Appendix 1 – search strategy employed in MEDLINE and adapted for EMBASE and the Cochrane Database

- 1. Exp Intestine, Large/
- 2. C?ecum.ti,ab
- 3. Colon\*.ti,ab
- 4. Rect\*.ti,ab
- 5. Colorectal.ti,ab
- 6. OR/1-5
- 7. Dysplas\*.ti,ab
- 8. Adenoma\*.ti,ab
- 9. Polyp\*.ti,ab
- 10. Adenomatous polyps/
- 11. Pseudopolyp\*.ti,ab
- 12. Neoplas\*.ti,ab
- 13. Lesion\*.ti,ab
- 14. OR/7-13
- 15. 6 AND 14
- 16. exp Colonic Neoplasms/
- 17. exp Colonic Polyps/
- 18. exp Rectal Neoplasms/
- 19. OR/15-18
- 20. exp Colonoscopy/
- 21. Colonoscop\*.ti,ab
- 22. Endoscop\*.ti,ab
- 23. Sigmoidoscop\*.ti,ab
- 24. OR/20-23
- 25. (Real adj time).ti,ab
- 26. (In adj vivo).ti,ab
- 27. Spectroscop\*.ti,ab
- 28. Endomicroscop\*.ti,ab
- 29. Chrom?endoscop\*.ti,ab
- 30. Fl?orosc\*.ti,ab
- 31. (Narrow adj band).ti,ab
- 32. Optical.ti,ab
- 33. (i adj scan).ti,ab
- 34. (colo?r AND enhancement).ti,ab
- 35. FICE.ti,ab
- 36. OR/25-35
- 37. Diagnos\*.ti,ab
- 38. Detect\*.ti,ab
- 39. Classif\*.ti,ab
- 40. Histolog\*.ti,ab
- 41. Assessment.ti,ab
- 42. Analysis.ti,ab
- 43. Characteri\*.ti,ab
- 44. OR/37-43
- 45. 19 AND 24 AND 36 AND 44
- 46. limit 45 to human

		Risk c	of Bias		Ар	plicabi	lity	
	Patient Selection	Index Test	Reference Standard	Flow and Timing	Patient Selection	Index Test	Reference Standard	Reason for <i>Unclear</i> or <i>High</i> Score
Ashktorab 2016	0	?	0	0	0	0	0	Unclear endoscopist experience with NBI technology prior to study
Ashktorab 2016	0		0	0	0	0	0	Unclear endoscopist experience with NBI technology prior to study
Basford 2014	0	0	0	0	0	0	0	-
Belderbos 2017		0	0	0	0	0	0	Unclear if consecutive patients were eligible, limited sampling of rectal hyperplastic polyps reduces adenoma prevalence
Buchner 2010			0	0	0	0	0	Unclear if consecutive patients were eligible, some lesions known to be adenomas prior to index test
Buchner 2010			0	0	0	0	0	Unclear if consecutive patients were eligible, some lesions known to be adenomas prior to index test
Canales-Sevilla 2010			0	0	0	0	0	Unclear if consecutive patients were eligible for study. Unclear endoscopist experience with NBI technology prior to study
Chan 2012	0	0	0	0	0	0	0	-
Chandran 2015	0	0	0	0	0	0	0	-
Dai 2013	0	0	0	0	0	0	0	-
dos Santos 2009		0	0	0	0	0	0	Unclear if consecutive patients were eligible
dos Santos 2010	0	0	0	0	0	0	0	-
dos Santos 2012	0	0	0	0	•	0	0	-
dos Santos 2017	0	0	0	0	•	0	0	-
dos Santos 2017	0	0	0	0	•	0	0	-
East 2008	0	0	0	0	•	0	0	-
Hewett 2012		0	0	0	•	0	0	Unclear if consecutive patients were eligible for study
Hewett 2012		0	0	0	0	0	0	Unclear if consecutive patients were eligible for study
Hoffman 2010	0	0	0	0	0	0	0	-
Hoffman 2010	0	0	0	0	0	0	0	-

