### PROFOUND

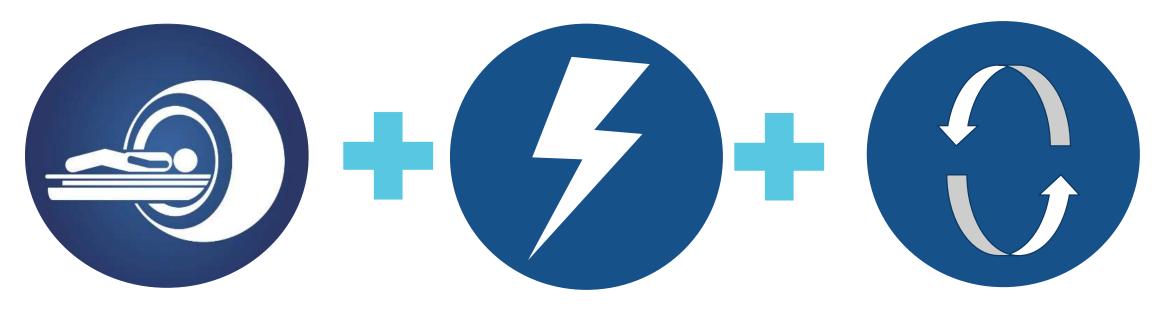
Customizable Incision-Free Ablation Therapies Men's and Women's Health

CORPORATE PRESENTATION | July 2019



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Creating Customizable Incision-Free Therapies By Combining Three Powerful Modalities



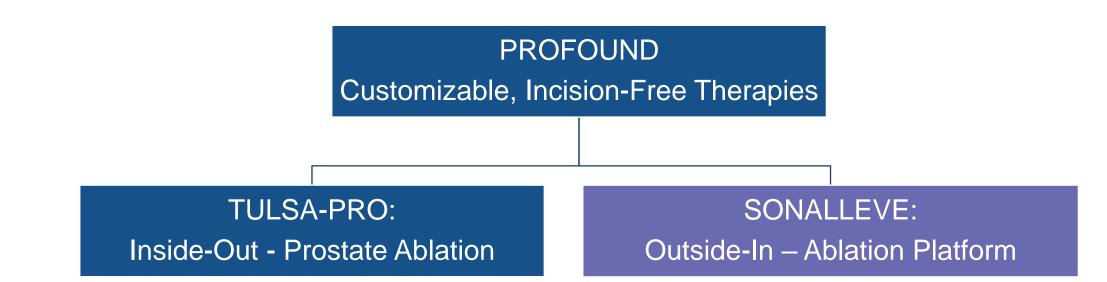
Real-time MRI imaging

Thermal ultrasound

Closed-loop temperature feedback control



#### **Customizable Incision-Free Therapies**



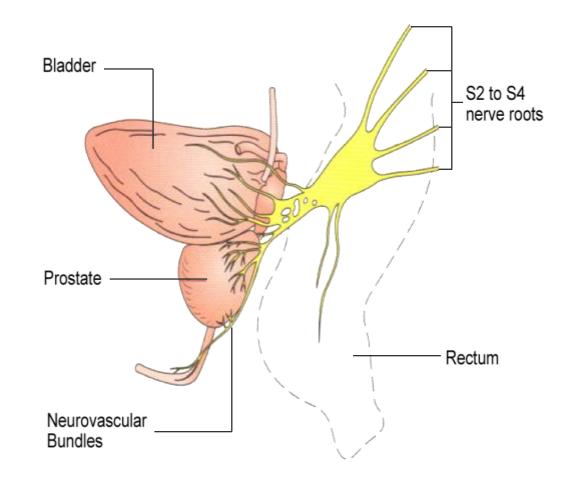


### TULSA-PRO°

CE Marked Filed For US FDA – May 2019 **'My life should not have to change'** 



#### Prostate Disease and Management

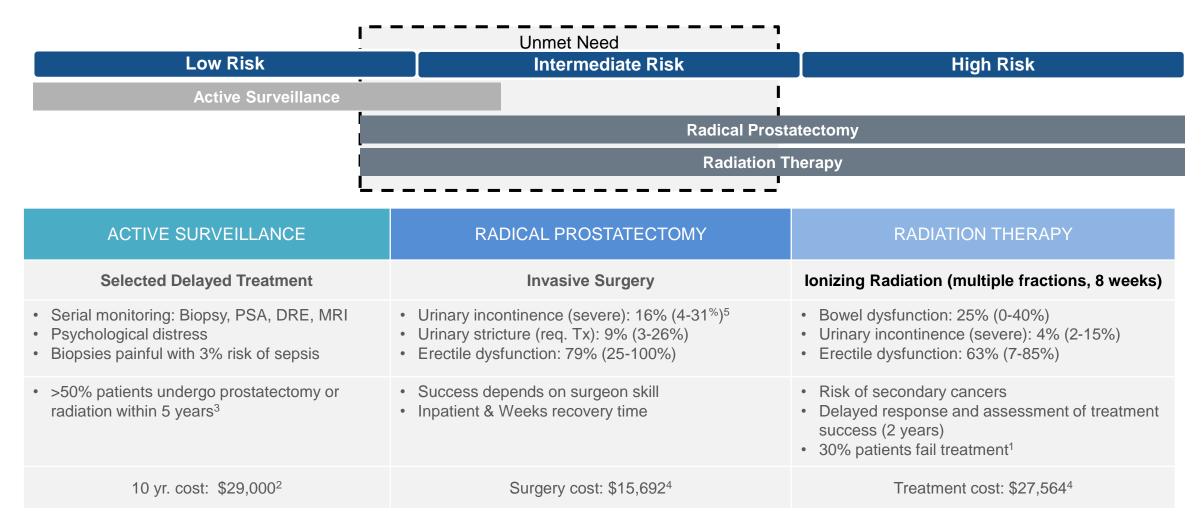




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Kirby (1997) An Atlas of Prostatic Diseases, The Encyclopedia of Visual Medicine Series.

