# Reducing the risk of early preterm birth: Omega-3 status test screening and targeted advice

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On behalf of the Omega-3 Implementation Group



### 15 million babies are born preterm each year



### Preterm birth (<37 weeks)

- 85% of all perinatal complications and death
- leading cause of death in children <5 years

### Early preterm birth (< 34 weeks)

- ~20% of preterm births
- ~75% occur spontaneously
- Observational and RCT evidence suggests omega-3 fatty acids extend the period of gestation and may reduce prematurity

### **Omega-3 marine fats and pregnancy duration**

- Omega-3 long chain polyunsaturated fatty acids (LCPUFA) docosahexaenoic acid (DHA) and eicosapentaenoic acid (EPA), are most commonly found in fish and fish oils
- Population comparisons in Denmark vs Faroe Islands
  Increased length of gestation in islanders with high fish intake
- Influence length of gestation by
  - counteracting pro-inflammatory prostaglandins
  - delaying the initiation of labour and cervical ripening
  - relaxing the myometrium



## **DOMINO Birth Outcomes** (800mg DHA, 100mg EPA)

	Omega-3 (n=1197)	Control (n=1202)
Early preterm birth <34 weeks	1.1%	<b>2.3%</b> p=0.03
Birth <37 weeks	5.6%	7.3% p=0.09
Post-term induction or post- term pre-labour C-section	18%	14% p<0.01

Omega-3 intervention was also associated with:

- Fewer admissions to neonatal intensive care
- Fewer cases of brain injury
- Fewer perinatal deaths



Makrides et al. JAMA 2010;304:1675-83 Zhou et al. AJCN 2012;95:1378-1384



# The path to reduce the risk of preterm birth...

 Synthesising all the available intervention studies in a systematic review and metaanalysis



 Conducting a new modern intervention study that aims to retain the benefit while minimising the risk





# The 2018 Cochrane Review

- 70 RCTs with 19,927 pregnant women (+23 on-going trials)
- Mainly from high income countries, but some from lowmiddle income countries
- Most trials included women with singleton pregnancies
- Test dose range 200mg 2,700mg/d ω-3 LCPUFA
- Mainly as fish oil capsules; also oil, fish, eggs, bars, dairy products
- Mainly throughout second half of pregnancy





# **Primary Outcomes of Cochrane Review**

Variable	Effect size
Birth <34 weeks	RR 0.58 (95% CI 0.44 to 0.77) <b>42% reduction</b> from 4.6% to 2.7% 11 trials with 5,409 women
Birth <37 weeks	RR 0.89 (95% CI 0.81 to 0.97) <b>11% reduction</b> from 13.4% to 11.9% 25 trials with 10,256 women
Birth >42 weeks	RR 1.61 (95% CI 1.11 to 2.33) <b>61% increase</b> from 1.6% to 2.6% 6 trials with 5,141 women





## Some Other Important Outcomes of the 2018 Cochrane Review

Variable	Effect size
Gestational length	MD 1.67 days (0.95 to 2.39), 41 trials with 12517 women
Pre-eclampsia	RR 0.84 (0.69 to 1.01), 20 trials with 8306 women
Perinatal death	RR 0.75 (0.54 to 1.03), 10 trials with 7416 women
Birth Weight	MD 76g (38 to 113), 42 trials with 11584 women
Low birth weight <2500g	RR 0.90 (0.82 to 0.99), 15 trials with 8449 women
SGA	RR 1.01 (0.90 to 1.13), 8 trials with 6907 women
LGA	RR 1.15 (0.97 to 1.36), 6 trials with 3722 women





# What's Important About The Cochrane Review?

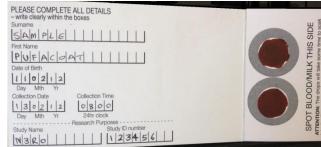
- Includes studies with low-risk, normal risk and high-risk women but almost all women have singleton pregnancies
- Dose: Most studies used >500mg  $\omega$ -3 LCPUFA/day
- Reporting biases may underestimate or overestimate omega-3 effect on prematurity and other adverse birth outcomes
- But, sensitivity analysis with high quality trials only indicate consistent effect of omega-3 supplementation on prematurity



