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Medical Innovations



Patent and Technology Transfer Enabler













Biolarvicide for Mosquito control





Domain: Insecticide

Name of Institute: ICMR-Vector Control Research Centre (VCRC)

Name of the inventor(s): Dr. Daniel Reegan

Brief about Technology: Highly effective against larvae of Culex, Anopheles, and Aedes mosquitoes, while ensuring exceptional safety for aquatic non-target species, beneficial insects, and mammals

Intended Use: Public Health Biolarvicide

USP: Eco-friendly cost-effective green product for use in mosquito control operations

Status:

- Manufacturing License received
- Approval received from Central Insecticide Board, Insecticide Registration Committee

Technology Transfer Status: Transferred to companies on Non-Exclusive basis

IP status: Indian Patent No. 222246 and 192055







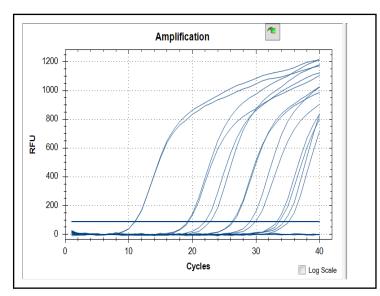






Nipah point of care assay





Nipah point of care assay

Domain: Diagnostic Assay/Kit

Name of Institute: ICMR-National Institute of Virology

Name of Inventor(s): Dr. Pragya Yadav

Brief about Technology: The Nipah real-time probe-based assay is an RT-qPCR method that detects Nipah virus RNA with high sensitivity and specificity using fluorescent hydrolysis probes targeting viral N gene. It provides rapid results within a few hours, enabling early diagnosis during outbreaks.

Intended Use: Enables rapid Nipah diagnosis, supporting timely patient care and containment of virus spread

USP: Rapid, highly sensitive and specific detection of Nipah viral RNA directly from clinical or field samples within 4 hours

Status: Manufacturing license received, Emergency use approval by CDSCO

Technology Transfer Status: Transferred to companies on Non-Exclusive basis









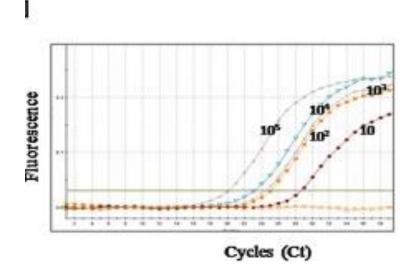




Kyasanur Forest Disease (KFD) Point of Care test







Domain: Diagnostic Assay/Kit

Name of Institute: ICMR-National Institute of Virology

Name of Inventor(s): Dr. Pragya Yadav, Dr AM Shete

Brief about Technology: The Kyasanur Forest Disease (KFD) real-time RT-PCR assay is a probe-based molecular diagnostic tool used for rapid and specific detection of KFD virus RNA. It works by amplification using target-specific primers and fluorescent hydrolysis probes, directed at the NS5 gene region. **Intended Use:** Facilitate rapid KFD diagnosis, enabling timely patient care and effective containment.

USP: Rapid, highly sensitive, and specific detection of KFD virus RNA directly from clinical or field samples within 4 hours

Status: Assay validation completed

Technology Transfer Status: Transferred to companies on Non-Exclusive basis













Single-tube multiplex RT-PCR for simultaneous detection of SARS-CoV-2, Influenza A, and B

Advanced Stage
Technology



Domain: Diagnostic Assay/Kit

Name of ICMR Institute: ICMR-National Institute of Virology (NIV)

Name of the Inventor: Dr. Varsha Potdar, Mrs. Veena Vipat and Mrs Sheetal Jadhav

Brief about Technology: TaqMan based Real Time PCR, Single tube multi virus detection assay (Four plex) Intended Use: Accurate detection of SARS CoV 2 and its variants (Till date XFG), along with influenza A subtypes H1N1/pdm09 and H3N2, as well as influenza B lineages Yamagata and Victoria

USP: Extensively used nationwide under the pan-India ILI/SARI VRDL network since 2021. All 72 VRDLs are currently utilizing the kit

Current status: Manufacturing License pending, Approved by CDSCO

Technology Transfer Status: Transferred to companies on Non-Exclusive basis

IP status: Filed (Indian Patent Application Number-202211052927)













Multiplex RT assay for simultaneous detection of Influenza A, Influenza B, RSV A/B, and SARS-CoV-2

Advanced Stage
Technology



Domain: Diagnostic Assay/Kit

Name of ICMR Institute: ICMR-National Institute of Virology (NIV)

Name of the Inventor: Dr Varsha Potdar, Mrs. Veena Vipat and Mrs. Sheetal Jadhav

Brief about Technology: TaqMan based Real Time PCR , Single tube multi virus detection assay (five plex)

Intended Use: Simultaneous detection of Influenza A,B,RSV and SARS CoV 2 in single assay

USP: Unlike commercial two-tube assays, the NIV assay offers single-tube, high-confidence detection of all five viruses—delivering accurate, cost-effective testing

Current Status: Test License received/Pilot Study/Pivotal Study

Technology Transfer Status: Transferred to companies on Non-Exclusive basis













Japanese Encephalitis (JE) Antigen Capture ELISA





Domain: Diagnostic Assay/Kit

Name of ICMR Institute: ICMR-National Institute of Virology (NIV)

Name of the Inventor: Dr. GN Sapkal, Dr. MM Gore, Dr. DT Mourya & Dr. PS Sathe

Brief about Technology: India's first indigenous JE antigen detection kit—cost-effective, sensitive, rapid, and user-friendly. Safe for BSL-2 labs and ideal for mosquito-based JEV surveillance to pinpoint high-risk regions

Intended Use: The JE Antigen Capture ELISA is a serological assay enables qualitative detection of JE antigen in mosquitoes, supporting identification of high-risk outbreak zones

USP: No JE antigen capture ELISA kit is commercially available for the detection of the virus.

