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CVac™ CAN-003 Clinical Trial Data

- CAN-003 data demonstrate a clinically significant and durable improvement in progression-free survival (PFS) for CVac patients versus standard of care (n=20; ovarian patients in second remission)
- CAN-003 data presented at European Cancer Congress in October, Society for Immunotherapy of Cancer in November, subject of Expert Panel Call in December.
- CVac's demonstrated an excellent tolerability and side effect profile in CAN-003
- Final PFS database for CAN-003 to be analyzed and expect to be presented/published in early 2014.

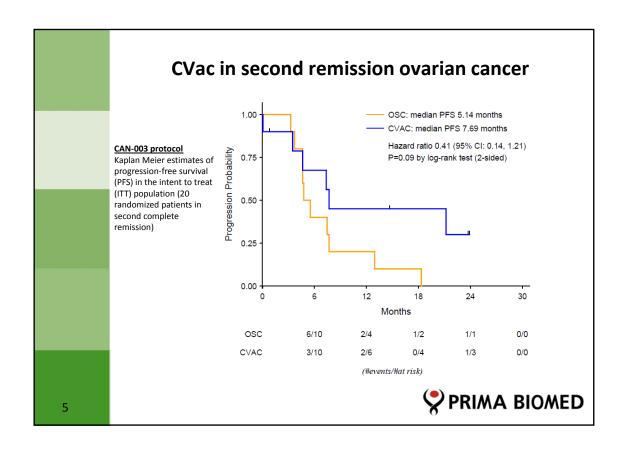


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Exciting path forward...

- Amended CAN-004 phase 2 protocol approved to start in multiple jurisdictions
- CVac[™] phase 2 ovarian cancer trial to start patient recruitment in 2Q 2014
- Important CVac data catalysts in 2014, 2015, 2016 for ovarian and pancreatic cancers
- Cash position A\$28.58 million (31 Dec. 2013) & multimillion Euro grant support in Germany
- Evaluation of value accretive business development transactions
- Leadership position in the emerging field of cancer immunotherapy

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CVac clinical trial design considerations

- Strong PFS signal in second remission epithelial ovarian cancer patient population
- Positive mucin 1-specific T cell response in CVac treated patients
- Importance of Overall Survival as primary endpoint:
 - Based on available reported data in cancer immunotherapy field
 - Endpoint of commercially approved cancer immunotherapies
 - Based on regulatory correspondence

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