



# NeoNat SCID-SMA Multiplex Real-Time PCR Kit

Semi-quantitative detection of TREC, KREC, SMN1 and SMN2



#### **BACKGROUND**

Thorough research in early detection of Severe Combined Immunodeficiency (SCID) and Spinal Muscular Atrophy (SMA) is vital for prevention of permanent disabilities or mortality of neonates. SCID is a congenital genetic disorder that causes life-threatening problems by supressing the immune system. This immunodeficiency occurs due to the lack of T cells, B cells and NK cells count. SMA is autosomal recessive disorder that causes neuromuscular disorder. This disorder is characterized by progressive degeneration and irreversible loss of lower motor neurons

### **Assay Description**

NeoNat SCID-SMA Multiplex is a Real-Time PCR kit for the screening of SCID by semi-quantitative determination of *TREC & KREC* and SMA by measuring *SMN1* & *SMN2* in newborn's DNA from DBS samples.

## **Assay features**

- Efficient workflow from DNA extraction to the result
- 3.2 mm DBS as input
- Ready to use reaction mix
- DBS controls with defined copy numbers
- DBS calibrators included in the kit
- Ouick turnaround time of ~2 hours
- Compatible with various Real-Time PCR instruments
- 100% clinical sensitivity and specificity



## **Genes Targeted**

Simultaneous amplification of multiple targets

Targets	Detection Channels
TREC (T-cell Receptor Excision Circle)	Texas Red
KREC (Kappa-deleting Recombination Excision Circle)	Cy5.5
SMN1 (Survival Motor Neuron 1)	FAM
SMN2 (Survival Motor Neuron 1)	VIC / HEX
B-Globin (Internal control)	Cy5

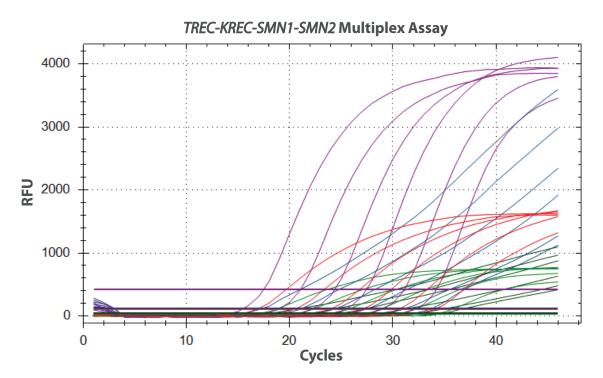
#### **Workflow**

Easy five steps from punching to result



## **Assay result and performance**

High linearity and PCR efficiency



#### **Limit of Detection**

- TREC & KREC 2.2 & 4.1 copies/μl
- *SMN1 & SMN2* 2.3 & 6.8 copies/µl

## **Ordering Information**

Product Code	Format
8100411	96 reactions
8100412	192 reactions
8100413	480 reactions





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