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**Impact of anti-hypertensive therapy in the sexual health of men and women: an analysis
from the SPRINT trial**

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Abstract

Background: Pharmacologic anti-hypertensive (HT) treatment reduces cardiovascular risk. However, many patients are non-adherent due to perceived or real concern about sexual-related side-effects.

Objectives: In a subset of the SPRINT (a randomized trial of intensive versus standard blood-pressure control) trial, we sought to investigate the impact of anti-HT treatment on sexual activities of men and women over time, and whether this impact varied with a more or less intensive anti-HT therapy.

Methods: Random-effects models for panel/longitudinal data.

Results: Among the 1268 men and 613 women included in this sub-study, 862 (68%) men and 178 (29%) women declared to be engaged in sexual activity of any kind. Compared with women and men not engaged in sexual activity, those engaged were younger (64 vs. 69yr for women and 65 vs. 75yr for men). Women had an overall low satisfaction with their sexual life but their sexual health was not affected by anti-HT therapy over time nor modified by an intensive treatment. Men's erections were slightly deteriorated over time (-0.1 to -0.2 points on a scale of 1 (worse) to 5 (best); $p < 0.05$), but were not aggravated by intensive anti-HT therapy ($p > 0.05$ for all).

Conclusions: Self-declared women's sexual health was not affected by an intensive anti-HT therapy. Men, reported a slight deterioration in the quality of their erections, irrespective of standard or intensive therapy. These findings may help reassuring patients about the sexual safety of intensive anti-HT therapy, therefore, potentially improving adherence to intensive therapy strategy.

Key-words: sexual health; hypertension; anti-hypertensive treatment.

Condensed abstract

In SPRINT (a randomized trial of intensive versus standard blood-pressure control), women had an overall low satisfaction with their sexual life but their sexual health was not affected by anti-hypertensive therapy over time nor modified by an intensive treatment. On the other hand, men's erections were slightly deteriorated over time, but were not furtherly aggravated by an intensive anti-hypertensive therapy. These findings have clinical implications as they may help reassuring patients about the sexual safety of intensive anti-hypertensive therapy, therefore, potentially improving treatment adherence.

Introduction

Arterial hypertension (HT) affects more than 25% of the adult population worldwide regardless of sex¹. Pharmacologic treatment of HT is key to reduce the cardiovascular risk associated with uncontrolled blood pressure (BP). However, many patients are not willing to take their anti-HT drugs due to concern about side-effects². Long-term treatment non-adherence is one of the major causes for uncontrolled BP and, consequently, increasing the morbidity and mortality risk (e.g., stroke, myocardial infarction, renal insufficiency, and death)³.

In clinical practice, sexual side-effects are frequently evoked as an important reason for not taking the anti-HT treatment. Particularly younger, sexually active men often complain of erectile dysfunction; whereas women report less sexual side effects⁴. Moreover, sexual side-effects may be under-recognized, and under-reported because of difficulties in approaching the problem both by the patient and the clinician⁵. Hypertension itself can also cause sexual dysfunction in the long-term which might be seen by patients as a far away, unlikely future; however, the treatment side-effects occur and are felt in the present, leading to poor treatment adherence and discontinuation – a “so-called” present-self vs. future-self conflict⁶.

In a subset of the SPRINT (a randomized trial of intensive versus standard blood-pressure control) trial⁷, patients were questioned about their sexual activities at baseline and throughout the follow-up using validated sexual health questionnaires. This study sought to investigate the impact of anti-HT treatment on self-declared sexual activities over time, and whether this impact could vary with a more or less intensive anti-HT therapy in men and women.

Methods

Trial oversight

SPRINT was sponsored by the NHLBI, with co-sponsorship by the National Institute of Diabetes and Digestive and Kidney Diseases, the National Institute of Neurological Disorders and Stroke, and the National Institute on Aging. The rationale and protocol for the trial have been previously published and are publicly available^{7,8}. All participants provided written informed consent to participate in the study. The authors of this manuscript had authorization from the National Institutes of Health (NIH)/National Heart, Lung, and Blood Institute (NHLBI) to perform the present analysis under the coordination of the Biologic Specimen and Data Repository Information Coordinating Center (BioLINCC; request #7509).

