Author	_	altered?	Topic (and Recommendation if relevant)		Actions taken
Karen Honson (individual) 11.10.20	Health, The Royal	input required	antithrombotic therapy  Aspirin plus ticagrelor commenced within 24 hours may be used in the short	In 2018 in the USA, the FDA began approving generic brands of ticagrelor and there are now a number of different brands available in America.  This is suggestive of a generic version of ticagrelor entering the Australian market at some stage.  Therefore, the issue of cost-effectiveness of the use of ticagrelor for the proposed draft recommendation would be worth revisiting once a generic version of ticagrelor is available on the Australian market. Furthermore, of great interest would be further research into the use of ticagrelor in stroke patients in light of the issues of the use of clopidogrel in patients deemed to have a CYP2C19 poor metaboliser status	
Prof Helen Dewey (individual) 12.10.20	Eastern Health Clinical School, Box Hill Hospital Director of Neurosciences		Acute antithrombotic therapy  Aspirin plus ticagrelor commenced within 24 hours may be used in the short term (first 30 days) in patients with minor ischaemic stroke or high-risk TIA to prevent stroke recurrence.	I disagree with this alternative being included in the guidelines when the therapy is currently 'off label' and prescription would be a significant cost for the patient. There is no doubt that there is evidence for the efficacy of this therapy but these are guidelines for the Australian environment and this therapy is not generally available at this time. The guideline could be updated once the therapy is approved for use for this indication.	indication. The cost differential is clearly noted

			(Johnston et al 2020 [137])		Australia and clopidogrel is not listed on the PBS as first line therapy for stroke.
•	QLD Statewide Stroke Clinical Network (SSCN) Multidisciplinary	NA	Cholesterol targets Limb weakness and management Aspirin plus ticagrelor use in TIA and minor	Overall, the Queensland clinicians involved in stroke care are very happy with the updates.  I can see the immense value these aids will bring in engaging with consumers and educating them regarding their choices in treatment. It would be particularly valuable to see something similar in the discharge planning guidelines, particularly for discharge medications and lifestyle change implementation.	Noted. No change required.
Kylie Jonasson (group)	Director- General, ACT Health	NA	All	Noted drafted updates	Noted. No change required.
22.10.20	Multidisciplinary				
Kate Jackson / NSW health (group) 28.10.20	Agency for Clinical Innovation Multidisciplinary		therapy  Aspirin plus ticagrelor commenced within 24 hours may be used in the short term (first 30 days) in patients with minor ischaemic stroke or high-risk TIA to prevent stroke recurrence. (Johnston et al 2020 [137])	et al (2016) [138] reported a non-statistical difference in the time to occurrence of stroke, MI or death within 90 days with ticagrelor alone compared to aspirin (HR 0.89, 95%CI 0.78-1.01, p=0.07). Ischaemic stroke occurred in 5.8% treated with ticagrelor vs 6.7% treated with aspirin (HR 0.87,	We have rephrased the last sentence to now say: "Single-agent ticagrelor was not superior to aspirin in patients with mild stroke or high risk TIA but may have similar bleeding risk.(Johnston et al. 2016 [138])"

Kate Jackson / Agency for NSW health (Clinical Innovation (group)  28.10.20  Multidisciplinary  Multidiscipl		Ta			<del></del>
Innovation  Multidisciplinary  In patients with ischaemic stroke, cholesterol lowering therapy should target LDL cholesterol to lowering therapy should target LDL cholesterol of attended to recommendation of attended to recommendation. These factors included:  "Clear observational data related to the relationship between LDL and stroke/CVD events"  "Relatively short duration of follow up (longer follow up would likely detect more events and potentially narrow the confidence intervals)  "The current levels align to existing national targets in Australia within PSS/MBS for additional agents that may be needed to reach target  "Adverse event rates are low so the balance clearly falls to desirable (lower CVD events) outcomes.  Further information about GRADE and the strength of the recommendations is listed in the				•	
Multidisciplinary  In patients with the GRADE estimang beaution of preventions. While the Grandshook. While the Grandshook. While the Grandshook. While the Grandshook of patients within the GRADE and free or receivents and put existing nation at the recommendation and further studies may change within the GRADE and the strength of the recommendations is listed in the	NSW health		lowering therapy		I
Multidisciplinary    Schaemic stroke, cholesterol lowering therapy should target LDL cholesterol < 1.8 mmol/L for secondary prevention of atherosclerotic conditions and the working group considered a range of factors (not just the confidence in the estimates of effects) in deciding to make this a strong rather than weak recommendations and the working droup considered a range of factors (not just the confidence in the estimates of effects) in deciding to make this a strong rather than weak recommendation. These factors included:    Clear observational data related to the relationship between LDL and stroke/CVD events   Relatively short duration of follow up (longer follow up would likely detect more events and potentially narrow the confidence intervals)  "The composite outcome was positive" "The corrent levels align to existing national targets in Australia within PBS/MBS for additional agents that may be needed to reach target "Adverse event rates are low so the balance clearly falls to desirable (lower CVD events) outcomes.  Further information about GRADE and the strength of the recommendations is listed in the	(group)	Innovation			, ,
