

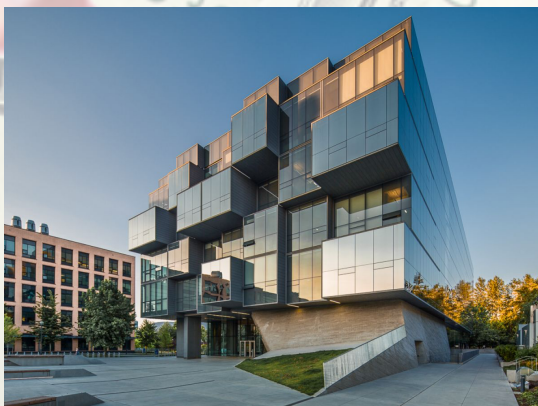
DIVING FOR

James McCormack
BSc (Pharm), PharmD
Professor
Faculty of Pharmaceutical Sciences
University of British Columbia
Vancouver, BC, Canada

MEDICAL PEARLS

therapeuticseducation.org
medicationmythbusters.com

TO GET A HANDOUT GO HERE
<http://therapeuticseducation.org/handouts>



- ☑ Entire salary comes through the UBC Faculty of Pharmaceutical Sciences - also some legal/educational work

- ☑ I have received no honorarium or research money from the drug industry in the last 25 or so years



- ☑ Premium podcast subscription Best Science (BS) Medicine podcast - therapeuticseducation.org

Guidelines would be awesome if they...

Were developed primarily by, and definitely for, the people that ultimately end up using them

Were a credible synopsis of the best available evidence presented in a way that clinicians could easily access and interpret

Allowed patient values and preferences to be taken into account

Wrong guidelines: why and how often they occur

**Primiano Iannone,¹ Nicola Montano,² Monica Minardi,³
James Doyle,³ Paolo Cavagnaro,⁴ Antonino Cartabellotta⁵**

“Unfortunately, depending on how their reliability is measured, up to 50% of guidelines can be considered untrustworthy. This carries serious consequences for patients’ safety, resource use and health economics burden.”

EBM 2017;22:1-3

Simplified lipid guidelines

Prevention and management of cardiovascular disease in primary care

G. Michael Allan MD CCFP Adrienne J. Lindblad ACPR PharmD Ann Comeau MN NP CCN(C) John Coppola MD CCFP
Brienne Hudson MD CCFP Marco Mannarino MD CCFP Cindy McMinis Raj Padwal MD MSc
Christine Schelstraete Kelly Zarnke MD MSc FRCPC Scott Garrison MD PhD CCFP Candra Cotton
Christina Korownyk MD CCFP James McCormack PharmD Sharon Nickel Michael R. Kolber MD CCFP MSc

Can Fam Phy 2015;61:857-67

CLINICAL PRACTICE GUIDELINES

Simplified guideline for prescribing medical cannabinoids in primary care

G. Michael Allan MD CCFP Jamil Ramji Danielle Perry Joey Ton PharmD Nathan P. Beahm PharmD
Nicole Crisp RN MN NP-Adult Beverly Dockrill RN Ruth E. Dubin MD PhD FCFP DCAPM Ted Findlay DO CCFP FCFP
Jessica Kirkwood MD CCFP Michael Fleming MD CCFP FCFP Ken Makus MD FRCPC Xiaofu Zhu MD FRCPC
Christina Korownyk MD CCFP Michael R. Kolber MD CCFP MSc James McCormack PharmD Sharon Nickel
Guillermina Noël MDes PhD Adrienne J. Lindblad ACPR PharmD

Can Fam Phy 2018;64:111-120

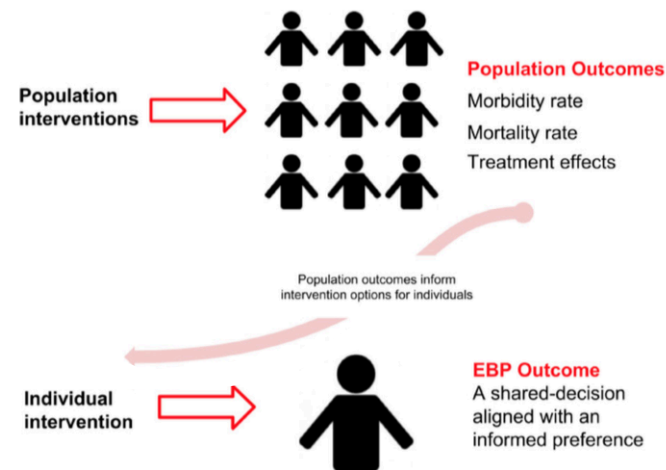


OPEN ACCESS

Shared decision is the only outcome that matters when it comes to evaluating evidence-based practice

James McCormack,¹ Glyn Elwyn²

“in the vast majority of circumstances, the only outcome of relevance for EBP is to measure whether a shared decision was made”



doi:10.1136/ bmjebm-2018-110922

BMJ June 2012



OVERDIAGNOSIS
Harming the healthy

MEDICATIONS

They can only really do 5 things - and only 2 of these are good

Help with symptoms

Reduce risk of future health issues

Cause side effects

Cost money

Be inconvenient

Have A Purpose

You are looking for numbers (%s)



In general who is it for - young/older, primary/secondary

TIME FRAME - 1 dose, 1 day, 1 week, 1 month, 1 year, 1 decade?

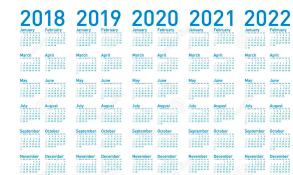
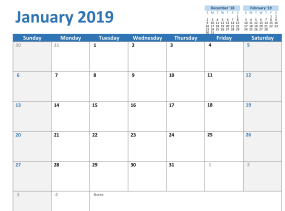
Is it for symptoms?

Clinically relevant endpoints

Is it for prevention?

CVD, fractures, exacerbations, infections

- anything as long as it isn't a surrogate marker (BP, cholesterol, glucose, FEV1, bone density)



Here is how I look if time is limited

(which it almost always is)



Google

ertugliflozin meta-analysis
ertugliflozin meta-analysis



Trusted evidence.
Informed decisions.
Better health.

English

Cochrane.org

Sign In

Title Abstract Keyword



Browse

Advanced search

Cochrane Reviews

Trials

Clinical Answers

About

Help

If no meta-analysis/systematic review - suggests not a lot of published studies

Progress in evidence-based medicine: a quarter century on

Benjamin Djulbegovic, Gordon H Guyatt

“Few clinicians would ever have the skill - or time - to conduct sophisticated assessment of the evidentiary basis for their practice”

Now - “directing clinicians to processed sources of evidence, and aiding decision making by advancing the science of trustworthy clinical practice guidelines that would be available to clinicians at the point of care delivery”

Lancet 2017;390:415–23

Key steps to communicating evidence

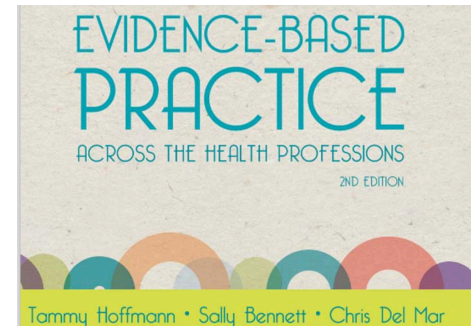
Understand the patient's (and family members') experiences and expectations.

