

Herramienta web para la integración de sistemas de predicciones inmunoinformáticas (REVSINE)

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Proyecto de residencias profesionales

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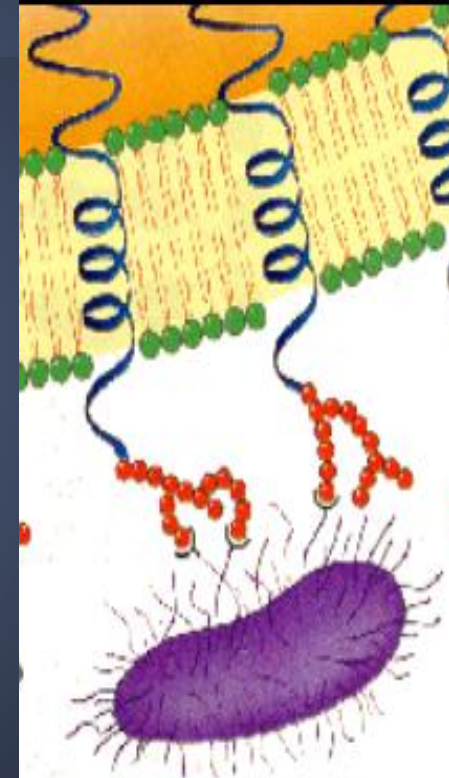
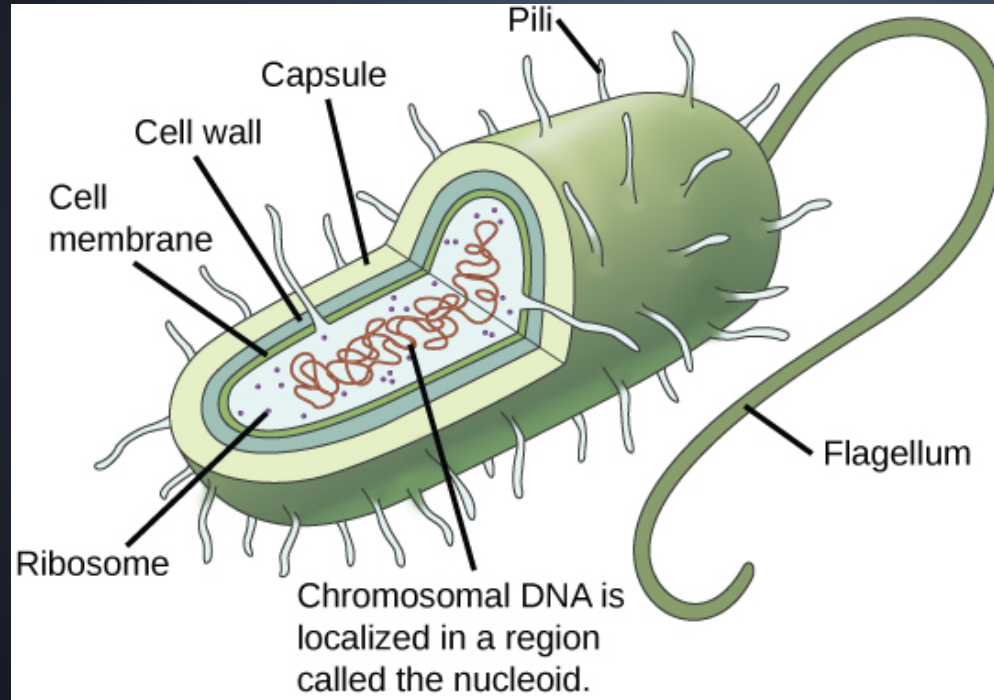
Introducción

- **Las vacunas tradicionales**
 - Empleando patógenos muertos
- **Finalidad de las vacunas:**
 - Respuesta para eliminar al patógeno.
 - Memoria inmune adaptativa.
 - Prevenir reinfecciones.

