EXHIBIT-16

TEN-POINT QUALITY ASSURANCE PROGRAM

1. SPECIFICATIONS

GUIDELINES:

U.S.P. AND N.F.

PROFESSIONAL GUIDANCE FROM DMMB

TECHNICAL DATA FROM OTHER SOURCES

REASONS FOR ADDITIONAL SPECIFICATION REQUIREMENTS:

ADVANCEMENT AND MODIFICATIONS TO U.S.P. AND N.F. ARE TIME CONSUMING

SPECIAL PACKAGING

DETERIORATION OF DRUGS DUE TO CONDITIONS AND STORAGE

STABILITY TESTS

ADVANCED INSTRUMENTATION INCREASING PURITY

2. PRE-AWARD SURVEY

ESTABLISH CAPABILITY AND QUALIFICATIONS

PLANT EVALUATION

3. PRE-AWARD SAMPLES

DETERMINE FIRM'S POTENTIAL

ANALYSIS OF SAMPLES