You Need a Thymatron® System IV to provide higher stimulus current, deeper cerebral stimulus penetration, and better seizure generalization.

You Need a Thymatron® System IV to provide the most effective form of bilateral brief pulse ECT with a 0.5 msec pulsewidth that minimizes side-effects and can be delivered at every stimulus dose including the maximum dose.

You Need a Thymatron® System IV for efficient and accurate stimulus titration.

“THYMATRON® Why Trust Anything Less?”
The Thymatron® System IV: Description and Specifications

ULTRABRIEF STIMULI: Ultrabrief stimuli need higher current for greater efficacy. If you still use the 800 mA maximum current of Mecta devices, to assure full efficacy you should upgrade to the 900 mA provided by Thymatron® instruments. Further, the Thymatron® System IV is the only instrument that delivers a 0.3 msec ultrabrief stimulus across the entire dosage range, 25% more than the Mecta.

EASIEST SEIZURE INDUCTION WITH THE THYMATRON SYSTEM IV: Thymatron® stimuli are 60% more effective for inducing seizures than the 800 mA Mecta stimuli (Chanpattana et al., 2001). This enormous difference results from the much higher stimulus dose of Thymatron® 900 mA current than 800 mA, yielding a larger volume of seizure foci in the brain (Swartz, 2006). This provides clinicians with a dose advantage in treating high-threshold geriatric patients. With the new age-adjusted titration method for Thymatron® instruments seizure threshold titration is easier, more precise, and requires fewer stimulations.

SPECIAL EEG SIGNAL PROCESSING MEASURES: provide continuous monitoring of level of consciousness and cortical activity; 95% Spectral Edge Frequency, Relative Delta Power, and Median Frequency (Billard V et al, 1997: A comparison of spectral edge, delta power, and bispectral index as EEG measure of alfentanil, propofol, and midazolam drug effect; Hans P et al, 2001: Effect of nitrous oxide on the bispectral index and the 95% spectral edge frequency during propofol-fentanyl anaesthesia; Sakai T et al, 1999: Hypnotic endpoints vs. the bispectral index, 95% spectral edge frequency and median frequency during propofol infusion with or without fentanyl).

STATE-OF-THE-ART 4-CHANNEL PRINTER STANDARD ON ALL UNITS allows you to monitor two channels of EEG, plus ECG and EMG (or, choose 4 channels of EEG), while providing hard-copy documentation for later reference.

SINGLE FRONT-PANEL DIAL lets you select the traditional Thymatron® functions plus important new ones, including Optimal Stimulus programs that automatically set the most efficient combination of stimulus parameters at every stimulus dose setting.

ELECTRONIC MEDICAL RECORD-KEEPING is simple with the included Genie™ IV EMR software. Patient treatment records created and stored with the Genie™ IV are easily incorporated into hospital database systems.

EXTENDED LOWER STIMULUS RANGE with pulsewidth and frequency to 0.25 or 0.3 msec and 10 Hz allows you to deliver stimuli up to 8 seconds long, to optimize treatment in accordance with research showing greater efficacy of short-pulsewidth, extended-duration stimuli (Iseberg et al, 1996).

EEG COHERENCE MEASURES of maximum sustained coherence, and time to peak coherence, interhemispheric cross-correlation measures reported to reflect seizure quality and clinical impact (Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998).

EEG AMPLITUDE measures of maximum sustained EEG power, and average seizure energy, with separate values for early, mid- and postical seizure phases, found by the Duke University group to be important correlates of seizure quality and efficacy (Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998).

HEART RATE MEASURES, including peak heart rate, a key measure of cerebral seizure duration and quality (Larson, Swartz & Abrams, 1984; Swartz, 1993; 1996; Swartz and Manly, 2000) that reflects the autonomic (brainstem) response to ECT. This is supplemented by continuous digital heart rate monitoring for safety and seizure generalization, with the result printed each second.

All of the above measures are automatically printed.

A POWERFUL 32-BIT INTERNAL COMPUTER employs Power Spectral Analysis to process and store up to 10 minutes of digitized EEG for the special features described here. You can send this data to your IBM PC-compatible computer via a rear-panel serial port for further comprehensive EEG analysis, using Somatics' proprietary Genie™ IV software.
Because each ECT treatment session is **STORED IN MEMORY**, you can retrieve it if you run out of paper during a treatment—just slip in another pad after the treatment and **press a button for a complete printout**.

