

PORTLAND SQUARE PTY LTD

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RS:KS
15th June, 1998

Mr Keith Smith
Manager
Registerable Devices Unit

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Dear Mr Smith,

The Margron hip replacement stem and neck components, Aust L No: 53760
are to be introduced to the European market.

I would be interested in obtaining a CE mark for the prosthesis, and your assistance in this regard
would be appreciated.

The device does not have USA F.D.A. approval, but the prosthesis has been inserted into
thirty-five patients over a fourteen month period and there have been no significant problems or
regulatory action taken against the Margron hip replacement components.

The device does not contain significant new technology and the device is not of human or animal
origin.

Yours faithfully,

RONALD SEKEL