#### Localized Prostate Cancer – Unmet Need in Standard of Care



Opportunity for patients with organ confined disease for less invasive, function preserving targeted therapies that do not preclude additional intervention if needed in the future



### MR-Guided TULSA – Closed Loop Temperature Control

## 1. Transurethral directional ultrasound ablation

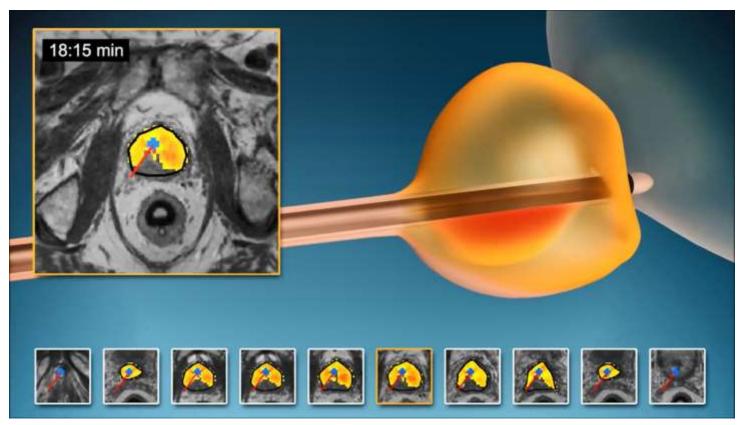
- Sweeping ultrasound, continuous rotation (no risk of cold spots between discrete sonications)
- Capable of treating large and small prostate volumes

# 2. Real-time MRI & Closed-loop thermal ablation

 Real-time temperature feedback provides millimeter accuracy

#### 3. Urethra and rectum cooled

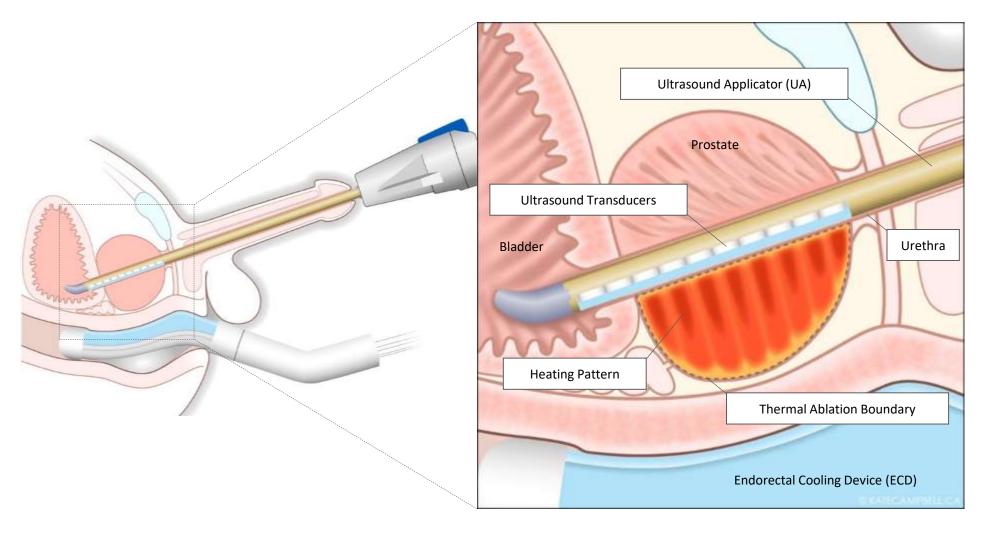
• Thermal protection of important anatomy