cholesterol lowering therapy should target LDL cholesterol 21.8 mmol/L for secondary prevention of atherosclerotic cordiovascular disease. (Amarenco et al 2020 [112])  al 2020 [112])  cholesterol 4.8 mmol/L for secondary prevention of atherosclerotic cordiovascular disease. (Amarenco et al 2020 [112])  al 2020 [112])  cholesterol 5.8 mmol/L for secondary prevention of atherosclerotic cordiovascular disease. (Amarenco et al 2020 [112])  al 2020 [112])  cholesterol 5.8 mmol/L for secondary prevention of atherosclerotic cordiovascular disease. (Amarenco et al 2020 [112])  cholesterol 6.8 mmol/L for secondary prevention of atherosclerotic cordiovascular disease. (Amarenco et al 2020 [112])  cholesterol 7.8 mmol/L for secondary prevention 6.8 mmol/L for secondary free for secon	(group)	Multidiaciplinary		recommendation might be more appropriate.	•
cholesterol lowering therapy should target LDL cholesterol < 1.8 mmol/L for secondary prevention of atherosclerotic CVD was significant. In GRADE mandsow, while the strong the secondary prevention of atherosclerotic cardiovascular disease. (Amarenco et al. 2020 [112])  2020 [112])  cholesterol < 1.8 mmol/L for secondary prevention of atherosclerotic cardiovascular disease. (Amarenco et al. 2020 [112])  and the secondary prevention of atherosclerotic cardiovascular disease. (Amarenco et al. 2020 [112])  and the secondary prevention of the secondary preventi	28.10.20	iviuitidiscipiiriary	ischaemic stroke,		change the effect estimate. This aligns with the
should target LĎL cholesterol < 1.8 mmol/L for secondary prevention of atherosclerotic CVD was significant. In GRADE we only refer to weak and strong recommendations and the working group considered a range of factors (not just the confidence in the estimates of effects) in deciding to make this a strong rather than weak recommendation. These factors included:  "Clear observational data related to the relationship between LDL and stroke/CVD events  "Relatively short duration of follow up (longer follow up would likely detect more events and potentially narrow the confidence intervals)  "The composite outcome was positive "The current levels align to existing national targets in Australia within PBS/MBS for additional agents that may be needed to reach target  "Adverse event rates are low so the balance clearly falls to desirable (lower CVD events) outcomes.  Further information about GRADE and the strength of the recommendations is listed in the			cholesterol		guidelines within the GRADE handbook. While
cholesterol < 1.8 mmol/L for secondary prevention of atherosclerotic CVD was significant. In GRADE we only refer to weak and strong recommendations and the working group considered a range of factors (not just the confidence in the estimates of effects) in deciding to make this a strong rather than weak recommendation. These factors included:  2020 [112])  (Clear observational data related to the relationship between LDL and stroke/CVD events  Relatively short duration of follow up (longer follow up would likely detect more events and potentially narrow the confidence intervals)  The composite outcome was positive  The current levels align to existing national targets in Australia within PBS/MBS for additional agents that may be needed to reach target  Adverse event rates are low so the balance clearly falls to desirable (lower CVD events) outcomes.  Further information about GRADE and the strength of the recommendations is listed in the			lowering therapy		it is true the confidence intervales for recurrent
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prevention of atherosclerotic cardiovascular disease. (Amarenco et al 2020 [112])  2020 [112])			mmol/L for		atherosclerotic CVD was significant. In GRADE
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Kate Jackson /				Broadly agree with this recommendation for	We agree focus on PRE is on strength and that
NSW health	Clinical			progressive resistance training, so long as the	has been reflected throughout. However, we
()	Innovation		For stroke	wording of the recommendation makes it clear that the	have added this into the recommendation to
(group)	NA ICE Cartalian			1 7 0	ensure this is clear and the recommendation
28.10.20	Multidisciplinary	У	weakness	STRENGTHENING, and not necessarily FUNCTION.	now reads:
20.10.20			repetitive practice		
			using assistive	Agree with the new recommendation for repetitive	For stroke survivors with reduced strength in
			_	training.	their arms or legs, progressive resistance
			constraint induced		training should be provided to increase
			movement	For cycling, there were only two studies included in	strength. (Dorsch et al. 2018 [73])
				the subgroup analysis in the de Sousa	
				systematic review – is this sufficient to make a	Regarding cycling, there are other
			•	recommendation?	recommendations made on small numbers of
			used to improve		studies and this is one important consideration
			•	For Electrical stimulation the subgroup analysis (in de	as to why we only made a weak
			• ,	Souza) included 2 studies that included ES only on	recommendation. We will continue to monitor
				the upper limb – does removing these studies change	
				the result of the sub-group analysis?	any new evidence in this regard.
