1. Data Abstraction Tool

Article information 1. What is the name of this study? (e.g. DART, Physician's Health Study) Omit if name of study is not provided. DART Physician's Health Study Maastricht Essential Fatty Acid Birth (MEFAB) Cohort The DIAMOND Study (DHA Intake And Measurement Of Neural Development) data using DIAMOND study POSGRAD NUHFAL Infant Fish Oil Supplementation Study (IFOS) Danish National Birth Cohort Groningen LCPUFA study DINO (Docosahexaenoic acid for the Improvement in Neurodevelopmental Outcome) DOMINO INFAT BeMIM (Belgrade-Munch Infant Milk Trial) GINI The Docosahexaenoic Acid to Optimise Mother Infant Outcome (DOMInO) Childhood Asthma Prevention Study Salmon in Pregnancy Study (SiPS) Other study names 2. Study Design Trial: Randomized Parallel (Omega-3 vs. Control; Omega-3 XX vs. X) Trial: Randomized Cross-over

•	Trial: Randomized Factorial Design
•	Observational: Prospective, longitudinal, comparative study
•	Observational: Nested Case Control
Oth	ner
	ar Response
3. F	unding source
	Industry funded
	Government funded
	Some authors employed by industry (companies that make the supplements)
	Funding or affiliations not reported
	Mulitple foundations and Societies
	None of the authors had any personal or financial conflicts of interest
	Trade group funded
	March of Dimes
	Manufacturer supplied product
	Some authors serve on scientific advisory boards for corporations
4. C	ner funding source country in which study conducted (where subjects live) Note that this might be different than the countries wher authors are based. Select "NR (not reported)" if it's truly unclear.
	US
	Canada
	Denmark
	Finland
	Germany
	Greece
	Italy

	Japan
	Netherlands
	Norway
	Sweden
	UK
	NR (not reported)
	Mexico
	Australia
	Taiwan
	Spain
	Hungary
	Turkey
	Bangladesh
	Thailand
	Serbia
	Multie-center study in hospitals of 11 countries in Europe
Oth	er audy Population
5. 31	Healthy infants or children
	Preterm infants
	rieleiii iiiaiis
la constitución de la constituci	Healthy pregnant women
	Postpartum women
	Breast-feeding women
	Low birth weight infants

Standard Deviation	Standard Deviation	Standard Deviation	Standard Deviation
1			
Mean age (Pregnant)	Mean age (Lactating mothers)	Mean age (Mothers)	Mean age (Infants)
2. Age at baseline	Moan ago /Loctoting	Moan ago (Mathara)	Moan ago (Infants)
up)	up)	up)	up)
completers (largest follow-	completers (largest follow-	completers (largest follow-	completers (largest follow-
# of	# of	# of	# of
_	_		_
# of withdrawals	# of withdrawals	# of withdrawals	# of withdrawals
omized	omized	omized	omized
# Enrolled/Rand	# Enrolled/Rand	# Enrolled/Rand	# Enrolled/Rand
	-	_	-
1. Sample size Pregnant	Lactating mothers	Mothers	Infants
II. Participant Info	ormation		
4	F		
7. Exclusion criteria			
)		
	V		
6. Inclusion criteria as de	fined in study		
children with family his	story of allergy		
П		.ge. o. do toloping double	•
Pregnant women who	se unhorn children were at h	nigh risk of developing asthma	1

4			
Age range (Pregnant)	Age range (Lactating mothers)	Age range (Mothers)	Age range (infants)
4			
3. Race/Ethnicity (Mother	r)		
White/European (spe	cify %)		
Black/African America	an/etc. (specify %)		
Asian (specify %)			
Hispanic (specify %)			
Inuit/Eskimo (specify	%)		
NR (specify %)			
varied by study site a	nd experimental group (speci	fy %)	
Minority (specify %)			
Other			
non black			
Puerto Rican/Latino			
Native Hawaiian or ot	her pacific ethnicity		
Other 4. Race/Ethnicity (Infant)			
White/European (spec			
Black/African America	an/etc. (specify %)		
Asian (specify %)			

Hisp	oanic (spe	ecify %)						
Inuit	t/Eskimo	(specify %)						
Not	Not reported (specify %)							
varie	ed by stud	dy site and experiment	al groups (specify %)					
Non	-white							
Other 5 Is has	eline hio	marker information re	enorted?					
Yes	_	Clear Response	sporteu:					
		ega-3 intake reported	12					
• Yes	• No	Clear Response		ional atudy, r	ologoo oton borol			
III. IIIU		Pregnant	Lactating mothers	Mothers	olease stop here]			
Start tim	e of							
Length (duration intervent								
Longest	follow-							
up time				V. Arms How to fill out this section: For controlled trials, Arm 1 should be the placebo/control group or lowest dose of intervention.				
up time V. Arn How to fi	ill out thi olled trial	s, Arm 1 should be the	placebo/control group	or lowest dose of int	ervention.			
up time V. Arn How to fi	ill out thi olled trial		placebo/control group	or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m	ill out thi olled trial	s, Arm 1 should be the	placebo/control group	or lowest dose of int	ervention.			
up time V. Arn How to fi	ill out thi olled trial nany arm	s, Arm 1 should be the	placebo/control group	or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m Arm 1: Nam	ill out thi olled trial nany arm	s, Arm 1 should be the	placebo/control group	or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m Arm 1: Nam Des	ill out thi olled trial nany arm ne	s, Arm 1 should be the		or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m Arm 1: Nam Des RCT: Arn	nany arm ne cription 1 (place	s, Arm 1 should be the		or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m Arm 1: Nam Des RCT: Arn Bran	ne cription name	s, Arm 1 should be the		or lowest dose of int	ervention.			
up time V. Arn How to fi For control 1. How m Arm 1: Des RCT: Arn Bran Man	ne cription name	s, Arm 1 should be the as are there? ebo or other control) (if applicable)		or lowest dose of int	ervention.			

	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	
Arm	2:	
	Name	
	Description	
RCT	: Arm 2 (placebo or other control)	
	Brand name (if applicable)	
	Manufacturer (specify)	
	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	
Arm	3:	
	Name	
	Description	
RCT	: Arm 3 (placebo or other control)	
	Brand name (if applicable)	
	Manufacturer (specify)	

	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	
Arm	4:	
	Name	
	Description	
RC1	F: Arm 4 (placebo or other control)	
	Brand name (if applicable)	
	Manufacturer (specify)	
	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	
Arm	5:	
	Name	
ECT	Description 7: Arm 5 (placebo or other control)	

	Brand name (if applicable)	
	Manufacturer (specify)	
	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	
	se indicate if there are any references from the studies reference list that we should p rence number from the article)	ull (indicate the
Arm	6:	
	Name	
	Description	
RCT	: Arm 6 (placebo or other control)	
	Brand name (if applicable)	
	Manufacturer (specify)	
	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	

	Maternal conditions	
	Infant conditions	
Arm	7:	
	Name	
	Description	
RCT	: Arm 7 (placebo or other control)	
	Brand name (if applicable)	
	Manufacturer (specify)	
	Purity data (specify)	
	Presence of other potentially active ingredients (e.g., arachidonic acid, vitamin E) (specify)	
	Storage conditions or other efforts to preserve product viability) (specify)	
	n-3 composition (e.g., grams or percent EPA, DHA per capsule) (specify)	
	Dose per day (e.g., 1 1gm capsule, twice a day) (specify)	
	If placebo, how was blinding achieved? (specify)	
	Maternal conditions	
	Infant conditions	