Development of a Target Product Profile (TTP) – Part 1

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Vaccines, from concept to implementation (GLHL 7209) – January 19, 2021



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Target Product Profiles (TPPs)

"The TPP is essentially an organized "wish-list" of characteristics, features, and attributes that one would like to see in a newly developed medical product once it reaches the market."

Lee BY and Burke DS. Constructing target product profiles (TPPs) to help vaccines overcome post-approval obstacles. Vaccine. 2010 Apr 1; 28(16): 2806–2809.



Preferred Product Characteristics (PPCs) and Target Product Profiles (TPPs)

- WHO's Initiative for Vaccine Research (IVR) develops
 PPCs and TTPs to guide vaccine developers.
- PPCs provide strategic guidance as to WHO's preferences for new vaccines in priority disease areas and are typically developed prior to phase III clinical trial planning.
- High priority vaccines include: Dengue, HPV, Malaria, Measles-Rubella, Pneumococcal conjugate, Polio (inactivated), Rabies, Rotavirus, Typhoid, Yellow fever.



Preferred Product Characteristics (PPCs) and Target Product Profiles (TPPs)

- TPPs provide technical guidance as to WHO's criteria in response to emergency or epidemic scenarios.
- Developed to facilitate to accelerated vaccine development.
- WHO TPPs are usually formulated early in product development, prior to phase I clinical testing.
- Both PPCs and TPPs articulate vaccine preferences from an LMIC perspective.



WHO PPCs and TPPs

- Developed with key stakeholders and end-users.
- Primary audience is any entity engaged in global vaccine product development.
- Compliance can facilitate WHO policy recommendation and prequalification.
- Can reduce the time between vaccine licensure and introduction into countries.



A Target Product Profiles

- Has the ultimate goal in mind.
- Identifies "optimal target" and a "minimal target."
- Early TPPs may propose estimated targets that can be refined based on manufacturing, nonclinical & clinical results.
- Typically, targets do not change unless the intended use of the product or standard of care changes, or in response to regulatory guidance.
- The final version of the TPP often serves as the precursor to the annotated product label.



TPP Elements

- 1. Indication for use
- 2. Target population
- 3. Contraindication
- 4. Safety/Reactogenicity
- 5. Efficacy
- 6. Dose regimen
- 7. Durability of protection
- 8. Route of administration
- 9. Coverage

- 10. Product stability and storage
- 11. Co-administration with other vaccines
- 12. Presentation
- 13. Production
- 14. Registration and Prequalification
- 15. Post-marketing surveillance



Indication for use

What do you want the vaccine to do ideally? What would be the minimum benefit that would be acceptable?

Preferred	Critical or Minimal
Immunization protects against	Immunization reduces the
COVID-19 infection.	severity of COVID-19 disease.



Target population

Age, gender, health profile, e.g. pregnant, lactating women

Preferred	Critical or Minimal
All ages and medical profiles.	Adults including the elderly.



Contraindications

What populations should be excluded for safety reasons, children, pregnant women, people w/ allergic conditions?

Preferred	Critical or Minimal
None.	Immune compromised patients.



Safety/Reactogenicity

Adverse events (AEs)

Preferred	Critical or Minimal
Mild, transient AEs	Safety and reactogenicity profile whereby
No serious AEs.	vaccine benefits outweigh safety risks.

Describe two or more safety risks that could result from a COVID-19 vaccine that you think would seriously limit the vaccine's use by the general public.



TPP - Element 5 Efficacy

To what extent does your vaccine work across different outcomes, e.g., preventing death, reducing hospital stay, reducing viral shedding?

Preferred	Critical or Minimal
> 70% efficacy (on population basis, including the elderly).	> 70% efficacy (population basis) and >50% efficacy in the elderly.



Dose regimen

Single/multiple dose, time between doses, booster requirements

Preferred	Critical or Minimal
Single-dose regimen.	No more than two dose regimen one month apart.



