista científica de oncología publica resultados de investigación entre el CIM y el Roswell Park con la vacuna cubana CIMAvax-EGF

5 ago. La prestigiosa revista científica *Frontiers in Oncology* publicó este 3 de agosto resultados de la investigación entre el CIM y el Instituto Roswell Park, de Nueva York, en Estados Unidos, con la vacuna terapéutica cubana contra el cáncer de pulmón CIMAvax-EGF.

Así dio a conocer en su cuenta de Twitter el Centro de Inmunología Molecular (CIM).

Se trata de un estudio en curso por la institución científica radicada en Nueva York, en el escenario más avanzado de los pacientes en la segunda línea del cáncer de pulmón de células no pequeñas.

“En este estudio no se usa la vacuna solamente, sino que es una combinación de CIMAvax-EGF® con Nivolumab, una droga inmunomoduladora. En este caso también demostramos que la combinación fue muy segura e incrementó la inmunogenicidad.

“Al mismo tiempo tenemos un conjunto de pacientes que está obteniendo una supervivencia muy larga. Ello nos brinda la posibilidad de identificar nuevos biomarcadores que nos permitan anticipar poblaciones que reciban un beneficio de la combinación de ambas drogas”, explicó recientemente a Cubadebate la doctora Tania Crombet, directora de Investigaciones Clínicas del CIM.

Más de 10 000 pacientes cubanos han sido beneficiados con esta vacuna en los diferentes escenarios de atención médica (primaria, hospitalares e institutos) del sistema nacional de Salud.

Los datos del mundo real de uso de CIMAvax-EGF® —como se le denomina a la investigación clínica una vez que se concluye el registro del producto—, confirman los datos de seguridad y efectividad que se habían generado en las investigaciones cubanas previas. De igual modo, los datos obtenidos por el Instituto Roswell Park.



Frontiers in Oncology es la tercera revista más citada en su campo con más de 155 000 citas y 60 millones de visitas en 17 000 artículos. La publicación de esta investigación confirma los resultados obtenidos en los estudios que se están conduciendo en Cuba actualmente con el fármaco y constituye un importante paso en la visibilización de la ciencia cubana.

Fuente: Cubadebate. Disponible en <https://bit.ly/3QHvfxD>

Cuarta dosis y nuevas vacunas

7 ago. Hemos evidenciado el gran triunfo de la ciencia con las vacunas frente a la COVID-19; se han aprobado varias vacunas en Europa, dos de ellas basadas en una nueva tecnología de ARN, de las empresas Moderna y Pfizer-BioNTech, dos empleando adenovirus (AstraZeneca-Oxford y Janssen-Cilag), una con virus inactivado (Valneva Austria) y otra con proteína recombinante (Novavax). Actualmente están en fase de estudio por la Agencia Europea del Medicamento, vacunas de las empresas Sanofi-GSK, Hipra (vacuna española que emplea dos proteínas recombinantes de las variantes alfa y beta), Moderna y Pfizer (que incluyen a la nueva variante Ómicron), así como muchas más en investigación.

► <https://www.ema.europa.eu/en/human-regulatory/overview/public-health-threats/coronavirus-disease-covid-19/treatments-vaccines/vaccines-covid-19/covid-19-vaccines-authorised>

Actualmente la pandemia en España ha pasado a una fase de estabilización, con una disminución de los casos activos y de personas hospitalizadas, y con una alta tasa de vacunación. Ante esta situación actual cabe preguntarse qué hacer en los próximos meses en relación a las vacunas. Los estudios científicos han demostrado que tanto la vacunación como haber pasado la infección inducen una respuesta inmunitaria que genera memoria, y que ésta puede ser duradera (no sabemos cuánto tiempo). Incluso la combinación infección-vacunación, ha mostrado generar mejor respuesta, al ver nuestra inmunidad al virus completo, actuando una diversidad de «soldados inmunitarios» y no solo los que ven la proteína S (presente en las vacunas disponibles). Esto también se observa al combinar vacunas distintas.

La memoria inmunitaria (tanto celular como humoral) se mantiene durante tiempo, pero los anticuerpos, esas proteínas tan específicas que pueden bloquear y neutralizar al virus, decaen tras varios meses. En una persona sana y joven que se vacunó y/o infectó previamente, si se contagia con una nueva variante, es previsible que tenga síntomas durante unos días, pero rápidamente resolverá la infección. Por el contrario, una persona vulnerable (mayor, con patologías, trasplantados, etcétera), si se contagia puede desarrollar



una enfermedad grave. La estrategia futura de vacunación debe por tanto centrarse en aquellos más vulnerables (ancianos y con alto riesgo de sufrir una enfermedad grave), pero no en la población general, como aseguran el Centro Europeo para el Control y Prevención de Enfermedades y la Agencia Europea del Medicamento.

► <https://www.ema.europa.eu/en/news/ecdc-ema-issue-advice-fourth-doses-mrna-covid-19-vaccines>

La pregunta que surge es si las personas con mayor riesgo, deben recibir una cuarta dosis de la misma vacuna, combinar vacunas ya autorizadas, o esperar a la aprobación otras nuevas, como las que incluyen ya a la variante ómicron. Los estudios realizados en Israel con cuatro dosis de Pfizer indican que se incrementa la protección frente a enfermedad grave/fallecimientos, pero que los anticuerpos vuelven a decaer pasados unos 4 meses, por lo que no parece ser la mejor estrategia a largo plazo. Los ensayos llevados a cabo combinando vacunas autorizadas han mostrado buenos resultados, así como también los de las nuevas vacunas, aún en fase de aprobación.

El contagio con el virus moviliza también a la inmunidad, por lo que una persona inoculada con 3 dosis y que ha sufrido la infección, su sistema inmunitario se ha activado ya 4 veces. Otro elemento importante: no cansar siempre a los mismos soldados inmunitarios y esperar 5-6 meses tras una infección para recibir una dosis de vacuna.

Fuente: La Voz de Galicia. Disponible en <https://bit.ly/3bNd2jz>

El ensayo clínico PANCOVID liderado por el Hospital La Paz de Madrid ha sido publicado en la revista Clinical Infectious Diseases

9 ago. Los resultados del ensayo clínico español PANCOVID, diseñado para evaluar la eficacia de diversas alternativas farmacológicas, y estrategias en el tratamiento de SARS-CoV-2 sintomático, han sido publicados en la revista Clinical Infectious Diseases, una de las más importantes de su campo. Se trata de uno de los ensayos académicos sobre estrategias de tratamiento de COVID-19 más grande realizado en España.

Los resultados del ensayo clínico español PANCOVID, diseñado por el Hospital Universitario La Paz - IIdiPAZ para evaluar la eficacia de diversas alternativas farmacológicas, y estrategias en el tratamiento de SARS-CoV-2 sintomático, han sido publicados en la revista Clinical Infectious Diseases, una de las más importantes de su campo.

