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Küsimus: Kas kõigil ravinaivsetel 2. tüüpi diabeeti põdevatel inimestel alustada ravi elustilisekkumisega või suukaudse antidiabeetilise ravimiga või mõlemaga, parema ravitulemuse saamiseks?

Kontekst:

Bibliograafia:

Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebakõla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			

Keskmene erinevus HbA1c algväärtusest (mean difference from baseline) kui langus kehakaalus oli  $\geq 5\%$  (järelkontroll: keskmene 12 kuud)

2 <sup>1,2,a,b</sup>	jälgimisuuringud	väga suur <sup>c</sup>	suur <sup>d</sup>	väike	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) algväärtusest = -0,91% (95% CI -2,3; 0,48)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus HbA1c algväärtusest (mean difference from baseline) kui langus kehakaalus oli  $< 5\%$  (järelkontroll: keskmene 12 kuud)

17 <sup>1,3,4,5,6,7,8,9,10,11,a,b,f</sup>	jälgimisuuringud	väga suur <sup>c</sup>	väike	väike	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) algväärtusest = -0,224% (95% CI -0,64; 0,19)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus HbA1c algväärtusest (mean difference from baseline) (järelkontroll: keskmene 12 kuud)

3 <sup>2,9,11,a,b</sup>	jälgimisuuringud	väga suur <sup>c</sup>	väike	väike	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) algväärtusest = -0,128% (95% CI -1,56; 1,31)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus üldkolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli  $\geq 5\%$  (järelkontroll: keskmene 12 kuud)

1 <sup>1,a,b</sup>	jälgimisuuringud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e,h</sup>	puudub	Keskmene erinevus (MD) = -15,1 mg/dl (95% CI -46,43; 16,23)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus üldkolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli  $< 5\%$  (järelkontroll: keskmene 12 kuud)

15 <sup>1,3,4,5,6,7,9,10,11,a,b,f</sup>	jälgimisuuringud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -4,39 mg/dl (95% CI -15,47; 6,69)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus üldkolesteroli (mg/dl) algväärtusest (mean difference from baseline) (järelkontroll: keskmene 12 kuud)

2 <sup>9,11,a,b</sup>	jälgimisuuringud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = 4,24 mg/dl (95% -64,36; 72,83)	 VÄGA MADAL	KRIITILINE
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Keskmene erinevus LDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli  $\geq 5\%$  (järelkontroll: keskmene 12 kuud)

Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebaköla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			
12	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -4,44 mg/dl (95% CI -61,49; 52,61)		KRIITILINE

Keskmene erinevus LDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli < 5% (järekontroll: keskmene 12 kuud)

14 3,4,5,6,7,9,10,11,a,b,f	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -0,67 mg/dl (95% CI -16,87; 15,53)		KRIITILINE
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Keskmene erinevus LDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) (järekontroll: keskmene 12 kuud)

3 2,9,11,a,b	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -0,62 mg/dl (95% CI -34,7; 33,47)		KRIITILINE
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Keskmene erinevus HDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli ≥ 5% (järekontroll: keskmene 12 kuud)

2 1,2,a,b	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = 3,76 mg/dl (95% CI -10,62; 18,15)		KRIITILINE
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Keskmene erinevus HDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli < 5% (järekontroll: keskmene 12 kuud)

15 1,3,4,5,6,7,9,10,11,a,b,f	jälgimisuurud	väga suur <sup>c</sup>	suur <sup>d</sup>	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = 1,22 mg/dl (95% CI -0,37; 2,82)		KRIITILINE
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Keskmene erinevus HDL-kolesteroli (mg/dl) algväärtusest (mean difference from baseline) (järekontroll: keskmene 12 kuud)

3 2,9,11,a,b	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>g</sup>	puudub	Keskmene erinevus (MD) = 0,55 mg/dl (95% CI -8,26; 9,37)		KRIITILINE
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Keskmene erinevus triglütseriidide (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli > 5% (järekontroll: keskmene 12 kuud)

2 1,2,a,b	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -35,11 mg/dl (95% CI -189,15; 118,91)		KRIITILINE
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Keskmene erinevus triglütseriidide (mg/dl) algväärtusest (mean difference from baseline) kui langus kehakaalus oli < 5% (järekontroll: keskmene 12 kuud)

13 1,3,4,5,6,7,9,10,11,a,b,f	jälgimisuurud	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmene erinevus (MD) = -16,9 (95% CI -88,97; 55,07)		KRIITILINE
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Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebaköla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			

Keskmise erinevuse triglütseriidide (mg/dl) algväärtusest (mean difference from baseline) (järelkontroll: keskmise 12 kuud)

