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Appendix A. Search Strategy

Database: Embase <1974 to 2013 October 03> Search Strategy:

- 1 retracted article/ (6992)
- 2 (random\$ or placebo\$ or single blind\$ or double blind\$ or triple blind\$).ti,ab. (964464)
- 3 (animal\$ not human\$).sh,hw. (3892889)
- 4 (book or conference paper or editorial or letter or review).pt. not exp randomized controlled trial/ (4026686)
- 5 1 or 2 (971306)
- 6 5 not (3 or 4) (788381)
- 7 exp cohort analysis/ (159852)
- 8 exp longitudinal study/ (65122)
- 9 exp prospective study/ (251233)
- 10 exp follow up/ (749090)
- 11 cohort\$.tw. (363883)
- 12 7 or 8 or 9 or 10 or 11 (1281760)
- 13 exp case-control study/ (90165)
- 14 (case\$ and control\$).tw. (421076)
- 15 13 or 14 (455349)
- 16 (case\$ and series).tw. (156704)
- 17 exp review/ (2051725)
- 18 (literature adj3 review\$).ti,ab. (210224)
- 19 exp meta analysis/ (76123)
- 20 exp "Systematic Review"/ (64783)
- 21 17 or 18 or 19 or 20 (2235229)
- 22 (medline or embase or pubmed or cinahl or amed or psychlit or psychinfo or scisearch or cochrane).ti,ab. (99517)
- 23 retracted article/ (6992)
- 24 22 or 23 (106462)
- 25 21 and 24 (78850)
- 26 (systematic\$ adj2 (review\$ or overview)).ti,ab. (66304)
- 27 (meta?anal\$ or meta anal\$ or metaanal\$ or metanal\$).ti,ab. (75185)
- 28 25 or 26 or 27 (159362)
- 29 (ae or si or to or co).fs. (3025512)
- 30 (safe or safety).ti,ab. (597602)
- 31 side effect\$.ti,ab. (238444)
- 32 ((adverse or undesireable or harm\$ or serious or toxic) adj3 (effect\$ or reaction\$ or event\$ or outcome\$)).ti,ab.

(406794)

- 33 exp adverse drug reaction/ (358463)
- 34 exp drug toxicity/ (77722)
- 35 exp intoxication/ (328776)
- 36 exp drug safety/ (221042)
- 37 exp drug monitoring/ (40454)
- 38 exp drug hypersensitivity/ (49258)
- 39 exp postmarketing surveillance/ (22410)
- 40 exp phase iv clinical trial/ (1496)

- 41 (toxicity or complication\$ or noxious or tolerability).ti,ab. (1146485)
- 42 exp postoperative complication/ (478856)
- 43 exp peroperative complication/ (19583)
- 44 or/29-43 (4747913)
- 45 retracted article/ (6992)
- 46 (random\$ or placebo\$ or single blind\$ or double blind\$ or triple blind\$).ti,ab. (964464)
- 47 (animal\$ not human\$).sh,hw. (3892889)
- 48 (book or conference paper or editorial or letter or review).pt. not exp randomized
- controlled trial/ (4026686)
- 49 45 or 46 (971306)
- 50 49 not (47 or 48) (788381)
- 51 exp cohort analysis/ (159852)
- 52 exp longitudinal study/ (65122)
- 53 exp prospective study/ (251233)
- 54 exp follow up/ (749090)
- 55 cohort\$.tw. (363883)
- 56 (51 or 52 or 53 or 54 or 55) not (47 or 48) (1147686)
- 57 exp case-control study/ (90165)
- 58 (case\$ and control\$).tw. (421076)
- 59 (57 or 58) not (47 or 48) (387052)
- 60 (case\$ and series).tw. (156704)
- 61 60 not (47 or 48) (136432)
- 62 exp review/ (2051725)
- 63 (literature adj3 review\$).ti,ab. (210224)
- 64 exp meta analysis/ (76123)
- 65 exp "Systematic Review"/ (64783)
- 66 62 or 63 or 64 or 65 (2235229)
- 67 (medline or embase or pubmed or cinahl or amed or psychlit or psychinfo or scisearch or cochrane).ti,ab. (99517)
- 68 retracted article/ (6992)
- 69 67 or 68 (106462)
- 70 66 and 69 (78850)
- 71 (systematic\$ adj2 (review\$ or overview)).ti,ab. (66304)
- 72 (meta?anal\$ or meta anal\$ or metaanal\$ or metanal\$).ti,ab. (75185)
- 73 (70 or 71 or 72) not (47 or 48) (79223)
- 74 kidney donor/ (6363)
- 75 living donor/ (16693)
- 76 kidney/ (259788)
- 77 (74 or 76) and 75 (3011)
- 78 (liv\$ and kidney and don\$).ti. (2260)
- 79 77 or 78 (4371)
- 80 73 and 79 (14)
- 81 50 and 79 (139)
- 82 56 and 79 (853)
- 83 59 and 79 (83)
- 84 61 and 79 (46)
- 85 80 or 81 or 82 or 83 or 84 (1032)
- 86 44 and 79 (2185)
- 87 86 not (3 or 4) (1680)
- 88 87 not 85 (1027)

Database: Ovid MEDLINE(R) <1946 to October Week 1 2013> Search Strategy:

- 1 meta analysis as topic/ (14064) 2 meta-analy\$.tw. (57155) 3 metaanaly\$.tw. (1274) 4 meta-analysis/ (51068) 5 (systematic adj (review\$1 or overview\$1)).tw. (46408) 6 exp Review Literature as Topic/ (7621) 7 or/1-6 (114404) 8 cochrane.ab. (33039) 9 embase.ab. (29462) 10 (psychlit or psyclit).ab. (1189) 11 (psychinfor or psycinfo).ab. (8174) 12 or/8-11 (47806) 13 reference list\$.ab. (11604) 14 bibliograph\$.ab. (11732) 15 hand search.ab. (865) 16 relevant journals.ab. (895) 17 manual search\$.ab. (2220) 18 or/13-17 (25483) 19 selection criteria.ab. (26028) 20 data extraction.ab. (10019) 21 19 or 20 (33588) 22 review/ (1912544) 23 21 and 22 (25897) 24 comment/ (533153) 25 letter/ (803396) 26 editorial/ (334975) 27 animal/ (5486090) 28 human/ (13631608) 29 27 not (28 and 27) (3957888) 30 7 or 12 or 18 or 23 (143113) 31 randomized controlled trials as topic/ (102017) 32 randomized controlled trial/ (387734) 33 random allocation/ (81475) 34 double blind method/ (131321) 35 single blind method/ (19455) 36 clinical trial/ (503981) 37 clinical trial, phase i.pt. (16097) 38 clinical trial, phase ii.pt. (26744) 39 clinical trial, phase iii.pt. (10077) 40 clinical trial, phase iv.pt. (985) 41 controlled clinical trial.pt. (89736) 42 randomized controlled trial.pt. (387734) 43 multicenter study.pt. (181196) 44 clinical trial.pt. (503981) 45 exp Clinical trials as topic/ (295298) 46 or/31-44 (954164) 47 (clinical adj trial\$).tw. (210053) 48 ((singl\$ or doubl\$ or treb\$ or tripl\$) adj (blind\$3 or mask\$3)).tw. (129006)
- 49 placebos/ (33702)

50 placebo\$.tw. (161007) 51 randomly allocated.tw. (15955) 52 (allocated adj2 random\$).tw. (18445) 53 47 or 48 or 49 or 50 or 51 or 52 (415598) 54 46 or 53 (1119857) 55 case report.tw. (183596) 56 case report.tw. (183596) 57 letter/ (803396) 58 historical article/ (299710) 59 55 or 56 or 57 or 58 (1275472) 60 54 not 59 (1096065) 61 exp cohort studies/ (1361688) 62 cohort\$.tw. (260919) 63 controlled clinical trial.pt. (89736) 64 epidemiologic methods/ (30937) 65 limit 64 to yr=1971-1983 (5365) 66 61 or 62 or 63 or 65 (1535709) 67 exp case-control study/ (660962) 68 (case\$ and control\$).tw. (312502) 69 67 or 68 (885537) 70 epidemiologic studies/ (6242) 71 (follow up adj stud\$).tw. (36469) 72 longitudinal.tw. (132878) 73 (observational adj stud\$).tw. (40485) 74 retrospective.tw. (251881) 75 cross sectional.tw. (156469) 76 cross-sectional studies/ (179104) 77 or/70-76 (663510) 78 exp Living Donors/ (10898) 79 exp Kidney/ (301075) 80 Kidney Transplantation/ (80731) 81 kidney.ti. (105471) 82 79 or 80 or 81 (403205) 83 78 and 82 (4883) 84 (liv\$ and kidney and don\$).ti. (1566) 85 83 or 84 (5223) 86 30 and 85 (52) 87 85 and 60 (418) 88 66 and 85 (1899) 89 85 and (69 or 77) (1415)

Appendix B. Unextracted and Excluded Studies

Peri/Post Nephrectomy Outcomes: Eligible Studies-Not extracted

- 1. Aboutaleb E, Herbert P, Crane J, et al. Mini-incision donor nephrectomy techniques: a systematic review. Experimental & Clinical Transplantation: Official Journal of the Middle East Society for Organ Transplantation. 2010 Sep;8(3):189-95. PMID 20716035. PP-OR-NO Extract
- 2. Adiyat KT, Tharun BK, Shetty A, et al. Comparison of three different techniques of extraction in laparoscopic donor nephrectomy. Indian Journal of Urology. 2013 Jul;29(3):184-7. PMID 24082437. *PP-OR-NO Extract*
- 3. Aguiar WF, Passerotti CC, Claro JF, et al. Mini-incisions by lombotomy or subcostal access in living kidney donors: a randomized trial comparing pain, safety, and quality of life. Clinical Transplantation. 2007 Mar-Apr;21(2):269-76. PMID 17425757. PP-OR-NO Extract
- 4. Ahmadi AR, Lafranca JA, Claessens LA, et al. Shifting paradigms in eligibility criteria for live kidney donation: a systematic review. Kidney international. 2014. *LT-SR-Extract*
- 5. Amirzargar MA, Babolhavaeji H, Hosseini SA, et al. The new technique of using the epigastric arteries in renal transplantation with multiple renal arteries. Saudi Journal of Kidney Diseases & Transplantation. 2013 Mar;24(2):247-53. PMID 23538346. PP-OR-NO Extract
- 6. Anderson KM, Lindler TU, Lamberton GR, et al. Laparoscopic donor nephrectomy: effect of perirenal fat upon donor operative time. Journal of Endourology. 2008 Oct;22(10):2269-74. PMID 18831674. *PP-OR-NO Extract*
- 7. Arai K, Nishiyama T, Hara N, et al. Retroperitoneoscopic donor nephrectomy with a gel-sealed hand-assist access device. BMC Urology. 2013;13:7. PMID 23374442. *PP-OR-NO Extract*
- 8. Ashraf HS, Hussain I, Siddiqui AA, et al. The outcome of living related kidney transplantation with multiple renal arteries. Saudi Journal of Kidney Diseases & Transplantation. 2013 May;24(3):615-9. PMID 23640649. *PP-OR-NO Extract*
- 9. Aull MJ, Afaneh C, Charlton M, et al. A randomized, prospective, parallel group study of laparoscopic versus laparoendoscopic single site donor nephrectomy for kidney donation. American Journal of Transplantation. 2014 Jul;14(7):1630-7. PMID 24934732. *PP-OR-NO Extract*
- 10. Bachmann A, Wolff T, Ruszat R, et al. Retroperitoneoscopic donor nephrectomy: a retrospective, non-randomized comparison of early complications, donor and recipient outcome with the standard open approach. European Urology. 2005 Jul;48(1):90-6; discussion 6. PMID 15967257. *PP-OR-NO Extract*
- 11. Bachmann A, Wyler S, Wolff T, et al. Complications of retroperitoneoscopic living donor nephrectomy: single center experience after 164 cases. World Journal of Urology. 2008 Dec;26(6):549-54. PMID WOS:000260856600005. *PP-OR-NO Extract*
- 12. Bargman V, Sundaram CP, Bernie J, et al. Randomized trial of laparoscopic donor nephrectomy with and without hand assistance. Journal of Endourology. 2006 Oct;20(10):717-22. PMID 17094745. *PP-OR-NO Extract*

- 13. Becker BN, Becker YT. Rehospitalization after living kidney donation. Clinical Journal of The American Society of Nephrology: CJASN. 2014 Feb;9(2):227-8. PMID 24458083. *PP-OR-NO Extract*
- 14. Bergman S, Feldman LS, Anidjar M, et al. "First, do no harm": monitoring outcomes during the transition from open to laparoscopic live donor nephrectomy in a Canadian centre. Canadian Journal of Surgery. 2008 Apr;51(2):103-10. PMID 18377750. PP-OR-NO Extract
- 15. Biglarnia A, Bergqvist D, Johansson M, et al. Venous thromboembolism in live kidney donors--a prospective study. Transplantation. 2008 Sep 15;86(5):659-61. PMID 18791446. *PP-OR-NO Extract*
- 16. Brockmann JG, Senninger N, Wolters HH. Living donor of the kidney-open-video. Langenbecks Archives of Surgery. 2007 May;392(3):219-25. PMID 17375320. *PP-OR-NO Extract*
- 17. Buresley S, Samhan M, Al-Mousawi M. Kuwait experience in laparoscopic donor nephrectomy: first 80 cases. Transplantation Proceedings. 2007 May;39(4):813-5. PMID 17524819. *PP-OR-NO Extract*
- 18. Cannon RM, Eng M, Marvin MR, et al. Laparoscopic living kidney donation at a single center: an examination of donor outcomes with increasing experience. American Surgeon. 2011 Jul;77(7):911-5. PMID 21944358. *PP-OR-NO Extract*
- 19. Chandak P, Kessaris N, Challacombe B, et al. How safe is hand-assisted laparoscopic donor nephrectomy?--results of 200 live donor nephrectomies by two different techniques. Nephrology Dialysis Transplantation. 2009 Jan;24(1):293-7. PMID 18711221. *PP-OR-NO Extract*
- 20. Cherif M, Ounissi M, Karoui C, et al. Short- and Long-Term Outcomes of Living Donors in Tunisia: A Retrospective Study. Transplantation Proceedings. 2010 Dec;42(10):4311-3. PMID WOS:000285732200105. *PP-OR-NO Extract*
- 21. Chin EH, Hazzan D, Herron DM, et al. Laparoscopic donor nephrectomy: intraoperative safety, immediate morbidity, and delayed complications with 500 cases. Surgical Endoscopy. 2007 Apr;21(4):521-6. PMID 17180288. *PP-OR-NO Extract*
- 22. Cho HJ, Choi YS, Bae WJ, et al. Another option for laparoscopic living donor nephrectomy: a single center experience comparing two-port versus hand-assisted technique. Journal of Endourology. 2013 May;27(5):587-91. PMID 23228097. PP-OR-NO Extract
- 23. Chow GK, Prieto M, Bohorquez HE, et al. Hand-assisted laparoscopic donor nephrectomy for morbidly obese patients. Transplantation Proceedings. 2002;34(2):728. PMID 2002171828. *PP-OR-NO Extract*
- 24. Crane C, Lam VW, Alsakran A, et al. Are there anatomical barriers to laparoscopic donor nephrectomy? ANZ Journal of Surgery. 2010 Nov;80(11):781-5. PMID 20969683. *PP-OR-NO Extract*
- 25. Crotty C, Tabbakh Y, Hosgood SA, et al. Systemic heparinisation in laparoscopic live donor nephrectomy. Journal of transplantation. 2013;2013:138926. PMID 24455192. *PP-OR-NO Extract*
- 26. Dahm F, Weber M, Muller B, et al. Open and laparoscopic living donor nephrectomy in Switzerland: a retrospective assessment of clinical outcomes and the motivation to donate. Nephrology Dialysis Transplantation. 2006 Sep;21(9):2563-8. PMID 16702206. *PP-OR-NO Extract*
- 27. Dinckan A, Dinc B, Turkyilmaz S, et al. Comparison of open and retroperitonoscopic donor nephrectomy in terms of lipid and protein peroxidation responses. Transplantation Proceedings. 2013 Nov;45(9):3214-9. PMID 24182787. *PP-OR-NO Extract*
- 28. Diner EK, Radolinski B, Murdock JD, et al. Right laparoscopic donor nephrectomy: the Washington Hospital Center experience. Urology. 2006 Dec;68(6):1175-7. PMID 17169641. *PP-OR-NO Extract*

- 29. Dolce CJ, Keller JE, Walters KC, et al. Laparoscopic versus open live donor nephrectomy: outcomes analysis of 266 consecutive patients. Surgical Endoscopy. 2009 Jul;23(7):1564-8. PMID 19263157. *PP-OR-NO Extract*
- 30. Dols LF, Kok NF, d'Ancona FC, et al. Randomized controlled trial comparing hand-assisted retroperitoneoscopic versus standard laparoscopic donor nephrectomy. Transplantation. 2014 Jan 27;97(2):161-7. PMID 24092379. *PP-OR-NO Extract*
- 31. Dols LF, Kok NF, Terkivatan T, et al. Optimizing left-sided live kidney donation: hand-assisted retroperitoneoscopic as alternative to standard laparoscopic donor nephrectomy. Transplant International. 2010 Apr 1;23(4):358-63. PMID 19886969. PP-OR-NO Extract
- 32. Duchene DA, Woodruff DY, Gallagher BL, et al. Successful outcomes of older donors in laparoscopic donor nephrectomy. Journal of Endourology. 2010 Oct;24(10):1593-6. PMID 20836718. *PP-OR-NO Extract*
- 33. Eroglu M, Guvence N, Kiper A, et al. Rib resection for live-donor nephrectomy. International Urology and Nephrology. 2005 December;37(4):675-9. PMID 2006043230. *PP-OR-NO Extract*
- 34. Fehrman-Ekholm I, Moller S, Steinwall J, et al. Single or double arteries in the remnant kidney after donation: influence on the long-term outcome of the donor. Transplantation Proceedings. 2009 Mar;41(2):764-5. PMID 19328974. *PP-OR-NO Extract*
- 35. Friedersdorff F, Werthemann P, Cash H, et al. Outcomes after laparoscopic living donor nephrectomy: comparison of two laparoscopic surgeons with different levels of expertise. BJU International. 2013 Jan;111(1):95-100. PMID 22757693. PP-OR-NO Extract
- 36. Friedman AL, Cheung K, Roman SA, et al. Early clinical and economic outcomes of patients undergoing living donor nephrectomy in the United States. Archives of Surgery. 2010 Apr;145(4):356-62; discussion 62. PMID 20404286. *PP-OR-NO Extract*
- 37. Genc V, Ozgencil E, Orozakunov E, et al. Pure laparoscopic versus open live donor nephrectomy: evaluation of health survey and graft functions. Transplantation Proceedings. 2011 Apr;43(3):791-4. PMID 21486599. *PP-OR-NO Extract*
- 38. Goh YS, Cheong PS, Lata R, et al. A necessary step toward kidney donor safety: the transition from locking polymer clips to transfixion techniques in laparoscopic donor nephrectomy. Transplantation Proceedings. 2014 Mar;46(2):310-3. PMID 24655950. *PP-OR-NO Extract*
- 39. Gorodner V, Horgan S, Galvani C, et al. Routine left robotic-assisted laparoscopic donor nephrectomy is safe and effective regardless of the presence of vascular anomalies. Transplant International. 2006 Aug;19(8):636-40. PMID 16827680. PP-OR-NO Extract
- 40. Greco F, Hoda MR, Alcaraz A, et al. Laparoscopic living-donor nephrectomy: Analysis of the existing literature. European Urology. 2010 October;58(4):498-509. PMID 2010493890. *PP-OR-NO Extract*
- 41. Guo FF, Shao ZQ, Yang WY, et al. Clinical analysis of living related renal transplantation with donors older than 50 years in China. Transplantation Proceedings. 2010 Sep;42(7):2471-6. PMID 20832526. *PP-OR-NO Extract*
- 42. Gupta M, Singh P, Dubey D, et al. A comparison of kidney retrieval incisions in laparoscopic transperitoneal donor nephrectomy. Urologia Internationalis. 2008;81(3):296-300. PMID 18931546. *PP-OR-NO Extract*
- 43. Gures N, Gurluler E, Berber I, et al. Comparison of the right and left laparoscopic live donor nephrectomies: a clinical case load. European Review for Medical & Pharmacological Sciences. 2013 May;17(10):1389-94. PMID 23740454. *PP-OR-NO Extract*

