

NUVA

Unified Nomenclature of Vaccines

Concept of valence

Definitions

VACCINE

ANTIGEN

Vaccine

A vaccine is an antigenic preparation, either similar to or derived from a pathogen, or composed of genetic material encoding the target antigen. It is administered to an individual to elicit a specific and active immune response that protects against the natural disease caused by the corresponding pathogen.

Antigen

An **antigen** (“antibody generator”) is any substance or molecule that is recognized by the immune system as foreign or potentially harmful, triggering an immune response. When an antigen enters the body, it stimulates the production of specific antibodies or activates immune cells to neutralize or eliminate the threat.

The concept of valence



Immunisation Schedule 2025, France

Vaccins contre :	Naissance	2 mois	3 mois	4 mois	5 mois	6 mois	11 mois	12 mois
Diphtérie (D), Tétanos (T), coqueluche acellulaire (Ca), Poliomyélite (P)		DTCaP		DTCaP			DTCaP	
<i>Haemophilus influenzae b</i> (Hib)		Hib		Hib			Hib	
Hépatite B (Hep B)		Hep B		Hep B			Hep B	
Pneumocoque (PnC) ¹		PnC		PnC			PnC	
Rotavirus		Rota	Rota	+/- Rota ²				
Méningocoque B ³			MnB		MnB			MnB
Méningocoque ACWY (vaccin conjugué)						MnACWY		MnACWY
Rougeole (R), Oreillons (O), Rubéole (R) ⁴								ROR 1

DTCaP = DTaP-IPV
PnC = PCV
ROR = MMR

MnACWY = MenACWY-CRM or MenACWY-TT
MnB = MenB-4C or MenB-FHbp

Immunisation Schedule, USA

COVID-19 vaccine	1vCOV-mRNA	Comirnaty/Pfizer-BioNTech COVID-19 Vaccine
		Spikevax/Moderna COVID-19 Vaccine
	1vCOV-aPS	Novavax COVID-19 Vaccine
Dengue vaccine	DEN4CYD	Dengvaxia
Diphtheria, tetanus, and acellular pertussis vaccine	DTaP	Daptacel Infanrix
<i>Haemophilus influenzae</i> type b vaccine	Hib (PRP-T) Hib (PRP-OMP)	ActHIB Hiberix PedvaxHIB
Hepatitis A vaccine	HepA	Havrix Vaqta

Vaccine scheduler, ECDC (Poland)

	Birth	Months								Years					
		2	3	4	5	6	7	13-15	16-18	5	6	10	12-13	14	18
Coronavirus disease (COVID-19)										COVID-19					
tuberculosis	BCG ³														
rotavirus infection		ROTA													
diphtheria		D	D	D					D		D			d	
tetanus		TT	TT	TT					TT		TT			TT	
pertussis		wcP ⁴	wcP ⁴	wcP ⁴					wcP ⁴		acP			acp	
poliomyelitis		IPV	IPV	IPV					IPV		IPV				
Haemophilus influenzae type b infection		Hib	Hib	Hib					Hib						

Vaccine records

B. C. G.

Date	Lieu	Méthode	Signature et cachet du médecin
10 NOV 1972	Delft	Scarf. 88	
29/3/82	OCC Monseigneur	51-jack +	Docteur F.X. FCHCAULT MÉDECINE GÉNÉRALE 2, rue H. Bonnet, 2 89530 CHAMPDOST T. 845.21.35
7/4/82	Booshe + lili		

VACCINATION ANTIVARIOLIQUE

Date	Lot du vaccin	Résultat	Signature et cachet du médecin
17 MAI 1973	JoSol	+	
14/7/83	64321	+	Docteur F.X. FCHCAULT MÉDECINE GÉNÉRALE 2, rue H. Bonnet, 2 89530 CHAMPDOST T. 845.21.35