3 <sup>2,9,11,a,b</sup>	jälgimisuurused	väga suur <sup>c</sup>	suur <sup>d</sup>	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = 9,07 mg (95% CI -117,39; 135,54)		KRIITILINE
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Keskmise erinevuse süstoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) kui langus kehakaalus oli > 5% (järelkontroll: keskmise 12 kuud)

2 <sup>1,2</sup>	jälgimisuurused	väga suur <sup>c</sup>	suur <sup>d</sup>	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = -5,24 mmHg (95% CI -13,77; 3,3)		KRIITILINE
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Keskmise erinevuse süstoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) kui langus kehakaalus oli < 5% (järelkontroll: keskmise 12 kuud)

12 <sup>1,3,4,5,6,9,10,a,b,f</sup>	jälgimisuurused	väga suur <sup>c</sup>	suur <sup>d</sup>	väga suur <sup>g</sup>	väike	puudub	Keskmise erinevus (MD) = -2,24 mmHg (95% CI -5,83; 1,34)		KRIITILINE
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Keskmise erinevuse süstoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) (järelkontroll: keskmise 12 kuud)

2 <sup>2,9</sup>	jälgimisuurused	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = -6,61 mmHg (95% CI -27,56; 14,34)		KRIITILINE
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Keskmise erinevuse diastoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) kui langus kehakaalus oli > 5% (järelkontroll: keskmise 12 kuud)

2 <sup>1,2,a,b</sup>	jälgimisuurused	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = -3,13 mmHg (95% CI -19,13; 12,87)		KRIITILINE
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Keskmise erinevuse diastoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) kui langus kehakaalus oli < 5% (järelkontroll: keskmise 12 kuud)

12 <sup>1,3,4,5,6,9,10,a,b,f</sup>	jälgimisuurused	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = -3,53 mmHg (95% CI -9,80; 2,73)		KRIITILINE
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Keskmise erinevuse diastoolse vereröhu (mmHg) algväärtusest (mean difference from baseline) (järelkontroll: keskmise 12 kuud)

2 <sup>2,9,a,b</sup>	jälgimisuurused	väga suur <sup>c</sup>	väike	väga suur <sup>g</sup>	suur <sup>e</sup>	puudub	Keskmise erinevus (MD) = -2,94 mmHg (95% CI -13,31; 7,44)		KRIITILINE
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HbA1c (%) keskmiste erinevuse (struktureeritud aerobne treening vs. kontroll) (järelkontroll: vahemik 12 nädalat kuni 52 nädalat)

Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebaköla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			
20 <sup>12,13,14,15,16,17,18,19,20,21,22,23,24,25,26,27,28,29,f,i</sup>	randomiseeritud uuringud	väga suur <sup>j</sup>	väga suur <sup>k</sup>	väike	väike	puudub	Keskmiste erinevus (ingl weighted mean difference) aerobne treening (n = 490) vs. kontroll (n = 455) = -0,73% (95% CI -1,06; -0,40)	 VÄGA MADAL	KRIITILINE

HbA1c (%) keskmiste erinevus (struktureeritud vastupidavustreening vs. kontroll) (järelkontroll: vahemik 16 nädalat kuni 39 nädalat)

4 <sup>17,29,30,31,j</sup>	randomiseeritud uuringud	väga suur <sup>j</sup>	väga suur <sup>i</sup>	väike	suur <sup>e</sup>	puudub	Keskmiste erinevus (ingl weighted mean difference) vastupidavustreening (n = 182) vs. kontrollrühm (n = 148) = -0,57% (95% CI -1,14; -0,01)	 VÄGA MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (kombineeritud treening vs. kontroll) (järelkontroll: vahemik 12 nädalat kuni 52 nädalat)

7 <sup>15,17,18,29,32,33,34,i</sup>	randomiseeritud uuringud	väga suur <sup>j</sup>	suur <sup>m</sup>	väike	väike	puudub	Keskmiste erinevus (ingl weighted mean difference) kombineeritud treening (n = 262) vs. kontroll (n = 222) = -0,51% (95% CI -0,79; -0,23)	 VÄGA MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (füüsilise aktiivsuse alane nõustamine koos toitumise ko-interventsiooniga vs. kontroll) (järelkontroll: vahemik 26 nädalat kuni 104 nädalat)

12 <sup>35,36,37,38,39,40,41,42,43,44,45,46,i</sup>	randomiseeritud uuringud	väga suur <sup>n</sup>	suur <sup>o</sup>	väike	väike	puudub	Keskmiste erinevus (ingl weighted mean difference) füüsilise aktiivsuse alane nõustamine koos toitumisalase ko-interventsiooniga (n = 3126) vs. kontroll (n = 3171) = -0,58% (95% CI -0,74; -0,43)	 VÄGA MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (füüsilise aktiivsuse alane nõustamine vs. kontroll) (järelkontroll: vahemik 12 nädalat kuni 52 nädalat)

