Supplementary Table 1. Characteristics of included studies divided into categories according to definition (physician-diagnosed, self-report and chronic widespread pain by questionnaire only) and birth cohort studies.

First author	Study	Base	Control	Diagnosis of	Duration	Incidence	Risk factors -Univariate	Risk factors	Quali
& year	design	population	group	Fibromyalgia	Follow-up &	per 1000		Multivariate	ty
					No. of new			Adjusted Hazard	
					cases			ratio (95% CI)	
Physician di	agnosis of Fil	bromyalgia reco	rded on datab	ase					•
Forseth	Prospective	Random	100 FM-free	ACR criteria	5.5 years	FM =5.83/1000	n/a	n/a	n/a
1997 * (1)	cohort	sample of 820	controls	including		person-years			
	study	females aged		tender point	21 cases CWP				
	Females	20±49 yr free		count by	12 had FM by	CWP =1% per			
	aged 20±49	of FM and		clinical	ACR criteria	annum or 10 per			
	yr	followed up		examination		1,000 person -			
		after 5.5 years			Allowing for	years			
		n=717 (82%			drop-outs,				
		response)			23/717 (3.2%)				
Weir	Retrospecti	Deseret	105,021 FM-	ICD-9-CM code	5 years	Males: 6.88 per	*Females		n/a
2006(2)	ve cohort	Mutual	free controls	729.1	N= 2595	1000 person-years	*older age – peak at 50-		
	study	Benefits			incident cases	Females: 11.28	60 years		
		Administration				cases per 1000			
		database				person-years	Associated diagnoses:		
		62,000				(9.01 for both	depression, anxiety,		
		working				sexes combine).	headache, IBS, CFS, RA,		
		people <65					SLE, diabetes. (not clear		
		years old					if these preceded FM)		
Chang 2015	Retrospecti	NHIRD	68,568	ICD-9-CM code	3 years	Depressed cohort:	*female	*Depression	high

(3)	ve cohort	1 million,	Depression-	729.1	N=1800	21.6 per 1000	*Depression	6.28(5.67–6.96)	
	study	17,142	free matched			person-years	*Migraine	*Migraine	
		patients with	controls		Depressed		*low back pain	1.63 (1.04–2.56)	
		depression			cohort:	Non-depressed	*allergic rhinitis	*low back pain	
					FMS 6.5%	cohort:		2.63 (2.18-3.18)	
					V	3.3 per 1000		*allergic rhinitis	
					1.0% non-	person-years	Patients with	1.44 (1.23-1.68)	
					depressed		depression developed		
							FM at earlier age than	Bidirectional:	
					aHR =6.28,		those without.	Patients with FMS	
					(95% CI 5.67-			had increased risk of	
					6.96)			developing	
								depression	
Wang 2017	Retrospecti	NHIRD	138,520	ICD-9-CM	3 ½ years	GERD cohort: 5.76	*GERD	*GERD	high
(4)	ve case	1 million	matched	729.1		per 1000 person-	*age	1.31 (1.19-1.45)	
	cohort	population:	non-GERD		FM onset cases	years	*NSAIDs	*Older Age	
	study	GERD cases	controls		n= 2584	vs	* Diabetes	1.02 (1.01-1.02)	
		n= 34,630				control cohort:	*Hypertension	*Peptic Ulcer	
						3.96 per 1000	*Hyperlipidemia	1.12 (1.02-1.23)	
						person-years	*Anxiety	*NSAID	
							*Sleep disorder	1.17 (1.07-1.27)	
						aHR of 1.44 (95%	*Stroke		
						CI 5 1.29-1.60)	*Peptic ulcer disease	Bidirectional :	
								Patients with FM had	
							Protective: *PPI	greater risk of	
								developing GERD	
							No difference by sex.		
Chen 2017	Retrospecti	NHIRD	4976	ICD-9-CM code	12 years	Osteomyelitis	*female	*female	high
(5)	ve case	1 million	matched	729.1 on at	N= 768	cohort: 25.4 per	*osteomyelitis	1.26 (1.09–1.46)	
	cohort	population:	Osteomyeliti	least 3		1000 person-years	*older age	*osteomyelitis	
	study	Osteomyelitis	s-free	occasions		V	*hypertension,	1.26 (1.06–1.49)	
		patients:	controls			Comparison	* IBS,	*older age	
		n= 1174				cohort: 18.8 per	*sleep disorder	1.48 (1.18–1.85)	