Current status: CDSO Manufacturing License received

Mode of Transfer (Exclusive/Non-exclusive): Transferred to a company on Non-Exclusive basis













Human Chandipura (CHPV) IgM ELISA





Domain: Diagnostic Assay/Kit

Name of ICMR Institute: ICMR-National Institute of Virology (NIV)

Name of the Inventor: Dr. GN Sapkal & Dr. VP Bondre

Brief about Technology: CHPV causes high fatality in children; early detection via IgM antibodies is vital. The BSL-2 safe kit supports surveillance to identify high-risk regions.

Intended Use: CHPV IgM Capture ELISA is intended for qualitative determination of IgM antibodies in serum / CSF of patients presenting clinical signs and symptoms consistent with CHPV encephalitis. The serological assay is designed for providing laboratory diagnosis of CHPV at sentinel hospitals and primary health centers

USP: First indigenous Human CHPV IgM ELISA kit which is cost effective, sensitive, rapid and user friendly.

Current status: Manufacturing License received

Mode of Transfer (Exclusive/Non-exclusive): Transferred to a company on Non-Exclusive basis



















Domain: Diagnostic Assay/Kit

Name of ICMR Institute: ICMR-National Institute of Virology (NIV)

Name of the Inventor: Dr. GN Sapkal, S.R.Vaidya, D.T. Mourya

Brief about Technology: This ELISA uses the inactivated Measles antigen & will provide a qualitative determination of Measles specific IgM in human serum. Which can be used for early diagnosis of measles.

Intended Use: The Anti-Measles IgM ELISA kit qualitatively detects Measles IgM antibodies in serum, aiding presumptive diagnosis during outbreaks and in clinical settings

USP: Indigenous kit, can be used for early diagnosis of Measles virus. The test is very user friendly ELISA kit which is cost effective, sensitive, rapid & user friendly.

Current status: Manufacturing License Received

Mode of Transfer (Exclusive/Non-exclusive): Transferred to a company on Non-Exclusive basis







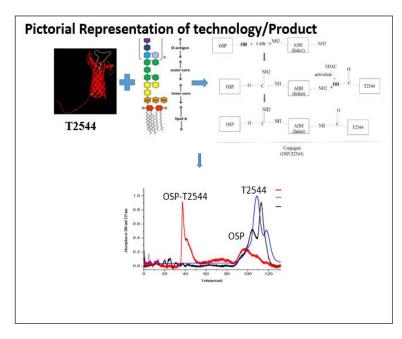






Glycoconjugate vaccine composition against Salmonella spp.





Domain: Vaccine

Name of Institute: ICMR NIRBI

Name of the inventor(s): Dr. Santasabuj Das, Dr. Risha Haldar, Dr. Amlanjyoti Dhar, Dr. Suparna Chakraborty

Brief about Technology: A glycoconjugate vaccine (OSP-T2544) comprising of O-specific polysaccharide (OSP) from *Salmonella* Typhimurium and the conserved outer membrane protein (T2544) of *Salmonella* Typhi/ Paratyphi A that induces humoral and cellular immunity and protects against S. Typhi, S. Paratyphi A, S. Typhimurium and S. Enteritidis infections in mice

Intended Use: Candidate vaccine against Typhoidal and non-Typhoidal Salmonella

USP: A multivalent conjugate vaccine which induces strong seroconversion, secretory intestinal IgA response, and lasting immunological memory, offering protection against key *Salmonella* strains – S. Typhi, S. Paratyphi A, S. Typhimurium and S. Enteritidis in mouse models.

Status: Proof of concept established through *in vitro* and *in vivo* experiments, functional prototype developed, technology transferred to industry

Technology Transfer Status: Transferred to a company on non-exclusive basis

IP Status: Filed (Indian Patent application number – 20231107021)













Rapid LAMP Detection Assay for Mpox Virus







Reaction	F3L-4	B6R-4	β-Actin	Observation	Inference
NTC	u,	4	4	All the rubes showing -Ve results	Test is valid
Sample DNA A		ur)		F3L-4: +Ve B6R-4: +Ve β-Actin: +Ve	Positive for Orthopoxvirus and Monkeypox
Sample DNA B	2			F3L-4: -Ve B6R-4:+Ve β-Actin:+Ve	Positive for Orthopoxvirus and Negative for Monkeypox
Sample DNA C		4		F3L-4: -Ve B6R-4:-Ve β-Actin:+Ve	Negative for Orthopoxvirus and Monkeypox
Sample DNA D	T T			F3L-4: +Ve B6R-4:-Ve β-Actin:+Ve	Inconclusive Repeat the test

Domain: Diagnostics (Assay/Kits)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Shyam Sundar Nandi (PI), Sonali Sawant, Yadav Pragya, Shete-Aich Anita,