# <u>Omega-3 LCPUFA to Reduce</u> the Incidence of Prematurity







- Blinded RCT in 6 centres
- **5544** women with singleton or multiple pregnancies
- Supplementation from <20 weeks until 34 weeks of gestation to reduce incidence of early preterm birth and prevent the need of obstetric intervention for post-term dates
- Broad based strategy few exclusions
- DBS fatty acid profiles at entry and 34 weeks



Makrides et al. NEJM 2019;381:1035-45

### **ORIP Results**

• Primary outcome - There was little difference between the DHA and control group on the incidence of early preterm birth (<34 weeks)

	Omega-3	Control	Adjusted RR	Adjusted
Primary Outcome	(n=2734)	(n=2752)	(95% CI)	P Value
Early Preterm Birth (<34w)	2.25%	1.98%	1.13 (0.79, 1.63)	0.50

### • Secondary birth outcomes

Outcome	Omega-3 (n=2734)	Control (n=2752)	Adjusted RR (95% CI)	Adjusted P Value
Pre-term Birth (<37w)	7.71%	8.93%	0.86 (0.72, 1.03)	0.11
Post term induction or pre-labour LSCS	5.16%	5.66%	0.91 (0.73, 1.14)	0.42
Gestational Age at birth (mean)	273.18	273.16	-	0.96



Makrides et al. NEJM 2019;381:1035-45





#### RRAD CROUCH HEALTH REPORTER

researchers 5500 studying fish oil supplements to prevent preterro births are rching for the "sweet spot to achieve the best effect for inlividual women. Research by the South Aus alian Health and Medical tesearch Institute's Women Theme has proved dements of omega-3 nd in fish oil - can prevent

ublished today in the New uals can know if they need England Journal of Medicine supplement and also how of a trial involving more than much they need. pregnant Australian "We are trying women found drawbacks with sweet snot women need to be using a universal "one dose fits at." Dr Best said. "If women are Lead nesearcher Dr Karen

Best said the latest results indicate the benefits for preventing preterm birth might be best supplement at all. achieved through a targeted supplementation strategy. She said the work is now focusing on developing a simple test to check the level of omega-3 in women, so individ-

week, or taking a different sup plement contain Adelaide mum-to-b ghan, 28, is 31 weeks pregnan with her first baby and st supports the research.

The work they are doing is fantastic "she said

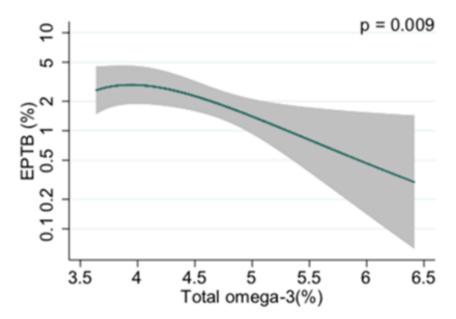
# Interpretation

- ORIP (largest RCT) contrasted most other studies in the Cochrane review
  - singleton pregnancies and/or
  - focused mainly on women with low intakes
- Further analyses for ORIP singleton pregnancies
  - Does baseline omega-3 status predict the risk of FPTB?
  - Does baseline omega-3 status modify the effect of omega-3 LCPUFA supplementation on EPTB?



# Does baseline omega-3 status predict the risk of early preterm birth?

 Risk of EPTB in singleton pregnancies in the control group (n=2545) by baseline omega-3 status in DBS



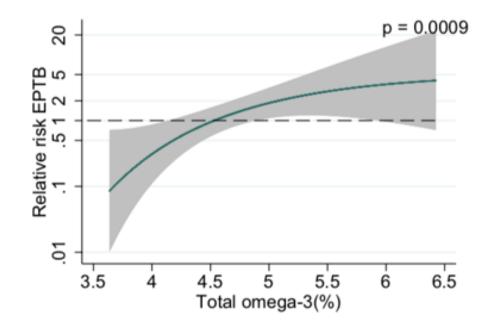
 Low n-3 status in early pregnancy is associated with a higher risk of EPTB



