

Current management of NASH

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Disclosures

Partner of HEPATOSCAN Medical Services® - Commercial interest in transient liver elastography using FibroScan®.

- Most common liver disease in western countries
- Hepatic manifestation of metabolic syndrome (MetS)

✓ World prevalence: 6-46%

✓ Obesity III: 90%

✓USA: 10-46%; NASH: 3-5%

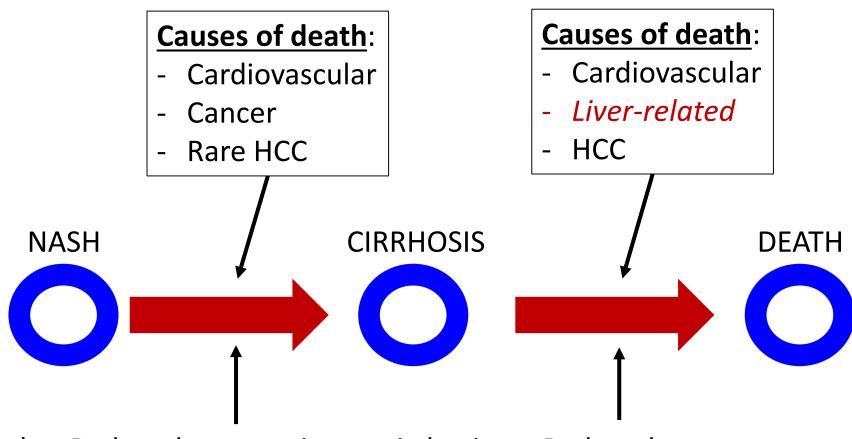
✓ Brazil: 19-35%

Vernon, et al. Aliment Pharmacol Ther 2011 Williams, et al. Gastroenterology 2011 Lazo, et al. Am J Epidemiol 2013 Parise, et al. 2003 Xarnikowski, et al. 2007

Major factors related to NAFLD:

- ✓ Obesity
- ✓ Type 2 diabetes (T2D)
- ✓ Metabolic syndrome (MetS)

