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Author Correction: USP4 positively regulates RLR-induced NF-κB activation by targeting TRAF6 for K48-linked deubiquitination and inhibits enterovirus 71 replication

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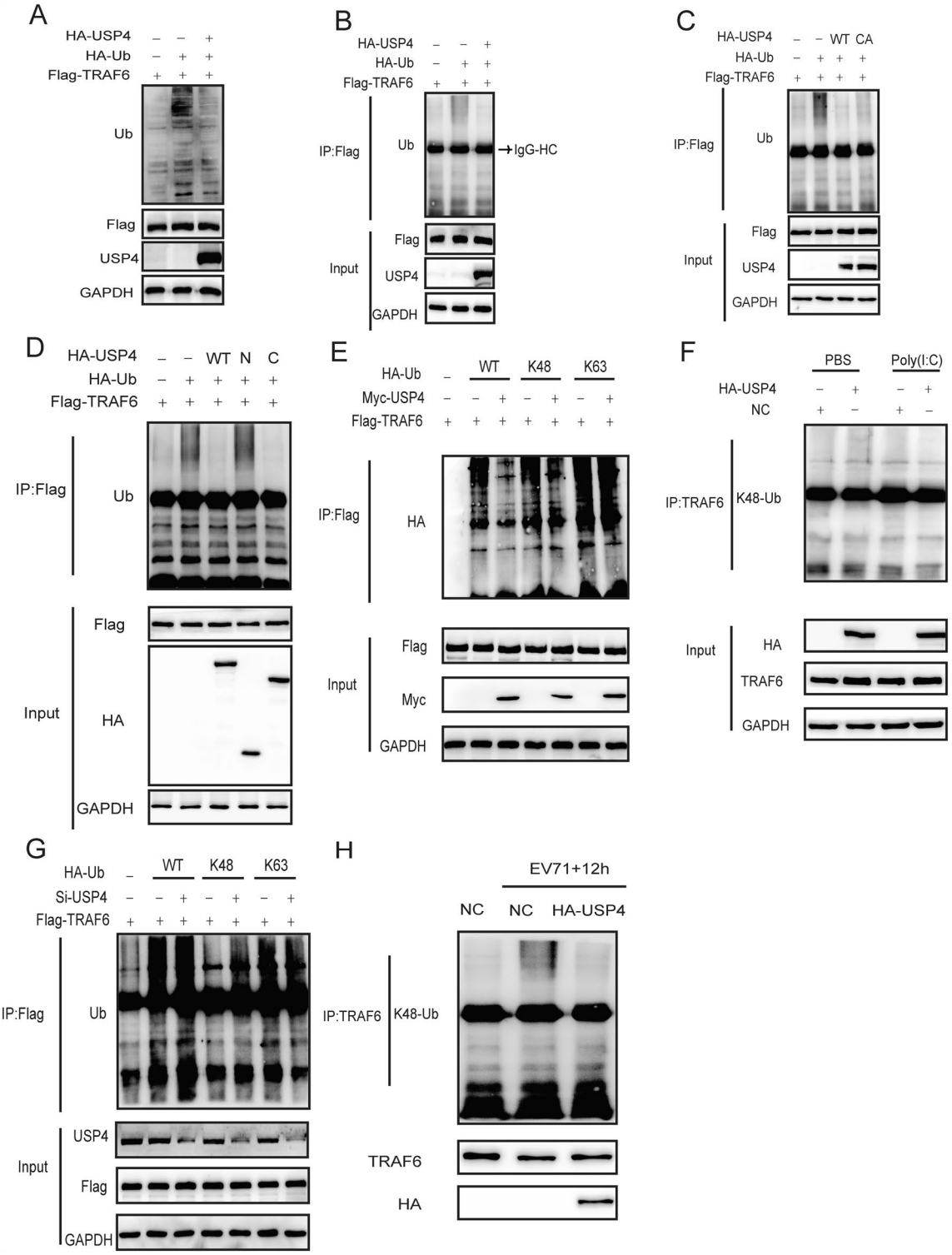
This Article contains an error.