		(mostly older men) **				1,000 person-years	* diabetes *anxiety *Depression *hyperlipidaemia	*hypertension 1.29 (1.07–1.55) * IBS 1.41 (1.01–1.97) *sleep disorder 1.48 (1.21–1.81) Exposure-response effect for severity of osteomyelitis	
Marrie 2012 * (6)	Retrospecti ve case cohort study	Manitoba Health and medical records data N= 4192 with Multiple Sclerosis (72% women)	20,940 Non-MS matched controls	ICD-9-729.1 ICD-10 M79.7 at 5 hospital visits	5 years minimum N= 359 Prevalence of FM was 2.4% by medical records and 3.5% a/c to self-report	1.2 per 1,000 person years v 0.9 per person-years in the general population. IRR=1.44 (95%CI:1.01–2.07)	*female *increasing age *MS Mean age of onset FM was 53.1 in MS patients and 52.6 in general population	*multiple sclerosis 1.44 (1.01-2.07)	high
Chen 2018 (7)	Retrospecti ve case cohort study	NHIRD 1 million: Incident cases of FM over 3 years n=61,613	18,040 -matched patients without IBD	ICD-9-CM code 729.1	12 years N= 1,106 of IBD group who developed FM N=2,606 in the non-IBD group,	IBD group: 32.7 per 1,000 person-years, Control cohort: 18.2 per 1,000 person-years	*older age – 40-59 group No sex difference in IBD group but in non-IBD cohort higher incidence in women compared to men Ulcerative colitis and Crohn's disease act as	*IBD 1.70 (1.59–1.83) *Medical Comorbidity In non- IBD cohort only: *Older age *Female IBD had relatively	high

							an independent risk	larger influence in	
							factors	the younger age	
								groups who had	
								fewer comorbidities	
I-Wen Penn	Retrospecti	LHID	69,680	ICD-9-CM code	12 years	Migraine cohort:	*Migraine	*Migraine	high
2019 (8)	ve cohort	17,420	controls	729.1)		7.01 per 1000	*Older age	1.51 (1.38 to 1.65)	
	study	migraine			N=2,834	Person Years	*Diabetes	*Older age	
		patients with					*Hypertension	1.02 (1.01 to 1.02)	
		no previous				Non-migraine	*Hyperlipidaemia	*Hyperlipidaemia	
		FM				cohort: 4.49 per	*Depression	1.15 (1.03 to 1.28)	
						1000 Person Years	*Anxiety		
							*Sleep disorder	Bidirectional but	
							*Coronary Heart	FM had stronger	
							Disease	predictive power for	
							*Irritable Bowel	the onset of	
							Syndrome	migraine	
							(Sex and chronic fatigue	than did migraine for	
							syndrome not	the onset of FM.	
							associated)		
Kivimaki	Prospective	4791 (540		Physician-	2 years	4.9 per 1000		*High workload,	fair
2004 (9)	cohort	male,		diagnoses FM		person years		2.5 (1.1–5.6),	
	study	4251 female)			47/4791			*low decision	
		Hospital						latitude,	
		employees						2.6 (1.2-5.9),	
		(59%						*workplace	
		response)						bullying	
								3.9 (1.3–11.8),	
Gallagher	Retrospecti	UK GPRD	n/a	Clinical	11 years	0.4 per 1000 in	*female sex.		n/a
et al 2004	ve case	2.4 million		diagnosis by		2000/2001	*age 40- 55yrs		
(10)	cohort			GP using READ					
	study			code.					
Collin et al	Electronic	All patients	n/a	GP recorded	13 years	Annual incidence	*Females had six-fold		n/a

2017 (11)	health records	with 1 st FM diagnosis over		diagnosis using READ	N=20,984	rate for FM diagnosis was 0.33	higher incidence of FM than men		
	cohort	13 year period		code.		per 1000	*Peak age = 40 and 49		
	study.	from 660		couc.		per 1000	years		
	GPRD	practices.					*lower SES		
Self-reporte	ed diagnosis o	of Fibromyalgia							
Mork 2012	Prospective	HUNT1 -		Self report –	11 years	incident FM,	*Sleep problems in	*sleep problems	fair
* (12)	cohort	12,350 women		physician		(2.4 per 1000	exposure-response	3.43 (2.26–5.19)	
	study	pain- free		diagnosed FM	327 new onset	person-years).	manner.		
		at baseline.							
		(68% followed					No diff re age.		
		up)							
Mork 2010	Prospective	HUNT1 -		Self report –	11 years	380 incident FM,	*little exercise .	*raised BMI	fair
* (13)	cohort	15,990 women		physician		(2.2 per 1000	*Raised BMI	1.70 (1.35–2.13)	
	study	pain- free		diagnosed FM	380 new onset	person-years)			
		at baseline.			CWP				
Choi 2010	Prospective	Adventist	Non-FM	Self report	20 years	1.72/1,000 per	*smoking	*smoking	high
(14)	cohort	Health study	women in	"Have you	N=136	year	*>1 prior surgery	2.37 (1.33-4.23)	
	study	(women)	study	been			*symptom control	*1 allergy	
		3156 included		diagnosed			medication ^a	1.61 (0.92–2.83)	
		in both and		with			*no of allergies (dose	*2+ allergies	
		second survey		fibromyalgia			response)	3.99 (2.31-6.88)	
		25 years apart		by a physician"			* prior peptic ulcer		
				Excluded			*prior hyperemesis	Not associated:	
				those with FM			gravidarum.	age, BMI,	
				before first				education,	
				survey			^a analgesia, anti-	employment &	
							indigestion medication,	marital status,	
							tranquilizers, hypnotics	hypertension,	
								breast cysts/lumps	
								or RA	
Chronic wid	espread pain	identified by q	uestionnaire o	nly					