VACCINATIONS - D.T. - COQ - POLIO

Date	Vaccin - Dose	Signature et cachet du médecin
26 JANV 1973	DT Coq + Polio	
28.8.73	DT Coq + Polio	
26 AVRIL 1973	DT Coq adjuv + Polio	
19 MAI 1974	DT Coq ad + Polio	
14.5.74	DT Coq ad + Polio	
12/7/76	Polio Sabin	
27/11/79	DT Vax + Polio Sabin	
7.8.83	DT Polio	
5/9/89	DT Polio	
15/2/94	DT Polio	
30/1/96	DT Polio	
9/9/98	DT Polio	

External Codes

30010001	30010146
30010015	30010155
30010027	30010169
30010042	30010174
30010058	30010188
30010073	30010203
30010084	30010217
30010096	30010226
30010107	30010239

From Raw Codes to Immunisation Decisions via Valence Interpretation



External Codes

30010001	30010146
30010015	30010155
30010027	30010169
30010042	30010174
30010058	30010188
30010073	30010203
30010084	30010217
30010096	30010226
30010107	30010239



Valence-based
interpretation engine



- Vaccine status
- Decision

Valence concept in NUVA: an Operational Objective

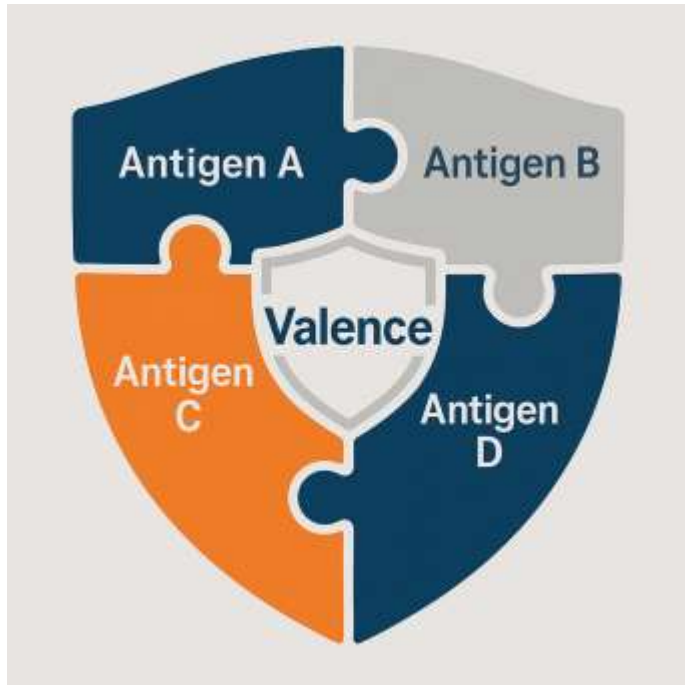
- The valence is the smallest functional unit of a vaccine, knowledge of which is necessary and sufficient to assess an individual's immunisation status against a specific infectious agent (or a subspecies thereof) and to plan the next vaccine dose, if appropriate.
- A valence represents an antigen, or a relevant group of antigens, that is specific to a given target infectious agent or subspecies (such as serotypes or shared proteins).

The concept of valence in NUVA was primarily introduced to facilitate and standardise the interpretation of vaccine histories by a vaccine decision support system, regardless of how the recorded vaccines are designated.

Valence in NUVA – Example : BEXSERO

- This vaccine contains 4 antigens
 - Recombinant *Neisseria meningitidis* group B NHBA fusion protein
 - Recombinant *Neisseria meningitidis* group B NadA
 - Recombinant *Neisseria meningitidis* group B fHbp fusion
 - Outer membrane vesicles (OMV) from *Neisseria meningitidis* group B
- This vaccine has one valence in the NUVA representation
 - 4CMenB: Four-component Meningococcal Group B Vaccine

Why does the valence not coincide with the antigens in this case?



Because precise knowledge of the four different antigens in this vaccine does not impact the evaluation of the immune protection of the person who received it, nor does it affect the determination of the next dose to be administered..