Jagadish Deshpande

Brief about Technology: This assay can be used for diagnosis of Monkeypox, where quantitative detection is not necessary. The assay specifically detects two genes of Monkeypox virus (genus specific B6R gene and species specific F3L gene)

Intended Use: This assay can be used for diagnosis of Monkeypox, where quantitative detection is not necessary. The assay specifically detects two genes of Monkeypox virus (genus specific B6R gene and species specific F3L gene)

USP: Indigenous, simple and cost-effective Point of care technology can be used in the fields for detection of *Monkeypox* virus

Status: Pivotal Study and Test License received

Technology Transfer Status: Transferred to companies on Non-Exclusive basis

IP Status: Filed (Indian Patent application number – 202211057074)

For Licensing Opportunity, Contact: patentmitra.hq@icmr.gov.in







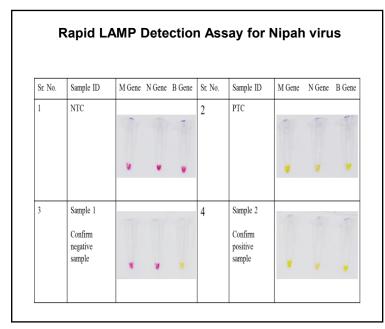






Rapid LAMP Detection Assay for Nipah virus





Domain: Diagnostics (Assay/Kits)

Name of Institute: ICMR-National Institute of Virology, Mumbai Unit

Name of the inventor(s): Shyam Sundar Nandi (PI), Sonali Sawant, Yadav Pragya, Shete-Aich Anita,

Jagadish Deshpande

Brief about Technology: Indigenously developed loop mediated isothermal amplification (LAMP) assay specifically detects two genes of Nipah virus (N and M gene). The results are interpreted visually by observing the color change. Sophisticated reading instrument not needed

Intended Use: The assay will be most preferred for diagnosis of Nipah virus infection where quantitative detection is not necessary

USP: Indigenous, simple and cost-effective Point of care technology can be used in the fields for detection of Nipah virus

Status: External validation at NIV, Pune

IP Status: Filed (Indian Patent application number – 202211066352)



















Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Pragya Yadav, Dr. AM Shete & Dr. DT Mourya

Brief about Technology: The assay is based on capturing the IgM antibodies on a microtiter plate using anti-human-IgM antibody. Unbound material will be removed by washing. Formation of anti-IgM antibodies complex would be detected using KFD antigen, anti KFD biotinylated IgG and Avidin-Horseradish peroxidise (AV-HRP) system.

Intended Use: For qualitative detection of anti KFD IgM antibodies in the serum of patients with suspected febrile illness & haemorrhages.

USP: Currently there is no diagnostic ELISA kit available for detection of anti-KFD IgM antibodies

Status: Test License received/ Pilot Study/Pivotal Study

Technology Transfer Status: Transferred to companies on Non-Exclusive basis













Anti-CCHF Human IgM ELISA





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Pragya Yadav, Dr. AM Shete and Dr. DT Mourya

Brief about Technology: This is the First indigenous anti-CCHF human IgM antibody detection kit which is cost effective, sensitive, rapid and user friendly. Kit uses gamma inactivated antigen, safe to handle in BSL-2 level laboratory for screening of human serum samples.

Intended Use:The kit is intended for qualitative detection of anti CCHF IgM antibodies in human serum samples. It can be used at any level of clinical set up, public health centres

USP:The kit uses gamma inactivated antigen, safe to handle in BSL-2 level of laboratory for screening of human serum samples. The kit is cost effective, sensitive, rapid and user friendly

Status: Test License received, Batch validation for CDSCO approval

Technology Transfer Status: Transferred to companies on Non-Exclusive basis







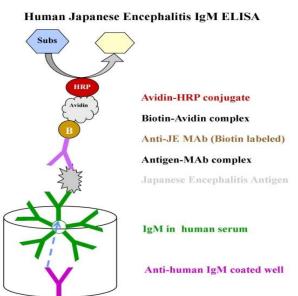






Diagnostic ELISA for detection of IgM antibodies against Japanese Encephalitis virus





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Kanchankumar Patil & Dr. Deepti Parashar

Brief about Technology: IgM antibodies in the patient's serum/ CSF (if present) and IgM from Positive Control are captured by anti-human IgM (μ chain specific) coated on to the solid surface (wells). In the next step, JE antigen (inactivated JE virus) is added which binds to captured human JE specific IgM Intended Use: Japanese Encephalitis (JE) IgM capture ELISA developed by ICMR-NIV is intended for qualitative determination of JE virus specific IgM antibodies in serum/ CSF of patients presenting clinical signs and symptoms consistent with JE.

USP: Indigenous diagnostic test which is being currently used in National Vector Borne Disease control program and cost effective as compared to currently available commercial tests.