Hagen	Prospective	HUNT study:		CWP by ACR	11 years	16.4 per 1000	*any headache	After adjustment	fair
012 (15)	cohort	1995/97 and		questionnaire	N=2494	person-years	*migraine	for age, gender,	
	study	2006/08:					* non-migrainous	BMI education,	
		41,766 eligible					headache	physical activity,	
		in both. Of						smoking, anxiety/	
		these, 26,197					Exposure-response	depression:	
		surveyed (63%					effect for frequency of		
		response)					headaches	*any headache	
		13,781 were					Bidirectional	>14x /month	
		CWP-free						2.7 (1.6-4.7)	
								*migraine	
								2.0 (1.6-2.4)	
Nundal	Prospective	Nord-	No CWP at	CWP by	11 years,	10.9 per 1000	*female	*female	high
014 (16)	cohort	Trøndelag	follow-up	questionnaire	12% developed	person years	*current pain	*Anxiety/depressio	
	study	Health Study		- ARC ACR	CWP:		*Anxiety	n 1.73(1.42-2.09)	
		19,192 free of		criteria		7.2 per 100 person	*depression,	*smoking	
		CWP (82%			8% (n = 915) of	years for pain-free	*smoking	1.45(1.30-1.62)	
		response)			those pain-free	but	*sleep problems	*alcohol regularly	
					at baseline		*BMI =underweight	0.79(0.64-0.97)	
					&	18.2 per 1000 py	*BMI=overweight/	*sleep problems	
					20% (n = 1412)	For those with	obese	1.49(1.33-1.67)	
					of those with	some pain at		*low BMI	
					some pain at	baseline	Moderate alcohol use	2.05 (1.24-3.39)	
					baseline .		was protective	*raised BMI	
								1.35 (1.18-1.55)	
Jhlig 2018	Prospective	2 nd and 3 rd	13,113 no	CWP by ACR	11 years -mean	Total population:	*insomnia	*Insomnia	fair
k Sivertsen	cohort	HUNT studies	insomnia	criteria by	N=1400	9.4 per 1000	*fewest and highest yrs	1.93,(1.40-2.66)	
014 (17,	study	Full data on		questionnaire	developed	person years	of education		
.8)		13,113 (23% of			CWP			Insomnia was risk	
		original		Compared to		17.8 per 1000	significant interaction	factor for 11 out of	
		sample)		clinical	19.6% of	person-years for	for education	18 conditions but	
	1			interview -	insomnia	insomnia cohort	level (P interaction =	only 3 had adjusted	

		316 had		adjusted	cohort	V	0.03. : In fully	OR >2: anxiety,	
		insomnia		agreement	v	9.3 per 1000	adjusted analyses	depression and FM	
				(kappa value)	10.2% of	person-years for	separated by education	: 2.05, (1.51–2.79)	
				for CWP = 0.48	control cohort	control cohort	level, RRs		
				(95% CI 0.38-			for CMSC were :		
				0.64).		Adj HR = 1.58	≤9 years 1.31 (1.0–1.65)		
						(1.26–1.98)	10-12yrs 0.96 (0.8–1.2)		
							≥13 yrs.1.25 (1.0–1.56)		
McCabe	Prospective	European		ACR by	4.3 years	16.5 per 1000	*High BMI		fair
2016 (19)	cohort	Male Ageing		questionnaire		person years	*Depressed		
	study **	study			151 (7.1%) new		*2 comorbidities		
		2118 men			CWP		*longer sit to stand		
		CWP-free at					time		
		baseline					*slow to walk 50 yards		
		followed up					*no alcohol		
		mean age = 59							
		years					No association with		
							vitamin D deficiency		
McBeth	Prospective	Random	CWP-free at	ACR criteria by	1 year	81 (5.8%) had	* female	*Illness behaviour	Fair
2001 (20)	cohort	sample of	follow-up	questionnaire		developed CWP	*older age	9.0 (3.7–22.2)	
	study	population			N=81		*anxiety/depression	*Somatic	
		1658 FM-free				58 per 1000	*Illness Behaviour *	symptoms	
		at baseline				person-years	*Somatic symptoms	3.3 (1.5–7.4)	
		(75% response					* pre-existing pain		
		rate at				8% of those with			
		baseline).				some pain at	†includes frequent visits		
						baseline	to GP		
		1480 (93%							
		response) 1				2% of pain-free			
		year later				cohort at baseline			
		(50% of							
		original							
		sample).							