Build partnerships.

Discuss the evidence, including a balanced discussion about uncertainties.

Present recommendations.

Check for understanding and agreement.



Risky Relative Adjectives

HOW

low is low

moderate is moderate

high is high



Evidence-based risk communication

“There is likely no single best method of communicating probabilities to patients but rather several good options with some better suited to certain risk scenarios.”

Ann Intern Med 2014;161:270-80

Recommended approaches

Need a time frame, main endpoints, ask what they know

GENERAL SUGGESTIONS - these are “relative”

use percentages (5%) or natural frequencies (5 out of 100) - BOTH?

use absolute terms

add bar graphs or icon arrays

use incremental risk format with icon arrays in the same array

- **avoid use of NNTs**

if use relative risks add baseline risks

Ann Intern Med 2014;161:270-80

Three “sobering” but very empowering concepts

SYMPTOMS

If a patient seems to be getting a benefit from a medication for symptoms they likely aren't

PREVENTION

If a patient is on a medication for risk reduction (BP, chol, glucose BMD) the benefit they are receiving is likely not large enough for them to make up for the cost, inconvenience and adverse effects

DOSE

If a patient is on a medication they are likely on too high a dose

It's all about figuring out

The Chance of “X”
WITH NO
TREATMENT/TEST
The Chance “X”
WITH
TREATMENT/TEST



Tools For Practice

TOOLS FOR PRACTICE

Total articles found: 233

#233 Drink Up: Increasing Fluid Intake to Prevent Recurrent UTIs

Author(s): Adrienne J Lindblad, Rodger Craig

Publication Date: April 15, 2019

Collection: Tools for Practice

Categories: General, Obstetrics-Gynecology, Urology

Clinical Question: Does increasing water intake prevent recurrent urinary tract infections (UTIs)?

Tags: water, water intake, low fluid, UTI, urinary track, infection, women, female, antibiotics, non-pregnant, premenopausal, cystitis, cranberry juice, vaginal estrogen, oral estrogens, antibiotic prophylaxis, .

 [View Article](#)



Begin Reflective Exercise

(to launch a reflective exercise, you must be logged into GoMainpro)

#232 Muscling out molluscum contagiosum: Which treatments work?

Author(s): Danielle Perry, G. Michael Allan, Nicolas Dugré

Publication Date: April 01, 2019

Collection: Tools for Practice

Categories: Dermatology, General, Infectious Disease

Clinical Question: How effective are commonly used therapies for molluscum contagiosum?

Tags: molluscum contagiosum, lesion, potassium hydroxide, cryotherapy, curettage, cantharidin, imiquimod, virus, immune system, infection, pediatric, immunocompetent, burn, self-limiting

 [View Article](#)



Begin Reflective Exercise

(to launch a reflective exercise, you must be logged into GoMainpro)

#231 Does an ASA a day really keep the doctor away?

Author(s): Paul Fritsch, Michael R Kolber

Publication Date: March 18, 2019

Collection: Tools for Practice

Categories: Cardiology, Gastroenterology, General, Oncology

Clinical Question: Is ASA effective for reducing cardiovascular events in patients without pre-existing cardiovascular disease?

Tags: ASA, cardiovascular, cardiovascular disease, elderly, diabetic, gastrointestinal, cancer, colon, CVD, transfusion, hemodynamic, circulatory system, primary prevention, aspirin, bleeding, bleeds

 [View Article](#)

<https://www.bmj.com/rapid-recommendations>

Dual vs single antiplatelet therapy



The BMJ Practice: [Dual antiplatelet therapy with aspirin and clopidogrel for acute high risk transient ischaemic attack and minor ischaemic stroke](#)

BMJ Research: [Clopidogrel plus aspirin versus aspirin alone for acute minor ischaemic stroke or high risk transient ischaemic attack](#)

Oxygen therapy for acutely ill medical patients



The BMJ Practice: [Oxygen therapy for acutely ill medical patients: a clinical practice guideline](#)

The Lancet research: [Mortality and morbidity in acutely ill adults treated with liberal versus conservative oxygen therapy \(IOTA\): a systematic review and meta-analysis](#)

MAGICapp: [Expanded version of the results](#)

Prostate cancer screening



The BMJ Practice: [Prostate cancer screening with prostate-specific antigen \(PSA\) test: a clinical practice guideline](#)

The BMJ research: [Prostate cancer screening with prostate-specific antigen \(PSA\) test: a systematic review and meta-analysis.](#)

BMJ Open research: [Values and preferences of men for undergoing prostate-specific antigen screening for prostate cancer: a systematic review](#)

The BMJ editorial: [What should doctors say to men asking for a PSA test?](#)

Comparing Treatment Options for Pain: The C-TOP Tool

Neuropathic Pain

Osteoarthritis Pain
Coming Soon

Back Pain
Coming Soon

Medication Options

Amitriptyline
(Elavil®)

Cannabinoids
(Nabiximols, nabilone, medical marijuana)

Duloxetine
(Cymbalta®)

Gabapentin
(Neurontin®)

High-Dose Opioids
(morphine, oxycodone)

Pregabalin
(Lyrica®)

All Treatments
(comparison)

Curious about capsaicin, botox, tramadol, carbamazepine, or venlafaxine for neuropathic pain?
[Click here to learn more.](#)

Meaningful Pain Relief from Amitriptyline

(30% reduction in pain scores)



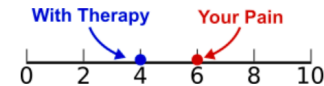
Amitriptyline Benefit	Placebo Benefit	No Benefit
25%	25%	50%

(ranges 13% to 45%)

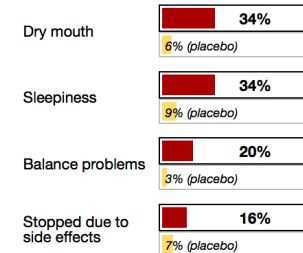
A typical placebo group response seen in pain studies is 25% but this can be adjusted in the [FAQ](#) section.