El ensayo ha sido liderado por los doctores José Ramón Arribas, jefe de Sección de Medicina Interna y enfermedades infecciosas, Alberto Borobia, coordinador de la Unidad de Ensayos Clínicos y Antonio Carcas del servicio de Farmacología Clínica, todos ellos pertenecen al CIBERINFEC. El objetivo principal del ensayo clínico PANCOVID ha sido proporcionar estimaciones fiables sobre los efectos de diferentes estrategias de tratamiento de COVID-19 sintomático.

En el estudio se investigó la eficacia de Emtricitabina/Tenofovir disoproxil fumarato (TDF/FTC) y de Baricitinib en pacientes, principalmente hospitalizados, con alto riesgo de COVID-19 grave durante un año. La investigación ha sido impulsada y promovida por el CIBER de Enfermedades Infecciosas y ha sido financiado por el Instituto de Salud Carlos III, en la convocatoria extraordinaria de ayudas para la investigación de la COVID-19.

Se incluyó a un total de 355 participantes, donde el criterio de valoración principal fue la mortalidad a los 28 días, que era del 3,1%. El riesgo relativo de mortalidad a los 28 días para los participantes tratados con TDF/FTC fue de 1,76% y del 0,42% para los tratados con Baricitinib. Estos resultados no sugirieron un efecto beneficioso de TDF/FTC en pacientes con COVID hospitalizados; sin embargo, son compatibles con el efecto beneficioso de Baricitinib ya establecido por otros ensayos clínicos. Los resultados del ensayo han servido para afianzar el papel de Baricitinib como terapia antiinflamatoria en pacientes con COVID grave.

Fuente: Comunidad de Madrid. Disponible en <https://bit.ly/3AkHRoX>

Prestigiosa revista europea publica artículo del desarrollo de la vacuna Abdala

9 ago. El Dr. Gerardo Enrique Guillén, director de Investigaciones Biomédicas del Centro de Ingeniería Genética y Biotecnología (CIGB) informó a través de su cuenta en Twitter que se publicó en revista de la Sociedad Europea de Biotecnología el artículo del desarrollo de la vacuna Abdala que comprende desde la construcción genética hasta la evaluación en modelos animales.

La publicación de este artículo en una revista de alto impacto constituye un reconocimiento a la calidad del desarrollo de la ciencia cubana, en tanto, estos resultados fueron arbitrados por expertos internacionales para su publicación, un proceso que se realiza a ciegas para los autores.

La revista *New Biotechnology* tiene un alto factor de impacto dentro de las publicaciones científicas, con un estimado de 6.49 en el 2021.

Abdala es un inmunógeno desarrollado por el CIGB, siendo una de las tres vacunas creadas por Cuba ante la pandemia.

La misma se ha aplicado como parte de la campaña de vacunación masiva anti COVID impulsada por la nación caribeña.

Fuente: Cubadebate. Disponible en <https://bit.ly/3JZN3lp>



OMS: Muertes por COVID-19 a nivel mundial disminuyeron nueve por ciento durante la última semana

10 ago. De acuerdo con el último reporte semanal de la OMS, publicado este miércoles, la cifra de fallecimientos como consecuencia del coronavirus disminuyó un 9 % durante la última semana. Asimismo, el número de nuevos casos se mantuvo estable.

La agencia de Naciones Unidas informó que en la última semana se produjeron más de 14 000 decesos por COVID-19 y casi siete millones de contagios.

Los casos aumentaron un 30 % en el Pacífico occidental y disminuyeron un 46 % en África.

En América y el Medio Oriente, la cifra de contagios se redujo en más del 20 %.

Los decesos aumentaron 19 % en Oriente Medio, pero cayeron más del 70 % en el continente africano, 15% en Europa y 10% en África.

La subvariante Ómicron BA.5 sigue predominando en el mundo y representa casi el 70 % de las secuencias de virus reportadas, según la OMS. Los datos indican que otras subvariantes de Ómicron parecen disminuir.

La Organización Mundial de la Salud aclaró que su evaluación de las tendencias de la COVID-19 continúa con limitaciones por el abandono de los países de una parte de los esfuerzos de vigilancia, pruebas y secuenciación, como consecuencia de la relajación de los controles de la pandemia.

Fuente: Cubadebate. Disponible en <https://bit.ly/3JRvl3o>



USS realiza estudios clínicos para el desarrollo de nuevas vacunas

10 ago. USS participa en 5 estudios clínicos para el desarrollo de nuevas vacunas. Se trata de nuevas vacunas contra la COVID-19, Influenza y Neumococo. A su vez, la institución trabaja en la conformación de un centro de estudios clínicos de referencia.

El avance de las enfermedades, la aparición de nuevos agentes infecciosos o la resistencia de algunos virus o bacterias a las terapias actuales, obliga al mundo de la ciencia a buscar nuevas alternativas de tratamiento.

En ese contexto, son muy importantes los estudios clínicos. Se trata de investigaciones que buscan probar la eficacia y seguridad de nuevos medicamentos, vacunas, dispositivos u otras fórmulas que se desarrollan con el fin de encontrar mejores formas de prevenir, diagnosticar o tratar una enfermedad.

Cada estudio clínico tiene diferentes etapas y un protocolo muy riguroso que se debe cumplir. En el caso de Chile, es el Instituto de Salud Pública (ISP) el organismo encargado de autorizar la realización de estos ensayos con el fin de proteger los derechos y el bienestar de los participantes y asegurar la calidad de los datos obtenidos en la investigación. Se requiere también de la aprobación por Comités de Ética Científica independientes, que aseguran que los estudios cumplan con todas las normas internacionales de buenas prácticas clínicas de investigación.

Centro USS

La Universidad San Sebastián participa en cinco estudios clínicos multicéntricos para probar nuevas vacunas. Se trata de vacunas contra el SARS-CoV-2, Influenza y Neumococo, tres microorganismos respiratorios que causan gran carga de morbilidad en el país.

“Aspiramos a ser un centro de referencia a nivel nacional para el desarrollo de estudios clínicos que impacten en resultados de pacientes, en la práctica clínica y la salud pública”, explica Pilar Espinoza, directora de investigación, postgrado y vinculación internacional de la Facultad de Ciencias de Ciencias para el Cuidado de la Salud.

La docente comenta que actualmente se está trabajando en la conformación de un Centro de Estudios Clínicos USS, de la mano de la Facultad de Medicina y Ciencia.

Estudios

La primera experiencia en esta área fue la participación de la USS en el estudio multicéntrico para evaluar la eficacia, seguridad e inmunogenicidad de la vacuna contra el virus SARS-CoV-2 (CoronaVac), liderado en Chile por la Pontificia Universidad Católica y el laboratorio Sinovac Biotech, donde se invitó a la Universidad a participar junto a otras instituciones.