14 <sup>47,48,49,50,51,52,53,54,55,56,57,58,f,i</sup>	randomiseeritud uuringud	väga suur <sup>n</sup>	suur <sup>p</sup>	väike	väike	puudub	Keskmiste erinevus (ingl weighted mean difference) füüsilise aktiivsuse alane nõustamine (n = 403) vs. kontroll (n = 367) = -0,16% (95% CI -0,50; 0,18)	 VÄGA MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) sekkumise pikkus varieerus 2 päevast 1 aastani

8 <sup>59,60,61,62,63,64,65,66,q,r</sup>	randomiseeritud uuringud	suur <sup>s</sup>	väike <sup>t</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 1050) vs. kontroll (n = 985) = -0,44% (95% CI -0,60; -0,29)	 KESKMINÉ	KRIITILINE
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HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) sekkumise pikkus < 6 nädalat

3 <sup>61,64,66,q,r</sup>	randomiseeritud uuringud	suur <sup>u</sup>	väike <sup>v</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 558) vs. kontroll (n = 508) = -0,42 (95% CI -0,68; -0,15)	 KESKMINÉ	KRIITILINE
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Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebakõla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			

HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) sekkumise pikkus 6 nädalat > x < 1 aasta

3 59,62,63,q,r	randomiseeritud uuringud	väga suur <sup>w</sup>	suur <sup>x</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 355) vs. kontroll (n = 337) = -0,43% (95% CI -0,74; -0,12)	⊕○○○ VÄGA MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) sekkumise pikkus 1 aasta

2 60,65,q,r	randomiseeritud uuringud	suur <sup>y</sup>	väike	suur <sup>z</sup>	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 137) vs. kontroll (n = 140) = -0,68% (95% CI -1,22; -0,14)	⊕⊕○○ MADAL	KRIITILINE
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HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) esmatasand

3 61,64,66,q,r	randomiseeritud uuringud	suur <sup>u</sup>	väike <sup>aa</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 558) vs. kontroll (n = 508) = -0,42% (95% CI -0,68; -0,15)	⊕⊕⊕○ KESKMINNE	KRIITILINE
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HbA1c (%) keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) kogukonnas

5 59,60,62,63,65	randomiseeritud uuringud	väga suur <sup>ab</sup>	väike <sup>ac</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 492) vs. kontroll (n = 477) = -0,48% (95% CI -0,70; -0,26).	⊕⊕○○ MADAL	KRIITILINE
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HbA1c keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) grupinõustamine

5 60,63,64,65,66,q,r	randomiseeritud uuringud	väga suur <sup>ad</sup>	väike <sup>ae</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 774) vs. kontroll (n = 719) = -0,47% (95% CI -0,66; -0,28).	⊕⊕○○ MADAL	KRIITILINE
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HbA1c keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) (ingl mean difference) individuaalne nõustamine

1 61,q,r	randomiseeritud uuringud	suur <sup>af</sup>	väike	väike	suur <sup>ag</sup>	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 71) vs. kontroll (n = 70) = -0,80% (95% CI -1,35; -0,25).	⊕⊕○○ MADAL	KRIITILINE
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HbA1c keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) HbA1c uuringu alguses < 9%

6 59,60,62,63,64,66,q,r	randomiseeritud uuringud	väga suur <sup>ah</sup>	väike <sup>ai</sup>	väike	väike	puudub	Keskmiste erinevus (ingl mean difference) käitumuslik sekkumine (n = 867) vs. kontroll (n = 803) = -0,40% (95% CI -0,55; -0,24).	⊕⊕○○ MADAL	KRIITILINE
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HbA1c keskmiste erinevus (ingl mean difference) (käitumuslik sekkumine vs. kontroll) HbA1c uuringu alguses ≥ 9%

Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebakõla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			
2 61,65,q,r 67,68,69,70,71,72,73,74,75,76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100,101,102,103,104,105,106,107,108,109,110,111,112,113,114,115,116,117,118,119,120,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	väike <sup>a)</sup>	suur <sup>z)</sup>	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) käitumuslik sekkumine (n = 183) vs. kontroll (n = 182) = -0,79% (95% CI -1,23; -0,34).	⊕⊕○○ MADAL	KRIITILINE

HbA1c muutus keskmiste erinevus (mean difference) (igasugune toitumisalane sekkumine vs. kontroll) (0-24 kuud)

