

*Witherspoon L, et al. A phase I study of an injectable lidocaine paste for spermatic cord block in men with chronic scrotal content pain*

**APPENDIX**

<b>Schedule of events</b>										
<b>Procedures</b>	Screening Part 1	Screening Part 2	<b>Day 0<sup>3</sup></b>	Day 1	Day 2–6	Day 7±1	Day 8–13	Day 14±2	Day 15–27	Day 28±3
Informed consent	X									
Demographics	X									
Medical history	X									
Concomitant medication	X		X	X		X		X		X
Physical examination and vital signs	X		X	X		X				X
Pain NRS (in clinic)	X	X	X	X		X				X
Lidocaine test injection		X								
CESI			X			X		(X) <sup>4</sup>		X
IIEF-5			X			X		(X) <sup>4</sup>		X
Injection of lidocaine study formulation			X							
Blood sample <sup>1</sup>		X	X	X		X				
Ultrasound			X			X				X
Telephone followup								X		
NRS diary (at home) <sup>2</sup>	X		X	X	X	X	X	X	X	
Adverse events evaluation			X	X		X		X		X

<sup>1</sup>Blood is drawn at the second screening for baseline electrolyte panel, kidney function and blood cell count and on day 0 (one-hour post injection,) on day 1 and on day 7±1 to determine lidocaine serum levels. <sup>2</sup>Subjects record their NRS three times a day for the first 14 days and then once daily for the remaining study period. <sup>3</sup>Day 0 is the study treatment day (7–21 days after injection of lidocaine test injection). <sup>4</sup>On day 14±2 subjects are asked the questions from CESI and IIEF-5 over the phone. CESI: chronic epididymitis symptom index; IIEF-5: International Index of Erectile Function; NRS: numerical rating scale.