Simmonds et al. BJOG 2020;127:975-81

# Does baseline omega-3 status modify the effect of supplementation on early preterm birth?

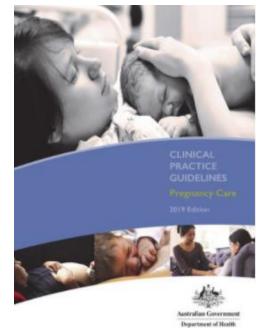
 Relative risk of EPTB (intervention vs. control) in singleton pregnancies by baseline total omega-3 status (n=5070)



Omega-3 status <4.2% + supplementation **reduced** risk of EPTB (RR 0.23, 0.07 - 0.79) Omega-3 status >4.9% + supplementation **increased** risk of EPTB (RR 2.27, 1.13 - 4.58) *Simmonds et al. BJOG 2020;127:975-81* 

### Summary ... & actions

- Women with low omega-3 status are at higher risk of prematurity and supplementing these women reduces their risk of EPTB
- Women with higher omega-3 status do not need extra omega-3 supplementation, and extra supplementation may increase their risk of early preterm birth
- Updated recommendations for omega-3 in pregnancy



"assess omega-3 status and supplement pregnant women with low omega-3"

New Omega-3 test for all pregnant women in South Australia - personalised advice based on omega-3 status



### How to implement into practice?

Patient Details	Ethnic Group: Caucasian	Aboriginal Asian A	frican-Caribbean
Family Name		Given Name(s)	incan-oanoosan
army rearrie		Civen Hamo(s)	
Date of Birth	UR Number	Medicare Number	
Date of Dittr	On Humber		
Address			
Suburb		State	Postcode
Clinical Details - Ma	Indatory		
	Constant Constant Constant	Neural Tube Defect only	] Omega-3 status (SAHMRI)
EDD/LMP / /	_ Cycle length (days)	Maternal weight (Kgs)	
GA Clinical weeks + days	on	/ /	
GA Ultrasound weeks + days	on	/ /	
Crown-rump length (CRL) mr	non	/ /	
Pregnancy: Singleton		]Yes No Age at egg retr	ieval/age of egg donor
	abetes (IDMM only) Yes No		Previous [] T21 [] T18/13
regnancy complications. Di			
Name of Imaging Practice:			
	sk assessment an Ultrasound requ		
Patient status at the time of the service a private patient in a private hospital or app a private patient in a recognised hospital a public patient in a recognised hospital an outpatient public of a recognised hospital	proved day hospital facility	Medicare Benefits (Section 20A of the Health the approved pathology practitioner who will re- eligible pathologist determinable service(s) esta Do Not send to My Health Record	Insurance Act 1973). I offer to assign my right to nder the requested pathology service(s) and any bilished as necessary by the practitioner.
an outpatient private of a recognised hospi	tal	X Patient signature	Date //
Privacy Disclosure SAMSAS requires the per the purpose of Risk assessment and Propram	sonal information contained in this request form for Audits. SAMSAS may therefore request copies of	Practitioner	's Use Only
ultrasound and cytogenetic reports from your	doctor in order to complete its testing and audits.	(Reason patie	nt cannot sign)
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	and URN on the sample label and reg		nd/or checking the patient's ID band
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Tel: Email: Signature: Request Date: / Deliver to: South Australian Maternal SA Pathology, Specimen f	/ Serum Antenatal Screening (SAMSAS) P Seception Area, Level 3, Royal Adelaide H 8161 8085 samsas,program@health.t	ospital. Port Rd ADELAIDE SA 5000.	is.html

- Collaboration with SA Pathology
- SA Maternal Serum Antenatal Screening Program covers 80-85% of women in SA with testing
   before 16 weeks
- No additional blood test and no cost to women or health service
- Omega-3 testing not currently offered by SA Pathology, so testing will be done at SAHMRI



