SUPPLEMENTARY INFORMATION

A co-culturing approach enables discovery and biosynthesis of a bioactive indole alkaloid metabolite

Fleurdeliz Maglangit ^{1,2}*, Qing Fang ¹, Kwaku Kyeremeh ³, Jeremy Sternberg ⁴, Rainer Ebel ¹, Hai Deng ¹*

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Figure S2: H-NMR of BE-13793C 1 (DMSO-*d*₆, 298K, 600MHz)



Figure S3: HSQC of BE-13793C $\mathbf{1}$ (CD₃OD, 298K, 600MHz)



Figure S4: HMBC of BE-13793C 1 (DMSO-*d₆*, 298K, 600MHz)



Figure S5: COSY of BE-13793C **1** (CD₃OD, 298K, 600MHz)

Table S1. Deduced functions of the ORFs in BE-13793C **1** (*bec*) biosynthetic gene cluster in *Streptomyces* sp. MA37 showing homology to the rebeccamycin (*reb*) BGC in *Lechevalieria aerocolonigenes* and staurosporine (*sta*) BGC in *Streptomyces* sp. TP-A0274

Gene	Deduced Function	AA	Staurosporine % Protein identity	Rebeccamycin % Protein identity
bec0	FAD-dependent oxidoreductase	508	53%	56%
becD	bis-indole formation	1095	60%	54%
becC	FAD-binding monooxygenase	540	62%	61%
becP	cytochrome P450	417	51%	55%
becR	LuxR family DNA-binding response regulator	221	44%	31%

Lechevalieria aerocolonigenes (rebeccamycin)



Figure S6: Biosynthetic gene cluster comparison of BE-13793C **1** in *Streptomyces* sp. MA37, rebeccamycin in *Lechevalieria aerocolonigenes*, and staurosporine in *Streptomyces* sp. TP-A0274



Figure S7. MIC curve of BE-13793C 1 against HT29 (ATCC HTB-38) colon cells