# Risk-benefit analysis attests to the importance of neonatal male circumcision to public health and individual well-being 

## Introduction

Evidence-based policy statements in the USA ${ }^{1,2}$ and Australia ${ }^{3}$ support circumcision, which is best done in the neonatal period. ${ }^{4}$ A thorough risk-benefit analysis is, however, required.

## Aim

To determine the overall medical benefit and risk of neonatal male circumcision.

## Methods

PubMed searches by 'circumcision’ and relevant keywords.

## Results

Table 1 shows increase in risk of various medical conditions conferred by lack of circumcision. Together, benefits of circumcision exceeded risks by 100 to $1 .{ }^{5}$ Over their lifetime 1 in 2 uncircumcised males will suffer a condition (some fatal) caused by retention of the foreskin. ${ }^{5}$ A meta-analysis showed substantial protection against urinary tract infections, finding these affected 1 in 3 uncircumcised males over their lifetime. ${ }^{6}$ The degree of protection was 10 -fold in infancy when risk of kidney damage is greatest. Circumcision protects against phimosis, paraphimosis, balanitis, sexually transmitted infections (e.g., oncogenic HPV [an epidemic], HSV-2, Trichomonas, mycoplasma, chancroid, syphilis and HIV), thrush, inferior hygiene, penile cancer (that affects 1 in 1,000 uncircumcised males over the lifetime) and prostate cancer. ${ }^{1,2,5}$
Circumcision protects the female partner(s) from cervical cancer, bacterial vaginosis and STIs ${ }^{1-5}$ (Table 1).
Risk of adverse events is $0.5 \%$, virtually all being minor and immediately and easily treatable with complete resolution ${ }^{1,2,5,7}$ (Table 1). Neonatal male circumcision is highly cost-effective. ${ }^{8}$ There are no long-term adverse effects on sexual function, sensitivity or pleasure; 9 if anything sex is better. Legal and ethical considerations also support neonatal male circumcision. ${ }^{10}$

## Conclusions

- The strong net benefit and low risk of neonatal male circumcision makes it comparable to childhood vaccination.
- Circumcision of baby boys should be offered routinely.
- Access should be facilitated and affordability assured by state and federal governments in the USA, UK, Australia and all other countries worldwide as an evidence-based public health imperative.


## References

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TABLE 1. Comprehensive risk-benefit analysis of neonatal male circumcision (ref ${ }^{5}$ )

Risks from not circumcising

| Condition | Increase in risk (95\% CI) | Rating of evidence | Percentage of uncircumcised affected |
| :---: | :---: | :---: | :---: |
| Urinary tract infections: age 0-1 year | $9.9(7.5-13)^{\text {Ref } 6}$ | 1++ | 1.3 |
| Urinary tract infections: age 1-16 years | 6.6 (3.3-13) ${ }^{\text {Ref6 }}$ | 1++ | 2.7 |
| Urinary tract infections: age $>16$ years | 3.4 (0.92-50) ${ }^{\text {Re f6 } 6}$ | 1+ | 28 |
| Urinary tract infections: lifetime | 3.6 (1.8-5.7) ${ }^{\text {Ref } 6}$ | 1+ | 32 |
| Pyelonephritis (infants) | $10^{\text {Ref6 }}$ | 2+ | 0.6 |
| - with concurrent bactaeremia | $20^{\text {Ref6 }}$ | 2+ | 0.1 |
| - hypertension in early adulthood | - | $2-$ | 0.1 |
| - end-stage renal disease in early adult | - | $2-$ | 0.06 |
| Candidiasis (thrush) | 2.5 (1.7-3.7) ${ }^{\text {Ref } 5}$ | 2+ | 10 |
| Prostate cancer | 1.2-2.0 ${ }^{\text {Ref5 }}$ | 2+ | 2-10 |
| Balanitis | 3.1 (1.9-5.0) ${ }^{\operatorname{Ref} 5}$ | 1+ | 10 |
| Phimosis | $100{ }^{5}$ | 1++ | 10 |
| High-risk HPV infection: RCT | 1.5 (1.1-2.0) ${ }^{\text {Ref } 5}$ | 1++ | 6 |
| High-risk HPV infection: meta-analysis | 2.7 (1.2-6.3) ${ }^{\text {Ref } 5}$ | 1+ | 10 |
| Herpes simplex virus type 2: RCT | $1.4(1.0-2.5)^{\text {Ref } 5}$ | 1++ | 4 |
| HSV-2: meta-analysis | 1.1 (1.0-1.3) ${ }^{\text {Ref } 5}$ | 1- | 1 |
| Genital ulcer disease | 2.0 (1.4-2.3) ${ }^{\operatorname{Ref} 5}$ | 1+ | 2 |
| Trichomonas vaginalis | 1.9 (1.0-3.6) $\operatorname{Ref} 5$ | 1+ | 0.5 |
| Mycoplasma genitalium | 1.8 (1.0-3.4) ${ }^{\text {Ref } 5}$ | 1++ | 1 |
| Chancroid | 0.1-1.1 ${ }^{\operatorname{Ref} 5}$ | 1++ | Low |
| Syphilis | 1.9 (1.2-2.9) ${ }^{\text {Ref } 5}$ | 2+ | Low |
| HIV (acquired heterosexually) | 2.4 (1.8-3.2) ${ }^{\text {Ref } 5}$ | 1++ | 0.3 |
| Penile cancer (lifetime) | >20 ${ }^{\text {Ref5 }}$ | 1++ | 0.1 |
| In female partner: |  |  |  |
| Cervical cancer | 2.4 (1.3-4.3) ${ }^{\text {Ref } 5}$ | 2++ | - |
| Chlamydia trachomatis | 5.6 (1.7-20) ${ }^{\text {Ref } 5}$ | 2+ | - |
| Herpes simplex virus type 2 | $2.2(1.4-3.6){ }^{\operatorname{Ref} 5}$ | 2+ | - |
| Trichomonas vaginalis | 1.9 (1.0-10) Ref 5 | 1++ | - |
| Bacterial vaginosis | $1.7(1.1-2.6)^{\text {Ref } 5}$ | 1++ | - |

Thus risk to an uncircumcised male of developing a condition requiring medical attention over their lifetime = very approximately 1 in 2.

Risks associated with neonatal circumcision

| Condition | Percentage <br> affected |
| :--- | :--- |
| Refs $1,2,5,7$ |  |

Thus risk of an easily treatable condition = approx. 1 in 200 and of a serious complication = 1 in 5000.

