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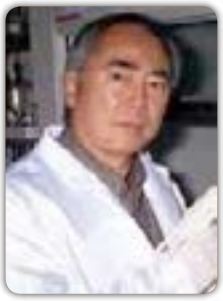

Douglas N Ishii

Colorado State University, USA

Why clinical trials that reduce glucose levels fail to prevent complications in diabetic patients: Tests support an alternative hypothesis for pathogenesis

Statement of Problem: Meta-analysis of outcomes on 34,533 Type 2 diabetic patients shows that intensive lowering of glucose levels does not prevent neuropathy, retinopathy, nephropathy, cardiovascular death, nor excess mortality. Nor does lowering of glucose levels prevent complications in approximately 40% of type 1 patients. Exposing patients to adverse effects from unbeneficial drugs is unjustified, yet remains standard therapy. The development of meaningful novel treatments awaits an alternative hypothesis for pathogenesis of diabetic complications.

Methodology & Theoretical Orientation: Insulin and insulin-like growth factors (IGFs) are neurotrophic factors. The inter-related hypotheses were developed that diminished insulin and IGF activities is the dominant cause of neurological complications, and that replacement of such activities should ameliorate diabetic complications irrespective of unabated hyperglycemia. These hypotheses were tested by infusing IGFs, insulin, or their combination into diabetic rats to determine whether neuropathy is alleviated under conditions in which hyperglycemia remains unabated.

Conclusion & Significance: IGF mRNA levels are reduced in peripheral nerves, brain and spinal cord in diabetes. Replacement IGF infusion prevented impaired sensory and motor nerve regeneration, hyperalgesia, abnormal ultrastructure in autonomic axons, loss of epidermal nerve fiber density, and poor gastric wound healing despite undiminished hyperglycemia. Tiny doses of insulin and/or IGF were infused into diabetic rat brains under conditions that did not reduce hyperglycemia. A decrease in total mRNA, protein, and DNA levels was associated with brain atrophy and impaired learning/memory in diabetic rats. Insulin and IGF i.c.v. infusion prevented all such disturbances despite unabated hyperglycemia. Insulin and IGFs are master switches controlling the levels of hundreds of proteins in brain; loss of protein regulation, not hyperglycemia, is proposed as the most likely pathogenic cause for diabetic complications. Governments should manufacture clinical grade IGF (off- patent). Clinical trials are urgently needed to test insulin/IGF therapy.

Recent Publications

1. Boussageon R, Bejan Angoulvant T, Saadatian Elahi M, Lafont S, Bergeonneau C, Kassai B, Erpeldinger S, Wright J M, Gueyffier F and Cornu C (2011) Effect of intensive glucose lowering treatment on all-cause mortality, cardiovascular death, and microvascular events in type 2 diabetes: meta-analysis of randomized controlled trials. British Medical Journal 343:d4169.
2. Serbedzija P, Madl J E and Ishii D N (2009) Insulin and IGF-I prevent brain atrophy and DNA loss in diabetes. Brain Research 1303:179-94.
3. Lupien S B, Bluhm E J and Ishii D N (2006) Effect of IGF-I on DNA, RNA, and protein loss associated with brain atrophy and impaired learning in diabetic rats. Neurobiology of Disease 21(3):487-495.
4. Ishii D N and Lupien S B (2003) Insulin-like growth factor replacement therapy for diabetic neuropathy: experimental basis. Experimental Diabetes Research 4(4):257-269.
5. Pulford B E and Ishii D N (2001) Uptake of circulating insulin-like growth factors (IGFS) into cerebrospinal fluid appears to be independent of the IGF receptors as well as IGF binding proteins. Endocrinology 142(1):213-220.

Biography

Douglas N Ishii received his BA in Biochemistry from University of California, Berkeley and PhD in Pharmacology from Stanford University Medical School and conducted Postdoctoral work in Neurobiology at Stanford. He became Assistant then Associate Professor of Pharmacology at Columbia University, New York City. He is a Professor of Biomedical Sciences at Colorado State University. He served on various scientific study sections for National Science Foundation, National Institutes of Health and The Juvenile Diabetes International Foundation. Press coverage on his laboratory's research on pathogenesis of diabetic neurological complications, and cause of brain atrophy in Alzheimer's disease, includes articles in Der Spiegel, Hong Kong Standard, NY Times, LA Times, Denver Post, Chicago Tribune, ABC News, Forbes News, USA Today, National Public Radio, and elsewhere. Nineteen patents were awarded based on this research.

Douglas.Ishii@colostate.edu

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