To be enrolled in the SPRINT trial, participants were required to meet all the following criteria: an age of at least 50 years, a systolic blood pressure, SBP of 130 to 180 mm Hg, and an increased risk of cardiovascular events (defined by one or more of the following: clinical or subclinical cardiovascular disease other than stroke; chronic kidney disease with an estimated glomerular filtration rate [eGFR] of 20 to 60 ml/min/1.73m² calculated with the use of the four-variable Modification of Diet in Renal Disease Equation⁹; a 10-year risk of cardiovascular disease \geq 15% on the basis of the Framingham risk score; or an age \geq 75 years). Patients with diabetes mellitus or prior stroke were excluded. Participants and study personnel were aware of the study-group assignments, but outcome adjudicators were not. The protocol encouraged, but did not mandate, the use of drug classes with the strongest evidence for reduction in cardiovascular outcomes¹⁰. Blood pressure was considered as the mean of three BP measurements at an office visit while the patient was seated after 5 minutes of quiet rest; the measurements were made with the use

of an automated measurement system (Model 907, Omron Healthcare). The preferred method was the automated device as it offers reduced potential for observer biases and decreased demand on staff in terms of training and effort in data collection. Medical records and electrocardiograms were obtained for documentation of events. Whenever clinical site staff became aware of a death, a standard protocol was used to obtain information on the event. A total of 9361 participants were randomized. The trial was stopped earlier than expected after analyses of the primary outcome exceeded the monitoring boundaries at two consecutive time-points. The median follow-up time was 3.26 years. The primary outcome was a composite of myocardial infarction (MI), other acute coronary syndromes (ACS), stroke, heart failure (HF), or cardiovascular death. Compared with a standard systolic blood pressure target of <140 mmHg and intensive strategy with a <120 mmHg, reduced the primary outcome (6.8% vs 5.2%; 1.65% per year vs. 2.19% per year; hazard ratio [HR] with intensive treatment, 0.75; 95% confidence interval [CI], 0.64 to 0.89; $p < 0.001$)⁷.

Sexual health questionnaires

Sexual health questionnaires were obtained in a subset of the SPRINT trial population (n=1881; 20% of the overall trial population), where 1268 were men and 613 women. The characteristics of these patients were similar to those of the overall trial population.

Supplemental Table 1. Throughout the trial, the most frequently prescribed anti-HT drugs were ACEi/ARBs and thiazides, followed by calcium-channel blockers and beta-blockers.

Supplemental Table 2.

For those engaged in sexual activities of any kind and who agreed to reply to the questionnaires (178 women and 862 men), the sexual health questions included an overall sexual life satisfaction question for both sexes, ranging from 1 (very dissatisfied) to 5 (very satisfied). Sexual health questionnaire for women was performed using a modified version

of the Female Sexual Function Inventory (FSFI)¹¹, and included questions about their sexual desire or interest, sexual arousal, vaginal lubrication during sexual activity, orgasm frequency, satisfaction with sexual relationship and emotional closeness with partner. These questions were classified from 1 (almost never satisfied or very dissatisfied) to 5 (almost always or very satisfied). We have also dichotomized these variables in half of the times or more (≥ 3 points) and less than half of the times (< 3 points).

For men the questionnaires were performed using the 5-item version of International Index of Erectile Function (IIEF-5) questionnaire¹², and included questions about confidence to get and keep an erection, if the erections were hard enough for penetration, and whether the erections could be maintained after penetration. These questions were also classified from 1 (almost never or very dissatisfied) to 5 (almost always or very satisfied) and we have also dichotomized these variables in half of the times or more (≥ 3 points) and less than half of the times (< 3 points).

These questionnaires are detailed in the ***Supplemental Material***.