Valence label

401 | EJ-I-Vero-3

Japanese encephalitis valence, Inactivated, Whole-virion, Derived from Vero cells, 3 µg

379 | COVID-mRNA-S-OMI-JN.1-30

COVID-19 valence, Omicron variant JN.1, mRNA based, Encoding SARS-CoV-2 spike protein, 30 µg

- ✓ Name of the disease or target microorganism
- ✓ More info on the target
- ✓ Vaccine type 1
- ✓ Vaccine type 2
- ✓ Vaccine type 3
- ✓ Dosage

The valence concept allows:

- To identify equivalent vaccines
- To interpret combined vaccines and calculate dose rank
- To take into account the level of information available
- To make decisions, either manually or with the help of a decision support system

Decomposition of Vaccines into Valences

Vaccines received:

01/02/2022 : BOOSTRIX TETRA

01/04/2022 : REVAXIS



Disease	Valence	No.
Pertussis	ap	1
Diphtheria	d	2
Tetanus	T	2
Polio	IPV	2

Antigen comparison

VAXELIS

- Diphtheria Toxoid: ≥ 20 IU
- Tetanus Toxoid: ≥ 40 IU
- Bordetella pertussis antigens:
 - Pertussis Toxoid (PT): 20 μ g
 - Filamentous Haemagglutinin (FHA): 20 μ g
 - Pertactin (PRN): 3 micrograms
 - Fimbriae Types 2 and 3 (FIM): 5 μ g
- Hepatitis B surface antigen: 10 μ g
- Poliovirus (Inactivated)
 - Type 1 (Mahoney): 40 D antigen units
 - Type 2 (MEF-1): 8 D antigen units
 - Type 3 (Saukett): 32 D antigen units
- Haemophilus influenzae type b polysaccharide:
 - Polyribosylribitol Phosphate: 3 μ g
 - Conjugated to meningococcal protein: 50 μ g

HEXYON

- Diphtheria Toxoid: ≥ 30 Lf
- Tetanus Toxoid: ≥ 10 Lf
- Bordetella pertussis antigens:
 - Pertussis Toxoid (PT): 25 μ g
 - Filamentous Haemagglutinin (FHA): 25 μ g
- Hepatitis B surface antigen: 10 μ g
- Poliovirus (Inactivated)
 - Type 1 (Mahoney): 29 D antigen units
 - Type 2 (MEF-1): 7 D antigen units
 - Type 3 (Saukett): 26 D antigen units
- Haemophilus influenzae type b polysaccharide:
 - Polyribosylribitol Phosphate: 12 μ g
 - Conjugated to Tetanus protein: 22-36 μ g