Meaningful Pain Relief

An example of a 30% reduction in pain scores is a decrease from 6 to 4 on a 10 point pain scale



Amitriptyline Harms



Other Considerations

- Typically taken at bedtime due to sleepiness effects
- Approximate cost (CAD) for 30-day supply (without dispensing fee): **\$1.50 to \$3.50**

<http://pain-calculator.com>

mystudies.org ~300 studies



Load studies



Study Results at Your Fingertips

You want to use evidence in your clinical practice from the landmark studies – those studies that change practice. Your patient comes in and asks you about the latest greatest study. How can you quickly and easily get all that information? **Let MyStudies help.**

You are at a presentation and you start to wonder if the presenter is really telling you everything you need to know about a study. Did they just present relative numbers? Did they only present the benefits with no mention of harms? Did they come up with conclusions that don't really match the results? **MyStudies can help.**

MyStudies beta

+ All 📁 ★ Unread A→Z Year Latest PMID, Title, Keyword, ... ✕

Filter Tags

#add-nptx-values-later #AEentered #checked #checkedAE #checkedJS

#dowewarthis #Jordanentered #not-checked #not-finished #not-working acarbose

ACE-inhibitor ACS acute-MI adenosine-antagonist alegitazar alendronate

alirocumab aliskirenalogipitin alteplase amiodarone amlodipine amoxicillin

angiography anti-platelets antioxidants antipsychotics aorticvalve apixaban ARBs

arrhythmias asa aspirin atenolol atrial-fibrillation beta-blocker bloodpressure

bococizumab budesonide CABG calcium calcium-channel-blocker Canakinumab

candesartan captopril cardiovascular CETPinhibitors chelation chlorothalidone

cholesterol clobifrate clopidogrel clopidogrelprasugrel COPD CRP dabigatran

dalcetrapib dalteparin degludec denosumab diabetes digoxin dronedarone dvt

elderly Empagliflozin enalapril enoxaparin ESRD estrogen evacetrapi

evolucumab exercise ezetimibe fibrates folicacid formoterol fractionalflowreserve

Gliblila gliargine glaucoma glitazones HDL heart-failure heartfailure heparin

homocysteine hormone HRT hyralazine hydrochlorothiazide hypertension

ibandronate indicatcel insulin intensive-BP-control intensive-glucose-lowering

intensive-lifestyle-interventio ipratropium irbesartan isosorbide-dinitrate kidney-disease

lasar liraglutide LMWH losartan LRTI mediterranean metformin metoprolol

mometasone multivitamin nephropathy neprilysin niacin nitrates

☆ REDUCE-IT

Cardiovascular Risk Reduction with Icosapent Ethyl for Hypertriglyceridemia.

The New England journal of medicine, 2019

☆ CABANA

Effect of Catheter Ablation vs Antiarrhythmic Drug Therapy on Mortality, Stroke, Bleeding, and Cardiac Arrest Among Patients With Atrial Fibrillation

JAMA, 2019

☆ AUGUSTUS

Antithrombotic Therapy after Acute Coronary Syndrome or PCI in Atrial Fibrillation.

The New England journal of medicine, 2019

apixaban
aspirin
wارفارين

☆ CREDESCENCE

Canagliflozin and Renal Outcomes in Type 2 Diabetes and Nephropathy

New England journal of medicine, 2019

☆ VITAL Omega-3

Martine n-3 Fatty Acids and Prevention of Cardiovascular Disease and Cancer.

The New England journal of medicine, 2018

☆ ODYSSEY OUTCOMES

Alirocumab and Cardiovascular Outcomes after Acute Coronary Syndrome.

The New England journal of medicine, 2018

#checked
alirocumab
PCSK9

☆ VITAL Vitamin D

Vitamin D Supplements and Prevention of Cancer and Cardiovascular Disease.

The New England journal of medicine, 2018

☆ ASCEND

Effects of Aspirin for Primary Prevention in Persons with Diabetes Mellitus

The New England journal of medicine, 2018

☆ DECLARE-TIMI 58

Dapagliflozin and Cardiovascular Outcomes in Type 2 Diabetes.

The New England journal of medicine, 2018

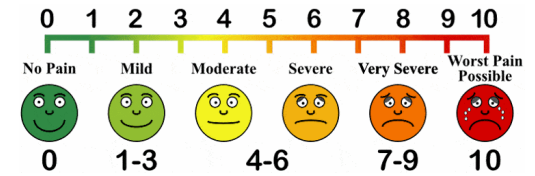
All the large RCTs evaluating the impact of glucose lowering medications on CVD Outcomes

RCTs evaluating the impact of medications on CVD outcomes in T2DM						
YEAR	NAME		MEDICATION	RESULT	OUTCOME CHANGED	ABSOLUTE DIFFERENCE/TIME
1970	UGDP	SU	tolbutamide (Orinase)	NEGATIVE	CVD mortality	↑ 8%/5 years
1971		BG	phenformin (DBI)	NEGATIVE	Mortality	↑ 6%/5-8 years
1976		SU	tolbutamide (Orinase)	NEGATIVE	Fatal MI	↑ 5%/5 years
1982		IN	insulin	NEUTRAL		
1998	UKPDS 33/34	IN,SU	insulin, chlorpropamide, glyburide/glibenclamide, glipizide	NEUTRAL		
1998		IN,SU,BG	metformin, insulin, chlorpropamide, glyburide/glibenclamide, glipizide	NEUTRAL except POSITIVE for metformin	Mortality MI	↓ 7%/11 years ↓ 6%/11 years
2003	STOP-NIDDM	OTH	acarbose (Precose)	POSITIVE	MI	↓ 1.5%/3 years
2005	PROACTIVE	GLIT	pioglitazone (Actos)	POSITIVE	MI	↓ 1.5%/3 years
2007	RECORD	GLIT	rosiglitazone (Avandia)	NEGATIVE	Heart failure	↑ 1%/4 years
2012	ORIGIN	IN	insulin	NEUTRAL		
2013	EXAMINE	DPP4	alogliptin (Nesina)	NEUTRAL		
2014	SAVOR-TIMI 53	DPP4	saxagliptin (Onglyza)	NEGATIVE	Heart failure	↑ 1%/2 years
2014	ALECARDIO	OTH	aleglitazar	NEUTRAL		
2015	ELIXA	GLP	lixisenatide (Adlyxin)	NEUTRAL		
2015	TECOS	DPP4	sitagliptin (Januvia)	NEUTRAL		
2015	EMPA-REG	GLIF	empagliflozin (Jardiance)	POSITIVE	Mortality Heart failure	↓ 2.5%/3 years ↓ 1.5%/3 years
2016	SUSTAIN 6	GLP	semaglutide (Ozempic)	POSITIVE	Combined outcome	↓ 2%/2 years
2016	LEADER	GLP	liraglutide (Victoza)	POSITIVE	Mortality Combined outcome	↓ 1%/4 years ↓ 2.5%/4 years
2017	CANVAS	GLIF	canagliflozin (Invokana)	POSITIVE	Combined outcome Heart failure Amputations	↓ 2%/3.5 years ↓ 1%/3.5 years ↑ 1%/3.5 years
2017	EXSCEL	GLP	exenatide (Byetta)	NEUTRAL		
2017	ACE	OTH	acarbose (Procoese)	NEUTRAL		
2017	Omarigliptin	DPP4	omarigliptin	NEUTRAL		
2018	HARMONY	GLP	albiglutide (Tanzeum)	POSITIVE	Combined outcome	↓ 2%/2 years
2018	CARMELINA	DPP4	linagliptin (Tradjenta)	NEUTRAL		
2018	DECLARE-TIMI 58	GLIF	dapagliflozin (Farxiga)	POSITIVE	Combined outcome (primarily heart failure)	↓ 1%/4 years
2019	REWIND	GLP	dulaglutide (Trulicity)	POSITIVE	Combined outcome Renal outcomes	↓ 1.5%/5.4 years ↓ 2.5%/5.4 years
2019	PIONEER 6	GLP (oral)	semaglutide (Ozempic)	POSITIVE	CVD mortality Mortality	↓ 1%/1.5 years ↓ 1.5%/1.5 years
2019	CREDENCE		canagliflozin (Invokana)	POSITIVE	Combined CVD outcome Combined renal outcome outcomes	↓ 2.5%/2.6 years ↓ 3%/2.6 years