- 44. Hadjianastassiou VG, Johnson RJ, Rudge CJ, et al. 2509 Living donor nephrectomies, morbidity and mortality, including the UK introduction of laparoscopic donor surgery. American Journal of Transplantation. 2007 November;7(11):2532-7. PMID 2007497988. *PP-OR-NO Extract*
- 45. Hakim NS. A fast and safe living donor "finger assisted" nephrectomy technique. International Surgery. 2007 Sep-Oct;92(5):304-7. PMID 18399104. *PP-OR-NO Extract*
- 46. Harper JD, Breda A, Leppert JT, et al. Experience with 750 consecutive laparoscopic donor nephrectomies--is it time to use a standardized classification of complications? Journal of Urology. 2010 May;183(5):1941-6. PMID 20303114. *PP-OR-NO Extract*
- 47. Hawasli A, Berri R, Meguid A, et al. Total laparoscopic live donor nephrectomy: a 6-year experience. American Journal of Surgery. 2006 Mar;191(3):325-9. PMID 16490540. *PP-OR-NO Extract*
- 48. Heimbach JK, Taler SJ, Prieto M, et al. Obesity in living kidney donors: clinical characteristics and outcomes in the era of laparoscopic donor nephrectomy. American Journal of Transplantation. 2005 May;5(5):1057-64. PMID 15816886. PP-OR-NO Extract
- 49. Helal I, Abdallah TB, Ounissi M, et al. Short- and long-term outcomes of kidney donors: a report from Tunisia. Saudi Journal of Kidney Diseases & Transplantation. 2012 Jul;23(4):853-9. PMID 22805410. *PP-OR-NO Extract*
- 50. Hofker HS, Nijboer WN, Niesing J, et al. A randomized clinical trial of living donor nephrectomy: a plea for a differentiated appraisal of mini-open muscle splitting incision and hand-assisted laparoscopic donor nephrectomy. Transplant International. 2012 Sep;25(9):976-86. PMID 22849958. *PP-OR-NO Extract*
- 51. Horgan S, Galvani C, Gorodner MV, et al. Effect of robotic assistance on the "learning curve" for laparoscopic hand-assisted donor nephrectomy. Surgical Endoscopy. 2007 Sep;21(9):1512-7. PMID 17287916. *PP-OR-NO Extract*
- 52. Hung CJ, Lin YJ, Chang SS, et al. Kidney grafts with multiple renal arteries is no longer a relative contraindication with advance in surgical techniques of laparoscopic donor nephrectomy. Transplantation Proceedings. 2012 Jan;44(1):36-8. PMID 22310572. *PP-OR-NO Extract*
- 53. Inoue T, Tsuchiya N, Narita S, et al. Laparoendoscopic single-site plus one trocar donor nephrectomy using the GelPort: initial clinical experience. Urology. 2013 Feb;81(2):308-12. PMID 23374790. *PP-OR-NO Extract*
- 54. Izquierdo L, Peri L, Alvarez-Vijande R, et al. Audit of an initial 100 cases of laparoscopic live donor nephrectomy. Transplantation Proceedings. 2010 Nov;42(9):3437-9. PMID 21094792. *PP-OR-NO Extract*
- 55. Jacobs SC, Cho E, Dunkin BJ, et al. Laparoscopic nephrectomy in the markedly obese living renal donor. Urology. 2000 Dec 20;56(6):926-9. PMID 11113733. *PP-OR-NO Extract*
- 56. Jacobs SC, Ramey JR, Sklar GN, et al. Laparoscopic kidney donation from patients older than 60 years. Journal of the American College of Surgeons. 2004 Jun;198(6):892-7. PMID 15194070. *PP-OR-NO Extract*
- 57. Jeon H, Johnston TD, Strup SE, et al. University of Kentucky experience with laparoscopic live donor nephrectomy using two different techniques. International Surgery. 2006 Nov-Dec;91(6):332-5. PMID 17256431. *PP-OR-NO Extract*
- 58. Johnson SR, Khwaja K, Pavlakis M, et al. Older living donors provide excellent quality kidneys: a single center experience (older living donors). Clinical Transplantation. 2005 Oct;19(5):600-6. PMID WOS:000231678900005. *PP-OR-NO Extract*

- 59. Kanashiro H, Falci R, Jr., Piovisan AC, et al. Subcostal mini incision: a good option for donor nephrectomy. Clinics (Sao Paulo, Brazil). 2010 May;65(5):507-10. PMID 20535369. *PP-OR-NO Extract*
- 60. Klop KW, Kok NF, Dols LF, et al. Can right-sided hand-assisted retroperitoneoscopic donor nephrectomy be advocated above standard laparoscopic donor nephrectomy: a randomized pilot study. Transplant International. 2014 Feb;27(2):162-9. PMID 24268098. *PP-OR-NO Extract*
- 61. Kocak B, Baker TB, Koffron AJ, et al. Laparoscopic living donor nephrectomy: a single-center sequential experience comparing hand-assisted versus standard technique. Urology. 2007 Dec;70(6):1060-3. PMID 18158014. *PP-OR-NO Extract*
- 62. Kocak B, Koffron AJ, Baker TB, et al. Proposed classification of complications after live donor nephrectomy. Urology. 2006 May;67(5):927-31. PMID 16698353. *PP-OR-NO Extract*
- 63. Kok NF, Alwayn IP, Lind MY, et al. Donor nephrectomy: mini-incision muscle-splitting open approach versus laparoscopy. Transplantation. 2006 Mar 27;81(6):881-7. PMID 16570012. *PP-OR-NO Extract*
- 64. Kok NF, Alwayn IP, Tran KT, et al. Psychosocial and physical impairment after mini-incision open and laparoscopic donor nephrectomy: A prospective study. Transplantation. 2006 Nov 27;82(10):1291-7. PMID 17130777. *PP-OR-NO Extract*
- 65. Kok NF, JN IJ, Schouten O, et al. Laparoscopic donor nephrectomy in obese donors: easier to implement in overweight women? Transplant International. 2007 Nov;20(11):956-61. PMID 17635838. *PP-OR-NO Extract*
- 66. Kok NF, Lind MY, Hansson BM, et al. Comparison of laparoscopic and mini incision open donor nephrectomy: single blind, randomised controlled clinical trial.[Reprint in Ned Tijdschr Geneeskd. 2007 Jun 16;151(24):1352-60; PMID: 17665628]. BMJ. 2006 Jul 29;333(7561):221. PMID 16847014. *PP-OR-NO Extract*
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Appendix C. Peri/Post-Surgical Outcomes: Supporting Tables

Table C1. Peri/Post-Surgical Outcomes: Characteristics of Included Systematic Reviews

Study	Donor Population	Literature Search	Comparison Population	No of participants (studies)	Length of Follow- up, years (range)	Age (mean)	Sex (% women)	Outcomes
Fonouni 2014	Donors undergoing open donor nephrectomy.	PubMed	Donors undergoing laparoscopic nephrectomy.	N= NR (15 studies)	NR	NR	NR	Warm ischemia time, operative blood loss, operative time, perioperative complications
Lafranca 2013	All studies that focused on outcomes of laparoscopic living donor nephrectomy by Body Mass Index of the donors.	MEDLINE, EMBASE and CENTRAL databases were searched from inception to Jan 2011	Living donors from alternative BMI categories	N=7338 donors (14 studies)	1 week – 11 years	NR	NR	Operation duration, conversion risk, estimated blood loss, length of stay, perioperative complications, difference in serum creatinine, decrease in GFR
Liu 2014	Donors undergoing left laparoscopic nephrectomy	PubMed, Embase, Cochrane, Web of Science from inception to July 2013	Donors undergoing right laparoscopic nephrectomy	N=32,426 (29 studies)	NR	NR	NR	Intra and Post- operative complications, delayed renal function, conversion, warm ischemia time, operative time, length of hospital stay
Wilson 2011	Donors undergoing laparoscopic donor nephrectomy	MEDLINE (Jan 1966 – Jan 2010), EMBASE (Jan 1980 – Jan 2010),, and CENTRAL (The	Donors undergoing open donor nephrectomy	N= 596 donors (6 studies)	NR	NR	NR	Analgesia requirements, duration of procedures, blood loss, perioperative complications, reoperations, warm ischaemia time, hospital stay

Study	Donor Population	Literature Search	Comparison Population	No of participants (studies)	Length of Follow- up, years (range)	Age (mean)	Sex (% women)	Outcomes
		Cocherane Library 2010, Issue 2) diatabases.						
Young 2008		Literature search in MELINE 1950-Jan, 2008; EMBASE 1980-Jan, 2008; CINAHL (1982-Jan, 2008; BIOSIS 1969-Jan, 2008, Cochrane Library.			< 1 year	NR	NR	Perioperative outcomes, reported clinical and intermediate outcomes
Yuan 2013	Donors undergoing open donor nephrectomy or standard laparoscopic nephrectomy.	MEDLINE, EMBASE and CENTRAL databases were searched from inception to Oct 2011	Donors undergoing standard laparoscopic nephrectomy or hand- assisted laparoscopic nephrectomy.	N=2243 donors (30 studies)	NR	NR	NR	Operative time, warm ischemia time, intraoperative blood loss, hospital stay and time to return to work

BMI= Body Mass Index; CICr= creatinine clearance; CMS= Center for Medicare and Medicaid Services; CV = Cardiovascular; ESRD = End-stage renal disease; GFR = Glomerular Filtration Rate; HTN = hypertension; IFG = Impaired Fasting Glucose; MetS = Metabolic Syndrome defined according to NCEP ATP III guidelines as the presence of 3 or more criteria: waist ci (1) waist circumference \geq 88 cm in women and \geq 102 cm in men; (2) hypertriglyceridemia

 $(\geq 150 \text{ mg/dL} [-1.69 \text{ mmol/L}]$ or treatment); (3) low high-density lipid in– cholesterol (HDL_C) (<50 mg/dL [_1.29 mmol/L] in women and <40 mg/dL [1.04 mmol/L] in men); (4) hyperglycemia ($\geq 100 \text{ mg/dL} [_5.6 \text{ mmol/L}]$); and (5) hypertension ($\geq 130/85 \text{ mm}$ Hg or treatment).; NDI= National Death Index; NR = Not reported; National Health and Nutrition Examination Survey (NHANES III) conducted between 1988 and 1994; OPTN= Organ Procurement and Transplantation Network; SBP = systolic blood pressure; SSDI= Social Security Death Master File

DvD= donor versus donor; DvND=donor versus non-dono

Systematic Review	<i>A priori</i> design	Duplicate extraction	Comprehen sive lit search	Publication status = inclusion criteria	Study list provided	Study characteristics provided	Scientific quality rated	Scientific quality used appropriately	Findings combining appropriate	Publication bias assessed	Conflicts of interest stated	Overall Quality Rating	Summary Statement
Fonouni 2014	No	No	No	No	Included: Yes Excluded: No	No	No	No	Yes	No	Yes	Low	This is a review of RCTs and meta-analyses attempting to assess the efficacy and safety of laparoscopic versus open live donor nephrectomy. It is unclear if this was a systematic review, since details of the review methods are very limited. The results seem to be summarized appropriately and the conclusions are based on findings that appeared largely consistent across the included studies.
Lafranca 2013	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	Yes	Yes	yes	Yes	Yes	Moderate/ High	This is a recent systematic review of observational studies attempting to assess the impact of BMI on laparoscopic live kidney donation. The evidence base is weak, but reasonable methods were used in the review.

Table C2. Peri/Post-Surgical Outcomes: AMSTAR Assessments of Included Systematic Reviews

Liu 2014	Yes	Yes	Yes	Yes	Included: Yes Excluded: No	Yes	Yes	Yes	Yes	Yes	Yes	High	This is a systematic review of left vs right laparoscopic live donor nephrectomy (1 RCT and 28 observational studies). The evidence base was weak but the methods of review are well documented and of high quality.
Wilson 2011	Yes	Yes	Yes	Yes	included - Yes excluded - Yes	Yes	Yes	Yes	Yes	No	Yes	High	This is a recent Cochrane systematic review of RCT and CCT studies attempting to assess the impact of laproscopic vs open live kidney donation. The evidence base and methods of the review are both well documented and of high quality.
Young 2008	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	No	Can't Answer	Yes	No	Yes	Moderate	This is a systematic review of observational studies attempting to assess the impact of live kidney donation among donors with isolated medical abormalities. The manuscript states that high quality methods were used but details are difficult to confirm regarding how study bias was incorporated into the review.

Yuan 2013	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	Yes	Yes	Yes	No	No	Moderate/ High	This is a recent systematic review of RCT and CCT studies attempting to assess the impact of laproscopic vs open livekidney donation. The evidence base and methods of the review are reasonable to answer the research question. Authors may have conflicts of interpet and publication bias
													interest and publication bias was suspected.

Table C3. Open versus Laparoscopic Nephrectomy: Peri/Post SurgicalOutcomes (1a)

Systematic	Intervention		Results		AMSTAR
Review	Arm 1	Arm 2	Arm 1	Arm 2	Assessment
PeriOperative C	Complications				
Fonouni 2014	Open Donor Nephrectomy (NR)	Laparoscopic Donor Nephrectomy (NR)	No quantitative sy reported. Data pre includes trials and reviews including	esented d systematic	Low/Moderate
Yuan 2013 12 studies	Open Donor Nephrectomy (569)	Laparoscopic Donor Nephrectomy (792)	OR (95% CI) 0.80 (0.56-1.14)		Moderate/High
Wilson 2011 5 trials	Open Donor Nephrectomy (264)	Laparoscopic Donor Nephrectomy (292)	RR (95% CI) 0.87 (0.47-1.59)		High
Operative Time					
Nephrectomy (NR)		Laparoscopic Donor Nephrectomy (NR)	No quantitative sy reported. Data pre includes trials and reviews including	esented I systematic	Low/Moderate
Yuan 2013 19 studies	Open Donor Nephrectomy (807)	Laparoscopic Donor Nephrectomy(1093)	MD (95% CI) 50.54 (32.66 to 68		Moderate/High
Wilson 2011 6 trials	Open Donor Nephrectomy (284)	Laparoscopic Donor Nephrectomy (312)	No quantitative sy reported. Six trials significantly longe time with laparoso	s show er operative	High
Intraoperative I	Blood Loss				
Fonouni 2014	Open Donor Nephrectomy (NR)	Laparoscopic Donor Nephrectomy (NR)	No quantitative synthesis reported. Data presented includes trials and systematic reviews including those trials.		Low/Moderate
Yuan 2013 8 trials	Open Donor Nephrectomy (352)	Laparoscopic Donor Nephrectomy (371)	MD (95% CI) -101.23 (-153.52	to -48.94)	Moderate/High
Wilson 2011 5 trials	Open Donor Nephrectomy (NR)	Laparoscopic Donor Nephrectomy (NR)	No quantitative sy reported. Four of similar blood loss	five trials show	High
Reoperation Fonouni 2014	Open Donor Nephrectomy (NR)	Laparoscopic Donor Nephrectomy (NR)	No quantitative sy reported. Data pro includes trials and reviews including	esented I systematic	Low/Moderate
Wilson 2011 6 trials	Open Donor Nephrectomy (284)	Laparoscopic Donor Nephrectomy (312)	RR (95% CI) 0.57 (0.09 to 3.64 ARI (laparoscopic	.)	High
Length of Hosp				•	
Fonouni 2014	ouni 2014 Open Donor Laparoscopic Dono Nephrectomy (NR) (NR)		No quantitative sy reported. Data pre includes trials and reviews including	Low/Moderate	
Yuan 2013 16 trials	Open Donor Nephrectomy (709)	Laparoscopic Donor Nephrectomy (972)	MD (95% CI) -1.27 (-1.72 to -0.	82)	Moderate/High
	1	L	1		1

Wilson et al 5 trials	Open Donor Nephrectomy (292)	Laparoscopic Donor Nephrectomy (237)	No quantitative synthesis reported. Three of five trials report shorter hospital stay with laparoscopic.	High
Time to Return t	o Work (days)			
Yuan 2013 16 trials	Open Donor Nephrectomy (435)	Laparoscopic Donor Nephrectomy (581)	WMD (95% CI) -16.35 (-23.00 to -9.71)	Moderate/High

ARI=absolute risk increase: ARR=absolute risk reduction; RR=relative risk; WMD=weighted mean difference;

Table C4. Standard Laparoscopic versus Hand-Assisted LaparoscopicNephrectomy: Peri/Post Surgical Outcomes (1b)

Systematic	Intervention		Results		AMSTAR
Review	Arm 1	Arm 2	Arm 1	Arm 2	Assessment
Peri-/post-ope	rative Complicati	ons			
Yuan 2013 7 trials	Laparoscopic Nephrectomy (183) Laparoscopic Nephrectomy (159)		OR (95% CI) 0.62 (0.27 to		Moderate/High
Operative Tim	e				
Yuan 2013 9 trials	Standard Laparoscopic Nephrectomy (221)	Hand-Assisted Laparoscopic Nephrectomy (221)	WMD (95% (-24.55 (-50.8		Moderate/High
Intraoperative	Blood Loss	-			
Yuan 2013 6 trials	Standard Laparoscopic Nephrectomy (160)	Hand-Assisted Laparoscopic Nephrectomy (134)	WMD (95% (-20.65 (-43.8		Moderate/High
Length of Hos	pital Stay (days)				·
Yuan 2013 6 trials	Standard Laparoscopic Nephrectomy (170)	tandard Hand-Assisted aparoscopic ephrectomy Nephrectomy		0.56)	Moderate/High

Systematic	Intervention		Results	•	AMSTAR
Review	Arm 1	Arm 2	Arm 1	Arm 2	Assessment
Peri-/post-ope	rative Complication	ons			
Liu 2014 21 studies	Left (1872)	Right (728)	OR (95% C 1.31 (0.89 t	,	Moderate/High
Liu 2014 16 studies	Left (1792)	Right (675)	OR (95% C 1.27 (0.86 t		Moderate/High
Operative Tim	e				
Liu 2014 14 studies	Left (2193)	Right (463)	WMD (95% 1.35 (-11.73		Moderate/High
Intraoperative	Blood Loss				
Liu 2014 15 studies	Left (2356)	Right (677)	WMD (95% 4.36 (-19.83	,	Moderate/High
Length of Hosp	oital Stay (days)	I	I		I
Liu 2014 11 studies			WMD (95% 0.05 (-0.08		Moderate/High

Table C5. Left versus right laparoscopic live donor nephrectomy (1c)

Systematic	Intervention		Results		AMSTAR
Review	Arm 1	Arm 2	Arm 1	Arm 2	Assessment
Operative Time	(minutes)			•	
Young 2008 3 studies	Older Donors (91)	Younger Donors (248)	WMD (95% CI) 11 (-3 to 25)		Moderate
Blood Loss (mi	lliliters)				
Young 2008 2 studies	Older Donors (56)	Younger Donors (90)	WMD (95% CI) 6 (-91 to 103)		Moderate
Length of Hosp	ital Stay (days)		1		I
Young 2008 3 studies	Older Donors (91)	Younger Donors (248)	WMD (95% CI) 0 (-1 to 1)		Moderate

Table C6. Peri/Post-Surgical Outcomes: Older versus Younger Donors (2a)

OR=odds ratio; RR= Risk ratio; WMD=weighted mean difference ^a-Donors with pre-existing isolated medical abnormalities including older age, obesity, hypertension, reduced glomerular filtration rate, proteinuria, microscopic hematuria and nephrolithiasis.

Table C7. Peri/Post-Surgical Outcomes: Obese versus Overweight andNormal Weight

Systematic	Intervention		Results		AMSTAR
Review	Arm 1	Arm 2	Arm 1	Arm 2	Assessment
Peri/post-opera	tive Complications			•	
Lafranca 2012 Systematic Review	Donors with high BMI= ≥30.0 (1442)	Donors with low BMI= <u><</u> 29.9 (4427)	RR (95% CI) 1.01 (0.75-1.36)		Moderate/High
Operative Time	(minutes)				
LaFranca 2012 8 studies	High BMI (380)	Low BMI (725)	WMD (95% CI) 16.91 (9.06 to 24	Moderate/High	
Blood Loss (mil	liliters)				
Lafranca 2012 7 studies	High BMI (284)	Low BMI (655)	WMD (95% CI) 34.46 (-6.73to 75	5.66)	Moderate/High
Length of Hosp	ital Stay (days)				
Lafranca 2012 10 studies	High BMI (1487)	Low BMI (4532)	WMD (95% CI) 0.18 (-0.02 to 0.39)		Moderate/High

RR= Risk ratio; WMD=weighted mean difference

^a-Donors with pre-existing isolated medical abnormalities including older age, obesity, hypertension, reduced glomerular filtration rate, proteinuria, microscopic hematuria and nephrolithiasis.