### **Omega-3 Status Test Results**

- Stand alone reports (with omega-3 recommendation) sent to requesting doctor
- Results also available on OACIS

- SA Pathology Report -

Omega-3 Status is ..... % total fatty acids xx weeks of gestation

Omega-3 Status <sup>1</sup>	Recommendation	
Omega-3 Status <3.7%	Take omega-3 fatty acid supplements until 37 weeks to reduce the risk of early preterm birth.         Suggested dose: 800 mg DHA and 100 mg EPA per day.         Typical suitable supplements include Infantem (algal oil supplement from Pharmamark) and Omega Brain (fish oil supplement from Blackmores)	
Omega-3 Status 3.7 to 4.3%	No action required. If already taking omega-3 fatty acids as part of a multivitamin and mineral supplement, this may continue.	
Omega-3 Status >4.3%	Stop any omega-3 fatty acid supplements. Risk of early preterm birth is low and additional omega- 3 supplementation may increase this risk.	

<sup>1</sup> Percentage of omega-3 fatty acid in serum



Omega-3 status is being measured as part of a research evaluation to determine its utility in identifying women with low omega-3 status and advising supplementation to reduce their risk of early preterm birth.

#### Evaluation of omega-3 screening program

It is important to assess how many women have low omega-3 levels and how many babies were born early to see if screening has reduced the number of premature births in South Australia. We will electronically link the omega-3 test results with birth data in a secure way, without identifying you or your baby in any way. You may decline to have your data linked without affecting you or your baby's care.

Opting out: If you do not want your data included in the statewide evaluation of the screening program, please telephone (08) 8128 4444 or email omega3@sahmri.com.

The evaluation has been approved by the Women's and Children's Health Network (WCHN) Human Research Ethics Committee (HREC/20/ WCHN/138). Should you wish to discuss the study with someone not directly involved, you may contact the executive secretary of the Human Research Ethics Committee, Mr Luke Fraser, WCHN (08) 8161 6521.

#### **Further Information**

sahmri.org/omega3
 Speak to your doctor or midwife



If you would like further information about the evaluation of the omega-3 screening program, please contact

omega3@sahmri.com
(08) 8128 4444

### Omega-3 screening to help prevent premature births





SA Pathology and the South Australian Health and Medical Research Institute (SAHMRI) are evaluating the effectiveness of omega-3 screening for pregnant women from 2021 to reduce the number of babies born prematurely.

#### ? Why Omega-3 Screening?

#### Blood collection

Omega-3 fats are nutrients commonly found in fish and algae. The right amount of omega-3 in your body during pregnancy can help you have a fullterm pregnancy. Babies born too soon (premature), particularly those born before 34 weeks of pregnancy, may have lengthy stays in hospital and may experience long-term health problems and delays in development.

Women expecting one baby who have low omega-3 levels in their blood are at higher risk of having their baby born prematurely. They are most likely to benefit from omega-3 supplements. Women who have sufficient omega-3 levels are already at lower risk of having a premature baby and so should not take additional omega-3 supplements.

Screening for omega-3 levels before 20 weeks of pregnancy will identify women who require omega-3 supplements and those who will not.

We do not yet have a screening test for women expecting twins or triplets. Your health professional will give you the SA Maternal Serum Antenatal Screening (SAMSAS) request form to take to a blood collection centre. Your omega-3 level will be measured using the blood collected for the SAMSAS program. The test is provided free to families.



#### **Omega-3 Results**

Your health professional will discuss your omega-3 test results with you.

