WHAT IS CLAIMED IS:

1. A cartridge for determining the presence and/or amount of a microbial contaminant in a sample, the cartridge comprising:

    a. a housing including an optical sample well, a fluid inlet port and a conduit fluidly connecting the fluid inlet port and the optical sample well;

    b. a pump mechanism associated with the housing and fluidly connected to the fluid inlet port, the conduit and the optical sample well;

    c. a mixing zone located along the conduit configured to include a dried composition including a hemocyte lysate.

2. A cartridge for determining the presence and/or amount of a microbial contaminant in a sample, the cartridge comprising:

    a. a housing including a cover section, a base section mechanically connected to the cover section, and a manifold section mechanically connected to the base section;

        i. the base section including two optical sample wells, a fluid inlet port and a conduit fluidly connecting the fluid inlet port and the optical sample well;

        ii. the conduit including an aspiration zone, a fluid restrictor, and a mixing zone;

        iii. the manifold section including a mixing portion configured to include a dried composition comprising a hemocyte lysate, the mixing portion corresponding to the mixing zone of the conduit; and

b. a pump mechanism associated with the housing and fluidly connected to the fluid inlet port, the conduit and the two optical sample wells.

3. The cartridge of claim 2, wherein the base section includes four optical sample wells, wherein two of the four optical sample wells are configured to include an agent representative of the microbial contaminant dried in the optical sample wells.

4. The cartridge of claim 3, wherein the manifold section further includes the agent representative of the microbial contaminant dried on a sample well portion of the manifold section.

5. The cartridge of claim 2, further comprising a chromogenic substrate dried on the mixing portion of the manifold section.

6. The cartridge of claim 2, wherein the pump mechanism is integrated into the base section, and the pump mechanism is a three-position syringe, wherein a first position creates a vacuum to introduce the sample into the aspiration zone, a second position provides transport from the aspiration zone, through the fluid restrictor, to the mixing zone, and a third position provides transport from the mixing zone to the sample wells.

7. The cartridge of claim 2, wherein the hemocyte lysate is limulus amoebocyte lysate.

8. The cartridge of claim 3, wherein the agent is a bacterial endotoxin.

9. A method for detecting the presence of a microbial contaminant in a sample, the method comprising:

a. introducing the sample into the fluid inlet port of the cartridge of claim 3, and transferring the sample to the conduit;

b. transferring the sample into the mixing zone of the conduit;

c. mixing the sample with the hemocyte lysate and the chromogenic substrate to generate a mixed sample;

d. transferring the mixed sample from the mixing zone to the sample well; and

e. measuring an optical property of the mixed sample in the optical sample well, wherein a change in the optical property is indicative of the presence of the microbial contaminant in the sample.

10. The method of claim 9, wherein the measuring the optical property is a change in absorbance of light at a preselected wavelength.

11. The method of claim 10, wherein the change in absorbance of light at a preselected wavelength is compared to a standard curve.

12. The method of claim 11, wherein the standard curve is an archived standard curve.

13. The method of claim 9, wherein the introducing the sample comprises creating a vacuum via the pump mechanism to introduce the sample into the fluid inlet port and the conduit.

14. The method of claim 9, wherein the transferring comprises transporting via the pump mechanism the sample from mixing zone to the optical sample well.

15. A method for detecting the presence of a microbial contaminant in a sample, the method comprising:

a. introducing the sample into the fluid inlet port of the cartridge of claim 2, and transferring the sample to the aspiration zone;

b. transferring the sample from the aspiration zone, through the fluid restrictor, to the mixing zone;

c. mixing the sample with the hemocyte lysate and the chromogenic substrate to generate a mixed sample;

d. transferring the mixed sample to the optical sample wells; and

e. measuring an optical property of the mixed sample in the optical sample wells, wherein a change in the optical property is indicative of the presence of the microbial contaminant in the sample.

16. The method of claim 15, wherein the measuring the optical property is a change in absorbance of light at a preselected wavelength.

17. The method of claim 16, wherein the change in absorbance of light at a preselected wavelength is compared to a standard curve.

18. The method of claim 17, wherein the standard curve is an archived standard curve.

19. The method of claim 15, wherein the introducing the sample comprises creating a vacuum via the pump mechanism to introduce the sample into the fluid inlet port and the aspiration zone, the transferring in b) comprising transporting via the pump mechanism the sample from the aspiration zone to the mixing zone; and the transferring in d) comprises transporting via the pump mechanism the mixed sample from mixing zone to the optical sample wells.