			For stroke	group analysis.	Regarding electrical stimulation - while de
			survivors with leg		Souza supplement does provide a breakdown
			weakness task		in UL and LL the effects were not that different
			specific training,		(SMD 0.37 vs 0.45) with two small studies in
					the UL leading to wide confidence intervals.
			repetitive practice		
			using cycling or		The group didn't feel there was a reason to
			electrical		expect differences in FES should be different
			stimulation may be		for muscles in the UL and LL and hence
			used to improve		presented the stronger, combined effect
			leg strength. (de		estimates.
			Sousa et		
			al 2018 [70])		

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Kate Jackson /			Shoulder pain	These shoulder recommendations seem to sit under a	·
NSW health	Clinical			heading of Central Post-Stroke Pain	heading from shoulder pain.
(0.00.10)	Innovation			(CPSP). It is considered that strapping intervention,	
(group)	NA 161-11 1 11		survivors with	and the electrical stimulation intervention are more	Regarding electrical stimulation -the data is
28.10.20	Multidisciplinary		shoulder pain,	likely to help some forms of Mechanical shoulder pain.	based on 4 studies -two of these used VAS but
			electrical	Even if the studies were 'non-discriminatory'	two others used Brief Pain Inventory 12 so the
			stimulation may be	recommending these therapies for CPSP, we believe,	outcome is a standardised mean difference
			used to	inappropriate.	(SMD). Thus 1.89 lower SMD is a significant
			manage pain. (Qiu	This applies to the Practice points as well. The data	and clinically meaningful difference.
			et al	for electrical stimulation to assist with	
			2019 [87])	pain is inadequate. The apparent improvements in	Regarding acupuncture we agree that the
				pain at a mean of less than 2.0 on the visual analogue	evidence is weak and effects small and this is
				scale (VAS) could be regarded as statistically	reflected in the evidence summary. However,
				significant but not clinically significant, where a shift of	we also agree that acupuncture should not be
				at least 2.0 on the VAS is usually required.	considered in isolation and have made
					changes to the recommendation and rationale
				Concerns raised in relation to the rationale to include	to reflect this. Regarding the minimally clinical
				any statement re acupuncture. Not only	important difference (MCID) there is a lack of
				does the degree of pain reduction not reach clinical	evidence to determine the threshold for this
				significance, but the studies are of low	specific to stroke. An additional sentence has
				quality, and there is no definition around what 'type' of	been inserted in the evidence summary to
				acupuncture is being recommended.	reflect this uncertainty and to also reflect in
				All the studies included in the systematic review had	other patient populations with shoulder
					problems (rotator cuff disease and post
				rehabilitation – it would be good to highlight/include	shoulder arthroplasty) the MCID was found to
					be -1.4cm and the group felt while the effect is
				, , ,	small it may still be worthwhile to some
				· · · · · · · · · · · · · · · · · · ·	patients.
Hannah Paal		NA	All	Nil	Noted. No change required.
(Group)	Stroke Clinical				
0.44.00	Network				
3.11.20	NA distribution also list and				
	Multidisciplinary				
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