# **AGU: Geophysical Research Letters**

# Keywords

energetic electron precipitation

plasmaspheric hiss

NO production

ozone loss

#### **Index Terms**

Ionosphere: Particle precipitation

Ionosphere: Wave/particle interactions

Atmospheric Composition and Structure: Middle atmosphere: composition and chemistry

Magnetospheric Physics: Energetic particles: precipitating

Radio Science: Radio wave propagation

#### **Abstract**

# Radiation belt electron precipitation due to geomagnetic storms: Significance to middle atmosphere ozone chemistry

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Geomagnetic storms triggered by coronal mass ejections and high-speed solar wind streams can lead to enhanced losses of energetic electrons from the radiation belts into the atmosphere, both during the storm itself and also through the poststorm relaxation of enhanced radiation belt fluxes. In this study we have analyzed the impact of electron precipitation on atmospheric chemistry (30-90 km altitudes) as a result of a single geomagnetic storm. The study conditions were chosen such that there was no influence of solar proton precipitation, and thus we were able to determine the storm-induced outer radiation belt electron precipitation fluxes. We use ground-based subionospheric radio wave observations to infer the electron precipitation fluxes at L = 3.2 during a geomagnetic disturbance which occurred in September 2005. Through application of the Sodankylä Ion and Neutral Chemistry model, we examine the significance of this particular period of electron precipitation to neutral atmospheric chemistry. Building on an earlier study, we refine the quantification of the electron precipitation flux into the atmosphere by using a time-varying energy spectrum determined from the DEMETER satellite. We show that the large increases in odd nitrogen (NO<sub>x</sub>) and odd hydrogen (HO<sub>x</sub>) caused by the electron precipitation do not lead to significant in situ ozone depletion in September in the Northern Hemisphere. However, had the same precipitation been deposited into the polar winter atmosphere, it would have led to >20% in situ decreases in O<sub>3</sub> at 65-80 km altitudes through catalytic HO<sub>x</sub> cycles, with possible additional stratospheric O<sub>3</sub> depletion from descending NO<sub>x</sub> beyond the model simulation period.

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