Symptom
Pearls

Symptoms



Scales - VAS, QOL, SGRQ - then what is the MICD

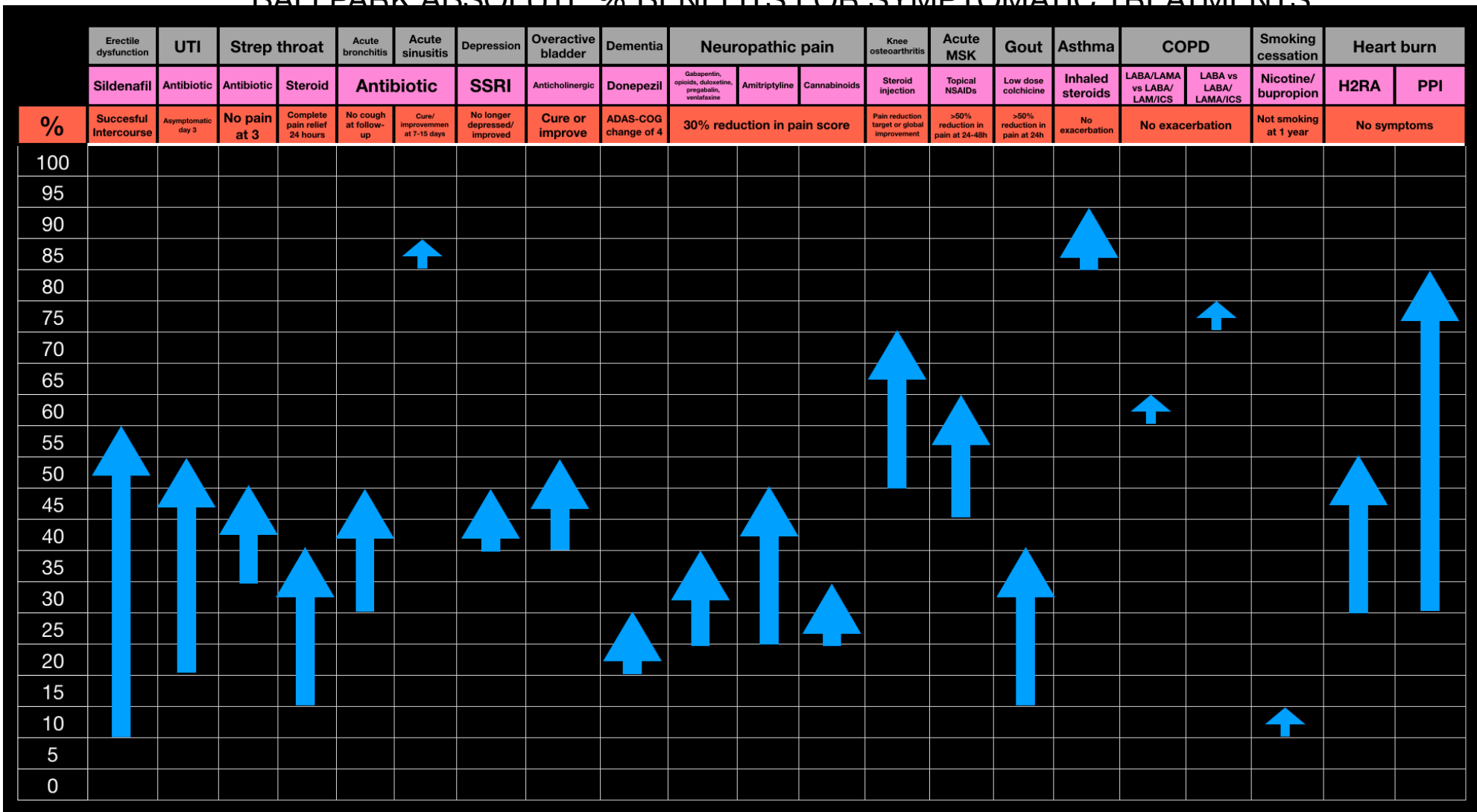
% of people who benefit in the treatment arm - that will be what you see in practice over placebo

% of people who benefit in the placebo arm - subtract that from the treatment to see how many actually benefit from the medication

Head-to-head studies are relatively uncommon

6-8 weeks	No longer depressed
Medication	50%
Placebo	40%
Medication benefit	$50 - 40 = 10\%$
If person responds, the chance it is the medication	$10 / 50 = 20\%$

BALL PARK ABSOLUTE % BENEFITS FOR SYMPTOMATIC TREATMENTS





Prevention Pearls

If you were to treat two risk factors (glucose, cholesterol, blood pressure) in 100 patients for a lifetime (40 years) how many people do you think would derive a clinical benefit?

0-10%
11-20%
21-30%
31-40%
41-50%
51-60%
61-70%
71-80%
81-90%
91-100%

Math 101

- actually grade 5

REMEMBER - X% of Y - “OF” means multiply

WHAT IS THE
ABSOLUTE
BENEFIT %?

	Relative benefit (%)				
BASELINE RISK (%)	10	15	20	25	30
10	1	1.5	2	2.5	3
15	1.5	~2.5	3	~4	4.5
20	2	3	4	5	6
30	3	4.5	6	~8	9

WHAT IS
THE NNT?

Absolute benefit	0.5%	1%	2%	3%	4%	5%	6%	7%	8%	9%	10%
NNT	200	100	50	33	25	20	~17	~14	~13	~11	10

Math 101 - actually grade 5

Math 101 - actually grade 5

REMEMBER - X% of Y - "OF" means multiply

WHAT IS THE ABSOLUTE BENEFIT %?

	Relative benefit (%)				
BASELINE RISK (%)	10	15	20	25	30
10					
15					
20					
30					

WHAT IS THE NNT?

