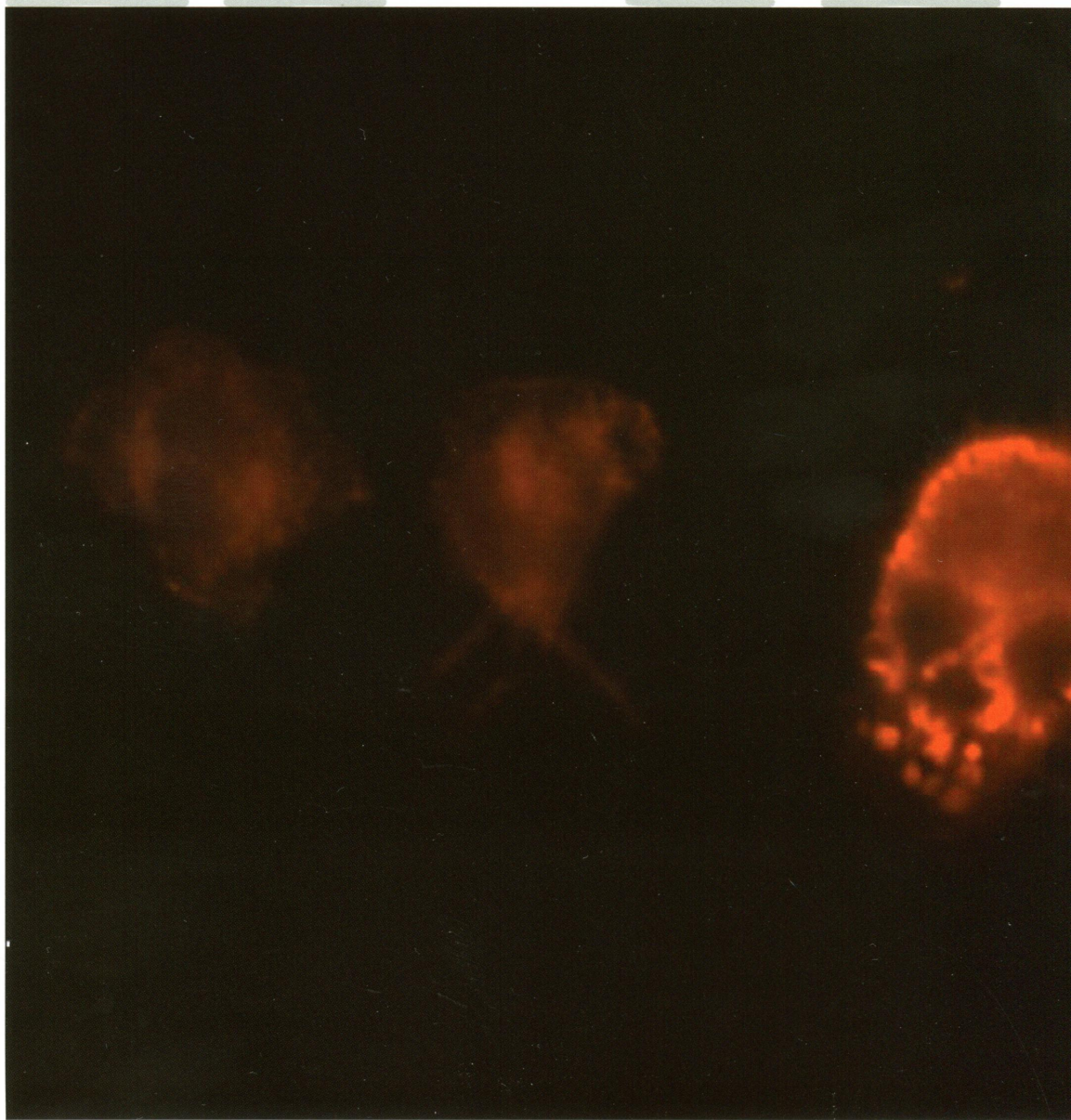


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# IAI

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**AUTHOR CORRECTIONS****Correction for Rao et al., Staphylococcal Enterotoxin B-Induced MicroRNA-155 Targets SOCS1 To Promote Acute Inflammatory Lung Injury**

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**Correction for Jensen et al., Timing of Toll-Like Receptor 9 Agonist Administration in Pneumococcal Vaccination Impacts Both Humoral and Cellular Immune Responses as Well as Nasopharyngeal Colonization in Mice**

Katrine M. Jensen, Jesper Melchjorsen,  
Frederik Dagnaes-Hansen, Uffe B. S.  
Sørensen, Rune R. Laursen, Lars  
Østergaard, Ole S. Søgaard, Martin  
Tolstrup

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*Cover photograph* (Copyright © 2014, American Society for Microbiology. All Rights Reserved.): Neutrophil migration to sites of infection is a critical component of the immune response. Migration is tightly regulated by numerous mechanisms, including specific timing of chemoattractant production and proapoptotic signals that limit the life spans of recruited neutrophils. Dysregulation of this response contributes to damaging inflammation and pathology, as observed in mice coinfecting with the helminth parasite *Heligmosomoides polygyrus* and the bacterial pathogen *Salmonella enterica* serovar Typhimurium. Image shows fluorescent actin staining of migrating neutrophils stimulated with fMLP, including a neutrophil that appears to be undergoing cell death (far right). (Photo courtesy of Rose Szabady, University of Massachusetts Medical School, Worcester, MA.) (For an article in this issue dealing with coinfection by a helminth and *S. Typhimurium*, see page 3855.)