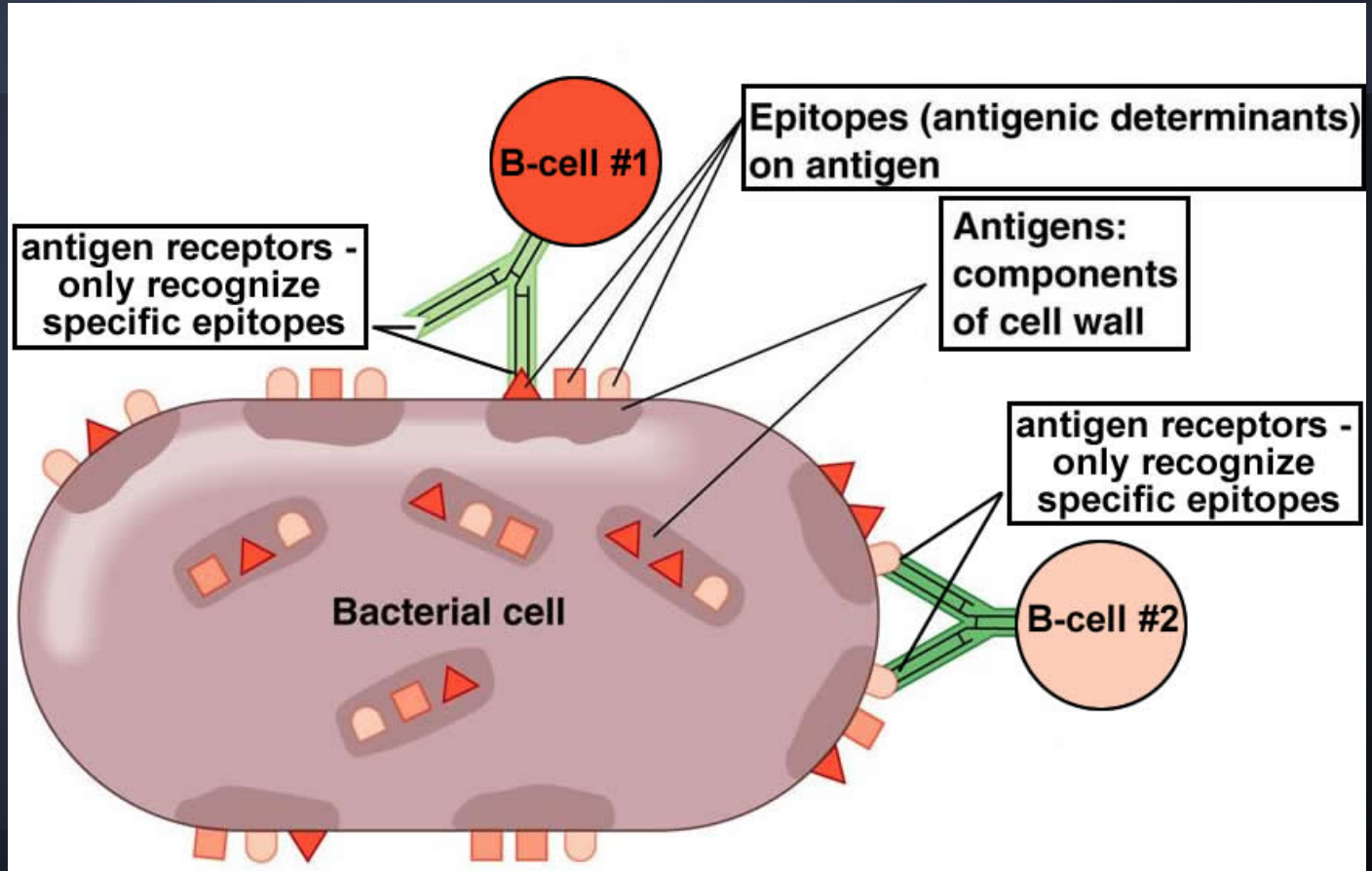


Introducción

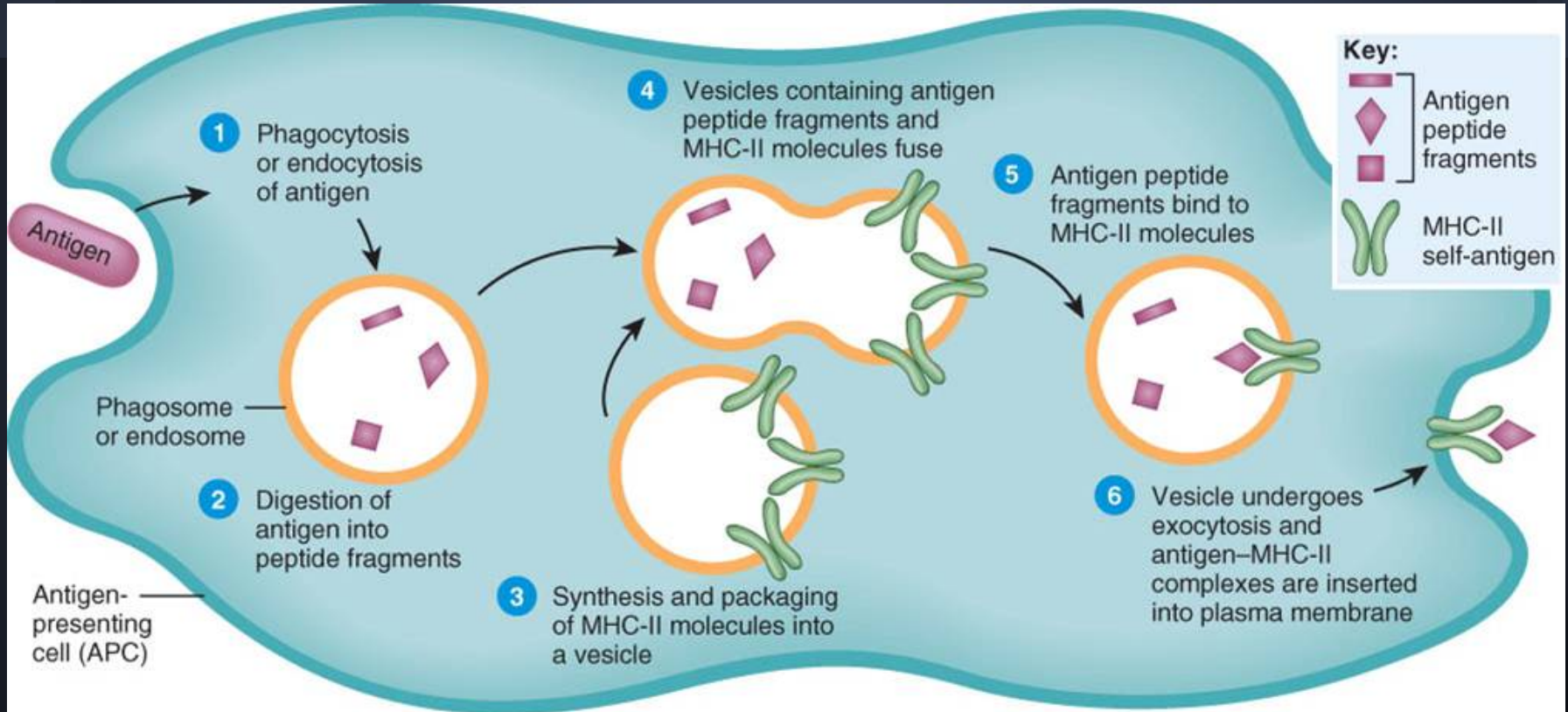
Patógenos: Patrones moleculares asociados a patógenos (PAMP's).