[illegible]

Ballpark Risks (CVD, fractures etc)

<https://therapeuticseducation.org/tools>

The Absolute CVD Risk/Benefit Calculator

Framingham QRISK®2-2014 ACC/AHA ASCVD

Heart disease + stroke + intermediate classification Heart attack + stroke

Age: 50 years

Gender: Male

Smoker: Yes

Diabetes: Yes

Systolic Blood Pressure: 120 mmHg

Total Cholesterol: 3 mmol/L

HDL Cholesterol: 1.3 mmol/L

Family History of Early CHD: 0%

Relative Benefit: 0%

Risk Time Period: 10 years

97.6% No event

2.4% Total with an event

0.0% Number who benefit from treatment

2.4% Baseline events using baseline factors alone

0.0% Additional events "caused" by risk factors

Switch to "Basic" View

cvdcalculator.com

Stroke Risk (CHA2DS2-VASc)

Age: 65-74

TIA or stroke (at any time in the past): No

Prior MI, peripheral artery disease, or aortic plaque: No

Female: No

CHA2DS2-VASc SCORE (0-9): 3

Major Bleeding Risk (HAS-BLED*)

Abnormal renal/liver function (creatinine <30 or bilirubin >2x ULN): No

Hypertension (SBP >160 mmHg): No

Abnormal liver function (bilirubin >2x ULN): No

History of major bleeding (any cause): No

HAS-BLED SCORE (0-9): 1

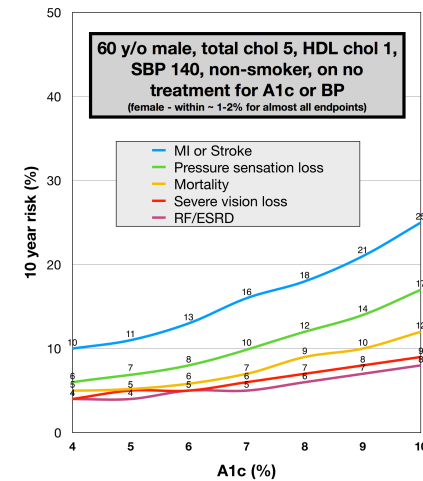
Which therapy options to HIDE?

Aspirin, Aspirin+Clopidogrel, Warfarin, Dabigatran, Rivaroxaban, Apixiban, Edoxaban

PERCENT PER YEAR

	annual risk of stroke/thromboembolism	annual risk of major bleeding (intracranial bleeding, bleeding requiring hospitalization, high degree of >20 g/L, or need for transfusion secondary to bleed)
NO THERAPY	4.3%	0.6%
ASPIRIN	3.4%	1.1%
WARFARIN	1.4%	2.2%
DABIGATRAN 110	1.4%	1.8%
DABIGATRAN 150	0.9%	2.2%
RIVAROXABAN	1.4%	2.2%
APIXABAN	1.1%	1.5%

sparctool.com



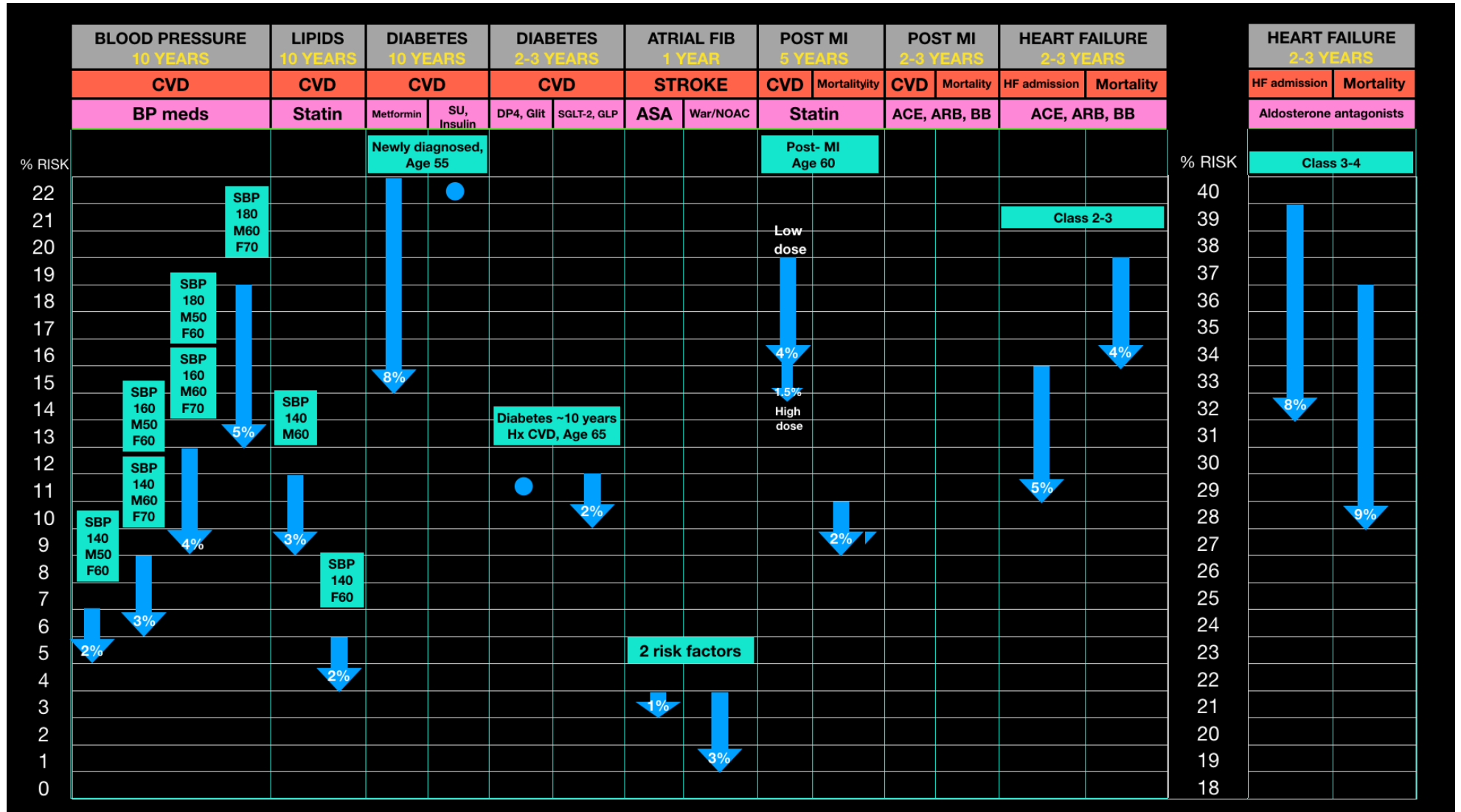
<https://sanjaybasu.shinyapps.io/recodesi/>

