Scott craniodigital syndrome is a condition that has only been found in two families. The manifestations include unusual head shape, growth and developmental delay, and mild webbing between the fingers and toes (syndactyly) Scott Craniodigital Syndrome With Mental Retardation is an extremely rare inherited disorder that is fully expressed in males only. However, females who carry a single copy of the disease gene (heterozygotes) may exhibit some of the symptoms associated with the disorder. The disorder has been reported in two separate families (kindreds) in the medical literature. Most of the symptoms are apparent at birth.