**PATENTED INDEPENDENT SAFETY MONITOR CIRCUIT** prevents the patient from receiving an excessive electrical dose regardless of the operation of the regular circuits.

**TRUE EMG RECORDING OF THE MOTOR SEIZURE.** Unlike simple movement detectors, the Thymatron® System IV's EMG can measure seizure muscle activity that is not visible to the naked eye, and which typically continues substantially longer than optically-detectable movements (Couture et al, 1988).

Because the special computer-automated programs of the Thymatron® System IV are stored on REPLACEABLE MICROCHIPS, updates are easily accomplished on-site via chip replacement. Somatics has already provided 4 advanced microchip upgrades for the System IV including: the ultrabrief 0.25 msec pulsewidth program, Genie™ IV computer software, real-time digital EEG monitoring. In comparison, any upgrades to the Mecta spectrum (there have been none) would have required return to the factory.

The **POSTICTAL SUPPRESSION INDEX** reports the degree of EEG flattening immediately following the seizure, which correlates with clinical efficacy (Nobler et al, 1993; Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998; Nobler et al, 2000). A recent study of the Thymatron™'s Postictal Suppression Index found that it significantly differentiated ECT remitters from non-remitters (Petrides et al, 2000). The authors concluded: “**higher PSI values (more abrupt ending of ictal EEG) are correlated with better clinical outcome of ECT in depression**”.

**COMPUTER DETERMINATION AND PRINTOUT OF EEG AND MOTOR SEIZURE DURATIONS.** The integral computer EEG analyzer continually measures the EEG and EMG and automatically prints the EEG and motor seizure durations with precision and reliability (Swartz et al, 1994; Krystal et al, 1995).

**JUST SET ACCORDING TO AGE AND TREAT.** Setting the Thymatron® System IV according to the patient's age facilitates easy selection of the stimulus charge.

Alternatively, **RAPID STIMULUS TITRATION** is facilitated with the Thymatron® System IV using a simple method-of-limits procedure (McCall et al, 1993; Rasmussen et al, 1994) that employs research based dose increments: 5, 10, 15, 25, 40, 80, and 100% Energy at your choice of pulsewidth.

(see next page for references)
REFERENCES


Alkire MT (1998): Quantitative EEG correlations with brain glucose metabolic rate during anesthesia in volunteers. Anesthesiology 89:323-33


NEW REMOTE TREAT HANDLE FOR THYMATRON®

You asked for a remote treat handle and here it is. You can press the TREAT button on this handle instead of reaching over to the Thymatron® itself: a simple thumb press safely triggers the stimulus for any electrode placement, including unilateral.

MICROSTIM™ PERIPHERAL NERVE STIMULATOR

This hand-held, solid state, peripheral nerve stimulator weighs only 7 oz. It applies a pulsed 0.2 msec square-wave stimulus through surface electrodes to precisely determine the point at which a safe degree of succinylcholine-induced muscle relaxation has been achieved. The operator has the option of selecting continuous (tetanus) or intermittent (twitch) stimulus modes. Battery powered (9 volt alkaline), it comes in a soft carrying case that clips to pocket or belt. 2.4” x 1.0” x 3.8”

#ENSI

NEW REMOTE TREAT HANDLE FOR THYMATRON®

You asked for a remote treat handle and here it is. You can press the TREAT button on this handle instead of reaching over to the Thymatron® itself: a simple thumb press safely triggers the stimulus for any electrode placement, including unilateral.

#EEDS

SOMATICS’ OWN DISPOSABLE SELF-STICK EEG/ECG/EMG ELECTRODES

Easy and quick to use, “the pregelled electrodes provided in the Thymatron DG starter kit . . . reduce preparation time” (Convulsive Therapy 2:53, 1986), compared to metal electrodes and ordinary disposable paper ECG electrodes. Their small size facilitates bifrontal or fronto-mastoid application without interfering with treatment electrode placement. Ideal for recording EEG, ECG, and EMG, they are conveniently packaged 5 per strip. Instantly adherent, they will remain in place throughout the seizure.
This sample ECT report of the Thymatron® System IV shows that the doctor set the % Energy dial to his patient's age of 45 years, yielding a 308 mC stimulus charge. The Optimal Stimulus Program selected a 0.3 msec pulsewidth, 70 Hz frequency stimulus delivered over 7.2 sec. Prior to stimulus administration the impedance measured a safe 1440 ohms, which dropped to 260 ohms during stimulus delivery.