59 67,68,69,70,71,72,73,74,75,76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100,101,102,103,104,105,106,107,108,109,110,111,112,113,114,115,116,117,118,119,120,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	suur <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 2674) vs. kontroll (n = 2208) = -0,35% (95% CI -0,43; -0,28)	⊕⊕○○ MADAL	
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HbA1c muutus keskmiste erinevus (ingl mean difference) (igasugune toitumisalane sekkumine vs. kontroll) (0-3 kuud)

35 68,70,73,74,75,76,77,78,79,80,81,82,83,84,85,89,91,93,94,98,99,100,102,103,105,108,109,111,112,113,114,118,119,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	suur <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1326) vs. kontroll (n = 1193) = -0,37% (95% CI -0,48; -0,26).	⊕⊕○○ MADAL	KRIITILINE
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HbA1c muutus keskmiste erinevus (ingl mean difference) (igasugune toitumisalane sekkumine vs. kontroll) (3-6 kuud)

26 67,69,70,71,78,81,86,88,90,91,95,96,97,98,101,103,104,106,107,110,115,117,120,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	suur <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1659) vs. kontroll (n = 1346) = -0,37% (95% CI -0,48; -0,26)	⊕⊕○○ MADAL	KRIITILINE
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HbA1c muutus keskmiste erinevus (ingl mean difference) (igasugune toitumisalane sekkumine vs. kontroll) (6-12 kuud)

16 67,77,78,87,92,96,97,98,99,104,106,115,116,117,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	väike <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1141) vs. kontroll (n = 775) = -0,40% (95% CI -0,52; -0,28)	⊕⊕⊕○ KESKMININE	KRIITILINE
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HbA1c muutus keskmiste erinevus (mean difference) (igasugune toitumisalane sekkumine vs. kontroll) (12-24 kuud)

5 72,77,87,88,117,al.am.an.ao	randomiseeritud uuringud	suur <sup>a)</sup>	suur <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 186) vs. kontroll (n = 147) = -0,14% (95% CI -0,63; 0,35)	⊕⊕○○ MADAL	KRIITILINE
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Kehamassi (kg) muutus keskmiste erinevus (igasugune toitumisalane sekkumine vs. kontroll (mean difference) (0-24 kuud)

54 67,68,69,70,71,72,73,74,75,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,94,95,96,98,99,100,101,102,103,105,106,107,108,109,110,112,113,114,115,116,117,118,119,120,al.am.an.ao.f	randomiseeritud uuringud	suur <sup>a)</sup>	väga suur <sup>a)</sup>	väike	väike	puudub	Keskmine erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 2445) vs kontroll (n = 2051) = -2,41kg (95% CI -2,96; -1,86).	⊕○○○ VÄGA MADAL	KRIITILINE
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Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebakõla	Töenduse kaudsus	Töenduse ebatäpsus	Muud kaalutlused			

Kehamassi (kg) muutus keskmiste erinevus (mean difference) (igasugune toitumisalane sekkumine vs. kontroll (0-3 kuud)

33 68,70,72,73,74,75,78,79,80,82,83,84,85,89,91,94,98,99,100,102,103,105,107,108,109,112,113,114,118,119,al,am,an,ao,ap,f	randomiseeritud uuringud	suur <sup>sp</sup>	väga suur <sup>aw</sup>	väike	väike	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1271) vs. kontroll (n = 1145) = -2,34kg (95% CI -2,99; -1,69).	 VÄGA MADAL	KRIITILINE
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Kehamassi (kg) muutus keskmiste erinevus (ingl *mean difference*) (igasugune toitumisalane sekkumine vs. kontroll (3-6 kuud)

24 67,69,70,71,72,78,81,86,88,90,91,95,96,98,101,103,106,107,110,115,120,al,am,an,ao,f	randomiseeritud uuringud	suur <sup>sp</sup>	väga suur <sup>ax</sup>	väike	väike	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1562) vs. kontroll (n = 1288) = -2,94kg (95% CI -3,92; -1,97).	 VÄGA MADAL	KRIITILINE
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Kehamassi (kg) muutus keskmiste erinevus (mean difference) (igasugune toitumisalane sekkumine vs. kontroll (6-12 kuud)

14 67,78,87,92,96,98,99,106,115,116,117,al,am,an,ao,f	randomiseeritud uuringud	suur <sup>sp</sup>	väga suur <sup>ay</sup>	väike	väike	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 1000) vs. kontroll (n = 704) = -2,27kg (95% CI -3,32; -1,21).	 VÄGA MADAL	KRIITILINE
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Kehamassi (kg) muutus keskmiste erinevus (igasugune toitumisalane sekkumine vs. kontroll (mean difference) (12-24 kuud)