Statistical analysis

In descriptive analyses, continuous variables are expressed as mean \pm standard deviation (SD) or median (percentile₂₅₋₇₅) based on their histogram distribution. Categorical variables are expressed as frequencies and proportions (%). Population description and comparison of men vs. women was performed using independent samples t-test for normally distributed continuous variables, Mann-Whitney test for skewed variables, and chi-square test for categorical variables. As the aim of our study was to assess the impact on anti-HT therapy on the sexual function of men and women throughout the follow-up, the evolution of the sexual scores over time, was performed using linear (for continuous) and logistic (for binary) random-effects models for panel data i.e., data with multiple observations per patient over

time (study visits: 0, 6, 12, 24, 36, and 48 months). The sexual scores were set as dependent variables and visit time as independent variable. To assess the effect of treatment (intensive vs. standard) a treatment-by-time interaction was included in the model. The individual anti-HT drugs used during the follow-up were retrieved from “free text” forms. To assess the impact of comorbidities on the sexual function scores, age, race, systolic blood pressure, body mass index, smoking history, number of anti-hypertensive agents, estimated glomerular filtration rate, presence of clinical or subclinical cardiovascular disease, total cholesterol levels, glucose levels, and urinary albumin-to-creatinine ratio were added to the panel data models in a multivariable fashion, retaining only the variables significantly associated with each individual sexual score in a stepwise backward fashion. Time-updated Cox models were used to study the association of erectile function and the study primary outcome adjusted on age, body mass index, chronic kidney disease and history of clinical or subclinical cardiovascular disease. All analyses were performed with the Stata® software (StataCorp. 2019. Stata Statistical Software: Release 16. College Station, TX: StataCorp LLC).

Results

Baseline patients` characteristics by sex

Among the 1268 men and 613 women included in this substudy, 862 (68%) men and 178 (29%) women declared to be engaged in sexual activity of any kind. Compared with women declaring not to be engaged in sexual activity of any kind, those who declared to be engaged in sexual activity were younger (64 vs. 69yr), more often current smokers, and with better renal function. **Table 1.** Compared with men not engaged in sexual activity of any kind, those who declared to be engaged in sexual activity were younger (65 vs. 71yr), more often black, with slightly higher body mass index, and with better renal function. **Table 1.**

Comparing men and women with sexual activity of any kind, women were slightly younger, more often black, more often current smokers, with higher HDL-cholesterol and less likely to be on a statin. **Table 1.**

Baseline sexual health

The main description of the questionnaires by sex is depicted in the **Table 2.** Women declared having an overall low satisfaction with their sexual life. Less than 50% of women declared to feel sexual desire or interest on at least half of the occasions and 30% said that they would almost never/never feel sexual desire; however, most declared to feel sexual arousal, become lubricated, and reach orgasm in at least half of the occasions they had sexual intercourse, most women were also satisfied with their relationship and emotional closeness during sexual activity. **Table 2.**

Most men said they felt at least moderately confident that they could get and keep an erection that the erections would be hard for penetration and maintained after penetration in most occasions. **Table 2.**

Sexual health over time by sex and treatment allocation

Women's sexual satisfaction, desire, arousal, lubrication, orgasm, and emotional closeness did not significantly change over time, neither did they differ by treatment allocation (i.e., they were not aggravated by intensive therapy). **Table 3 & Figure 1.**

Men's overall sexual satisfaction was maintained over time and did not differ by treatment group. However, they declared that their erections (particularly hard erections and their maintenance during intercourse) were slightly deteriorated over time (-0.1 to -0.2 points; $p < 0.05$), but they were not aggravated by intensive anti-HT therapy ($p > 0.05$ for all).

