## Supplementary table 2. Results of round 1 of the Delphi process

Domain and statements	Agree	Somewhat	Somewhat	Disagree
		agree	disagree	
1. Epidemiology of MAFLD and CKD				
1.1 The prevalence of CKD in individuals with MAFLD is higher	82%	18%		
compared to that in the non-MAFLD population.				
1.2 MAFLD is an independent risk factor for CKD in patients with	72%	28%		
T2DM, even after adjustment for common risk factors for CKD.				
1.3 MAFLD is an independent risk factor for CKD in patients	60%	38%	2%	
without T2D, even after adjustment for common risk factors for				
CKD.				
1.4 MAFLD is associated with a greater risk of CKD than patients	50%	34%	16%	
with liver fat but without evidence of systemic metabolic				

dysregulation.				
1.5 MAFLD is associated with an increased incidence of CKD.	82%	18%		
1.6 MAFLD is associated with an increased risk of kidney disease in	30%	46%	18%	6%
childhood.				
1.7 CKD increases the risk of overall mortality among patients with	74%	24%	2%	
MAFLD.				
2. Severity of MAFLD and CKD				
2.1 The presence of metabolic steatohepatitis (MESH) on liver	48%	44%	6%	2%
histology is independently associated with a higher prevalence of				
CKD than simple steatosis.				
2.2 The presence of MESH on liver histology is independently	46%	44%	8%	2%
associated with a higher incidence of CKD than simple steatosis.				
2.3 MAFLD with advanced fibrosis (stage F3/4) has a higher	64%	34%	2%	

prevalence of CKD than MAFLD without advanced fibrosis (stage				
F0-F2).				
2.4 MAFLD with advanced fibrosis (stage F3/4) has a higher	52%	46%	2%	
incidence of CKD than MAFLD without advanced fibrosis (stage				
F0-F2).				
2.5 Advanced liver fibrosis in patients with MAFLD is	56%	40%	4%	
independently associated with an increased risk of incident CKD in				
patients with T2D.				
2.6 Liver stiffness measured by transient elastography is	40%	46%	12%	2%
independently associated with an increased presence of albuminuria.				
3. Mechanisms linking MAFLD with CKD				
3.1 MAFLD and CKD share multiple risk factors such as abdominal	90%	10%		
obesity, insulin resistance, dyslipidemia, hypertension and				

dysglycemia.				
3.2 The MAFLD-associated genetic polymorphisms PNPLA3	30%	54%	14%	2%
rs738409 variant, HSD17B13 variant and TM6SF2 variant are				
associated with CKD.				
3.3 Gut microbiota is linked to both MAFLD and CKD.	48%	40%	10%	2%
3.4 Metabolic dysfunction is an important mechanistic link between	86%	14%		
MAFLD and CKD.				
4. Managing and treating MAFLD and CKD				
4.1 Lifestyle intervention including a hypocaloric diet and regular	74%	22%	4%	
physical exercise is associated with improvements in both MAFLD				
and CKD.				
4.2 Cardiometabolic risk factors should be treated in patients with	96%	4%		
MAFLD and CKD.				

4.3 The use of antihypertensive treatment (if required) is important	82%	18%		
in MAFLD for decreasing risk of CKD.				
4.4 Screening for MAFLD should be undertaken in patients with	54%	40%	4%	2%
CKD.				
4.5 Patients with MAFLD and CKD should ideally be treated in a	90%	8%	2%	
multidisciplinary team setting.				