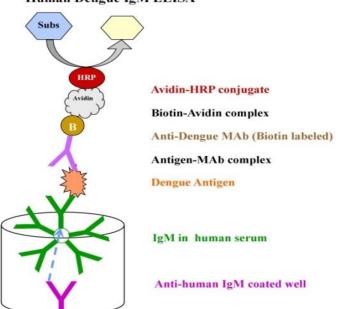




Diagnostic ELISA for detection of IgM antibodies against Dengue virus

Mid stage Technology

Human Dengue IgM ELISA



Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Kanchankumar Patil & Dr. Deepti Parashar

Brief about Technology: IgM antibodies in the patient's serum (if present) and IgM from Positive Control are captured by anti-human IgM (μ chain specific) coated on to the solid surface (wells). In the next step, DEN antigen (inactivated dengue virus) is added which binds to captured human dengue specific IgM. **Intended Use:** Dengue IgM capture ELISA developed by ICMR-NIV is intended for qualitative determination of Dengue virus specific IgM antibodies in serum of patients presenting clinical signs and symptoms consistent with Dengue.

USP: Indigenous diagnostic test which is being currently used in National Vector Borne Disease control program and cost effective as compared to currently available commercial tests

Status: Internal validation done









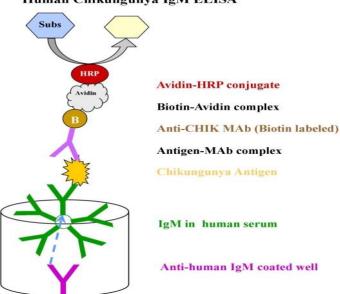




Diagnostic ELISA for detection of IgM antibodies against Chikungunya virus

Mid stage Technology





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Kanchankumar Patil & Dr. Deepti Parashar

Brief about Technology: IgM antibodies in the patient's serum (if present) and IgM from Positive Control are captured by anti-human IgM (μ chain specific) coated on to the solid surface (wells). In the next step, CHIK antigen (inactivated chikungunya virus) is added which binds to captured human chikungunya specific IgM

Intended Use: CHIK IgM capture ELISA developed by ICMR-NIV is intended for qualitative determination of chikungunya virus specific IgM antibodies in serum of patients presenting clinical signs and symptoms consistent with chikungunya

USP: Indigenous diagnostic test which is currently being used in National Vector Borne Disease control program and cost-effective as compared to currently available commercial tests.

Status: Third party validation done













HEV IgM Rapid Test







Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Tejaswini Mahesh Deshmukh, Ms. Manisha Tukaram Dudhmal, Dr. Kavita Satish Lole

Brief about Technology: HEV IgM Rapid Test is an IgM capture solid-phase lateral-flow immunochromatographic assay. All IgM class Abs present in test sample will be captured by anti-human IgM Abs immobilized at Test Line (TL). Presence of HEV specific IgM Abs can be differentially detected by colloidal gold labeled anti-HEV MAbs complexed with recombinant HEV protein immobilized within the device & can be visualized as pink/purple lines after completion of the assay.

Intended Use: The test enables effective hepatitis E diagnosis in low-resource, field, and point-of-care settings, including clinics, mobile units, and emergency or minimal-infrastructure labs.

USP: HEV IgM Rapid Test has been developed in IgM capture format & sensitively & specifically detects HEV infections. Its performance is comparable to laboratory based ELISAs (commercial & in-house). The test requires minimum 10µl serum/plasma sample & gives results within 15 min

Status: Third party validation done













Shigella vaccine





Domain: Vaccine

Name of Institute: ICMR-National Institute for Research in Bacterial Infections (NIRBI)

Name of the inventor(s): Dr. Hemanta Koley Dr Soma Mitra, Dr Santa Sabuj Das, Dr Mk Chakrabarti

Brief about Technology: The vaccine is developed to reduce the burden of disease and mortality caused by enteric pathogen like Shigella. This vaccine has been designed for protection against Six Shigella strains (S.sonnei, S.dysenteriae i, S. Boydii, and S. flexneri 2a, 3a, and 6)

Intended Use: For prevention of Shigella infections

USP: The vaccine is based Hexavalent strategy that can collectively provide the necessary broad spectrum protection needed to achieve a vaccine of global utility

Status: Third party validation complete

IP Status: Indian patent - IN 502420







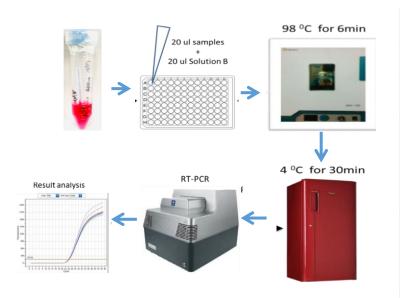






Method for preparation of sample for nucleic acid extraction free PCR





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute of Cancer Prevention and Research (NICPR)

Name of the inventor(s): Dr R Suresh Kumar, Dr Yashawardhan, Mr Ravi Shankar, Mr Babar Ali, Dr Shalini

Brief about Technology: The developed solution along with specific methodological processes like incubation at particular temperature and time period can help performing

PCR from sample directly

Intended Use: Real time PCR with out Nucleic acid extraction

USP: This is a rapid, extraction-free real-time PCR method using simple chemicals and incubation, ideal for liquid biopsies, with minimal time, consumables, and suitability for resource-limited settings

Status: Inhouse validation complete

IP Status: Filed (Indian Patent application number –202211041619)









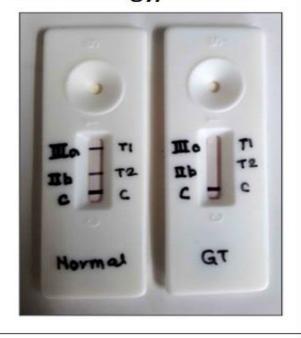




Rapid diagnostic POC for Glanzmann's thrombasthenia (GT)



Pictorial Representation of technology/Product



Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR - National Institute of Immunohematology (NIIH)

Name of the inventor(s): Dr. Shrimati Shetty & Dr. Tejasvita Gaikwad

Brief about Technology: For the first time, a point of care (POC) test has been successfully developed for detecting Glanzmann's thrombasthenia. This novel, rapid and simple test can be widely used for diagnosing GT cases without using a sophisticated flow cytometer or a platelet aggregometer. It is a lateral flow test strip.