Table 4 & Figure 2. All therapeutic regimens could have had a negative impact on the

quality of the erections, but those without thiazides and/or beta blockers could have been less impactful. **Supplemental Table 3.**

Association with clinical features and comorbid conditions

In men, older age (>65yr) and a body mass index above 30 Kg/m², were associated with lower overall sexual satisfaction and lower quality of erections. Patients with clinical or subclinical cardiovascular disease and those with impaired renal function had more difficulty in maintaining erections. **Supplemental Table 4.** Women taking a higher number of anti-HT drugs were less satisfied with their sexual life, and older women (>65yr) had less sexual desire and became less often lubricated during sexual activity. **Supplemental Table 4.**

Outcome associations

Among men, 117 primary outcome events occurred. In adjusted models, the sexual/erectile function was not significantly associated with the study primary outcome. **Supplemental Table 5.** Among women, 31 primary outcome events occurred. Feeling sexual desire or interest more often was independently associated with a lower rate of the study primary outcome (HR [95%CI] =0.62 [0.41-0.94], p =0.025), but the other sex-related questions were not. **Supplemental Table 5.**

Discussion

This study shows that in a population with arterial hypertension and a moderate-to-high cardiovascular risk (but without diabetes or prior stroke), most women declared that they were not engaged in any sexual activity while most men declared they were. Women's sexual desire was low but once they were engaged in sexual activities their sexual satisfaction was at least moderate in most occasions and was not deteriorated with intensive anti-HT therapy over time. Most men were confident that they could get and keep

an erection, but the maintenance of hard erections was slightly diminished throughout the trial, albeit not aggravated by intensive therapy, suggesting that anti-HT treatment may impair the quality of erections but that is not dependent on the intensity of the anti-HT treatment.

Sexual dysfunction might be more prevalent in women than in men and is associated with poor physical and emotional health as well as poor overall well-being¹³. Although the sexual problems tend to increase with age, these seem to affect men in particular, still lubrication problems may also be aggravated with women's age¹⁴. This study included mostly elderly people (mean age 67yr for women and 68yr for men) with moderate-high cardiovascular risk and hypertension, which is a population with a particularly high-risk for experiencing sexual dysfunction¹⁴. While most women declared that they were not engaged in any sexual activity, most men declared they were. The lack of sexual interest in older women has been reported, while most men could have stated to have more sexual interest due to fear of judgements about their "masculinity"^{14, 15}. Interestingly, women that stated to have a sexual activity of any kind reported to have an overall (at least) moderate sexual satisfaction. In a report using the baseline data of SPRINT, blood pressure was not associated with sexual activity in women, which was more related to psychosocial factors¹⁶, and, at baseline, no single class of anti-HT medication was associated with sexual dysfunction in women¹⁷. Furthermore, our study shows that women did not experience any deterioration of their sexual health during the follow-up nor their sexual health was influenced by the allocation to an intensive anti-HT therapy regimen. These findings have clinical relevance, because they suggest that regardless of the baseline sexual health in women (that may be related to many other factors rather than blood pressure)

this is unlikely to be deteriorated with an intensive treatment and should not preclude clinicians to try a systolic blood pressure goal of 120 mmHg.

Men were overall confident of their sexual performance, but many reported to have erectile problems at baseline and to have difficulties in maintaining an erection during sexual intercourse. These findings are compatible with other reports, stating that erectile dysfunction is common in men in their 60s and even more in men aged 70 and older, affecting more than 50% of men in these age ranges^{18, 19}. In patients with hypertension, erectile dysfunction has been independently associated with adverse cardiovascular outcomes²⁰. In our report, the associations between erectile dysfunction and the study primary outcome did not reach statistical significance; probably due to the small size and few events in this subsample. In a report from SPRINT, baseline blood pressure was not associated with sexual activity in men either²¹. Our study suggests that erectile function (hard and sustained erections, in particular) may be aggravated with anti-hypertensive treatment over time; however, erectile dysfunction was not furtherly deteriorated with intensive anti-HF treatment for a SBP goal of 120 mmHg, which suggests that erectile dysfunction is not dependent on the intensity of the anti-HT therapy. In a report from SPRINT, the erectile dysfunction induced by a more intensive BP lowering strategy was found to be more marked in non-Hispanic blacks compared with non-Hispanic whites, differences that disappeared in the adjusted models²². In our report, race was not independently associated with any sexual/erection question in the fully adjusted models.