Valence comparison

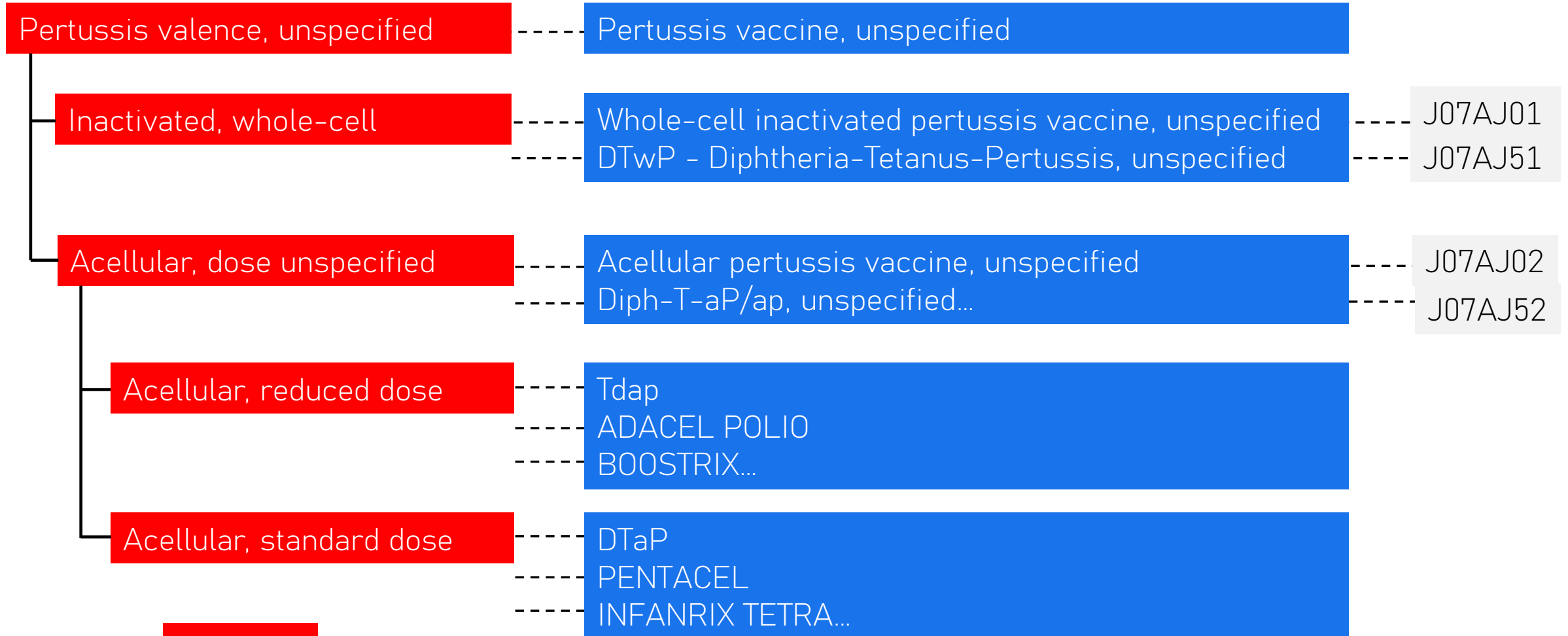
VAXELIS

- D Diphtheria valence, toxoid, standard dose
- T Tetanus valence, toxoid
- aP Pertussis valence, cellular, standard dose
- rHBsAg-10 Hepatitis B valence, recombinant HBs, 10 µg
- IPV Polio valence, trivalent, inactivated, whole-virus, injectable
- PRP-OMP Hib valence, PRP conjugated to OMP of *Neisseria meningitidis* B

HEXYON

- D Diphtheria valence, toxoid, standard dose
- T Tetanus valence, toxoid
- aP Pertussis valence, cellular, standard dose
- rHBsAg-10 Hepatitis B valence, recombinant HBs, 10 µg
- IPV Polio valence, trivalent, inactivated, whole-virus, injectable
- PRP-T Hib valence, PRP conjugated to tetanus toxoid

