



# Maternal supplementation with n-3 long chain polyunsaturated fatty acids during perinatal period alleviates the metabolic syndrome disturbances in adult hamster pups fed a high-fat diet after weaning

Submitted by [claire.leroy](#) on Wed, 04/29/2015 - 14:29

Titre	Maternal supplementation with n-3 long chain polyunsaturated fatty acids during perinatal period alleviates the metabolic syndrome disturbances in adult hamster pups fed a high-fat diet after weaning
Type de publication	Article de revue
Auteur	Kasbi-Chadli, Fatima [1], Boquien, Clair-Yves [2], Simard, Gilles [3], Ulmann, Lionel [4], Mimouni, Virginie [5], Leray, Véronique [6], Meynier, Anne [7], Ferchaud-Roucher, Véronique [8], Champ, Martine [9], Nguyen, Patrick [10], Ouguerram, Khadija [11]
Editeur	Elsevier
Type	Article scientifique dans une revue à comité de lecture
Année	2014
Langue	Anglais
Date	Juil. 2014
Numéro	7
Pagination	726-733
Volume	25
Titre de la revue	The Journal of Nutritional Biochemistry
ISSN	0955-2863
Mots-clés	hamster [12], Metabolic syndrome [13], mitochondrial function [14], n-3 LC-PUFA [15], perinatal nutrition [16]

Perinatal nutrition is thought to affect the long-term risk of the adult to develop metabolic syndrome. We hypothesized that maternal supplementation with eicosapentaenoic acid and docosahexaenoic acid during pregnancy and lactation would protect offspring fed a high-fat diet from developing metabolic disturbances. Thus, two groups of female hamsters were fed a low-fat control diet, either alone (LC) or enriched with n-3 long chain polyunsaturated fatty acids (LC-PUFA) (LO), through the gestational and lactation periods. After weaning, male pups were randomized to separate groups that received either a control low-fat diet (LC) or a high-fat diet (HC) for 16 weeks. Four groups of pups were defined (LC-LC, LC-HC, LO-LC and LO-HC), based on the combinations of maternal and weaned diets. Maternal n-3 LC-PUFA supplementation was associated with reduced levels of basal plasma glucose, hepatic triglycerides secretion and postprandial lipemia in the LO-HC group compared to the LC-HC group. Respiratory parameters were not affected by maternal supplementation. In contrast, n-3 LC-PUFA supplementation significantly enhanced the activities of citrate synthase, isocitrate dehydrogenase and  $\alpha$ -ketoglutarate dehydrogenase compared to the offspring of unsupplemented mothers. Sterol regulatory element binding protein-1c, diacylglycerol O-acyltransferase 2, fatty acid synthase, stearoyl CoA desaturase 1 and tumor necrosis factor  $\alpha$  expression levels were not affected by n-3 LC-PUFA supplementation. These results provide evidence for a beneficial effect of n-3 LC-PUFA maternal supplementation in hamsters on the subsequent risk of metabolic syndrome. Underlying mechanisms may include improved lipid metabolism and activation of the mitochondrial oxidative pathway.

Résumé en anglais

URL de la notice <http://okina.univ-angers.fr/publications/ua10542> [17]

DOI [10.1016/j.jnutbio.2014.03.003](https://doi.org/10.1016/j.jnutbio.2014.03.003) [18]

Lien vers le document <http://linkinghub.elsevier.com/retrieve/pii/S0955286314000576> [19]

Titre abrégé J Nutr Biochem

---

## Liens

- [1] [http://okina.univ-angers.fr/publications?f\[author\]=18612](http://okina.univ-angers.fr/publications?f[author]=18612)
- [2] [http://okina.univ-angers.fr/publications?f\[author\]=18613](http://okina.univ-angers.fr/publications?f[author]=18613)
- [3] <http://okina.univ-angers.fr/gi.simard/publications>
- [4] [http://okina.univ-angers.fr/publications?f\[author\]=18614](http://okina.univ-angers.fr/publications?f[author]=18614)
- [5] [http://okina.univ-angers.fr/publications?f\[author\]=18615](http://okina.univ-angers.fr/publications?f[author]=18615)
- [6] [http://okina.univ-angers.fr/publications?f\[author\]=18616](http://okina.univ-angers.fr/publications?f[author]=18616)
- [7] [http://okina.univ-angers.fr/publications?f\[author\]=18617](http://okina.univ-angers.fr/publications?f[author]=18617)
- [8] [http://okina.univ-angers.fr/publications?f\[author\]=18618](http://okina.univ-angers.fr/publications?f[author]=18618)
- [9] [http://okina.univ-angers.fr/publications?f\[author\]=18619](http://okina.univ-angers.fr/publications?f[author]=18619)
- [10] [http://okina.univ-angers.fr/publications?f\[author\]=18620](http://okina.univ-angers.fr/publications?f[author]=18620)
- [11] [http://okina.univ-angers.fr/publications?f\[author\]=18621](http://okina.univ-angers.fr/publications?f[author]=18621)
- [12] [http://okina.univ-angers.fr/publications?f\[keyword\]=16787](http://okina.univ-angers.fr/publications?f[keyword]=16787)
- [13] [http://okina.univ-angers.fr/publications?f\[keyword\]=6480](http://okina.univ-angers.fr/publications?f[keyword]=6480)
- [14] [http://okina.univ-angers.fr/publications?f\[keyword\]=16786](http://okina.univ-angers.fr/publications?f[keyword]=16786)
- [15] [http://okina.univ-angers.fr/publications?f\[keyword\]=16785](http://okina.univ-angers.fr/publications?f[keyword]=16785)
- [16] [http://okina.univ-angers.fr/publications?f\[keyword\]=16784](http://okina.univ-angers.fr/publications?f[keyword]=16784)
- [17] <http://okina.univ-angers.fr/publications/ua10542>
- [18] [http://dx.doi.org/10.1016/j.jnutbio.2014.03.003](https://dx.doi.org/10.1016/j.jnutbio.2014.03.003)
- [19] <http://linkinghub.elsevier.com/retrieve/pii/S0955286314000576>