Según explicó Espinoza, esto “nos permitió darnos cuenta de que contamos con el know how necesario para convertirnos en un centro de referencia, al mismo tiempo que empezamos a ser requeridos por parte de otras instituciones para hacer este trabajo”.

Actualmente la Universidad participa en otros cuatro Ensayos Clínicos Fase III, es decir buscan responder si la nueva intervención bajo estudio es no inferior o superior a la convencional, explica el Dr. Carlos Pérez, Decano de la Facultad de Medicina y Ciencia e investigador responsable de estos estudios junto con la profesora Espinoza.

Uno de ellos es el estudio de una nueva vacuna tetravalente contra la Influenza de Sinovac Biotech que pretende evaluar la inmunogenicidad y seguridad del producto farmacéutico en comparación con otra vacuna tetravalente, disponible en el mercado, para ser usada por personas mayores de tres años y que es patrocinado por la PUC y Sinovac Biotech (Chile).

Otros dos estudios son para evaluar la seguridad, la tolerancia e inmunogenicidad de una nueva vacuna contra el neumococo, en diferentes grupos de la población.

Espinoza agrega que se está en conversaciones con Sinovac Biotech (Chile) para comenzar próximamente el estudio de una nueva vacuna contra el SARS-CoV-2, variante Ómicron.

“Nuestro objetivo es generar conocimiento científico que responda a las problemáticas de salud relevantes de la población, de manera de socializar estos resultados con la comunidad académica y científica a nivel local e internacional. Estamos capacitados para entregar guía experta, soporte metodológico, logístico y estructural para el desenvolvimiento de estudios clínicos”, concluye Pilar Espinoza.

Fuente: IPSUSS. Disponible en <https://bit.ly/3Pqlsd0>





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Estrategia de búsqueda: Vaccine in the title or abstract AND 20220801:20220811 as the publication date 44 records

1.[WO/2022/162205](#)VACCINE COMPOSITION FOR BREAKING SELF-TOLERANCE

WO - 04.08.2022

Clasificación Internacional [C07K 16/24](#) Nº de solicitud PCT/EP2022/052154 Solicitante BAYER ANIMAL HEALTH GMBH Inventor/a ILG, Thomas

The present invention relates to a vaccine composition for breaking self-tolerance against a self-protein of a host, in particular for breaking self-tolerance against endogenous cytokines, in particular against the endogenous IL-4, IL-5, IL-13, IL-31 and IL-33 proteins in an animal host. The vaccine composition of the

invention contains a polyprotein, a DNA encoding for the polyprotein and/or an RNA encoding for the polyprotein and one or more immunostimulatory oligonucleotides. The polyprotein comprises at least two self-protein segments of the host and one or more T-cell epitopes of non-host origin in between and/or adjacent to the at least two self-protein segments. The present invention further concerns the use of the vaccine composition for the prevention and/or treatment of diseases including the prevention and/or treatment of a pruritic condition and/or an allergic condition. In another aspect, the present invention provides a method for detecting the presence of autoantibodies against self-proteins that can be generated with the vaccine composition of the invention.

2.WO/2022/162204VACCINE COMPOSITION FOR BREAKING SELF-TOLERANCE

WO - 04.08.2022

Clasificación Internacional C07K 16/24 Nº de solicitud PCT/EP2022/052153 Solicitante BAYER ANIMAL HEALTH GMBH Inventor/a ILG, Thomas

The present invention relates to a vaccine composition for breaking self-tolerance against a self-protein of a host, in particular for breaking self-tolerance against endogenous cytokines in an animal host. The vaccine composition of the invention contains a polyprotein, a DNA encoding for the polyprotein and/or an RNA encoding for the polyprotein and one or more immunostimulatory oligonucleotides. The polyprotein comprises at least two self-protein segments of the host and one or more T-cell epitopes of non-host origin in between and/or adjacent to the at least two self-protein segments. The present invention further concerns the use of the vaccine composition for the prevention and/or treatment of diseases including the prevention and/or treatment of a pruritic condition and/or an allergic condition. In another aspect, the present invention provides a method for detecting the presence of autoantibodies against self-proteins that can be generated with the vaccine composition of the invention.

3.20220241390VACCINE COMPOSITION COMPRISING RECOMBINANT PROTEIN OF STAPHYLOCOCCUS AUREUS ATTENUATED ENTEROTOXIN AND CYTOTOXIN

US - 04.08.2022

Clasificación Internacional A61K 39/085 Nº de solicitud 17686606 Solicitante REPUBLIC OF KOREA (ANIMAL AND PLANT QUARANTINE AGENCY) Inventor/a Dong Chan MOON

The present invention relates to a vaccine composition comprising a *Staphylococcus aureus* attenuated enterotoxin protein and cytotoxin protein, and more particularly to a *Staphylococcus aureus* enterotoxin protein, a *Staphylococcus aureus* cytotoxin protein, a vaccine composition for prevention of bovine mastitis, comprising the *Staphylococcus aureus* enterotoxin protein and *Staphylococcus aureus* cytotoxin protein and a method for preventing bovine mastitis comprising administering the vaccine composition to a bovine. The *Staphylococcus aureus* enterotoxin protein, the *Staphylococcus aureus* cytotoxin protein according to the present invention, and the vaccine composition comprising the proteins as an antigen can be used so that even vaccines comprising several antigens rather than all kinds of antigens show the excellent effects of prevention and treatment of bovine mastitis against all kinds of *Staphylococcus aureus* enterotoxin and cytotoxin having high incidence in Korea, thereby being more economically used for industrial purposes. Further, the vaccine composition for prevention of bovine mastitis, comprising the *Staphylococcus aureus* enterotoxin protein and *Staphylococcus aureus* cytotoxin protein according to the present invention has an excellent safety and bovine mastitis prevention and treatment effect even in the high CFU *Staphylococcus aureus* challenge test so that the composition can be variously utilized in *Staphylococcus aureus* vaccine and prevention related fields in future.

4.20220241409WATER SOLUBLE ADJUVANT AND COMPOSITION CONTAINING SAME

US - 04.08.2022

Clasificación Internacional A61K 39/39 Nº de solicitud 17600903 Solicitante Sumitomo Dainippon Pharma Co., Ltd. Inventor/a Hitoshi Ban

The present invention relates to a compound useful as a vaccine adjuvant for cancer vaccine, a preparation process thereof, a pharmaceutical composition comprising the compound, and use of the compound as a vaccine adjuvant for cancer vaccine.