### TULSA-PRO – Prostate Ablation From The Inside Out

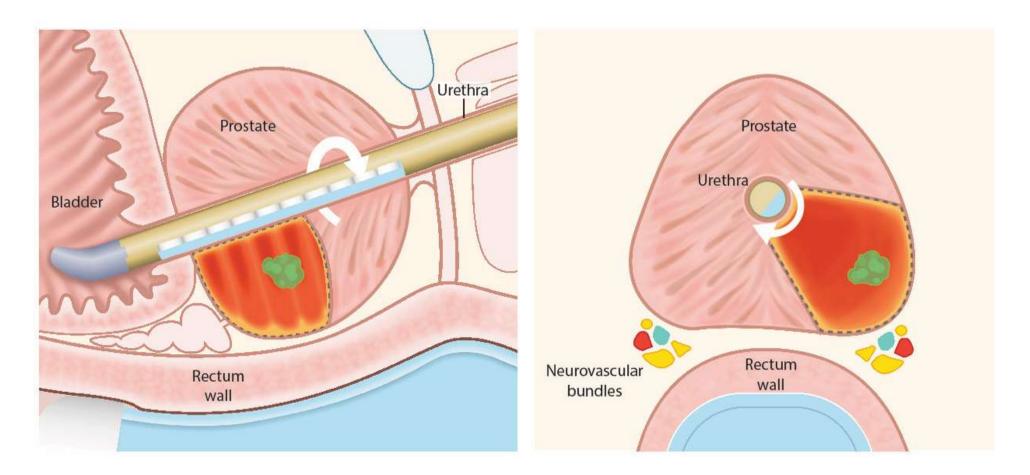
Whole Gland Ablation





### TULSA-PRO – Targeted Ablation

Partial Gland Ablation

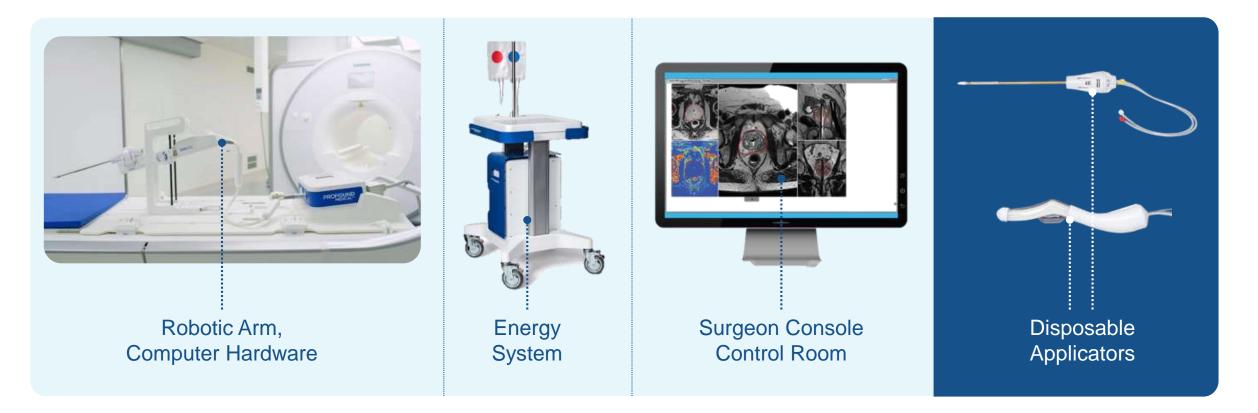






Equipment

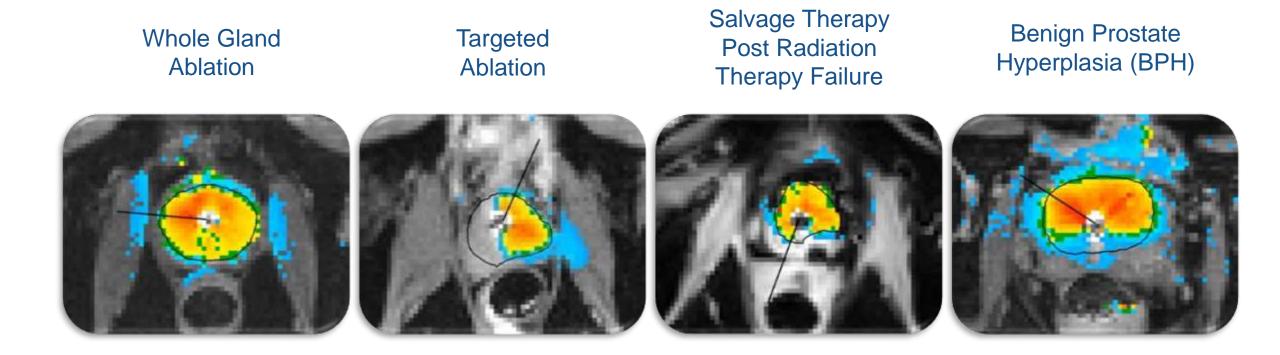
#### Compatible with MR from leading companies – Philips and Siemens





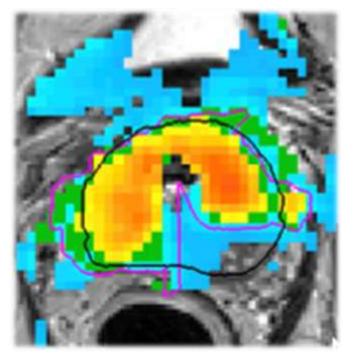
### TULSA-PRO – A Three In One Device For Ablation

Customizable, Predictable, Incision-Free



#### Example Prostate Tissue Ablation of Transition Zone & Suspicious Lesion

**20% of men over 50, 60% of men over 60 have BPH** Profound technology specially suitable for large prostates >80 CC



Patient with BPH and early stage lesion



### TACT – TULSA-PRO Clinical Trial Design

Pivotal study of whole-gland ablation in a clinically-significant patient population

#### **Study Population**

- n = 115, 13 clinical sites, 5 countries
- 45 80 years old
- Low (33%) & intermediate risk (67%) prostate cancer

#### **Ablation Treatment Plan**

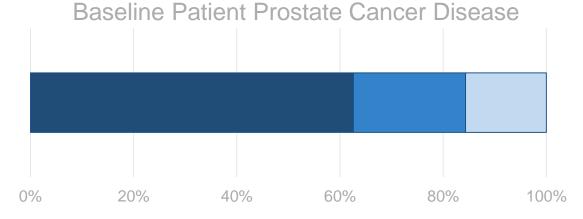
- Treatment intent was whole-gland ablation with sparing of the urethra and urinary sphincter
- Recommended by FDA to determine substantial equivalence with predicate devices and comparison with standard of care

#### **Primary Endpoints (12 months)**

- Safety: Frequency and severity of adverse events
- Efficacy: PSA reduction  $\ge 75\%$  (in > 50% of patients)

#### Secondary Endpoints (to 5 years)

- Prostate volume reduction at 1 year
- Prostate biopsy at 1 year in all patients
- Multi-parametric MRI at 1 year (Central Radiology Lab, Cleveland Clinic)
- Functional Disability: EPIC, IIEF, IPSS





Clinically significant GG2+ disease



Low volume GG1 disease (Very Low Risk)