In SPRINT (overall and in this subpopulation) the most commonly used anti-HT drug was angiotensin converting enzyme inhibitors/angiotensin receptor blockers (ACEi/ARBs) (in over 70% of the participant-episodes), followed by thiazide-type diuretics (over 50%), calcium channel blockers (CCBs) (in almost 50%), and beta-blockers (up to 40%). The

combination of ACEi/ARBs plus thiazides was used in 53% of the observations in men. Other anti-HT agents such as mineralocorticoid receptor antagonists (MRAs) and central acting drugs, were used in a much lower frequency (<10%). The use of ACEi/ARBs and CCBs has been reported as not being associated with erectile dysfunction, with many reports showing a “neutral” effect of these agents in erectile function²³⁻²⁵; ARBs in particular by blocking the vasoconstrictive action of angiotensin II may positively affect erectile function and are thus regarded as a first-line treatment in hypertensive patients with erectile dysfunction²⁶⁻²⁹. On the other hand, thiazides, beta-blockers (with the possible exception of nebivolol)³⁰, MRAs, and central-acting anti-HT agents may impair erectile function³¹⁻³⁵. Unfortunately, for an adequate BP control one often requires anti-HT drug combinations, which often include a diuretic to increase sodium excretion³⁶. In SPRINT this was no exception with more than half of men having a thiazide diuretic prescribed in association with other drugs, most frequently an ACEi/ARB. Nonetheless, consistently with prior studies, our observational data suggest that anti-HT regimens without beta-blockers or thiazides may be less impactful on erectile function.

Limitations

The impact of each individual drug on erectile function is hard to ascertain, because most patients take several medications that change over time and the individual effects of each drug cannot be assessed in a reliable fashion. However, the observation that an intensive anti-HT therapy does not furtherly deteriorate erectile function is reassuring and may help clinicians guiding their treatment strategies i.e., intensifying anti-HT treatment can improve outcomes and does not seem to further impact erectile function. The use of other medications (e.g., over the counter phosphodiesterase-5 inhibitors) could not be accounted for because was not systematically recorded in the dataset; however, it is expected that the

use of other (non anti-HT) medications that could impact sexual function was used similarly (i.e., at random) between the intensive and standard anti-HT therapy groups, and therefore does not impact our findings comparing the effect of intensive vs. standard anti-HT therapy on sexual function. The small number of events in this sexual questionnaire subsample (furtherly subdivided in men and women) likely lacked enough power to study the associations between sexual function and outcomes in a more robust fashion.

Conclusion

In SPRINT, women`s sexual health was maintained over time and was not affected by intensive anti-HT therapy. Men, declared having a slight deterioration in the quality of their erections over time, without further aggravation by intensive anti-HT therapy. These findings may help reassuring patients about the sexual safety of intensive anti-HT therapy, therefore, potentially improving adherence to intensive therapy strategy.

Clinical perspectives

Women had an overall low satisfaction with their sexual life but their sexual health was not affected by anti-hypertensive therapy over time nor modified by an intensive treatment.

Men`s erections were slightly deteriorated over time but were not aggravated by an intensive anti-hypertensive therapy.

These findings may help reassuring patients about the sexual safety of intensive anti-hypertensive therapy and improve treatment adherence.

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Disclosures

The content of this manuscript is solely the responsibility of the authors and does not necessarily represent the official views of the National Institutes of Health (NIH), the Department of Veterans Affairs, or the U.S. Government.