As a result of an error during the figure assembly, in Figure 6B, the panels *Flag* and *GAPDH* of the "Input" are overlapping with the *Flag* and *GAPDH* panels of the "Input" in Figure 6C.

The corrected Figure 6 and its accompanying legend appears below as Figure 1.


A
HA-USP4 – – +
HA-Ub – + +
Flag-TRAF6 + + +
Ub
Flag
USP4
GAPDH
B
HA-USP4 – – +
HA-Ub – + +
Flag-TRAF6 + + +
IP:Flag
Ub
→IgG-HC
Input
Flag
USP4
GAPDH
C
HA-USP4 – – WT CA
HA-Ub – + + +
Flag-TRAF6 + + + +
IP:Flag
Ub
Input
Flag
USP4
GAPDH
D
HA-USP4 – – WT N C
HA-Ub – + + + +
Flag-TRAF6 + + + + +
IP:Flag
Ub
Input
Flag
HA
GAPDH
E
HA-Ub – WT K48 K63
Myc-USP4 – – + – + – +
Flag-TRAF6 + + + + + + +
IP:Flag
HA
Input
Flag
Myc
GAPDH
F
PBS Poly(I:C)
HA-USP4 – + – +
NC + – + –
IP:TRAF6
K48-Ub
Input
HA
TRAF6
GAPDH
G
HA-Ub – WT K48 K63
Si-USP4 – – + – + – +
Flag-TRAF6 + + + + + + +
IP:Flag
Ub
Input
USP4
Flag
GAPDH
H
EV71+12h
NC NC HA-USP4
IP:TRAF6
K48-Ub
TRAF6
HA

◂**Figure 1.** USP4 removes K48-linked polyubiquitination from TRAF6. (**A**) Western blot analysis of HEK293T cells transfected with Flag-TRAF6, HA-USP4, and HA-Ub for ubiquitination levels; GAPDH was used as a loading control. (**B,C**) HEK293T cell lysates transfected with Flag-TRAF6, HA-Ub, together with either wild-type HA-USP4 (WT) or mutated HA-USP4 (CA) were collected and immunoprecipitated with anti-Flag agarose beads. The eluted immunocomplexes was then subjected to SDS-PAGE analysis with anti-ubiquitin antibody. (**D**) Lysates from HEK293T cells transiently expressing Flag-tagged TRAF6 and HA-tagged ubiquitin, together with either HA-tagged USP4 or its truncated constructs (N or C) were immunoprecipitated with anti-Flag agarose beads; the eluted protein complexes was subjected to western blot analysis with anti-ubiquitin antibody. (**E**) HEK293T cells transfected with Flag-TRAF6, Myc-USP4, together with either WT HA-Ub or its mutants (K48 or K63) were harvested and subjected to immunoprecipitation with anti-Flag agarose beads followed by SDS-PAGE analysis with HA antibody. (**F**) Lysates from HEK293T cells transfected with HA-USP4 or a control vector followed by treatment with poly (I:C) (HMW; 30 μg/ml) for 8 h were subjected to immunoprecipitation with an anti-TRAF6 antibody. This was followed by western blot analysis of eluted immunocomplexes with K48 linkage-specific ubiquitin antibodies. (**G**) HEK293T cells transfected with Flag-TRAF6, control siRNA or USP4-specific siRNA, together with either WT HA-Ub or its mutants (K48 or K63) were harvested and subjected to immunoprecipitation with anti-Flag agarose beads followed by SDS-PAGE analysis with anti-ubiquitin antibody. (**H**) Western blot analysis of RD cells transfected with control plasmid or HA-USP4 plasmid for 48 h, followed by treatment with EV71 (MOI = 0.5) for 12 h. The cell lysates were subjected to immunoprecipitation with anti-TRAF6 antibody, followed by western blot analysis of eluted immunocomplexes with K48 linkage-specific ubiquitin antibodies.