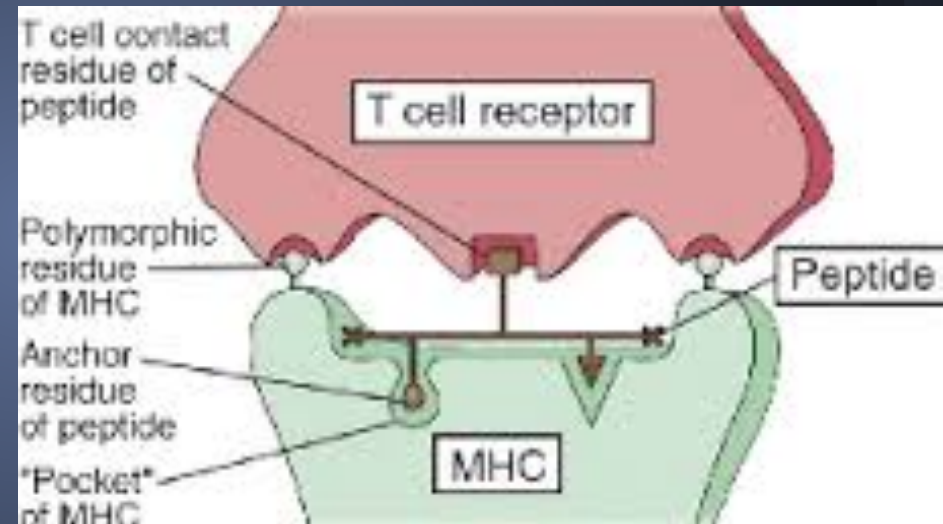
Epitopes



Procesamiento y presentación de epitopes

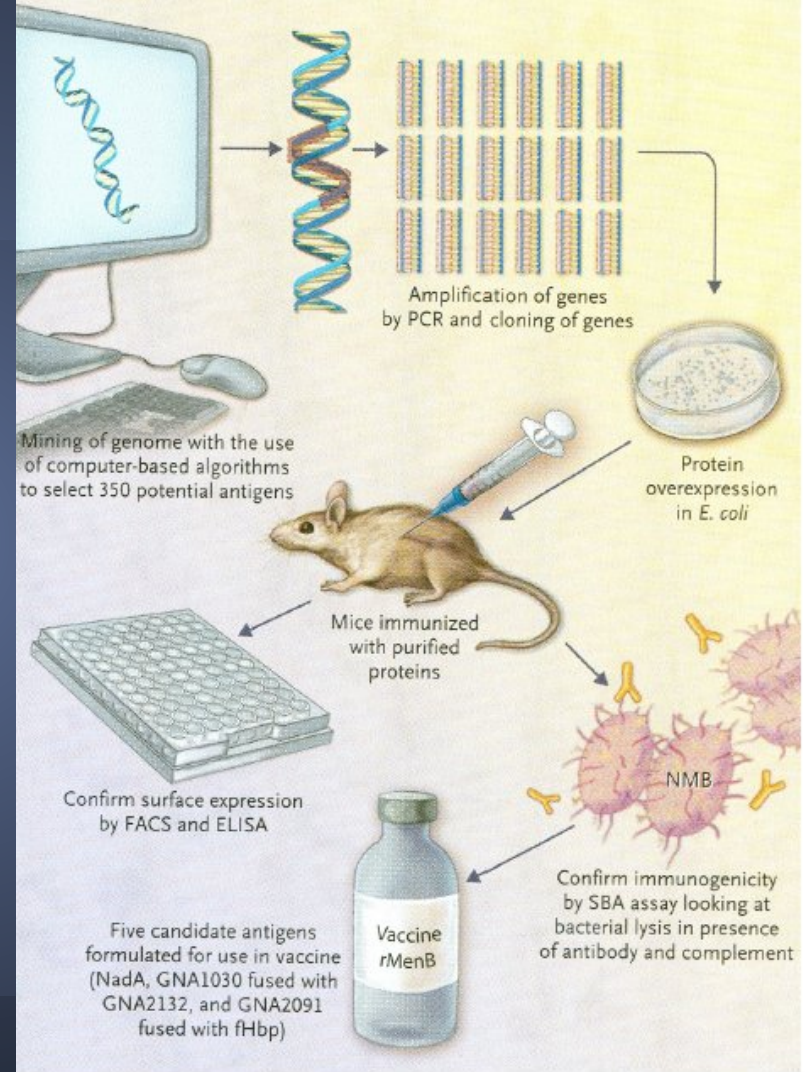


Respuesta inmunológica



Antecedentes

- Estrategias computacionales
 - Extensas bases de datos
 - Modelado matemático
- Herramientas inmunoinformáticas
 - Localización subcelular
 - Unión de péptidos a moléculas MHC I
 - Unión de péptidos a moléculas MHC II
 - Localización de epitopos de