Hierarchical representation of valences

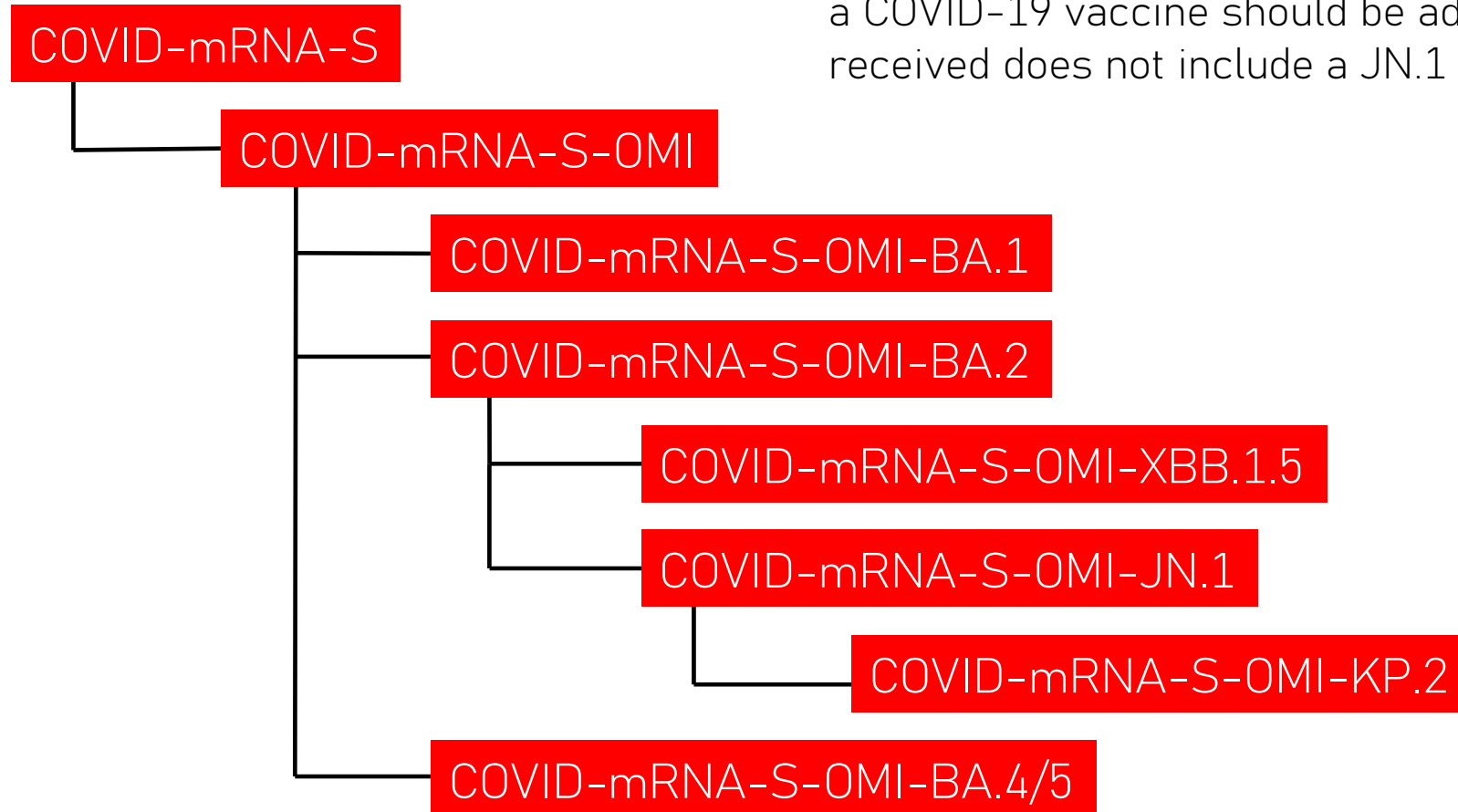


Valences
Vaccines
ATC codes

Enables the inclusion of vaccines that are not fully identified.
Illustrated here with the pertussis valences.

COVID-19 hierarchical valences

An example of a rule enabled by the hierarchisation:
a COVID-19 vaccine should be administered if the last dose
received does not include a JN.1 or KP.2 valence.



Applications

LOOKING FOR EQUIVALENT VACCINES
PRESCRIPTION

Mapping Tool

Language

EN



Vaccine name

REPEVAX



or pick a code

Code system



Code



	Code	Name	Diseases	Valences
	VAC0029	REPEVAX	Diphtheria Pertussis Poliomyelitis Tetanus	d ap IPV T

<https://nuva.syadem.com/>

Equivalent vaccines inferred from valence content

REPEVAX

ap | Pertussis valence, acellular, reduced dose
d | Diphtheria valence, toxoid, reduced dose
IPV | Polio valence, trivalent, inactivated, whole-virus, injectable
T | Tetanus valence, toxoid

Equivalent

Generalized

Specialized

i These are the vaccine with exactly the same valences as the original vaccine.

