

Title: DNA Elements for Constitutive Androstane Receptor- and Pregnane X Receptor-mediated Regulation of Bovine *CYP3A28* Gene

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S7 Table. bPXR-mediated transactivation of *CYP3A28* proximal promoter (PP) and fragment 3 (F3) using rifampicin (RIF).

| Mean luciferase activity* \pm SD | | |
|------------------------------------|----------|-------------------------|
| pGL4.10 | pCI-neo | 100 |
| | bPXR | 140.53 \pm 30.24 |
| | bPXR+RIF | 133.73 \pm 52.57 |
| XREM | pCI-neo | 933.39 \pm 509.69 |
| | bPXR | 1432.97 \pm 842.79 |
| | bPXR+RIF | 2112.60 \pm 972.68 |
| PP | pCI-neo | 3282.56 \pm 3038.26 |
| | bPXR | 4618.43 \pm 3841.19 |
| | bPXR+RIF | 4329.15 \pm 3575.36 |
| PP+F3 | pCI-neo | 26100.24 \pm 21617.61 |
| | bPXR | 30173.12 \pm 22813.48 |
| | bPXR+RIF | 32464.00 \pm 24524.25 |

*C3A cells were transfected with the control reporter pCMV β (150 ng/well), each reporter plasmids or *CYP3A4*-XREM-luc (XREM, 50 ng/well) and either bPXR expression plasmids or pCI-neo empty vector (25 ng/well). After transfection, cells were treated with vehicle (0.1% DMSO) or rifampicin (RIF, 10 μ M) for 24 hours, and reporter activities were measured. Firefly luciferase activities were normalized with β -galactosidase activities. Data are expressed as mean luciferase activities \pm SD (n = 3).