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Reiter Syndrome

National Cancer Institute

Source

National Cancer Institute. *Reiter Syndrome*. NCI Thesaurus. Code C34975.

A rare, reactive inflammatory disorder seen following bacterial infection. It predominantly affects males aged 20-40. Individuals with HLA-B27 antigen are estimated to have a 50 % increased risk. The disorder is characterized by arthritis, conjunctivitis, uveitis, iritis and ulceration of the oral cavity, genitals and volar surfaces of the hands and feet. The clinical course is self-limited with resolution of clinical disease usually within six months of onset.