DuPont 2015 Commercialization Plan

May	MDMs continue planning with their SPMs and Notified Body(s)
June – August	MDMs complete their risk assessments, any additional testing and required regulatory notifications (Notified Body for Class III devices, minor change notification in Japan)
July	DuPont completes submission to U.S. FDA, Health Canada and Notified Bodies
August	DuPont publishes 1-Year Real-Time Industry Summary Report
September	**Expected U.S. FDA Affirmation of Functional Equivalence (Submission + 8 weeks)**
	DuPont begins to confirm new orders with Tyvek® Transition material
October	SPMs begin to receive Tyvek® Transition material (U.S. FDA Affirmation + 4 weeks)
December	DuPont expects to transition the majority of manufacturing to Tyvek® Transition material*

*Note: Complete transition may take up to 12-18 months as inventories throughout the value chain are depleted.

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