

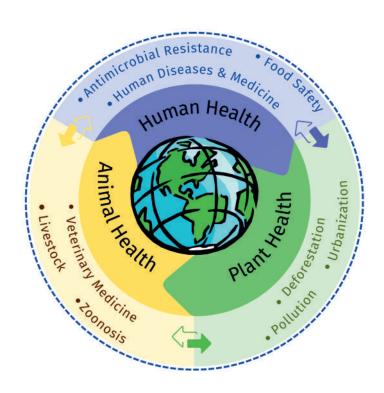




AMR and One Health

AMR continues to severely affect global health and the economy. It represents one of the leading public health threats of the $21^{\rm st}$ century, which is exacerbated by the overuse and misuse of antibiotics worldwide. The unabated spread of AMR could also result in several bacterial pathogens becoming more lethal. It is responsible for causing 700,000 deaths annually. It is predicted that more than 10 million people will die per year due to AMR infections by 2050.

The SARS-CoV-2 pandemic has further intensified the AMR crisis due to the atrocious use of antibiotics in treating COVID-19 patients. There is also increasing recognition of the relationships between human and animal health, food safety, and environmental sectors in both the evolution and solution to the AMR problem. It is therefore essential to take the One Health approach to effectively address the AMR issue.





Our Mission

"Access to Essential Vaccines and Novel Biologics Should Not Only be the Privilege of Just the Affording Few"

Health equity for all





Our Vision

Enabling One Health through Bio-innovation

AffordableAccessible

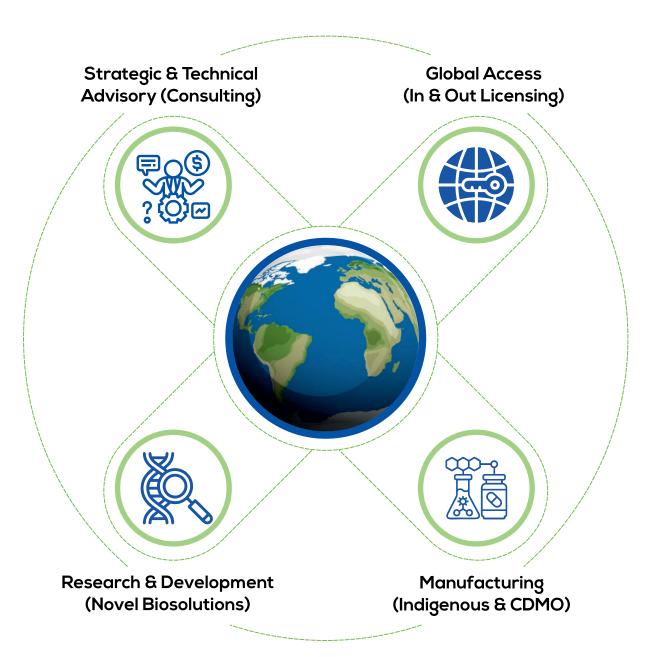
Acceptable





TechInvention's Approach

Our Multipronged Approach to One Health

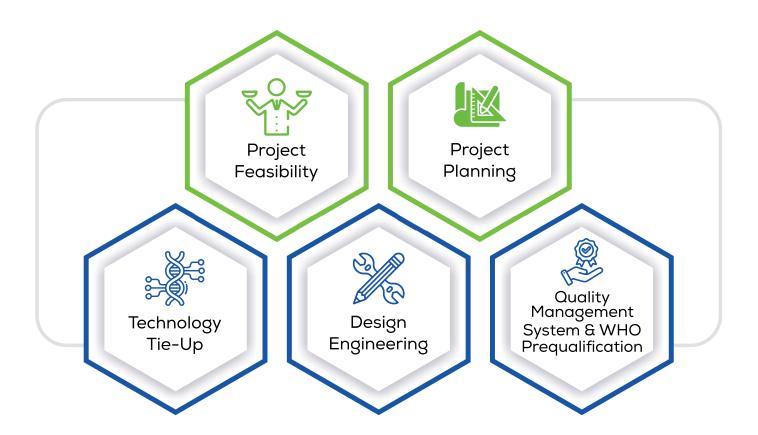






We offer our expertise and support in capacity building to enable local manufacturing of biologics and vaccines in LMICs.

Our key areas are:

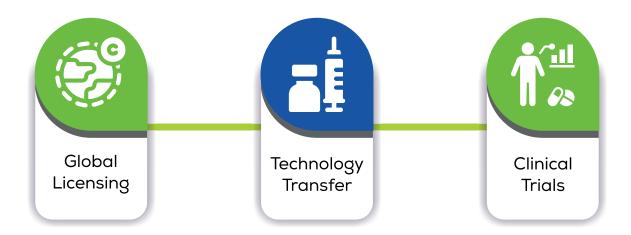




2 Global Access (In & Out Licensing)

Through our network of partners in 20+ countries, we can help in providing compliant and affordable technology and also support you with local clinical trials, regulatory licensure, IP services (patent drafting, patent filings, freedom to operate analysis, non-infringement opinion, prior art analysis, patentability analysis, and prosecution of patent application in India and other countries), and supplies of vaccines and novel biologics.

