Xeroderma pigmentosum study tests artificial antisense oligonucleotides as therapeutic

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Locations of the ERCC4/XPF intron variants identified in the Japanese XP-F cases. (A) Estimated structures of pre-mRNA products resulting from the ERCC4/XPF intron variants. Cryptic intron fragments identified in the patients’ mRNA are shown in blue lines. (B) cDNA sequences of the ERCC4/XPF exons 1 to 2 boundary in 48BR (normal) and XP43NG (XP-F). The 5’ cryptic intron 1 fragment is shown in blue letters. (C) A cDNA sequence of the ERCC4/XPF
Genetic researchers at Nagoya University, Japan, have delved into the genetic underpinning of a rare skin condition affecting children that is unusually common in Japan.

In the paper, "Deep intronic founder mutations identified in the ERCC4/XPF gene are potential therapeutic targets for a high-frequency form of xeroderma pigmentosum," published in *PNAS*, the team finds a potential therapeutic target for the disorder with artificial antisense oligonucleotides.

Xeroderma pigmentosum (XP) is a rare genetic disorder that can cause heightened sensitivity to sunlight and an increased risk of skin tumors due to a deficiency in the DNA repair system responsible for processing sunlight-induced photolesions.
XP patients commonly experience severe skin problems, including photosensitivity, dry skin, pigmentation abnormalities, and a heightened risk of skin cancer. Some cases may also exhibit neurological symptoms.

The clinical manifestations of XP vary depending on the affected genes and types of mutations. While every form of XP exhibits some of the pathologies, XP-F patients can have most or all of the disease manifestations at once.

The worldwide prevalence of XP is rare at approximately 3 in a million. In Japan, the rates are much higher, 1 out of 22,000. Of XP cases, XP-F incidents are around 1% globally and 4% in Japan, making XP-F 66 times more common in Japan than the global average.

Partly because XP is such a rare disorder, it remains under-studied, and treatment options are limited. There is a need for improved understanding, diagnosis, and potential therapeutic targets for XP, especially for the rarest variants like XP-F.

The study was conducted on a Japanese XP cohort (cohort size identified only as "largest") and identified 17 XP-F cases, all of which had one of two ERCC4/XPF gene variants. The first variant is a Japanese founder mutation that accounts for approximately 10% of all Japanese XP cases and causes incorrect pre-mRNA splicing. The second mutation induces alternative polyadenylation.

Both mutations result in reduced ERCC4/XPF gene expression, leading to a significant reduction in XPF protein expression and DNA repair deficiency in patients' cells.

The team attempted to correct the abnormal splicing events in patient cell samples through artificial antisense oligonucleotides (ASOs). ASOs
work by binding to specific mRNA sequences. They can induce degradation, modulation of splicing, prevention of translation, or in this case, interfere with alternative mRNA processing events, which can inhibit or upregulate the production of downstream proteins.

After treatment with ASOs, the XPF protein expression was recovered to the normal level, indicating that the ASOs efficiently restored the mRNA expression. The study demonstrates that antisense oligonucleotides specifically designed for these mutations can restore XPF protein expression and DNA repair capacity in cells from XP-F patients.

While ASOs are currently being developed and tested for a variety of genetically based diseases, the application in the current study illustrates how genetic studies can lead the way in finding therapeutic targets for even the rarest of diseases.


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