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1 A novel homozygous nonsense mutation in *ITGB4* gene causes epidermolysis bullosa in
2 Mouton Vendéen sheep

3

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9 Junctional Epidermolysis Bullosa (EB) is a severe genetic skin disorder in humans and animals
10 where any mechanical trauma can cause painful blisters. In sheep, this recessive disease
11 affecting limb extremities and mucous membranes causes perinatal death of affected lambs and
12 was previously associated to mutation in *LAMC2* and *ITGB4* genes in German black-headed
13 mutton and Spanish Churra sheep, respectively [1, 2]. Multiple observations of EB cases were
14 recently reported in the French Mouton Vendéen meat sheep breed. Skin biopsies of 6 affected
15 dead lambs and when available, blood samples from the parents (n = 7) and unaffected full-sib
16 lambs (n = 4) were collected for genomic DNA extraction. Based on the hypothesis of a
17 recessive transmission of a deleterious variant, we have performed an Illumina HiSeq paired-
18 end WGS (ENA accession number PRJEB37460; BWA alignment, ≈10X coverage) of 2
19 unrelated EB-affected lambs (supposed homozygous carriers) and 1 unaffected full-sib
20 (supposed heterozygous or non-carrier). The strategy focused on 13 candidate genes already
21 associated with EB or skin fragility syndrome in several species, and mainly belonging to the
22 collagen, laminin, integrin and keratin families. After GATK variant analysis of these 13 genes
23 fitting with the mutation segregation hypothesis, we have identified a novel SNP (OAR11_v4.0,
24 g.54799925G>A) in the exon 23 of the *ITGB4* gene of the integrin family whose variant allele
25 causes a premature stop codon (p.Arg885*). By a specific RFLP assay (DdeI digestion, Figure

26 1), we have determined that all EB-affected lambs were homozygous for this variant allele,
27 their parents were heterozygous and the full-sibs were either heterozygous or non-carrier, fitting
28 well with a recessive mutation segregation (Figure 1). Through a large set of genotyping of
29 renewal ewe lambs from the same annual cohort, the population allele frequency was estimated
30 at 6.3% (Table 1). The same frequency was observed for service rams (5.2%) and adult ewes
31 (6.0%) picked-up randomly in commercial flocks (Table 1). The discovery of a new mutation
32 in ovine *ITGB4* enhances the role of this gene in EB etiology and is an important molecular
33 tool to improve the selection scheme management of the Mouton Vendéen breed to limit the
34 dissemination of this disease.

35 **Acknowledgments**

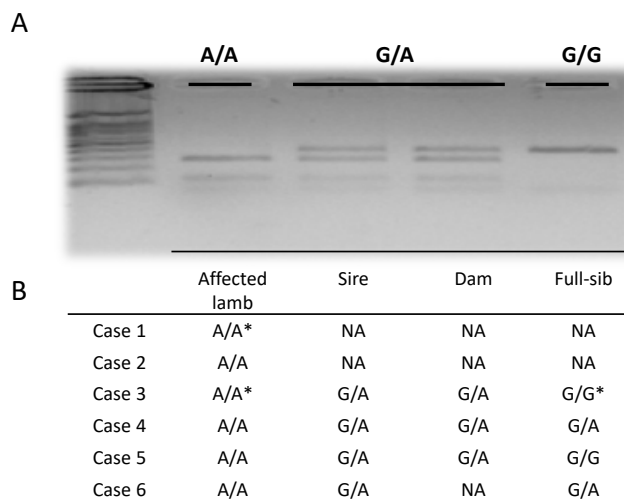
36 We thank Charline Rousseau and Pénélope Fournier from the Mouton Vendéen selection
37 organization, for their precious help in the planning of blood sampling. We are grateful to the
38 breeders who made their animals available for this study. LC was supported by a PhD grant
39 funded by APIS-GENE through the Proligen project.

40 **References**

- 41 1. Mömke et al. (2011) PloS ONE 6, e18943
- 42 2. Suarez-Vega et al. (2015) PloS ONE 10, e0126416

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44 Figure 1. Genotyping of EB cases and their relatives. (A) Specific RFLP results (DdeI
 45 digestion) for OAR11_v4.0, g.54799925G>A mutation in *ITGB4* for homozygous A/A case
 46 n°3, its heterozygous sire and dam (G/A) and non-carrier full-sib (G/G). (B) Genotyping
 47 results of the initial sampling from 6 cases (*whole-genome sequenced animals, NA not
 48 available).



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50 Table 1. Genotyping of g.54799925G>A in *ITGB4* in the Mouton Vendéen population.

	Number of animals	Number of heterozygous carriers	Allele frequency
Renewal ewe lambs	1335	168	6.3%
Adults ewes	1683	203	6.0%
Service rams	1200	126	5.2%

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