Appendix D. Long-Term Outcomes: Supporting Tables

Table D1. Long-Term Outcomes: Characteristics of Included Systematic Reviews

Study	Donor Population	Literature Search	Comparison Population	No of participants (studies)	Length of Follow- up, years (range)	Age (mean)	Sex (% women)	Outcomes
Ahmadi 2014	Extended criteria live donors with the following characteristics: 1. older age, 2. overweight and obesity, 3. hypertension, 4. vascular anomalies/multiplicity, 5. women of childbearing age, and 6. minors as donors	Literature search through November 2013	Donors without the listed characteristics or matched non-donors	Older donor age N=90,027 (38 studies) Obesity N=5924 (22 studies) HTN N=81,497 (7 studies) Vascular multiplicity N=14,878 (48 studies)	0-10 0-5 1-20 0-10	NR	NR	All clinically relevant outcomes by age, BMI, hypertension, vascular multiplicity are discussed but not summarized.
Boudville 2006	Included studies involving 10 or more normotensive adults who donated a kidney and in whom blood pressure was assessed at least 1 year later. 48 studies from 28 countries followed 5145 donors.	Literature search Medline and EMBASE 1966-Nov 2005	Healthy non- donor controls in 12 studies	N=5145 donors (48 studies)	Mean 7 (median 6, range = 1- 25 years)	41 (at donation)	58%	SBP, use of antihypertensives, HTN
Clemens 2006	Included any English language study where psychological function was assessed using questionnaires in 10 or more donors after nephrectomy.	Literature search in Medline, EMBASE, Web of Science, Psych INFO, Sociological Abstracts and CIAHL databases from 1969	Non-donor controls (general population, medical outpatients, potential donors, healthy individuals, family members of the recipients	N=5139 donors (51 studies)	Mean 4 years (range 1 week to 37 years from donation)	Mean 42 years	61%	Social function Self-concept Body image Psychological function Quality of life

Study	Donor Population	Literature Search	Comparison Population	No of participants (studies)	Length of Follow- up, years (range)	Age (mean)	Sex (% women)	Outcomes
		through July 2006	in 29 studies					
Garg 2006	Included studies with 10 or more healthy adults donated a kidney, and proteinuria, or glomerular filtration rate (GFR) was assessed at least 1 year later. 48 studies from 27 countries followed 5048 donors.	Literature search Medline and EMBASE 1966-Nov 2005	Healthy non- donor controls in 11 studies	N=5048 donors (48 studies)	Mean 7 (median 6, range 1- 25)	41 (at donation)	NR	GFR, proportion of donors with GFR< 60 ml/min, proteinuria
Young 2008				(30 studies) from 13 countries				

BMI= Body Mass Index; CICr= creatinine clearance; CMS= Center for Medicare and Medicaid Services; CV = Cardiovascular; ESRD = End-stage renal disease; GFR = Glomerular Filtration Rate; HTN = hypertension; IFG = Impaired Fasting Glucose; MetS = Metabolic Syndrome defined according to NCEP ATP III guidelines as the presence of 3 or more criteria: waist ci (1) waist circumference ≥88 cm in women and ≥102 cm in men; (2) hypertriglyceridemia (≥150 mg/dL [_1.69 mmol/L] or treatment); (3) low high-density lipid in– cholesterol (HDL_C) (<50 mg/dL [_1.29 mmol/L] in women and <40 mg/dL [1.04 mmol/L] in men); (4) hyperglycemia (≥100 mg/dL [_5.6 mmol/L]); and (5) hypertension (≥130/85 mm Hg or treatment).; NDI= National Death Index; NR = Not reported; National Health and Nutrition Examination Survey (NHANES III) conducted between 1988 and 1994; OPTN= Organ Procurement and Transplantation Network; SBP = systolic blood pressure; SSDI= Social Security Death Master FileDvD= donor versus donor; DvND=donor versus non-

		-											
Systematic Review	<i>A priori</i> design	Duplicate extraction	Comprehen sive lit search	Publication status = inclusion criteria	Study list provided	Study characteristics provided	Sicentific quality rated	Scientific quality used appropriately	Findings combining appropriate	Publication bias assessed	Conflicts of interest stated	Overall Quality Rating	Statement Statement
Ahmadi 2014	No	Yes	Yes	NA	included - No excluded - No	No	Yes	Can't Answer	Can't Answer	Can't Answer	Yes	Moderate	This is a recent systematic review of observational studies attempting to assess the impact of individually applying a number of extended criteria for living kidney donation (i.e. older age, obesity, hypertension, vascular anomalies and women of childbearing age). The manuscript states that high quality methods were used, but it is very brief on both methods and results so it is difficult to confirm.
Boudville 2006	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	Som e asse ssm ent, but not docu ment ed	Yes	Yes	No	Yes	Moderate	This is a systematic review of observational studies attempting to assess the impact of live kidney donation on hypertension. The evidence base is weak, but reasonable methods were used in the review.

Table D2. Long-Term Outcomes: AMSTAR Assessments of Included Systematic Reviews

Clemens 2006	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	Yes ^a	Can't Answer	Can't Answer	No	No	Moderate/ Low	This is a systematic review of observational studies attempting to assess the impact of live kidney donation on psychosocial health. The quality of the evidence base for this review was low and there is limited information about how the potential bias in these studies was handled.
Garg 2006	No	No	Yes	Yes	included - Yes excluded - No	Yes	No	Can't Answer	Yes	No	No	Moderate	This is a systematic review of observational studies attempting to assess the impact of live kidney donation on donor kidney function. The evidence base is weak, but reasonable methods were used in the review.
Young 2008	No	Yes	Yes	Yes	included - Yes excluded - No	Yes	No	Can't Answer	Yes	No	Yes	Moderate	This is a systematic review of observational studies attempting to assess the impact of living kidney donation among donors with isolated medical abnormalities. The manuscript states that high quality methods were used but details are difficult to confirm regarding how study bias was incorporated into the review.

^a Some assessment, but not documented

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
Berger 2011 United States	LKDs from OPTN database aged ≥70 donating between 1990 and 2010 linked to SSDI for death (n=219)	DvND: NDs are NHANES-III participants without contraindications to donation, matched on age, BMI, SBP, education, ethnicity and smoking history. (n=219)	1, 5, and 10 years	0	D:72.1 ND:NR	45	Mortality (survival at 5 and 10 years)
Chandran 2014 UCSF, United States	Living donors who donated at UCSF from 11994-12/2007 with impaired fasting glucose> 100 mg/dl who were alive and agreed to participate (n=45)	Living donors with normal fasting glucose matched to inpaired glucose donors for age, sex, race and year of donation who agreed to participate (n=45).	10.4 years	31% of donors with impaired fasting glucose participated in the study	47	58%	Estimated GFR, albumin/creatinine, diabetes.
Cherikh 2011 United States	Living donors with ESRD identified through OPTN and CMS ; 1987-2003 (n=126)	All living donors identified through OPTN (n=56,458)	9.8	NR	D with ESRD: 38.4 D controls : 38.8 (at donation)	D with ESRD post donation: 42 All LKD:57.2	ESRD by sex, race
Clemens 2011 Australia, Canada, Scotland	Living donors recruited from 9 transplant centers in Canada, Australia and Scotland; 1970-2007 (n=203)	Healthy non-donors suggested by donor participants, no renal disease, HTN, diabetes, CVD, pulmonary disease, cancer (104)	5.5 median (3.8- 8.4)	D: 52% (203/421) C:39.6% (104/172)	D: 44 C: 40	D: 62 C: 63	Psychosocial (SF- 36), 15 D and feeling thermometer
Cuevas- Ramos 2011	Living donors with Metabolic Syndrome (MetS) (n=28)	DvD: Living donors without MetS (112)	MetS: 4(2.1-5.8) w/o MetS: 12(8.2- 15.7)	61.9% (140/358)	D with MetS:41.2 D without MetS:36.0	D with MetS:46.4 D without MetS:58.9	GFR, proteinuria
Dols 2011 Netherlands	Living donors aged > 60; 1994-2006 (n=117)	Living donors aged <60 (422)	5.5	NR	D <u>></u> 60: 65 D<60: 46	D <u>></u> 60: 59 D<60: 56	Mortality, CV Mortality, HTN, proteinuria, GFR
Doshi 2013 United States	African American donors in Detroit, MI 1993-2006 (n=103)	Matched controls from CARDIA study (235) without contraindications to donation matched by age, gender,	D: 6.8 (2.3) C: 6.4 (2.2)	39.8% (103/171)	D:35(8) C:34(6)	D: 63% C:63%	HTN, GFR, proteinuria

Table D3. Long-Term Outcomes: Study Characteristics of Included Studies

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
		SBP, and duration of follow- up					
El-Agroudy 2007 Egypt	Living related kidney donors who donated 1976-2002 (n=339) Age groups Sex (male vs female)	Egyptian general population	10.7 (4.9) (5-30 years)	75.5% (339/1400)	47.8	61.9%	HTN, GFR, proteinuria, diabetes
Fehrman- Ekholm 2011 Sweden	Living kidney donors; 1965-2005 (n=573)	Present multiple regression with age as an independent variable with SBP and GFR as outcomes.	14(2-43)	48.4% (573/1110)	47.4	59	SBP, GFR, mortality
Garg 2012 (fractures) Canada	Adult Ontario donors; 1992-2009 (n=2015)	Non-donors from administrative healthcare dataset without medical conditions that would preclude donation. Matched on age, sex, rural or urban residence, income at time of nephrectomy and assigned index date. (N=20150)	D: 6.9 (3.8-11.0) C: 6.6 (3.5-10.7)	NR (0.8% of donors with history of fragility fracture before donation.	43	60	Fractures (lower and upper extremities)
Garg 2008 (CV) Canada	Living donors between 1993-2005 in Ontario, Canada. (n=1278)	DvD: age, sex Non-donors from administrative healthcare dataset without medical conditions that would preclude donation. Matched on age, sex, rural or urban residence, income at time of nephrectomy and assigned index date. (N=20280)	Mean 6.2 (SD 3.2) (range 1-13)	37.1% (1278/2033)	41	60	Mortality, CV events, HTN
Garg 2012 Canada	Living donors between 1992-2009 in Ontario, Canada. (n=2028)	DvD: age, sex DvND: overall, age, sex Matched on age, sex, rural or urban residence and income at time of nephrectomy.	Median 6.5 (max 17.7 years)	NR	43 at donation 50 at follow-up start	60	Mortality, CV events
Gibney 2007	Searched UNOS for	All living donors: African	17.6 (time	NA	32 at donation	36	ESRD

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
United States	patients who had donated a kidney and were now on the waiting list for a kidney. (102); 1993-2005 African American (45 (44%), Caucasian 41 (40%), Hispanic 11(12%), Asian 2 (2%), Native American 1(1%) donors on transplant waiting list	American 8889(14%), Caucasian 42,419(68%), Hispanic 7375(12%), Asian 1879(3%), Native American 487(0.8%).	between donation and wait list)				
Gibney 2008 United States	Searched UNOS for patients who had donated a kidney and were now on the waiting list for a kidney. (n=126) ; 1988-2006	DvD: African American (50) donors who need kidney transplant versus White (54) donors.	NR	NA	31	35	ESRD
Gracida 2003 Mexico	Living donors between 1992 and 2001, "normal" donors (n=422)	Donors with HTN (16) (defined as under control with diet and or 1 medication), high cholesterol (62), obesity (BMI >30) (81), age > 60 (6)	6.7	NR	34.5	49	HTN, kidney function (creatinine, GFR)
Gross 2013 United States	Living donors (n=2455) at three major transplant centers in the United States; 1963-2005	National Health Measurement Survey (NHMS) results African American Health Project	17	29.3% (2455/3470 donors who were contacted)	58	61	Psychosocial
Ibrahim 2009 Pregnancy Minnesota, USA	Women who donated a kidney at UofMN 1963- 2007 (n=2102), 1589 responded, 1085 reported 3213 pregnancies. Post-donation pregnancy only 317 in 141 D with post donation pregnancy	Pre-donation pregnancies (n=92519) in 846 D with pre- donation pregnancy only + 204 in 98 D with pre and post donation pregnancies Post-donation pregnancies 317 in 141 D with post- donation pregnancies only +	Women donated 1963-2007, questionnaires sent 2003-2007	2102 women donated, 180 did not respond, 333 were not contacted	39	100	Pregnancy outcomes: Adverse maternal outcomes (HTN, diabetes, preeclampsia), fetal loss, prematurity

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
	only + 173 in 98 D with pre and post donation pregnancies.	173 in 98 D with pre and post donation pregnancies.					
Ibrahim 2009 United States	Living kidney donors from the U of M 1963- 2007 (N=3698) linked to death master file. For a random sample (n=255) from 2003-2007 other outcomes (HTN, GFR, albumin/cr) were collected	NHANES sample matched to donors by age, sex, race, ethnic group, BMI, at the time of the measurement of GFR. US population norms served as control for SF-12 and 36 results. Donors of opposite sex and other BMI categories served as references for comparison	For GFR subgroup 12.2(9.2)	14.3% of the 1785 contacted donors presented for iohexol GFR measurement	52.9	62.1	Mortality and ESRD rate for all donors, GFR, HTN, proteinuria, quality of life for subset
Johnson 1999 United States	Living kidney donors at the University of Minnesota;1984- 1996(N=524)	between groups. General US population scores Scores of patients who have CHF and patients who are depressed	NR	40	41	61	Psychosocial
Karakayali 1998 Turkey	Living kidney donors (n=102)	DvD: Female donors (57) Male donors (45)	Mean 10.2 (range 8mos. – 22 years)	32	41	55.9	GFR, CKD
Lam 2012 Canada	All Ontario LKD 1992- 2009 linked to administrative database for outcomes of acute dialysis (n=2027)	DvND: Ontario database, healthiest non-donors without claims for conditions that preclude donation prior to assigned index date. Matched1;10 on age, sex, rural or urban residence and income at time of nephrectomy. N=20227	Median 6.6 (max 17.7 years) D:6.6 ND: 6.5	7.1% (5.5% left Ontario, 1.5% on non- donors and 0.6% of donors died,)	43	60	Acute dialysis during any hospital stay
Lee 2007 Korea	Living kidney donors who donated 1990-2001 and had GFR data after	Compares LKDs with GFR >60 (normal) (78) to those with GFR <60 (CKD-GFR)	Median 7.4 (range 4.5-14.3) years	86.2 % (104/756 participated)	42.5	42	GFR, HTN, Proteinuria

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
	50 months of follow-up (n=104) Donors Age >50 (n=29) Donors with HTN (n=6) 1 st degree relatives (n=28)	(26) at last follow-up. Donors Age <50 (75) Donors w/o HTN (98) Non 1 st degree relatives (76)					
Lentine 2010 United States	Living donors 1987-2007 from OPTN who had post-donation nephrectomy benefits with a private US health insurer at some point from 2000 to 2007 (n=4650)	Black donors (13.1%) White donors (76.3%) Hispanic donors (8.2%) Also unselected NHANES 2005-2006 participants stratified by race and ethnicity	Median time from donation to end data file for individual 7.7 years	NR Study sample N=4650, all donors in OPTN=86107	37.2	54.6	ESRD, HTN, Diabetes, CKD diagnosed from insurance claims
Lentine 2012 United States	Living donors 1987-2007 from OPTN who had post-donation nephrectomy benefits with a private US health insurer at some point from 2000 to 2007 (n=4650)	General insurance beneficiaries matched by gender and age, follow-up limited to the shortest	Median time from donation to start of insurance 4.9 years and to the end of insurance 7.7 years	Study sample N=4650	37.2	54.6	Depression
Lentine 2014 United States	Living donors 1987-2008 from OPTN who had post-donation nephrectomy Medicare billing claims 2000-2008 (n=4,007)	Living donors 1987-2007 from OPTN who had post- donation nephrectomy benefits with a private US health insurer at some point from 2000 to 2007 (4650) from prior study	Median time from donation to end of insurance 6.0 years	NR	54.8	60	HTN, Diabetes, CKD, proteinuria diagnosed from insurance claims
Mac Donald 2014 UMN, USA	Living donors at the Uof Minnesota 1963-2012 who were < 18 years of age at donation (n=39)	Living donors at the Uof Minnesota 1963-2012, 18-30 years of age at donation, matched to to adolescent donors on gender, relation to the recipient, BMI at donation, MDRD eGFR, year of donation (128)	Mean Adolescent D follow-up: 31.8+8.0 years Mean Adult D follow-up: 29.2+10.3 years	NR, 39/42 adolescent donors included in the analyses	Adolescent D: 17.1+0.7 Adult D: 24.2+3.6	Adolescent D: 43.6 Adult D: 51.6	Mortality, eGFR, proteinuria, HTN, Diabetes

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
Mjoen 2011 Norway	Living kidney donors 1963-2007 in Norway who responded to questionnaires (n=1508), responses from 1414 used in analyses	Unselected non- institutionalized population 16-80 years from Akershus county in Norway (6800)	Median 12.6 years	24% (1508/2269 responded)	46	60.5	Quality of Life
Mjøen 2013 Norway	Living kidney donors who donated 1963-2007 in Norway after exclusion of donors with BP>140/70, BMI>30, age >70 or <20, microalbuminuria, or eGFR<70 ml/min (n= 1,901).	Matched to healthy participants from the HUNT population study, only subjects with BP<140/90, BMI<30, those without diabetes, CVD, or HTN (use of BP meds) were included (32,621)	D: median 15.1 (range 1.5-43.9) years C: median 24.9 (range 0.1-26.0) years	0 (no loss to follow-up)	D: 46.0 C: 37.6	D: 59 C: 53.1	Mortality CV Mortality ESRD
Muzaale 2014	OPTN Donors (n=96,217)who donated 4/1/1994-11/30/2011 linked to CMS to ascertain ESRD status (maintenance dialysis, placement on waiting list or receipt of transplant)	NHANES III participants after excluding those with contraindications to kidney donation matched on age, sex, race, education, BMI, smoking, SBP (20,024) linked to CMS to ascertain ESRD status.	D:7.6 (3.9-11.5) C:15 (13.7-15)	NA	40.2	59	ESRD
Okamoto, 2010 Cross- sectional Japan	Glucose intolerant donors who donated 1985-2008, n= 71 (diabetic n=21, impaired glucose tolerance m=44)	Non-glucose intolerant donors (373)	DM D: 7.3 <u>+</u> 5.9 GIT D: 8.4 <u>+</u> 6.8 No GIT D: 10.7 <u>+</u> 6.8	7.9%	54	63	Mortality
Reese 2014 Retrospective United States	OPTN /UNOS donors ≥ 55 years at donation, who donated 1996-2006 and had a matched non- donor (3368) used for death outcome analysis, donors with Medicare used for CVD outcome	Participants in the Health and Retirement Study (NIH, nationally representative sample of adults > 50 years of age in the US) without HTN, diabetes, CVD, pulmonary disease, psychological or neurological	Median 7.8(IQR 5.1-10.2)	3368/5152 (65.4%) donors were matched 3368/7319 (46.0%) eligible non- donors were	51	59	Death, death or CVD event (ischemic cardiac disease, congestive heart failure, stroke, PVD), diabetes

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
	analysis (1312)	condition, BMI <40 who rated their health as good, very good or excellent who were matched to donors by index date, race, sex, neighborhood poverty level, BMI (3368).		matched for death outcomes			
Reisaeter 2009 Cross- Sectional Norway	Linked the Norwegian Renal Registry with the Medical Birth Registry of Norway to identify 326 donors with 726 pregnancies, 106 post- donation pregnancies;1967-2002	Medical Birth Registry of Norway Pre-donation pregnancies N=620 Random sample of birth registry 1% (N=21511)	NR	0	27	100	Chronic hypertension, gestational hypertension, preeclampsia, Mortality (stillbirths)
Segev 2010 United States	OPTN Donors who donated 1994-2009 linked to Social Security Death Master File (n=80,347)	NHANES III participants after excluding those with contraindications to kidney donation matched on age, sex, race, educational background, history of cigarette smoking, pre- operative body mass index (BMI), and preoperative systolic blood pressure (SBP) (n=9364)	6.3	0.001%	NR	58.5	Mortality
Storsley 2010 Canada	Aboriginal donors donating 1970- 2007(n=38)	Randomly selected white donor controls (n=76).	AD:14.6 <u>+</u> 9.3 WD:13.4 <u>+</u> 9.5	9%	AD: 32.0 WD: 40.0	Ad:61 WD:52	Death, ESRD, HTN, GFR
Thomas 2013 Canada	All Ontario LKD 1992- 2009 linked to administrative database for outcomes of kidney stones (n=2019)	DvND: Ontario database, healthiest non-donors without claims for conditions that preclude donation prior to assigned index date and no h/o kidney stones. Matched1;10 on age, sex, rural or urban residence and	Median 8.4 (max 19.7years) D: 8.8 ND: 8.4	<7%	43	60	Kidney stones (claims for surgical interventions)