4 87,88,117,al,am,an,ao,f	randomiseeritud uuringud	suur <sup>sp</sup>	väike <sup>az</sup>	väike	suur <sup>ag</sup>	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) igasugune toitumisalane sekkumine (n = 103) vs. kontroll (n = 102) = -2,14kg (95% CI -3,34; -0,93).	 MADAL	KRIITILINE
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HbA1c muutus keskmiste erinevus (ingl *mean difference*) (madala süsivesikusisaldusega ehk LC diet vs. kontroll)

9 69,75,78,84,85,95,96,113,120,ao	randomiseeritud uuringud	suur <sup>sp</sup>	väike <sup>ba</sup>	väike	väike	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) madala süsivesikusisaldus diet ehk LC diet (n = 220) vs. kontroll (n = 213) = -0,44% (95% CI -0,58; -0,31)	 KESKMINNE	KRIITILINE
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HbA1c muutus keskmiste erinevus (ingl *mean difference*) (madala rasvasisaldusega ehk LF diet vs. kontroll)

16 71,74,86,88,89,90,99,100,103,104,106,112,117,118,119,ao,f	randomiseeritud uuringud	suur <sup>sp</sup>	suur <sup>bb</sup>	väike	väike	puudub	Keskmiste erinevus (ingl <i>mean difference</i> ) madala rasvasisaldusega diet ehk LF diet (n = 598) vs. kontroll (n = 478) = -0,40% (95% CI -0,59; -0,20)	 MADAL	KRIITILINE
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HbA1c muutus keskmiste erinevus (ingl *mean difference*) (kõrge valgusisaldusega ehk HP diet vs. kontroll)

Töendatuse astme hinnang							Mõju	Töendatuse aste	Olulisus
Uuringute arv	Uuringukavand	Nihke töenäosus	Töenduse ebakõla	Töenduse kaudsus	Töenduse ebätäpus	Muud kaalutlused			
5 79,80,83,109,114,ao	randomiseeritud uuringud	suur <sup>ap</sup>	väga suur <sup>bc</sup>	väike	väike	puudub	Keskmine erinevus (ingl mean difference) kõrge valgusisaldusega dieet ehk HP dieet (n = 50) vs. kontroll (n = 39) = -0,50% (95% CI -1,01; 0,00)	 VÄGA MADAL	KRIITILINE

HbA1c muutus keskmiste erinevus (ingl mean difference) (toidu asendamine ingl meal replacement vs. kontroll)

4 82,87,102,108,ao	randomiseeritud uuringud	suur <sup>ap</sup>	suur <sup>bd</sup>	väike	väike	puudub	Keskmine erinevus (ingl mean difference) toidu asendamine ingl meal replacement (n = 198) vs. kontroll (n = 167) = -0,56% (95% CI -0,91; -0,22)	 MADAL	KRIITILINE
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HbA1c muutus sekkumiste erinevus (ingl mean difference) (madal glükeemiline indeks vs. kontroll)

3 100,105,115,ao	randomiseeritud uuringud	suur <sup>ap</sup>	väike <sup>be</sup>	väike	väike	puudub	Keskmine erinevus (ingl mean difference) madal glükeemiline indeks (n = 90) vs. kontroll (n = 90) = -0,09% (95% CI -0,38; 0,19)	 KESKMININE	KRIITILINE
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Hüpoplükeeemia sagedus (%) dapaglifloosi vs. platseebo (järelkontroll: vahemik 12 nädalat kuni 24 nädalat)

4 bf,bg	randomiseeritud uuringud	suur <sup>bh</sup>	väike	väike	väike	puudub	dapaglifloosi monoteraapia n = 882 vs. platseebo n = 251. Hüpoplükeeemia sagedus platseebo 2,0% (n = 251), dapaglifloosi 2,5mg 2,5% (n = 321), dapaglifloosi 5mg 2,2% (n = 316), dapaglifloosi 10mg 2,9% (n = 245)	 KESKMININE	KRIITILINE
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Cl: usaldusintervall

## Selgitused

a. Franz et al. (2015). Lifestyle Weight-Loss Intervention Outcomes in Overweight and Obese Adults with Type 2 Diabetes: A Systematic Review and Meta-Analysis of Randomized Clinical Trials