Bibliography

1. Kearney, P. M.; Whelton, M.; Reynolds, K.; Muntner, P.; Whelton, P. K.; He, J., Global burden of hypertension: analysis of worldwide data. *Lancet* **2005**, *365* (9455), 217-23.
2. Burnier, M., Medication adherence and persistence as the cornerstone of effective antihypertensive therapy. *Am J Hypertens* **2006**, *19* (11), 1190-6.
3. Durand, H.; Hayes, P.; Morrissey, E. C.; Newell, J.; Casey, M.; Murphy, A. W.; Molloy, G. J., Medication adherence among patients with apparent treatment-resistant hypertension: systematic review and meta-analysis. *J Hypertens* **2017**, *35* (12), 2346-2357.
4. Prisant, L. M.; Carr, A. A.; Bottini, P. B.; Solursh, D. S.; Solursh, L. P., Sexual dysfunction with antihypertensive drugs. *Arch Intern Med* **1994**, *154* (7), 730-6.
5. Viigimaa, M.; Vlachopoulos, C.; Lazaridis, A.; Doumas, M., Management of erectile dysfunction in hypertension: Tips and tricks. *World J Cardiol* **2014**, *6* (9), 908-15.
6. Hershfield, H. E., Future self-continuity: how conceptions of the future self transform intertemporal choice. *Ann N Y Acad Sci* **2011**, *1235*, 30-43.
7. Wright, J. T., Jr.; Williamson, J. D.; Whelton, P. K.; Snyder, J. K.; Sink, K. M.; Rocco, M. V.; Reboussin, D. M.; Rahman, M.; Oparil, S.; Lewis, C. E.; Kimmel, P. L.; Johnson, K. C.; Goff, D. C., Jr.; Fine, L. J.; Cutler, J. A.; Cushman, W. C.; Cheung, A. K.; Ambrosius, W. T., A Randomized Trial of Intensive versus Standard Blood-Pressure Control. *N Engl J Med* **2015**, *373* (22), 2103-16.
8. Ambrosius, W. T.; Sink, K. M.; Foy, C. G.; Berlowitz, D. R.; Cheung, A. K.; Cushman, W. C.; Fine, L. J.; Goff, D. C., Jr.; Johnson, K. C.; Killeen, A. A.; Lewis, C. E.; Oparil, S.; Reboussin, D. M.; Rocco, M. V.; Snyder, J. K.; Williamson, J. D.; Wright, J. T., Jr.; Whelton, P. K., The design and rationale of a multicenter clinical trial comparing two strategies for control of systolic blood pressure: the Systolic Blood Pressure Intervention Trial (SPRINT). *Clin Trials* **2014**, *11* (5), 532-46.
9. Levey, A. S.; Bosch, J. P.; Lewis, J. B.; Greene, T.; Rogers, N.; Roth, D., A more accurate method to estimate glomerular filtration rate from serum creatinine: a new prediction equation. Modification of Diet in Renal Disease Study Group. In *Ann Intern Med*, United States, 1999; Vol. 130, pp 461-70.
10. Chobanian, A. V.; Bakris, G. L.; Black, H. R.; Cushman, W. C.; Green, L. A.; Izzo, J. L., Jr.; Jones, D. W.; Materson, B. J.; Oparil, S.; Wright, J. T., Jr.; Roccella, E. J., The Seventh Report of the Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure: the JNC 7 report. *Jama* **2003**, *289* (19), 2560-72.

