

## Anti-Human parainfluenza virus (HPIV) nucleoprotein (NP) antibody, mouse monoclonal (PINP-01) $\,$

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| Product code   | 65-180   |
| Size   | 100 μg   |
| Storage  | -20℃   |
| Concentration  | 1.0 mg/ml  |
| Buffer   | PBS(-) with 50% glycerol   |
| Purity   | Purified IgG fraction with protein A from hybridoma cell culture medium  |
| Immunogen  | Viral core fraction prepared from purified Human parainfluenza virus (HPIV type 3)   |
| Isotype  | mouse IgG2a κ  |
| Reactivity   | Reacts with the HPIV nucleocapsid protein (nucleoprotein, NP)  |
| Validation   | Specificity has been validated by western blotting   |
| Application  | 1.Western blotting: x1/200-400 (Fig.1)   |
|  | 2.Immunofluorescence: x1/400 (Fig.2)   |
| Background   | Human Parainfluenza Virus Type 3 (HPIV-3), a non-segmented, negative-strand RNA virus, causes lower respiratory illness in newborns, infants and older people. The genome of HPIV-3 encodes at least six proteins: the nucleoprotein (NP), phosphoprotein (P), RNA-dependent RNA polymerase (L), matrix protein (M), and two virus-host cell interaction glycoproteins, fusion protein (F) and hemagglutinin-neuraminidase (HN). The polypeptide of these antigens was identified by immunoprecipitation, immunoblotting and several serological analyses. The P, HN, NP, F and M antigens have a molecular weight of 84kDa, 72kDa (572 a.a.), 68kDa (515 a.a.), 53kDa and 40kDa polypeptide, respectively. The NP was shown to be structurally related to the major nucleocapsid protein. |
| Data Link  | UniProtKB: P06159 (NCAP_PI3H4)   |
| Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC |  |

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PROCEDURES. NOT FOR MILITARY USE.



**Data Images:** 65-180 Anti-Human parainfluenza virus (HPIV) nucleoprotein (NP) antibody, mouse monoclonal (PINP-01)

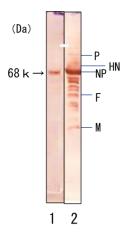


Fig.1. Identification of protein in HPIV-3 by Western blotting (WB)

Lane 1: HPNP-01 monoclonal antibody (MAb)

Lane 2: anti-HPIV-3 mouse serum

The virus purified from the culture supernatant of infected cells was applied to SDS-PAGE and WB. The MAb and antiserum were used at 1/200 and 1/400 dilution, respectively. The HRP-conjugated goat anti-mouse IgG was used at 1/2,000 as the second antibody and visualized by DAB (3,3'-Diaminobenzidine). A 68 kDa band was identified as HPIV NP.

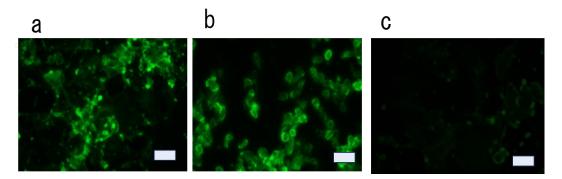


Fig.2. Detection of HPIV-3 protein infected in cells by immunofluorescence staining

(a) Vero cells were infected with HPIV on a slide glass and fixed with ethanol. (b) HPIV-infected Vero cells were treated with trypsin, smeared on the preparate and fixed with ethanol. (c) Uninfected Vero cells on a slide glass were fixed with ethanol. The MAb was used at 1/200 dilution. As the second antibody, FITC-conjugated rabbit anti-mouse IgG was used at 1/2,000 dilution. Bar maker represents 20µm.

References This antibody has not yet been used in publication.

**Related product**: 65-182 Anti-Human parainfluenza virus (HPIV) neutralizing antibody, mouse monoclonal (PIHN-01)