b. Algsest oli tegemist RCT-dega. LOOK AHEAD (2007) intensiivne elustiilisekkumine (n=2570) vs toe pakkumine ja diabeedialane koolitus (n=2575); Metz (2000) ettevalmistatud toidukavad (n=149) vs tavaravi (n=153); Wolf (2004) elustiilialane juhtumikorraldus (n=74) vs tavaravi (n=73); Li (2005) soja baasil sõogi asendamine (n=46) vs individualiseeritud toidukava (n=36); West (2007) MI (n=109) vs mitte midagi (n=108); Brehm (2009) MUFA (n=52) vs kõrge süsivesikusisaldusega dieet (n=43); Davis (2009) madala süsivesikusisaldusega (n=55) vs madala rasvasisaldusega dieet (n=50); Espósito (2009) Vahemere dieet (n=108) vs madala rasvasisaldusega dieet (n=107); Larsen (2011) madala rasva ja kõrge valgusisaldusega (n=53) vs madala rasva kõrge süsivesikusisaldusega dieet (n=46); Krebs (2012) madala rasva ja kõrge valgusisaldusega (n=207) vs madala rasva kõrge süsivesikusisaldusega dieet (n=212); Gulbrand (2012) madala rasvasisaldusega (n=31) vs madala süsivesikusisaldusega dieet (n = 30)

c. Meta-analüüs autorid pole uuringute kvaliteeti hinnanud, põhjendades et kaasati ainult RCT. Kasutatud küll RCT-de andmeid kuid ümbergrueperimise töölt on antud analüüs uuringu kavandiks enne ja pärast sekkumist analüüs.

d. Uuringute usaldusintervallid ei kattu

e. Lai usaldusvahemik

f. võrdluste arv

g. Meid huvitavad olulised kardiovaskulaarsed sündmused, tegemist aga kardiovaskulaarset riski peegeldava näitajaga

h. väike valim

i. Umpierre et al. (2011) Physical Activity Advice Only or Structured Exercise Training and Association With HbA1c Levels in Type 2 Diabetes: A Systematic Review and Meta-analysis

j. Kokku käsitles struktureeritud treeninguid 23 RCT. Süsteematiilise ülevaate ja meta-analüüs autorid hindasid nihke riski. Hinnatud 23-st RCT 15-l on ebasele kuidas toimus randomiseerimine, 19 RCT puhul oli ebasele kas uuritavate uurimisrühmadesse jaotumine oli pimendatud (allocation concealment), ainult 2 RCT raporteeris ravikavatsusanalüüs (intention to treat) tulemusi, ülejäänud uuringu lõpetanute tulemusi.

k. Aeroobne treening: I<sup>2</sup> = 92,8%; p <0,001

l. Vastupidavstreening: I<sup>2</sup> = 92,5%; p <0,001

m. Kombineeritud treening: I<sup>2</sup> = 67,5%; p < 0,005

n. Füüsiline aktiivsuse alast nõustamist käsitleti 24 RCT-s. Süsteematiilise ülevaate ja meta-analüüs autorid hindasid uuringutes nihke riski. Hinnatud 24-st RCT 15-l oli ebasele kuidas toimus randomiseerimine, 20 RCT puhul oli ebasele kas uuritavate uurimisrühmadesse jaotumine oli pimendatud (allocation concealment), ainult 4 RCT raporteeris ravikavatsusanalüüs (intention to treat) tulemusi, ülejäänud uuringu lõpetanute tulemusi.

o. Füüsiline aktiivsuse alane nõustamine koos toitumisalase ko-interventsiooniga: I<sup>2</sup> = 57,5%; p = 0,007

p. toitumisalane nõustamine: I<sup>2</sup> = 61,2%; p = 0,001

q. Adolfsson et al (2007): gruupi põhine võimestamine vs diabeedi tavaravi; Brown et al. (2002) diabeedi enesejuhtimise alane sekkumine vs tavaravi; Davies et al. (2008) struktureeritud gruupi põhine diabeedi õppemöödu programm (DESMOND) vs tavaravi; Deakin et al. (2006) X-PERT programm vs individuaalne vastuvõtt; Rachmani et al. (2005) patsiendi osalusprogramm vs tavalik konsultatsioon; Sarkadi et al. (2003) diabeedilise õppemöödu programm vs kontroll; Whitemore et al. (2004) õe poolne juhendamine vs standardravi

r. Ontario HTA (2009). Behavioural Interventions for Type 2 Diabetes An Evidence-Based Analysis

s. Meta-analüüs autorid hinnanud nihke riski. Osades uuringutes ebasele uuritavate rühmadesse jagamise pimendamine (allocation concealment), ebasele kas tulemi hindajad olid pimendatud (4 RCT)

t. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 7,64; df = 7; p = 0,37; I<sup>2</sup> = 8%

u. Meta-analüüs koostajad hinnanud uuringute kvaliteet mõõdukaks

v. Tau<sup>2</sup> = 0,02; Chi<sup>2</sup> = 2,27; df = 2; p = 0,025; I<sup>2</sup> = 28%

w. Meta-analüüs koostajad hinnanud uuringute kvaliteedi madalaks

x. Tau<sup>2</sup> = 0,04; Chi<sup>2</sup> = 3,80; df = 2; p = 0,15; I<sup>2</sup> = 47%