11. Rosen, R.; Brown, C.; Heiman, J.; Leiblum, S.; Meston, C.; Shabsigh, R.; Ferguson, D.; D'Agostino, R., Jr., The Female Sexual Function Index (FSFI): a multidimensional self-report instrument for the assessment of female sexual function. *J Sex Marital Ther* **2000**, *26* (2), 191-208.
12. Rosen, R. C.; Cappelleri, J. C.; Smith, M. D.; Lipsky, J.; Pena, B. M., Development and evaluation of an abridged, 5-item version of the International Index of Erectile Function (IIEF-5) as a diagnostic tool for erectile dysfunction. *Int J Impot Res* **1999**, *11* (6), 319-26.
13. Laumann, E. O.; Paik, A.; Rosen, R. C., Sexual dysfunction in the United States: prevalence and predictors. *Jama* **1999**, *281* (6), 537-44.
14. Laumann, E. O.; Nicolosi, A.; Glasser, D. B.; Paik, A.; Gingell, C.; Moreira, E.; Wang, T., Sexual problems among women and men aged 40-80 y: prevalence and correlates identified in the Global Study of Sexual Attitudes and Behaviors. *Int J Impot Res* **2005**, *17* (1), 39-57.
15. Moreira, E. D., Jr.; Kim, S. C.; Glasser, D.; Gingell, C., Sexual activity, prevalence of sexual problems, and associated help-seeking patterns in men and women aged 40-80 years in Korea: data from the Global Study of Sexual Attitudes and Behaviors (GSSAB). *J Sex Med* **2006**, *3* (2), 201-11.
16. Foy, C. G.; Newman, J. C.; Berlowitz, D. R.; Russell, L. P.; Kimmel, P. L.; Wadley, V. G.; Thomas, H. N.; Lerner, A. J.; Riley, W. T., Blood Pressure, Sexual Activity, and Dysfunction in Women With Hypertension: Baseline Findings From the Systolic Blood Pressure Intervention Trial (SPRINT). *J Sex Med* **2016**, *13* (9), 1333-1346.
17. Thomas, H. N.; Evans, G. W.; Berlowitz, D. R.; Chertow, G. M.; Conroy, M. B.; Foy, C. G.; Glasser, S. P.; Lewis, C. E.; Riley, W. T.; Russell, L.; Williams, O.; Hess, R., Antihypertensive medications and sexual function in women: baseline data from the SBP intervention trial (SPRINT). *J Hypertens* **2016**, *34* (6), 1224-31.
18. Selvin, E.; Burnett, A. L.; Platz, E. A., Prevalence and risk factors for erectile dysfunction in the US. *Am J Med* **2007**, *120* (2), 151-7.
19. Ayta, I. A.; McKinlay, J. B.; Krane, R. J., The likely worldwide increase in erectile dysfunction between 1995 and 2025 and some possible policy consequences. *BJU Int* **1999**, *84* (1), 50-6.
20. Bohm, M.; Baumhakel, M.; Teo, K.; Sleight, P.; Probstfield, J.; Gao, P.; Mann, J. F.; Diaz, R.; Dagenais, G. R.; Jennings, G. L.; Liu, L.; Jansky, P.; Yusuf, S., Erectile dysfunction predicts cardiovascular events in high-risk patients receiving telmisartan, ramipril, or both: The ONgoing Telmisartan Alone and in combination with Ramipril Global Endpoint Trial/Telmisartan Randomized Assessment Study in ACE iNtolerant subjects with cardiovascular Disease (ONTARGET/TRANSCEND) Trials. *Circulation* **2010**, *121* (12), 1439-46.
21. Foy, C. G.; Newman, J. C.; Berlowitz, D. R.; Russell, L. P.; Kimmel, P. L.; Wadley, V. G.; Thomas, H. N.; Lerner, A. J.; Riley, W. T., Blood Pressure, Sexual Activity, and Erectile Function in Hypertensive Men: Baseline Findings from the Systolic Blood Pressure Intervention Trial (SPRINT). *J Sex Med* **2019**, *16* (2), 235-247.
22. Foy, C. G.; Newman, J. C.; Russell, G. B.; Berlowitz, D. R.; Bates, J. T.; Burgner, A. M.; Carson, T. Y.; Chertow, G. M.; Dumas, M. N.; Hughes, R. Y.; Kostis, J. B.; Buren, P. V.; Wadley, V. G., Effect of Intensive vs Standard Blood Pressure Treatment Upon Erectile Function in Hypertensive Men: Findings From the Systolic Blood Pressure Intervention Trial. *J Sex Med* **2020**, *17* (2), 238-248.
23. Suzuki, H.; Tominaga, T.; Kumagai, H.; Saruta, T., Effects of first-line antihypertensive agents on sexual function and sex hormones. *J Hypertens Suppl* **1988**, *6* (4), S649-51.
24. Omvik, P.; Thaulow, E.; Herland, O. B.; Eide, I.; Midha, R.; Turner, R. R., Double-blind, parallel, comparative study on quality of life during treatment with amlodipine or enalapril in mild or moderate hypertensive patients: a multicentre study. *J Hypertens* **1993**, *11* (1), 103-13.
25. Kroner, B. A.; Mulligan, T.; Briggs, G. C., Effect of frequently prescribed cardiovascular medications on sexual function: a pilot study. *Ann Pharmacother* **1993**, *27* (11), 1329-32.
26. Llisterri, J. L.; Lozano Vidal, J. V.; Aznar Vicente, J.; Argaya Roca, M.; Pol Bravo, C.; Sanchez Zamorano, M. A.; Ferrario, C. M., Sexual dysfunction in hypertensive patients treated with losartan. *Am J Med Sci* **2001**, *321* (5), 336-41.