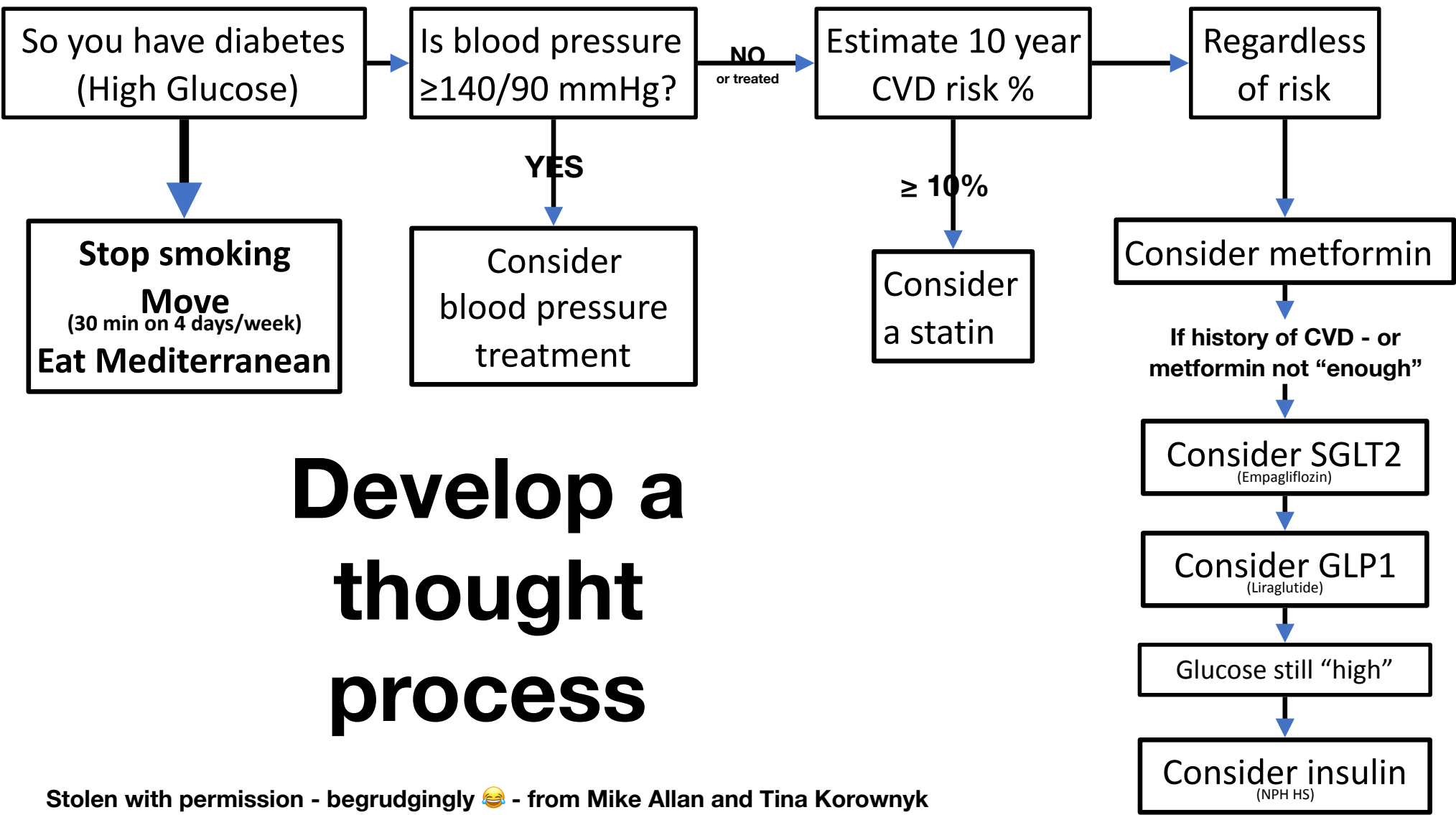
RISK FACTORS t-score	Zero			One			Two		
	-1.5	-2.5	-3.5	-1.5	-2.5	-3.5	-1.5	-2.5	-3.5
Female									
50	4	5/1	9/4	6	8/2	14/7	8	12/3	21/11
60	7	10/2	16/6	10/1	14/3	23/9	14/1	20/5	32/14
70	9/1	13/3	21/7	12/1	18/4	30/11	16/2	25/6	41/16
80	13/3	18/6	29/14	17/6	26/12	40/24	24/10	35/20	52/37
Male									
50	4	5/2	11/6	5	8/3	16/10	8/1	12/5	24/16
60	6/1	9/3	15/8	8/1	12/4	21/11	12/2	18/6	29/17
70	6/2	10/4	16/8	9/3	14/6	22/13	12/4	19/10	31/20
80	7/3	11/5	16/9	11/5	16/9	23/16	15/9	22/15	32/25

BALLPARK RELATIVE % BENEFITS FOR PREVENTATIVE TREATMENTS

	Lifestyle	Cholesterol	Blood pressure	Glucose	A fib	Heart failure	Osteoporosis	Flu	Zoster	HPV
RRR%	Cardiovascular events				Stroke	Mortality	Fractures	Infection		
100	Stopping smoking (obviously no RCTs)									
95									Zoster Vaccine	HPV Vaccine
90										
85										
80										
75								Flu vaccine		
70										
65					Warfarin/NOACS					
60										
55										
50			Blood pressure diabetes							
45										
40										
35		Statins		Metformin?						
30	Mediterranean		Blood pressure				Bisphos, monoclonal			
25	Physical Activity				Aspirin	ACEI, BB, Aldo antag				
20										
15		PCSK9 Monoclonal antibodies		SGLT2, GLP						
10	Aspirin						Calcium Vitamin D			
5		Ezetimibe								
0		Fibrate, niacin		DPP4, SU, insulin, glitazone						

BALLPARK ABSOLUTE % BENEFITS FOR PREVENTATIVE TREATMENTS





**Develop a
thought
process**

Stolen with permission - begrudgingly 😂 - from Mike Allan and Tina Korownyk

A Reasonable Side Effect List

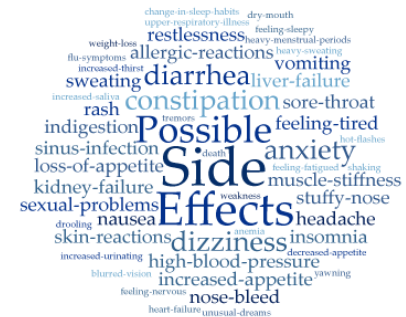
An unsolvable problem?



They are not captured well/completely/understandably in studies - but likely the best we have

Rarely can we figure out rare side effects

Many monographs, books, studies, websites just list a variety of symptoms, often with no numbers, no context, no idea of the duration, severity, frequency, statistical significance?



