Role Description: RA & QA Referent for the US Market

Role Entails:

- US FDA registration of medical devices and raw materials for pharmaceutics.
- US EPA registration of disinfectants.
- Product classification of imported/exported goods.
- US OEM supplier certifications.
- US OEM product approvals.
- Maintenance and improvement of storage and distribution procedures in the US.
- Fielding US customer complaints.
- Providing regulatory solutions to US customers.
- Providing support on quality and regulation procedures, including updates on laboratory work methods.

Role Requirements:

- BA in life sciences, biotechnology, materials science and engineering.
- Familiarity with biology and microbiology laboratories.
- Experience in EPA and FDA product registration.
- Knowledge and understanding of quality regulations (EPA, 21 CFR part 820, cGMP).
- Fluency in Hebrew and English (additional languages an advantage).
- Proficiency in Microsoft OFFICE applications.
- Intellectual acuity, meticulousness, and broad vision.
- Flexible work hours.
- Residence in Israel, Northern region.
- Full-time employment.
- Car owner.

Reporting to:

- VP Quality Assurance & Regulatory Affairs.

Please send CV to: jobs@bioind.com
For additional information please contact Ms. Anat Elbaz, VP Human Resources 052-2479622