Study Design Country	Donor Population Data Source (n=number analyzed)	Comparison(s) Comparison Population Data Sources (n=number analyzed)	Length of Follow-up mean or median years (range)	Attrition-% who did not participate), (n participants/ N total)	Age (mean)	Sex (% women)	Outcomes
Thomas 2014 Canada	All Ontario LKD 1992- 2009 linked to administrative database for outcomes of GI bleeding (n=2009)	income (n=20190) DvND: Ontario database, healthiest non-donors without claims for conditions that preclude donation or GI bleeding episodes prior to assigned index date. Matched1;10 on age, sex, rural or urban residence and income at time of nephrectomy. (n=20090)	8.4, max 19.7 D: 8.8 ND:8.4	8.5% 6.5% moved from Ontario, 2% died.	42	60	GI bleed
Tsai 2013 Taiwan Von Zur-	105 LKDs; 1983-2011 in Taiwan	Donors with lower eGFR at donation (n=NR) Female donors (n=60) Male donors (n=45)	5.4 <u>+</u> 4.9	NR	46.3	60	ESRD, CKD, proteinuria
Von Zur- Muhlen 2014 Sweden	455 LKDs; 1974-2008 in Sweden, 395 LKDs who agreed to participate in the study	Female vs male donors Pre-donation GFR, age, BMI	11 <u>+</u> 7	18.5%	49	58	eGFR, HTN, proteinuria
Wafa, 2011 Egypt	Consecutive live donors between 1976 and 2008 (n=2000)	8 donors who developed ESRD	NR	NR	30.9	25	ESRD

BMI= Body Mass Index; CICr= creatinine clearance; CMS= Center for Medicare and Medicaid Services; CV = Cardiovascular; D= Donors; ESRD = Endstage renal disease; GFR = Glomerular Filtration Rate; GI= Gastrointestinal; HTN = hypertension; IFG = Impaired Fasting Glucose; MetS = Metabolic Syndrome; ND= Non-Donors; NDI= National Death Index; NR = Not reported; National Health and Nutrition Examination Survey (NHANES III) conducted between 1988 and 1994; OPTN= Organ Procurement and Transplantation Network; SBP = systolic blood pressure; SSDI= Social Security Death Master File; DvD= donor versus donor; DvND=donor versus non-donor

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Berger 2011 Retrospectively ided prospective registry cohort with matched non-donor controls	mortality	Moderate (all older donors were included, controls were matched based on NHANES data)	Low	Low	Differential follow-up start - at donation for D vs at data collection for non-donors. Also, despite matching, donors likely are more carefully selected compared to non-donors.	Moderate risk of bias. This study has better design than many, matched healthy controls. Despite the effort, control population likely differs from donor pool.
Berger 2011, retrospective, USA	Mortality	Comparison group appropriately selected and matched to study group; significant difference in baseline characteristics on some parameters (no of females, BMI>30, and prevalence of HTN). (high)	Statistical analysis appropriate though study not powered for subgroup interaction effect. (high)	Very low percentage (<5%) of attrition. (low)		Moderate risk of bias due to differences in baseline characteristics , and lack of power for subgroup interaction effect.
Chhandran 2014 Retrospective matched cohort	eGFR, albuminuria, diabetes, HTN	Moderate (donors who participated differed from those who didn't)	Low	High	Mached donors differed from impaired glucose donors (lower BMI, lower BP predonation).	High to moderate risk of bias
Cherikh 2011 (AJT). Retrospective	ESRD	moderate to high. Comparison to	Relatively short follow up of avg 9.8 yrs.	low	Avg duration of follow up 9.8 yrs. May be short to completely assess ESRD risks.	moderate to high risk of bias. No

Table D4. Long-Term Outcomes: Risk of Bias Assessments of Included Studies

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
cohort of donors between 1987 and 2003. Affect of age and ethnicity.		USRDS 2009 annual data report				appropriate control group.
Cherikh 2011 Restrospective cohort study of LKDs and risk for ESRD	ESRD	Low (Single arm study including only LKDs as defined by known registry data)	Low (Robust measurement s for ESRD from multiple data sources, manual confirmation of data in some instances)	Low (No documentation of the extent of missing data, however likely low as patients cannot be "lost to follow-up" in these government datasets)	Νο	Low risk of bias. This retrospective study utilized multiple data sources with robust methods and did not attempt a comparison to a healthy non- donor population due to issues with selection bias.
Clemens 2011, Canada, Scotland, Australia, retrospective cohort	Psychosocial outcomes based on SF 36, 15D, feeling thermometer	High (17% of all donors participated), controls suggested by donors but were not able to donate	High, donors who responded might have been better of than the rest	High	There was correlation between D and ND controls were suggested by donors and are likely their family members.	High risk of bias, high attrition, outcomes are subjective. High -
Clemens, 2011 Retrospective cohort in Canada, Australia and Scotland	SF-36 and Feeling Thermometer	High - 44% of eligible donors participated, Controls were suggested by the donors themselves.	Moderate	Unclear - attrition is tough to assess in a one-time survey	Retrospective nature = potential recall bias	responders aren't likely entirely reflective of the entire sample, not all validated instruments used.

Author Year			Detection			Overall Risk
Study Design Cuevas-Ramos 2011 Retrospective Cohort Mexico	Outcomes eGFR, Albuminuria	Selection Bias High - interim report only, but >20% of full sample included and no way of knowing how they compare to the full cohort.	Bias Moderate - 5 vear f/u	Attrition Bias	Other Sources of Bias	of Bias Moderate to high - selection bias, high attrition
Cuevas-Ramos 2011 Retrospective	Low eGFR	Moderate to high - age and gender possible confounding factors, enrollment bias	Unclear/high - not clear if statisticians were blinded to data	Low		Moderate risk of selection bias between the two groups due to confounding factors, enrollment bias
Dols 2011 Retrospective cohort study of the impact of age and LKD	eGFR, proteinuria	Unclear (No documentation of baseline co- morbidities between each group, could older patients have more chronic illness, less strict selection criteria?)	Low	Unclear (16% lost to follow-up at 1 year in both groups, no formal disclosure of lost to follow-up for later years)	Surgical technique differences by time period, more younger patients were included from earlier years before laparoscopic donation was standard of care	Moderate risk of bias. This retrospective study was a single center cohort and did not disclose if there were differential co- morbidities between groups that could explain their results aside from age alone.
Dols 2011 (AJT). Prospective cohort of consecutive donors compared by age.Compared donors older or	Estimated GFR	High. Not comparing elderly donors to age matched controls.	Moderate- high. Short term follow up (median 5.5yrs)	low. Data on 539 consecutive donors.	Renal function based on estimated GFR. May be less accurate in elderly compared to younger population (MDRD used).	High risk of bias. No age matched comparison group.

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
younger then 60.						
Doshi, 2013 Retrospective cohort	eGFR, albimin, creatinine	High - matching not entirely successful, eGFR and insurance status not well matched	Moderate - short follow-up at 6.8 years	High - only 69% of eligible donor participated		Moderate - high attrition, matching not entirely successful
Doshi, 2013 United States	Renal function, HTN, diabetes, proteinuria	Low (best possible selection from a prospective cohort, comorbidities excluded and non-donors matched by age, gender, BP, time of follow-up.	Low	Moderate	Low	Low to Moderate Risk of Bias.
El-Agroudy 2007 Egypt Retrospecitve study	HTN by sex groups; serum creatinine, proteinuria, HTN, diabetes by age group; diabetes, CV events for the general cohort.	High, only 24.2% of the cohort had prospective follow-up.	Unclear	High	High, general population comparison, only age and sex matched.	High risk of bias (high attrition, possible selective outcome reporting, poorly matched control population).
El-Agroudy, 2007 Egypt Retrospective cohort	HTN, serum creatinine and proteinuria, by age	High - 25% of potential participants		High (75%)		High

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Fehrman-Ekholm 2011 (NDT). Cross-sectional retrospective sutdy in sweden.	GFR, measured and estimated	High. Donor function compared to previous studies of donors.	low	No data on deceased donors. 10% of donors with no data	Some donors deceased and 10% lost not included. Participation of 70% potential donors.	High risk of bias. No reported data of comparison group.
Fehrman-Ekholm 2011 Cross sectional study of LKDs and renal function over time	eGFR	High (Cross sectional study, survival bias given they excluded deceased patients from analysis, non- participation rate is moderately high)	Low	Low	No	Moderate to high risk of bias. This is a cross sectional study with a potential for a high degree of selection bias and immortal time bias which may tremendnously impact their results.
Garg 2008 (Transplantation) . Retrospective cohort sutdy of living donors in Ontario Canada. Donors between 1993 and 2005.	Composite of time to death or first major cardiovascular event.	moderate to high. Non- donors from general population.	Donors more frequently followed then general population	low	Short follow up of 6.2 years (range of 1-13).	moderate risk of bias. General population control group.
Garg 2008 Retrospective cohort study of LKDs and risk of cardiac disease	Time to death or major CV event	Unclear (Excellent study design with 5:1 matched controls on age, sex, income, and healthcare utilization but no information on control co- morbidities)	Low	Low	No	Low to moderate risk of bias. This is an excellently designed registry study using multiple data sources and decent controls, although they lacked co- morbidity

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
						information on
						controls. The comparison
						group could
						include
						patients that
						could have
						conditions that would
						excluded them
						from being a
						donor.
						Low risk of
						bias. This study added
						on the
		Low (Excellent				previous study
		study similar to				by making the
		previous 2008				controls more
		study, but higher sample				robust and comparable to
Garg 2012		size and more				the health of
Retrospective		variables used				someone who
cohort study of	Time to death	in matching				would be a
LKDs and risk of	or major CV	controls to				candidate for
cardiac disease Garg 2012	event	donors)	Low	Low	No	donation.
(BMJ).						
Retrospective			Follow up			
population based		moderate:	mean 6.5 yrs.			
matched cohort		matched	May be to			
study of donors in Ontario	Composite of time to death or	donors from "the healthiest	short for detection of			Moderate risk of bias.
Canada, Donors	first major	segment of the	composite			General
between 1992	cardiovascular	general	primary end			population
and 2009.	event.	population".	point	low	Poor reliability mentioned in cause of death	control group.
Garg 2012		Moderate (all	Authors state			Moderate risk
Retrospectively ided prospective	Lower and	donors in Ontario were	that database codes for			of bias due to claim - based
administrative	upper extremity	included and	fractures are			definition of
database donor	fragility	controls were	sensitive. Both		Non-donor characteristics are defined by	non-donor
cohort with	fractures	matched based	D and ND had	Moderate.	claims (some have low sensitivity)	characteristics

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
matched non- doner controls.		on claims)	similar criteria for fracture diagnosis. Non-blinded. High.			(unknown other predictors of fractures, such as smoking, fitness, falling tendency etc).
Garg 2012, retrospective, Canada.	Fractures	Comparison group appropriately selected and matched to study group; no significant difference in baseline characteristics. (low)	Statistical analysis appropriate; sample size significantly large detect effect. Subgroups specified a priori (low)	Study based on database containing all the variables needed by the researchers so no missing data issues. (low)	Yes. Outcomes mainly assessed from claims data which may not reflect true clinical outcomes. Also a single center study	Moderate bias, mainly due to study based on claims data which may or may not reflect true clinical outcome.
Gibney, 2007 United States	GFR, ESRD by race	Uncertain – no between group comparisons by characteristics.	Moderate	Difficult to say in a chart review design	Outcomes are assessed entire through registry data.	Moderate to high
Gibney 2007 United States Retrospective	Low eGFR/ESRD (Race of donors on wait list for transplantation advanced CKD/ESRD in donors)	High/unclear - not entirely clear whether AA donor group and Caucasian donor group were well matched in terms of confounding factors	High	Unclear		Moderate to high risk of bias - probable significant confounding factors between the two outcomes groups
Gibney 2008 (Trans Pro). Retrospective study of donors with ESRD compared to all donors between 1988 and 2006	Characteristics of donors with ESRD	moderate. Both groups as donors but no baseline characteristic comparisons beyond age and race.	High. No information reported on donors with ESRD who were not listed or deceased with ESRD.	moderate.	Comparison groups of different time frames.	high risk of bias. May be differences in baseline characteristics between donors with and without ESRD.

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Gibney 2008 Descriptive study of LKDs and age/sex/race and risk of ESRD	Frequencies of age/sex/race for LKDs on waitlist versus all LKDs	Low	Low	Unclear (see next comment)	The inclusion of LKDs from differential time periods (waitlist only assessed from 1996- 2006) and not having any other data to control for possible confounding makes interpretation of the results of this study very challenging.	Moderate risk of bias. The study design is descriptive in nature and the groups were chosen in very arbitrary way. A correlation was observed with no control for confounding factors.
Gracida, 2003		Low	Low	Unclear	F/u time 6.7 years, SDs not reported. Authors mention	High
Gross 2013 (AJT). Observational cross sectional survey of living donors between 1963-2005 at 3 centers	Quality of life based on SF-36 survey	high. Control group unselected from US population. Not medically matched.	moderate. Poor response rate	moderate. Only 50% of eligible donors contacted, of these 2/3rd's returned questionnaire	Non-participants in survey differed from participants.	Moderate risk of bias due to missing eligible donors, comparison to general population in US which are not medically matched.
Gross 2013 Cross-sectional study examining health related QOL in LKDs	Health-related QOL	High (Cross sectional design with only 27% of all donors enrolled, high initial exclusion rate with potential of survival bias)	Low	Low (survey participation rate was high at 97% of those who consented)	No	High (results could be entirely explained by selection bias given the sheer number of donors who were initially excluded).

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Ibrahim 2009 (NEJM). Single center retrospective cohort, donors between 1963 and 2007.	Survival and ESRD	Moderate to high, comparison to general population. Survival compared to life tables.	low	low	no	moderate to high risk of bias. No appropriate control group.
Ibrahim 2009 Long term consequences of kidney donation	Death, ESRD, eGFR, hypertension, QOL	Low (random selection of contacted donors within a stratified scheme lowers impact of selection bias, NHANES controls)	Low	Low	Νο	Low (well- designed study with random subpopulation from thousands of known donors, robust NHANEs controls)
Ibrahim 2009 Retrospective cohort of women who donated at the U of MN	Pregnancy outcomes such as maternal complications, fetal loss, prematurity	High, 75% responded to questionnaires	High, based on recall	High	Pre-donation pregnancies were more remote than post-donation (16. yrs vs 2.5 yrs from survey)	High risk of bias, though response rate is high, there were non- responders. Also, outcomes were based on recall many years later.
Ibrahim 2009, retrospective, USA.	Pregnancy outcomes.	Unclear if comparison group appropriate; no significant difference in baseline characteristics. (unclear)	Statistical analysis appropriate though many confounders not adjusted for, and ack of an internal control group.(high)	Missing data handled by imputation; responders differed from non- responders on some parameters.(unclear)	Yes. Possibility of both response and recall bias. (high)	High, due to inappropriate selection of comparison group, and high possibility of response and recall bias).
Johnson 1999 Cross sectional	Quality of life/psychosocia	Unclear - comparison	High	High - survey - cross sectional -		High risk of bias - survey

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
	l outcomes	group (?)		~60% response rate		with only 60%
		Low - Within				response rate
Karakayali, 1998	Hypertension by age	donor comparisons	Unclear	Unclear		Moderate
Lam 2012 Retrospectively ided prospective administrative database donor cohort with matched non- doner controls.	Acute dialysis (procedure codes)	Moderate(all donors in Ontario were included and controls were matched based on claims)	High. Database codes for AKI are not very sensitive, though dialysis is easier to identify	Moderate, higher proportion of non- donors died.	Non-donor characteristics are defined by claims (some have low sensitivity)	Moderate risk of bias. Acute dialysis is a rare event, might not have the power to see the difference. Also bias due to claim-based definition of non-donor characteristics
Lam 2012, retrospective, Canada.	Acute dialysis	Comparison group appropriately selected and matched to study group; no significant difference in baseline characteristics. (low)	Statistical analysis appropriate; sample size significantly large detect effect. However, study not powered for subgroup interaction effect though this could be negated by the large sample size(low)	Study based on database containing all the variables needed by the researchers so no missing data issues. (low)	Yes. Outcomes mainly assessed from claims data which may not reflect true clinical outcomes. Also a single center study	Moderate bias. Good study design and statistical analysis, but significant bias might result if claims data do not match significantly with clinical outcome. Also, study not powered for subgroup interaction effect.
Lee 2007 Retrospective	Low eGFR (eGFR <60 mL/min)	Moderate - low eGFR group with significantly greater age	High	Unclear		Moderate to high - significant age difference between the

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
						two groups being compared
Lentine, 2010 Retrospective	Cardiovascular disease, Hypertension, diabetes, CKD	Moderate – administrative database comparison group	Moderate			Moderate
Lentine, 2012 Retrospective	Depression	Moderate – administrative database comparison group	Moderate			Moderate
Lentine, 2014 Retrospective	HTN, CKD, Diabetes	Moderate, donots with insurance were selected	Moderate, CKD claims have low sensitivity	Low		Moderate
Mac Donald, 2014	Mortality, eGFR, proteinuria, HTN, diabetes	Moderate, younger donors were compared to older donors, longer follow-up of younger donors	Moderate, not all donors had creatinine measurement s and proteinuria measurement s thought the follow-up	High for lab outcomes: 23/39 and 88/128 donors had creatinine and HTN data available in 109 donors (subgroup distribution not given)	None	Moderate
Mjoen 2011 Cross-sectional study of LKDs and QOL	QOL	Low (high participate rate overall, limited survival bias)	Unclear (control group was a US population norm compared to a Norwegian donor population)	Low	No	Low (high participation rate given it's a cross- sectional survey design, possible bias from using a US control population)
Mjoen 2011 (AJT). Cross sectional study of donors between 1963 and 2007	Quality of life based on SF-36 survey	high. Control group unselected from Norwegian population. Not	moderate: 76 % response rate from donors in collection	low to moderate	No pre-donation comparison to see change with donation in SF-36	moderate- high. Comparison group not matched.