27. Dusing, R., Effect of the angiotensin II antagonist valsartan on sexual function in hypertensive men. *Blood Press Suppl* **2003**, *2*, 29-34.
28. Della Chiesa, A.; Pfiffner, D.; Meier, B.; Hess, O. M., Sexual activity in hypertensive men. *J Hum Hypertens* **2003**, *17* (8), 515-21.
29. Baumhake, M.; Custodis, F.; Schlimmer, N.; Laufs, U.; Bohm, M., Improvement of endothelial function of the corpus cavernosum in apolipoprotein E knockout mice treated with irbesartan. *J Pharmacol Exp Ther* **2008**, *327* (3), 692-8.
30. Baumhake, M.; Schlimmer, N.; Buyukafsar, K.; Arikian, O.; Bohm, M., Nebivolol, but not metoprolol, improves endothelial function of the corpus cavernosum in apolipoprotein e-knockout mice. *J Pharmacol Exp Ther* **2008**, *325* (3), 818-23.
31. Fogari, R.; Zoppi, A.; Poletti, L.; Marasi, G.; Mugellini, A.; Corradi, L., Sexual activity in hypertensive men treated with valsartan or carvedilol: a crossover study. *Am J Hypertens* **2001**, *14* (1), 27-31.
32. Brixius, K.; Middeke, M.; Lichtenthal, A.; Jahn, E.; Schwinger, R. H., Nitric oxide, erectile dysfunction and beta-blocker treatment (MR NOED study): benefit of nebivolol versus metoprolol in hypertensive men. *Clin Exp Pharmacol Physiol* **2007**, *34* (4), 327-31.
33. Williams, G. H.; Croog, S. H.; Levine, S.; Testa, M. A.; Sudilovsky, A., Impact of antihypertensive therapy on quality of life: effect of hydrochlorothiazide. *J Hypertens Suppl* **1987**, *5* (1), S29-35.
34. Grimm, R. H., Jr.; Grandits, G. A.; Prineas, R. J.; McDonald, R. H.; Lewis, C. E.; Flack, J. M.; Yunis, C.; Svendsen, K.; Liebson, P. R.; Elmer, P. J., Long-term effects on sexual function of five antihypertensive drugs and nutritional hygienic treatment in hypertensive men and women. Treatment of Mild Hypertension Study (TOMHS). *Hypertension* **1997**, *29* (1 Pt 1), 8-14.
35. Baumhake, M.; Schlimmer, N.; Kratz, M.; Hackett, G.; Jackson, G.; Bohm, M., Cardiovascular risk, drugs and erectile function--a systematic analysis. *Int J Clin Pract* **2011**, *65* (3), 289-98.
36. Burnier, M.; Bakris, G.; Williams, B., Redefining diuretics use in hypertension: why select a thiazide-like diuretic? *J Hypertens* **2019**, *37* (8), 1574-1586.

Figure 1. Women: change in lubrication and satisfaction during sexual intercourse over time and by treatment (intensive vs. standard)

A) Change over time

B) Change by treatment

Legend: No significant differences observed (all $p > 0.05$).

Figure 2. Men: change in the quality of erections during sexual intercourse over time and by treatment (intensive vs. standard)

A) Change over time

B) Change with treatment

Caption: Slight decrease in the quality of erections over time ($p < 0.05$ for all time-points, except 48 months where the estimates are imprecise due to low number of patients) but not by intensive treatment (vs. standard) ($p > 0.05$ for all comparisons).