5.[WO/2022/162370](#) ANTI-VIRAL THERAPEUTIC

WO - 04.08.2022

Clasificación Internacional [C07K 16/08](#) N° de solicitud PCT/GB2022/050217 Solicitante UNIVERSITY COLLEGE CARDIFF CONSULTANTS LIMITED Inventor/a STANTON, Richard

The invention relates to an anti-viral composition comprising at least one, and ideally a plurality of, monoclonal antibodies, or fragments thereof; an immunogenic agent, vaccine or pharmaceutical composition comprising the afore anti-viral composition; said anti-viral composition, immunogenic agent, vaccine or said pharmaceutical composition for use in the treatment of or prevention of a viral infection; use of said anti-viral composition in the manufacture of a medicament to treat or prevent a viral infection; a combination therapeutic for use in the treatment or prevention of a viral infection comprising said anti-viral composition, immunogenic agent, vaccine or pharmaceutical composition in combination with at least one other therapeutic agent; and a method of treating or preventing a viral infection comprising administering said anti-viral composition, immunogenic agent, vaccine or said pharmaceutical composition to an individual having, or suspected of having, a viral infection.

6.[20220241389](#) VACCINE FOR THE PREVENTION OF BREAST CANCER RELAPSE

US - 04.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 17574178 Solicitante The Henry M. Jackson Foundation for the Advancement of Military Medicine, Inc. Inventor/a George E. PEOPLES

The invention features methods to induce and maintain a protective cytotoxic T-lymphocyte response to a peptide of the HER2/neu oncogene, E75, with the effect of inducing and maintaining protective or therapeutic immunity against breast cancer in a patient in clinical remission. The methods comprise administering to the patient an effective amount of a vaccine composition comprising a pharmaceutically acceptable carrier, an adjuvant such as recombinant human GM-CSF, and the E75 peptide at an optimized dose and schedule. The methods further comprise administering an annual or semi-annual booster vaccine dose due to declining E75-specific T cell immunity. The invention also features vaccine compositions for use in the methods.

7.[WO/2022/163902](#) VACCINE COMPOSITION FOR PREVENTING HUMAN INFECTIOUS SARS CORONAVIRUS AND ALLEVIATING INFECTION SYMPTOMS

WO - 04.08.2022

Clasificación Internacional [C12N 15/86](#) N° de solicitud PCT/KR2021/003046 Solicitante LIBENTECH CO.,LTD. Inventor/a JANG, Hyun

The present invention relates to a vaccine composition for preventing human infectious SARS coronavirus (SARS-CoV-2, COVID-19) and alleviating infection symptoms. The vaccine composition, of the present invention, comprising a recombinant Newcastle disease virus, which expresses a SARS-CoV-2 RBD protein on the surface thereof, or an antigen purified therefrom induces immune responses that can fight COVID-19 infection, and thus can be effectively used as a vaccine for preventing and treating SARS-CoV-2 infection.

8.[WO/2022/161969](#) IMMUNOGENIC COMPOSITION AND VACCINE CONTAINING CHLAMYDIA SSP. SURFACE ANTIGENS AND ITS USE

WO - 04.08.2022

Clasificación Internacional [A61K 39/118](#) N° de solicitud PCT/EP2022/051669 Solicitante MEDIZINISCHE HOCHSCHULE HANNOVER Inventor/a KLOS, Andreas

The present invention relates in a first aspect to an immunogenic composition comprising at least three Chlamydia ssp. surface antigens selected from the group of PmpA, PmpD, PmpG, PmpH and antigen Ctad1. Further, the present invention relates to the immunogenic composition comprising CDN as adjuvant, in particular, c-diAMP. Moreover, a pharmaceutical composition comprising the immunogenic composition according to the present invention is provided as well as a vaccine comprising said immunogenic composition. The vaccine is particularly useful in eliciting an immune response against Chlamydia ssp. in an animal, including a human, in particular, for use in treating or preventing the infection by Chlamydia ssp. The vaccine, pharmaceutical composition or immunogenic composition may be administered mucosally, preferably is administered at least three times.

9. 20220241395 INACTIVATING PATHOGENS AND PRODUCING HIGHLY IMMUNOGENIC INACTIVATED VACCINES USING A DUAL OXIDATION PROCESS

US - 04.08.2022

Clasificación Internacional [A61K 39/145](#) N° de solicitud 17497810 Solicitante Najít Technologies, Inc.
Inventor/a Ian J. Amanna

Provided are surprisingly effective methods for inactivating pathogens, and for producing highly immunogenic vaccine compositions containing an inactivated pathogen rendered noninfectious by exposure to a Fenton reagent, or by exposure to a Fenton reagent or a component thereof in combination with a methisazone reagent selected from the group consisting of methisazone, methisazone analogs, functional group(s)/substructure(s) of methisazone, and combinations thereof. The methods efficiently inactivate pathogens, while substantially retaining pathogen antigenicity and/or immunogenicity, and are suitable for inactivating pathogens, or for the preparation of vaccines for a wide variety of pathogens with genomes comprising RNA or DNA, including viruses and bacteria. Also provided are highly immunogenic inactivated vaccine compositions prepared by using any of the disclosed methods, and methods for eliciting an immune response in a subject by administering such vaccine compositions.

10. WO/2022/162177 CHLAMYDIA TRACHOMATIS ANTIGENIC POLYPEPTIDES AND USES THEREOF FOR VACCINE PURPOSES

WO - 04.08.2022

Clasificación Internacional [C07K 16/28](#) N° de solicitud PCT/EP2022/052104 Solicitante INSERM (INSTITUT NATIONAL DE LA SANTÉ ET DE LA RECHERCHE MÉDICALE) Inventor/a LEVY, Yves Chlamydiae are intracellular bacterial pathogens responsible for a variety of infections. The inventors have set up candidate vaccines against Chlamydia trachomatis. In particular, the inventors have identified specific epitopes to be included in vaccine candidates thanks to in silico analysis of the amino-acid sequence of these proteins to map predicted MHC-I and -II epitopes by online software (NetMHC-4.0 and NetMHCII-2.3) and peptide binding prediction software. B cell epitopes were also mapped using online software (BepiPred-2.0 and Discotope). Finally, the inventors have generated some specific CD40 or Langerin antibodies comprising one or more identified epitope(s) of the present invention and that are suitable for vaccine purposes. Therefore, the present invention relates to Chlamydia trachomatis (Ct) antigenic polypeptides and uses thereof for vaccine purposes.

11. WO/2022/163647 ORAL CORONAVIRUS INFECTION VACCINE

WO - 04.08.2022

Clasificación Internacional [C12N 1/21](#) N° de solicitud PCT/JP2022/002677 Solicitante NATIONAL UNIVERSITY CORPORATION KOBE UNIVERSITY Inventor/a SHIRAKAWA, Toshiro

Provided is a vaccine that can be administered orally against coronavirus infection. A vaccine that can be administered orally against coronavirus infection can be provided by transformed bifidobacteria designed so as to present some or all of the proteins that constitute the coronavirus on the bifidobacterium surface layer. Transformed bifidobacteria designed so as to present some or all of the proteins that constitute the

coronavirus on the bifidobacterium surface layer induce humoral immunity and cellular immunity by oral administration and can suppress any increase in severity of pneumonia, etc., even after viral infection.