Análisis independientes

About CELLO

CELLO v.2.5: subCELLular Localization predictor

ORGANISMS

☒ Gram negative

☐ Gram positive

☐ Eukaryotes

SEQUENCES

☐ DNA

☒ Protein

Paste the query sequences in FASTA format below

<1888086|Gensark|outer_membrane|extracellular|Autotransporter||major_ring-forming_surface_protein|BPECU097|MTKISDVQENP|LNKKNSSLSLNKRPFP|LIATTLPASSLP|NADAGAGAP|NAESITVY|NQAINATATV|SOH|GNATYTTTGATTTCTATAP|TAP|EVEV|GATV|VMFTY|DQ|AN|AQN|GAGAP|NL|NP|G|G|ASS|TACTYTT|NL|SGAAN|AL|G|NL|L|G|A|N|AT|L|TN|T|N|G|S|A|S|G|P|V|N|N|K|D|AT|N|AT|T|S|G|G|AT|N|L|V|T|N|T|S|G|T|N|L|C|T|D|K|Q|L|P|H|S|L|N|G|T|A|V|T|G|Q|A|P|A|V|L|T|G|I|S|T|G|G|I|N|V|T|F|E|K|T|N|G|G|I|A|G|N|T|G|S|L|N|N|V|T|F|E|Q|G|V|H|T|G|V|C|A|S|T|G|O|N|T|L|U|F|G|A|T|G|A|T|G|G|T|L|E|N|G|I|T|H|T|N|V|N|S|P|L|T|A|T|T|P|A|A|G|G|P|A|N|A|T|P|Q|S|N|S|A|G|G|G|V|N|L|P|N|F|A|K|E|T|P|A|K|A|P|A|F|A|N|I|T|A|T|N|G|A|N|V|P|T|G|G|L|N|A|L|T|S|T|L|Q|C|I|N|T|L|V|N|T|N|I|V|T|N|P|L|L|T|G|V|V|T|P|N|A|S|N|T|L|P|H|N|T|S|T|G|D|A|N|T|N|G|V|V|N|D|N|A|G|S|G|Q|A|T|S|H|T|N|P|H|L|N|L|K|E|G|S|L|A|G|A|G|A|N|A|K|A|S|Q|C|I|Q|V|L|N|F|R|G|S|A|N|A|L|T|N|G|T|G|A|T|L|V|N|T|T|L|A|H|P|R|Q|A|Q|V|N|V|T|V|G|N|S|S|A|N|V|L|E|A|P|N|A|S|A|T|T|Y|G|V|T|G|G|T|S|H|Y|N|W|G|S|T|S|V|L|P|A|N|A|N|D|I|P|T|T|I|G|A|T|S|T|L|V|D|A|P|G|G|P|N|D|L|A|G|K|L|V|T|Y|N|G|T|N|L|S|H|N|T|L|Q|G|L|Y|G|S|P|R|A|P|A|L|L|A|K|A|K|O|S|H|E|R|A|T|P|C|E|P|L|N|V|L|N|S|N|G|T|S|P|G|S|G|N|P|N|I|T|E|G|V|A|V|S|T|A|L|T|N|Q|A|T|T|N|H|N|T|S|V|N|L|V|K|S|D|S|V|L|T|A|S|E|H|N|Q|L|T|N|N|L|N|G|A|L|L|N|G|S|A|G|T|G|G|V|A|N|H|L|J|A|S|N|G|I|N|G|A|A|V|T|P|Q|S|V|S|F|H|N|Q|A|N|T|A|L|Q|N|T|Z|L|A|T|G|G|S|N|P|P|H|L|T|V|Q|A|N|S|A|T|T|A|S|D|Q|N|A|S|L|S|O|N|A|L|P|K|Y|V|N|A|N|G|H|G|A|G|G|O|B|A|T|N|G|N|F|S|S|L|G|I|D|T|S|N|V|V|T|C|T|E|P|F|O|R|E|S|P|H|P|N|K|S|T|O|N|L|Y|H|G|G|T|E|R|A|N|V|A|T|V|N|E|G|S|A|N|V|T|V|S|V|S|F|D|P|A|K|L|A|V|N|A|V|G|A|V|E|T|N|A|N|A|G|H|T|P|A|L|L|O|S|I|P|L|G|L|Q|T|G|S|T|S|H|G|S|G|Q|A|N|Q|V|T|T|Y|T|S|Q|A|N|T|S|E|A|N|L|A|T|A|L|A|S|H|Y|L|L|A|K|I|S|L|N|N|G|L|S|N|P|H|G|S|P|H|N|K|

Or upload from file:

Seleccionar archivo

No se eligió archivo

Reset

Submit

NetMHCIIpan 3.0 Server

NetMHCIIpan 3.0 server predicts binding of peptides to MHC class II molecules. The predictions are available for all three human MHC class II isotypes, HLA-DP, HLA-DP and HLA-DQ. NetMHCIIpan 3.0 server can produce predictions for peptides of 9 - 19 amino acids in length.

Two submission types are handled - the list of peptides or a protein sequence in FASTA format. The server provides a possibility for the user to choose MHC molecule in the MHC protein sequence of interest. For HLA-DP and HLA-DQ molecules alpha and beta chains can be chosen separately from the list. Also, for HLA-DP, HLA-DQ and molecule should be uploaded separately. For HLA-DQ molecule, the sequence should be validated.