Durability of protection

Lifetime immunity, time-limited immunity, need for regular boosters

Preferred	Critical or Minimal
Lifetime.	Confers protection for at least 6 months.



Route of administration

injectable, nasal spray, oral delivery

Preferred	Critical or Minimal
Oral.	Injectable.



CoverageMonovalent, multivalent

Preferred	Critical or Minimal
Multivalent.	Monovalent.

Your vaccine protects against a single variant or multiple variants.



Product stability and storage

Shelf life, temperature requirements, preservatives

Preferred	Critical or Minimal
Shelf life > 24 months at -20 °C.	Shelf life > 12 months at -20 °C.
Stability > 6 months at 2-8°C.	Stability >8 hours at 2-8°C.



Product stability and storage

"There are few immunization issues more important than the appropriate storage and handling of vaccines."



https://blogs.unicef.org/blog/getting-childrenimmunized-going-the-extra-mile-in-car/

14 page chapter from WHO/PAHO

https://www.paho.org/immunization/toolkit/resources/partner-pubs/ebook/Chapter5-Vaccine-Storage-and-Handling.pdf?ua=1

Vaccine Vial Monitor

Symbol	Explanation	Stage
□ ✓	The inner square is lighter than the outer circle. If the expiry date has not passed, USE the vaccine.	I
	As time passes the inner square is still lighter than the outer circle. If the expiry date has not passed, USE the vaccine.	II
Discard point: the color of the inner square matches that of the outer circle. DO NOT USE the vaccine.		III
×	Beyond the discard point: inner square is darker than the outer circle. DO NOT USE the vaccine.	IV

From WHO (www.who.int).

https://s3.amazonaws.com/gpeitk/reference links/en/Pakist Guide Book for AICs.pdf

Which is the most heat sensitive vaccine?

- 1. Hepatitis B
- 2. Rabies
- 3. Oral Polio
- 4. Measles/Mumps/Rubella



Co-administration with other vaccines

Interference with COVID-19 vaccine or the other vaccines

Preferred	Critical or Minimal
Can be given with other vaccines	Must be used as a stand
without affecting immunogenicity,	alone vaccine.
safety or efficacy of the vaccines.	



Presentation

Liquid/lyophilized, mono-dose/multi-dose, dosage volume

Preferred	Critical or Minimal
Liquid product in mono-dose or	Liquid or lyophilized product,
multidose (10-20) presentations.	mono or multidose, diluent
Maximal dosage volume 0.5mL.	provided. Dosage 0.5mL.



Production

Number of doses of vaccine to be produced annually.

Preferred	Critical or Minimal
9 billion doses	350 million doses



Registration and Prequalification

Type of license- e.g. is it experimental

WHO	US FDA
WHO prequalified and/or meets	Emergency Use Authorization,
criteria for EUAL (Emergency Use	Expanded Access Program.
Assessment & Listing Procedure.	

EUA requires detailed information on efficacy, short term safety and manufacturing processes.



Post-marketing Surveillance

Monitoring safety, effectiveness, herd immunity, potential emergency of vaccine resistant mutants

WHO and National Regulatory Authorities

Post-marketing surveillance will include assessment of serious adverse effects, effectiveness and emergence of vaccine resistant SARS-CoV-2 mutants in accordance with national regulatory authorities and the WHO prequalification requirements.

Questions?



Go to Part 2



WHO PPCs and TPPs

PPCs and TPPs shape vaccine product development

Although not formal WHO guidelines, PPC and TPP documents have oversight from WHO's Product Development for Vaccines Advisory Committee, which in turn informs WHO's Strategic Advisory Group of Experts on Immunization (SAGE) on the status of vaccine development. However, a policy recommendation for vaccine use will ultimately depend on a multitude of factors in addition to product specific attributes, including affordability, cost-effectiveness and the programmatic considerations such as feasibility of implementation and anticipated coverage. Broader perspectives such as equity and community acceptability are also important for health policy.

https://www.who.int/immunization/research/ppc-tpp/en/