Omega-3 test result	Health professional advice
Less than 3.7%	<ul> <li>If you have a low omega-3 level (less than 3.7%), your health professional will recommend you take omega-3 supplements (fish oil or algal oil) to reduce your risk of a premature birth:</li> <li>You can take supplements after receiving omega-3 results up until 37 weeks of pregnancy.</li> <li>The suggested dose is 800 mg of DHA and 100 mg of EPA every day.</li> <li>Omega-3 supplements you can take include: Infantem* (Pharmamark) and Omega Brain (Blackmores).</li> <li>PLEASE NOTE: If you are already on prescription medication to stop your blood clotting (like heparin), speak with your doctor before taking omega-3 supplements.</li> </ul>
Between 3.7 and 4.3%	<ul> <li>If you have a moderate omega-3 level:</li> <li>You do not need to do anything different from what you usually do.</li> <li>If you are already taking omega-3 fatty acids as part of a multivitamin and mineral supplement or an omega-3 supplement, you can continue to do so.</li> </ul>
Above 4.3%	<ul> <li>If you have a sufficient omega-3 level:</li> <li>Your risk of a premature birth is low and additional omega-3 supplements may increase this risk.</li> <li>Your health professional will advise that you stop taking any omega-3 supplements.</li> </ul>

\*Algal oil supplement of DHA and EPA.

#### Omega-3 blood sample and cost

No additional blood sample is required as omega-3 analysis will be performed on serum collected as a part of the established SAMSAS program. The omega-3 fatty acid analyses will be performed at no cost to the patient or health service.

### How to order the Omega-3 status test

- Discuss the omega-3 status test and refer woman to the Information for Families sheet.
- Order using the updated SAMSAS request form.
- 3 Tick the omega-3 status test on the SAMSAS request form.
- 4 Refer woman to the Privacy Disclosure on the SAMSAS request form.
- 5 The omega-3 status test results will be reported to the requesting provider as a standalone report and will be available on OACIS.

### Evaluation of Omega-3 status testing program

This SA Pathology SAHMRI collaboration will assess the feasibility and reach of identifying women with low omega-3 status, providing appropriate advice and ultimately assessing success in reducing the rates of early preterm birth. This will be done by deidentified linkage of the omega-3 status test results with relevant pregnancy outcome data. Women who do not want their data linked will need to contact (08) 8128 4444 or email omega3@sahmri.com.

The evaluation has been approved by the Women's and Children's Health Network Human Research Ethics Committee (HREC/20/WCHN/138). Should you wish to discuss the study with someone not directly involved, you may contact the executive secretary of the Human Research Ethics Committee, Mr Luke Fraser, WCHN (08) 8161 6521.

#### Omega-3 status test for prematurity risk

SA Maternal Serum Antenatal Screening (SAMSAS) Program

Information for health professionals



#### **Further Information**

For further information regarding omega-3 status testing or results interpretation:

Visit sahmri.org/omega3

Call the omega-3 status test hotline 0438 273 155





SA Pathology, through the SA Maternal Serum Antenatal Screening (SAMSAS) program, and the South Australian Health and Medical Research Institute (SAHMRI) are evaluating serum omega-3 testing for women with singleton pregnancies in South Australia from 2021.

#### Why do omega-3 status testing?

Women with a singleton pregnancy and low omega-3 status (concentration) in their blood are at higher risk of early preterm birth and most likely to benefit from omega-3 supplementation. Screening before 20 weeks' gestation will identify women who require omega-3 supplementation.

The latest National Health and Medical Research Council and Department of Health National Pregnancy Care guidelines recommend assessing omega-3 fatty acid status and supplementing pregnant women with low omega-3 intakes.

#### There is high quality evidence for omega-3 status testing

 A Cochrane systematic review of 70 randomised controlled trials of almost 20,000 women with mainly singleton pregnancies indicated that omega-3 supplementation from early-mid pregnancy until birth reduces the risk of early preterm birth by 42% (from 46 per 1000 to 27 per 1000 births) and preterm birth by 11% (from 134 per 1000 to 119 per 1000 births)<sup>1</sup>.

 As the Cochrane review included many studies that were conducted before prenatal supplements with low dose omega-3 were commonly taken by women, a large randomised trial, with 5540 women, was undertaken to test if a universal supplementation strategy would be successful with contemporary practice. Overall this trial showed that supplementation of all women would not be effective<sup>2</sup>, but secondary analyses revealed that reductions in early preterm birth are best achieved by a targeted approach to omega-3 supplementation based on omega-3 status before 20 weeks<sup>3</sup>. Women with low omega-3 status are at higher risk of early birth and more likely to benefit from supplementation<sup>2,3</sup>.