McBeth	Prospective	Random	1322 CWP-	ACR criteria	1 year	6% had developed	*heavy manual work	*Illness behaviour	fair
2003 (21)	cohort	sample of	free at	but without TP	N=81 had CWP	CWP	*kneeling	2.1(1.2-3.9)	
(same	study	population	follow-up	examination			*repetitive wrist work	*repetitive wrist	
sample as		1403 FM-free					*Illness Behaviour +	Work	
McBeth		at baseline					*Somatic symptoms	1.8(1.2-2.7)	
2001)		and followed					*Fatigue	*any pain at	
		up at 1 year.						baseline	
		(91%					†includes frequent visits	2.1(1.3-3.3)	
		response)					to GP		
McBeth	Prospective	Altrincham		28 (11%) had	1 year	n/a	* depression	After adjustment	fair
2007	cohort	pain study -		new onset			*poor sleep,	for age, sex,	
Substudy of	study	267 at high		CWP			*threatening life events	depression,	
Mcbeth		risk of CWP -						disturbed sleep,	
2001		agreed to HPA					High cortisol:	traumatic life	
(22)		measurement					*post dexamethasone	events and pain	
		58%response -					*evening saliva	status:	
		241 followed						*High level of	
		up					*low morning cortisol	cortisol post	
							_	dexamethasone	
							Independent and	*Low cortisol in	
							additive predictors of	morning saliva	
							CWP	*high cortisol	
								evening saliva	
Gupta	Prospective	3171 adults	CWP-free at	CWP by ACR	15 months	10.2% developed	*pain (not chronic) at	* non-chronic pain	Fair
2007	cohort	aged 25–65	follow-up	criteria by		CWP at follow-up	baseline	at baseline	
(23)	study	yrs free of		questionnaire	N=324 (10.2%)		*somatic symptoms	6.1 (4.3, 8.6)	
		CWP were				= 81.6 per 1000	*illness behaviour	*Somatic	
		followed-up				person years	*health anxiety	symptoms	
		15 months					*anxiety	1.8 (1.1, 3.1)	
		(82%					*depression	*Illness behaviour	
		response) but					*sleep disturbance	3.3 (2.3, 4.8)	
		75% of whole					*stressful life events	*sleep disturbance	
		cohort						2.2 (1.6, 3.2)	

							No difference by age	All showed an	
							and sex. No effect of	exposure-response	
							SES.	relationship and	
								their effect on risk	
								was additive	
Gupta 2007	Prospective	768 CWP- free	CWP-free at	CWP by ACR	15 months	11.3% developed	Pain threshold at	An increased	fair
(24)	cohort	Of which 463	follow-up	criteria by	231 subjects at	CWP at follow-up	baseline did <u>not</u> predict	sensitivity to	
	study	were invited		questionnaire	baseline free of		CWP onset	pressure pain in	
Subset of		and 267 (58%)		- Selected	CWP, 26/231	(high risk group)		subjects with	
above		agreed, of		"high risk"	had new CWP		Tender point count did	chronic widespread	
study		whom 254		group	(11.3%)		not predict new CWP	pain is therefore	
		(54%)						likely to be a	
		provided full						secondary	
		data. 231						phenomenon	
		followed-up						rather than the	
		15 months						antecedent of pain.	
McBeth	Prospective	North	WP-free at	WP by ACR	3 years	For whole cohort:	*female	*older age	fair
2014 (25)	cohort	Staffordshire	follow-up	criteria by		61 per 1000	*few years education	0.97 (0.96-0.99)	
	study	Osteoarthritis		questionnaire	N=800/4326	person years:	*high/low BMI	*baseline pain	
		Project					*moderate/low SES	1.2 (1.1-1.3)	
		Data on 4326			3526	For pain free: 26	*low social participation	*anxiety	
		at baseline			No pain at	per 1000 person-	*no alcohol	1.4 (1.0-2.1)	
		(>50 years)			baseline: 7.7%	years	*financial strain,	*physical	
		and 3-yr			(121/1562)		*obese	impairment	
		follow up (46%			developed WP		*anxiety or depression,	1.2 (1.0-1.5)	
		response)				For cohort with	*lower physical and	*cognitive	
		800 were			Regional pain	regional pain:	mental HRQOL	complaint	
		CWP- free at			at baseline:	82 per 1000	*osteoarthritis	1.3 (1.0-1.6)	
		baseline			24.6%	person-years	*comorbidities	*nonrestorative	
					(679/2764)		*cognitive impairment	sleep	
					developed WP		*sleep problems	1.9 (1.2-2.8)	
								*diffuse OA	
								1.7 (1.3-2.1)	

