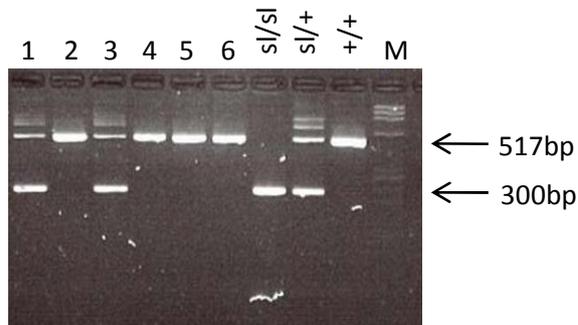


| | |
|-------------|----------------------------|
| Gene symbol | <i>Ednrb</i> ^{sl} |
|-------------|----------------------------|

| primer | Sequence(5'-3') | Size | target gene | accession |
|--------------------------|--|-------|-------------|-------------|
| rSl-del-F rSl-del-R | cctcctggactagaggttcc acgacttagaaagctacact | 517bp | Ednrb | NM_017333.1 |
| rSl-wild-F rSl-wild-R | AGGCATTAATGGGAATGTCC ACTCCAGTCTGATGCGTTCC | 300bp | Ednrb | NM_017333.1 |
| comment | | | | |

| PCR condition | | | | | | | |
|----------------|-----|--|----------|--------|-----------|-------|-----------------|
| Taq polymerase | | BIOTAQ™ HS DNA Polymerase (Bioline Reagents Ltd., UK) | | | | | |
| Thermal cycler | | PC-808(ASTEC) | | | | | |
| PCR buffer | | Ampdirect Plus | | | | | |
| | | first denature | denature | anneal | extension | cycle | final extension |
| PCR | °C | 94 | 94 | 60 | 72 | 35 | 72 |
| | min | 3 | 0.5 | 0.5 | 1 | | 3 |
| comment | | PCR method; Blood was applied to FTA®card (GE Healthcare UK Ltd.,UK) and dried. 1.5mm FTA disc was removed from the bloody-stained region on FTA®card. Untreated sample discs were placed directly in 15 μ L PCR mixture containig 1 × Ampdirect®Plus, 0.2 μ M each primer and 0.4 units of BIOTAQ™ HS DNA Polymerase. | | | | | |

Electrocataphoresis ;4% agarose gel 300V, 30min



| | |
|---------------------------|--|
| Strain harboring mutation | AR- <i>Ednrb</i> ^{sl} /Okkm , LE.AR- <i>Ednrb</i> ^{sl} /Okkm |
|---------------------------|--|