NAFLD clinical scenarios

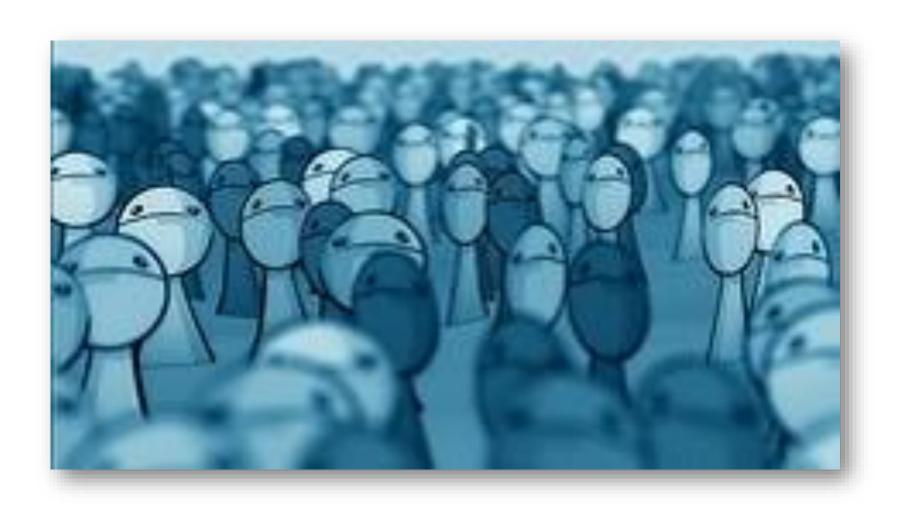


Goals: Reduced progression to cirrhosis

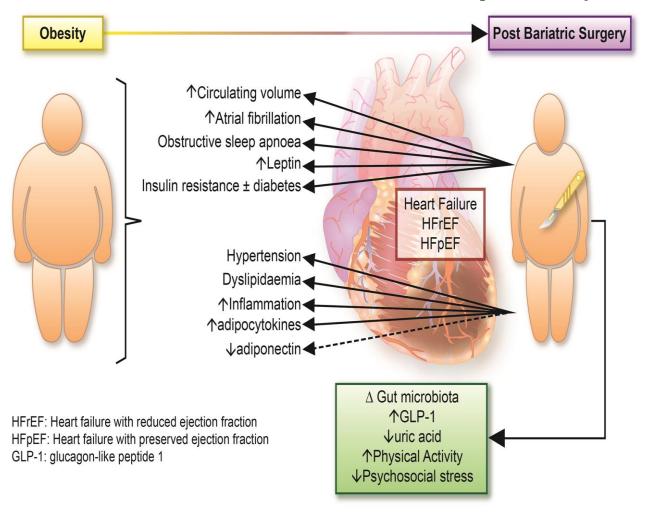
Reduced outcomes



Diet and lifestyle changes



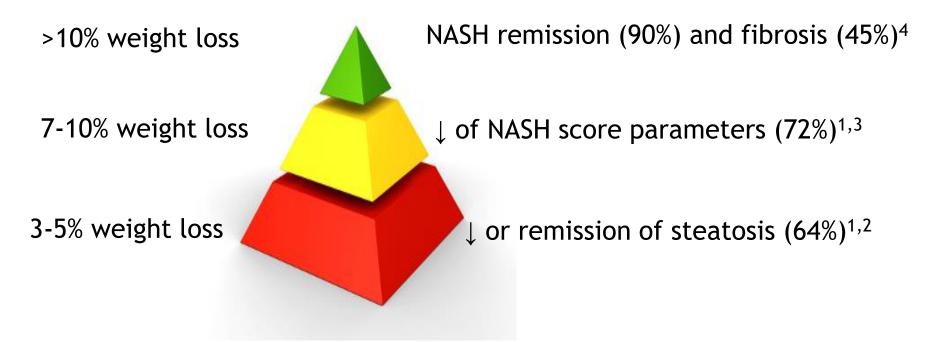
Weight loss benefits key outcomes associated with excess adiposity



Finer N. European Heart Journal, 2019

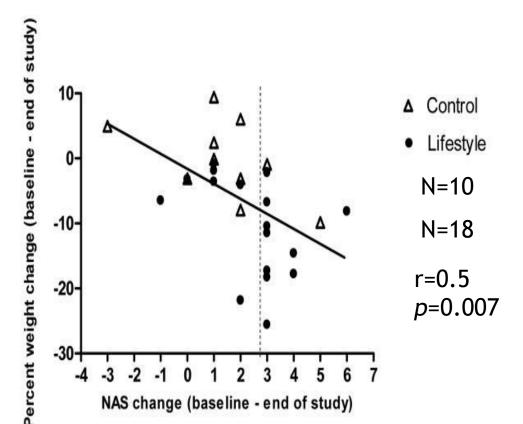
Weight loss: the cornerstone of the treatment

3 randomized and 1 cohort studies (N = 293)



- 1. Harrison et al. Hepatology 2009; 2. Wong et al. JHepatol. 2013
- 3. Promrat et al. Hepatology 2010; 4. Vilar-Gomez et al. Gastroenterology 2015

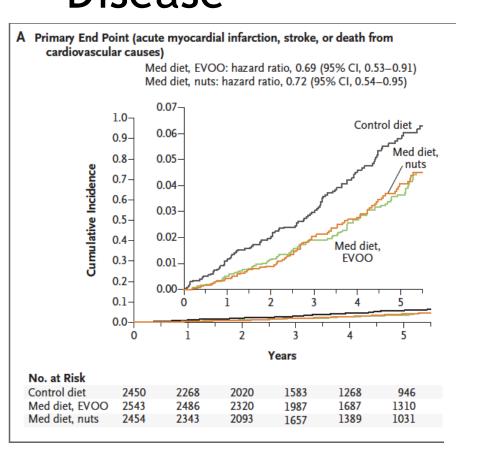
Effect of combined diet and exercise on liver histology in NASH



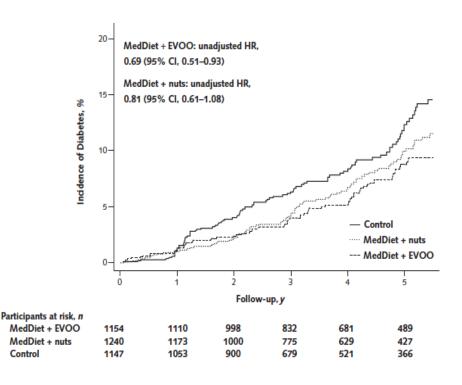
NAS change (baseline - end of study)