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
compared to general population in Norway.		medically matched.	period			Donors may be more healthy then comparison general population group
Mjoen 2013 (KI). Retrospective study of events 1963-2007 in Norwegian donors compared to potentially eligible kidney donor control group	ESRD, cardiovascular and all-cause mortality	moderate. Age of controls younger (46 vs 37.6 yrs). Also slightly lower BMI and systolic BP.	low	low	All donors and controls from single country. No data on renal function.	low to moderate: Study has in general longer follow up times and large number of events then others
Mjoen 2013 Retrospective cohort study of LKDs and long term outcomes	death, ESRD	High (Inclusion of donors prior to 1985, control population was not matched with donors to any variable, ie age/sex)	Low	Unclear (use of imputation for missing data)	Νο	Moderate (Lack of matching by key variables between donors and controls is a key limitation of this study; attempts to control this with regression, but matching would have been a better study design)
Muzaale 2014 (JAMA). Retrospective cohort matched to NHANES III	Cumulative	Moderate: screened non- donor population derives from	moderate: Short follow		Compared groups not of concurrent years.	Moderate risk of bias: moderate. Control population
healthy non- donors	incidence and risk of ESRD	NHANES III. May not all	up, mean of 7.6 years	low	Donors were 1994-2011 and matched controls between 1988-1994.	may differ from donor

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
		have been approved donors.				pool.
						Low (Well- designed study that took into account differential follow-up when comparing
Muzaale 2014 Retrospective registry study of LKDs and risk of	ESRD	Low (Matched control population from NHANES, donors are from				donors to non- donors, matched NHANES population, large N with enormous
ESRD Okamoto 2010 Cross sectional	Mortality and ESRD	registry data) Moderate - age difference between to groups	Low	Low High - survey - cross sectional	No	power). Moderate to high risk of bias - age difference between the two groups, responder bias
Okamoto, 2010 Japan Cross-sectional survey	Survival, ESRD, HTN	GI group is older, non-GI group is more female	High	Moderate	Not detected	High risk of bias
Reese 2014, USA	Survival, Death or CVD, diabetes	Moderate – 65% of eligible donors and 46% of eligible non-donors were matched and used in	Low for death High for CVD as they used Medicare claims	Low – all donors included in the analysis were followed for outcomes	Not detected	Low to moderate risk of bias for death; Moderate to high risk of bias for CVD

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
		analysis of mortality. Matched donors were younger, less likely to be AA, more likely to come from neighborhoods with less poverty				outcomes.
	Pregnancy outcomes such as maternal complications,				Women who have deliveries after donation	High risk of bias. Comparison between pregnancies pre-and post- donation is biased by difference in mother characteristics , with older age at post- pregnancy being a predictor of poor pregnancy outcomes. Comparison of donor outcomes to non-donor is biased by selection: donors are healthier than random sample of
Reisaeter 2009,Norway	fetal loss, prematurity	Low	Low	Low	are older at delivery, increasing probability of complications.	women and should have

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
						better outcomes.
Reisaeter 2009, retrospective, Norway.	Pregnancy outcomes.	Appropriate selection of comparison group, though description of baseline characteristics not extensive.(High) Administrative	Statistical analysis test used (Fisher's) did not allow for adjusting of confounding, though a more sophisticated test GLMM (regression analysis) employed but ony used for two groups in this regard. (high)	No mention of missing data or attrition. (unclear)	Yes. Possible selective reporting of outcomes. (high)	High due to baseline characteristics of mothers not described thoroughly enough to explore possible differences that might affect pregnancy outcomes.
Segev, 2010	Death, ERSD	comparison group	Low	Low		Low- Moderate
Storsley, 2010	Mortality, renal function, HTN, diabetes, proteinuria			High		High

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Storsley 2010, Canada, Manitoba	Death, ESRD, renal function, HTN, diabetes, proteinuria among Aboriginal donors and white donors (controls)	High (data available 31/38 AD and 64/76 WD)	Low	High	High, white donors were chosen other than matched Aboriginal non-donors, gen population outcomes are also different between A and W individuals. Unclear if donation modifies this risk.	High risk of bias
Thomas 2013 Canada Retrospectively	Stones	High, donors screened for donation. Controls were screened likely due to symptoms	Low to Moderate	Moderate	Moderate, non-donor's characteristics are defined by claims.	Moderate Risk of Bias
Thomas 2013 Canada Retrospectively ided prospective administrative database donor cohort with matched non- donor controls.	Kidney stones requiring surgical procedures and hospital encounters for kidney stones	High, donors had greater screening for history of prior kidney stones (in clinic) vs controls were screened based on claims over a certain time frame	Moderate, Claims for kidney stones requiring procedures or hospital visit not validated.	Moderate	None detected.	Moderate
Thomas 2014 Retrospectively ided prospective administrative database donor cohort with matched non- donor controls.	GI bleeding risk (ICD-9 codes)	Moderate(all donors in Ontario were included and controls were matched based on claims)	High, GI B identified based on ICD- 9 codes, though they have PPV of 86%	Moderate	Non-donor characteristics are defined by claims (some have low sensitivity)	Moderate risk of bias due to claim - based definition of non-donor characteristics as well as claim based definition of the outcomes, though should not be differential between donors and

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
						non-donors.
Thomas 2014, retrospective, Canada	GI bleeding	Comparison group appropriately selected and matched to study group; no significant difference in baseline characteristics. (low)	Statistical analysis appropriate; sample size significantly large detect effect (low)	Study based on database containing all the variables needed by the researchers so no missing data issues. (low)	Yes. Outcomes mainly assessed from claims data which may not reflect true clinical outcomes. Also a single center study	Low to moderate bias. Good study design and statistical analysis, but significant bias might result if claims data do not match significantly with clinical outcome.
Callaua	Gibleeding	(10W)	Low -			outcome.
Tsai, 2013 Taiwan Retrospective	GFR, ESRD	Low – includes their entire cohort of donors	Statistical methods appear appropriate	Low – attrition not an issue with a retrospective chart review.	Small sample size, short f/u time	Low to moderate
Tsai 2013						Moderate risk of bias - probable confounding factors between the two outcomes groups which were not
Retrospective	Low eGFR	Moderate	Low	Unclear		measured
Von Zur-Muhlen 2014,						
Retrospective,						
Sweden	eGFR, BP, proteinuria	Moderate	Low	Moderate	No control group, donors compared to donors	Moderate

Author Year Study Design	Outcomes	Selection Bias	Detection Bias	Attrition Bias	Other Sources of Bias	Overall Risk of Bias
Wafa 2011 (exp						
and clin Tx). Retrospective						
cohort of donors						
at single center						
in Egypt. NO						
COMPARISON						
GROUP. Did						
make comment on no added						
risks over			low. Easily			
general			define end	low. Data on 2000		
population.	ESRD	Moderate	point of ESRD	consecutive donors		Moderate
						High (Serious
						concern about
						how they
				High (Measurement		measured ESRD in their
				on ESRD rate		population
				depends on		while the
				complete follow-up		authors claim
				data for ESRD		they have
				development on all		follow-up data
				donors, unclear if		on all their
				they actually have that as they don't		patients, how do they know
				disclose their lost-		they captured
				to-follow up rate.		all the ESRD
	Descriptive			Authors claim they		in the donor
Wafa 2011	study (rate of			saw all of their		population?
Cross-sectional	ESRD may not			donors, but do they		No linkage to
descriptive study of LKDs who	be accurate as no data on lost-	Low (not	Low (not	see 2000 post- donor patients		ESRD
developed ESRD	to-follow up)	Low (not applicable)	applicable)	annually?)	No	registries is a huge problem)

Table D5. Long-term Outcomes of Living Kidney Donation: Living Kidney Donors compared to healthynon-donors

Study, Year,	Mean	Inte	rvention/Control	(n)		Results		ROB
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
Mortality								
Mjoen, 2013 Norway	D: 15.1 N-D: 24.9	Donors (1,901)	Non-Donors (32,621)		n/N (%) 224/1901 (11.7) HR ^a (950 1.30 (1.11 p=0.0	to 1.52)		Low-Moderate
Garg, 2012 Canada	Median 6.5	Donors (2,028)	Non-Donors (20,800)		n/N (%) 16/2,028 (0.8) RR:0. 95% CI: 0.2	n/N (%) 365/20,280 (1.8) .44		Low
Segev, 2010 United States	Up to 12 years	Donors (at 12 years: 10,436)	Non-Donors (at 12 years: 127)		Kaplan-Meir curves sug in matched controls at No values provided, diff 12 yea Log-rank	ggest mortality higher 5-12 years follow-up. erence may be 1% at ars.		Low-Moderate
Cardiovascular	Events							
Mjoen, 2013 Norway	D: 15.1 N-D: 24.9	Donors (1,901)	Non-Donors (32,621)		Cardiovascular Deaths n/N (%) 68/1901 (3.6)	Cardiovascular Deaths n/N (%) 688/32621 (2.1)		Low - Moderate
					HR*(95 ⁰ 1.40 (1.03 p=0.0	% CI) to 1.91)		
Garg, 2012 Canada	Median 6.5	Donors (2,028)	Non-Donors (20,280)		Major cardiovascular events n/N (%) 26/2,028 (1.3)	Major cardiovascular events n/N (%) 287/20,280 (1.4)		Low
					RR (959 0.91 (0.61	% CI) to 1 35)		
ESRD	1	L	1 1		0.01 (0.01	10 1.00		
Muzaale, 2014 United States	At 15 years post- donation	Donors (8,781)	Non-Donors (50,124)		30.8 per 10,000 (CI: 24.3-38.5) p<.001	3.9 per 10,000 (CI: 0.8-8.9)		Low - Moderate
Mjoen, 2013	D: 15.1	Donors	Non-Donors		n/N (%)	n/N (%)		Low-Moderate

Study, Year,	Mean		vention/Control			Results		ROB
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
Norway	N-D: 24.9	(1,901)	(32,621)		9/1,901 (0.5) HR ^a (9) 11.38 (4.3 p<0	7 to 29.63)		
Renal Function								
Garg, 2006 SR	(189) WMD (95% Cl) -10 (-15 to -6)					Low		
Acute Kidney In								
Lam, 2012 Canada	D: median 6.9 ND: Median 6.5	Donors (2,027)	Non-Donors (20,270)		Acute Dialysis 1/2,027 (0.05%) 6.5/100,000 person- years RR (95 0.58(0.0			Moderate
Proteinuria						/		
Garg, 2006 SR	7	Donors (129) Donors (67)	Non-Donors (59) Non-Donors (51)		WMD (66 (2 <u>Microlabuminuria_n/N</u> (%)	n/N (%)		Low
						2/52(3.9) 5% CI) 2 to 12.6)	-	
Hypertension			.					
Boudville, 2006 SR Psychosocial	Min 5	Donors (157)	Non-Donors (128)		Systolic Bloo WMD(9 6 (1.6	95% CI)		Moderate
Clemens,2011 Canada, Scotland, Australia	Median 5.5	Donors (203)	Non-Donors (104)			<u>15D QOL score</u> Mean (SD) 0.94 (0.06)		High

Study, Year,	Mean	Int	tervention/Control ((n)		ROB		
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
					p= 0.46	85 (10)		
Other: Fragility f	ractures							
Garg, 2012 Canada	Median 6.5	Donors (2,015)	Non-Donors (20,150)		Rate/10,000 person- years (95% CI) 16.4 (11.1 to 24.2)	Rate/10,000 person-years (95% CI) 18.7 (16.5 to 21.1)		Moderate
					Model-based Rate 0.88(0.58			
Other : GI Bleed								
Thomas, 2014 Canada	Median 8.4	Donors (2,009)	Non-Donors (20,090)		Rate/10,000 person- years (95% CI) 18.5/10,000 person- years	Rate/10,000 person-years (95% CI) 14.9/10,000 person-years		Moderate
					Model-based Rate 1.24(0.85	e Ratio (95% CI) 5-1.81)		
					Time to first event (ho blee HR (95 ⁰ 1.25(0.87	<u>d)</u> % CI)		
Other: Kidney St	lones				1.23(0.07	-1.79)		
Thomas, 2013 Canada	Median:8.4	Donors (2,019)	Non-Donors with no evidence of kidney stones (20,190)		<u>Kidney Stones with</u> <u>surgical intervention</u> 8.3/10,000 person- years	<u>Kidney Stones with</u> <u>surgical</u> <u>intervention</u> 9.7/10,000 person- vears		Moderate
	Kidney Stones with surgical intervention RR (95% CI) 0.85 (0.47 to 1.53)		% CI)					
					Kidney stones with hospital encounters 12.1/10,000 person-	Kidney stones with hospital encounters 16.1/10.000		
					years	person-years		
					Kidney stones with h RR (95' 0.75 (0.45	% CI)		

CI= Confidence Interval; HR= Hazard Ratio; NS= Not statistically significant; QOL= Quality of Life; RR=Risk Ratio; SD= Standard Deviation; SR= Systematic Review; WMD= Weighted Mean Difference ^aHazard ratio adjusted for age, gender, year of inclusion, systolic BP, smoking, and BMI after multiple imputation.

^b-Studies in this SR described method of GFR estimation as timed urine creatinine clearance, use of inulin or radioisotopes or a predictive equation for GFR.

Table D6. Long Term Living Kidney Donation Outcomes – Older donors versus Older Healthy Non-donor Controls

		Int	tervention/Control (n)			Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Mortality								I
Mjoen, 2013 Norway	D: 15.1 year N-D 24.9 years	Donors (1,901)	Non-Donors (32,621)			HR (95% CI) Age, years 1.10 (1.10 to 1.11)		Low to Moderate
Reese, 2014 United States	7.84 years	Donors age <u>></u> 55 (3368)	Non-Donors age <u>≥</u> 55 (3368)		0.90 (0	n/N (%) 152/3368(4.5%) 5.6 deaths per 1000 person/years 95% CI) 0.71-1.15)		Low to Moderate
		Donors age <u>></u> 60 (1648)	Non-Donors age <u>></u> 60 (1648)			95% CI) -0.95), p=0.03		
Berger, 2011 United States	5 years	Donors age >70 (219)	Non-donors age >70 (219)		<u>Survival</u> n/N (%) (95% CI) 209/219 (95.8) (91.4-98.1)	<u>Survival</u> n/N(%) (95% CI) 201/219 (91.8) (87.3-94.7)		Moderate
	10 years				<u>Survival</u> n/N(%) (95% CI) 22/219 (90.0) (83.5-94.0)	<u>Survival</u> n/N(%) (95% CI) 160/219 (73.0) (65.6-79.0) HR (95% CI)		-
Segev, 2010 United States	Up to 12 years	Donors (80,347)	Non-Donors (80,347)		control No numerical valu >60: matched control years, then curves 50-59: Donors have	0.37 (0.21-0.65) s suggest mortality hig s at 5-12 years follow- ues provided, differenc rols have higher death intersect e higher mortality until ave higher death rate	up. e at 12 years: rate until 12	Moderate

		Int	ervention/Control (n)			Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					years 18-39: matched co years	ontrols have higher de ontrols have higher de due to very large sam	ath rate at 5-12	
Cardiovascular Out	comes							
Reese, 2014 United States	7.84 years	Donors age <u>></u> 55 (1312/3368) with Medicare	Non-Donors age ≥55 (1312/3368) with Medicare			n HR (95% CI) 1.20), p=0.70		Low- Moderate
		Donors age <u>></u> 60 (?/1648)	Donors age <u>≥</u> 60 (?/ 1648)			D events or death, 0.72		
Garg, 2012	Median 6.5 (max 17.7)	Donors age <u><</u> 55 (1741)	Non-Donors age <55 (17410)		Death censored CV event n/N (%) 18/1741(1.4)	Death censored CV event n/N (%) 181/17410(1.4)	P for interaction 0.48	Moderate
						5% CI) 0.60-1.5)		
		Donors age <u>></u> 55 (287)	Non-Donors age <u>></u> 55 (2870)		Death censored CV event n/N (%) 8/287(4.4)	Death censored CV event n/N (%) 106/2870(6.4)		
						5% CI) 0.3-1.4)		
Psychosocial	<u> </u>						1	T
Clemens, 2011 Australia, Canada, Scotland	5.5 years (median)	Donor Age <u>></u> 43 (NR)	Non-Donors Age <u>≥</u> 43 (NR)		SF-36 Mental component summary Mean (SD) 54 (8) p=NS	SF-36 Mental component summary Mean (SD) 56 (6)		High
Other: Diabetes	7.04	Denera	Non Denero			250/ (1)		Low Medarate
Reese, 2014 United States	7.84 years	Donors age <u>></u> 55 (1312/3368) with Medicare	Non-Donors age <u>></u> 55 (1312/3368) with Medicare			95% CI) 1.32), p=0.80		Low-Moderate

		Inte	ervention/Control (n)			Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Other: Fragility Fr	actures	I				1	1	
Garg, 2012 Canada	6.9 years	Donors age <u>></u> 55 (285)	Non –Donors age ≥55 (2,850)		n/N (%) 8/285 Event rate/10,000 person years 43.2	n/N (%) 70/2850 Event rate/10,000 person years 39.5 5% CI)		Moderate
					1.14 (0.5 Age did not influ between donati	6 to 2.35) ence association on and fractures		
					AHR (95%) CI for for every 5 year i dona	raction 0.5 r a fragility fracture increase in age at ation: 05-1.54)		
Other: GI Bleeding	g							
Thomas, 2014 Canada	Median 8.4 years	Donors age <u>></u> 40 (1,190)	Non-Donors age <u>≥</u> 40 (11,900)			n/N (%) 209/11900 Event rate/10,000 person years 20.3 5% CI) 9 to 1.80)		Low to Moderate
Other: Kidney Sto	nes				1.10 (0.7	5 10 1.007		
Thomas, 2013	Median 8.4 years	Donors age >40 (1203)	Non-Donors age <u>></u> 40 (12,030)			Kidney stones with surgical intervention n/N (%) 107/12,030 Event rate per 10,000 – 10.4 5% CI) 9 to 1.78)		Moderate
					RR (95% CI) for surgical interver older	kidney stone with htion per 5 years r age: 90-1.50)		

		Inte	ervention/Control (n)			Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					0.68 (0.3 RR (95% CI) fo	$\frac{\text{Kidney Stones}}{\text{with Hospital}}$ $\frac{\text{Encounters}}{(\%)}$ 176/12,030 Event rate per 10,000 = 17.0 5% CI) 8 to 1.22) or kidney stone 9 to 5 years older 0.82-1.26)		

CI= ; OR=Odds Ratio; RR= Risk Ratio; SBP= Systolic Blood Pressure; SD= Standard Deviation

	Inte	rvention/Conti	rol (n)				
Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Median: 5.5	Donors <u>></u> 60 (117)	Donors age<60		n/N (%) 3/117 (2.5)	n/N (%) 9/422 (2.1)	_	Moderate
		(422)					
6.8	Donors <u>></u> 55 (287)	Donors age <55 (1741)		n/N (%) 13/287 (4.5)	n/N (%) 29/1741(1.6)		Moderate
D<18: 31.8+8 D 18-30: 29.2+10. 3	Donors <18 (39)	Donors 18- 30 (128)		n/N (%) 2/39 (5.1) P>0.99	n/N (%) 8/128 (6.2)		Moderate
Median: 6.3	Donors age>60 (3017)	Donors age 50-59 (13439)	Donors age 40-49 (24375) Donors age 18-39 (39516)	12 year mortality 9.4% HR (95% CI) 9.4 (7.3 to 12.1)	12 year mortality 3.5% HR (95% CI) 3.3 (2.6 to 4.1)	12 year mortality 1.3% HR (95% CI) 1.6 (1.2-2.0) reference	Low-Moderate
ents			•				
Median 6.5	Donors age <u>></u> 55 (287)	Donors age <55 (1741)		Event rate. 10,000 person years 4.4	Event rate. 10,000 person years 1.4		Moderate
Median 7.7		Donors (4650)		Age (per year) Medical Claims		
•							
Median 7.6 years	Donors age <u>></u> 60 (4,039)	Donors age 50-59 (16,840)	Donors age 40-49 (28,994)	Cum Incidence of ESRD at 15 years per 10,000 (95%CI) 70.2(30.4-161.8)	Cum Incidence of ESRD at 15 years per 10,000 (95%CI) 54.6 (34.8-85.4)	Cum Incidence of ESRD at 15 years per 10,000 (95%CI) 17.4(10.1-30.0)	Moderate
	Follow- Up Median: 5.5 6.8 D<18: 31.8+8 D 18-30: 29.2+10. 3 Median: 6.3 Median: 6.5 Median 7.7	Mean Follow- UpArm 1Follow- UpDonors ≥60 (117)Median: 5.5 Donors ≥55 (287)D<18: $31.8+8$ D 18-30: 29.2+10. 3Donors <18 (39)	Mean Follow- UpArm 1Arm 2Median: 5.5 Donors $\geq 60 (117)$ Donors age<60 (422) 6.8 Donors $\geq 55 (287)$ Donors age $<55 (1741)$ D<18: $31.8+8$ D 18-30: $29.2+10.$ 3 Donors $<18 (39)$ Donors 18- $30 (128)$ Median: 6.3 Donors $age>60(3017)$ Donors age $50-59$ (13439)Median 6.5 Donors $age \ge 55$ (287)Donors age $<55 (1741)$ Median 7.7 Donors $29.2+10.$ Donors $30 (128)$ Median 6.3 Donors $age \ge 55$ (287)Donors age $<55 (1741)$ Median 7.7 Donors $255 (1741)$ Donors age $<55 (1741)$ Median 7.7 Donors $2000000000000000000000000000000000000$	Follow- UpDonors ≥60 (117)Donors age<60 (422)Median: 5.5≥60 (117)Donors age <55 (287)	$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$	$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$

Table D7. Long Term Living Kidney Donation Outcomes – Older versus Younger Donors

		Inte	rvention/Conti	ol (n)		Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
				Donors age 18-39 (46,344)			Cum Incidence of ESRD at 15 years per 10,000 (95%CI) 29.4(21.4-40.2)	
Gibney, 2008 United States	NR	White Donors age <35 (17,281)	White Donors age ≥35 (38,463)		Donors on Transplant Wait-List n/N (%) 35/17,281 (0.2) RR (9	Donors on Transplant <u>Wait-List</u> n/N (%) 19/38,463 (.05) 5% CI)	_	High
		AA Donors age <35 (5,061)	AA Donors age <u>></u> 35 (5,268)			5 to 7.16) <u>Donors on Transplant</u> <u>Wait-List</u> n/N (%) 6/5,268 (.11)		
						5% CI) 5 to 17.89)		
Renal Function		1						1
Tsai, 2013 Taiwan	Mean: 5.4 years		Donors (105)			regression model predicting tial age: 0.999 (0.965-1.033		Low-moderate
Dols, 2011 Netherlands	5.5 years	Donors <u>≥</u> 60 (117)	Donors <60 (422)		MDRD eGFR <60mL/min n/N (%) 94/117 (80) p<0.001	MDRD eGFR <60mL/min n/N (%) 131/422 (31)		Moderate
Fehrman-Ekholm, 2011 Sweden	14 years		Donors (573)		Multiple	regression predicting eGFF urrent age: -0.6559 (0.0571		High
Lentine, 2010 United States	Median 7.7 years		Donors (4650)	Chronic Kidney Disease Adjusted hazard ratio (95% CI) Age (per year) Medical Claims 1.04 (1.03-1.06) p<0.05		1)	Moderate
MacDonald 2014 United States	D<18: 28.4 years	Donors <18 (23)	Donors 18- 30 (88)		<u>eGFR</u> <u>66.7+10.9</u> ml/min/1.73m ²	eGFR 66.5+16.8 ml/min/1.73m ²		Moderate