The prediction values are given in nm IC50 values and as % Rank to a set of 200,000 random natural peptides. The user has a choice of setting the threshold for defining strong Rank. Strong and weak binding peptides will be indicated in the output. The output can also be sorted based on predicted binding affinity as well as filtered on the user-specific.

The project is a collaboration between CBS, [JMI](#), and [LIS](#).

View the [version history](#) of this server. All the previous versions are available on line, for comparison and reference.

Instructions

Output format

Article abstract

SUBMISSION

Type of input

Fasta

Paste a single sequence or several sequences in [FASTA](#) format into the field below:

or submit a file in [FASTA](#) format directly from your local disk:

Seleccionar archivo

No se eligió archivo

NetMHCpan 2.8 Server

NetMHCpan server predicts binding of peptides to any known MHC molecule using artificial neural networks (ANNs). The method is trained on more than 150,000 query molecules. Predictions can be made for HLA-A, B, C, E and G alleles, as well as for non-human primates, mouse, cattle and pig. Further, the user can upload full length restricted peptides from any given protein of interest.

Version 2.8 has been trained on extended data set including 10 prevalent HLA-C and 7 prevalent BoLA MHC-I molecules.

Predictions can be made for 8-14 mer peptides. Note, that all non 9mer predictions are made using approximations. Most HLA molecules have a strong preference for binding. The prediction values are given in nm IC50 values and as % Rank to a set of 200,000 random natural peptides. For alleles distant to the MHC molecules included in the training set, the prediction values are approximations.

The project is a collaboration between CBS, [JMI](#), and [LIS](#).

View the [version history](#) of this server. All the previous versions are available on line, for comparison and reference.

Instructions

Output format

Article abstract

SUBMISSION

Type of input

Fasta

Paste a single sequence or several sequences in [FASTA](#) format into the field below:

BepiPred 1.0 Server

BepiPred 1.0 server predicts the location of linear B-cell epitopes using a combination of a hidden Markov model and a propensity scale method.

Instructions

Output format

Data

SUBMISSION

Paste a single sequence or several sequences in [FASTA](#) format into the field below:

Submit a file in [FASTA](#) format directly from your local disk:

Seleccionar archivo

No se eligió archivo

Score threshold for epitope assignment

0.35

Submit

Clear fields

Restrictions:

At most 2000 sequences and 200,000 amino acids per submission; each sequence not less than 10 and not more than 6000 amino acids.

Confidentiality:

The sequences are kept confidential and will be deleted after processing.

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Análisis de resultados laboratorio

NetMHC1



NetMHCII



Tabla de traslape

```
# NetMHCpan version 2.8
# Input is in FSA format
# Peptide length 9,10
HLA-A01:01 : Estimated prediction accuracy 0.853 (using nearest neighbor HLA-A01:01)
```

pos	HLA	peptide	Identity	1-LogSK(aff)	affinity(nM)	NRank	BindLevel
114	HLA-A*01:01	QGSPIADNY	MAP1609C	0.539	146.17	0.17	<= SB
93	HLA-A*01:01	NFPIFENY	MAP1609C	0.504	215.03	0.25	<= SB
274	HLA-A*01:01	SSNLKPDQY	MAP1609C	0.502	217.75	0.25	<= SB
40	HLA-A*01:01	FSPRLPVEY	MAP1609C	0.351	1120.65	0.80	<= WB
139	HLA-A*01:01	FLTSELPSY	MAP1609C	0.352	1491.37	0.80	<= WB
0	HLA-A*01:01	MTLSEKIVR	MAP1609C	0.323	1524.49	1.00	<= WB
296	HLA-A*01:01	NANQIHWEY	MAP1609C	0.318	1594.56	1.00	<= WB
175	HLA-A*01:01	AVNPQDQTY	MAP1609C	0.301	1920.82	1.00	<= WB
140	HLA-A*01:01	TLSELPSYLA	MAP1609C	0.257	3106.04	1.50	<= WB
140	HLA-A*01:01	LTSELPSYL	MAP1609C	0.246	3504.76	2.00	<= WB
205	HLA-A*01:01	LAKDAGGY	MAP1609C	0.233	4080.24	2.00	<= WB
141	HLA-A*01:01	TSSELPSYLA	MAP1609C	0.230	4157.81	2.00	<= WB
204	HLA-A*01:01	GLAKDAGGY	MAP1609C	0.210	4841.76	3.00	<= WB
110	HLA-A*01:01	PIVGQSSPY	MAP1609C	0.205	5454.75	3.00	<= WB
91	HLA-A*01:01	LTDLPAFHY	MAP1609C	0.193	6180.38	3.00	<= WB
118	HLA-A*01:01	YADWQDQY	MAP1609C	0.190	6590.88	3.00	<= WB
113	HLA-A*01:01	QGSPIADNY	MAP1609C	0.180	7146.34	3.00	<= WB
92	HLA-A*01:01	IVNPAFENY	MAP1609C	0.179	7241.56	3.00	<= WB
297	HLA-A*01:01	ANGIHWEY	MAP1609C	0.178	7288.73	4.00	<= WB
166	HLA-A*01:01	MSGSSAMILA	MAP1609C	0.178	7304.02	4.00	<= WB
241	HLA-A*01:01	QNTLPLQY	MAP1609C	0.176	7437.22	4.00	<= WB
118	HLA-A*01:01	YADWQDQY	MAP1609C	0.176	7437.83	4.00	<= WB
0	HLA-A*01:01	MTLSEKIVR	MAP1609C	0.171	7824.19	4.00	<= WB
154	HLA-A*01:01	CRGSSAMLA	MAP1609C	0.170	7978.45	4.00	<= WB
275	HLA-A*01:01	SNLKPDQY	MAP1609C	0.169	8027.69	4.00	<= WB
176	HLA-A*01:01	AVNPQDQTY	MAP1609C	0.167	8234.95	4.00	<= WB
168	HLA-A*01:01	GSSAMILAV	MAP1609C	0.166	8260.57	4.00	<= WB

```
# NetMHCpan version 3.0
# Input is in FSA format
# Peptide length 15
# Threshold for strong binding peptides (IC50) 50.00 nM
# Threshold for weak binding peptides (IC50) 500.00 nM
# Threshold for strong binding peptides (Rank) 0.5%
# Threshold for weak binding peptides (Rank) 2%
```