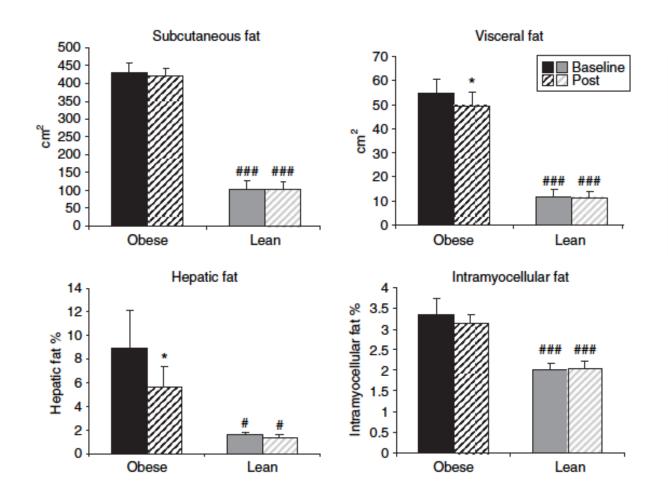
- 48 week diet and exercise intervention
- Steatosis improves with minimal wt loss but benefits to Inflammation/ballooning may require >7% body wt loss
- No effect on fibrosis

Control



N = 7447Median FU 4,8y





N = 29 Hispanic adolescents (15 obese/14 lean) 12-week controlled aerobic exercise program

Inf	tra-	he	pat	ic	fat
			~~		

	Exercise	Con	trol S	td. Mean Di		W-1	
Study	n	n		IV, Fixed, 95% CI		Weight	
Hallsworth 2011	11	8	_	0		3.0%	
Keating 2015 Group 1	12	12			-	5.3%	
Keating 2015 Group 3	12	12			-	5.6%	
Sullivan 2012	12	6			\neg	4.0%	
Keating 2015 Group 2	12	12			-	6.0%	
Lee 2013 (Aerobic vs control)	16	12				7.1%	
Lee 2012 (Resistance vs control)	16	13			_	7.5%	
Lee 2012 (Aerobic vs centrel)	16	13			-	7.6%	
Pugh 2013	6	5			+	2.9%	
Zelber-Sagi 2014	33	31		-		17.6%	
Lee 2013 (Resistance vs control)	16	12			-	7.6%	
Johnson 2009	12	7		_		5.0%	
Larson-Meyer 2008	12	12		-		6.9%	
Shojaee-Moradie 2007	10	7		_	-	4.7%	
Shah 2009	9	9		_	\rightarrow	5.2%	
Tamura 2005	7	7		-	-	4.0%	
Total (95% CI)	-0.69 [-0.90	, -0.48]		. <	>	100.0%	
			-4	-2	ó	2	4
			Favours	exercise		Favours contr	ol

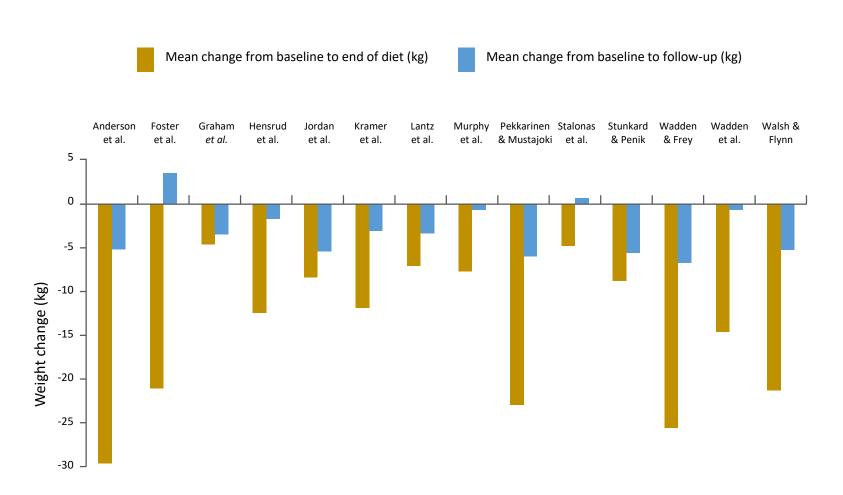
N= 1644
Physical activity-only
systematic review of 28
randomized clinical
trials