Our key areas are:





We facilitate in-licensing and out-licensing partnerships that bring together companies with complementary competencies to strengthen pipelines, achieve economies of scale, advance technologies, and introduce commercial products to market.





Clinical Trial Pipeline- Vaccine Registration in India

Vaccine candidate	Study title
Cholera	A phase III, open label, multicentre, parallel group, randomised clinical study to compare the immunogenicity and safety of Euvichol-Plus® with Shanchol™ in healthy adults and children above the age of one year.
Varicella	A prospective, randomised, single blind, parallel, active controlled, multicentre, non-inferiority phase III study to evaluate the immunogenicity and safety of live attenuated varicella vaccine of Sinovac Biotech Ltd., China compared to Variped® (MSD Pharmaceuticals Pvt. Ltd., India) in healthy paediatric subjects in India.
Tetanus, Diphtheria, and Pertussis	A multicentre, randomised, observer-blind, non-inferiority, phase III study to evaluate the immunogenicity and safety of Boostagen RED™ by BioNet-Asia compared to Adacel® by Sanofi Pasteur Ltd. in healthy subjects aged 4-65 years.
Influenza	A randomised, multicentre, phase III, comparative clinical trial to evaluate the safety and immunogenicity of quadrivalent influenza vaccine manufactured by SPbSRIVS and Influvac® Tetra of Abbott in healthy adult human subjects.

Vaccine candidate	Clinical trial related document preparation and submission to CDSCO	Approval of clinical trial by SEC at CDSCO	Clinical trial NOC and conduct of clinical trial	Clinical study report submission to CDSCO	Approval of clinical study report	Marketing authorization
Cholera						
Varicella						
Tetanus, Diphtheria, and Pertussis						
Influenza						



3 Manufacturing (Our Indigenous Pipeline & CDMO)

To better serve the biopharmaceutical global market and enable scale-up of our indigenous R&D products, Virchow Biotech Pvt. Ltd. and TechInvention Lifecare Pvt. Ltd. formed a joint venture VirchowTech, dedicated to the scale-up and commercial production for vaccines and biologics. The business generates synergy by leveraging optimal use of the extensive resources to enable capacity building and affordability.

VirchowTech is home to an ISO 13485:2012 and WHO-GMP-certified state-of-the-art manufacturing facility with a highly innovative and motivated research team and a manufacturing staff who will not settle for anything less than the highest quality. The manufacturing set up encompasses production units, testing laboratories, research and development facilities, warehouse, and packaging divisions.



1SO - International Organization for Standardization, WHO - World Health Organization, GMP - Good manufacturing practice







4 Research & Development Pipeline (Novel Biosolutions)

TechInvention's indigenous portfolio aims to tackle infectious diseases and the prevailing global concern of antimicrobial resistance, thereby contributing to the ultimate goal of One Health. Our multipronged approach encompasses diagnostics, vaccines, and novel biotherapeutics such as monoclonal antibodies and lysins.

Vaccines

Recombinant Platform Technology					
Target pathogen	R & D	Proof-of-concept studies	Preclinical studies		
SARS-CoV-2					
Human Papillomavirus (VLP)					
MERS-CoV					
Meningococcal serogroup B					

Conjugate Platform Technology						
Target pathogen	R & D	Proof-of-concept studies	Preclinical studies			
Streptococcus pneumoniae						
Meningogoccal serogroup ACYW						



Biotherapeutics

Monoclonal Antibodies - Phage Display Technology					
Target pathogen	R & D	Proof-of-concept studies	Preclinical studies		
SARS-CoV-2					
Chikungunya virus					
Cytomegalovirus					
Dengue virus					

Phage-encoded lysins - Recombinant Technology					
Target pathogen	R & D	Proof-of-concept studies	Preclinical studies		
Uropathogenic E. coli					
A. baumannii					
P. aeruginosa					





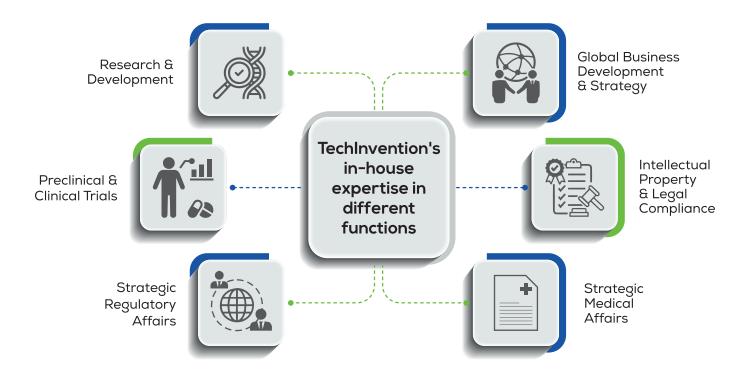
Diagnostics

Extraction kits	R & D	Validation	Manufacturing	Commercialization
DNA				
RNA				
Plasmid				
Gel				
Rapid detection kits	R & D	Validation	Manufacturing	Commercialization
T. gondii				
Human Papillomavirus				