Intended Use: Detecting Glanzmann's thrombasthenia

USP: The kit is a a novel, specific, and sensitive strip-based point-of-care test for platelet function defects, offering rapid results within 20 minutes and enabling diagnosis of Glanzmann's Thrombasthenia (GT) even at PHC level, with a working cost of ₹120

Status: Internal and External validation completed

IP Status: Filed (Indian Patent application number -202211003648)









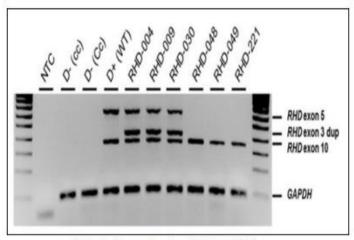




Indian RhD genotyping assay



Pictorial Representation



Gel photograph of multiplex PCR

Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR - National Institute of Immunohematology (NIIH)

Name of the inventor(s): Dr Swati Kulkarni, NIIH, Dr Yann Fichou, Establishment Français Du Sang (EFS)

Inserm, Brest (France)

Brief about Technology: The multiplex PCR assay determining RHD genotype and detecting most common weak D phenotype in Indian population. The method comprising detecting the presence of RHD exon 5 &10 along with duplication of exon 3 of the RHD gene from DNA sample. The presence of a duplication of exon 3 of the RHD gene in the DNA sample is indicative of a weak D type 150 phenotype. **Intended Use:** For identifying most common weak D variant in Indian population and confirming the RhD status of apparently RhD negative & D variants individuals

USP: This a Simple Multiplex PCR assay for RhD genotyping. This assay will help in identifying the most common weak D mutation in Indians i.e. Weak D type 150 (60-85%).

IP Status: Filed (Indian Patent application number –201917040421)









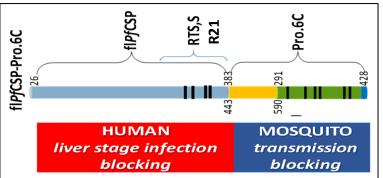


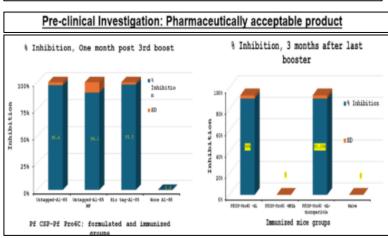


Mid stage

Chimeric recombinant multi-stage vaccine for preventing *Plasmodium* falciparum infection and its community transmission **Technology**

(AdFalciVac)





Domain: Vaccine

Name of Institute: ICMR-RMRCBB

Name of the inventor(s): Dr. Susheel K Singh, Dr. Subhash Singh, Dr. Agam P Singh, Dr. Sanghamitra Pati

Brief about Technology: A recombinant multistage vaccine (AdFalciVac) against P. falciparum, produced in Lactococcus lactis and a pharmaceutically acceptable carrier for protection against infection in humans, as well as reducing the community transmission

Intended Use: The vaccine has been developed to offer both protection against *Plasmodium falciparum* infection in humans and interruption of community transmission, unlike the existing two vaccines that can partially prevent infection in humans but cannot stop transmission in the community.

USP: Rational targeting of two most vulnerable stages of parasite (i.e. pre-erythrocytic stage and Sexual stage of malaria), Stable & functional chimera between antigens, Protection against both infection in humans and controlling vector-borne community transmission.

Status: Third party validation ongoing

IP Status: Filed (Indian Patent Application - 202411095679)







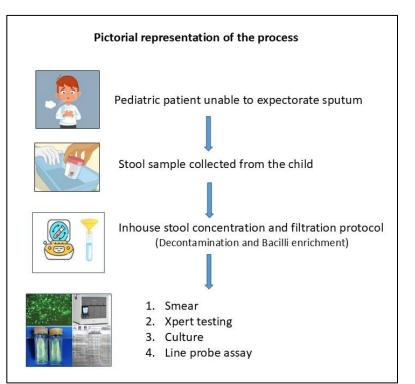






Stool NAAT for Pediatric TB detection





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR - National Institute for Research in Tuberculosis

Name of the inventor(s): Dr. Priya Rajendran

Brief overview of Technology: The technology involves a novel in-house stool concentration protocol optimized for pediatric samples, enabling the detection of *Mycobacterium tuberculosis* (MTB) and drug resistance markers through nucleic acid amplification tests (NAAT), Line Probe Assay (LPA), and Xpert Ultra, as well as phenotypic tests such as smear microscopy and culture methods.

Intended Use: Detection of MTB and drug resistance in pediatric stool samples

USP: Non-invasive method of MTB detection and its drug resistance in pediatric stool samples using a standardized concentration method for molecular diagnostics.

