

Publication

Antisense Oligonucleotide-Mediated Transcript Knockdown in Zebrafish

JournalArticle (Originalarbeit in einer wissenschaftlichen Zeitschrift)**ID** 4519618**Author(s)** Pauli, Andrea; Montague, Tessa G.; Lennox, Kim A.; Behlke, Mark A.; Schier, Alexander F.**Author(s) at UniBasel** [Schier, Alexander](#) ;**Year** 2015**Title** Antisense Oligonucleotide-Mediated Transcript Knockdown in Zebrafish**Journal** PloS one**Volume** 10**Number** 10**Pages / Article-Number** e0139504**Mesh terms** Animals; Embryonic Development, genetics; Feasibility Studies; Female; Gene Knock-down Techniques; Male; Morpholinos, genetics, pharmacology; Oligonucleotides, Antisense, genetics, pharmacology; RNA, Long Noncoding, genetics; RNA, Messenger, antagonists & inhibitors, genetics; Transcription, Genetic; Zebrafish, embryology, genetics; Zebrafish Proteins, genetics; Zygote

Antisense oligonucleotides (ASOs) are synthetic, single-strand RNA-DNA hybrids that induce catalytic degradation of complementary cellular RNAs via RNase H. ASOs are widely used as gene knockdown reagents in tissue culture and in *Xenopus* and mouse model systems. To test their effectiveness in zebrafish, we targeted 20 developmental genes and compared the morphological changes with mutant and morpholino (MO)-induced phenotypes. ASO-mediated transcript knockdown reproduced the published loss-of-function phenotypes for *oep*, *chordin*, *dnd*, *ctnnb2*, *bmp7a*, *alk8*, *smad2* and *smad5* in a dosage-sensitive manner. ASOs knocked down both maternal and zygotic transcripts, as well as the long noncoding RNA (lncRNA) MALAT1. ASOs were only effective within a narrow concentration range and were toxic at higher concentrations. Despite this drawback, quantitation of knockdown efficiency and the ability to degrade lncRNAs make ASOs a useful knockdown reagent in zebrafish.

Publisher PUBLIC LIBRARY SCIENCE**ISSN/ISBN** 1932-6203**URL** <https://journals.plos.org/plosone/article?id=10.1371/journal.pone.0139504>**edoc-URL** <https://edoc.unibas.ch/74125/>**Full Text on edoc** No;**Digital Object Identifier DOI** 10.1371/journal.pone.0139504**PubMed ID** <http://www.ncbi.nlm.nih.gov/pubmed/26436892>**ISI-Number** WOS:000362499200030**Document type (ISI)** Journal Article