# Allele: HLA-DPA1*01:03:DPB1*01:01									
pos	Allele	peptide	Identity	Pos	Core 1-LogSK(aff)	Affinity(nM)	NRank	BindingLevel	
131	HLA-DPA1*01:03:DPB1*01:01	CTYMETLTSLP	MAP1609C	3	TYMETLTSLP	0.783	10.45	0.50	<= SB
130	HLA-DPA1*01:03:DPB1*01:01	ACTYMETLTSLP	MAP1609C	4	TYMETLTSLP	0.777	11.14	0.80	<= SB
132	HLA-DPA1*01:03:DPB1*01:01	STYMETLTSLP	MAP1609C	2	TYMETLTSLP	0.760	13.41	0.80	<= SB
129	HLA-DPA1*01:03:DPB1*01:01	ACCTYMETLTSLP	MAP1609C	5	TYMETLTSLP	0.747	15.48	1.50	<= SB
128	HLA-DPA1*01:03:DPB1*01:01	ACCTYMETLTSLP	MAP1609C	6	TYMETLTSLP	0.740	16.63	1.50	<= SB
133	HLA-DPA1*01:03:DPB1*01:01	TYMETLTSLPY	MAP1609C	1	TYMETLTSLP	0.739	16.47	1.50	<= SB
134	HLA-DPA1*01:03:DPB1*01:01	TYMETLTSLPSY	MAP1609C	0	TYMETLTSLP	0.711	22.69	3.00	<= SB
126	HLA-DPA1*01:03:DPB1*01:01	MYETLTSLPSLA	MAP1609C	4	FLTSELPSY	0.669	35.30	5.00	<= WB
128	HLA-DPA1*01:03:DPB1*01:01	MYETLTSLPSLA	MAP1609C	4	FLTSELPSY	0.668	35.30	5.00	<= WB
65	HLA-DPA1*01:03:DPB1*01:01	PAFMYQSLSVTR	MAP1609C	2	FEWYQSL	0.658	41.52	5.00	<= WB
97	HLA-DPA1*01:03:DPB1*01:01	PEMYQSLSVTR	MAP1609C	0	FEWYQSL	0.636	51.25	7.00	<= WB
84	HLA-DPA1*01:03:DPB1*01:01	TPMRYQSSQVY	MAP1609C	3	FEWYQSL	0.649	44.49	6.00	<= WB
137	HLA-DPA1*01:03:DPB1*01:01	ETLTSELPSYLA	MAP1609C	2	FLTSELPSY	0.626	57.08	8.00	<= WB
195	HLA-DPA1*01:03:DPB1*01:01	PGFYVLSALLP	MAP1609C	5	FVRSNLP	0.601	69.02	9.00	<= WB
172	HLA-DPA1*01:03:DPB1*01:01	QKQSTYMETLTSLP	MAP1609C	4	CTYMETLTSLP	0.608	69.63	9.00	<= WB
205	HLA-DPA1*01:03:DPB1*01:01	AEFLNPFVRSNLP	MAP1609C	6	FVRSNLP	0.607	69.92	9.00	<= WB
178	HLA-DPA1*01:03:DPB1*01:01	HPQFTYAGLSALLP	MAP1609C	3	QFTYAGLS	0.606	70.70	9.00	<= WB
180	HLA-DPA1*01:03:DPB1*01:01	QFTYAGLSALLP	MAP1609C	4	YAGLSALL	0.604	72.43	9.00	<= WB
69	HLA-DPA1*01:03:DPB1*01:01	ETNPAWYQSLSV	MAP1609C	4	FEWYQSL	0.598	76.34	10.00	<= WB
181	HLA-DPA1*01:03:DPB1*01:01	QFTYAGLSALLP	MAP1609C	3	YAGLSALL	0.593	81.73	15.00	<= WB
207	HLA-DPA1*01:03:DPB1*01:01	LEFVRSNLPQDA	MAP1609C	4	FVRSNLP	0.581	83.05	15.00	<= WB
268	HLA-DPA1*01:03:DPB1*01:01	LEFVRSNLPQDA	MAP1609C	3	FVRSNLP	0.587	87.40	15.00	<= WB
4	HLA-DPA1*01:03:DPB1*01:01	SENVNWRGLVLA	MAP1609C	3	VNWRGLL	0.569	90.07	15.00	<= WB
3	HLA-DPA1*01:03:DPB1*01:01	LSKVNWRGLVLA	MAP1609C	4	FLTSELPSY	0.574	90.07	15.00	<= WB
138	HLA-DPA1*01:03:DPB1*01:01	LSKVNWRGLVLA	MAP1609C	3	FLTSELPSY	0.576	98.19	15.00	<= WB
2	HLA-DPA1*01:03:DPB1*01:01	LSKVNWRGLVLA	MAP1609C	5	VNWRGLL	0.570	104.35	15.00	<= WB
5	HLA-DPA1*01:03:DPB1*01:01	EVNTPNVLQDA	MAP1609C	2	VNWRGLL	0.566	111.68	15.00	<= WB
114	HLA-DPA1*01:03:DPB1*01:01	QGSPIADWQDQY	MAP1609C	3	PIADWQDA	0.561	115.01	15.00	<= WB
113	HLA-DPA1*01:03:DPB1*01:01	QGSPIADWQDQY	MAP1609C	4	PIADWQDA	0.557	120.88	15.00	<= WB
263	HLA-DPA1*01:03:DPB1*01:01	AEFLNPFVRSNLP	MAP1609C	3	ELFNPVRS	0.555	123.26	15.00	<= WB
112	HLA-DPA1*01:03:DPB1*01:01	QGSPIADWQDQY	MAP1609C	5	PIADWQDA	0.552	125.11	15.00	<= WB
124	HLA-DPA1*01:03:DPB1*01:01	QGSPIADWQDQY	MAP1609C	5	MYQSSAM	0.551	125.11	15.00	<= WB