Papageorgi ou 2002	Prospective cohort	2334 in original		CWP a/c ACR definition	7 years	6.4% developed CWP over 7 years	* Some pain at baseline	female:	fair
(26)	study	survey: 1329 CWP- free at baseline followed up 7 years later		(questionnaire only)	75/1156 developed CWP 13/562 2.3% of pain-free and 62/594=10.4% of "some pain"	= 9.1 per 1000 person years Of pain –free =3.3 per 1000 person years Of "some pain" cohort= 14.8 per 1000 person years	2% and 10% of subjects with no pain and "other pain" at baseline had developed CWP after seven years, respectively. No sex difference	*Fatigue 4.8 (2.2 to 10.8) *Dry eyes 3.5 (1.5 to 7.9) Age >50 yrs 3.0 (1.4 to 6.1)	
Andorsen 2017 (27)	Prospective cohort study	Tromso Longitudinal study N= 4,496 pain free at baseline and followed-up (66% response)	No CWP at follow-up	Severe musculo- skeletal pain in 3 regions for more than 3 months	13 years N=441	7.5 per 1000 per year	*female gender, *current smoking, *poor self-perceived health, *low educational level, *high BMI *Physically inactive (not in multivariate analysis) *Mental health complaints	*female gender 1.46 (1.29-1.66) *current smoking 1.33 (1.16-1.52) *poor self- perceived health 1.62 (1.30-2.02) *few yrs education 1.73 (1.46-2.05) *high BMI 1.39 (1.10-1.77) Mental health complaints (men only) 2.03 (1.18-3.50)	Fair
Bergman 2002 & 2004 (28, 29)	Prospective cohort study	1852 Participated in 1995 and 1998 (78% response 1150 pain-free	No CWP at follow-up	CWP by ACR criteria for postal survey	3 years N= 101 new onsets of CWP: 25/1150	7.2 per 1000 person-years for pain-free at baseline	*Initial chronic regional pain *Older age (>59 years) *lack of social support *Immigrant status	*older age 3.1 (1.47-6.7) * family history of chronic pain 1.9 (1.1-3.1)	fair

		at baseline 463 had chronic regional pain			(2.17%) of pain-free 76/463 (16.4%) of those with regional pain	54.6 per 1000 person-years for those with regional pain at baseline.	*current smoker *regular alcohol protective Not associated: sex, SES, educational level, housing area.	* >6 painful regions of pain 12.1 (4.5-33) Protective factors: *regular alcohol 0.42 (0.2-0.85) *personal social support 0.49 (0.28-0.85) Exposure response effect - number of painful sites.	
Aili 2018 (30)	Prospective cohort study	EPIPAIN Study Participants without sleep problem in 1995/8 Followed up after 5 years N=1249 (83%) and 18 yrs, n=791 (52% response) Age 20- 74years	Pain-free at follow-up	CWP by questionnaire - ACR criteria	5 & 18 years New CWP: 89 (7%) at 5 yr follow-up 103 (13%) at 18 years follow-up	Over 5 years Incidence= 14.2 per 1000 person-years Over 18 years: 7.2 per 1000 person-years	*early awakening *non-restorative sleep *initiating sleep *maintaining sleep *fatigue *no of pain regions *poor mental health No difference by sex at 5 yr, but more female at 18years	At 5 yrs: * 4 sleep problems 2.18 (1.04–4.56) *6 pain sites 7.78 (3.13–19.26) *manual work 2.34 (1.23-4.44) *fatigue 2.68 (1.21 – 5.96) At 18 years f-up: *Female *all 4 sleep problems 2.29 (1.01–5.18) *no of pain sites 8.24 (2.88–23.63) Sleep and fatigue influenced CWP onset	fair

