

FDA accepts Merck's sBLA for Keytruda six-week dosing schedule for melanoma and multiple other indications

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Merck is seeking FDA approval of a 400 mg Q6W dose infused over 30 minutes for KEYTRUDA indications in melanoma, classical Hodgkin lymphoma, primary mediastinal large B-cell lymphoma,

The post [FDA accepts Merck's sBLA for Keytruda six-week dosing schedule for melanoma and multiple other indications](#) appeared first on [Pharmaceutical Business review](#).