264	HLA-DPA1*01:03:DPB1*01:01	PAELNPFVRSNLP	MAP1609C	2	ELFNPVRS	0.540	144.60	32.00	<= WB
96	HLA-DPA1*01:03:DPB1*01:01	MYQSSAMILAV	MAP1609C	1	VNWRGLL	0.537	149.91	32.00	<= WB
111	HLA-DPA1*01:03:DPB1*01:01	VQGSPIADWQDA	MAP1609C	6	PIADWQDA	0.536	151.14	32.00	<= WB
261	HLA-DPA1*01:03:DPB1*01:01	ANPFIENPQDQY	MAP1609C	4	AFPIENPQ	0.534	154.82	32.00	<= WB
269	HLA-DPA1*01:03:DPB1*01:01	ENPFIENPQDQY	MAP1609C	2	FVRSNLP	0.533	155.81	32.00	<= WB
92	HLA-DPA1*01:03:DPB1*01:01	PEMYQSLSVTR	MAP1609C	6	FEWYQSL	0.530	162.27	32.00	<= WB
262	HLA-DPA1*01:03:DPB1*01:01	MPAEFLNPFVRSN	MAP1609C	3	AEFLNPFV	0.529	163.60	32.00	<= WB
115	HLA-DPA1*01:03:DPB1*01:01	QGSPIADWQDQY	MAP1609C	6	PIADWQDA	0.525	165.50	32.00	<= WB
91	HLA-DPA1*01:03:DPB1*01:01	QNTLPLQDQY	MAP1609C	6	FEWYQSL	0.511	196.48	32.00	<= WB
260	HLA-DPA1*01:03:DPB1*01:01	GAPMYQSLSVTR	MAP1609C	6	FEWYQSL	0.510	196.48	32.00	<= WB
182	HLA-DPA1*01:03:DPB1*01:01	ETYAGLSALLP	MAP1609C	2	YAGLSALL	0.506	209.49	32.00	<= WB
1	HLA-DPA1*01:03:DPB1*01:01	TLSELPSYLA	MAP1609C	2	VNWRGLL	0.506	209.72	32.00	<= WB
73	HLA-DPA1*01:03:DPB1*01:01	HPQFTYAGLSALLP	MAP1609C	4	QFTYAGLS	0.502	217.95	32.00	<= WB
96	HLA-DPA1*01:03:DPB1*01:01	MYQSSAMILAV	MAP1609C	6	FEWYQSL	0.500	218.05	32.00	<= WB
73	HLA-DPA1*01:03:DPB1*01:01	PAVYLLQDQDQY	MAP1609C	3	YLLQDQDA	0.497	232.05	32.00	<= WB
72	HLA-DPA1*01:03:DPB1*01:01	PAVYLLQDQDQY	MAP1609C	3	YLLQDQDA	0.496	242.65	32.00	<= WB
270	HLA-DPA1*01:03:DPB1*01:01	MYQSSAMILAV	MAP1609C	1	FVRSNLP	0.489	251.06	32.00	<= WB
71	HLA-DPA1*01:03:DPB1*01:01	GRVYLLQDQDQY	MAP1609C	1	YLLQDQDA	0.475	273.53	32.00	<= WB
74	HLA-DPA1*01:03:DPB1*01:01	AVYLLQDQDQY	MAP1609C	2	YLLQDQDA	0.477	285.97	32.00	<= WB
80	HLA-DPA1*01:03:DPB1*01:01	THREMYQDQDQY	MAP1609C	2	MYQSSAM	0.473	282.22	32.00	<= WB
126	HLA-DPA1*01:03:DPB1*01:01	CKAQSTCTYMETLTSLP	MAP1609C	5	CTYMETLTSLP	0.475	290.97	32.00	<= WB
79	HLA-DPA1*01:03:DPB1*01:01	NSRYVYLLQDQDA	MAP1609C	2	YLLQDQDA	0.473	291.23	32.00	<= WB
361	HLA-DPA1*01:03:DPB1*01:01	HWYQAGLSNLP	MAP1609C	2	MYQSSAM	0.464	330.30	32.00	<= WB
139	HLA-DPA1*01:03:DPB1*01:01	FLTSELPSYLA	MAP1609C	6	FLTSELPSY	0.462	330.30	32.00	<= WB
259	HLA-DPA1*01:03:DPB1*01:01	GAPMYQSLSVTR	MAP1609C	6	VNWRGLL	0.459	362.02	32.00	<= WB
261	HLA-DPA1*01:03:DPB1*01:01	GAPMYQSLSVTR	MAP1609C	6	AEFLNPFV	0.457	362.02	32.00	<= WB
802	HLA-DPA1*01:03:DPB1*01:01	SNYQAGLSNLP	MAP1609C	3	VNWRGLL	0.441	421.23	32.00	<= WB
76	HLA-DPA1*01:03:DPB1*01:01	FLTSELPSYLA	MAP1609C	6	FLTSELPSY	0.439	449.88	32.00	<= WB
271	HLA-DPA1*01:03:DPB1*01:01	FVRSNLPQDQY	MAP1609C	0	FVRSNLP	0.425	454.01	50.00	<= WB
116	HLA-DPA1*01:03:DPB1*01:01	SPVYLLQDQDQY	MAP1609C	2	YLLQDQDA	0.424	454.01	50.00	<= WB
183	HLA-DPA1*01:03:DPB1*01:01	YAGLSALLPQDQY	MAP1609C	1	YAGLSALL	0.424	507.96	50.00	<= WB
41	HLA-DPA1*01:03:DPB1*01:01	QNTLPLQDQY	MAP1609C	2	QNTLPLQD	0.413	536.96	50.00	<= WB
258	HLA-DPA1*01:03:DPB1*01:01	LOGMNAEFLNPFV	MAP1609C	3	ANPFIENP	0.413	587.88	50.00	<= WB