12. [4036226](#) ABGESCHWÄCHTES AFRIKANISCHES SCHWEINEPESTVIRUS UND SEINE VERWENDUNG ALS IMPFSTOFF

EP - 03.08.2022

Clasificación Internacional [C12N 7/00](#) N° de solicitud 21305138 Solicitante AGENCE NAT CHARGEÉE DE LA SECURITE SANITAIRE DE L'ALIMENTATION DE L'ENVIRONNEMENT ET DU TRAVAIL Inventor/a BLOT LE POTIER MARIE-FRÉDÉRIQUE

The present invention relates to an attenuated African Swine Fever (ASF) virus, wherein :• genes MGF 360-12L, 360-13L, 360-14L, 505-2R, 505-3R are deleted or are interrupted or mutated such that the genes are not transcribed and/or translated,• ORF of ASFV_G_ACD_00520 is deleted or is interrupted or mutated such that it is not transcribed and/or translated, and• genes MGF 505-1R et 505-4R are truncated, compared to the genome of the corresponding unattenuated virus. The present invention also refers to a vaccine comprising the attenuated ASF virus, and its use in preventing African Swine Fever in a subject. The present invention also relates to an in-vitro method for obtaining the attenuated ASF virus, which comprises at least one step of thermal-attenuation of a virulent ASF virus strain selected among Georgia 2007/1, Pig/HLJ/2018, a strain of ASF virus of genotype II or a genetically close ASF virus strain, and amplification by inoculation of Specific-Pathogen-Free pigs and selecting said attenuated ASF virus. The present invention refers to an in vitro method for the differential detection of the attenuated ASF virus and of the corresponding non-attenuated ASF virus as well.

13. [4034148](#) BIOMATERIALBASIERTER ANTIGENFREIER IMPFSTOFF UND VERWENDUNG DAVON
EP - 03.08.2022

Clasificación Internacional [A61K 38/19](#) N° de solicitud 20868441 Solicitante HARVARD COLLEGE Inventor/a NAJIBI ALEXANDER J

Disclosed herein are vaccine compositions and method to use the same. The compositions and methods disclosed herein provide means to prevent and/or treat a variety of cancers.

14. [4034158](#) BIOLOGISCH ABBAUBARE NANOKOMPLEX-IMPFSTOFFE, VERFAHREN ZUR UNTERDRÜCKUNG DER HEPATITIS-B-VIRUS-REPLIKATION UND DER HEPATITIS-B-VIRUS-OBERFLÄCHENANTIGENSEKRETION

EP - 03.08.2022

Clasificación Internacional [A61K 39/29](#) N° de solicitud 20868095 Solicitante ASCENDO BIOTECHNOLOGY INC Inventor/a HUANG PING-YEN

A hepatitis B virus (HBV) vaccine includes an HBV core antigen (HBcAg) and/or HBV surface antigen (HBsAg) formulated in nanocomplexes. The nanocomplexes contain chitosan and g-PGA. These nanocomplexes containing HBc/sAg, chitosan, and g-PGA can induce more balanced T helper cells (Th1 and Th2) polarization than can a conventional vaccine with an alum adjuvant. HBc/s-NC of the invention can elicit high levels of antibodies against HBsAg, a rapid elimination of HBsAg, and a slow decrease of HBeAg, indicating a phenomenon of HBsAg seroconversion. Thus, HBc/s-NC can overcome immune tolerance caused by chronic HBV infection to re-establish host immunity leading a functional cure.

15. [2603362](#) Immunogenic compositions against enteric diseases and methods for its preparation thereof
GB - 03.08.2022

Clasificación Internacional [A61K 39/112](#) N° de solicitud 202204276 Solicitante SERUM INSTITUTE OF INDIA PVT LTD Inventor/a RAJEEV MHALASAKANT DHERE

The present disclosure relates to novel immunogenic monovalent and multivalent polysaccharide-protein conjugate vaccine compositions comprising a polysaccharide selected from *Salmonella* serovar strains *S. typhi*; *S. paratyphi A*; *S. typhimurium* and *S. enteritidis* and alternative improved methods of

polysaccharide fermentation, polysaccharide purification, polysaccharide-protein conjugation and stable formulation. The present disclosure further relates to methods for inducing an immune response in subjects against *Salmonella typhi* and non-*typhi* related diseases and/or for reducing or preventing *Salmonella typhi* and non-*typhi* related diseases in subjects using the compositions disclosed herein. The vaccine elicits bactericidal antibodies and is useful for prevention of gastroenteritis, enteric and typhoid fever.

16. [4035676](#) IMPFSTOFFZUSAMMENSETZUNGEN

EP - 03.08.2022

Clasificación Internacional [A61K 39/12](#) N° de solicitud 21386011 Solicitante UNIV OXFORD INNOVATION LTD Inventor/a

The invention describes vaccine compositions containing particles having a polypeptide shell and a water-immiscible core. The polypeptide shell may comprise one or more pathogenic antigen proteins and/or one or more adjuvant polypeptides. Administration of the composition generates an immune response to the polypeptide contained in the shell. Adjuvant may be comprised in the water-immiscible core of the particle. The particles are therefore useful in methods of vaccination.

17. [WO/2022/162398](#) VACCINE COMPOSITIONS

WO - 04.08.2022

Clasificación Internacional [A61K 39/12](#) N° de solicitud PCT/GB2022/050253 Solicitante OXFORD UNIVERSITY INNOVATION LIMITED Inventor/a CARLISLE, Robert

The invention describes vaccine compositions containing particles having a polypeptide shell and a water-immiscible core. The polypeptide shell may comprise one or more pathogenic antigen proteins and/or one or more adjuvant polypeptides. Administration of the composition generates an immune response to the polypeptide contained in the shell. Adjuvant may be comprised in the water-immiscible core of the particle. The particles are therefore useful in methods of vaccination.

18. [20220241421](#) REDUCED FOAMING VACCINE COMPOSITIONS

US - 04.08.2022

Clasificación Internacional [A61K 47/26](#) N° de solicitud 17580896 Solicitante Abic Biological Laboratories Ltd. Inventor/a Noel Yves Henri Jean Genin

The present invention relates to novel stable compressed vaccine composition comprising at least one anhydrous antigenic component comprising a stabilizer susceptible to foaming when the composition is mixed with liquid diluent; and an effective amount of a sugar alcohol.