								independently
Birth cohor	t studies							
Vandenker khof 2011 (31)	Prospective cohort study	1958 British Birth Cohort Study At age 45 years. N=was 8572 in present study	Participants free of CWP at follow up	ACR criteria by Questionnaire	n/a	Prevalence: 1056 (12%) had CWP at age 45 years	at 33 years: both sexes: *Raised BMI *unemployed *smoking *physical exertion at work *low SES * Other marital status *alcohol - protective women only: *fruit and veg rarely *frequent fatty foods	both sexes: *Raised BMI *unemployed *smoking *physical exertion at work Women: *fruit and veg rarely *alcohol protective Men: *marital status not single
Jones 2007 (32)	Prospective cohort study	1958 British Birth Cohort Study N=10,453: 7,470 participated at age 45 years (71.5%).	Participants free of CWP at follow up	ACR criteria by Questionnaire	n/a	Prevalence: 915 (12.2%) had CWP aged 45 years No difference between men (11.8%) and women (12.7%)	*Multiple symptoms aged 7 years (vomiting, bilious attacks, periodic abdominal pain and frequent headaches or migraine)	*Multiple symptoms during childhood As proportion of children exposed was small, the population attributable risk associated with childhood symptom reporting was low.
Jones 2009	Prospective	1958 British	Participants	ACR criteria by	n/a	Prevalence:	Childhood traumas:	Prior to 7 years of

(33)	cohort study	Birth Cohort Study N= 9377 participants followed up at age 45 yrs, 7571 participated (71.5% response)	free of CWP at follow up	Questionnaire		12.3% had CWP at 45 yrs was and there was no difference between men (11.8%) and women (12.7%)	*hospitalisation as child after traffic accident *prolonged separation from mother/ in institutional care <7 yrs *Maternal death before age 7 yrs *family financial difficulties	age: *family financial difficulties *maternal death *periods in local authority care .
Pang 2010 (34)	Prospective cohort study	Birth Cohort Study N= 14,936 participants followed up at age 45 yrs, 8572	Participants free of CWP at follow up	ACR criteria by Questionnaire	n/a	12.9% of females and 11.7% of males		After adjustment for gender, social class, childhood symptoms and adult: * persistent behaviour problems at 7, 11 and 16 years reported by teacher (RR 2.14; 95% Cl 1.43, 3.21)
Gale 2012 (35)	Prospective cohort study	1958 British Birth Cohort Study N= 6902 participants followed up at age 45 yrs, 7571 participated (71.5%	Participants free of CWP at follow up	ACR criteria by Questionnaire	N=993	Prevalence: 14.4% had CWP at 45 yrs was and there was no difference between the sexes (51.9% were women)	*lower IQ measured at age 11 years. As mentioned above (Macfarlane 2009) CWP associated with low SES, and at age 42 years, with no academic qualifications, smoking, higher BMI &	After adjustment for confounders : *lower IQ measured at 11 years Linear relationship between IQ age 11 years and risk of

Prospective cohort study Prospective cohort			response)					psychological distress.	CWP at 45 years	
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			NB dose response	
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^{*=} women only in the sample

NHIRD = National Health Insurance Research Database (Taiwan)
GPRD = UK General Practice Research Database
IRR = Incidence rate ratio
FM=Fibromyalgia

^a= tranquilizers, sleep medications, laxatives, pain killers or pep pills

b= incidence is 4 times higher than prevalence in this study

LHID = Longitudinal Health Insurance Database (Taiwan), NHIRD=National Health Insurance Research Database (Taiwan)

IRR= incidence rate ratio

ICPC= International Classification of Primary Care (from Wonca International Classification Committee).

NCDS= National Child Develoment Study

- 1. Forseth KO, Gran JT, Husby G. A population study of the incidence of fibromyalgia among women aged 26-55 yr. British journal of rheumatology. 1997;36(12):1318-23.
- 2. Weir PT, Harlan GA, Nkoy FL, Jones SS, Hegmann KT, Gren LH, et al. The incidence of fibromyalgia and its associated comorbidities: a population-based retrospective cohort study based on International Classification of Diseases, 9th Revision codes. Journal of clinical rheumatology: practical reports on rheumatic & musculoskeletal diseases. 2006;12(3):124-8.
- 3. Chang MH, Hsu JW, Huang KL, Su TP, Bai YM, Li CT, et al. Bidirectional Association Between Depression and Fibromyalgia Syndrome: A Nationwide Longitudinal Study. The journal of pain: official journal of the American Pain Society. 2015;16(9):895-902.