### Prostate Ablation Efficacy – PSA

#### **PSA Primary efficacy endpoint resolutely met**

- Primary endpoint of PSA reduction ≥75% was achieved in 110 of 115 (96%)
- Median (IQR) PSA reduction was 95% (91-98%)
- Median PSA nadir was 0.34 (0.12-0.56) ng/ml

|                         | Pre-Treatment | 1 Month     | 3 Month     | 6 Month     | 12 Month    | PSA NADIR   |
|-------------------------|---------------|-------------|-------------|-------------|-------------|-------------|
| Ν                       | 115           | 113         | 115         | 115         | 115         | 115         |
| Median                  | 6.26          | 0.53        | 0.46        | 0.53        | 0.53        | 0.34        |
| IQR                     | 4.65 - 7.95   | 0.30 – 1.19 | 0.17 – 0.95 | 0.20 - 1.00 | 0.28 – 1.25 | 0.12 - 0.56 |
| Average                 | 6.72          | 0.90        | 0.77        | 0.77        | 0.93        | 0.51        |
| T-Test against baseline |               | <0.001      | <0.001      | <0.001      | <0.001      | <0.001      |

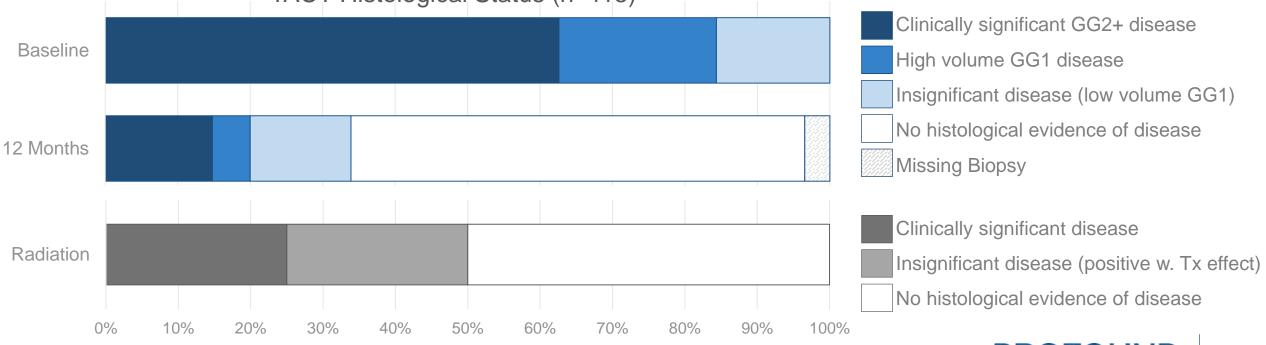
Missing values are interpolated using the LVCF method for the first time point after treatment.



### TACT Clinical Data As Presented At AUA – May 2019

#### TACT Biopsy Outcomes (1-year, 10-core TRUS, High Sampling Density 0.4 cc / core)

- Only 4 of 115 follow-up biopsies are missing, all due to patient refusal
- Among men with pre-treatment intermediate-risk GG2 disease, 54 of 68 (79%) were free of GG2 disease
- Of men with one-year biopsy data, 72 of 111 (65%) had complete histological response and were free of any disease
- 41% (16 of 39) of positive biopsies were clinically insignificant (Very Low Risk)
- Multivariate Analysis: Among men w. pre-Tx GG2 disease and w/o calcifications at screening, **51 of 60 (85%)** were free of GG2 disease



TACT Histological Status (n=115)

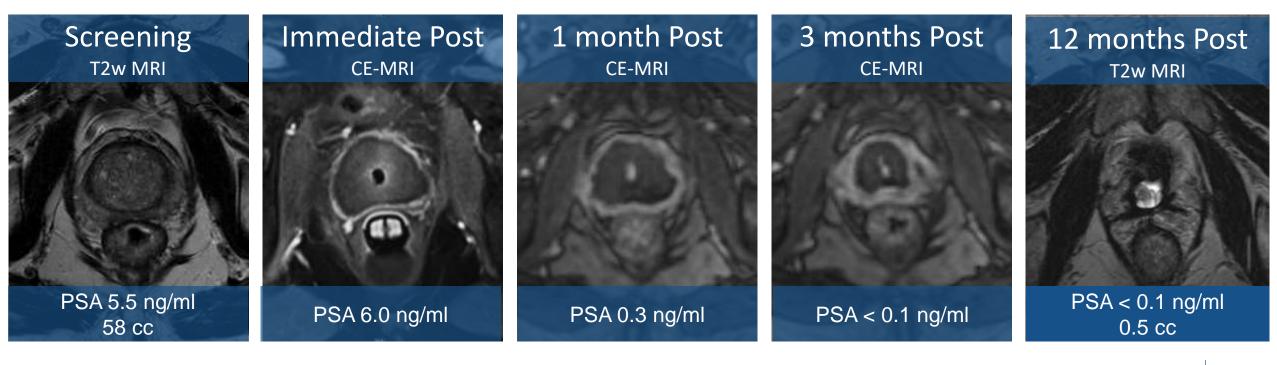
### Prostate Ablation Efficacy – Volume Reduction on MRI

Prostate Volume significantly reduced demonstrating effective prostate ablation

- Median perfused prostate volume decreased from 41 cc to 4 cc, on MRI at 1 year (interim analysis by local radiologists)
- Prostate volume reduction to be re-assessed by Central Radiology Core Lab, as per TACT protocol
- Prostate ablation confirmed on Contrast Enhanced MRI immediately after TULSA and during follow-up

Follow-up Prostate MRI predicts clinically significant disease on biopsy

 Multivariate Analysis: Absence of PIRADS ≥ 3 lesion at 1-year multi-parametric MRI has 92% Negative Predictive Value for absence of GG2 disease on 1-year biopsy (interim analysis by local radiologists, to be re-assessed by Central Radiology Core Lab)

