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## Endo H/ PNGase F Assay for JESS Automated Western Blot

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Charis Ma<sup>1</sup>, Yu Chen<sup>1</sup>, sidranse<sup>1</sup>

<sup>1</sup>National Human Genome Research Institute



Charis Ma

National Institutes of Health

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**Protocol status:** Working

**We use this protocol and it's working**

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### Abstract

This protocol details an optimized Endo H and PNGase F digestion assay that produces samples that can be run with the JESS Automated Western Blot System.



## Materials

10X Glycoprotein Denaturing Buffer (NEB, Catalog# B0701S)

10X GlycoBuffer 3 (NEB, Catalog# B0701S)

10X GlycoBuffer 2 (NEB, Catalog# B0701S)

10% NP-40 (NEB, Catalog# B0701S)






EndoH (NEB, Catalog# P0703S)

PNGase F (NEB, Catalog# P0704S)







10K Amicon Ultra Centrifugal Filter (Millipore, UFC5010BK)










## Glycoprotein Denaturation

- 1 In an eppendorf tube, add  40  $\mu\text{g}$  protein,  2  $\mu\text{L}$  10X Glycoprotein Denaturing Buffer, and water for a total reaction volume of  20  $\mu\text{L}$  . Mix well.
- 2 Denature glycoprotein by heating reaction at  100  $^{\circ}\text{C}$  for  00:10:00 . 10m
- 3 Split reaction volume into 2 eppendorf tubes with 10  $\mu\text{L}$  in each tube. One tube will be used for the EndoH reaction, while the other will be used for the PNGaseF reaction.

## Endo H








- 4 Add the following to one reaction tube from step 3:  2  $\mu\text{L}$  10X GlycoBuffer 3,  2  $\mu\text{L}$  Endo H,  6  $\mu\text{L}$  water. Total volume will be  20  $\mu\text{L}$  .
- 5 Mix gently.
- 6 Incubate reaction at  37  $^{\circ}\text{C}$  for  01:00:00 1h

## PNGase F

- 7 Add the following to the second reaction tube from step 3:  2  $\mu\text{L}$  10X GlycoBuffer 2,  2  $\mu\text{L}$  NP-40,  6  $\mu\text{L}$  water,  1  $\mu\text{L}$  PNGaseF. Total volume will be  21  $\mu\text{L}$  .
- 8 Mix gently.
- 9 Incubate reaction at  37  $^{\circ}\text{C}$  for  01:00:00 . 1h

## Desalting and Washing for JESS Automated Western Blot System (Bio-Techne) preparation





- 10 Add  480  $\mu\text{L}$  and  479  $\mu\text{L}$  of 0.1X Sample Buffer (Bio-Techne) to the Endo H and PNGaseF reaction tubes respectively to make the final volume  500  $\mu\text{L}$  .
- 11 Add all  500  $\mu\text{L}$  of each reaction to a 10K Amicon Ultra Centrifugal Filter
- 12 Spin at  14000 x g for  00:20:00 . You will recover a final volume of  20  $\mu\text{L}$  , and the protein concentration recovered will be ~ [M] 1 mg/mL .

20m

## JESS Sample Preparation

5m

- 13 Add 4 parts of the washed Endo H/ PNGaseF sample from step 12 with 1 part of 5X Fluorescent Master Mix (Bio-Techne).
- 14 Vortex to mix
- 15 Incubate sample at  95  $^{\circ}\text{C}$  for  00:05:00 .
- 16 Load into JESS microplate (Bio-techne) to be detected by the Automated Western Blot System.

5m

## Protocol references

<https://www.neb.com/en-us/protocols/2012/10/18/endo-hf-protocol>

<https://www.neb.com/en-us/protocols/2014/07/31/pngase-f-protocol>