^{**=}predominantly or only men in the sample

- 4. Wang JC, Sung FC, Men M, Wang KA, Lin CL, Kao CH. Bidirectional association between fibromyalgia and gastroesophageal reflux disease: two population-based retrospective cohort analysis. Pain. 2017;158(10):1971-8.
- 5. Chen JH, Muo CH, Kao CH, Tsai CH, Tseng CH. Increased Risk of New-Onset Fibromyalgia Among Chronic Osteomyelitis Patients: Evidence From a Taiwan Cohort Study. The journal of pain: official journal of the American Pain Society. 2017;18(2):222-7.
- 6. Marrie RA, Yu BN, Leung S, Elliott L, Warren S, Wolfson C, et al. The incidence and prevalence of fibromyalgia are higher in multiple sclerosis than the general population: A population-based study. Multiple sclerosis and related disorders. 2012;1(4):162-7.
- 7. Chen JH, Chen HJ, Kao CH, Tseng CH, Tsai CH. Is Fibromyalgia Risk Higher Among Male and Young Inflammatory Bowel Disease Patients? Evidence from a Taiwan Cohort of One Million. Pain physician. 2018;21(3):E257-e64.
- 8. Penn IW, Chuang E, Chuang TY, Lin CL, Kao CH. Bidirectional association between migraine and fibromyalgia: retrospective cohort analyses of two populations. BMJ open. 2019;9(4):e026581.
- 9. Kivimaki M, Leino-Arjas P, Virtanen M, Elovainio M, Keltikangas-Jarvinen L, Puttonen S, et al. Work stress and incidence of newly diagnosed fibromyalgia: prospective cohort study. Journal of psychosomatic research. 2004;57(5):417-22.
- 10. Gallagher AM, Thomas JM, Hamilton WT, White PD. Incidence of fatigue symptoms and diagnoses presenting in UK primary care from 1990 to 2001. Journal of the Royal Society of Medicine. 2004;97(12):571-5.
- 11. Collin SM, Bakken IJ, Nazareth I, Crawley E, White PD. Trends in the incidence of chronic fatigue syndrome and fibromyalgia in the UK, 2001-2013: a Clinical Practice Research Datalink study. Journal of the Royal Society of Medicine. 2017;110(6):231-44.
- 12. Mork PJ, Nilsen TI. Sleep problems and risk of fibromyalgia: longitudinal data on an adult female population in Norway. Arthritis and rheumatism. 2012;64(1):281-4.
- 13. Mork PJ, Vasseljen O, Nilsen TI. Association between physical exercise, body mass index, and risk of fibromyalgia: longitudinal data from the Norwegian Nord-Trondelag Health Study. Arthritis care & research. 2010;62(5):611-7.
- 14. Choi CJ, Knutsen R, Oda K, Fraser GE, Knutsen SF. The association between incident self-reported fibromyalgia and nonpsychiatric factors: 25-years follow-up of the Adventist Health Study. The journal of pain: official journal of the American Pain Society. 2010;11(10):994-1003.
- 15. Hagen K, Linde M, Steiner TJ, Zwart JA, Stovner LJ. The bidirectional relationship between headache and chronic musculoskeletal complaints: an 11-year follow-up in the Nord-Trondelag Health Study (HUNT). European journal of neurology. 2012;19(11):1447-54.
- 16. Mundal I, Grawe RW, Bjorngaard JH, Linaker OM, Fors EA. Psychosocial factors and risk of chronic widespread pain: an 11-year follow-up study--the HUNT study. Pain. 2014;155(8):1555-61.
- 17. Uhlig BL, Sand T, Nilsen TI, Mork PJ, Hagen K. Insomnia and risk of chronic musculoskeletal complaints: longitudinal data from the HUNT study, Norway. BMC musculoskeletal disorders. 2018;19(1):128.