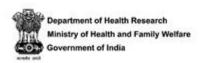
Status: Pilot study completed and large-scale study ongoing













CRISPR-Cas based TB detection system

Mid stage Technology

Pictorial Representation of technology/ Product



RapidBact: DNA Extraction kit

(Patent Application No. 202511026919)



Glow-TB_{PCR}: PCR-CRISPR-Cas based detection kit

(Patent Application No. 202411043686)



RapidGlow: Incubation-cumfluorescence detection device (Patent Application No. 202411016083)



RGLOW: Data acquisition and result analysis software (Copyright applied)

Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR -Regional Medical Research Centre, Dibrugarh (RMRCNE)

Name of the inventor(s): Dr. Md. Atique Ahmed

Brief about Technology: The system uses heat lysis and column-based DNA extraction (RapidBact), followed by multiplex PCR and CRISPR-Cas12a detection (Glow-TBPCR). Cas12a, guided by TB-specific gRNAs, triggers fluorescence upon target detection. The RapidGlow device enables incubation and real-time signal detection, supported by RGLOW software for automated analysis

Intended Use: Diagnosis of tuberculosis

USP: Low-cost, rapid DNA extraction from $60\,\mu l$ sputum; multiplex detection of 3 TB genes; ultrasensitive (LOD ~1 copy/ μl); portable, high-throughput, and cost-effective alternative to centralized diagnostics.

Technology Transfer Status: Transferred to a company on Non-Exclusive basis

IP Status: Filed (Indian Patent application number –202411016083, 202411043686, 202511026919)







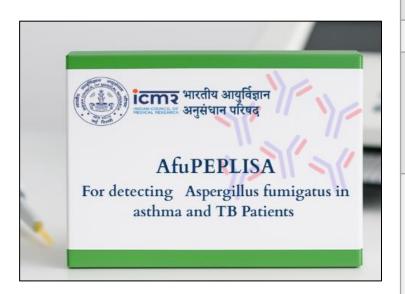






Immunodiagnostic kit for detecting Aspergillus fumigatus in asthma and TB Patients (AFuPEPLISA)





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute for Research in Reproductive & Child Health (NIRRCH)

Name of the inventor(s): : Dr Taruna Madan Gupta

Brief about Technology: The AFuPEPLISA Immunodiagnostic Kit presents a strong market potential in the early detection of Aspergillus fumigatus among patients with bronchial asthma and pulmonary tuberculosis—conditions that are increasingly prevalent in developing regions.

Intended Use: Detection of Aspergillus fumigatus in the sera of patients with bronchial asthma and pulmonary tuberculosis

USP: Aspergillus fumigatus complicates bronchial asthma and pulmonary tuberculosis, with elevated IgE and IgG levels aiding in serodiagnosis. Early detection is crucial to prevent bronchiectasis and fibrosis. Current diagnostic tests like ImmunoCAP, Immulite 2000, and Platellia Aspergillus IgG ELISA show variable sensitivity due to differences in antigen quality, assay matrices, and immobilization methods, leading to inconsistent results across platforms

Status: GMP finalized, third party validation complete

IP status: Filed (Indian patent application- 202411008706)







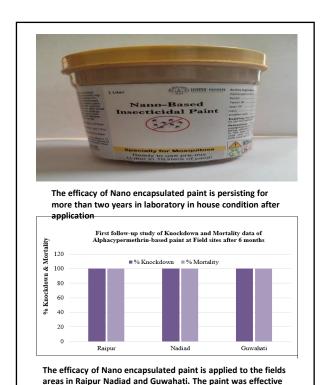






Nano-emulsion based insecticidal paint





and 100% mortality of mosquitoes observed after 6 months

study continued

Domain: Insecticide

Name of Institute: ICMR-National Institute of Malaria Research (NIMR)

Name of the inventor(s): Dr Himmat Singh

Brief about Technology: Slow releasing insecticidal paint using Nanoencapsulation technique

Intended Use: Insecticide

USP: This is targeted application, has long-lasting protection, reduced environmental impact, ease of application, wide area application and cost effective

Status: In house validation complete

IP Status: Filed (Indian patent application- 202411054306)

For Licensing Opportunity, Contact: patentmitra.hq@icmr.gov.in













Diagnostic Assay kit for common enteric viruses





Domain: Diagnostic (Assay/Kit)

Name of Institute: ICMR-National Institute for Research in Bacterial Infections (NIRBI)

Name of the inventor(s): Dr Mamta Chawla-Sarkar, Dr. Santasabuj Das

Brief about Technology: A multiplex, 2-tube Real-time qRT-PCR assay (Taqman-based assay)

Intended Use: The kit allows multiple enteric viruses causing childhood gastroenteritis by Real Time PCR

USP: Single kit for detection of multiple enteric viruses

Status: Third party validation complete

Technology Transfer Status: Transferred to a company on non-exclusive basis













Single tube multiplex method for simultaneous detection of Influenza A,B,C and D viruses

Early stage Technology



Domain: Diagnostics (Assay/kits)

Name of Institute: ICMR-National Institute of Virology (NIV)

Name of the inventor(s):Dr Varsha Potdar, Mrds Veena Vipat and Mrs Sheetal Jadhav

Brief about Technology: TaqMan based Real Time PCR, Single tube influenza multi virus detection assay (Four plex)

Intended Use: Accurate detection of diverse Influenza types which may have zoonotic potential in single tube assay .

