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UNITED STATES DISTRICT COURT

DISTRICT OF NEVADA

MEDTRAK VNG, INC., a Nevada corporation,

Plaintiff,

v.

ACUNETX, INC., a Nevada corporation, and
CHAPIN HUNT, an individual,

Defendants.

CASE NO. 2:12-CV-853

**[PROPOSED] TEMPORARY
RESTRAINING ORDER**

THE COURT, having reviewed and considered Plaintiff's Application for Temporary Restraining Order ("Application"), the exhibits filed in connection therewith, and the pleadings of this case, and having held a hearing on the same on _____, 2012, in which the Court entertained the arguments of counsel and representatives for each of MEDTRAK VNG, INC, ACUNETX, INC., and CHAPIN HUNT, finds and orders as follows:

1. On or about March 23, 1994, the FDA issued a 510(k) Registration (FDA 510(k) Number K925111) for the VNG device at issue. This FDA 510(k) Registration was originally issued to Eye Dynamics, Inc. ("EDI"), AcuNetx's predecessor in interest.

2. Under the terms of a 2004 Software Ownership Agreement, the VNG Software used in relation to the VNG device was assigned and sold to MedTrak and licensed back to EDI/AcuNetx to facilitate continued manufacture of the systems. Plaintiff, MedTrak, has