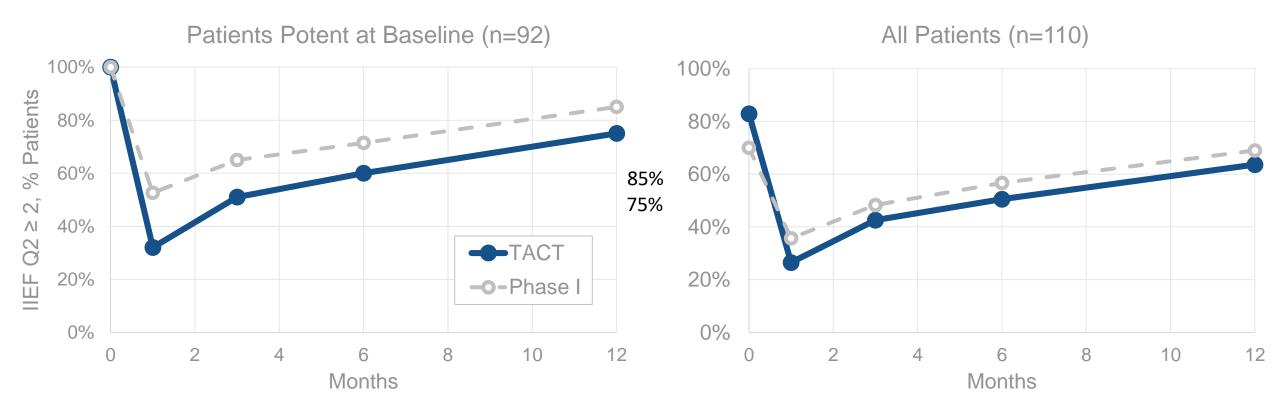




### TACT Erectile Function – As Presented At AUA – May 2019

#### **Erectile Function, at one year:**

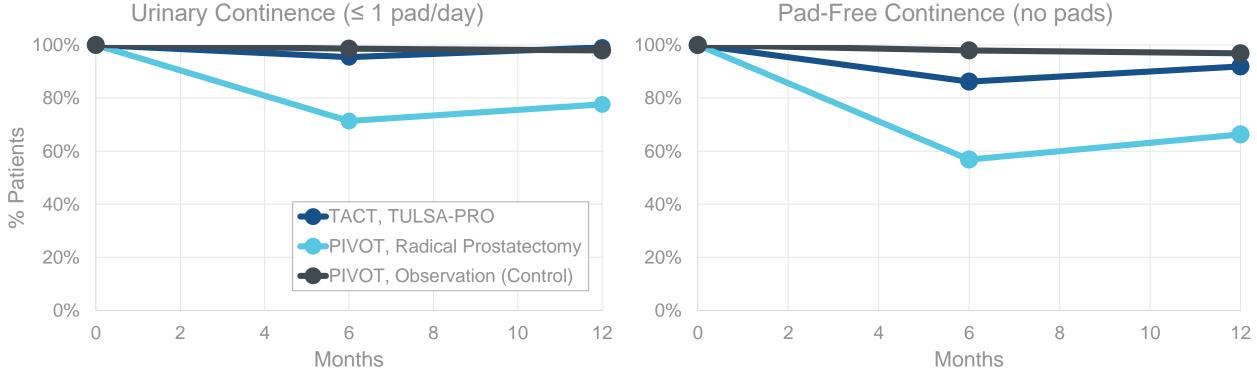
- 23% surgeon-assessed moderate erectile dysfunction (CTCAE Grade 2, intervention such as medication indicated)
- 0% any occurrence of severe erectile dysfunction (CTCAE Grade 3, intervention such as medication not helpful)
- 75% (69/92) of previously potent patients maintained erections sufficient for penetration
- Trend and recovery similar to Phase I



### **Urinary Incontinence** – Context to PIVOT

#### Urinary Incontinence (Pad use), at one year:

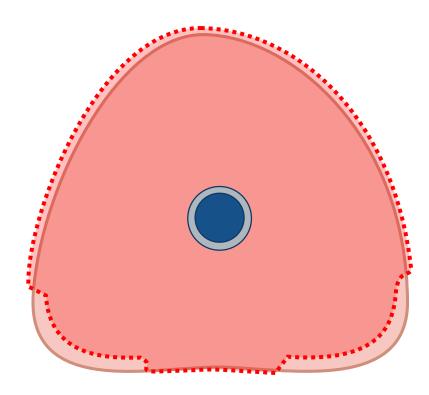
- TULSA Urinary Continence (≤ 1 pad/day) similar to Observation (control) arm of PIVOT study
- TULSA Pad-Free Continence (no pads) only 5%-points lower than Observation (control) arm of PIVOT study •
- TULSA continence outcomes markedly superior to Radical Prostatectomy arm of PIVOT study •
- PIVOT: Wilt et al, The New England Journal of Medicine, 2017 •



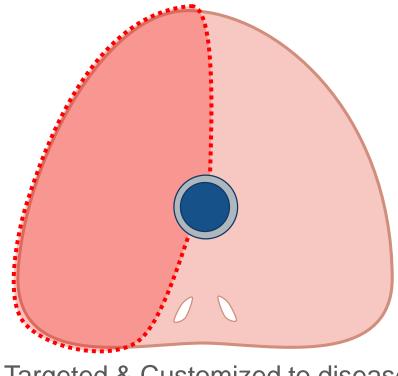
Pad-Free Continence (no pads)