Tabla Transposición epitopos de células T presentados por MHC I y MHC II

Proteína	MHC I Epitopo Cel T				MHC I Epitopo Cel T			
	Secuencia	Posición	Alélio	Unión	Secuencia	Posición	Alélio	Unión
MAP1609C	DQFIYAGSL	180-188	BoLA-T2c	WB	HPDQFIYAGSLALL	178-192	H-2-IAb	SB
	LLVGAAAV	14-22	BoLA-T2c	WB	CPRLVVGAAAVTLP	11-25	H-2-IAb	SB
	YQGLSVIM	101-109	BoLA-T2c	WB	FEWYQGLSVIMPV	97-111	H-2-IAb	WB
	NAAGGHNAV	284-292	BoLA-T2c	WB	FQDAYNAAGGHNAV	279-293	H-2-IAb	SB
	SMGSSAMI	165-173	BoLA-T2c	WB	AVGISMGSSAMILA	161-175	H-2-IAb	WB

- Integración en una sola plataforma
- Resultados en formatos homogéneos y científicos

Justificación

- El presente trabajo aportaría una solución informática al área de vacunología.
- Aumentaría la velocidad y facilidad de selección de epitopes candidatos a vacunas.
- Evitaría los procesos de edición para formatos de presentación de resultados en tablas.

Objetivo general

Desarrollar una aplicación web integradora que permita predecir la ubicación subcelular de proteínas de patógenos y los péptidos de unión a células T y B, cuyos resultados se despliegan en formatos de artículos científicos.

Objetivo particulares

1. Analizar los mecanismos computacionales de las herramientas inmunoformáticas seleccionadas.
2. Construir una sola interfaz web integradora de las herramientas inmunoinformáticas.
3. Programar el análisis e interpretación de los datos.
4. Diseñar y desplegar los resultados en formatos de artículos científicos.

Materiales y métodos

- Materiales
 - Linux, IDE Eclipse, lenguajes; PHP, HTML y JavaScript.
 - Librería cURL.
- Métodos
 - Se programó la interfaz de usuario y se aseguró la integración de cuatro herramientas inmunoinformáticas disponibles en la web:
 - CELLO (cello.life.nctu.edu.tw)
 - NetMHCI (www.cbs.dtu.dk/services/NetMHCI/)
 - NetMHCII (www.cbs.dtu.dk/services/NetMHCII/)
 - BepiPred (<http://www.cbs.dtu.dk/services/BepiPred/>)

Materiales y métodos

Métodos

- Se programó un intérprete homogéneo en PHP.
 - Expresiones regulares
- Se diseñaron las tablas de resultado en HTML.
- Se programó la creación automática de las tablas utilizando PHP.

**REVSINE es una herramienta
informática para dar soporte a
la vacunología**



<http://posgrado.itlp.edu.mx/revsine/index.php>

Conclusiones

1. Se logró la integración de herramientas inmunoinformáticas.
2. Sus resultados pueden ser interpretados fácilmente con programación.
3. Es posible utilizar las respuestas individuales para armar tablas en formato científico.

Trabajo Futuro

1. Agregar otras herramientas inmunoinformáticas y generar nuevas tablas de reporte.
2. Control de usuarios y registro de experimentos.
3. Descargar los servidores y ejecutarlos de forma local.
4. Utilizar cómputo paralelo para agilizar los resultados.
5. Agregar la espera de resultados de forma interactiva.
6. Agregar una versión de descarga para Windows.

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REVSINE

<https://github.com/jlsalazar/revxine.git>