Pharmaceutical Benefits Advisory Committee

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DELETION OF PHARMACEUTICAL BENEFIT ITEMS EFFECTIVE 1 January 2020

Following is the advice of the Pharmaceutical Benefits Advisory Committee concerning the deletion of items from the declaration under subsections 85(2) and 85(2A) of the *National Health Act 1953*, with effect from the above date.

Item

PBAC Advice

Verteporfin

The sponsor has requested the deletion of Visudyne® from the PBS. The sponsor stated that the product will be transferred to them (from Novartis Pharmaceuticals), effective 1 November 2019. After this date, registered supply of the product will cease, as the small volumes of supply are not commercially viable. The product will remain available to the small patient population via unlicensed supply.

The PBAC noted that aflibercept and ranibizumab are PBS-listed for the same indication (subfoveal choroidal neovascularisation (CNV) due to macular degeneration) and there were very few services of verteporfin in 2018-19. The PBAC had no objections to this deletion request.

John Paul Director Health Technology Assessment Section Office of Health Technology Assessment Technology Assessment and Access Division 29 November 2019