本公司提供的电子版本说明书仅供参考，实验请以收到的纸质手册为准。

Anti-Human MAGEL2 Polyclonal Antibody

多克隆抗体

产品货号: PA10379

第三版

描述: Prader-Willi syndrome (PWS) is caused by the loss of expression of imprinted genes in chromosome 15q11-q13 region. Affected individuals exhibit neonatal hypotonia, developmental delay, and childhood-onset obesity. Necdin (NDN), a gene involved in the terminal differentiation of neurons, localizes to this region of the genome and has been implicated as one of the genes responsible for the etiology of PWS. This gene is structurally similar to NDN, is also localized to the PWS chromosomal region, and is paternally imprinted, suggesting a possible role for it in PWS.

抗原: Synthetic peptide of human MAGEL2

配方:

如何使用:加1ml超纯水重溶

稳定性: -20 ° C保存条件下，冻干粉,保质期为五年；液体，保质期为两年。

稀释液:PBS (pH7.4)， 1% BSA

应用:WB 1～5 μg/ml.

特异性: Expressed in placenta, fetal and adult brain. Not detected in heart and small intestine, very low levels in fibroblasts. Not expressed in brain of a Prader-Willi patient.1 Publication