- 18. Sivertsen B, Lallukka T, Salo P, Pallesen S, Hysing M, Krokstad S, et al. Insomnia as a risk factor for ill health: results from the large population-based prospective HUNT Study in Norway. Journal of sleep research. 2014;23(2):124-32.
- 19. McCabe PS, Pye SR, Beth JM, Lee DM, Tajar A, Bartfai G, et al. Low vitamin D and the risk of developing chronic widespread pain: results from the European male ageing study. BMC musculoskeletal disorders. 2016;17:32.
- 20. McBeth J, Macfarlane GJ, Benjamin S, Silman AJ. Features of somatization predict the onset of chronic widespread pain: results of a large population-based study. Arthritis and rheumatism. 2001;44(4):940-6.
- 21. McBeth J, Harkness EF, Silman AJ, Macfarlane GJ. The role of workplace low-level mechanical trauma, posture and environment in the onset of chronic widespread pain. Rheumatology (Oxford, England). 2003;42(12):1486-94.
- 22. McBeth J, Silman AJ, Gupta A, Chiu YH, Ray D, Morriss R, et al. Moderation of psychosocial risk factors through dysfunction of the hypothalamic-pituitary-adrenal stress axis in the onset of chronic widespread musculoskeletal pain: findings of a population-based prospective cohort study. Arthritis and rheumatism. 2007;56(1):360-71.
- 23. Gupta A, Silman AJ, Ray D, Morriss R, Dickens C, MacFarlane GJ, et al. The role of psychosocial factors in predicting the onset of chronic widespread pain: results from a prospective population-based study. Rheumatology (Oxford, England). 2007;46(4):666-71.
- 24. Gupta A, McBeth J, Macfarlane GJ, Morriss R, Dickens C, Ray D, et al. Pressure pain thresholds and tender point counts as predictors of new chronic widespread pain in somatising subjects. Annals of the rheumatic diseases. 2007;66(4):517-21.
- 25. McBeth J, Lacey RJ, Wilkie R. Predictors of new-onset widespread pain in older adults: results from a population-based prospective cohort study in the UK. Arthritis & rheumatology (Hoboken, NJ). 2014;66(3):757-67.
- 26. Papageorgiou AC, Silman AJ, Macfarlane GJ. Chronic widespread pain in the population: a seven year follow up study. Annals of the rheumatic diseases. 2002;61(12):1071-4.
- 27. Andorsen OF, Ahmed LA, Emaus N, Klouman E. A prospective cohort study on risk factors of musculoskeletal complaints (pain and/or stiffness) in a general population. The Tromso study. PloS one. 2017;12(7):e0181417.
- 28. Bergman S, Herrstrom P, Jacobsson LT, Petersson IF. Chronic widespread pain: a three year followup of pain distribution and risk factors. The Journal of rheumatology. 2002;29(4):818-25.
- 29. Bergman S, Jacobsson LT, Herrstrom P, Petersson IF. Health status as measured by SF-36 reflects changes and predicts outcome in chronic musculoskeletal pain: a 3-year follow up study in the general population. Pain. 2004;108(1-2):115-23.
- 30. Aili K, Andersson M, Bremander A, Haglund E, Larsson I, Bergman S. Sleep problems and fatigue as predictors for the onset of chronic widespread pain over a 5- and 18-year perspective. BMC musculoskeletal disorders. 2018;19(1):390.
- 31. Vandenkerkhof EG, Macdonald HM, Jones GT, Power C, Macfarlane GJ. Diet, lifestyle and chronic widespread pain: results from the 1958 British Birth Cohort Study. Pain research & management. 2011;16(2):87-92.

- 32. Jones GT, Silman AJ, Power C, Macfarlane GJ. Are common symptoms in childhood associated with chronic widespread body pain in adulthood? Results from the 1958 British Birth Cohort Study. Arthritis and rheumatism. 2007;56(5):1669-75.
- 33. Jones GT, Power C, Macfarlane GJ. Adverse events in childhood and chronic widespread pain in adult life: Results from the 1958 British Birth Cohort Study. Pain. 2009;143(1-2):92-6.
- 34. Pang D, Jones GT, Power C, Macfarlane GJ. Influence of childhood behaviour on the reporting of chronic widespread pain in adulthood: results from the 1958 British Birth Cohort Study. Rheumatology (Oxford, England). 2010;49(10):1882-8.
- 35. Gale CR, Deary IJ, Cooper C, Batty GD. Intelligence in childhood and chronic widespread pain in middle age: the National Child Development Survey. Pain. 2012;153(12):2339-44.
- 36. Bendayan R, Cooper R, Muthuri SG. Lifetime cigarette smoking and chronic widespread and regional pain in later adulthood: evidence from the 1946 British birth cohort study. BMJ open. 2018;8(8):e021896.
- 37. Muthuri SG, Kuh D, Bendayan R, Macfarlane GJ, Cooper R. Chronic physical illness in early life and risk of chronic widespread and regional pain at age 68: evidence from the 1946 British birth cohort. Pain. 2016;157(10):2382-9.
- 38. Markkula RA, Kalso EA, Kaprio JA. Predictors of fibromyalgia: a population-based twin cohort study. BMC musculoskeletal disorders. 2016;17:29.