ORIGINAL ARTICLE

Marine n-3 Fatty Acids and Prevention of Cardiovascular Disease and Cancer

This is a !@#\$\$%
5.3 year study

Table S7. Hazard Ratios (HR) and 95% Confidence Intervals (CI) for Safety and Adverse Events by Randomized Assignment to Omega-3 Fatty Acids (n-3) compared to Placebo

	No. of Events				
	n-3 (N = 12,933)	Placebo (N = 12,938)			
Outcome			HR	95% CI	P-value
Monitored safety conditions					
Gastrointestinal bleeding	370	374	0.99	0.86-1.14	0.89
Blood in urine	919	874	1.06	0.96-1.16	0.25
Easy bruising	3443	3399	1.02	0.97-1.07	0.48
Frequent nosebleeds	465	491	0.95	0.83-1.07	0.40
Kidney failure or dialysis	85	88	0.97	0.72-1.30	0.82
Other symptoms and side effects					
Stomach upset or pain	4887	4843	1.01	0.97-1.05	0.72
Nausea	3558	3550	1.00	0.96-1.05	0.94
Constipation	5184	5111	1.01	0.97-1.05	0.51
Diarrhea	5599	5580	1.00	0.97-1.04	0.77
Skin rash	3331	3367	0.99	0.94-1.03	0.58
Bad taste in mouth	2240	2245	1.00	0.95-1.06	0.92
Increased burping	2217	2158	1.03	0.97-1.10	0.29

How are these not 100%

	A	B	N	O	P	Q	R	S	T	U	V
1		Drug class (ENG)	Statins		Proton pump inhibitor		Proton pump inhibitor		Proton pump inhibitor		Proton p
2		Drug class (FR)	Statines		Inhibiteur de la pompe à protons		Inhibiteur de la pompe à protons		Inhibiteur de la pompe à protons		Inhibiteu
3		Drug name (generic - ENG)	Lovastatin		Lansoprazole		Esomeprazole		Omeprazole		Dexlanso
4		Other drug synonyms (ENG)	Apo-Lovastatin; Co Lovastatin; Pro-		Apo-Lansoprazole; Mylan-Lansoprazole; pr		ACT Esomeprazole; Apo-Esomeprazole; Myli		Apo-Omeprazole; Auro-Omeprazol		Dexilant
5		Drug name (generic - FR)	Lovastatine		Lansoprazole		Esomeprazole		Omeprazole		Dexlanse
6		Other drug synonyms (FR)	Apo-Lovastatin; Co Lovastatin; Pro-		Apo-Lansoprazole; Mylan-Lansoprazole; pr		ACT Esomeprazole; Apo-Esomeprazole; Myli		Apo-Omeprazole; Auro-Omepra		Dexilant
7		Drug name (brand)	Mevascor		Prevacid		Nexium		Lozac, Olex		Dexilant
8	Side effect category		%	Category	%	Category	%	Category	%	Category	%
9		Headache	7.7*	1-10%	2.1*	1-10%	2.1*	1-10%	2.1*	1-10%	2.1*
10		Seizure		0		0		0		0	
11	Neurologic	Pins and needles sensation	1	1-10%				0		0	
12		Insomnia		0				0		0	
13		Trembling						0		0	
14		Angina		0 >1		1-10%	>1	1-10%	>1	1-10%	>1
15		Suicidal thoughts		0			0	0		0	
16		Depression		0			0	0		0	
17		Strange dreams		0			0	0		0	
18		Confusion		0			0	0		0	
19	Psychiatric	Anxiety		0	1.7	1-10%	1.7	1-10%	1.7	1-10%	1.7
20		Difficulty falling asleep	1.9	1-10%	2.1	1-10%	2.1	1-10%	2.1	1-10%	2.1
21		Anxiety disorders	2.3	1-10%	<1	0,1-1%	<1	0,1-1%	<1	0,1-1%	<1
22		Nervousness		0			0	0		0	
23		Feeling jittery						0		0	
24		Difficulty breathing		0	1.1	1-10%	1.1	1-10%	1.1	1-10%	1.1
25	Respiratory	Throat or nose infection		0	8.5	1-10%	8.5	1-10%	8.5	1-10%	8.5
26		Coughing		0			0	0		0	
27		Fainting						0		0	
28		Cold fingers & toes						0		0	
29		High blood pressure		0	2.6	1-10%	2.6	1-10%	2.6	1-10%	2.6
30		Flushing		0			0	0		0	
31		High blood pressure		0			0	0		0	
32	Cardiovascular	Dizziness when standing up		0			0	0		0	
33		Chest pain		0	>1	1-10%	>1	1-10%	>1	1-10%	>1
Display a menu CAMILLE Sheet Explore											



Your Medication's Side Effects

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Language: [English \(EN\)](#)

[I think I have a side effect](#)

[Search for another medication](#)

Atorvastatin (Lipitor)

This medications is generally used to treat the following conditions:

- None reported
- Constipation

— Diarrhea

— Headache

— Liver damage

— Muscle and/or joint pain

— Nausea or vomiting

— Stomach ache
- High blood sugar

People aged over 65 should be aware that this medication increases the risk of:

- Diarrhea

— Falls

Costs



Generic Name	Brand name	Strength	Dosing	90 Day Cost (unless otherwise noted)	Coverage
HYPOGLYCEMIC AGENTS					
Biguanides					
Metformin	Glucophage	500mg	2 BID	\$30	BC / IA covered
Metformin SR	Glumetza SR	1000mg	2 QD	\$255	NC by BC or IA
Sulfonylureas					
Glyburide	Diabeta	5mg	BID	\$25	BC / IA covered
Gliclazide, Gliclazide MR	Diamicon/MR	80mg/30mg MR	BID, 2 QD MR	\$30	BC / IA covered
Meglitinides					
Repaglinide	Gluconorm	1mg	TID	\$35	BC / IA covered
Dipeptidylpeptidase-4 Inhibitors (DPP-4)					
Linagliptin	Trajenta	5mg	QD	\$265	SA req'd for BC and IA
Saxagliptin	Onglyza	5mg	QD	\$295	SA req'd for BC and IA
Sitagliptin	Januvia	100mg	QD	\$310	SA req'd for BC and IA
Sodium Glucose Cotransporter 2 (SGLT2) Inhibitors					
Empagliflozin	Jardiance	10mg	QD	\$270	SA req'd for BC and IA
Canagliflozin	Invokana	100mg	QD	\$280	SA req'd for BC and IA
Glucagon-like Peptide 1 Agonist (GLP-1)					
Liraglutide	Victoza	1.2mg SQ	QD	\$575	NC by BC or IA
Liraglutide	Victoza	1.8mg SQ	QD	\$855	NC by BC or IA
Insulin (Prices may vary between pharmacies, relative differences likely consistent. Max allowable price for 1500 Units of penfill insulin)					
Regular insulin	Novolin Toronto/ Humulin R	100U/mL	As dir	\$60	BC / IA covered
Long-acting insulin	Novolin NPH/Humulin N	100U/mL	As dir	\$65	BC / IA covered
Rapid-acting insulin	Novorapid/Humalog	100U/mL	As dir	\$75	BC / IA covered
Basal insulin (Glargine)	Basaglar	100U/mL	As dir	\$90	BC covered, NC by IA
Basal insulin (Glargine)	Toujeo	300U/mL	As dir	\$110	NC by BC or IA
Basal insulin (Glargine)	Lantus	100U/mL	As dir	\$115	BC / IA covered
Basal insulin (Detemir)	Levemir	100U/mL	As dir	\$130	BC / IA covered
OBESITY					
Orlistat	Xenical	120mg	TID	\$505	NC by BC or IA
Liraglutide	Saxenda	3mg SQ	QD	\$1,165	NC by BC or IA
LEGEND: BC = Alberta Blue Cross, IA = Indian Affairs, NC = Not covered, SA = special authorization, SR = sustained release, OTC = over the counter, SQ = subcutaneous injection, SS=Social Services					