19. [4034153](#) MODIFIZIERTES VESIKULÄRES STOMATITIS-VIRUS-GLYCOPROTEIN UND VERWENDUNGEN DAVON ZUR BEHANDLUNG VON HIRNTUMOREN

EP - 03.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 20775645 Solicitante UNIV LOUVAIN Inventor/a VANDERMEULEN GAËLLE

The present invention relates to a vaccine for treating and/or preventing a brain tumor. More particularly, the invention relates to a modified vesicular stomatitis virus glycoprotein (VSV-G) comprising at least one tumor antigen, or a fragment thereof, for use in preventing and/or treating a brain tumor in an individual in need thereof, when administered before a surgery intended to remove all or part of the tumor, such as, a tumor resection. The inventors have shown that vaccination of individual with a brain tumor with a vaccine comprising a nucleic acid sequence encoding a modified VSV-G according to the invention may be combined to a tumor resection in order to ameliorate the prognostic of said individuals.

20. [20220241387](#) HORN FLY PROTEIN AS ACTIVE ANTIGEN IN ANTI-HORN FLY VACCINE FOR PROTECTION OF BOVINES AGAINST HORN FLY INFESTATIONS

US - 04.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 17166433 Solicitante The United States of America, as represented by the Secretary of Agriculture Inventor/a FELICITO GUERRERO Antigenic polypeptides derived from a naturally occurring horn fly protein, and nucleic acid molecules encoding the polypeptides, are described. The polypeptides elicit an immune response which, in turn, produces detrimental effects in horn flies feeding on vaccinated cattle. Thus, the present disclosure provides a novel horn fly vaccine.

21. [WO/2022/161495](#) RECOMBINANT SARS-COV-2 VACCINE

WO - 04.08.2022

Clasificación Internacional [C12N 15/50](#) N° de solicitud PCT/CN2022/075134 Solicitante GENESAIL BIOTECH (SHANGHAI) CO., LTD. Inventor/a LIANG, Min

Provided are SARS-CoV-2 vaccines useful for treating or protecting a subject from infection by SARS-CoV-2. The SARS-CoV-2 vaccines contain nucleic acids encoding at least one type of S protein variant, and other factors useful for treating or protecting a subject from infection by SARS-CoV-2, or expression vectors or viruses containing such nucleic acids, or host cells containing such nucleic acids or expression vectors.

22. [WO/2022/162539](#) AN IMMUNOGEN

WO - 04.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud PCT/IB2022/050661 Solicitante UNIVERSITY OF PRETORIA Inventor/a MILLAR, Robert Peter

This invention relates to an immunogen comprising a gonadotropin releasing hormone (GnRH) peptide sequence, a kisspeptin peptide sequence and a stimulant of raising an immune response, such an immunogen for use in a method to regulate the release of hormones in a vertebrate including modulation of reproductive hormones, to reduce fertility in a vertebrate and to treat hormone-dependent diseases including hormone-dependent tumours including prostate tumours, breast, ovary and endometrial tumours, benign hyperplasia including benign prostatic hyperplasia and uterine fibroids, endometriosis, polycystic ovarian disease, infertility, sexual dysfunction and any disorder that would benefit from an increased or decreased GnRH-dependent activity and a vaccine formulation comprising the immunogen. The invention also relates to the use of the immunogen in the preparation of a medicament for use in a method to regulate the release of hormones in a vertebrate.

23. [4034550](#) IMPFTHERAPIE FÜR RAN-PROTEINERKRANKUNGEN

EP - 03.08.2022

Clasificación Internacional [C07K 14/47](#) N° de solicitud 20869039 Solicitante UNIV FLORIDA Inventor/a RANUM LAURA

Aspects of the disclosure relate to compositions and methods for eliciting (or enhancing) anti-repeat-associated non-ATG (RAN) protein antibody expression or production in a subject. Administration of the compositions according to the methods of the present disclosure may in some embodiments result in decreased levels of RAN protein expression and/or aggregation. Such compositions and methods may therefore be useful for the treatment of diseases and disorders known to be associated with RAN proteins.

24. [4035673](#) TRANSDERMALER IMPFSTOFF

EP - 03.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 21386013 Solicitante UNIV OXFORD INNOVATION LTD Inventor/a

The invention describes transdermal vaccines which contain ultrasound responsive particles comprising a polypeptide shell. The surface of the particle has one or more indentations which are generally able to entrap a gas bubble. The particles are capable of generating inertial cavitation on exposure to ultrasound.

The particles can be delivered transdermally, and can comprise antigen protein and/or adjuvant within the particle structure. The particles are therefore useful in methods of vaccination using transdermal delivery routes.

25.[WO/2022/162397](#) TRANSDERMAL VACCINE

WO - 04.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud PCT/GB2022/050252 Solicitante OXFORD UNIVERSITY INNOVATION LIMITED Inventor/a HETTINGA, Johanna

The invention describes transdermal vaccines which contain ultrasound responsive particles comprising a polypeptide shell. The surface of the particle has one or more indentations which are generally able to entrap a gas bubble. The particles are capable of generating inertial cavitation on exposure to ultrasound. The particles can be delivered transdermally, and can comprise antigen protein and/or adjuvant within the particle structure. The particles are therefore useful in methods of vaccination using transdermal delivery routes.

26.[20220241393](#) PORCINE REPRODUCTIVE AND RESPIRATORY SYNDROME VACCINE VIRUS

US - 04.08.2022

Clasificación Internacional [A61K 39/12](#) N° de solicitud 17618625 Solicitante Elanco US, Inc. Inventor/a Stephen Qitu WU

The present invention relates to modified, live Porcine Reproductive and Respiratory Syndrome viruses. Viruses were genetically analyzed and selected based on phylogenetic grouping for modification by repeated passage in tissue culture. The modified, live viruses were assessed for the ability to provide protective immunity to heterologous viruses. The modified, live viruses are useful in vaccines, particularly in vaccines which can treat infection of swine by multiple heterologous viruses.

27.[4035677](#) FALTUNGSPROMOTOREN UND IHRE VERWENDUNG ZUR HERSTELLUNG UND STABILISIERUNG VON POLYPEPTIDEN

EP - 03.08.2022

Clasificación Internacional [A61K 39/215](#) N° de solicitud 21154077 Solicitante MAX PLANCK GESELLSCHAFT Inventor/a

The present invention relates to the recombinant production of a protein of interest in a prokaryotic host cell wherein the protein of interest is obtained in a correctly folded and stable form. The protein of interest may be a difficult-to-make polypeptide for use as a vaccine or a pharmaceutical. The protein of interest is co-expressed with or fused to a 'fold promoter', which may be a VHH antibody recognizing the said protein.

