

## **Product datasheet for AP12453PU-N**

## **U2AF1L5** (Center) Rabbit Polyclonal Antibody

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** IHC, WB

Recommended Dilution: ELISA: 1/1,000.

Western Blot: 1/50-1/100.

Immunohistochemistry: 1/50-1/100.

Reactivity: Human
Host: Rabbit
Isotype: Ig

Clonality: Polyclonal

**Immunogen:** This antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide

selected from the Center region of human U2AF1.

**Specificity:** This antibody detects U2AF1/U2AF35 (Center).

**Formulation:** PBS with 0.09% (W/V) Sodium Azide as preservative.

State: Purified

State: Liquid purified Ig fraction.

**Concentration:** lot specific

**Purification:** Saturated Ammonium Sulfate (SAS) precipitation followed by dialysis against PBS.

Conjugation: Unconjugated

Storage: Store the antibody undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

**Gene Name:** U2 small nuclear RNA auxiliary factor 1 like 5

**Database Link:** Entrez Gene 102724594 Human

Q01081



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## U2AF1L5 (Center) Rabbit Polyclonal Antibody - AP12453PU-N

**Background:** U2AF1 belongs to the splicing factor SR family. U2 auxiliary factor, comprising a large and a

small subunit, is a non-snRNP protein required for the binding of U2 snRNP to the pre-mRNA branch site. This gene encodes the small subunit which plays a critical role in both

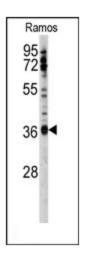
constitutive and enhancer-dependent RNA splicing by directly mediating interactions

between the large subunit and proteins bound to the enhancers.

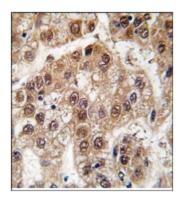
Synonyms: U2AFBP

**Note: Molecular weight:** 27872 Da (35 kDa subunit).

## **Product images:**



Western blot analysis of anti-U2AF1 Antibody (Center) in Ramos cell line lysates (35ug/lane). U2AF1 (arrow) was detected using the purified Pab.



Formalin-fixed and paraffin-embedded human hepatocarcinoma tissue reacted with U2AF1 antibody (Center), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.