Intended Use: Simultaneous detection of Influenza A, B, C, and D to identify zoonotic infections and track animal-to-human spillover event

USP: This will be unique product useful for One health approach

Status: In-house validation complete

Technology Transfer Status: Not Transferred







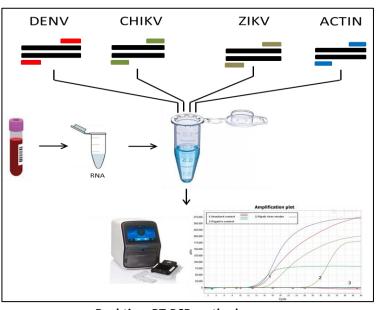






Single tube multiplex real-time RT-PCR method for simultaneous detection of Dengue, Chikungunya and Zika viruses

Early stage Technology



Real-time RT-PCR method

Domain: Diagnostic kits/Assay

Name of Institute: ICMR-National Institute of Virology

Name of the inventor(s): Dr. Alagarasu Kalichamy, Dr. Deepti Parashar, Mr. Mahadeo Kakade, Dr. Gajanan Sapkal, Dr. Anukumar, Dr. Ashok M

Brief about Technology: Customized primers and probes with actin control to ensure RNA integrity and precise amplification. Optimized multiplex setup enables rapid, sensitive detection of co-circulating arboviruses with minimal reagent usage

Intended Use: Simultaneous Detection of DENV, CHIKV, and ZIKV in a Single-Tube Multiplex RT-PCR Assay in clinical samples

USP: Multiplex assay with a stably expressing endogenous internal control in a single reaction

Status: In-house validation complete with sensitivity of 96.1% for dengue, 97.2% for chikungunya and specificity of 100% for all three viruses. Concurrent results in QCMD panel for Zika







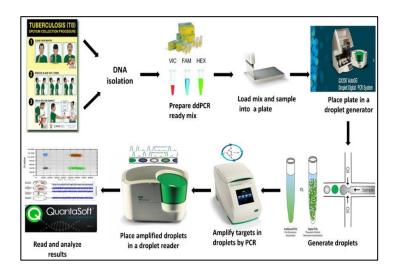






Detection of *Mycobacterium* tuberculosis-derived circulating cell-free DNA (ccfDNA) in plasma for early TB diagnosis

Early stage
Technology



Domain: Diagnostic kits/Assay

Name of Institute: ICMR-National Institute for Research in Tuberculosis (NIRT)

Name of the inventor(s): Dr. Luke Elizabeth Hanna, Mr. Manohar Nesakumar, Ms. Evangeline Ann Daniel

Brief about Technology: Optimized dual target droplet digital PCR assay

Intended Use: Early diagnosis of Extrapulmonary TB, Asymptomatic/subclinical TB, Symptomatic clinically diagnosed/possible TB, Individuals at high risk of developing TB

USP: This ultrasensitive, highly specific plasma-based test detects tuberculosis across its full spectrum, even in diagnostically challenging cases.

Status: In-house validation complete







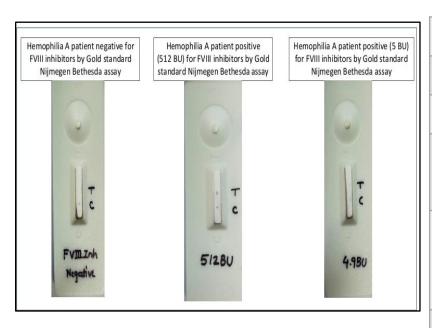






Factor VIII inhibitor Point of care test

Early stage Technology



Factor VIII inhibitor Point of care test kit

Domain: Diagnostics (Assay/kits)

Name of Institute: ICMR- National Institute of Immunohaematology (NIIH)

Name of the inventor(s): Dr. Rucha Patil, Sharda Shanbhag, Sweta Kaskar, Moni Singh

Brief about Technology: The technology utilizes gold nanoparticles conjugated with Factor VIII antigen to

specifically detect FVIII inhibitors in plasma

Intended Use: To detect FVIII inhibitors in hemophilia A patients

USP: It is a first-of-its-kind LFIA uses gold nanoparticles to rapidly detect FVIII inhibitors in plasma within 20 minutes, requiring minimal equipment and no technical expertise. Cost-effective (~₹100), user-friendly, and compatible with Emicizumab therapy, it outperforms conventional assays in speed, accessibility, and affordability

Status: Completed and internal validation performed with >90% sensitivity and specificity for > 5BU (high titer inhibitors)

IP status: Filed (Indian Patent application number- 202511083527)







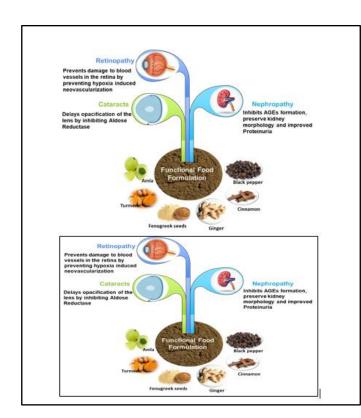






Prophylactic composition for diabetes-associated complications

Early stage
Technology



Domain: Functional Food

Name of Institute: ICMR-National Institute of Nutrition (NIN)

Name of the inventor(s): GB Reddy, SS Reddy, KK Kalahasti, S Pandarinath, M Nagaraju

Brief about Technology: The invention relates to two functional food compositions aimed at preventing diabetes/hyperglycemia-associated complications: cataract, retinopathy, and nephropathy.