# TULSA-PRO – Optimize Treatment Design – Maximize Efficacy, Minimize Side Effects



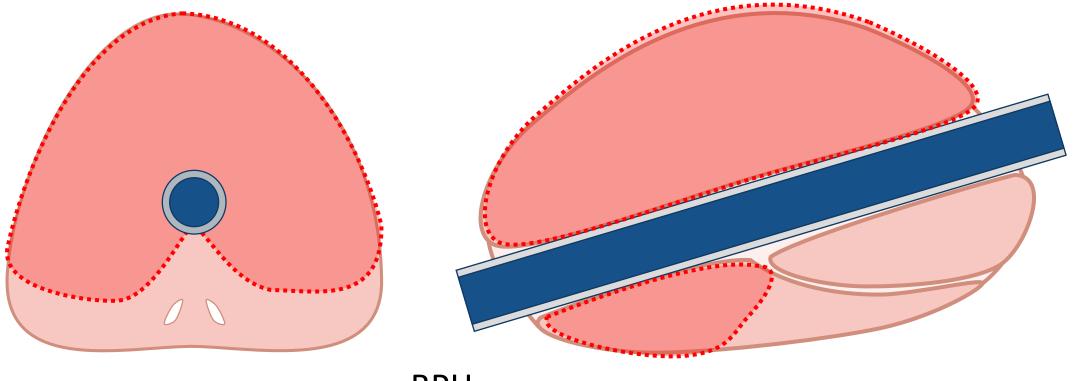
**Bilateral Sparing** 



Targeted & Customized to disease and anatomy



### TULSA-PRO – Real World Clinical Approach







#### Real World Context and Outcomes

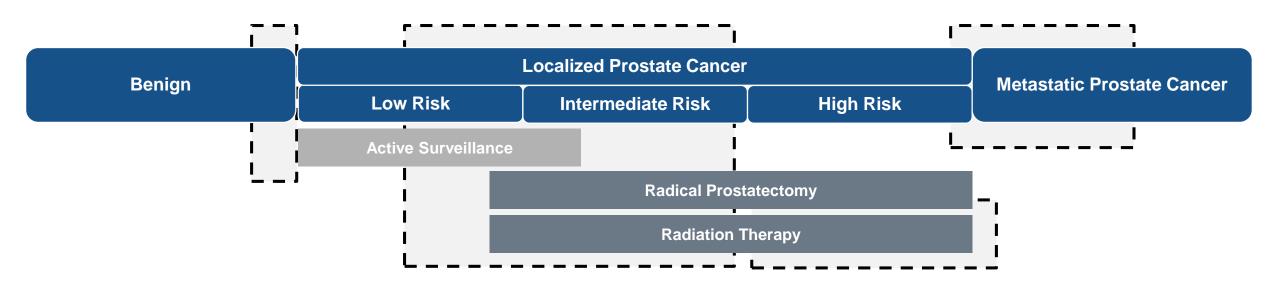
|   | Prostatectomy <sup>1-4</sup>   | Radiation <sup>1-5</sup>  | HIFU <sup>6-8</sup>   | TULSA (TACT)   |  |
|---|--|---|---|--|--|
| Biopsy /<br>Histology   | 16 – 24% Pos. Surg. Margin<br>(Meta-Analysis, Tewari <i>et al</i> 2012)<br>10 – 15% Pos. Surg.<br>Margin<br>(RCT, Yaxley <i>et al</i> 2016)<br>24% Pos. Surg. Margin<br>(ProtecT, Hamdy <i>et al</i> 2016)   | <ul> <li>50% Negative<br/>(Complete response)</li> <li>25% Insignificant disease<br/>(Positive w. treatment effect)</li> <li>25% Positive clinically<br/>significant Pca<br/>(Meta-Analysis Page 5, Approx. No.)</li> </ul> | 59 – 61% Negative<br>(Complete response, FDA IDE<br>Studies DEN150011 & K153023,<br>Intent to treat analysis)<br>63% Negative, after 40%<br>having repeat HIFU and<br>39% ADT (n=774, Crouzet <i>et al</i><br>2013) | 65% Negative<br>(Complete response)<br>14% Insignificant disease<br>(GG1, ≤2 cores, < 50% CCL)<br>21% Positive clinically<br>significant Pca |  |
| Erectile<br>Dysfunction<br>erections insufficient for<br>penetration                      | 79%<br>(Range: 25 – 100%)  | 63%<br>(Range: 7 – 85%)   | 58%<br>(Range: 38 – 67%)  | 20% – 25% - Grade 2<br>medication indicated.<br>No Grade 3 ED  |  |
| Urinary<br>Incontinence<br>moderate to severe   | 15%<br>(Range: 0 – 50%)  | 4%<br>(Range: 2 – 15%)  | 3%<br>(Range: 3 – 22%)  | 2.6% - Grade 2 pads<br>indicated. No Grade 3<br>Incontinence   |  |
| Urethral Stricture<br>moderate to severe  | 9%<br>(Range: 3 – 26%)   | 2%<br>(Range: 1 – 9%)   | 35%<br>(Range: 9 – 35%)   | 2.6%   |  |
| <b>GI Toxicity</b> , moderate<br>to severe diarrhea,<br>urgency, incontinence,<br>fistula | 15%<br>(Range: 0 – 24%)  | 25%<br>(Range: 0 – 40%)   | 7%<br>(Range: 1 – 21%)  | No GI Toxicity   |  |
| References  | <ol> <li>Thompson (Chair) <i>et al,</i> AUA prostate cancer clinical guideline update panel, J Urol 2007</li> <li>Resnick et al, Prostate Cancer Outcomes Study (PCOS), NEJM 2013</li> <li>Potosky <i>et al</i>, Prostate Cancer Outcomes Study (PCOS), J NCI 2004</li> <li>Elliott <i>et al</i>, CaPSURE database, J Urol 2007</li> </ol> |   | 5. Budaus <i>et al</i> , Review, Eur Urol 20012<br>6. FDA IDE Study K153023<br>7. FDA IDE Study DEN150011<br>8. Crouzet <i>et al</i> , Whole-gland HIFU, Eur Urol 2014  |  |  |

