## Different genome-wide transcriptome responses of *Nocardioides simplex* VKM Ac-2033D to phytosterol and cortisone-21-acetate

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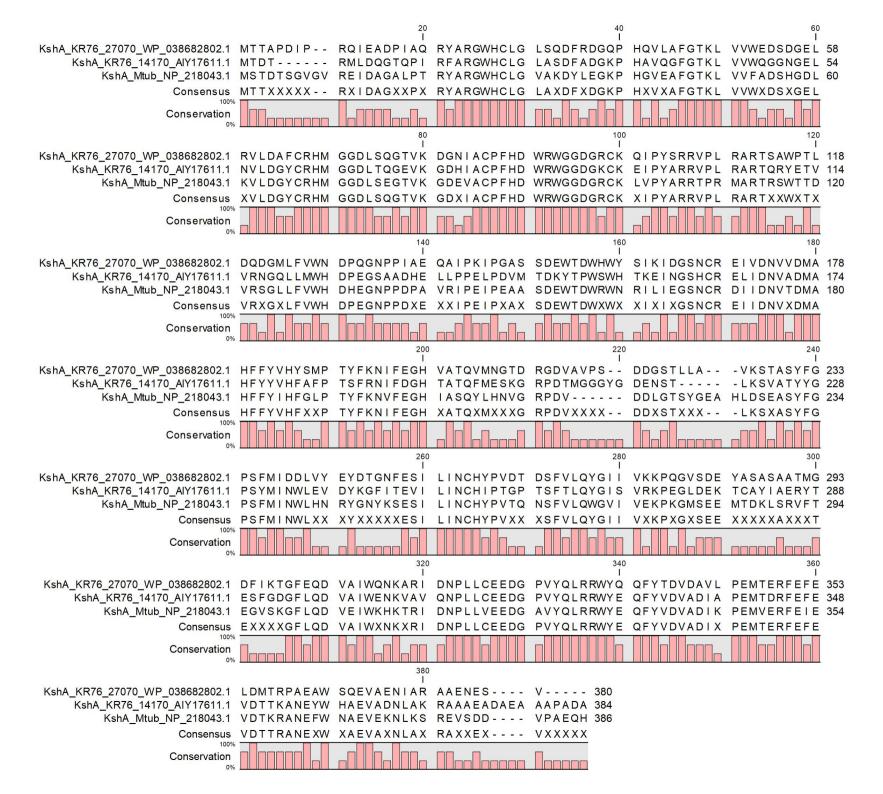
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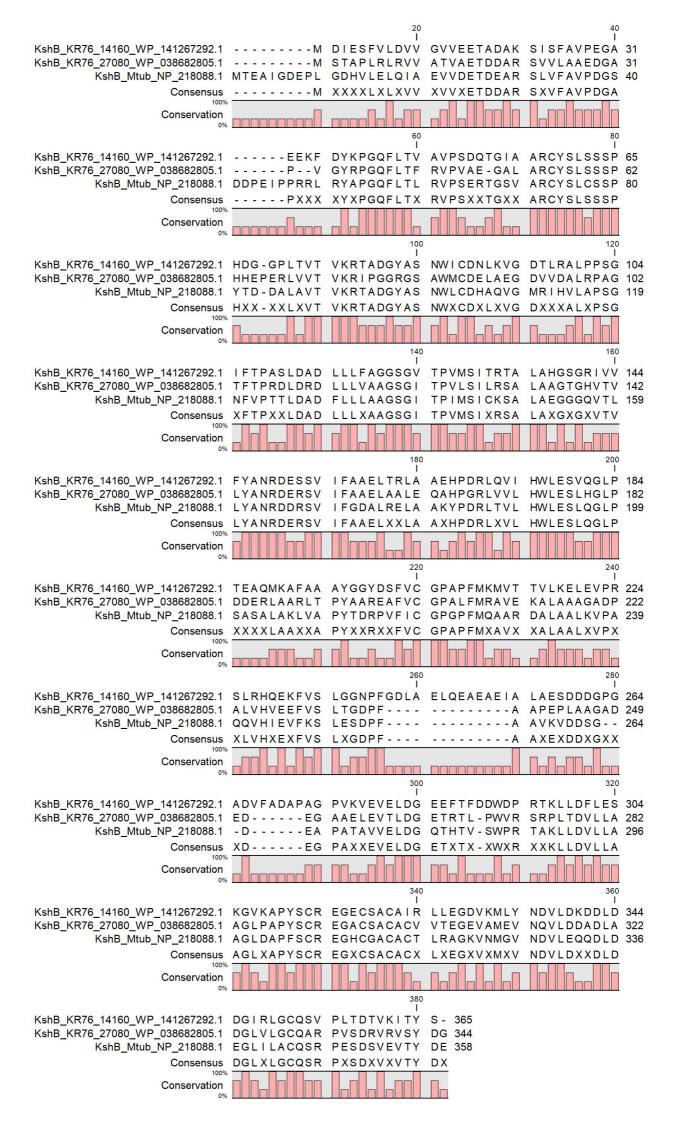
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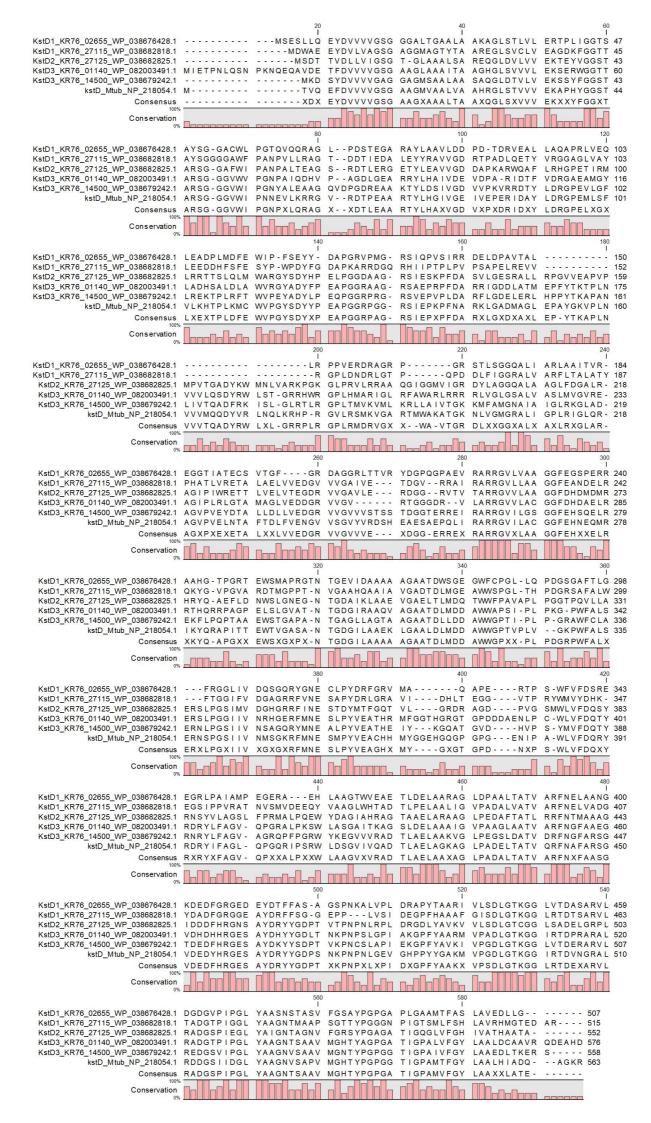
## Supplementary Figure S1. KshA alignment