Heterogeneity: $Chi^2 = 21.22$, df = 15 (P = 0.13); $I^2 = 29\%$

Test for overall effect: Z = 6.43 (P < 0.00001)

The effect of physical activity on hepatic liver fat content was more prominent in young patients and patients with a higher baseline BMI

Maintaining weight loss is challenging



N = 540 FU = 7y

 NAFLD patients followed in a outpatients clinic of University Hospital - Federal University of Rio de Janeiro (HUCFF-UFRJ)

√7% weight maintenance

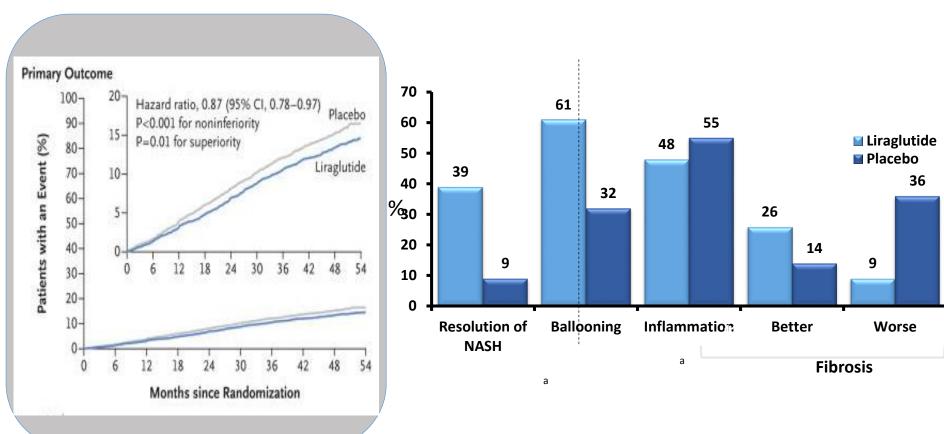
√49% increased weight

✓ < 10% of total with reduction over 7% of initial weight
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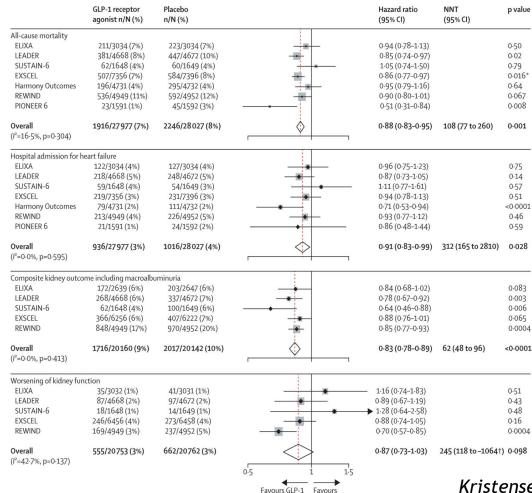
Management of obesity

Mechanism of action				Mean percentage weight loss		Advantages	Disadvantages
		USA	European Union	Placebo	Drug		
hentermine; 15–30 mg orally	Sympathomimetic	For short- term use	No	Not stated in label	Not stated in label	Inexpensive	Side-effect profile; no long-term data*
Orlistat; 120 mg orally three times a lay before meals	Pancreatic lipase inhibitor	Yes	Yes	-2.6%†	-6.1%†	Not absorbed; long-term data*	Modest weight loss; side-effect profile
orcaserin; 10 mg orally twice a day	5-HT _x serotonin agonist with little affinity for other serotonergic receptors	Yes	No	-2.5%	-5.8%	Mild side-effects; long-term data*	Expensive; modest weight loss
Phentermine/ topiramate ER; 7-5 mg/ 46 mg or 15 mg/ 92 mg orally ndicated as rescue (requires titration)	Sympathomimetic anticonvulsant (GABA receptor modulation, carbonic anhydrase inhibition, glutamate antagonism)	Yes	No	-1-2%	-7.8% (mid- dose) -9.8% (full dose)	Robust weight loss; long-term data*	Expensive; teratogen
Naltrexone SR/ bupropion SR; 12 mg/360 mg orally (requires itration)	Opioid receptor antagonist; dopamine and noradrenaline reuptake inhibitor	Yes	Yes	-1.3%	-5-4%	Reduces food craving; long-term data*	Moderately expensive; side-effect profile
iraglutide; 3·0 mg Injection (requires itration)	GLP-1 receptor agonist	Yes	Yes	-3%	-7·4% (full dose)	Side-effect profile; long-term data*	Expensive; injectable

