IL-1ß Activity Predicts Investment in Present Versus Delayed Outcomes Emily K. Corrigan, Maggie Kleiser, Mary Eliza Baker, Jeffrey Gassen, Gary W. Boehm, Marjorie L. Prokosch, Randi P. Proffitt Leyva,





- Life history theory predicts that individuals living in environments with a relatively high extrinsic mortality risk will favor more immediate reproduction and an enhanced preference for present versus future rewards (Stearns, 1992; Kaplan & Gangestad, 2005).
- Individuals' life history strategies should also be calibrated to internal cues bearing on one's somatic condition, which also plays an important role in determining one's mortality risk (Rickard, Frankenhuis, & Nettle, 2014).
- Interleukin-1 beta (IL-1 β) is a proinflammatory cytokine that plays a key role in regulating local and systemic inflammatory processes after injury and immune challenge (Dinarello, 2011) and is involved in the body's response to both physical and psychosocial stressors (Goshen & Yirmiya, 2009).
- We predicted that IL-1 β may be a key internal marker of somatic condition, playing an important role in decisions about how much to invest in immediate versus delayed outcomes, both at the behavioral and cellular level.

Analysis 1

- We examined the relationship between serum IL-1 β levels and investment in present versus delayed outcomes.
- Participants provided answers to Delaying Gratification Inventory (DGI; Hoerger et al., 2011), Mini-K (Figueredo et al., 2014), Future Orientation scale (FO; Steinberg et al., 2009), and Barratt Impulsiveness Scale (BIS; Patton et al., 1995).
- We predicted that higher levels of serum IL-1 β would be associated with temporal discounting and an overall faster life history strategy.

| | Serum | | | De |
|---------------|-------|--------|---------------|----|
| | IL-1β | Mini-K | BIS-11 | Gr |
| Future | | | | |
| Orientation | 34* | .43*** | 58*** | • |
| Delayed | | | | |
| Gratification | 37** | .48*** | 56*** | |
| | | | | |
| RI2-11 | .42** | 49*** | | |
| Mini K | 21 | | | |

comes

Note. $*p \le .05$, $**p \le .01$, and $***p \le .001$

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Results

• Serum IL-1 β was related to a more present focus, inability to delay gratification, and greater global impulsivity. The relationship between serum IL-1 β and a faster life history strategy as measured by the Mini-K was trending towards, but did not reach significance (p = .17).

We next sought to explore if characteristics known to negatively impact somatic condition would predict serum IL-1^β levels, leading to a preference for present outcomes.

Analysis 2

- Guided by previous literature, we selected five established somatic stressors as predictors of IL-1β activity: Body Mass Index (BMI; Flegal et al., 2013) Childhood Stress (Simmons & Bernstein, 1982) Adult Stress (Prior et al., 2016) Childhood Illnesses (Bozzoli et al., 2008) Adult Illnesses (Klevens et al., 2007)
- Individually, none of these predictors were significantly correlated with serum IL-1 β levels (*ps* > .21).
- The allostatic load literature, however, describes the cumulative effect of environmental stressors on the body (e.g., see Schulkin, 2004). With this in mind, we computed a summative somatic damage composite using Z scores of each variable listed above so that a higher score would represent greater somatic stress.

Results

We found that our cumulative allostatic load variable predicted investment in present over future outcomes, mediated through increased serum levels of IL-1 β .



Direct Effects: $\beta s < .02$ (.06) Indirect Effects: $\beta s > .02 (.01)^*$

Analysis 3

- We next looked to explore if the relationship between IL-1 β would remain after controlling for these sources of somatic stress.
- After controlling for all of these variables, the relationship between serum IL-1 β and all measures of preference for investment in present outcomes (DGI, FO, & BIS) holds (ps < .03).



Additional Preliminary Analyses

We have preliminary data suggesting that factors harmful to somatic condition individually predict elevated release of IL-1β by peripheral blood-derived mononuclear cells (PBMCs) in vitro. Please ask the presenter for additional information.



- serum IL-1 β .
- rise

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Discussion and Future Directions Serum IL-1β predicts investment in present over delayed outcomes. Consistent with the prediction that IL-1 β is a marker of one's somatic condition, factors known to negatively impact bodily health together predict levels of

The relationship between serum IL-1 β and investment in the present remains after controlling for antecedents to somatic damage. Previous literature has suggested that the role of internal and external factors in determining IL-1β-related outcomes might be moderated by variants of the $IL1\beta$, which warrants further investigation (Baune et al., 2010).

It appears that the ability for elevated serum IL-1 β to promote present temporal focus exhibits a path independence, such that the primary effect of IL-1 β on preference for present outcomes is not sensitive to the factors which determine its

Selected References