The EEG seizure lasted 48 seconds. Peak seizure amplitude was reached at 31 sec, with a mid-ictal amplitude of 264 µV, a Maximum Sustained Power of 77841 µV², and an Average Seizure Energy Index of 72 V² reflecting strong seizure intensity.

Peak Interhemispheric Coherence reached at 33 sec was consistent with the seizure amplitude peak at 31 sec. The Maximum Sustained Coherence value of 95% reflected synchronous participation of both hemispheres in the seizure. The rapid drop of EEG seizure amplitude to 10 µV postictally yielded a high Postictal Suppression Index of 96%. Power Spectral Analysis was not enabled.

In summary, the record shows a synchronous, high-intensity, well-developed, and well-generalized EEG seizure pattern with a strong midictal phase, pronounced postictal suppression, and a substantial tachycardia response—which is to say, an ECT-induced seizure of high expected clinical efficacy (Abrams, 2002)

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**GENIE™ IV ELECTRONIC PATIENT DATABASE AND EEG MONITORING SYSTEM**

Designed to meet your clinical and research needs, the Genie™ IV enables you to enter complete patient information at each treatment for storing, printing or incorporating into a hospital-based electronic patient database system.

Equally important is the Genie™ IV’s comprehensive real-time display of up to 4 channels of EEG, ECG, and EMG on a PC screen (not included), allowing you to monitor and then store each treatment session.

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**GENIE™ IV Patient Information Data File/Printout**

Date: 12-16-05  Name: Laurenz Smarba  Age: 58  Sex: M  
Somewhat improved but still has insomnia & poor appetite  
Oriented, alert, coherent and cooperative  
ECT #3 (R-UNI x 1)  Anesthesia: Dr. Jones  ECT: Dr. Smith  
Atropine 0.2 mg - Brevital 50 mg - Succinylcholine 40 mg  
Thymatron IV  85% Energy (LOW 0.5 program)  
Moderately strong seizure-symmetrical, well developed  
Good heart rate response with rapid return to baseline  
No complications, quick recovery  
Recommendation for ECT #4: same as above
DOES YOUR ECT DEVICE DELIVER THE DOSE YOU SPECIFY?  
DO YOU TRAIN DOCTORS OR NURSES IN ECT QUALITY?

Device malfunction can cause ineffective ECT treatments or excessive side-effects. Now you can check your ECT device yourself with Somatics’ easy-to-use, patented ECTOBRAIN™ II, which performs the same current output check professional engineers use. A single button press instantly tells you if your ECT device is operating safely—providing reassurance and peace of mind. ECTOBRAIN™ II works with any Thymatron.™

ECTOBRAIN™ II also features a Patient Simulator mode that generates EEG, ECG, and EMG signals derived from real patients for testing up to 4 channels of your monitor/printer tracing display and for training and demonstration purposes. Both good- and poor-quality seizures can be selected.

The good-quality seizure shows a high amplitude EEG followed by electrical silence at termination, with a pronounced tachycardia response and a high-amplitude EMG that terminates shortly before EEG termination. The poor-quality recording exhibits a low-amplitude abortive-type EEG seizure lasting only 10 sec, followed by continued but lower-amplitude EEG fluctuations after termination; there is no tachycardia response, and an initial low-amplitude EMG response lasts only a few seconds.

A device checkup can cost $600 to $800 but real costs are more. How often does the question arise in treating a difficult patient whether the ECT device is stimulating properly or the EEG tracing recording correctly? Most ECT units sent to us for presumed malfunction have nothing wrong with them! ECTOBRAIN™ II can quickly determine whether or not the device is working. It can reveal problems in technique (e.g., recording electrode application) that are correctable on site or with user-replaceable parts (e.g., lead wires). Just connect the stimulus and recording cables and press the TREAT button as for a patient.

The chart recorder of your ECT device will display samples of EEG, ECG, and EMG tracings as described above. The printed report will show the values of the stimulus parameters and other printed variables of your ECT device, including the measured stimulus charge output in mC.

Satisfaction guaranteed by Somatics’ 30-day unconditional full-satisfaction trial period. 5-year warranty on parts and labor.

Special price when ordered together with a Thymatron® System IV.

