

A QUIEN CORRESPONDA

Por medio de la presente inform a ustedes respecto a la documentación solicitada para participar en vuestra licitación, presentamos la información de GMP del fabricante del principio activo obtenido de las bases de datos de U.S. Food and Drug Administration (FDA). En este sitio se indica la información de los sitios de fabricación que se encuentran registrados y auditados por la FDA.

Sin otro particular, le saluda atentamente.

ELISA CARVAJAL GARCIA

Asesor Técnico

ETEX Farmacéutica Ltda

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