28.[WO/2022/162165](#) FOLD PROMOTERS AND THEIR USE FOR THE PRODUCTION AND STABILIZATION OF POLYPEPTIDES

WO - 04.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud PCT/EP2022/052087 Solicitante MAX-PLANCK-GESELLSCHAFT ZUR FÖRDERUNG DER Inventor/a GÖRLICH, Dirk

The present invention relates to the recombinant production of a protein of interest in a prokaryotic host cell or eukaryotic host cell wherein the protein of interest is obtained in a correctly folded and stable form. The protein of interest may be a difficult-to-make polypeptide for use as a vaccine or a pharmaceutical. The protein of interest is co- expressed with or fused to a 'fold promoter', which may be a VHH antibody recognizing the said protein.

29.[WO/2022/165340](#) UNIVERSAL VACCINES AGAINST IMMUNOGENS OF PATHOGENIC ORGANISMS THAT PROVIDE ORGANISM-SPECIFIC AND CROSS-GROUP PROTECTION

WO - 04.08.2022

Clasificación Internacional [A61K 35/76](#) Nº de solicitud PCT/US2022/014572 Solicitante AEGLE BIOTECH Inventor/a LUBIT, Beverly, W.

The present disclosure provides, in part, a priming and boosting vector-based platform to develop vaccines against viral pathogens that is tailored to elicit a broad T cell response targeting conserved viral epitopes while including helper T cell (TH) epitopes and an adjuvant to achieve a balanced immune response consisting of both cellular immunity, coupled with a broad neutralizing antibody response in the design of a candidate universal vaccine to HIV or a human coronaviruses, e.g., SARS-CoV-2. The universal vaccines are prepared against an immunogen of an infectious pathogenic virus comprising at least one nucleic acid polynucleotide comprising an open reading frame encoding at least one polypeptide antigen or an immunogenic fragment thereof, wherein the polypeptide antigen, or the immunogenic fragment thereof, comprises a conserved internal protein that is enriched in T cell recognition antigens. The effectiveness of the priming and boosting platform is tested in humanized mouse models: a transgenic mouse model that expresses the hACE2 gene under the control of the human cytokeratin 18 promoter and a humanized mouse model comprising a fully functional human immune system.

30. [20220241398](#) LYSSAVIRUS ANTIGEN CONSTRUCTS

US - 04.08.2022

Clasificación Internacional [A61K 39/205](#) Nº de solicitud 17591421 Solicitante GLAXOSMITHKLINE BIOLOGICALS SA Inventor/a KATHRYN HASHEY

Nucleic acid based vaccine constructs encoding Lyssaviral antigens are useful in preventing and treating diseases. Self-amplifying RNA molecules encoding Lyssaviral antigens provide potent and long-lasting immunity.

31. [20220242867](#) NOVEL IMIDAZOPYRIMIDINE COMPOUNDS AND USES THEREOF

US - 04.08.2022

Clasificación Internacional [C07D 487/04](#) Nº de solicitud 16764171 Solicitante Dana-Farber Cancer Institute, Inc. Inventor/a Ofer Levy

The present disclosure provides compounds of Formula (I), and pharmaceutically acceptable salts, solvates, hydrates, polymorphs, co-crystals, tautomers, stereoisomers, isotopically labeled derivatives, prodrugs, and compositions thereof. The compounds described herein are used as enhancers and/or modifiers of an immune response (e.g., innate and/or adaptive immune response), and are useful in treating and/or preventing a disease, as adjuvants in a vaccine for the disease, (e.g., proliferative disease, inflammatory disease, autoimmune disease, infectious disease, or chronic disease), or as stand alone anti-infective or immune response modifying agents. Also provided in the present disclosure are pharmaceutical compositions, kits, methods, and uses including or using a compound described herein.

32. [20220242925](#) NOVEL PEPTIDES AND COMBINATION OF PEPTIDES FOR USE IN IMMUNOTHERAPY AGAINST SMALL CELL LUNG CANCER AND OTHER CANCERS

US - 04.08.2022

Clasificación Internacional [C07K 14/47](#) Nº de solicitud 17716026 Solicitante Immatics Biotechnologies GmbH Inventor/a Andrea MAHR

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

33. [20220241410](#) VACCINE ADJUVANTS BASED ON TLR RECEPTOR LIGANDS

US - 04.08.2022

Clasificación Internacional [A61K 39/39](#) Nº de solicitud 17613414 Solicitante THE UNIVERSITY OF MONTANA Inventor/a Helene Bazin-Lee

Lipidated oxoadenines of formula (I) are TLR7/8 receptor ligands useful for modulating immune responses. The compounds may have therapeutic application in the treatment of cancer, infectious diseases, allergy, or autoimmune disorders.

34. [20220241403](#) Therapeutic Vaccine for Hepatitis B Virus (HBV) using the HBV Core Antigen

US - 04.08.2022

Clasificación Internacional [A61K 39/29](#) Nº de solicitud 17721554 Solicitante UNIVERSITY OF WASHINGTON Inventor/a Edward A. CLARK

Provided herein are compositions of CD1280 binding proteins and a Hepatitis B virus core antigen (HBcAg) and/or a Hepatitis B virus E antigen (HBeAg), or antigenic fragments or mutants thereof, attached to the CD180 binding protein, and methods for using the compositions to treat or limit the development of hepatitis-B virus (HBV)-related disorders.

35. [20220241391](#) METHODS OF BLOCKING ASFV INFECTION THROUGH INTERRUPTION OF CELLULAR AND VIRAL RECEPTOR INTERACTIONS

US - 04.08.2022

Clasificación Internacional [A61K 39/12](#) Nº de solicitud 17535545 Solicitante Dalu CHEN Inventor/a Dalu CHEN

A method of preventing and treating viral infections in animals (and preferably ASFV in porcine), by inhibiting viral ligand interactions with critical cellular receptors that are involved either directly (endocytosis and/or macropinocytosis) or indirectly (phagocytosis of RBCs that have been aggregated by viral interactions) with cellular entry in an animal, and preventing and treating the viral infection in the animal. A method of treating a viral infection in an individual with a virus that is both lysogenic and lytic. A composition for treating a viral infection in an individual with a virus that is both lysogenic and lytic. A vaccine for preventing viral infection, including whole and/or partial domains of proteins of both a lysogenic and lytic phase of a virus.

36. [4034548](#) CORONAVIRUSIMPFSTOFFE UND VERWENDUNGEN DAVON

EP - 03.08.2022

Clasificación Internacional [C07K 14/165](#) Nº de solicitud 20923712 Solicitante GUANGZHOU ARGORNA BIOPHARMACEUTICALS CO LTD Inventor/a ZHANG BILL BILIANG

This disclosure relates to coronavirus vaccines and uses thereof. In one aspect, the disclosure provides a nucleic acid vaccine, comprising a sequence encoding a spike protein or fragment thereof derived from a coronavirus.

