1	Reactive oxygen species formed in organic lithium-oxygen
2	batteries
3	- Supporting Information -
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17 SI-1 Influence of reduction pulse length

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19 A Pt-ME with an active radius of $r_{\rm T} = 12.5 \,\mu m$ was used. Figure SI-1 shows multi-cycle CVs 20 at the ME after the ME and GDE had been at OCP for 15 min. Prior to that incubation time 21 and recording of the CV, the GDE performed ORR at $E_{GDE} = -0.7$ V for 120 s (Fig. SI-1A), 240 s (Fig. SI-1B) and 360 s (Fig. SI-1-1C). After 120 s a broad peak around $E_{ME} = 0.3$ V is 22 23 noticeable in the CV recorded at the ME. With increasing duration of the reduction pulse at 24 the GDE, two peaks develop at $E_{ME} = 0.15$ V and $E_{ME} = 0.45$ V in the subsequently recorded 25 CV at the ME. It is very clear that the ME accumulates a soluble compound formed during the 26 reduction pulse as increasing reduction pulse length leads to more detected products. Because 27 the ME and the GDE are not in physical contact, this compound must diffuse across the 28 interelectrode distance of $(10 \pm 2) \,\mu\text{m}$.

Surprisingly, the first peak disappears in the following cycles while the second remains. Fig. SI-1D compares the first positive half cycles of Figs. SI-1A to SI-1C. This behavior is not fully understood yet and needs further investigation.

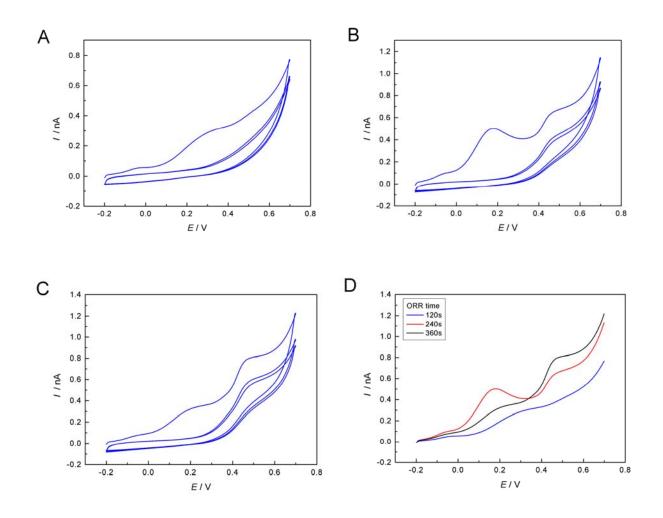


Fig. SI-1 CVs recorded at the ME after the GDE and ME were held at OCP for 15 min. Prior to that incubation time the GDE performed an ORR at $E_{GDE} = -0.7$ V for A) 120 s B) 240 s C) 360 s. D) Comparison of the first

34 positive going half cycle of panels A-C.