TILLOMED Laboratories Ltd.

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30th June 2011

To whom it may concern

This is to certify that Dr Vandana Jolad was a full-time employee of Tillomed Laboratories Limited at the Department of Regulatory Affairs and Pharmacovigilance from **June 2005 to June 2011** reporting to myself, the undersigned. Overall Dr Jolad has been prompt, efficient with the below duties and achievements and various other tasks assigned to her.

- Preparation and submission of applications, renewals and variations to the relevant authorities in a timely and accurate manner. Follow-up with MHRA/EU regulatory authorities and maintenance of licences. Keep up-to-date with the latest guidelines and legislation.

- Writing of expert non-clinical, quality reports and standard operating procedures. Writing, reviewing and advising on SmPC's, user tested leaflets and packaging.

- Direct negotiations and liaison with third party companies globally and other departments of the company concerning quality and submission applications. Involved actively in audit inspections and key interface for regulatory related QA and technical activities, related agreements and contracts.

- Preparation of CTD dossiers from old format and submissions for MRP/national applications.

- **From 2009 to 30th June 2011** due to business development activities, Dr Jolad has been premised at the Mumbai offices of Maneesh Pharma as Head of Regulatory for Tillomed.

Musharraf M Ginai
Managing Director