

The Foundation for Angelman Syndrome Therapeutics (FAST) is so very grateful to the Morgan Family, our largest private donors, and to all of their friends and supporters for raising and donating to FAST the sum of \$24,000.00 for Angelman Syndrome research. Less than three years ago, it was discovered that a therapeutic for Angelman Syndrome is no longer just a hope and dream, but a reality; and that children now suffering from this disorder may one day have the ability to speak, live free of seizures and have the chance to lead an independent life. The only challenge we face today is in raising the money necessary to fund the research that will lead us to this therapeutic. Largely because of the Morgan family, FAST is able to begin funding this vital research now and is currently accepting applications from scientists focused on curing this devastating disorder.

The Morgan family has also worked tirelessly to increase awareness of Angelman Syndrome and the FAST organization could not be more appreciative. To learn more about Angelman Syndrome, please visit www.CureAngelman.org and to read more about all the hard work and effort the Morgan family has undertaken, please visit http://www.cureangelman.org/newsletter/NewsLetter_March2010.pdf?page=6

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