KshA from *M. tuberculosis* consists of conservative Rieske domain (amino acid residues 24-153), where C67 and C86 are responsible for one Fe<sup>2+</sup> ion binding, and H69 and H89 — for another [71]. The catalytic domain (amino acid residues 154-374) contains conserved amino acid residues V176, D178, H181, H186, Q204, Y232, M238, N240, N257, F301, D304 and W308. In *KR76 27070* of *N. simplex*, N240 is replaced with D240 in this protein.



## Supplementary Figure S2. KshB alignment

There are three domains in KshB containing a number of conservative amino acid residues, namely, a flavin binding domain RxYSL (72-76 in product of Rv3571 *M. tuberculosis*), a NAD binding domain GGIGITP or AGSGITP (135-141 in product of Rv3571 *M. tuberculosis*), and a 2Fe2S-cluster binding domain Cx<sub>4</sub>Cx<sub>2</sub>Cx<sub>29</sub>C (305-344 in product of Rv3571 *M. tuberculosis*) [85]. All three domains were also found in ortholog of KshB in *N. simplex* with two



## Supplementary Figure S3. KstD alignment

The important residues of KstD2 from *R. erythropolis* SQ1 [86] are the following: FAD-binding domain (12-35 in product of Rv3537 of *M. tuberculosis*), consensus GSG(A/G)(A/G)(A/G)X<sub>17</sub>E; Ser325 (318A in product of Rv3537 of *M. tuberculosis* or T in other actinomycetal KSTD enzymes, i.e., small residue at this position); and Thr503 (497 in product of Rv3537 of *M. tuberculosis*). *KR76\_27125* of *N. simplex* contains Ala in position corresponding to 318A in *M. tuberculosis* and 325S in *R. erythropolis*; but FAD-binding domain is different from consensus: GSGT-GX<sub>17</sub>E.