GLP-1 receptor agonists have the potential for cardio-metabolic as well as liver-benefits



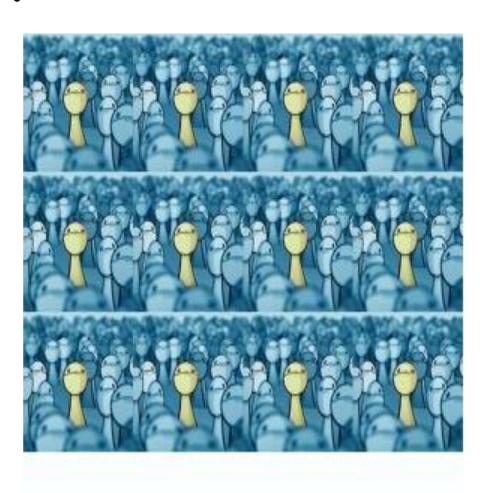
GLP-1 agonists reduce all cause mortality, hospitalization for heart failure and improve renal status in DM



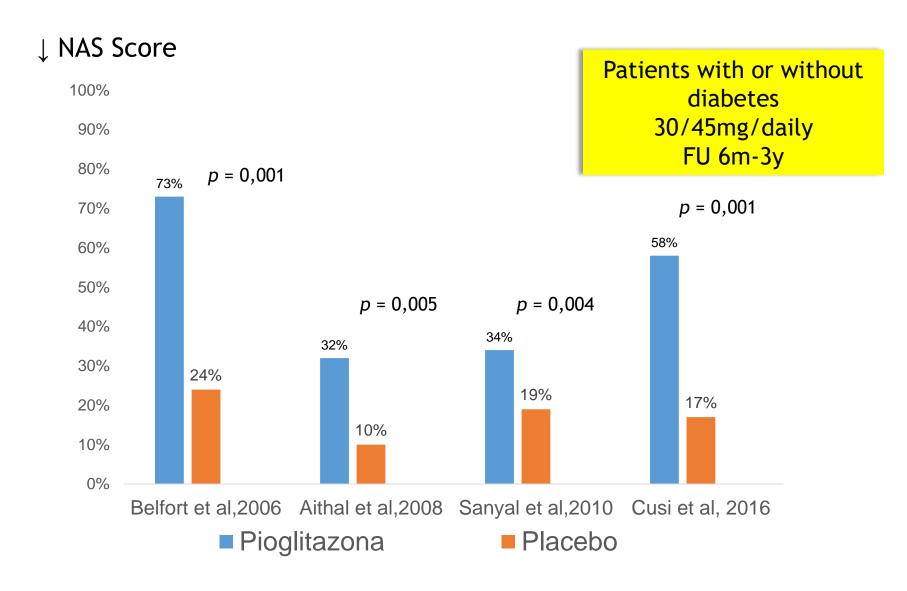
receptor agonist placebo

	N	Follow-up (m)	NAFLD Outcomes
Mathurin et al, 2009 *	381	50	48% NASH improvement Early stage fibrosis at 5y
Taitano et al,2015	160	31	90% NASH remission 60% regression of fibrosis
Lassailly et al,2015	109	12	85% NASH remission 34% regression of fibrosis
Manco et al,2016	93	12	100% NASH remission 90% regression of fibrosis

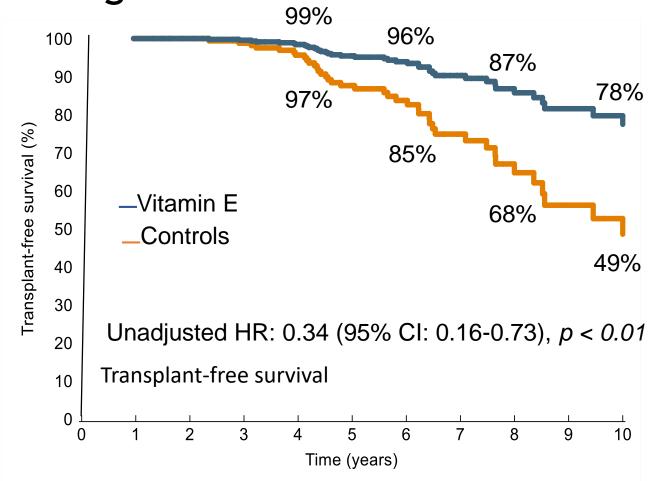
Which would be the best candidates for pharmacological treatment for NAFLD?