Intended Use: A molecular target-based functional food composition for ameliorating diabetes/ hyperglycemia associated complications

USP: This product has demonstrated preventive and therapeutic efficacy against diabetes and its complications in preclinical studies.

Status: In-house validation complete







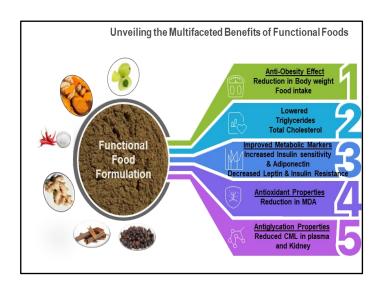






Anti-obesogenic functional food and method of preparation thereof

Early stage
Technology



Domain: Functional food

Name of Institute: ICMR-National Institute of Nutrition (NIN)

Name of the inventor(s): GB Reddy, SS Reddy, KK Kalahasti, S Pandarinath

Brief about Technology: The anti-obesogenic functional food due to its anti-inflammatory, antiglycation, AR inhibition, antioxidant, and immunomodulatory properties with no side effects provides a sustainable solution to managing obesity and improving the outcomes of obesity-related complications. It is cost-effective, contains easily available and regularly consumed dietary agents

Intended Use: A molecular target-based functional food composition for the management of obesity and associated complications

USP: This anti-obesogenic functional food is made of commonly consumed and affordable dietary ingredients, offers a safe, multi-targeted approach to sustainably manage obesity and its related complications through anti-inflammatory, anti-glycation, AR inhibitory, anti- oxidant, and immunomodulatory effects

Status: Preclinical studies complete

IP Status: Filed (Indian patent Application - 202411045411)













Light-responsive in situ hydrogel for intraocular drug delivery

Early stage
Technology



Domain: Therapeutics

Name of Institute: Department of Pharmacy, Birla Institute of Technology and Sciences

(BITS), Hyderabad Campus

Name of the inventor(s):Prof. Nirmal Jayabalan, Sai Shreya Cheruvu

Brief about Technology: This technology involves a UV-triggered intravitreal gel depot designed for sustained drug release, minimizing injection frequency in chorioretinal conditions and providing a promising alternative for patients unresponsive to anti-VEGF therapy

Intended Use: Spironolactone-loaded, long-acting, light-responsive in situ hydrogel: a novel therapy beyond anti-VEGF for treating diabetic retinopathy.

USP: The technology enables up to 90 days of sustained drug release with enhanced biocompatibility, reducing injection frequency and improving safety and patient adherence.

Status: Preclinical studies completed













Natamycin loaded peptide conjugated nanomicelles (PeNatcel)

Early stage Technology



Domain: Therapeutics

Name of Institute: Department of Pharmacy, Birla Institute of Technology and Sciences (BITS), Hyderabad Campus; Shantilal Shanghvi Cornea Institute, KAR Campus, LV Prasad Eye Institute, Hyderabad

Name of the inventor(s): Prof. Nirmal Jayabalan, Priyadarshini Sathe, Dr. Prashant Garg

Brief about Technology: This innovation encapsulates natamycin, a poorly water-soluble antifungal, into peptide-transporter-targeted nanomicelles. The design significantly enhances the solubility, corneal permeation, and bioavailability, thus delivering superior therapeutic outcomes

Intended Use: Topical eye drop for treating fungal keratitis that achieves bioavailability with lower drug doses and fewer instillations, leading to higher patient compliance and faster recovery

USP: With a reduced dose (0.6%) and dosing frequency (4×/day), the formulation achieves 13× higher transcorneal permeation and 2× longer pre-corneal residence, enhancing patient compliance

Status: Preclinical studies completed













Your vaccine development journey starts here – Pathogen isolates available for material transfer

Name of the Pathogen	Targeted Disease		
SARS-Cov2	COVID		
Influenza A virus (H1N1 and H3N2)	Influenza		
Influenza B virus	Influenza		
Measles virus (Morbillivirus)	Measles		
Mumps virus	Mumps		
Nipah virus	Nipah virus infection		
Rubella virus	Rubella		
Zika virus	Zika virus infection		
Crimean-Congo haemorrhagic fever (CCHF) virus	Crimean-Congo haemorrhagic fever		
Chandipura Virus	Acute encephalitis syndrome (AES)		















Patents Available For Commercialization from the ICMR portfolio

ICMR has portfolio of **55 publications** available for licensing, which includes **45 patents** and **10 pending applications**. These are organized across **five key technology areas**:

Therapeutics: We have 16 patents/patent applications in this category. This is our largest area of focus, reflecting our commitment to advanced treatment solutions.

Diagnostics: We have 12 patents/patent applications in this category, showcasing our tools for precision healthcare.

Devices: We have 6 patents/patent applications in this category for targeted opportunities in medical devices.

Vaccines: We have 3 patents/patent applications in this category which are focused on immunization.

Miscellaneous: We have 18 patents/patent applications in this category showcasing a diverse range of versatile technologies.

Scan for detailed patent portfolio:

• To explore these technologies, pathogen isolates, and our patent portfolio available for licensing, please visit our website <u>patentmitra.icmr.org.in</u> or scan the QR code below