		Inte	vention/Cont	rol (n)		Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Árm 3	Arm 1	Arm 2	Arm 3	ROB
	D 18-30: 30.4 years				<u>eGFR<60 (%)</u> 26.1 P=0.19	<u>eGFR<60 (%)</u> 40.9		
					<u>eGFR<45 (%)</u> 4.3 P=0.97	<u>eGFR<45 (%)</u> 4.5		
					OR (95% CI) of eGF 0.53(0.21-1.34) adjus ethnicity, BMI, eGFR	sted for age, gender,		
Ibrahim, 2009 United States	12.2 years		Donors (255)		Ag	exol GFR <60 mL/min/1.73n Odds Ratio (95% CI) e, per year: 1.15 (1.08-1.21)	Moderate
El-Argoudy, 2007 Egypt	10.7 years	Donors age 51-69 (44)	Donors age 36-50 (120)	Donors age 21-35 (175)	Serum Creatinine mg/dL Mean (SD) 0.8 (1.2) Range: 0.6-5.4 p=0.01	<u>Serum Creatinine</u> mg/dL Mean (SD) 1.0 (1.1) Range: 0.6-4.0	Serum Creatinine mg/dL Mean (SD) 1.0 (3.0) Range: 0.5-1.2	High
Lee, 2007 Korea	Median: 5.4 years		Donors (104)			eGFR<60 mL/min per 1.73 Odd Ratio (95% CI) le, per year 1.06 (1.01-1.10)		High
Gracida, 2003 Mexico	6.7 years	Donors age >60 (81)	Normal donors ^e (422)		<u>GFR (mL/min)^b</u> 71	<u>GFR (mL/min)^b</u> 78.5		High
Von Zur-Muhlen 2014 Sweden	11 <u>+</u> 7	Done	or Age			orrelated with lower eGFR (p<0.0001)		Moderate
Proteinuria	1		1	1				
Dols, 2011 Netherlands	5 years	Donors age <u>></u> 60 (64)	Donors age<60 (206)			n/N (%) 8/206(3.9) 95% CI) 63 to 8.14)	-	Moderate
	10 years	Donors age <u>></u> 60 (15)	Donors age<60 (94)		n/N (%) 0/15 RR (9	n/N (%) 6/94 (6.4) 5% CI) 03 to 7.72)	_	1
El-Argoudy, 2007 Egypt	10.7	Donors age 51-69 (44)	Donors age 36-50 (120)	Donors age at 21-35 (175)	Proteinuria mg/24h, mean (SD) 141(53)	Proteinuria mg/24h, mean (SD) 133(49)	Proteinuria mg/24h mean (SD) 127(48)	High

		Inte	rvention/Conti	ol (n)					
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB	
					p=0.5				
MacDonald 2014 United States	D<18: 31.8 D 18-30: 29.2	Donors <18 (39)	Donors 18- 30 (128)		Proteinuria (≥1+ on random dipstick) (%): 15.4, P=0.80	Proteinuria (≥1+ on random dipstick) (%): 14.1		Moderate	
						teinuria adol vs adult: 57-3.03) ^c			
Hypertension	1								
Dols, 2011 Netherlands	5.5 years	Donors <u>></u> 60 (117)	Donors <60 (422)		n/N (%) 12/117 (10%) p=0.56	n/N (%) 25/422 (6%)		Moderate	
					RR (9 1.73 (0.9				
Fehrman-Ekholm, 2011 Sweden	14 years		Donors (573)		Multiple regression predicting SBP: β (SE), p ? Current age: Systolic BP 0.527 (0.068) p=0.0000 Diastolic BP 0.033(0.041) p=0.42				
Lentine, 2010 United States	Median 7.7 years		Donors (4,650)		ted HR (95% Cl) Age (per ye <u>Medical Claims</u> 1.06 (1.06-1.07) p<0.05 <u>Drug-Treated</u> 1.06 (1.05-1.07) p<0.05		Moderate	
Ibrahim 2009 United States	12.2 years		Donors (255)			Hypertension requiring medication OR (95% CI) Age, per year: 1.09 (1.04-1.13)			
El-Argoudy, 2007 Egypt	10.7 years	Donors age 51-69 44	Donors age 36-50 120	Donors age 21-35 175	<u>Hypertension requiring</u> <u>medication</u> 1med: N=15 2med: N=7 3med: N=1 RR (95% CI) 6.09 (3.48-10.68)	Hypertension requiring medication 1med: N=28 2med: N=11 3med: N=1 RR (95% CI) 3.89 (2.25–6.71)	<u>Hypertension</u> requiring medication 1med: N=8 2med: N=6 3med: N=1	High	
MacDonald 2014 United States	D<18: 31.8 D 18-30: 29.2	Donors <18 (39)	Donors 18- 30 (128)		Hypertension requiring medication (%): 35.9, P=0.70	Hypertension requiring medication 39.4		Moderate	
						<u>HTN adol vs adult:</u> /1-2.61) ^c .			

		Inte	rvention/Contro	ol (n)		Results		
Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Psychosocial		-	I I					
Gross, 2013 United States	17 years		Donors (2,455))		tal Health HRQoL Impairme ear increase in age at dona OR (95%CI) 0.74 (0.65 to 0.85)		High
Lentine, 2012 United States	Median 7.7 years		Donors (4650)		Depression Age at o		Moderate	
Clemens, 2011 Australia, Canada, Scotland	Median 5.5	Donor Age <u>≥</u> 43 Total I	Donor Age <43 N = 203		<u>SF-36 Mental</u> component summary Mean (SD) 54 (8) p=NS	<u>SF-36 Mental</u> <u>component summary</u> Mean (SD) 52 (9)		High
Mjoen, 2011 Norway	Median 15.1	Ľ	Donors (71/1,37	7)	Older age at donation	risk of having doubt	Moderate	
Johnson, 1999 United States	NR	Donors age <u>></u> 40 Total №	Donors age <40 N = 524		SF-36 Mental health score ^d Mean (SD) 81 (NR)	SF-36 Mean Mental health score Mean (SD) 80 (NR)		High
Diabetes								•
Lentine, 2010 United States	Median 7.7		Donors (4,650)		Adjusted HR (95% CI) Age (per year) <u>Medical Claims</u> 1.05 (1.03-1.06) <u>Drug-Treated</u> 1.05 (1.03-1.07)			
MacDonald 2014 United States	D<18: 31.8 year D 18-30: 29.2 years	Donors <18 (39)	Donors 18- 30 (128)		Diabetes Requiring Medication (%): 5.1, P=0.19 <u>OR (95% CI) of diat</u> <u>0.61(0.15</u>	Diabetes requiring Medication 12.5 Detes adol vs adult:		Moderate
Other – Fragility Fra	actures				· · · · · · · · · · · · · · · · · · ·			•
Garg, 2012 Canada	6.9	Donors age <u>></u> 55 years (285)	Donors age <55 years (1,730)		No. of events/ No. at risk 8/285 Event rate/10,000	No. of events/ No. at risk 17/1730 Event rate/10,000		Low

Study, Year, Country	Mean Follow- Up	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					person years 43.2	person years 12.7		
						5% CI) 4 to 6.55)		
Other – GI Bleed						· · · · ·		
	Median 8.4 years	Donors <u>></u> 40 (1190)	Donors <40 years (819)		n/N (%) 25/1,190 (2.1) Event rate/10,000 person years 23.4	n/N (%) 10/819 (1.2) Event rate/10,000 person years 11.9		Moderate
		RR (95% CI) 1.72 (0.83 to 3.56)						
Other - Kidney Stone	es							
	Median 8.4 years	Donors <u>≥</u> 40 (1203)	Donors <40 years (816)		Kidney Stones with Surgical Intervention n/N (%) 10/1,203 (.83) Event rate/10,000 person years 9.4	<u>Kidney Stones with</u> Surgical Intervention n/N (%) 6/816 (.73) Event rate/10,000 person years 6.8		Moderate
					RR (9	5% CI)		
						1 to 3.09)		
					Kidney Stones with Hospital Encounters n/N (%)	Kidney Stones with Hospital Encounters		
					12/1,203 (.99) Event rate/10,000	n/N (%) 11/816 (1.3) Event rate/10,000		
					person years 11.1	person years 13.5		
					RR (95% CI) 0.74 (0.33 to 1.67)			

CI= ; NR= Not Reported; NS= Not statistically different; OR=Odds Ratio; RR= Risk Ratio; SBP= Systolic Blood Pressure; SD= Standard Deviation

Table D8. Long Term Living Kidney Donation Outcomes – Male versus Female Comparisons in Donorsversus Healthy Non-Donors

Mortality Median Norway Median C: 24 9 years Median C: 24 9 years Male sex predicting all-cause mortality in a cohort of donors (1901) and controls (32.621) Adjusted HR (95% CI) ^a 1.52 (1.41-1.65) Moderate Segev. 2010 United States Up to 12 years Male Donors (33,380) Male Non-Donors (33,380) Kaplan-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Low-Moderate Up to 12 years Female Donors (46,967) Female Non- Donors (46,967) Kaplan-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Low-Moderate Cardiovascular Outcomes Female Donors (812) Male Non-Donors (8120) Death (8120) Death censored CV event n/N (%) 20/812(3) Death censored CV event n/N (%) 22/12/16(2,4) P for interaction 0.1 Low-Moderate Female Donors (1216) Female Non-Donors (1216) Female Non-Donors (1216) Male Non-Donors (1216) Death censored CV event n/N (%) 22/12/16(2,4) Death censored CV event n/N (%) 22/12/16(2,4) Death censored CV event n/N (%) 22/12/16(2,4) Death censored CV event n/N (%) 22/12/16(2,4) Death censored CV event n/N (%) 22/12/16(2,4)	Study, Year,	Mean	In	tervention/Control (n)					
Mjoen, 2013 Norway Median D: 15.1 years Male sex predicting all-cause mortality in a cohort of donors (1901) and controls (32,621) Adjusted HR (95% CI) ^a Moderate Segev, 2010 United States Up to 12 years Male Donors (33,380) Male Non-Donors (33,380) Kapian-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Low-Moderate Up to 12 years Female Donors (46,967) Female Non- Donors (46,967) Female Non- Donors (46,967) Kapian-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Low-Moderate Cardiovascular Outcomes Female Donors (8120) Female Non-Donors (8120) Emale Non-Donors (8120) Male Non-Donors (8120) Death censored CV event n/N (%) 20/812(3.3) P for interaction 0.1 Low-Moderate Female Donors (1216) Female Non-Donors (1216) Female Non-Donors (1216) Male Non-Donors (12160) Death censored CV event n/N (%) 22/1216(2,4) P for interaction 0.1 Low-Moderate	Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Norway years C: 24.9 years D: 15.1 years donors (1901) and controls (32,621) 1.52 (1.41-1.65) Segev, 2010 United States Up to 12 years Male Donors (33,380) Male Non-Donors (33,380) Male Non-Donors (33,380) Kaplan-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001	Mortality								
United States years (33,380) (33,380) (33,380) matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Up to 12 years Female Donors (46,967) Female Non- Donors (46,967) Kaplan-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001		D: 15.1 years C: 24.9	Male sex predi donors	cting all-cause mortality (1901) and controls (32	in a cohort of ,621)	Ac	ljusted HR (95% C 1.52 (1.41-1.65)	I) ^a	Moderate
Up to 12 years Female Donors (46,967) Female Non- Donors (46,967) Kaplan-Meir curves suggest mortality higher in matched controls at 5-12 years follow-up. No numerical values provided, difference at 12 years around 1% Log-rank P<.001 Cardiovascular Outcomes Male Donors (812) Male Non-Donors (812) Death (8120) Death censored CV (8120) Death censored CV event n/N (%) 20/812(3.3) Death censored CV event n/N (%) 20/812(0.3) P for interaction 0.1 Low-Moderate Female Donors (1216) Female Non-Donors (12160) Female Non-Donors (12160) Death censored CV event n/N (%) 20/812(3.3) Death censored CV event n/N (%) 22/1216(2.4) P for interaction 0.1 Low-Moderate						matched co	ontrols at 5-12 year values provided, d years around 1%	s follow-up.	Low-Moderate
Cardiovascular Outcomes Garg, 2012 Canada Median 6.5 (max 17.7) Male Donors (812) Male Non-Donors (8120) Death (8120) Death censored CV event n/N (%) 20/812(3.3) Death censored CV event n/N (%) 367/8120 (6.2) P for interaction 0.1 Low-Moderate HR (95% Cl) 0.50 (0.30-0.80) Female Donors (1216) Female Non-Donors (12160) Female Non-Donors (12160) Death censored CV event n/N (%) 22/1216(2.4) Death censored CV event n/N (%) 22/1216(2.4) HR (95% Cl) 0.90 (0.60-1.40) Death censored CV event n/N (%) 22/12160 (2.8) Death censored CV event n/N (%) 22/12160 (2.8)						matched co			
Canada (max 17.7) (812) (8120) censored CV event n/N (%) 20/812(3.3) censored CV event n/N (%) 367/8120 (6.2) interaction 0.1 HR (95% Cl) 0.50 (0.30-0.80) Female Donors (1216) Female Non-Donors (12160) Female Non-Donors (12160) Death censored CV event n/N (%) 22/1216(2.4) Death censored CV event n/N (%) 22/12160 (2.8) HR (95% Cl) (2.8) HR (95% Cl) 0.90 (0.60-1.40) 0.1 HR (95% Cl) 0.90 (0.60-1.40) HR (95% Cl) 0.90 (0.60-1.40)	Cardiovascular O	utcomes							•
Female Donors (1216) Female Non-Donors (12160) Death Censored CV event n/N (%) 22/1216(2.4) Death censored CV event n/N (%) 243/12160 (2.8) HR (95% CI) 0.90 (0.60-1.40) 0.50 (0.30-0.80)						censored CV event n/N (%)	censored CV event n/N (%)	interaction	Low-Moderate
(1216) (12160) (12160) (12160) (12160) (12160) (12160) (22/1216(2.4)) (2.8) HR (95% CI) (0.90 (0.60-1.40)									
HR (95% CI) 0.90 (0.60-1.40)						censored CV event n/N (%)	censored CV event n/N (%) 243/12160		
Mjoen, 2013 Median Male sex predicting ESRD in donors (1901) versus Adjusted Hazard Ratio (95% CI) ^a Moderate	ESRD	Modion	Malo aoy prodi	oting ESPD in donors (1		Adjusts	d Hazard Datia (0)		Modorato

Study, Year,	Mean	Inte	rvention/Control (n)			Results		
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Norway	D: 15.1 C: 24.9		controls (32,621)		0.	90 (0.43-1.88)		
Psychosicial	77		Mala was danama		Dennesien	Democratica	Г Г	Madausta
Lentine, 2012 US	7.7 years	Male donors (2111)	Male non-donors (2111)		Depression diagnosis per 100 person- years 3.1	Depression diagnosis per 100 person- years 4.7		Moderate
		Female donors (2539)	Female Non- donors (2539)		Depression diagnosis per 100 person- years 6.6	Depression diagnosis per 100 person- years 9.2		
Fragility Fracture			1			-		
Garg, 2012 Canada	Median 6.5	Male Donors (805)	Male Non-Donors (8050)		n/N (%) 8/805 Event rate/10,000 person years 12.9 RR (95% 1.0 (0.5 t		P for interaction 0.7	Moderate
		Female Donors (1210)	Female Non- Donors (12100)		n/N (%) 17/1210 Event rate/10,000 person years 18.8 RR (95% 0.8 (0.5 t	n/N (%) 197/12100 Event rate/10,000 person years 22.4 6 CI)		
GI Bleeding					0.0 (0.0 (• • • • •		
Thomas, 2014 Canada	Median 8.4 years	Male Donors (808)	Male Non-Donors (8080)		n/N (%) 12/808 Event rate/10,000	n/N (%) 135/8080 Event rate/10,000	P value for interaction 0.2	Moderate

Study, Year,	Mean	Inte	rvention/Control (n)			Results			
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB	
					person years 15.7	person years 17.9			
		Female Donors (1201)	Female Non- Donors (12010)		n/N (%) 23/1201 Event rate/10,000 person years 20.1	n/N (%) 140/12010 Event rate/10,000 person years 12.9			
Kidney Stones									
Thomas, 2013 Canada	Median 8.4 years	Male Donors (806)	Male Non-Donors (8060)		n/N (%) Kidney stones with surgical intervention 7/806 Event rate/10,000 person years 9.1 n/N (%) Hospitalizations for Kidney stones 7/806 Event rate/10,000 person years 9.1	n/N (%)Kidney stones with surgical intervention 72/8060 Event rate/10,000 person years 13.7 n/N (%) Hospitalizat ions for Kidney stones 178/8060 Event rate/10,000 person years 14.2	P for interaction 0.4 P for interaction 0.03	Moderate	
		Female Donors (1213)	Female Non- Donors (12130)		n/N (%) Kidney stones with surgical intervention	n/N (%)Kidney stones with surgical			

Study, Year,	Mean	Inter	vention/Control (n)			Results		
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					9/1213 Event rate/10,000 person years 7.7	intervention 77/12130 Event rate/10,000 person years 7.0		
					n/N (%) Hospitalizations for Kidney stones 16/1213 Event rate/10,000 person years 7.7	n/N (%) Hospitalizat ions for Kidney stones 122/12130 Event rate/10,000		
						person years 7.0		

CI=Confidence Interval ; ESRD= End-Stage Renal Disease; HR= Hazard Ratio ^a adjusted for age, sex, year of inclusion, blood pressure, BMI and smoking status

Study, Year,	Mean	Interv	vention/Co	ntrol (n)		Results				
Country	Follow-Up (years)	Arm 1	Arm 2 Arm		Arm 1	A	Arm 2 Arm 3		ROB	
Mortality										
Tsai 2013 Taiwan	Mean: 5.4 years	Male Donors (45)	Femal Donors (n/N (%) 1/45(2.2)		N (%) D (1.7)		Low-moderate	
Segev 2010	Median: 6.3	Male Donors	Femal		12 year mortality 2	2.7%	<u>12 ye</u>	ar mortality 1.9%	Low-Moderate	
United States	years	(33,380)	donors (46,967			Male	5% CI) ^a : e Sex 5 to 2.0)			
Cardiovascular E	vents	I								
Lentine 2010 United States	Median 7.7 years		Donors (46	50)		Male Medial	R (95% CI) [♭] ∋ sex Claims 3.10) p<0.05		Moderate	
Garg 2012 Canada	Median 6.5 (max 17.7)	Female Don (1216)	ors M	ale Donors (812)	Death censored CV event n/N Death censored ((%) (%)		nsored CV event n/N (%) 20/812(3.3)	Low-Moderate		
					HR of dath censred major cardiovascular event HR(9 0.57(0.26-1.23)		event HR(95% CI):			
ESRD										
Tsai 2013 Taiwan	Mean: 5.4 years	Male Donors (45)	Femal Donors (n/N (%) 1/45(2.2)		N (%) 0 (1.7)		Low-moderate	
Cherikh 2011 United States	9.8 years	Male Donors (24,146)	Femal Donor (32,312	s	n/N (%) 73/24,146 (0.30) RR (95	53/32,3	N (%) 312 (0.16)	-	High	
Wafa 2011	NR (>5	Male Donors	Femal	e	2.24 (1.30 n/N (%)		N (%)		Moderate- High	
Egypt	years)	(953)	Donor (1047	s	6/953 (.62) RR (95 3.29 (0.67	2/10 % CI)	47 (.19)	-		
Gibney 2008 United States	NR	White Male Donors (23,413)	White Fer Donor (32,33	s	White Male Donors on transplant waiting list n/N (%) 42/23,413 (.18)	White Dor transpla	e Female hors on ant waiting list N (%)		Moderate-High	