y. Meta-analüüs koostajad hinnanud nihke riski. Osades uuringutes on uuritavate rühmadesse jagamise pimendamine ebasele (allocation concealment), ebasele kas tulemite hindajad olid pimendatud, ebasele kas oli kasutatud ravikavatsusanalüüs (intention to treat)

z. Ühe uuringu valimi moodustasid ainult Hispaania päritolu uuritavad

aa. Tau<sup>2</sup> = 0,02; Chi<sup>2</sup> = 2,27; df = 2; p = 0,25; I<sup>2</sup> = 28%

ab. Meta-analüüs koostajad hinnanud uuringute kvaliteeti madalaks

ac. Tau<sup>2</sup> = 0,01; Chi<sup>2</sup> = 4,49; df = 4; p = 0,34; I<sup>2</sup> = 11%

ad. Meta-analüüs koostajad hinnanud uuringute kvaliteeti. Kolmel uuringul mõõdukas ja kahel madal

ae. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 4,25; df = 4; p = 0,37; I<sup>2</sup> = 6

af. Süsteematiilise ülevaate ja meta-analüüs koostajad hinnanud uuringu kvaliteediks mõõdukas

ag. väike valim, lai usaldusvahemik

ah. Meta-analüüs koostajad hinnanud kaasatud RCT-de kvaliteeti. Kolme kaasatud uuringu hinnang mõõdukas ja kolme kaasatud uuringu hinnang madal.

ai. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 0,01; df = 1, p = 0,93; I<sup>2</sup> = 0%

aj. Meta-analüüs koostajad hinnanud nihke riski. Osades uuringutes on uuritavate rühmadesse jagamise pimendamine ebasele (allocation concealment), ebasele kas ühes RCT-s oli kasutatud ravikavatsusanalüüs (intention to treat), nihke risk ühes RCT-s tingituna pikast jälgimisperioodist (8 aastat)

ak. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 0,01; df = 1; p = 0,93; I<sup>2</sup> = 0%

al. Al-Shokri 2012 MNT vs toitumisalane tavaravi, Anderson-Loftin 2005 LF vs tavaravi/kontroll/diabeedi alane õpe, Andrews 2011 toitumisalane nõustamine vastavalt UK juhendile vs tavaravi, Azadbakht 2011 LF/DASH vs kontroll, Barnard 2009 LF vegan vs ADA juhend, Bowen 2016 kohandatud taldrikureegel & SV lugemise gruupi nõustamine vs kontroll, Carter 2016 HP/LF vs toiduenergia vähendamine, Cheskin 2008 portseljanit kontrolliv toidu asendamine vs ADA juhend, Coppel 2010 Euroopa diabeedijuhend vs kontroll/meditsiiniline vaatlus, Daly 2006 LC vs

tasakaalustatud toitumine, Dyson 2007 LC vs tasakaalustatu toitumine UK juhend, Elhayany 2010 LC Vahemere vs ADA 2003 juhend, Elhayany 2010 traditsiooniline Vahemere vs ADA 2003, Evangelista 2009 HP vs AHA tavadieet, Evangelista 2009 tavaline valgusisaldus vs AHA tavadieet, Foster 2009 portsjonit kontrolliv dieet vs diebedi alane õpe ja tugi, Franz 1995 MNT vs toitumisalane tavaravi, Gannon 2004 HP vs ADA & AHA dieet, Gannon 2003 HP ADA & AHA dieet

am. Goday 2016 VLC ketogeenne vs ADA madal kaloraaž, Goldstein 2011 LC vs ADA piiratud kaloraaž, Imai 2011 "jurvili enne SV" vs vahetusel põhinev toitumine, Itsipoilos 2011 Vahemere vs tavalitoitumine/kontroll, Jönsson 2009 Paleo vs diabedi dieet, Jung 2014 traditsiooniline Korea vs kontroll, Kahleova 2011 taimetoitus vs diabedi dieet/kontroll, Kaplan 1987 vahetusel põhinev dieet vs kontroll, Kattelmann 2010 toitumisalane õpe vs ADA, Kondo 2014 kalal põhinev vs kontroll, Koo 2010 LF & madal kaloraaž vs kontroll, Laitinen 1993 intensiivne toitumisalane õpe vs tavaravi/terviseõpetus, Li 2016 LF & kõrge kiudainesisaldusega vs tavaravi, Liu 2015 toituminalane sekkumine vs tavaravi, Luger 2013 HP vs tavaline dieet, Ma 2008 madal GI vs ADA, Mesci 2010 lihtsaid SV piirav toitumine vs diabedi diabeet, Miller 2002 toitumisalane õpe vs kontroll, Muchiri 2016 toitumisalane õpe vs tavaravi, O'Neill 2016 Kaalujäljigjad vs madal kaloraaž & kõrge kiudainesisaldus