	Code	Name	Valences
<i>i</i>	VAC0736	TRIAXIS POLIO	d ap IPV T
<i>i</i>	VAC0585	TdapIPV - Diphtheria-Tetanus-Pertussis-Polio vaccine, low dose, unspecified	d ap IPV T
<i>i</i>	VAC0568	BOOSTRIX-POLIO	d ap IPV T
<i>i</i>	VAC0043	BOOSTRIX-TETRA	d ap IPV T
<i>i</i>	VAC0639	ADACEL POLIO	d ap IPV T
<i>i</i>	VAC0734	POLIO BOOSTRIX	d ap IPV T

Equivalent vaccines inferred from valence content

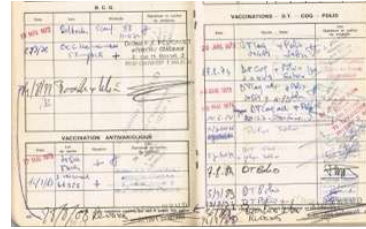
INFANRIXTETRA

aP | Pertussis valence, acellular, standard dose
D | Diphtheria valence, toxoid, standard dose
IPV | Polio valence, trivalent, inactivated, whole-virus, injectable
T | Tetanus valence, toxoid

Code	Name	Valences			
VAC0347	QUADRACEL	D	aP	IPV	T
VAC0350	QUATRO-VIRELON	D	aP	IPV	T
VAC0569	INFANRIX-IPV	D	aP	IPV	T
VAC0490	KINRIX	D	aP	IPV	T
VAC0037	TETRAVAC-ACELLULAIRE	D	aP	IPV	T
VAC0735	POLIOINFANRIX	D	aP	IPV	T
VAC0544	TETRAXIM	D	aP	IPV	T
VAC0551	DTaPIPv vaccine, standard dose, unspecified	D	aP	IPV	T
VAC0776	TETRABIK	D	aP	IPV	T
VAC0066	DTCP PASTEUR	D	aP	IPV	T
VAC0515	INFANRIX POLIO	D	aP	IPV	T

Valences: support for vaccine recommendation

Child
6 year-old



Vaccine history



Vaccine interpretation

Last valence received at the age of 6:
ap Pertussis valence, acellular, reduced dose



Valence-based recommendation

aP Pertussis valence, acellular, standard dose at the age of 11

The concept of valence could be a means of prescribing a vaccine using a generic name, without reference to a specific brand name.

Perspective



Classification of valences

1. Live

1.1. Live attenuated

1.2. Live viral vector

1.2.1. Replicating viral vector

1.2.2. Non-replicating viral vector

1.3. Chimeric live

2. Non-live

2.1. Nucleic acid

2.2. Whole inactivated

2.3. Subunit

Non-live valences

2.1. Nucleic acid

2.4.1. DNA

2.4.2. RNA

2.2. Inactivated

2.2.1. Whole-cell (bacterial)

2.2.2. Whole-virion (viral)

2.2.3. Split (viral)

2.3. Subunit

2.3.1. Toxoid

2.3.1. Polysaccharide unconjugated

2.3.2. Polysaccharide conjugated

2.3.3. Purified protein

2.3.4. Recombinant protein – non-VLP

2.3.5. Recombinant protein – VLP

2.3.6. Multi-antigen protein

INFANRIX HEXA : 6 diseases, 10 antigens

Valence	Class	Disease
D	Toxoid	Diphtheria
T	Toxoid	Tetanus
aP	Multi-antigen protein	Pertussis
IPV	Inactivated whole-virion	Polio
rHBsAg-10	Recombinant protein	Hepatitis B
PRP-T	Polysaccharide conjugated	<i>H. Influenzae</i> b



Diphtheria (toxoid)-Tetanus (toxoid)-Pertussis (Multi-antigen protein) – Polio (Inactivated whole-virion) – Hepatitis B (Recombinant protein) – Hib (Polysaccharide conjugated) vaccine,

Conclusion

- The valence concept enables seamless navigation across heterogeneous vaccine coding systems.
- It empowers users with more precise, interoperable, and automated vaccination decision-making.
- Designed to be adaptable, the valence model can evolve across levels of abstraction and shifting public health contexts - without disrupting vaccine code integrity.