Randomized Studies with Pioglitazone

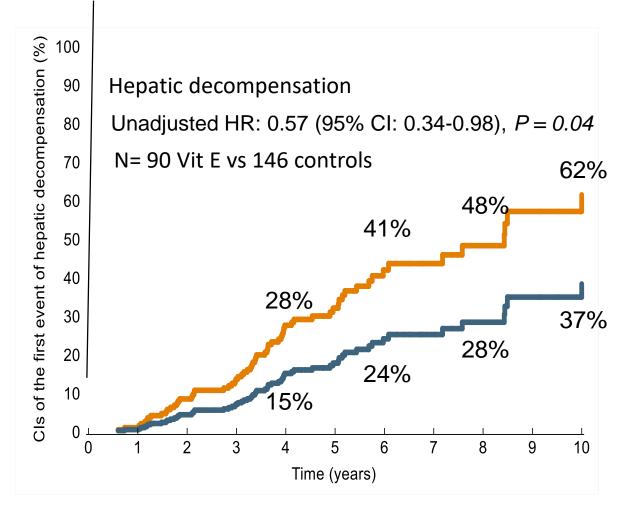


Vitamin E improves transplant free survival and decompensation rates in NASH with stage 3 or 4 fibrosis



Vitamin E improves transplant free survival and decompensation rates in NASH with

F3/4



N = 132377 35-79y FU = 8y Non diabetic individuals

✓ A total of 6,555 incident diabetes (3,734 men and 2,821 women) were identified, on average, over 5.8 years of follow-up.

√The risk of incident diabetes was significantly associated with
NAFLD [HR=2.08 (men) and 2.65 (women)].

✓ Elevated ALT, AST, GGT and ALP were also significantly associated with the increased risk of diabetes.

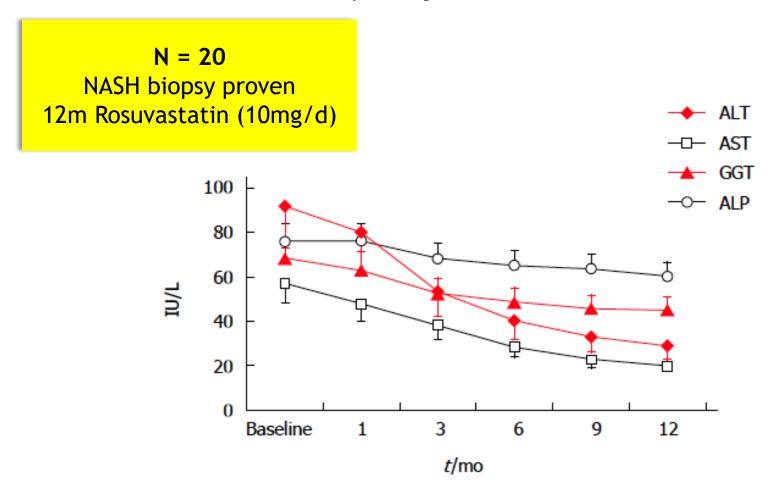
NAFLD - Nonalcoholic Fatty Liver Disease and diabetes

Metformin

- √First-line drug in the treatment of diabetes and prevention of prevention of prevention of prevention.
- ✓ No proven benefit in histological parameters
- ✓ Anti-Tumor Effect Limited Human Data (Retrospective Study)

Haukeland, et al. Scand J Gastroenterol 2009 Shields, et al. Therap Adv Gastroenterol 2009 Bhalla, et al. Cancer Prev Res 2012 Zhang, et al. J Clin Endocrinol Metab 2012

NAFLD - Nonalcoholic Fatty Liver Disease and dyslipidemia



Take home messages

- NAFLD is a highly prevalent disease.
- There is no approved drugs.
- Diet and lifestyle changes are essential.
- Pioglitazone and Vitamin E still play a role.
- Statins are underused in NAFLD patients.