### TULSA-PRO – Customizable, Predictable, Incision-free

|                               | Prostatectomy                   | Radiation                                       | TULSA  |
|-------------------------------|---------------------------------|---|--|
| Outcome                       | Predictable                     | 2 year follow-up required                       | Predictable, NPV 95%                             |
| Treatment type                | Whole gland                     | Whole gland,<br>Limited customization           | Customized                                       |
| Throughput,<br>Procedures/day | 2 typically,<br>3 if longer day | Multiple sessions - 5 to<br>40 over 4 - 8 weeks | 4 in a routine day, Consistency, higher possible |
| Patient recovery              | Weeks                           | Deterioration over time                         | 2 days   |



### Broader & Deeper use of TULSA for Prostate Disease



#### BPH

- Large and Very Large
   Prostates
- Preservation of ejaculatory function
- Combined with targeted cancer ablation
- Prophylactic ablation of suspicious MRI lesion

#### Customized Targeted Ablation (25% - 99%)

- Targeted and customized to any size prostate and disease
- Large ablations (wide margins, not too focal, 25% - 99% ablation)

#### **Recurrence after Radiation**

 Localized recurrences have limited options, and morbidity is high

#### **Palliative Locally Advanced**

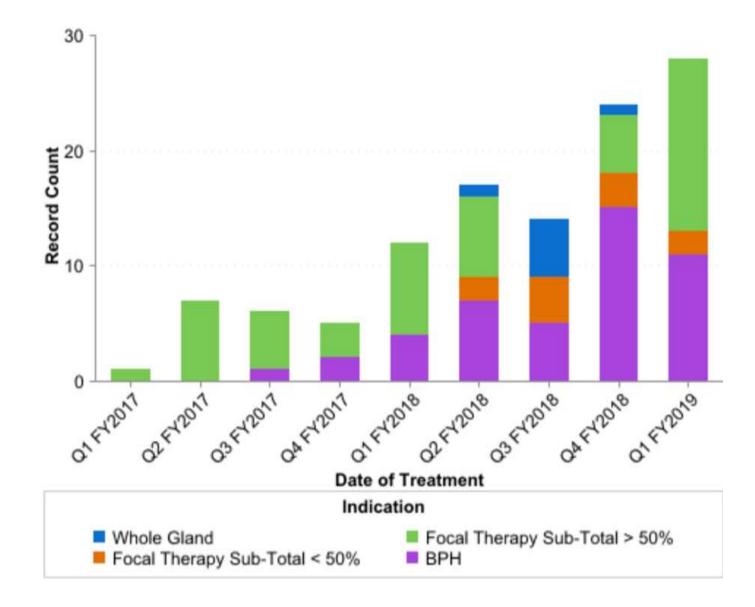
 Severe urinary symptoms including BOO with retention and/or intractable hematuria

#### Oligometastatic

- Benefit to locally treat prostate
- Often radio-recurrent



### TULSA-PRO In Commercial Use – Example From Europe



- Initiated Q1-2017
- Methodically increased usage
- Discovered potential to treat BPH patients – Q3-2017
- Streamlined procedure routinely 4 patients per day
- Increased utilization rate in 2019



#### TULSA-PRO Total Addressable Market: Pre-reimbursement

| <ul> <li>New Prostate Cancer Diagnosis (US + Canada)</li> </ul>  | 180,000 <sup>1</sup> |
|--|----------------------|
| • BPH, Prostates, surgical candidates, Unusual shapes (US + Canada)  | 400,000 <sup>2</sup> |
| <ul> <li>Total Opportunity, # of patients</li> </ul>   | 580,000              |
| <ul> <li>Total Addressable Market, patient paid is 5 -10% of total opportunity</li> </ul>  | 29,000 - 58,000      |
| <ul> <li>Add selected International markets (UK, Germany, Japan)</li> </ul>  | 14,500 - 29,000      |
| <ul> <li>Total patient pay addressable market # of patients</li> </ul>   | 43,500 - 87,000      |
| <ul> <li>Addressable market, \$4,000 per patient<br/>(includes: disposable + amortized capital + service)</li> </ul>                                   | \$174 – 348 M        |
| <ul> <li>Achievable share in X years, 25% (&lt;11,000 patients per year)<br/>TULSA Installed base = 110 at treatment rate 100 patients/year</li> </ul> | \$43.5 – 87 M        |

#### References:

- 1. Prostate cancer: 175,000 new prostate cancer diagnosed each year in US according to American Cancer Society
- 2. BPH: 300,000 surgeries based upon CMS data, + 1% of 10 Million BHP patients in United Stated + Canada



Prevalence Market Opportunity – Prostate Cancer

| Estimated Active Surveillance (AS) Population | 5.5 million |         |         |         |
|---|-------------|---------|---------|---------|
| Addressable AS Market –<br>Patient Pay (5%)   | ~ 275,000   |         |         |         |
| Penetration of AS Patient Pay<br>Population   | 5%          | 10%     | 20%     | 30%     |
| Patient Pay AS Market                         | \$55 M      | \$110 M | \$220 M | \$330 M |



