





### Supplementary Figure 1

Abnormal slow-wave initiation: mismatched frequencies in adjacent gastric regions (non-continuous recording); example from diabetic gastroparesis (ID#6). **A.** Position diagram relating to (B). **B.** A persistent disorganized pattern of aberrant initiation was recorded at the mid corpus ( $4.5 \pm 0.3$  SD c/min). Representative electrograms are shown from the three distinct regions indicated in (A). The gross spatial disorganization precluded activation mapping; however, data animation demonstrated scattered focal activities, multiple propagating waves, and colliding wavefronts (*SuppFigure1i.wmv*). **C.** Position diagram relating to (E-F); the **recording** array was then relocated distally in the same patient. **D.** Example electrograms from eight electrodes positioned as indicated in (E), demonstrating lower frequency activity ( $2.3 \pm 0.1$  SD c/min;  $P < 0.0001$  vs corpus). **E.** Isochronal maps for representative waves (*i*) and (*ii*); intervals = 2 s. The distal activity was spatially dissociated from the disorganized proximal activity. Irregular focal activities occurred periodically in the distal field, colliding with the proximal wavefronts (*i*). At other times, the proximal activity successfully entrained the whole field (*ii*). The time stamps are referenced to the accompanying animation (*SuppFigure1ii.wmv*).

### Supplementary Figure 2

Abnormal slow-wave conduction in a patient with idiopathic gastroparesis (ID#10) and severe ICC depletion ( $0.3 \pm 0.6$  SD bodies / field). **A.** Position diagram. **B.** Electrograms (from the positions indicated in C) demonstrate regular activity of normal frequency ( $3.4 \pm 0.2$  SD c/min). **C.** A consistently stable but highly deranged propagation pattern was observed for the recorded duration (maps *i-ii*, see also animation *SuppFigure2.wmv*), including conduction block (grey bar), circumferential propagation distal to the block, and retrograde propagation with colliding wavefronts.

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Comparison of patient and experimental variables between the present study and a previous study of human subjects with normal stomachs, who were mapped and analyzed using similar methods <sup>16</sup>, and in whom no abnormal slow-wave events were observed intra-operatively.

\* ([www.smoothmap.org](http://www.smoothmap.org)).

### Supplementary Table 2

Individual patient data from the study cohort. *n.d.: no data.*

Supplementary Table 1

	<b>Cohort with Normal Stomachs</b> <sup>16</sup>	<b>Gastroparesis Cohort (current study)</b>
<i>No. Patients</i>	12	12
<i>Age (median, range)</i>	50 yrs (21-60)	42 yrs (30-60)
<i>Sex (M:F)</i>	7:5	6:6
<i>Time of Mapping</i>	Immediately following incision	Immediately following incision
<i>Duration of Mapping (mean ± SD)</i>	11.5 ± 3.9 min / pt	13.4 ± 4.6 min / pt
<i>Anaesthetic Regimen</i>	Prophylactic antibiotics, benzodiazepine premedication, an epidural anesthetic, a short-acting intravenous opiate, muscle relaxant (atracurium or suxamethonium), propofol, isoflurane or sevoflurane.	Prophylactic antibiotics, benzodiazepine premedication, a short-acting intravenous opiate, muscle relaxants (suxamethonium or rocuronium), propofol, desflurane.
<i>Off-line Filtering Methods</i>	2 Hz low-pass Butterworth filter	Savitzky-Golay and moving median filters <sup>19</sup>
<i>Analysis and Isochronal Mapping Methods</i>	Manual analyses in SmoothMap v3.05*	FEVT, REGROUPS, and SIV methods (performed in GEMS v1.2 <sup>18,20,21</sup> with manual review)
<i>Velocity and Amplitude Calculation Methods</i>	Algorithms in SmoothMap v3.05*	Algorithms in GEMS v1.2 <sup>18,19,23</sup>

Supplementary Table 2

<b>ID</b>	<b>Aetiology</b>	<b>Sex</b>	<b>Age</b>	<b>BMI</b>	<b>TSS</b>	<b>GET (4-hr Retention)</b>	<b>ICC Count (mean, SD)</b>	<b>Figures / Animations</b>
1	Diabetic	M	32	28	13	27%	n.d.	<i>Figure 5;</i> <i>Figure5i-ii.wmv</i>
2	Diabetic	M	39	27	n/a	41%	n.d.	<i>Figure 6;</i> <i>Figure6i.wmv</i>
3	Diabetic	M	32	21	14.5	22%	1.7 ± 1.5	<i>Figure 1</i> <i>Figure1.wmv</i>
4	Diabetic	M	30	21	17.5	74%	2.1 ± 1.3	<i>Figure 6;</i> <i>Figure6ii.wmv</i>
5	Diabetic	F	50	32	18	27%	3.3 ± 2.0	<i>Figure 4;</i> <i>Figure4i-iii.wmv</i>
6	Diabetic	M	42	n.d.	14	47%	2.5 ± 1.8	<i>Supp. Fig 1;</i> <i>SuppFig4i-ii.wmv</i>
7	Diabetic	F	38	n.d.	17.5	14%	3.9 ± 2.5	-
8	Diabetic	M	62	44	20	25%	2.0 ± 1.6	-
9	Idiopathic	F	58	16	14	19%	n.d.	<i>Figure 5</i>
10	Idiopathic	F	62	23	16	75%	0.3 ± 0.6	<i>Supp. Fig 2;</i> <i>SuppFig2.wmv</i>
11	Idiopathic	F	34	30	20	15%	2.5 ± 2.1	-
12	Idiopathic	F	58	27	15.5	19%	2.2 ± 1.7	-
<b>Median:</b>	-	-	<b>42</b>	<b>27</b>	<b>16</b>	<b>26%</b>	<b>2.2</b>	-

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