Hong 2012	0	0	0	0	0	0	0	
Hong 2012	0	0	0	0	0	0	0	
Ikematsu 2015	0	0	0	0	0	0	0	
lwatate 2015	0		0	0	0	0	0	Ur
lwatate 2015	0		0	0	0	0	0	Ur
Kaltenbach 2015	0	0	0	0	0	0	0	
Kaltenbach 2015	0	0	0	0	0	0	0	
Kang 2015	0	0	0	0	0	0	0	
Kang 2015	0	0	0	0	0	0	0	
Kim 2011	0	0	0	0	0	0	0	
Klare 2016	0	0	0	0	0	0	0	
Kuiper 2011	0		0	0	0	0	0	Le
Kuiper 2012	0	0	0	0	0	0	0	
Kuruvilla 2015	0	0	0	0		0	0	I
Ladabaum 2013		0	0	0	0	0	0	
Lee 2011	0	0	0	0	0	0	0	
Lee 2011	0	0	0	0	0	0	0	
Liu 2008		0	0	0	0	0	0	Und
Longcroft- Wheaton 2011	0	0	0	0	0	0	0	
Longcroft- Wheaton 2012	0	0	0	0	0	0	0	
Longcroft- Wheaton 2012	0	0	0	0	0	0	0	
Machida 2004		0	0	0	0	0	0	Und
Okamoto 2011		0	0	0		0	0	Uno we
Paggi 2012	0	0	0	0	O	0	0	
Paggi 2015	0	0	0	0	0	0	0	
Pigo 2013	0	?	0	0	0	0	0	Unc
				-		_	_	

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nclear if all endoscopists had sufficient experience with NBI prior to study	
nclear if all endoscopists had sufficient experience with NBI prior to study	
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esions also assessed with AFI, likely biasing interpretation of the NBI	
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High incidence of SSAs (17%) unrepresentative of routine clinical practice

Unclear if consecutive patients were eligible for study

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Unclear how patients were selected and if consecutive patients were eligible for study

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Unclear how patients were selected and if consecutive patients were eligible for study

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Unclear how patients were selected and if consecutive patients were eligible for study. High incidence of adenomas (95%) not representative of practice

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Unclear if all endoscopists had sufficient experience with iSCAN prior to study

Pohl 2009	0		0	0	0	0	0	Er
Pohl 2016	0	0	0	0	0	0	0	
Pohl 2016	0	0	0	0	0	0	0	
Rastogi 2011	0	0	0	0	0	0	0	
Rath 2015	0		0	0	•	0	0	U
Rees 2017	0	0	0	0	0	0	0	
Ren 2012	0		0	0	0	0	0	Un
Repici 2013	0	0	0	0	0	0	0	
Rex 2009	0	0	0	0	0	0	0	
Rogart 2008	0	0	0	0	0	0	0	
Rogart 2011	0	0	0	0	0	0	0	
Rogart 2011	0	0	0	0	0	0	0	
Rotondano 2012	0	0	0	0	?	0	0	Hig
Sakamoto 2012	0		0	0		0	0	chr iı
Salazar Muente 2012		0	0	0	0	0	0	
Sano 2009	0	0	0	0	0	0	0	
Sano 2015			0	0	0	0	0	U
Schachschal 2014	0	0	0	0	0	0	0	
Seref Koksal 2014	0		0	0	0	0	0	E
Shahid 2012		0	0	0	0	0	0	Un
Singh 2011			0	0	0	0	0	ι
Singh 2013	0		0	0	0	0	0	
Sola-Vera 2015	0	0	0	0	0	0	0	
Szura 2016	8	?	0	0	0	0	0	ι
Takeuchi 2014	0	0	0	0	0	0	0	
Takeuchi 2015	0	0	0	0	0	0	0	

Endoscopists were often inexperienced with FICE prior to the study starting

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Unclear if endoscopists had sufficient experience with iSCA

N prior to study

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nclear if endoscopists had sufficient experience v	with NBI prior
to study	

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High incidence of adenomas (90%) not representative of clinical practice

Half of the lesions had just been assessed by dye chromoendoscopy, likely biasing the interpretation of NBI. High incidence of adenomas (89%) not representative of practice

Unclear if consecutive patients were eligible for study

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Unclear if consecutive patients were eligible for study. IC dye was used for detection and may have biased NBI

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Endoscopists had no experience using NBI prior to the study starting

Unclear how patients selected and if consecutive patients were eligible for the study

Unclear how patients were selected, likely not consecutive. Unlikely sufficient endoscopist experience with NBI

Unlikely sufficient endoscopist experience with NBI

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Unclear how patients were selected, likely not consecutive. Unlikely sufficient endoscopist experience with NBI

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Takeuchi 2015	$\mathbf{O} \mathbf{O} \mathbf{O} \mathbf{O}$	$\mathbf{O} \mathbf{O} \mathbf{O}$
Togashi 2009	$\bigcirc \bigcirc $	$\mathbf{O} \mathbf{O} \mathbf{O}$
Van den Broek 2009	$\mathbf{O} \bigcirc \mathbf{O} \mathbf{O}$	$\mathbf{O} \mathbf{O} \mathbf{O}$
Wallace 2014	$\mathbf{O} \mathbf{O} \mathbf{O} \mathbf{O}$	$\mathbf{O} \mathbf{O} \mathbf{O}$
Wallace 2014	$\mathbf{O} \mathbf{O} \mathbf{O} \mathbf{O}$	$\mathbf{O} \mathbf{O} \mathbf{O}$
Yoo 2011	$\bigcirc \bigcirc $	00
Zhou 2011		$\mathbf{O} \mathbf{O} \mathbf{O}$