Molecular diagnostic kits	R & D	Validation	Manufacturing	Commercialization
MERS-CoV				
Varicella				
Influenza				
SARS-CoV-2 + MERS-CoV + Influenza				
SARS-CoV-2 + MERS-CoV				
Influenza + SARS-CoV-2				
Chikungunya virus				
Dengue virus				
SARS-CoV-2				
Malaria				
Tropical fever panel				



Expertise



Culture

TechInvention has been certified for two consecutive years (2021 & 2022) as Flace Great Place To Work®, which is the gold standard and unrivaled worldwide Work when it comes to employer brand recognition.







Why Work with TechInvention?



India has highest number of WHO PQ vaccines and manufacturers.

TechInvention has arrangements with most of them including WHO PQ manufacturers from other developing countries

concept to commerciallidation

iensulling time bound flawless et al.



TechInvention's facility designs are highly compliant and optimizes capital and operational costs making products affordable



Hand holding all through the project



Strong non-infringing intellectual property



Access to recombinant platform technologies for targeting WHO critical category of priority pathogens



Access to other recombinant and conjugate vaccines and technologies



Deployment & training of technical manpower



Access to biotherapeutics such as monoclonal antibodies and lysins for infectious diseases

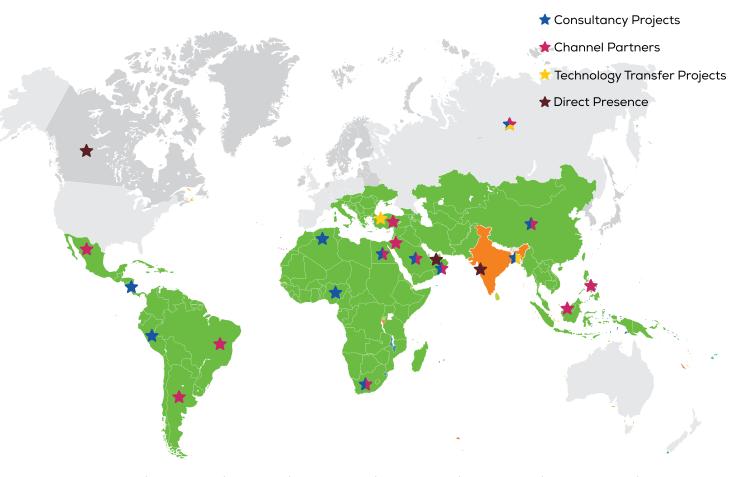


Experienced team with prowess and proficiency in commercial scale manufacturing of diagnostics, vaccines, & biotherapeutics





Global Projects and Partners



















































































Our Stellar 1st

- 1. Enabled setting up the 1^{st} state-of-the-art vaccine manufacturing facility in Central America from concept-to-commercialization
- 2. Set-up of the $1^{\rm st}$ R & D centre & technology transfer of rHBsAg vaccine to Hacetteppe University, Ankara, Turkey
- 3. Secured 1st long term supply contract of a human vaccine NUPCO (Saudi Arabia) from a BRICS country
- 4. Developing 1st novel paediatric combination vaccine (Hexa) with a cross border partnership from 2 BRICS countries
- 5. Enabled 1st technology transfer of Hib conjugate vaccine into a CIS country along with facility design
- 6. Bringing in to India, the world's first and only oral cholera vaccine in LDPE unidose
- 7. Enabled manufacturing of India's $1^{\rm st}$ Hepatitis A vaccine at Biological E. Ltd., Hyderabad
- 8. Bringing the world's 1^{st} recombinant pertussis vaccine to India
- 9. Certain novel serotypes being worked upon for the $1^{\rm st}$ time in the world for pneumococcal conjugate vaccine & human papillomavirus vaccine



Highlights

- Strategic portfolio encompassing diagnostics, vaccines, and biotherapeutics for dreaded infectious diseases
 - 2 Platform technologies that confers affordability, safety, and efficacy
 - A certified 'Deep Tech Pioneer' by the Hello Tomorrow organization of Paris
 - 20+ Institutional Biosafety Committee (IBSC) approvals for ongoing initiatives
 - 5 15+ Research abstracts acceptance in global conferences of repute
 - 6 15+ Patents filed
 - 7 2 Publications in high impact factor journals of global repute
 - 8 BIRAC's BIG Scheme-19 awardee
- 9 2021 SME Excellence awardee in healthcare by SME Chamber of India
- Active contribution to the G20 presidency goal of One Health with its entire R&D portfolio based around the One Health approach to acquire the UN's Sustainable Development Goal 3





Memberships















































Notes



Notes



The health of soil, plant, and man is one and indivisible.

- Albert Howard



- www.techinvention.biz
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