37. [4034253](#) ZUSAMMENSETZUNGEN UND VERFAHREN ZUR ERHÖHUNG DER WIRKSAMKEIT VON IMMUNTHERAPIEN UND IMPFSTOFFEN

EP - 03.08.2022

Clasificación Internacional [A61P 35/00](#) Nº de solicitud 20869832 Solicitante UNIV MICHIGAN REGENTS Inventor/a MOON JAMES J

This invention relates generally to compositions and methods for increasing the efficacy of immunotherapies and vaccines. In particular, the present invention relates to elevating the richness and diversity of a subject's gut microbiome through administration of an agent (e.g., fiber containing prebiotic agent (e.g., epigallocatechin gallate (EGCG), fucoidan, potato starch, oligofructose and inulin)) (e.g., melatonin) with an immunotherapy or vaccine. Such compositions and methods are useful for treating cancer, infectious pathogens, autoimmune diseases, neurological disorders, and/or obesity.

38. [4034159](#) NEOGLYCOKONJUGATE ALS IMPFSTOFFE UND THERAPEUTISCHE WERKZEUGE

EP - 03.08.2022

Clasificación Internacional [A61K 39/385](#) N° de solicitud 20867004 Solicitante KORANEX CAPITAL
Inventor/a SHIAO TZE CHIEH

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

39.[4035675](#)NEUARTIGE PEPTIDE UND KOMBINATION AUS PEPTIDEN ZUR VERWENDUNG IN DER IMMUNTHERAPIE GEGEN OVARIALKARZINOM UND ANDERE KARZINOME

EP - 03.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 22151693 Solicitante IMMATICS BIOTECHNOLOGIES GMBH Inventor/a SCHUSTER HEIKO

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

40.[20220241401](#)VACCINES AGAINST GENITAL HERPES SIMPLEX INFECTIONS

US - 04.08.2022

Clasificación Internacional [A61K 39/245](#) N° de solicitud 17550272 Solicitante Board of Supervisors of Louisiana State University and Agricultural and Mechanical College Inventor/a Konstantin G. Kousoulas The present invention provides vaccines for treating or preventing a herpes simplex virus infection and methods of using and making the vaccine. Further provided are recombinant herpes simplex virus genomes, recombinant viruses, and immunogenic compositions.

41.[2920140](#)Anticuerpos neutralizantes del virus de la inmunodeficiencia humana (VIH)

ES - 01.08.2022

Clasificación Internacional [A61K 39/00](#) N° de solicitud 19156884 Solicitante Theraclone Sciences, Inc. Inventor/a Chan-Hui, Po-ying

42.[2022205209](#)Novel peptides and combination of peptides for use in immunotherapy against esophageal cancer and other cancers

AU - 04.08.2022

Clasificación Internacional [C07K 14/47](#) N° de solicitud 2022205209 Solicitante Immatics Biotechnologies GmbH Inventor/a Fritsche, Jens

The present invention relates to peptides, proteins, nucleic acids and cells for use in immunotherapeutic methods. In particular, the present invention relates to the immunotherapy of cancer. The present invention furthermore relates to tumor associated T-cell peptide epitopes, alone or in combination with other tumor associated peptides that can for example serve as active pharmaceutical ingredients of

vaccine compositions that stimulate anti-tumor immune responses, or to stimulate T cells ex vivo and transfer into patients. Peptides bound to molecules of the major histocompatibility complex (MHC), or peptides as such, can also be targets of antibodies, soluble T-cell receptors, and other binding molecules.

43.[WO/2022/160017](#)SARS-COV-2 VACCINE ANTIGENS

WO - 04.08.2022

Clasificación Internacional [A61K 39/215](#) N° de solicitud PCT/AU2022/050050 Solicitante GRIFFITH UNIVERSITY Inventor/a GOOD, Michael

The present disclosure provides immunogenic compositions and methods of inducing an immune response and/or preventing, treating or ameliorating an infection, disease or condition associated with SARS-CoV-2 in a subject with such immunogenic compositions.

44.[WO/2022/162312](#)PEPTIDES AND USES THEREOF

WO - 04.08.2022

Clasificación Internacional [A61K 8/64](#) N° de solicitud PCT/FR2022/050145 Solicitante SCHWEIGHOFFER, Fabien Inventor/a SCHWEIGHOFFER, Fabien

The present application relates to compounds and their use in health or cosmetics. The application more particularly relates to peptides that can be used in vaccine strategy as well as for the cosmetic or therapeutic treatment of any tissue or organ. The invention covers peptides as well as compositions containing same, the preparation and uses thereof, in particular in humans.

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

Results Search in US Patent Collection db for: (ABST/vaccine AND ISD/20220801->20220811), 21 records.

PAT. NO.	Title
1	11,407,829 LAG3 binding peptides
2	11,407,819 Compositions and use of a fibrinogen binding motif present in EFB and COA for therapeutics and vaccines against <i>Staphylococcus aureus</i>
3	11,407,810 Peptides and combination of peptides for use in immunotherapy against various tumors
4	11,407,809 Peptides and combination of peptides for use in immunotherapy against various tumors
5	11,407,808 Peptides and combination of peptides for use in immunotherapy against various tumors
6	11,407,807 Peptides and combination of peptides for use in immunotherapy against various tumors
7	11,407,798 Peptides and combination of peptides for use in immunotherapy against lung cancer, including NSCLC, SCLC and other cancers
8	11,406,706 Lipid nanoparticle vaccine adjuvants and antigen delivery systems
9	11,406,701 Vaccination of immunocompromised subjects

10	11,406,698	Vaccine compositions
11	11,406,697	Methods and compositions for treating glioma and medulloblastoma brain tumors using the zika virus
12	11,406,691	AMH-INH-GNIH tri-expression gene vaccine of improving fecundity of animals, preparation method and application
13	11,401,319	Peptides and combination of peptides for use in immunotherapy against esophageal cancer and other cancers
14	11,401,310	Peptides and combination of peptides for use in immunotherapy against ovarian cancer and other cancers
15	11,401,267	Substituted benzyl-triazole compounds for Cbl-b inhibition, and further uses thereof
16	11,400,170	Methods for treating cancers
17	11,400,151	Methods for improving immunological response in vaccinated animals
18	11,400,149	Ebola virus and Marburg virus glycoprotein mucin-like domain replacement expression system used as new vaccine approaches
19	11,400,147	Pneumococcal capsular saccharide conjugate vaccine
20	11,400,108	Cyclic dinucleotides as agonists of stimulator of interferon gene dependent signalling
21	11,400,094	2H-pyrazolo[4,3-d]pyrimidine compounds as toll-like receptor 7 (TLR7) agonists and methods and uses thereof

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