Unclear how patients were selected, likely not consecutive

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Unlikely sufficient endoscopist experience with NBI prior to study start. Some lesions previously assessed with AFI, likely biasing interpretation

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Unclear how patients were selected. High incidence of adenomas (93%) not representative of clinical practice

Unclear how patients were selected, likely not consecutive. Unlikely sufficient endoscopist experience with NBI

		Risk o	f Bias		Арр	olicabi	lity	
	Patient Selection	Index Test	Reference Standard	Flow and Timing	Patient Selection	Index Test	Reference Standard	Reason for <i>Unclear</i> or <i>High</i> Score
Apel 2006	0	?		0	0	0	0	Unclear endoscopist experience with chromoendoscopy. Pathologists were not blinded to intra-operative prediction
Averbach 2003			0	0	0	0	0	Unclear how patients were selected. Unclear endoscopist experience with chromoendoscopy prior to study start
Axelrad 1996	0	0	0	0	0	0	0	-
Bianco 2006	0		0	0	0	0	0	Unclear endoscopist experience with chromoendoscopy prior to study start
de Palma 2006		0	0	0	0	0	0	Unclear how patients were selected, likely not consecutive
dos Santos 2009			0	0	0	0	0	Unclear how patients were selected. Lesion chromoendoscopy interpretation was biased by just having performed FICE
dos Santos 2010	0	0	0	0	0	0	0	-
dos Santos 2012	0	0	0	0	0	0	0	-
Eisen 2002			0	0	0	0	0	Some patients only had sigmoidoscopy, changing the prevalence of lesion histological subtypes. Unclear endoscopist experience with chromoendoscopy
Fu 2004	0	0	0	0	0	0	0	-
Hurlstone 2004			0	0	0	0	0	Unclear if consecutive patients eligible. Unclear if sufficient endoscopist experience with chromoendoscopy
Ince 2007		0	0	0	0	0	0	Unclear how patients were selected, likely not consecutive
Kato 2006		0	0	0		0	0	Unclear how patients were selected. High incidence of adenomas (88%) not representative of clinical practice
Kiesslich 2001	0		0	0	0	0	0	Unclear endoscopist experience with chromoendoscopy prior to study start
Kohut 2009	0	0	0	0	0	0	0	-
Konishi 2003	0	0	0	0	0	0	0	-
Konishi 2003	0	0	0	0	0	0	0	-
Liu 2003			0	0	0	0	0	Unclear indications for colonoscopy. Unclear endoscopist experience with chromoendoscopy prior to study start
Liu 2008			0	0	0	0	0	Unclear how patients were selected, likely not consecutive. Lesion chromoendoscopy interpretation was biased by just having performed FICE. Unclear endoscopist experience
Ljubicic 2001			0	0	0	0	0	Unclear if consecutive patients eligible. Unclear if sufficient endoscopist experience with chromoendoscopy

Longcroft- Wheaton 2011	0		0	0	0	0	0	Chr
Longcroft- Wheaton 2013	0	0	0	0	0	0	0	
Longcroft- Wheaton 2013	0	0	0	0	0	0	0	
Machida 2004			0	0	0	0	0	ι
Pohl 2009	0		0	0	0	0	0	Unc
Sakamoto 2012	0		0	0		0	0	C be
Togashi 1999	0		0	0	0	0	0	Unc
Togashi 2006	0		0	0	0	0	0	Unc
Togashi 2009			0	0	0	0	0	с
Tung 2001	0	0	0	0	0	0	0	
Urban 2005			0	0	0	0	0	U

Chromoendoscopy interpretation was biased by just performing
FICE

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Unclear if consecutive patients eligible. Chromoendoscopy interpretation was biased by just performing NBI

Inclear endoscopist experience with chromoendoscopy prior to study start

Chromoendoscopy interpretation was biased by just having been analysed by NBI. High incidence of adenomas (89%) not representative of practice

Unclear endoscopist experience with chromoendoscopy prior to study start

Unclear endoscopist experience with chromoendoscopy prior to study start

Unclear if consecutive patients were eligible. Lesion chromoendoscopy interpretation was biased by just having performed FICE

Unclear if consecutive patients eligible. Unclear if sufficient endoscopist experience with chromoendoscopy

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