an. Nicholson 1999 LC & taimetoitus vs kontroll, Nuttall 2008 HP vs AHA, Pedersen 2007 portsjoni kontroll vs toitumisalane tavaravi, Pi-Sunyer 1999 toidu asendamine vs ADA & vahetamine, Rock 2014 LF vs tavaravi, Rock 2014 LC vs tavaravi, Saslow 2017 Ketogeenne vs ADA taldrikuuregel, Sato LC vs Jaapani diabeediliidi kaloraaži vähendamisel põhinev, Shirai 2013 madala kaloraažiga & osaliselt toidu asendamine vs madala kaloraažiga, Talib 1997 grupipõhine toitumisalane nõustamine vs diabeedialane õpe, Trico 2016 HP & HF enne vs vähene kaloraaži vähendamine, Visek 2014 madal GI vs diabedi dieet, Williams 1998 väga madala kaloraažiga dieet (1 päev) vs käitumisteraapia, Williams 1998 väga madala kaloraažiga dieet vs käitumisteraapia, Yamada 2014 LC vs madala kaloraažiga dieet, Yang 2017 toitumisalane õpe vs Korea diabeedialane õpe, Yip 2001 vedel toidu asendamine vs ADA vahetus, Yusof 2009 madal GI vs tavaline vahetus, Ziemer 2003 vs vahetus

ao. Cradock (2017) Diet Behavior Change Techniques in Type 2 Diabetes: A Systematic Review and Meta-analysis

ap. Süsteematiilise ülevaate ja meta-analüüs koostajate poolt hinnatud nihke riski. Nihke risk valdavalt ebaselge "unclear"

aq. Tau<sup>2</sup> = 0,04; Chi<sup>2</sup> = 124,40; df = 58; p < 0,001 I<sup>2</sup> = 53%

ar. Tau<sup>2</sup> = 0,05; Chi<sup>2</sup> = 88,94; df = 34, p < 0,001; I<sup>2</sup> = 62%

as. Tau<sup>2</sup> = 0,03; Chi<sup>2</sup> = 45,01; df = 25; p = 0,008; I<sup>2</sup> = 44%

at. Tau<sup>2</sup> = 0,02; Chi<sup>2</sup> = 24,04; df = 15; p = 0,06; I<sup>2</sup> = 38%

au. Tau<sup>2</sup> = 0,19; Chi<sup>2</sup> = 12,35; df = 4; p = 0,01; I<sup>2</sup> = 68%

av. Tau<sup>2</sup> = 3,27; Chi<sup>2</sup> = 486,26; df = 53; p < 0,001; I<sup>2</sup> = 89%

aw. Tau<sup>2</sup> = 2,56; Chi<sup>2</sup> = 203,73; df = 32; p < 0,001; I<sup>2</sup> = 84%

ax. Tau<sup>2</sup> = 4,74; Chi<sup>2</sup> = 319,17; df = 23; p < 0,001; I<sup>2</sup> = 93%

ay. Tau<sup>2</sup> = 3,15; Chi<sup>2</sup> = 110,87; df = 13; p < 0,001; I<sup>2</sup> = 88%

az. Tau<sup>2</sup> = 0,41; Chi<sup>2</sup> = 4,10; df = 3; p = 0,25; I<sup>2</sup> = 27%

ba. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 2,94; df = 8; p = 0,94; I<sup>2</sup> = 0%

bb. Tau<sup>2</sup> = 0,09; Chi<sup>2</sup> = 45,02; df = 15; p = < 0,001; I<sup>2</sup> = 67%

bc. Tau<sup>2</sup> = 0,23; Chi<sup>2</sup> = 17,17; df = 4; p = 0,002; I<sup>2</sup> = 77%

bd. Tau<sup>2</sup> = 0,07; Chi<sup>2</sup> = 6,70; df = 3; p = 0,08; I<sup>2</sup> = 55%

be. Tau<sup>2</sup> = 0,00; Chi<sup>2</sup> = 0,64; df = 2; p = 0,73; I<sup>2</sup> = 0%

bf. Ptaszynska A, Johnsson KM, Parikh SJ, de Bruin TW, Apanovitch AM, List JF. Safety profile of dapagliflozin for type 2 diabetes: pooled analysis of clinical studies for overall safety and rare events. *Drug Saf.* 2014 Oct;37(10):815-29

bg. Monotherapy = NCT00528372, Low-dose monotherapy = NCT00736879, Monotherapy phase II = NCT00263276, Monotherapy (Japan) = NCT00972244

bh. pole hinnatud

## Viited

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