#### TULSA-PRO Total Addressable Market – Additional Clinical Studies

| Title  | Purpose                         | Status/Comments  |
|--|---------------------------------|--|
| Radio-Recurrent<br>Cancer/Palliative Care                                | Inclusion in NCCN<br>guidelines | <ul><li>EU validation trial in progress</li><li>US study in H1-2020</li></ul>                          |
| Prostate Cancer:<br>Focal/Disease Targeted<br>Therapy. MR visible tumors | Adoption,<br>Reimbursement      | <ul> <li>Registry – EU H2-2019</li> <li>Active discussions in with Advisory<br/>Board in US</li> </ul> |
| Prostate Cancer  | Reimbursement                   | <ul> <li>Active discussions with Advisory<br/>Board in US now that TACT is<br/>complete</li> </ul>     |
| BPH, focus on surgical candidates  | Adoption,<br>Reimbursement      | <ul> <li>Validation studies – TURKU, ALTA.</li> <li>Initiate US study in H1-2020</li> </ul>            |



#### SONALLEVE

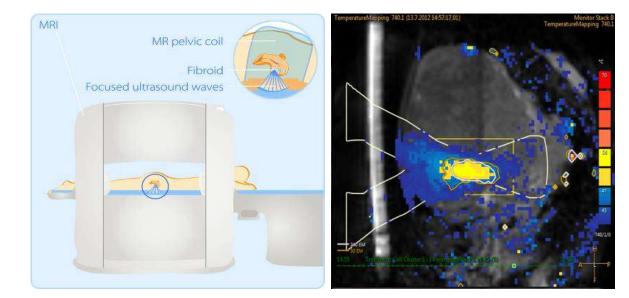
#### Technology platform for:

- Uterine Fibroid Treatment
- Bone Metastasis Pain
- Pediatric bone
- Hyperthermia

Over 200 publications from leading US and European clinicians and hospitals

CE Marked CFDA Approved







### Uterine Fibroid

Symptom Relief & Durability

In normal commercial use, over 85% of patients experienced sustained symptom improvement

| Months         | Patients available for | Symptom improvement |            |        |  |
|----------------|------------------------|---------------------|------------|--------|--|
| post-procedure | follow-up              | Improved            | No relief  | Worse  |  |
| 3 months       | 105                    | 90 (85.7%)          | 14 (13.3%) | 1 (1%) |  |
| 6 months       | 99                     | 92 (92.9%)          | 7 (7.1%)   | 0      |  |
| 12 months      | 89                     | 78 (87.6%)          | 11 (12.4%) | 0      |  |

#### Durability of the therapeutic effect compared to other uterine preserving treatments

| Need for alternative treatment    | @ 12 month | @ 24 month  | References |
|-----------------------------------|------------|-------------|------------|
| Myomectomy                        | 10.6 %     | 13-16.5 %   | 1,2,3,4    |
| UAE (Uterine Artery Embolization) | 7-10 %     | 12.7-23.7 % | 5,6,7      |
| MR-HIFU/MRgFUSNPV >60%            | 6 %        | 13 %        | 8          |

"Volumetric MR-guided high-intensity focused ultrasound ablation of uterine fibroids: treatment speed and factors influencing speed," M. J. Park, Y. S. Kim, B. Keserci, H. Rhim, and H. K. Lim, Eur Radiol, vol. 23, no. 4, pp. 943–950, Apr. 2013. 1. Gorny KR, Woodrum DA et al. Magnetic resonance–guided focused ultrasound of uterine leiomyomas: review of a 12-month outcome of 130 clinical patients. J Vasc Interv Radiol 2011 2. Subramanian S, Clark MA, Isaacson K. Outcome and resource use associated with myomectomy. Obs & Gyn.2001; 98: 583-587 3. Nezhat FR, Roemisch M, et al. Recurrence rate after laparoscopic myomectomy. Am Assoc Gynecol Laparosc. 1998;5: 237-240 4. Rossseti et al. Long term results of laparoscopic myomectomy: recurrence rate in comparison with abdominal myomectomy. Hum Reprod. 2001;16:770-774 5. Doridot et al. Recurrence of leiomyomata after laparoscopic myomectomy. J Am Assoc Gynecol Laparosc. 2001;8: 495-500 6. Spies JB, Bruno J, et al. Long-term outcome of uterine artery embolization of leiomyomata. Obstet Gynecol. 2005; 106: 933-939 7. Goodwin SC, Spies JB, et al. Uterine artery embolization for treatment of leiomyomata: long-term outcomes from FIBROID registry. Obstet & Gynecol. 2008; 111: 22-32 8. Sharp HT. Assessment of new technology in the treatment of idiopathic menorrhagia and uterine leiomyomata. Obstet Gynecol. 2006;108: 990–1003



- 1. Data compelling as presented
- 2. Focus on Asia
  - Reference site in S. Korea, treating 200 patients per year
  - Top tier hospitals in China. First site led by the President of Radiological Society of China



#### Sonalleve – Platform

- Additional applications
  - 1. Pain management
  - 2. Osteoid Osteoma
  - 3. Pancreatic cancer
  - 4. Hyperthermia
  - 5. Neuro-modulation
- Strategy Partner with Cologne and the FUS Foundation to continue to develop clinical data. Deploy recurring revenue business model for all new clinical applications



#### Summary

- 1. TULSA Applied for US FDA 510(k) clearance
- 2. Business model is capital efficient
  - Tulsa focus on US at key teaching hospitals and private clinics
  - Sonalleve focus on Asia
- 3. Patient-pay TAM \$50 100 Million per year
- 4. Potential to expand TAM by 10X following reimbursement
- 5. Future investments
  - Efficient sales team
  - Market expanding clinical trials
  - Continued product evolution





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Customizable Incision-Free AblationTherapies Men's and Women's Health | Oncology