Table D9. Long Term Living Kidney Donation Outcomes – Male versus Female Donors

Study, Year,	Mean	Interv	vention/Control ((n)		Results				
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB		
						12/32,331 (.04)				
					RR (95 4.83 (2.54					
	NR	African American Male Donors (4,545)	African American Female Donors (5,784)		AA Male Donors on <u>Transplant Wait-List</u> n/N (%) 29/4,545 (.63)	AA Female Donors on Transplant Wait- List n/N (%) 21/5,784 (.36)		Moderate-High		
					RR (95 1.76 (1.00					
Muzaale 2014 United States	15 years post- donation	Male Donors (39449)	Female Donors (56768)		Cumulative Incidence of ESRD at 15 years per 10,000 (95% CI) 44.1(22.9-59.1)	Cumulative Incidence of ESRD at 15 years per 10,000 (95% CI) 21.1(14.9-29.9)		Low-Moderate		
Renal Function					111(22:0 00:1)	21.1(11.0 20.0)				
Tsai 2013 Taiwan	Mean: 5.4 years	Male Donors (45)	Female Donors (60)		eGFR<60 n/N (%) 20/45(44.4)	eGFR<60 n/N (%) 16/60 (26.7)		Moderate		
					RR (95 1.67 (0.98	3 to 2.84)				
Lentine 2010 United States	Median 7.7 years		Donors (4650)		Ň	sted Hazard Ratio (95% Male sex /ledical Claims for CKD 1.64 (1.16-2.34)p<0.05	CI)			
Ibrahim 2009 United States	Mean 12.2 years		Donors (255)		lohex	ol GFR <60 mL/min/1.7 Female sex OR (95% CI) [°] .11 (1.11-8.67) p=0.003		Moderate		
Lee 2007 Korea	Median: 5.4 years		Donors (104)		MDRD e	GFR<60 mL/min per 1. Male sex OR (95% CI) 0.65 (0.17-2.45) p=0.52		Moderate-High		
Karakayali 1998	10.2 years	Male Donors (45)	Female Donors (57)		GFR 81.6 <u>+</u> 10.2 P not provided	GFR 79.4 <u>+</u> 12.3		Moderate		

Study, Year,	Mean	Inter	vention/Control	(n)		Results		
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Von Zur-Muhlen 2014 Sweden	11 <u>+</u> 7 years	Male Donors (166)	Female Donors (229)		MDRD eGFRml/min/1.73m ² 69 <u>+</u> 13 P<0.01	MDRD eGFRml/min/1.73m 65 <u>+</u> 12		Moderate
Proteinuria								
Tsai 2013 Taiwan	Mean: 5.4 years	Male Donors (45)	Female Donors (60)		>150 mg/day or >1+ on UA n/N (%) 7/45(15.6) RR (95 1.87 (0.63			Moderate
Ibrahim 2009 United States	Mean 12.2 years	Female sex pre	dicting Albuminu	ria (255)		Albuminuria Odds Ratio (95% CI) .31 (0.12-0.79) p=0.01		Moderate
Hypertension								
Tsai 2013 Taiwan	Mean: 5.4 years	Male Donors (45)	Female Donors (60)		>140/90 mgHg n/N (%) 6/45(13.3) RR (95 2.00 (0.60			Moderate
Lentine 2010 United States	Median 7.7		Donors (4650)	I	M Adjus	lale sex predicting HTN sted Hazard Ratio (95% Medial Claims 1.13 (0.98-21.31) Drug-Treated .21 (1.03-1.43) p<0.05		Moderate
El-Argoudy 2007 Egypt	10.7 years	Male Donors (129)	Female Donors (201)		HTN (>140/90) 17.8% P=0.03	HTN (>140/90) 24.7%		High
Psychosocial	1	1	Г	1	1			1
Johnson 1999 United States	NR	Male Donors (204)	Female Donors (320)			more likely to find the c ion more stressful. (SF OR (95% CI) 1.8 p=0.1		High
Mjoen 2011, Norway	12.6 years	Male Donors 544 (39.5%)	Female Donors 833(60.5%)		Male donors were no doubt toward Adjusted ^d Ol	t more likely to have ds donation		Moderate

Study, Year,	Mean	Inter	vention/Control (n)		Results		
Country	Follow-Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					1.1 (0.1	7-1.8)		
Lentine 2012 US	4.9 years	Male donors (2111)	Female donors (2539)		Depression diagnosis per 100 person-years 3.1	Depression diagnosis per 100 person-years 6.6		Moderate
Diabetes								
Lentine 2010 United States	Median 7.7 years		Donors (4650)		Male A	e sex predicting Diabete Adjusted HR (95% CI) Medial Claims 0.91 (0.68-1.22) Drug-Treated 1.10 (0.73-1.66)	S	Moderate
Other – Fragility	Fractures				•			
Garg 2012 Canada	6.9 years	Male Donors (805)	Female Donors (1210)		n/N (%) 8/805 (.99) Event rate/10,000 person years 12.9	n/N (%) 17/1,210 (1.4) Event rate/10,000 person years 18.8		Moderate
					RR (95			
					0.71 (0.31	to 1.63)		
Other - GI Bleed								
Thomas 2014 Canada	Median 8.4 years	Male Donors (808)	Female Donors (1,201)		n/N (%) 12/808 (1.5) Event rate/10,000 person years 15.7	n/N (%) 23/1,201 (1.9) Event rate/10,000 person years 20.1		Moderate
					RR (95 0.78 (0.39			

CI=Confidence Interval ; HTN= Hypertension ; OR=Odds Ratio; RR= Risk Ratio ^a adjusted for sex, time since donation, current BMI, creatinine level at donation, smoking status and systolic and diastolic blood pressure.

^b calculated by means of multivariate Cox regression

^c adjusted for age, sex, year of inclusion, blood pressure, BMI and smoking status ^dAdjusted for age, time since donation, relationship, medical problems, graft loss in recipient.

Table D10. Long Term Living Kidney Donation Outcomes: African American Donors versus HealthyAfrican American Non-Donors

Study, Year,	Follow-		ervention/Control (n)					
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Mortality								
Segev 2010 United States	Median: 6.3	African American Donors	African American Non-donors		matched black non donors starting at	suggest mortality hig donors compared to 2 years of follow-up, I k P<0.001	black	Low-Moderate
		White Donors	White Non- donors		matched white non- donors starting at	suggest mortality hig donors compared to 2 years of follow-up, l k P<0.001	white	
ESRD								
Muzaale 2014 United States	Median 7.6 years	African American Donors 12,387 (13)	African American Non-donors 12,387 (13)		Absolute risk of ESRD per 10,000 (95% CI) 74.7 (47.8-105.8)	Absolute risk of ESRD per 10,000 (95% Cl) 23.9 (1.6-62.4)	RR (95% CI) 3.0 (1.9 to 4.7) Absolute risk increase per 10,000 50.8(p<0.001)	Low-Moderate
		Hispanic Donors 12,061 (13)	Hispanic Non- Donors 12,061 (13)		Absolute risk of ESRD per 10,000 (95% CI) 32.6 (17.9-59.1)	Absolute risk of ESRD per 10,000 (95% Cl) 6.7 (0.0-15.0)	RR (95% Cl) 4.7 (2.1 to 10.7) Absolute risk increase per 10,000 25.9 (p=0.002)	
		White Donors 71,769 (75)	White Non- Donors 71,769 (75)		Absolute risk of ESRD per 10,000 (95% CI) 22.7 (15.6-30.1)	Absolute risk of ESRD per 10,000 (95% CI) 0(0.0-0.0)	RR (95% Cl) 45.0 (2.7 to 741.8) Absolute risk increase per 10,000 22.7 (p<0.001)	
Renal Function								-
Doshi 2013 USA	6.8	African American Donors (103)	African American Non-donors (235)		Change in creatinine 0.9 ± 0.2 to 1.2 ± 0.3 mg/dL Change in eGFR 109 ± 20 to 77 ± 19 mL/min/1.73 m2	e Change in creati (stable) 0.9 ± 0.2 mg/dL Change in eGFF (stable) 109 ± 17 mL/mir	<60 mL/min/1 .73 m2 RR (95% CI) 75	

Study, Year,	Follow-	Inte	ervention/Control (n)					
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
						m2	1236)	
					Proportion with eGFR <60 mL/min/1.73 m2 16 (15.5%) eGFR<45 mL/min/1.73 m2 6(6%)	Proportion with eGFR <60 mL/min/1.73 m2 0 eGFR<45 mL/min/1.73 m2 0		
					eGFR<30 mL/min/1.73 m2 0	eGFR<30 mL/min/1.73 m2		
Proteinuria					0	0		
Doshi, 2013 United States	Mean D: 6.8 years ND: 6.4 years	African American Donors (103)	African American Non-donors (235)		Urinary Albumin mean (SD) 15lg/mg (41) p= 0.06 Microalbuminuria	Urinary Albumin Mean (SD) 7lg/mg (11) Microalbuminuria		Moderate
	-				n/N (%) 6/103 (5.8)	n/N(%) 9/235 (3.8)		
						95%CI) 55 to 4.16)		
Hypertension		1					1	
Doshi, 2013 United States	Mean D: 6.8 ND: 6.4	African American Donors (103)	African American Non-donors (235)		HTN (BP>140/90 or medications) n/N (95% CI) 42/103(40.8)	HTN (BP>140/90 or medications) n/N(95%) 42/235(17.9)		Moderate
					22.9% (12. RR of HTN: 2	difference 2 to 33.6%) 2.3(1.6 to 3.3) (1.7-3.4) ^a		
Diabetes	•							
Doshi, 2013 United States	Mean D: 6.8 ND: 6.4	African American Donors (103)	African American Non-donors (235)		Diabetes 2 (1.9%)	Diabetes 4 (1.7%) 5% CI)		Moderate
	ND. 0.4	(105)	(200)			1 to 6.13)		

BP= Blood Pressure; D= Donors; HTN= Hypertension; ND=Non-Donors

^aadjusted for baseline differences in medical insurance and eGFR.

Study, Year,	Follow-	Inte	ervention/Control (n	1)		Results		
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Mortality								
Segev 2010 United States	Median: 6.3 years	Black donors (10,505)	Hispanic donors (9,846)	White donors (9,846)	HR (95% CI) 1.3 (1.0-1.6) 12 year mortality 2.8%	HR (95% CI) 0.6 (0.4-0.9)	Reference 12 year mortality 1.7%	Low-Moderate
Storsley 2010 Canada	14 years	Aboriginal Donors (38)	White Donors (76)		n/N (%) 4/38 (10.5) p= NS RR (95%	n/N (%) 6/76 (7.9)	-	High
					1.33 (0.40 t			
Cardiovascular Ev	vents							
Lentine 2010 United States	Median 7.7 years	Black Donors (609)	Hispanic Donors (381)	White (non- Hispanic) Donors (3548)	<u>Cardiovascular</u> <u>Disease</u> Adjusted RR ^a (95%CI) Medical Claims 1.15(0.63-2.11)	Cardiovascular Disease Adjusted RR ^a (95% CI) Medical Claims 0.91(0.37-2.26)	Reference	Moderate
ESRD					, , , ,			
Cherikh 2011 United States	9.8 years	Black donors (7,333)	White donors (40,398)		n/N (%) 59/7,333 (0.8) RR (95% 6.02 (4.16 t		-	Moderate
Muzaale 2014 United States	Median 7.6years	Black Donors n (%) 12,387 (13)	Hispanic Donors n (%) 12,061 (13)	White Donors N (%) 71,769 (75)	Cum Incidence of ESRD at 15 years per 10,000 (95% CI) 74.7 (47.8-105.8)	Cum Incidence of ESRD at 15 years per 10,000 (95% CI) 32.6 (17.9-59.1)	Cum Incidence of ESRD at 15 years per 10,000 (95% CI) 22.7 (15.6- 30.1)	Low-Moderate
Lentine 2010 United States	Median 7.7 years	Black Donors (271)	Hispanic Donors (179)	White (non- Hispanic) Donors (1,786)	<u>CKD requiring</u> <u>dialysis</u> n/N (%) 2/271 (0.7) p=0.02 RR(95% CI) ^b	<u>CKD requiring</u> <u>dialysis</u> n/N (%) 1/179 (0.5) p=0.10 RR(95% CI) ^b	CKD requiring dialysis n/N (%) 0/1786 (0)	Moderate

Table D11. Long Term Living Kidney Donation Outcomes: Comparison between Donor Racial Subgroups

Study, Year,	Follow-	Inte	ervention/Control (n)		Results		
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					13.19 (1.20 to 144.95)	29.78 (1.22 to 728.47)		
Storsley 2010 Canada	14 years	Aboriginal Donors (38)	White Donors (76)		n/N (%) <u>1/38 (2.6)</u> RR (95% 5.92 (0.25 to		-	High
Gibney 2007 United States	Mean 17.6 years	African American Donors* n (%) 8889 (14)	Hispanic Donors n (%) 7375 (12)	White Donors n (%) 42,419 (68)	<u>On Transplant Wait-</u> <u>list</u> n/N(%) 45/8,889 (0.5) RR (95% Cl) ^a 5.24 (3.43 to 7.99)	<u>On Transplant</u> <u>Wait-list</u> n/N(%) 11/7,375 (.14) RR (95% CI) ^a 1.54 (0.79 to 3.0)	On <u>Transplant</u> <u>Wait-list</u> n(%) 41(.09)	Moderate-High
Renal Function	•	•	•	•		i i i		
Lentine 2010 United States	Median 7.7 years	Black Donors Medicare Insured (271)	Hispanic Donors Medicare Insured (179)	White (non- Hispanic) Donors Medicare Insured (1,786)	AHR (95% CI) of Medical Claims for <u>CKD</u> 2.32(1.48-3.62)	AHR (95% CI) of Medical Claims for CKD 1.90(1.05-3.43)	Reference	Moderate
Lentine 2014 United States	Median 7.7 years	Black Donors (325) Medicare-Insured	Hispanic Donors (228) Medicare Insured	White (non- Hispanic) Donors (3,342) Medicare Insured	<u>CKD</u> Adjusted HR ^a (95% CI) 1.84 (1.37-2.47)	<u>CKD</u> Adjusted HR ^a (95% CI) 1.13 (0.75-1.70)	Reference	Moderate
		Black Donors (609) Privately Insured	Hispanic Donors (381) Privately Insured	White (non- Hispanic) Donors (3548) Privately Insured	<u>CKD</u> Adjusted HR ^a (95% CI) 2.32 (1.48-3.62)	<u>CKD</u> Adjusted HR ^a (95% CI) 1.90 (1.05-3.43) p<0.001	Reference	
Storsley 2010 Canada	14 years	Aboriginal Donors (31)	White Donors (64)		<u>eGFR (MDRD)</u> ml/min, mean (SD) 77 (17)	eGFR (MDRD) ml/min, mean (SD) 67 (13)		High
					Adjusted mean difference (95% CI ^c 5.9(-0.6-12.5)	Ref		
Proteinuria								
Lentine 2014	Median 6.0 years	Black Donors (325) Medicare-Insured	Hispanic Donors (228)	White (non- Hispanic)	Adjusted HR ^a (95% CI)	Adjusted HR ^a (95% CI)	Reference	Moderate

Study, Year,	Follow-	Inte	ervention/Control (n)		Results		
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
			Medicare Insured	Donors (3342) Medicare Insured	2.44 (1.45-4.11)	0.98 (0.40-2.44)		
		Black Donors (609) Privately Insured	Hispanic Donors (381) Privately Insured	White (non- Hispanic) Donors (3548) Medicare Insured	Adjusted HR ^a (95% CI) 2.27 (1.32-3.89)	Adjusted HR ^a (95% Cl) 1.47 (0.67-3.26)	Reference	
Storsley 2010 Canada	14 years	Aboriginal Donors (29)	White Donors (57)		>0.3g/day or abnormal alb/cr n/N (%) 6/29(21) RR (95%	>0.3g/day or abnormal alb/cr n/N(%) 2/57(4) 6 Cl)	-	High
Hypertension					5.89 (1.27 to	o 27.41)		
Lentine 2014	Median 6.0 years	Black Donors (325) Medicare-Insured	Hispanic Donors (228) Medicare Insured	White (non- Hispanic) Donors (3342) Medicare Insured	Adjusted HR ^a (95% CI) 1.41 (1.17-1.70)	Adjusted HR ^a (95% CI) 1.11 (0.95-1.46)	Reference	Moderate
		Black Donors (609) Privately Insured	Hispanic Donors (381) Privately Insured	White (non- Hispanic) Donors (3548) Medicare Insured	Adjusted Hazard Ratio ^a (95% CI) 1.52 (1.23-1.88)	Adjusted Hazard Ratio ^a (95% CI) 1.36 (1.04-1.78)	Reference	
Lentine 2010 United States	Median 7.7 years	Black Donors (609)	Hispanic Donors (381)	White (non- Hispanic) Donors (3548)	Adjusted RR ^a (95% CI) Drug-Treated 1.31 (1.02-1.68)	Adjusted RR ^a (95% CI) Drug-Treated 1.03 (0.73-1.46)	Reference	Moderate
Storsley 2010 Canada	14 years	Aboriginal Donors (31)	White Donors (64)		HTN (>140/90) 13(42%) Adjusted OR 6.3 (1.8-22.1) HTN >10 yrs post donation 13/21 (62%) P=0.001 HTN >20 yrs post donation	HTN (>140/90) 12(18%) Ref HTN >10 yrs post donation 7/38(18%) HTN >20 yrs post donation		High

Study, Year,	Follow-	Inte	ervention/Control (n	ı)		Results		
Country	Up (years)	Arm 1	Arm 2	Árm 3	Arm 1	Arm 2	Arm 3	ROB
					11/11 (100%) P<0.0001	3/16(19%)	-	
					# of meds 1: 9 (29%) 2: 2 (6.5%) 3: 2 (6.5%) P= NS	# of meds 1: 6 (9.3%) 2: 4 (6.2%) 3: 2 (3.1%)		
Psychosocial	•						•	•
Gross 2013 United States	17 years	White Donors (2282)	Black/African American Donors (113)		compared to participa	n. (p=0.0007). White n African Americans 4). Americans scored erceptions domain of	donors likely to (p=0.0034) and d higher on the f the SF-36	High
Lentine 2012 United States	Median 7.7 years		Donors (4650)		Depression diag	nosis by claims HR	race or Hispanic	Moderate
Diabetes		•						
Lentine 2014 United States	Median 6.0 years	Black Donors (325) Medicare-Insured	Hispanic Donors (228) Medicare Insured	White (non- Hispanic) Donors (3342) Medicare Insured	Adjusted HR ^a (95% Cl) 1.57 (1.16-2.12)	Adjusted HR ^a (95% Cl) 2.13 (1.56-2.92)	Reference	Moderate
		Black Donors (609) Privately Insured	Hispanic Donors (381) Privately Insured	White (non- Hispanic) Donors (3548) Pirvately Insured	Adjusted HR ^a (95% CI) 1.64 (1.07-2.51)	Adjusted HR ^a (95% CI) 1.86 (1.12-3.10)	Reference	
Lentine 2010 United States	Median 7.7 years	Black Donors (609)	Hispanic Donors (381)	White (non- Hispanic) Donors (3548)	Adjusted HR ^a (95% CI) Medical Claims 1.52 (1.00-2.30) Drug-Treated 2.31 (1.33-3.98)	Adjusted HR ^a (95% CI) Medical Claims 1.65(1.00-2.74) Drug-Treated 2.94(1.57-5.51)	Reference	Moderate
Storsley 2010 Canada	14 years	Aboriginal Donors (31)	White Donors (64)		Overall 6(19%) p=0.005 >20 yrs of follow-up	Overall 1(2%) >20 yrs of follow-	-	High

Study, Year,	Follow-	Inte	ervention/Control (n	1)		Results		
Country	Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
					5/11(45%) P=0.007	up 1/15(6%)		

BP= Blood Pressure; CI=Confidence Interval; ESRD= End-Stage Renal Disease; GFR= Glomerular Filtration Rate; HR= Hazard Ratio; MDRD= Modification of Diet in Renal Disease, OR=Odds Ratio; RR= Risk Ratio, SD= Standard Deviation

^a adjustment method unclear ^bRisk Ratio is calculated with White donors as the comparison.

^c Sample size for other races too small. No statistical differences.

