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Background: Wiskott-
Aldrich syndrome

(WAS) is an X-linked
immunodeficiency,

characterized by

microthrombocytopenia,

eczema and recurrent

infections. More than

441 patient mutations

have been described all

over the world, mainly

based on Caucasian and

Japanese people. There

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have been few reported cases involving Chinese WAS patients. Objective: We i

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