### Trouble-Shooting with the ECTOBRAIN™ II

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* See specified sections of Ectobrain™ II manual on [www.thymatron.com](http://www.thymatron.com) downloads page

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This ground-breaking new text edited by internationally-recognized ECT expert Dr. Conrad M. Swartz comprehensively covers the scientific basis and clinical practice of ECT as well as the latest nonconvulsive electrical and magnetic brain stimulation therapies. The many expert contributors from around the world illustrate compellingly that ECT is now a mainstream psychiatric treatment. The wealth of new and surprising information it contains is certain to provide ECT practitioners with much enjoyable "brain stimulation".

Call David Mirkovich at 1(800)642-6761 for our very attractive price.
SPECIAL LIMITED-TIME TRADE-IN OFFER
Somatics is pleased to offer a substantial trade-in amount towards a new Thymatron® System IV for any model or age MECTA ECT machine or Thymatron® ECT instrument, without exception. Call us at (800) 642-6761 for the details--offer expires April 30, 2012.

SOMATICs SUPPORTS & SERVICES EVERY THYMATRON® IT HAS SOLD
Unlike our competitors, Somatics has always provided support and service for every Thymatron® it has ever sold, including several that are now a quarter-century old. Just e-mail or call us as indicated below to obtain a fair and reasonable quote for servicing your old faithful Thymatron®.

SOMATICs THYMATRON® INSTRUMENTS IMPORTANT RESEARCH TOOLS
Since the Thymatron® was first introduced in 1983 hundreds of research studies have appeared in the medical literature using a Thymatron® instrument. Prominent among these is the series of publications by the multi-hospital CORE research group, a consortium of academic psychiatric centers. In a series of important articles over the last decade the CORE group used Thymatron® ECT instruments to demonstrate the striking efficacy of ECT in the treatment of psychotic depression (Petrides et al, 2001), determine that age had a strong positive association with the response to bilateral ECT (O’Connor et al, 2001), show that DSM III melancholic features are unreliable predictors of ECT response (Fink et al, 2007), find that unipolar and bipolar depressives respond equally well to ECT (Bailine et al, 2010), and report that, although fewer black than white depressed patients received ECT, there was no overall racial difference in treatment response (Williams et al, 2008). Hundreds of other studies used a Thymatron® instrument to demonstrate, among other things, that ECT given twice a week was equally effective as three times a week, but with fewer cognitive side-effects (Lerer et al, 1995).

Antidepressant potency of high-dose right unilateral ECT was equal to bilateral ECT (Abrams et al, 1991). Caffeine lengthened seizure duration but did not change the convulsive threshold (McCall et al, 1993). Bilateral ECT did not yield any evidence for brain damage as measured by levels of neuron-specific enolase and S-100 protein (Agelink et al, 2001). ECT was nearly four times more effective than transcranial magnetic stimulation (TMS) for major depression (Eranti et al, 2007).

None of 7 patients with intracranial masses were neurologically adversely affected by ECT (Rasmussen et al, 2007). In 28 severely depressed patients given a course of unilateral ECT, only responders showed elevations of N-acetylaspartate, suggesting that ECT exhibits positive neurotrophic effects (Michael et al, 2003).

In 32 consecutive patients seizure durations automatically reported by the Thymatron instrument correlated highly with determinations made by trained physicians (Rosenquist et al, 1998).

REFERENCES

The Choice is Easy (and Smart!)
Thymatron® Why Trust Anything Less?
"Thymatron® System IV: The most advanced ECT device technically and operationally, with demonstrated superior safety and clinical effectiveness."

SAFE, TIME-SAVING DISPOSABLES FOR ECT

THYMAPAD™ Adherent Stimulus Electrodes

Thymapads™ are much faster and easier to use than the old-fashioned disk, headstrap, and jelly method.

They remain exactly where applied and have no exposed metal surfaces to cause accidental shocks. There’s no mess to clean up afterwards, nothing to wash, dry, or sterilize, no sticky hands - just remove them and discard.

Thymapads™ flexibly conform to the surface of the head and fit all Mecta machines too.

VENTIL-A™ Mouth Protector

The Ventil-A™’s thick 100% closed-cell foam construction protects all the teeth. Fits easily under any anesthesia mask and features a non-collapsible air channel for free flow oxygen. One-piece design for dimensional stability and looped end for fast and easy insertion/removal. One size fits >98% of adults.

Both of these single-use ECT aids (US Patent 6039046) save the time and expense of washing and sterilization and eliminate the risk of cross-infection that occurs with re-usable products.