<https://www.acfp.ca/wp-content/uploads/2018/03/ACFPPrisingDoc2018.pdf>

Inconvenience

Get the prescription



Fill the prescription



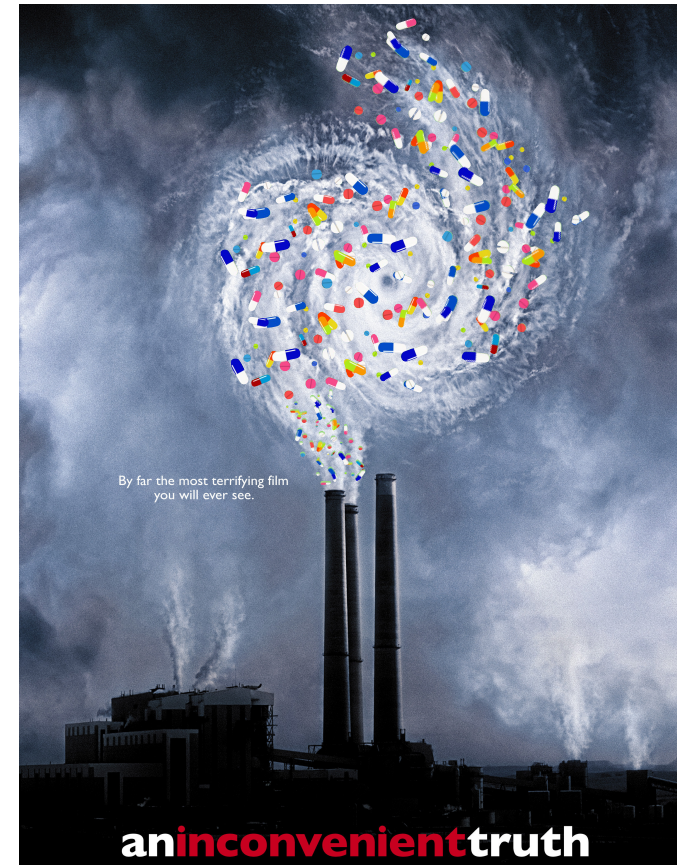
Pay for the prescription



Take the prescription

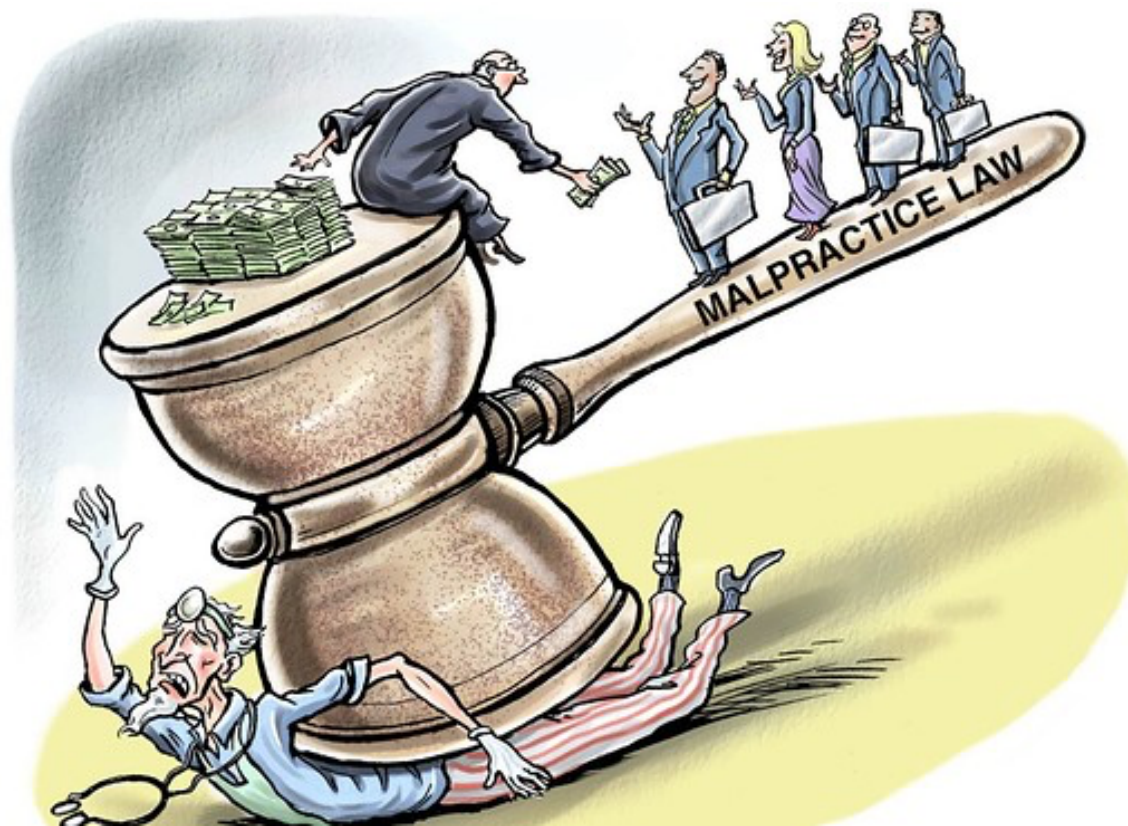


Labelling/worry



On a scale of 0-10, how much do you worry about legal issues if you don't follow guidelines?

0
1
2
3
4
5
6
7
8
9
10



RESEARCH ARTICLE

Open Access

Can shared decision-making reduce medical malpractice litigation? A systematic review

Marie-Anne Durand^{1,2*}, Benjamin Moulton^{3,4,5}, Elizabeth Cockle², Mala Mann⁶ and Glyn Elwyn^{1,7}

“There is insufficient evidence to determine whether or not shared decision-making and the use of decision support interventions can reduce medical malpractice litigation. Further investigation is required.”

Two or more reasonable treatment or screening options

Shared decision-making model

Defensive medicine model

ADVERSE OUTCOME OCCURS

Choice made does **NOT**
MEET the “standard of care”

Choice made **MEETS** the
“standard of care”

Choice made **MEETS**
the “standard of care”

Choice made does **NOT**
MEET the “standard of care”

Discussion
NOT
documented

Discussion
documented
in notes

Decision
aid used

Discussion
NOT
documented

Discussion
documented
in notes

Decision
aid used

**Plaintiffs lawyer argues risks and
benefits should have been discussed**

Low to
medium
risk

No medico
legal
protection

No medico
legal
protection

Medium
risk

Low
risk

Low to
medium
risk

Low
risk

Low
risk

Defensive model (guidelines/standard of care)

NEVER get to a low litigation risk

Low to
medium
risk

Reducing litigation risk 2 THINGS to DO

Shared decision-making model

- 1) Use a decision aid
- 2) Document decision

Low
risk