1 Middleton P, Gomersall JC, Gould JF, Shepherd E, Olsen SF, Makrides M. Omega-3 fatty acid addition during pregnancy. Cochrane Database Syst Rev. 2018;11:CD003402.

2 Makrides M, Best K, Yelland L, McPhee A, Zhou SJ, Quinlivan J, et al. A randomized trial of prenatal omega-3 fatty acid supplementation and preterm delivery (ORIP trial). New England Journal of Medicine. 2019;381:1035-45. https://doi.org/10.1056/nejmoa1816832

3 Simmonds LA, Sullivan TR, Skubicz M, Middleton PF, Best KP, Yelland LN, et al. Omega-3 fatty acid supplementation in prognancybaseline omega-3 status and early preterm birth: exploratory analysis of a randomised controlled trial (ORIP). BJOG. 2020;27(8):975-961. https://doi.org/10.1111/1471-0528.16168.

4 Percentage of total omega-3 fatty acid status in serum

#### Omega-3 status test results: how to advise women

Omega-3 test result <sup>3,4</sup>	Recommendation
Less than 3.7%	For a low status, <b>take omega-3 fatty acid</b> <b>supplements</b> after receiving the omega-3 results until 37 weeks to reduce the risk of early preterm birth.
	Suggested dose: 800 mg DHA and 100 mg EPA per day
	Typical suitable supplements include Infantem* (Pharmamark) and Omega Brain (Blackmores)
Between 3.7 and 4.3%	For a moderate omega-3 status, <b>no action required.</b> If already taking omega-3 fatty acids as part of a multivitamin and mineral supplement or a standalone supplement, this may continue.
Above 4.3%	For a sufficient status, <b>stop any omega-3 fatty</b> acid supplements.
	Risk of early preterm birth is low and additional omega-3 supplementation may increase this risk.

\*Algal oil supplement of DHA and EPA.

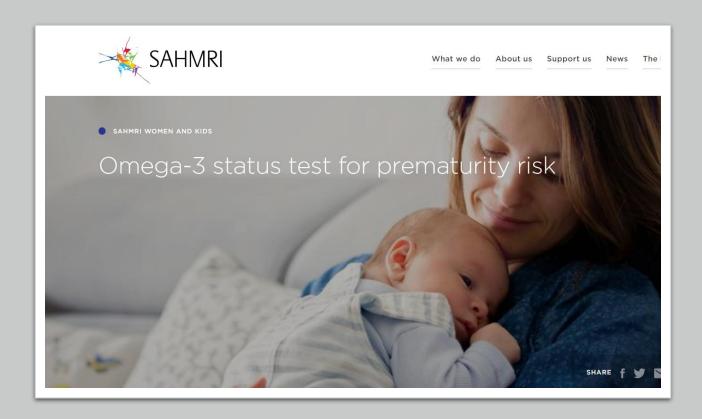
Contraindications to omega-3 fatty acid supplements

- Omega-3 fatty acid supplements should be avoided for women requiring heparin because of possible additive anti-coagulant effects.
- Low dose aspirin can be taken with omega-3 fatty acid supplements.
   Recent randomised trials of omega-3 fatty acid interventions have included women on low dose aspirin without increase in adverse events.

### What next?

- Go live date 19<sup>th</sup> April 2021
- Updated SAMSAS forms on SA Pathology website and hard copy
- Resources for more information
  - SAHMRI website
  - Health professional hotline/email
  - Consumer email
  - Newsletters
  - In-services as required
- Ongoing Evaluation
  - deidentified linkage of the omega-3 status test results with pregnancy outcome to see if specific advice can reduce the rate of EPTBs in SA.
  - Partnership with APBPA to consider extension nationally and internationally





Further Information

- Visit sahmri.org/omega3
- Call the omega-3 hotline 0438 273 155
- For order forms call the SAMSAS Program (08) 816 7285