Study, Year	Mean Follow-	Intervention/C Pre-Donation			Results	Results		
	Up (Years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	-
Mortality				1		_		
Mjoen 2013 Norway	D: 15.1 year N-D 24.9 years	Donors (1,901)	Non-Donors (32,621)		1.0	HR (95% 0 BMI, kg/m 1 (0.99 to 1.03	1 ²	Moderate
Cardiovascular E			1	•	I			
Mjoen 2013 Norway	D: 15.1 year N-D 24.9 years	Donors (1,901)	Non-Donors (32,621)		1.	HR (95% 0 BMI, kg/m .03(1.00-1.07),	1 ²	Moderate
ESRD		•	•		•			
Mjoen 2013 Norway	D: 15.1 year N-D 24.9 years	Donors (1,901)	Non-Donors (32,621)		1.1	HR (95% 0 BMI, kg/m 3 (0.96 to 1.32	1 ²	Moderate
Renal Function				<u> </u>				
Gracida 2003 Mexico	Mean 6.7	Obese (BMI >30 kg/m ²) Donors (81)	Normal Donors ^a (422)		GFR (mL/min) ^a 83.9 SD: NR	^a : GFR (mL/min) 78.5 SD: NR		High
Ibrahim 2009 United States	Mean 12.2		Donors (255)	I		ol GFR <60 mL OR (95% (, per unit: 1.12 p=0.02	/min/1.73m ² CI)	Moderate
Von Zur_Muhlen 2014 Sweden	11 <u>+</u> 7		Donors (375)				correlated with lower (P<0.0001)	Moderate
Hypertension								
Ibrahim 2009 United States	Mean 12.2		Donors (255)			tension requirin R (95% CI) BMI 1.12 (1.04-1 p=0.003	, per unit: .21)	Moderate
Gracida 2003 Mexico	Mean 6.7	Obese (BMI >30 kg/m ²) Donors (81)	Normal Donors (422)		Mean Arterial Pressure (MAP) 91.2 mmHg	Mean Arter Pressure (MAP) 88.2 mmH		High

Table D12. Long Term Living Kidney Donation Outcomes – Obese Donors versus non-obese donors

Psychosocial								
Gross 2013 United States	Mean 17	Donors with BMI >=35 ^b (102)	Donors with BMI 30 – 34.9 (329)	Donors with BMI 25 – 29.9 (883)	Physical HRQoL impairment ^c : OR (95% CI): 4.32 (2.37-7.87)	Physical HRQoL impairment ^c : OR (95% CI): 2.85 (1.84-4.42)	Physical HRQoL impairment ^c : OR (95% CI): 1.84 (1.31-2.65)	Moderate-High

BMI= Body Mass Index; CI=Confidence interval; GFR= Glomerular Filtration Rate; HRQoL= Health-Related Quality of Life (higher score=higher quality); OR=Odds Ratio ^a method of GFR estimation of measurement not reported ^b reference group BMI <25

^c Physical HRQoL impairment defined as PCS > - 1 SD below sex-by-age norms

dAdjusted for age, gender, year of inclusion, systolic BP, smoking and multiple imputations for missing values performed

Table D13. Long Term Living Kidney Donation Outcomes – Donors with lower renal function versus donors with normal renal function

Study, Year, Country	Mean follow-up		n/Control (n) ation GFR		Results		ROB
	(years)	Arm 1 Arm	2 Arm 3	Arm 1	Arm 2	Arm 3	-
Renal Function							
Lee 2007 Korea	Median: 5.4 years Range: 4.5-14.25	Donor	s (104)	Per unit change ir	min per 1.73 m²)	Moderate to High	
Von Zur_Muhlen 2014 Sweden	11years	Donor	s (375)	Lower measured GFR eGFR a	High		
Tsai 2013 Taiwan	Mean: 5.4 years	MDRD eGFR (ml/min per 1.73 m ²)≥ <90	MDRD eGFR (ml/min per 1.73 m ²) <u>></u> 90	Median time to CKD 3.55 years Median time to CKD AHR (95% CI) of developing CKD (eGFR<60 ml/min/1.73m2) per 1ml/min/1.73m2 if eGFR at donation: 0.95(0.92-0.99), p=0.021 AHR (95% CI) of developing CKD (eGFR<60 ml/min/1.73m2) per 1 mg/dL of serum creatinine at donation: 0.044(0.0-4.36), p=0.183		Low to Moderate	
Proteinuria					•		
Von Zur_Muhlen 2014 Sweden	11years	Donor	Donors (375)		Lower measured GFR at donation was not correlated to urine albumin creatinine ratio at follow-up.		
Hypertension		•					•
Von Zur_Muhlen 2014 Sweden	11years	Donor	Donors (375)		Lower measured GFR at donation was correlated to Mean Arterial Pressure at follow-up.		

CG= Cockcroft-Gault; CI= Confidence Interval; CKD= Chronic Kidney Disease; eGFR= estimated Glomerular Filtration Rate; MDRD= Modification of Diet in Renal Disease; OR= Odds Ratio; SD= Standard Deviation

Study,	Mean	Ir	ntervention/Control (n)			Results		ROB
Year, Country	Follow- Up (Years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
Mortality								
Okamoto 2010 Japan	7.3 years	Glucose Intolerant Donors ^a (65)	Normal glucose tolerance Donors (330)			n/N(%): 14/330 (4.1%) 95% CI) 32 to 3.68)		Moderate to High
ESRD								
Okamoto 2010 Japan	7.3 years	Glucose Intolerant Donors ^a (65)	Normal glucose tolerance Donors (330)			n/N (%) 2/330 (0.6) 95% CI) 5 to 20.65)		Moderate to High
Renal funct								
Chandran 2014 United States	Mean: 10.2	Impaired fasting glucose Donors ^b (45)	Normal fasting glucose Donors (45)		MDRD eGFR (mL/min/1.73m ²) mean (SD): 70.7 (16.1) p=.21	MDRD eGFR (mL/min/1.73m ²) mean (SD): 67.3 (16.6)		Moderate
Okamoto 2010	7.3	Glucose Intolerant Donors ^c (65)	Normal glucose tolerance Donors (330)		Renal Dysfunction self-report n/N(%): 5/65(7.7%)	Renal Dysfunction self-report n/N(%): 22/330(6.7%) P=0.690		Moderate to High
Proteinuria	-							
Chandran 2014 United States	Mean: 10.2	Impaired fasting glucose Donors ^b (45)	Normal fasting glucose Donors (45)		Albumin/creatine ratio (mg/g) mean (SD): 9.76 (23.6) p=.29	Albumin/creatine ratio (mg/g) mean (SD): 5.91 (11)		Moderate
Diabetes								
Chandran 2014 United States	Mean: 10.2	Impaired fasting glucose Donors ^o (45)	Normal fasting glucose Donors (45)		Diabetes n/N(%): 7/46(15.6%) p=.06	Diabetes n/N(%): 1/45 (2.2%)		Moderate
Okamoto 2010	Mean: 7.3	Glucose Intolerant Donors ^c (65)	Normal glucose tolerance Donors (330)		Diabetes by self- report n/N(%): 14/65(21.5%) Diabetes on meds	Diabetes by self- report n/N(%): 8/330 (2.4%) P<0.0001 Diabetes on meds		Moderate- High

Table D14. Long Term Living Kidney Donation Outcomes – Donors with impaired glucose tolerance versus donors with normal glucose tolerance

Study,	Mean	li	ntervention/Control (n)			Results		ROB
Year, Country	Follow- Up (Years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
					n/N(%): 17/65(26.2%)	n/N(%): 0/330(0%) P=NR		
Hypertensio	n							
Chandran 2014 United States	Mean: 10.2	Impaired fasting glucose Donors ^b (45)	normal fasting glucose Donors (45)		HTN- n/N(%): 16/45 (35.6%) p=.16	HTN n/N(%): 10/45 (22.2%)		Moderate
Okamoto 2010 Japan	7.3 years	Glucose Intolerant Donors ^a (65)	Normal glucose tolerance Donors (330)			BP>140/90 n/N(%): 73/330 (22.1%) 95% CI) 86 to 2.03)		Moderate to High
		Glucose Intolerant Donors ^a (65)	Normal glucose tolerance Donors (330)		HTN-(on medication) n/N(%): 9/65 (13.8%)	HTN-(on medication) n/N(%): 37/330 (11.2%)		
						95% CI) 63 to 2.43)		

CI= Confidence Interval; HTN= Hypertension; RR= Risk Ratio

^a Glucose Intolerant Donors were those with fasting blood sugar= \geq 110 mg/dL and 120-min blood sugar= \geq 140 mg/dL; and/or diagnosed Diabetes.

Table D15. Long Term Living Kidney Donation Outcomes – Donors with metabolic syndrome versusdonors without metabolic syndrome

Study, Year,	Mean		ntervention/Contro	l (n)		Results		ROB
Country	Follow- Up (Years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	
Renal Function								
Cuevas-Ramos 2011 Mexico	Median 5 years	Donors with MetS ^a (28)	Donors without MetS (112)		MDRD eGFR (mL/min/1.73 m ²) Mean (SD): 66.3 (12.7)	<u>MDRD eGFR</u> (mL/min/1.73 m ²) Mean (SD): 71.8 (16.2)		Moderate to High
Proteinuria								
Cuevas-Ramos 2011 Mexico	Median 5 years	Donors with MetS (28)	Donors without MetS (112)		Albuminuria mg/d mean(SD): 0.5 (0.6)	Albuminuria mg/d mean(SD): 0.2 (0.5)		Moderate to High

MetS= Metabolic Syndrome; SD= Standard Deviation

^a MetS defined as having 3 or more of three criteria 1) waist circumference of >88 cm in women or >102 cm in men; 2)

hypertriglyceridemia; 3) hyperdemia; 4) hyperglycemia; and 5) hypertension (>130/85).

Table D16. Long Term Living Kidney Donation Outcomes – Hypertensive donors versus normotensive donors

		Ir	tervention/Control (n)		Results		
Study, Year, Country	Follow- Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Mortality								
Mjøen 2013 Norway	D:15.1 years ND: 24.9 years	Donors (1,901)	Non-Donors (32,621)		Adjusted	<u>SBP</u> d HR ^f (95% CI) per 1 r 1.00 (1.00-1.01)	nmHg SBP	Low to Moderate
				Cardiovascula	ar Mortality			
Mjøen 2013 Norway	D:15.1 years ND: 24.9 years	Donors (1,901)	Non-Donors (32,621)			<u>SBP</u> d HR ^f (95% CI) per 1 1 1.01 (1.00-1.02)	mmHg SBP	Low to Moderate
				ESR	D			
Mjøen 2013 Norway	D:15.1 years ND: 24.9 years	Donors (1,901)	Non-Donors (32,621)		Adjuste	<u>SBP</u> d HR ^f (95% CI) per 1 ۱ 1.01 (1.00-1.06)	mmHg SBP	Low to Moderate
Renal Function								
Gracida 2003 Mexico	Mean: 6.7	Hypertensive Donors ^a (16)	Normal Donors ^b (422)		GFR (mL/min) ^c mean (SD): 78.1 (NR)	GFR (mL/min) ^c mean (SD): 78.5(NR)		High
Lee 2007 Korea	Median: 5.4 years	Hypertensive Donors ^d (6)	Normotensive Donors (98)		CKD ^e n/N(%): 4/6 (67%) RR	CKD ^e n/N (%) 22/98 (22%) (95% CI) 1.51 to 5.83)	-	Moderate to High
			Donors (104)		Predicti	Moderate to High		

CKD=Chronic Kidney Disease; GFR= Glomerular Filtration Rate; HTN= Hypertension; SD= Standard Deviation ^a-readily controlled with diet or one drug

^b-normal donors were those under 60, non-obese without hypertension, elevated uric acid or 'high cholesterol'.

^c - method of GFR measurement or estimation not reported
 ^{d-} blood pressure >140/90
 ^e - MDRD-GFR of less than 60 mL/min per 1.73 m²

^f - After adjustment for age, gender, year of inclusion, smoking, BMI

Table D17. Long Term Living Kidney Donation Outcomes – Donors Related to Recipients versus Not-Related

		Interv	vention/Control (n)			Results		
Study, Year, Country	Mean Follow- Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
Death or cardiova	Iscular events							
Garg 2008 Canada	6.2 years	Genetically related living donors (1,261)	Unrelated living donors (767)		1.2% HR (95% CI) 0.9 (0.5-3.0)	1.6% Reference		Low to Moderate
Garg 2012 Canada	Median 6.5 years	Genetically related living donors (1,261)	Unrelated living donors (767)		Event rate 1000 person years 1.6 HR (95% CI) 0.8 (0.5-1.4)	Event rate 1000 person years 1.9 HR (95% Cl) 0.9 (0.4-1.8) P for interaction= 0.87		Low to Moderate
ESRD								•
Muzaale 2014 United States	Median 7.6 years	Biological relationship to recipient	Non-biological relationship to recipient		Cum Incidence of ESRD at 15 years per 10,000 (95% CI) 34.1 (26.9- 43.3)	Cum Incidence of ESRD at 15 years per 10,000 (95% CI) 15.1 (8.7-26.3)		Low to Moderate
Renal Function			1			l l		
Lee 2007 Korea	Median: 7.4 years	Donors 1 st degree relatives (28)	Donors non 1 st degree relatives (76)		MDRD eGFR<60 mL/min per 1.736 m ² n/N(%) 5/28(18%) P=0.31	MDRD eGFR<60 mL/min per 1.736 m ² n/N(%) 21/76(28%)		Moderate-High
Hypertension		<u> </u>	II		I	<u> </u>		
Garg 2008 Canada	6.2 years	Genetically related living donors (1,261)	Unrelated living donors (767)		15.9% HR (95% CI) 1.0 (0.7-1.3)	17.3% Reference		Low to Moderate
Psychosocial	•	•			· · ·			-
Gross 2013 United States	17 years	Donor first degree relative of a recipient	Donor not a first degree			degree relative of a rec vith OR(95% CI) 0.54(0		Moderate-High

		Inter	vention/Control	(n)				
Study, Year, Country	Mean Follow- Up (years)	Arm 1	Arm 2	Arm 3	Arm 1	Arm 2	Arm 3	ROB
		(1,768)	relative of a recipient (687)		p<0.0025 c	f physical HRQOL	_ impairment.	
Johnson 1999 United States	NR	Relatives other than first degree (26)	First degree relatives (398)		more likely to sa	er than first degree by they regretted do o 1 st degree relative P=0.06	nating compared	High
Mjoen 2011, Norway	12.6 years	[Donors (71/1377)		Being an unrelated donor is associated with increased risk of having doubt towards donation Adjusted* OR (95% CI) 2.2 (1.2 to 3.9)			Moderate
Lentine 2012 US	4.9 years	Biologically related to recipient (3780)	Spouse/partner of recipient (353)	Not biologically related or spouse to recipient (520)	Depression diagnosis per 100 person- years 4.9 Reference	Depression diagnosis per 100 person- years 5.0 1.16(0.79-1.71)	Depression diagnosis per 100 person- years 5.8 1.30(0.93-1.81)	Moderate

*Adjusted for age, time since donation, gender, medical problems, graft loss in recep.

Study, Year,	Population	Interver	tion/Control (n)	Re	sults	ROB
Country		Arm 1	Arm 2	Arm 1	Arm 2	
Miscarriage						
Ibrahim 2009 United States	Donors with pre or post	Post-donation pregnancy (317)	Pre-donation pregnancy (2,519)	n/N (%) 42/317 (13.2)	n/N (%) 207/2,519 (8.2)	High
pregnancies(987)					95% CI) 18 to 2.20)	
	Donors with pre and post donation	Post-donation pregnancy (173)	Pre-donation pregnancy (204)	n/N (%) 36/173 (21)	n/N (%) 33/204 (16)	
pregnancy (98)				95% CI) 84 to 1.97)		
Stillbirth/Fetal De	eath					
Ibrahim 2009 Donors with pre or United States post	Post-donation pregnancy (317)	Pre-donation pregnancy (2,519)	n/N (%) 1/317 (.32)	n/N (%) 13/2,519 (.52)	High	
	pregnancies(987)			RR (9 0.61 (0.1		
	Donors with pre and post donation		Pre-donation pregnancy (204)	n/N (%) 1/173 (.58)	n/N (%) 2/204 (1.0))	
	pregnancy (98)			RR (95% CI) 05 to 6.45)	
Reisaeter 2009 Norway	Donors with pre- or post- donation	Post-donation pregnancy (106)	Pre-donation pregnancy (620)	n/N (%) 3/106 (2.8)	n/N (%) 7/620 (1.1)	High
,	pregnancy (326)			RR (95% CI) 10 to 1.52)	
Prematurity	•		· · ·	\$, , , , , , , , , , , , , , , , , , ,	
Ibrahim 2009, United States	Donors with pre or post pregnancies	Post-donation pregnancy (317)	Pre-donation pregnancy (2,519)	n/N (%) 20/317 (6)	n/N (%) 95/2,519 (3.7)	High
	(987)			RR (95% CI) 05 to 2.67)	
	Donors with pre and post donation	Post-donation pregnancy (173)	Pre-donation pregnancy (204)	n/N (%) 15/173 (8.7)	n/N (%) 15/204 (7.4)	
	pregnancies (98)			RR (۱ ۱.18 (۵.		
Reisaeter 2009 Norway	Donors with pre- or post- donation pregnancy (326)	Post-donation pregnancy (106)	Pre-donation pregnancy (620)	<22 weeks n/N (%) 1/106 (1)	<22 weeks n/N (%) 2/620 (0.3)	High
				RR (9 2.92 (0.2		
				<37 weeks n/N (%)	<37 weeks n/N (%)	

Table D18. Long Term Living Kidney Donation Outcomes – Post Donation Pregnancy-related Outcomes

Study, Year,	Population	Interven	tion/Control (n)	Res	sults	ROB	
Country		Arm 1	Arm 2	Arm 1	Arm 2		
				10/106 (9.8)	44/620 (7.5)		
				BR (9	95% ci)	—	
					9 to 2.56)		
Gestational Hype	ertension						
Ibrahim 2009	Donors with pre or	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%):	High	
Jnited States	post	pregnancy (317)	(2,519)	22/317(6.9)	16/2,519 (0.6)		
	pregnancies(987)				5% CI) 0 to 20.58)		
	Donors with pre	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%)		
	and post donation	pregnancy (173)	(204)	6/173 (3.5)	1/204 (0.5)		
	pregnancies (98)			RR (9			
D	D	Destates			6 to 58.20)		
Reisaeter 2009 Norway	Donors with pre- or post- donation	Post-donation pregnancy (106)	Pre-donation pregnancy (620)	n/N (%) 3/106 (2.8)	n/N (%) 11/620 (1.8)	High	
Norway	pregnancy (326)	pregnancy (100)	(020)		5% CI)		
	F - 3 5 (7				5 to 5.62)		
Gestational Diab	etes						
Ibrahim 2009	Donors with pre or	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%)	High	
United States			(2,519)	12/317 (3.8)	19/2,519 (0.8)		
	(987)				5% CI) 6 to 10.24)		
	Donors with pre Post-d	Post-donation	Pre-donation pregnancy	n/N (%):	n/N (%)		
	and post donation	pregnancy (173)	(204)	1/173 (0.6%)	1/204 (0.5)		
	pregnancies (98)				5% CI)		
Dragolomnoio				1.18 (0.0	7 to 18.71)		
Preeclampsia	Demonstrativity and an	Dest densitien			··· (N1 (0/)	Lint	
Ibrahim 2009 United States	Donors with pre or post pregnancies	Post-donation pregnancy (317)	Pre-donation pregnancy (2519)	n/N (%) 21/317(6.6)	n/N (%) 22/2519 (0.9)	High	
	(987)	prognancy (orr)	(2010)		5% CI)		
	. ,				2 to 13.63)		
	Donors with pre	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%)		
	and post donation pregnancies (98)	pregnancy (173)	(204)	6/173 (3.5)	1/204 (0.5) 5% CI)		
	pregnancies (90)		7.07 (0.8				
Reisaeter 2009	Donors with pre- or	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%):	High	
Norway	post- donation	pregnancy (106)	(620)	6/106 (5.7) ^a	16/620 (2.6) ^a	Ŭ	
	pregnancy (326)			p=0.026	1 les e diverte ell		
				(95% Cl) 2.19 (0.8			
Proteinuria		1		2.19 (0.0		I	

Study, Year,	Population	Interven	tion/Control (n)	Res	sults	ROB
Country		Arm 1	Arm 2	Arm 1	Arm 2	
Ibrahim 2009	Donors with pre or	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%):	High
United States	post pregnancies	pregnancy (317)	(2519)	13/317 (4.1)	25/2519 (1.0)	
	(987)			RR (95	5% CI)	
				4.13 (2.13	3 to 7.99)	
	Donors with pre	Post-donation	Pre-donation pregnancy	n/N (%)	n/N (%):	
	and post donation	pregnancy (173)	(204)	8/173 (4.6)	4/204 (1.5)	
	pregnancies (98)			RR (95	5% CI)	
				2.36 (0.72	2 to 7.70)	
Low birth weight						
Reisaeter 2009	Donors with pre- or	Post-donation	Pre-donation pregnancy	<500g	<500g	High
Norway	post- donation	pregnancy (106)	(620)	n/N (%)	n/N (%)	
	pregnancy (326)			1/106 (0.9)	3/620 (0.5)	
				RR (95	5% CI)	
				1.95 (0.20) to 18.57)	
				500-2500g	500-2500g	
				n/N (%):	n/N (%)	
				8/106 (7.5)	34/620 (5.5)	
				RR (95	5% CI)	
				1.38 (0.6	5 to 2.89)	

CI= Confidence Interval; RR=Risk Ratio ^a-adjusted for maternal age, birth order, and year of birth