

CALIFORNIA STATE SCIENCE FAIR 2014 PROJECT SUMMARY

Name(s)	Project Number
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	34864
Project Title	
Break the AGE Barrier! Inhibit Advanced Glycation End products to	
Combat Atherosclerosis, Cancer and Diabetic Disorders	
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Abstract	
Objectives/Goals	of approvalarosis
Advanced Glycation End-products (AGEs) have been identified as the root cause of a herosclerosis, cancer, Alzheimer's and diabetic neuropathy, retinopathy and nephropathy. AGEs after the structures and	
functions of vital proteins and lipids. The objective of this project is to identify conditions for AGE	
formation, identify solutions for inhibiting AGE formation and to by by the apol	secial breakthrough cure
for many life-threatening diseases. The objective is to test and compare the inh additives containing phenols, anthocyanins, chelators and GLUT1 monopolizer	bitory effects of 9 natural
additives containing phenols, anthocyanins, chelators and GLU I'l manopolizers on AGE formation. Methods/Materials	
AGE formation due to non-enzymatic endogenous (endo) and exogenous (exo)	protein glycation (PG) and
lipid peroxidation (LPO) was simulated with 4 in vitro tests (5 trials ineach test) with and without equal	
concentrations of the 9 additives. PG tests used ribose, maltose, fructose, lactose and glucose sugars. Endo-PG tests were conducted at 37 deg. C with collagen and sugars, Exo-PG tests were conducted at	
Endo-PG tests were conducted at 37 deg. C with collagen and sugars Exo-PG tests were conducted at	
100, 80 and 60 deg. C with lysine and sugars. A home made smartprione spectrophotometer (constructed with a DVD diffraction grating and injection molded plastic parts) and Beer Lambert's Law were used to compute solution absorbance. LPO tests used saff ower oil (morpounsaturated fatty acid-MUFA) and olive oil (polyunsaturated fatty acid-PUFA). Fenton's Reagent created reactive oxygen species in the oils.	
compute solution absorbance. LPO tests used safflower sil (propounsaturated fatty acid-MUFA) and olive	
oil (polyunsaturated fatty acid-PUFA). Fenton's Reagent created reactive oxygen species in the oils.	
I I lodometric titration was used to compute their peroxide values (PV). Changes 1	n absorbance values,
reaction rates, PV and IC(50) values (versus the control) were used to rank the additives.	
Results Ribose produced the most AGE (\$ 2% por the malase) Ascorbic acid inhibited PG the best (68%	
endo 62.6% exo) followed by bluberty PUFA produced 73.9% more AGEs	than MUFAs Resveratrol
Ribose produced the most AGEs (55.2% more than malrose). Ascorbic acid inhibited PG the best (68% endo, 62.6% exo), followed by oluberty. PUFAs produced 73.9% more AGEs than MUFAs. Resveratrol inhibited LPO the best (58.5% endo, 64.2% exo), followed by carnosine and tocopherol. However, the anomalies, niacinamide for PG and black arrant for LPO, didn't inhibit or increased AGE formation in	
anomatics, machaniae for rol and blackes range in the of the reased AOE formation in	
some cases. AGE formation increased at higher temperatures. PG increased and	LPO decreased with
increasing alkalinity. Conclusions/Discussion	
This project has identified a potential break hrough treatment for AGE inhibition	on to combat cancer.
atherosclerosis, Alzheimer's and other diabetic disorders. The AGE inhibitory p	properties of all the
additives, except blackcurrent and macinamide, have been identified.	-
Summary Statement	
My project aims to identify several inexpensive AGE inhibitors, discover sever	al critical factors for
controlling AGE formation, and provide breakthrough remedies for several life	-threatening diseases.
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Help Received	
My science teacher, Mrs. Makhijani provided valuable guidance. My parents purchased all the materials	
and provided encouragement.	