## Supplemental Table S1. ACR Components at Week 16

|  |  | Deucravacitinib |  |
| :---: | :---: | :---: | :---: |
| Component | Placebo $n=66$ | $\begin{gathered} 6 \mathrm{mg} \text { QD } \\ \mathrm{n}=70 \end{gathered}$ | $\begin{gathered} 12 \mathrm{mg} \text { QD } \\ \mathrm{n}=67 \end{gathered}$ |
| Tender joint count |  |  |  |
| Baseline, mean (SD) | 16.9 (9.8) | 18.1 (10.3) | 19.4 (11.8) |
| Change from baseline, mean (SD) ${ }^{\text {a }}$ | -4.6 (9.7) | -9.3 (9.7) | -12.2 (10.2) |
| \% change from baseline, mean (SD) ${ }^{\text {a }}$ | -25.9 (51.2) | -49.0 (51.9) | -60.4 (35.9) |
| Swollen joint count |  |  |  |
| Baseline, mean (SD) | 10.5 (7.7) | 11.9 (7.0) | 11.3 (9.0) |
| Change from baseline, mean (SD) ${ }^{\text {a }}$ | -4.3 (8.0) | -7.7 (5.8) | -8.5 (9.1) |
| \% change from baseline, mean (SD) ${ }^{\text {a }}$ | -39.8 (63.3) | -68.4 (33.1) | -67.7 (44.1) |
| Patient Global Assessment of disease activity |  |  |  |
| Baseline, mean (SD) | 66.2 (15.8) | 68.2 (16.8) | 63.6 (15.6) |
| Change from baseline, mean (SD) ${ }^{\text {a }}$ | -13.4 (23.5) | -28.7 (23.1) | -27.6 (25.8) |
| \% change from baseline, mean (SD) ${ }^{\text {a }}$ | -18.1 (37.9) | -43.0 (34.0) | -40.4 (49.2) |


| Patient Global Assessment of pain |  |  |  |
| :--- | :---: | :---: | :---: |
| Baseline, mean (SD) | $64.9(18.2)$ | $63.6(21.7)$ | $63.8(15.9)$ |
| Change from baseline, mean (SD) | a | $-13.8(21.5)$ | $-25.3(26.1)$ |
| \% change from baseline, mean (SD) | $-27.5(25.0)$ |  |  |
| Physician Global Assessment of disease activity | $-20.2(35.0)$ | $-36.9(39.1)$ | $-42.0(37.3)$ |
| Baseline, mean (SD) | $63.8(14.8)$ | $68.2(14.7)$ | $63.3(16.1)$ |
| Change from baseline, mean (SD) | $-19.9(21.8)$ | $-33.6(23.0)$ | $-32.2(25.0)$ |
| \% change from baseline, mean (SD) | $-29.2(34.7)$ | $-49.1(32.1)$ | $-47.7(35.2)$ |
| HAQ-DI total score |  |  |  |
| Baseline, mean (SD) | $1.3(0.6)$ | $1.3(0.6)$ | $1.3(0.6)$ |
| Change from baseline, mean (SD) | $-0.1(0.4)$ | $-0.4(0.5)$ | $-0.4(0.6)$ |
| \% change from baseline, mean (SD) | $-10.6(31.2)$ | $-36.0(39.0)$ | $-30.8(52.6)$ |
| High-sensitivity C-reactive protein | $20.4(39.1)$ | $17.6(23.6)$ | $16.5(21.7)$ |
| Baseline, mean (SD) | $-3.3(22.6)$ | $-14.2(24.5)$ | $-10.9(22.8)$ |
| Change from baseline, mean (SD) | $-40.0(54.0)$ | $-9.1(144.0)$ |  |
| \% change from baseline, mean (SD) | $36.0(221.3)$ | -4.0 |  |

All calculations were performed on individual patient data. Change from baseline $=$ Week 16 value - baseline value.
$\%$ change from baseline $=\left((\text { week } 16 \text { value }- \text { baseline value })^{*} 100\right) /$ baseline value.
ACR, American College of Rheumatology; HAQ-DI, Health Assessment Questionnaire-Disability Index; QD, once daily; SD, standard deviation
${ }^{\text {a }}$ Calculated for patients with valid Week 16 data (placebo, $\mathrm{n}=58$; 6 mg QD, $\mathrm{n}=63 ; 12 \mathrm{mg}$ QD, $\mathrm{n}=60$ ).
${ }^{\mathrm{b}}$ Calculated for patients with valid Week 16 data (placebo, $\mathrm{n}=58$; 6 mg QD, $\mathrm{n}=62$; 12 mg QD, $\mathrm{n}=60$ ).
${ }^{c}$ Calculated for patients with valid Week 16 data (placebo, $n=57 ; 6 \mathrm{mg}$ QD, $\mathrm{n}=63 ; 12 \mathrm{mg}$ QD, $\mathrm{n}=59$ ).
${ }^{d}$ Calculated for patients with valid Week 16 data (placebo, $n=58 ; 6 \mathrm{mg}$ QD, $\mathrm{n}=62 ; 12 \mathrm{mg}$ QD, $\mathrm{n}=59$ ).
${ }^{e}$ Calculated for patients with valid Week 16 data (placebo, $n=57 ; 6 \mathrm{mg}$ QD, $\mathrm{n}=62 ; 12 \mathrm{mg}$ QD, $\mathrm{n}=60$ ).

